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Proposed Guidelines for Carcinogen Risk Assessment

Office of Research and Development
U.S. Environmental Protection Agency
Washington, DC

1 **PROPOSED GUIDELINES FOR CARCINOGEN RISK ASSESSMENT**

2 FRL

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4 **AGENCY:** U.S. Environmental Protection Agency

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6 **ACTION:** Notice of Availability and Opportunity to Comment on Proposed Guidelines for
7 Carcinogen Risk Assessment

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9 **SUMMARY:** The U.S. Environmental Protection Agency (EPA) is today publishing a
10 document entitled *Proposed Guidelines for Carcinogen Risk Assessment* (hereafter "Proposed
11 Guidelines"). These Proposed Guidelines were developed as part of an interoffice guidelines
12 development program by a Technical Panel of the Risk Assessment Forum within EPA's Office
13 of Research and Development. These Proposed Guidelines are a revision of EPA's 1986
14 *Guidelines for Carcinogen Risk Assessment* (hereafter "1986 cancer guidelines") published on
15 September 24, 1986 (51 FR 33992). When final, these guidelines will replace the 1986
16 guidelines.

17 In a future *Federal Register* notice, the Agency intends to publish for comment how it
18 will implement the Proposed Guidelines once they are finalized. The plans will propose and seek
19 comment on how the Guidelines will be used for Agency carcinogen risk assessment and, in
20 particular, will address the impact of the Guidelines on the Agency's existing assessments, and
21 any mechanisms for handling reassessments under finalized Guidelines.

22
23 **DATES:** The Proposed Guidelines are being made available for a 120-day public review and
24 comment period. Comments must be in writing and must be postmarked by [insert date 120
25 days after date of publication in the *Federal Register*]. See Addresses section for guidance on
26 submitting comments.

27
28 **FOR FURTHER INFORMATION, CONTACT:** Technical Information Staff, Operations
29 and Support Group, National Center for Environmental Assessment-Washington

Office, telephone: 202-260-7345. Email inquiries may be sent to
guidelines-cancer@epamail.epa.gov.

ADDRESSES:

The Proposed Guidelines will be made available in the following ways:

1) the electronic version will be accessible on EPA's Office of Research and Development home page on the Internet at <<http://www.epa.gov/ORD/>>

2) 3½" high-density computer diskettes in Wordperfect 5.1 format will be available from ORD Publications, Technology Transfer and Support Division, National Risk Management Research Laboratory, Cincinnati, OH; telephone: 513-569-7562; fax: 513-569-7566. Please provide the EPA No. (EPA/600/P-92/003Ca) when ordering.

3) This notice contains the full draft document. In addition, copies of the draft will be available for inspection at EPA headquarters and regional libraries, through the U.S. Government Depository Library program, and for purchase from the National Technical Information Service (NTIS), Springfield, VA; telephone: 703-487-4650, fax: 703-321-8547. Please provide the NTIS PB No. (PB96-157599) (\$35.00) when ordering.

Submitting Comments

Comments on the Proposed Guidelines may be mailed or delivered to the Technical Information Staff (8623), NCEA-WA/OSG, U.S. Environmental Protection Agency, 401 M Street, S.W., Washington, DC 20460. Comments should be in writing and must be postmarked by the date indicated. Please submit one unbound original with pages numbered consecutively, and three copies. For attachments, provide an index, number pages consecutively with the comment, and submit an unbound original and three copies.

Please note that all technical comments received in response to this notice will be placed in a public record. For that reason, commenters should not submit personal information (such as medical data or home address), Confidential Business Information, or information protection by copyright. Due to limited resources, acknowledgments will not be sent.

SUPPLEMENTARY INFORMATION: In 1983, the National Academy of Sciences (NAS)/National Research Council (NRC) published its report entitled *Risk Assessment in the Federal Government: Managing the Process* (NRC, 1983). In that report, the NRC recommended that Federal regulatory agencies establish "inference guidelines" to ensure consistency and technical quality in risk assessments and to ensure that the risk assessment process was maintained as a scientific effort separate from risk management. The 1986 cancer guidelines were issued on September 24, 1986 (51 FR 33992). The Proposed Guidelines published today continue the guidelines development process. These guidelines set forth principles and procedures to guide EPA scientists in the conduct of Agency cancer risk assessments and to inform Agency decisionmakers and the public about these procedures.

Both the 1986 guidelines and the current proposal contain inference guidance in the form of default inferences to bridge gaps in knowledge and data. Research conducted in the past decade has elucidated much about the nature of carcinogenic processes and continues to provide new information. The intent of this proposal is to take account of knowledge available now and to provide flexibility for the future in assessing data and employing default inferences, recognizing that the guidelines cannot always anticipate future research findings. Because methods and knowledge are expected to change more rapidly than guidelines can practicably be revised, the Agency will update specific assessment procedures with peer-reviewed supplementary, technical documents as needed. Further revision of the guidelines themselves will take place when extensive changes are necessary.

Since 1986, the EPA has sponsored several workshops about revising the cancer guidelines (U.S. EPA, 1989b, 1989c, 1994a). The Society for Risk Analysis conducted a workshop on the subject in connection with its 1992 annual meeting (Anderson et al., 1993). Participants in the most recent workshop in 1994 reviewed an earlier version of the guidelines proposed here and made numerous recommendations about individual issues as well as broad recommendations about explanations and perspectives that should be added. Most recently, the Committee on the Environment and Natural Resources of the Office of Science and Technology Policy reviewed the guidelines at a meeting held on August 15, 1995. The EPA appreciates the efforts of all participants in the process and has tried to address their recommendations in this proposal.

1 In addition, the recommendations of the NRC (1994) in *Science and Judgment in Risk*
2 *Assessment* have been addressed. Responses to these recommendations are given generally in
3 Appendix B as well as being embodied in the Proposed Guidelines. Responses that explain the
4 major default assumptions adopted under these guidelines and the policy for using and departing
5 from these default assumptions appear in Section 1.3.

6 The Science Advisory Board also will review these Proposed Guidelines at a meeting to
7 be announced in a future *Federal Register* notice. Following these reviews Agency staff will
8 prepare summaries of the public and SAB comments. Appropriate comments will be
9 incorporated, and the revised Guidelines will be submitted to the Risk Assessment Forum for
10 review. The Agency will consider comments from the public, the SAB, and the Risk
11 Assessment Forum in its recommendations to the EPA Administrator.

12 13 **Major Changes from the 1986 Guidelines**

14 ***Characterizations***

15
16 Increased emphasis on providing characterization discussions for the hazard, dose
17 response, and exposure sections is part of the proposal. These discussions will summarize the
18 assessments to explain the extent and weight of evidence, major points of interpretation and
19 rationale, and strengths and weaknesses of the evidence and the analysis, and to discuss
20 alternative conclusions and uncertainties that deserve serious consideration (U.S. EPA, 1995).
21 They serve as starting materials for the risk characterization process which completes the risk
22 assessment.

23 24 ***Weighing Evidence of Hazard***

25 A major change is in the way hazard evidence is weighed in reaching conclusions about
26 the human carcinogenic potential of agents. In the 1986 cancer guidelines, tumor findings in
27 animals or humans were the dominant components of decisions. Other information about an
28 agent's properties, its structure-activity relationships to other carcinogenic agents, and its
29 activities in studies of carcinogenic processes was often limited and played only a modulating
30 role as compared with tumor findings. In this proposal, decisions come from considering all of

1 the evidence. This change recognizes the growing sophistication of research methods,
2 particularly in their ability to reveal the modes of action of carcinogenic agents at cellular and
3 subcellular levels as well as toxicokinetic and metabolic processes. The effect of the change on
4 the assessment of individual agents will depend greatly on the availability of new kinds of data
5 on them in keeping with the state of the art. If these new kinds of data are not forthcoming from
6 public and private research on agents, assessments under these guidelines will not differ
7 significantly from assessments under former guidelines.

8 Weighing of the evidence includes addressing the likelihood of human carcinogenic
9 effects of the agent and the conditions under which such effects may be expressed, as these are
10 revealed in the toxicological and other biologically important features of the agent.
11 (Consideration of actual human exposure and risk implications are done separately; they are not
12 parts of the hazard characterization). In this respect, the guidelines incorporate
13 recommendations of the NRC (1994). In that report, the NRC recommends expansion of the
14 former concept of hazard identification, which rests on simply a finding of carcinogenic
15 potential, to a concept of characterization that includes dimensions of the expression of this
16 potential. For example, an agent might be observed to be carcinogenic via inhalation exposure
17 and not via oral exposure, or its carcinogenic activity might be secondary to another toxic effect.
18 In addition, the consideration of evidence includes the mode(s) of action of the agent apparent
19 from the available data as a basis for approaching dose response assessment.

21 ***Classification Descriptors***

22 To express the weight of evidence for carcinogenic hazard potential, the 1986 cancer
23 guidelines provided summary rankings for human and animal cancer studies. These summary
24 rankings were integrated to place the overall evidence in classification groups A through E,
25 Group A being associated with the greatest probability of human carcinogenicity and Group E
26 with evidence of noncarcinogenicity in humans. Data other than tumor findings played a
27 modifying role after initial placement of an agent into a group.

28 These Proposed Guidelines take a different approach, consistent with the change in the
29 basic approach to weighing evidence. No interim classification of tumor findings followed by
30 modifications with other data takes place. Instead, the conclusion reflects the weighing of

evidence in one step. Moreover, standard descriptors of conclusions are employed rather than letter designations, and these are incorporated into a brief narrative description of their informational basis. The narrative with descriptors replaces the previous letter designation. The descriptors are in three categories: "known/likely," "cannot be determined," or "not likely." For instance, using a descriptor in context, a narrative could say that an agent is *likely* to be carcinogenic by inhalation exposure and *not likely* to be carcinogenic by oral exposure. The narrative explains the kinds of evidence available and how they fit together in drawing conclusions, and points out significant issues/strengths/limitations of the data and conclusions. Subdescriptors are used to further refine the conclusion. The narrative also summarizes the mode of action information underlying a recommended approach to dose response assessment.

In considering revision of the former classification method, the Agency has examined other possibilities that would retain the use of letter and number designation of weights of evidence. The use of standard descriptors within a narrative presentation is proposed for three primary reasons. First, the proposed method permits inclusion of explanations of data and of their strengths and limitations. This is more consistent with current policy emphasis on risk characterization. Second, it would take a large set of individual number or letter codes to cover differences in the nature of contributing information (animal, human, other), route of exposure, mode of action, and relative overall weight. When such a set becomes large—10 to 30 codes—it is too large to be a good communication device, because people cannot remember the definitions of the codes so they have to be explained in narrative. Third, it is impossible to predefine the course of cancer research and the kinds of data that may become available. A flexible system is needed to accommodate change in the underlying data and inferences, and a system of codes might become out of date, as has the one in the 1986 cancer guidelines.

Dose Response Assessment

The approach to dose response assessment calls for analysis that follows the conclusions reached in the hazard assessment as to potential mode(s) of action. The assessment begins by analyzing the empirical data in the range of observation. When animal studies are the basis of the analysis, the estimation of a human equivalent dose utilizes toxicokinetic data, if appropriate and adequate data are available. Otherwise, default procedures are applied. For oral dose, the

1 default is to scale daily applied doses experienced for a lifetime in proportion to body weight
2 raised to the 0.75 power. For inhalation dose, the default methodology estimates respiratory
3 deposition of particles and gases and estimates internal doses of gases with different absorption
4 characteristics. These two defaults are a change from the 1986 cancer guidelines which provided
5 a single scaling factor of body weight raised to the 0.66 power. Another change from the 1986
6 guidelines is that response data on effects of the agent on carcinogenic processes are analyzed
7 (nontumor data) in addition to data on tumor incidence. If appropriate, the analyses of data on
8 tumor incidence and on precursor effects may be combined, using precursor data to extend the
9 dose response curve below the tumor data. Even if combining data is not appropriate, study of
10 the dose response for effects believed to be part of the carcinogenic influence of the agent may
11 assist in thinking about the relationship of exposure and response in the range of observation and
12 at exposure levels below the range of observation.

13 Whenever data are sufficient, a biologically based or case-specific dose response model
14 is developed to relate dose and response data in the range of empirical observation. Otherwise,
15 as a standard, default procedure, a model is used to curve-fit the data. The lower 95%
16 confidence limit on a dose associated with an estimated 10% increased tumor or relevant
17 nontumor response (LED_{10}) is identified. This generally serves as the point of departure for
18 extrapolating the relationship to environmental exposure levels of interest when the latter are
19 outside the range of observed data. The environmental exposures of interest may be measured
20 ones or levels of risk management interest in considering potential exposure control options.
21 Other points of departure may be more appropriate for certain data sets; as described in the
22 guidance, these may be used instead of the LED_{10} . Additionally, the LED_{10} is available for
23 comparison with parallel analyses of other carcinogenic agents or of noncancer effects of agents
24 and for gauging and explaining the magnitude of subsequent extrapolation to low-dose levels.
25 The LED_{10} , rather than the ED_{10} (the estimate of a 10% increased response), is the proposed
26 standard point of departure for two reasons. One is to permit easier comparison with the
27 benchmark dose procedure for noncancer health assessment—also based on the lower limit on
28 dose. Another is that the lower limit, as opposed to the central estimate, accounts for uncertainty
29 in the experimental data. The issue of using a lower limit or central estimate was discussed at a
30 workshop held on the benchmark procedure for noncancer assessment (Barnes et al., 1995) and

1 at a workshop on a previous version of this proposal (U.S. EPA, 1994b). The latter workshop
2 recommended a central estimate; the benchmark workshop recommended a lower limit.

3 The second step of dose response assessment is extrapolation to lower dose levels, if
4 needed. This is based on a biologically based or case-specific model if supportable by
5 substantial data. Otherwise, default approaches are applied that accord with the view of mode(s)
6 of action of the agent. These include approaches that assume linearity or nonlinearity of the
7 dose response relationship or both. The default approach for linearity is to extend a straight line
8 to zero dose, zero response. The default approach for nonlinearity is to use a margin of exposure
9 analysis rather than estimating the probability of effects at low doses. A margin of exposure
10 analysis explains the biological considerations for comparing the observed data with the
11 environmental exposure levels of interest and helps in deciding on an acceptable level of
12 exposure in accordance with applicable management factors.

13 The use of straight line extrapolation for a linear default is a change from the 1986
14 guidelines which used the "linearized multistage" (LMS) procedure. This change is made
15 because the former modeling procedure gave an appearance of specific knowledge and
16 sophistication unwarranted for a default. The proposed approach is also more like that employed
17 by the Food and Drug Administration (U.S. FDA, 1987). The numerical results of the straight
18 line and LMS procedures are not significantly different (Krewski et al., 1984). The use of a
19 margin of exposure approach is included as a new default procedure to accommodate cases in
20 which there is sufficient evidence of a nonlinear dose response, but not enough evidence to
21 construct a mathematical model for the relationship. (The Agency will continue to seek a
22 modeling method to apply in these cases. If a modeling approach is developed, it will be subject
23 to peer review and public notice in the context of a supplementary document for these
24 guidelines.)

25 The public is invited to provide comments to be considered in EPA decisions about the
26 content of the final guidelines. After the public comment period, the EPA Science Advisory
27 Board will be asked to review and provide advice on the guidelines and issues raised in
28 comments. EPA asks those who respond to this notice to include their views on the following:
29

(1) The proposed guidance for characterization of hazard, including the weight of evidence descriptors and weight of evidence narrative which are major features of the proposal. There are three categories of descriptors: "known/likely," "cannot be determined," and "not likely" which are further refined by subdescriptors. It is felt that these three descriptors will satisfactorily delineate the types of evidence bearing on carcinogenicity as they are used with subdescriptors in the context of a narrative of data and rationale. However, an issue that has been discussed by external peer reviewers and by EPA staff is whether the descriptor-subdescriptor called "cannot be determined—suggestive evidence" should become a separate, fourth category called "suggestive." The EPA may choose this course in the final guidelines and requests comment. In considering this issue, commenters may wish to refer not only to Sections 2.6.2. and 2.7.2. which cover the descriptors and narrative, but also to case study example #6 in Section 2.6.3. and example narrative #2 in Appendix A of the proposal. EPA asks commenters on this question to address the rationale (science as well as policy) for leaving the categories of descriptors as proposed or making the fourth category. How might the coverage of a "suggestive" category be defined in order to be most useful?

(2) The use of mode of action information in hazard characterization and to guide dose response assessment is a central part of the proposed approach to bringing new research on carcinogenic processes to bear in assessments of environmental agents (Sections 1.3.2., 2.3.2., 2.5., 3.1.). The appropriate use of this information now and in the future is important. EPA requests comment on the treatment of such information in the proposal, including reliance on peer review as a part of the judgmental process on its application.

(3) Uses of nontumor data in the dose response assessment and the methodological and science policy issues posed are new to these guidelines (Sections 1.3.2., 3.1.2.). EPA requests comment on both issues.

(4) Dose response assessment is proposed to be considered in two parts—range of observed data and range of extrapolation (Section 3.1.). The lower 95% confidence limit on a dose associated with a 10% response (tumor or nontumor response) is proposed as a default

1 point of departure, marking the beginning of extrapolation. This is a parallel to the benchmark
2 procedure for evaluating dose-response of noncancer health endpoints (Barnes et al.,1995). An
3 alternative is to use the central estimate of a 10% response. Another alternative is to use a 1%,
4 instead of a 10%, response when the observed data are tumor incidence data. Does the generally
5 larger sample size of tumor effect studies support using a 1% response as compared with using
6 10% for smaller studies? Are there other approaches for the point of departure that might be
7 considered?

8
9 (5) Discussions of default assumptions and other responses to the 1994 NRC report
10 *Science and Judgment in Risk Assessment* appear in Section 1.3.1. and Appendix B of the
11 proposal, respectively. Comments are requested on responses to the NRC recommendations and
12 how the guidelines as a whole address them.

13
14 _____
15 Date

16 _____
17 Carol M. Browner
Administrator

CONTENTS

LIST OF FIGURES	14
1. INTRODUCTION	15
1.1. PURPOSE AND SCOPE OF THE GUIDELINES	15
1.2. ORGANIZATION AND APPLICATION OF THE GUIDELINES	15
1.2.1. Organization	15
1.2.2. Application	16
1.3. USE OF DEFAULT ASSUMPTIONS	17
1.3.1. Default Assumptions	18
1.3.2. Major Defaults	24
1.3.2.1. <i>Is the Presence or Absence of Effects Observed in a Human Population Predictive of Effects in Another Exposed Human Population?</i>	25
1.3.2.2. <i>Is the Presence or Absence of Effects Observed in an Animal Population Predictive of Effects in Exposed Humans?</i>	26
1.3.2.3. <i>How Do Metabolic Pathways Relate Across Species?</i>	29
1.3.2.4. <i>How Do Toxicokinetic Processes Relate Across Species?</i>	29
1.3.2.5. <i>What Is the Correlation of the Observed Dose Response Relationship to the Relationship at Lower Doses?</i>	30
1.4. CHARACTERIZATIONS	35
2. HAZARD ASSESSMENT	38
2.1. OVERVIEW OF HAZARD ASSESSMENT AND CHARACTERIZATION	38
2.1.1. Analyses of Data	38
2.1.2. Cross-Cutting Topics for Data Integration	38
2.1.2.1. <i>Conditions of Expression</i>	38
2.1.2.2. <i>Mode of Action</i>	39
2.1.3. Presentation of Results	39
2.2. ANALYSIS OF TUMOR DATA	40
2.2.1. Human Data	40
2.2.1.1. <i>Types of Studies</i>	41
2.2.1.2. <i>Criteria for Assessing Adequacy of Epidemiologic Studies</i>	42
2.2.1.3. <i>Criteria for Causality</i>	46
2.2.1.4. <i>Assessment of Evidence of Carcinogenicity from Human Data</i> .	47
2.2.2. Animal Data	48
2.2.2.1. <i>Long-Term Carcinogenicity Studies</i>	48
2.2.2.2. <i>Other Studies</i>	54
2.2.3. Structural Analogue Data	55
2.3. ANALYSIS OF OTHER KEY DATA	56
2.3.1. Physicochemical Properties	56

CONTENTS (continued)

1	2.3.2. Structure-Activity Relationships	56
2	2.3.3. Comparative Metabolism and Toxicokinetics	57
3	2.3.4. Toxicological and Clinical Findings	59
4	2.3.5. Mode of Action-Related Endpoints and Short-Term Tests	60
5	2.3.5.1. <i>Direct DNA Effects</i>	60
6	2.3.5.2. <i>Secondary DNA Effects</i>	60
7	2.3.5.3. <i>Nonmutagenic and Other Effects</i>	61
8	2.3.5.4. <i>Criteria for Judging Mode of Action</i>	62
9	2.4. BIOMARKER INFORMATION	62
10	2.5. MODE OF ACTION--IMPLICATIONS FOR HAZARD	
11	CHARACTERIZATION AND DOSE RESPONSE	64
12	2.6. WEIGHT OF EVIDENCE EVALUATION FOR POTENTIAL HUMAN	
13	CARCINOGENICITY	68
14	2.6.1. Weight of Evidence Analysis	68
15	2.6.2. Descriptors for Classifying Weight of Evidence	83
16	2.6.3. Case Study Examples	85
17	2.7. PRESENTATION OF RESULTS	105
18	2.7.1. Technical Hazard Characterization	105
19	2.7.2. Weight of Evidence Narrative	107
20		
21	3. DOSE RESPONSE ASSESSMENT	110
22	3.1. DOSE RESPONSE RELATIONSHIP	110
23	3.1.1. Analysis in the Range of Observation	111
24	3.1.2. Analysis in the Range of Extrapolation	113
25	3.1.3. Use of Toxicity Equivalence Factors and Relative Potency Estimates	118
26	3.2. RESPONSE DATA	119
27	3.3. DOSE DATA	121
28	3.3.1. Interspecies Adjustment of Dose	121
29	3.3.2. Toxicokinetic Analyses	122
30	3.3.3. Route-to-Route Extrapolation	123
31	3.3.4. Dose Averaging	125
32	3.4. DISCUSSION OF UNCERTAINTIES	125
33	3.5. TECHNICAL DOSE RESPONSE CHARACTERIZATION	126
34		
35	4. TECHNICAL EXPOSURE CHARACTERIZATION	129
36		
37	5. RISK CHARACTERIZATION	130
38	5.1. PURPOSE	130
39	5.2. APPLICATION	131
40	5.3. PRESENTATION OF RISK CHARACTERIZATION SUMMARY	132
41	5.4. CONTENT OF RISK CHARACTERIZATION SUMMARY	132
42		

CONTENTS (continued)

1	APPENDIX A	134
2		
3	APPENDIX B	147
4		
5	APPENDIX C	154
6		
7	REFERENCES	159

LIST OF FIGURES

Figure 1-1.	Decisions on Dose Response Assessment Approaches for the Range of Extrapolation	34
Figure 1-2.	Risk Characterization	37
Figure 2-1.	Factors for Weighing Human Evidence	71
Figure 2-2.	Factors for Weighing Animal Evidence	75
Figure 2-3.	Factors for Weighing Other Key Evidence	78
Figure 2-4.	Factors for Weighing Totality of Evidence	81
Figure 3-1.	Graphical Presentation of Data and Extrapolations	117

1. INTRODUCTION

1.1. PURPOSE AND SCOPE OF THE GUIDELINES

These guidelines revise and replace United States Environmental Protection Agency (EPA) Guidelines for Carcinogen Risk Assessment published in 51 FR 33992, September 24, 1986. The guidelines provide EPA staff and decisionmakers with guidance and perspectives to develop and use risk assessments. They also provide basic information to the public about the Agency's risk assessment methods.

The guidelines encourage both regularity in procedures to support consistency in scientific components of Agency decisionmaking and innovation to remain up-to-date in scientific thinking. In balancing these goals, the Agency relies on input from the general scientific community through established scientific peer review processes. The guidelines incorporate basic principles and science policies based on evaluation of the currently available information. As more is discovered about carcinogenesis, the need will arise to make appropriate changes in risk assessment guidance. The Agency will revise these guidelines when extensive changes are due. In the interim, the Agency will issue special reports, after appropriate peer review, to supplement and update guidance on single topics, (e.g., U.S. EPA, 1991b)

1.2. ORGANIZATION AND APPLICATION OF THE GUIDELINES

1.2.1. Organization

Publications of the Office of Science and Technology Policy (OSTP, 1985) and the National Research Council (NRC, 1983, 1994) provide information and general principles about risk assessment. Risk assessment uses available scientific information on the properties of an agent¹ and its effects in biological systems to provide an evaluation of the potential for harm as a consequence of environmental exposure to the agent. Risk assessment is one of the scientific analyses available for consideration, with other analyses, in decisionmaking on environmental

The term "agent" refers generally to any chemical substance, mixture, or physical or biological entity being assessed, unless otherwise noted.

1 protection. The 1983 and 1994 NRC documents organize risk assessment information into four
2 areas: hazard identification, dose response assessment, exposure assessment, and risk
3 characterization. This structure appears in these guidelines, which additionally emphasize
4 characterization of evidence and conclusions in each part of the assessment. In particular, the
5 guidelines adopt the approach of the NRC's 1994 report in adding a dimension of
6 characterization to the hazard identification step. Added to the identification of hazard is an
7 evaluation of the conditions under which its expression is anticipated. The risk assessment
8 questions addressed in these guidelines are:

9 C For hazard--Can the agent present a carcinogenic hazard to humans, and if so, under
10 what circumstances?

11 C For dose response--At what levels of exposure might effects occur?

12 C For exposure--What are the conditions of human exposure?

13 C For risk--What is the character of the risk? How well do data support conclusions
14 about the nature and extent of the risk?

16 **1.2.2. Application**

17 The guidelines apply within the framework of policies provided by applicable EPA
18 statutes and do not alter such policies. The guidelines cover assessment of available data. They
19 do not imply that one kind of data or another is prerequisite for regulatory action concerning any
20 agent. Risk management applies directives of regulatory legislation, which may require
21 consideration of potential risk, or solely hazard or exposure potential, along with social,
22 economic, technical, and other factors in decisionmaking. Risk assessments support decisions,
23 but to maintain their integrity as decisionmaking tools, they are not influenced by consideration
24 of the social or economic consequences of regulatory action.

25 Not every EPA assessment has the same scope or depth. Agency staff often conduct
26 screening-level assessments for priority-setting or separate assessments of hazard or exposure for
27 ranking purposes or to decide whether to invest resources in collecting data for a full assessment.
28 Moreover, a given assessment of hazard and dose response may be used with more than one
29 exposure assessment that may be conducted separately and at different times as the need arises in
30 studying environmental problems in various media. The guidelines apply to these various

1 situations in appropriate detail given the scope and depth of the particular assessment. For
2 example, a screening assessment may be based almost entirely on structure-activity relationships
3 and default assumptions. As more data become available, assessments can replace or modify
4 default assumptions accordingly. These guidelines do not require that all of the kinds of data
5 covered here be available for either assessment or decisionmaking. The level of detail of an
6 assessment is a matter of Agency management policy regarding the applicable decisionmaking
7 framework.

8 9 **1.3. USE OF DEFAULT ASSUMPTIONS**

10 The National Research Council, in its 1983 report on the science of risk assessment
11 (NRC, 1983), recognized that default assumptions are necessarily made in risk assessments
12 where gaps exist in general knowledge or in available data for a particular agent . These default
13 assumptions are inferences based on general scientific knowledge of the phenomena in question
14 and are also matters of policy concerning the appropriate way to bridge uncertainties that
15 concern potential risk to human health (or, more generally, to environmental systems) from the
16 agent under assessment.

17 EPA's 1986 guidelines for cancer risk assessment (EPA, 1986) were developed in
18 response to the 1983 NRC report. The guidelines contained a number of default assumptions.
19 They also encouraged research and analysis that would lead to new risk assessment methods and
20 data and anticipated that these would replace defaults. The 1986 guidelines did not explicitly
21 discuss how to depart from defaults. In practice, the agency's assessments routinely have
22 employed defaults and, until recently, only occasionally departed from them.

23 In its 1994 report on risk assessment, the NRC supported continued use of default
24 assumptions (NRC, 1994). The NRC report thus validated a central premise of the approach to
25 risk assessment that EPA had evolved in preceding years--the making of science policy
26 inferences to bridge gaps in knowledge--while at the same time recommending that EPA
27 develop more systematic and transparent guidelines to inform the public of the default inferences
28 EPA uses in practice. It recommended that the EPA review and update the 1986 guidelines in
29 light of evolving scientific information and experience in practice in applying those guidelines,
30 and that the EPA explain the science and policy considerations underlying current views as to

the appropriate defaults and provide general criteria to guide preparers and reviewers of risks assessments in deciding when to depart from a default . Pursuant to this recommendation, the following discussion presents descriptions of the major defaults and their rationales. In addition, it presents general policy guidance on using and departing from defaults in specific risk assessments.

1.3.1. Default Assumptions

The 1994 NRC report contains several recommendations regarding flexibility and the use of default options:

- Ⓒ EPA should continue to regard the use of default options as a reasonable way to deal with uncertainty about underlying mechanisms in selecting methods and models for use in risk assessment.
- Ⓒ EPA should explicitly identify each use of a default option in risk assessments.
- Ⓒ EPA should clearly state the scientific and policy basis for each default option.
- Ⓒ The Agency should consider attempting to give greater formality to its criteria for a departure from default options in order to give greater guidance to the public and to lessen the possibility of ad hoc, undocumented departures from default options that would undercut the scientific credibility of the Agency's risk assessments. At the same time, the Agency should be aware of the undesirability of having its guidelines evolve into inflexible rules.
- Ⓒ EPA should continue to use the Science Advisory Board and other expert bodies. In particular, the Agency should continue to make the greatest possible use of peer review, workshops, and other devices to ensure broad peer and scientific participation to guarantee that its risk assessment decisions will be based on the best science available through a process that allows full public discussion and peer participation by the scientific community.

In the 1983 report (p. 28), NAS defined the use of "inference options" (default options) as a means to bridge inherent uncertainties in risk assessment. These options exist when the assessment encounters either "missing or ambiguous information on a particular substance" or "gaps in current scientific theory." Since there is no instance in which a set of data on an agent

1 or exposure is complete, all risk assessments must use general knowledge and policy guidance to
2 bridge data gaps. Animal toxicity data are used, for example, to substitute for human data
3 because we do not test human beings. The report described the components of risk assessment
4 in terms of questions encountered during analysis for which inferences must be made. The
5 report noted (p. 36) that many components ". . . lack definitive scientific answers, that the degree
6 of scientific consensus concerning the best answer varies (some are more controversial than
7 others), and that the inference options available for each component differ in their degree of
8 conservatism. The choices encountered in risk assessment rest, to various degrees, on a mixture
9 of scientific fact and consensus, on informed scientific judgment, and on policy determinations
10 (the appropriate degree of conservatism). . . ." The report did not note that the mix varies
11 significantly from case to case. For instance, a question that arises in hazard identification is
12 how to use experimental animal data when the routes of exposure differ between animals and
13 humans. A spectrum of inferences could be made, ranging from the most conservative, or risk
14 adverse one that effects in animals from one route may be seen in humans by another route, to an
15 intermediate, conditional inference that such translation of effects will be assumed if the agent is
16 absorbed by humans through the second route, to a nonconservative view that no inference is
17 possible and the agent's effects in animals must be tested by the second route. The choice of an
18 inference, as the report observed, comes from more than scientific thinking alone. While the
19 report focused mainly on the idea of conservatism of public health as a science policy rationale
20 for making the choice, it did not evaluate other considerations. These include such things as the
21 matters of time and resources and whether the analysis is for an important decision required to
22 be made soon or is simply a screening or ranking effort. For a screening analysis, one might
23 make several "worst case" inferences to determine if, even under those conditions, risk is low
24 enough that a problem can be eliminated from further consideration. In the above discussion
25 concerning inferences about route-to-route extrapolation, one might use the most conservative
26 one for screening.

27 These revised guidelines retain the use of default assumptions as recommended in the
28 1994 report. Generally, these defaults remain public health conservative, but in some instances,
29 they have been modified to reflect the evolution of scientific knowledge since 1986.

1 In addition, the guidelines reflect evaluation of experience in practice in applying
2 defaults and departing from them in individual risk assessments conducted under the 1986
3 guidelines. The application and departure from defaults and the principles to be used in these
4 judgments have been matters of debate among practitioners and reviewers of risk assessments.
5 Some observers believe that in practice EPA risk assessors have been too resistant to considering
6 departures; others question whether proposed departures have been adequately supported. Some
7 cases in which departures have been considered have been generally accepted, while others have
8 been controversial. The guidelines here are intended to be both explicit and more flexible than
9 in the past concerning the basis for making departures from defaults, recognizing that expert
10 judgment and peer review are essential elements of the process.

11 In response to the recommendations of the 1994 report, these guidelines call for
12 identification of the default assumptions used within assessments and for highlighting significant
13 issues about defaults within characterization summaries of component analyses in assessment
14 documents. As to the use of peer review to aid in making judgments about applying or departing
15 from defaults, we agree with the NRC recommendation. The Agency has long made use of
16 workshops, peer review of documents and guidelines, and consultations as well as formal peer
17 review by the Science Advisory Board (SAB). In 1994, the Administrator of EPA published
18 formal guidance for peer review of EPA scientific work products that increases the amount of
19 peer review for risk assessments as well as other work, as a response to the NRC report and to
20 SAB recommendations (U.S. EPA, 1994b).

21 The 1994 NRC report recommended that EPA should consider adopting principles or
22 criteria that would give greater formality and transparency to decisions to depart from defaults.
23 The report named several possible criteria for such principles (p. 7): ". . . [P]rotecting the public
24 health, ensuring scientific validity, minimizing serious errors in estimating risks, maximizing
25 incentives for research, creating an orderly and predictable process, and fostering openness and
26 trustworthiness. There might be additional relevant criteria. . . ." The report indicated, however,
27 that the committee members had not reached consensus on a single criterion to address the key
28 issue of how much certainty or proof a risk assessor must have in order to justify departing from
29 a default. Appendix N of the report contains two presentations of alternative views held by
30 some committee members on this issue. One view, known as "plausible conservatism,"

1 suggested that departures from defaults should not be made unless new information improves the
2 understanding of a biological process to the point that relevant experts reach consensus that the
3 conservative default assumption concerning that process is no longer plausible. The same
4 criterion was recommended where the underlying scientific mechanism is well understood, but
5 where a default is used to address missing data. In this case, the default should not be replaced
6 with case-specific data unless it is the consensus of relevant experts that the proffered data make
7 the default assumption no longer plausible. Another view, known as the "maximum use of
8 scientific information" approach, acknowledged that the initial choice of defaults should be
9 conservative but argued that conservatism should not be a factor in determining whether to
10 depart from the default in favor of an alternate biological theory or alternate data. According to
11 this view, it should not be necessary to reach expert consensus that the default assumption had
12 been rendered implausible; it should be sufficient that risk assessors find the alternate approach
13 more plausible than the default.

14 The EPA is not adopting a list of formal decision criteria in the sense of a checklist based
15 on either view. It would not be helpful to generate a checklist of uniform criteria for several
16 reasons. First, risk assessments are highly variable in content and purpose. Screening
17 assessments may be purposely "worst case" in their default assumptions to eliminate problems
18 from further investigation. Subsequent risk assessments based on a fuller data set can discard
19 worst-case default assumptions in favor of plausibly conservative assumptions and progressively
20 replace or modify the latter with data. No uniform checklist will fit all cases. Second, a
21 checklist would likely become more a source of rote discussion than of enlightenment about the
22 process.

23 Instead, these guidelines use a combination of principles and process in the application of
24 and departure from default assumptions. The guidelines provide a framework of default
25 assumptions to allow risk assessment to proceed when current scientific theory or available case-
26 specific data do not provide firm answers in a particular case, as the 1983 report outlined. Some
27 of the default assumptions bridge large gaps in fundamental knowledge which will be filled by
28 basic research on the causes of cancer and on other biological processes, rather than by agent-
29 specific testing. Other default assumptions bridge smaller data gaps that can feasibly be filled

1 for a single agent, such as whether a metabolic pathway in test animals is like (default) or unlike
2 that in humans.

3 The decision to use a default, or not, is a choice considering available information on an
4 underlying scientific process and agent-specific data, depending on which kind of default it is.
5 Generally, if a gap in basic understanding exists, or if agent-specific data are missing, the default
6 is used without pause. If data are present, their evaluation may reveal inadequacies that also lead
7 to use of the default. If data support a plausible alternative to the default, but no more strongly
8 than they support the default, both the default and its alternative are carried through the
9 assessment and characterized for the risk manager. If data support an alternative to the default
10 as the more reasonable judgment, the data are used. (This framework of choices is not wholly
11 applicable to screening assessments. As mentioned above, screening assessments may
12 appropriately use "worst case" inferences to determine if, even under those conditions, risk is
13 low enough that a problem can be eliminated from further consideration.)

14 Scientific peer review, peer consultative workshops and similar processes are the
15 principal ways determining the strength of thinking and generally accepted views within the
16 scientific community about the application of and departure from defaults and about judgments
17 concerning the plausibility and persuasiveness of data in a particular case. The choices made are
18 explicitly discussed in the assessment, and if a particular choice raises a significant issue, it is
19 highlighted in the risk characterization.

20 The discussion of major defaults in these guidelines together with the explicit discussion
21 of the choice of inferences within the assessment and the processes of peer review and peer
22 consultation will serve the several goals stated in the 1994 report. One is to encourage research,
23 since results of research efforts will be considered. Another is to allow timely decisionmaking,
24 when time is a constraint, by supporting completion of the risk assessment using defaults as
25 needed. Another is to be flexible, using new science as it develops. Finally, the use of public
26 processes of peer consultation and peer review will ensure that discipline of thought is
27 maintained to support trust in assessment results.

28 Experience has shown that the most difficult part of the framework of choices is the
29 judgment of whether a data analysis is both biologically plausible and persuasive as applied to
30 the case at hand. There is no set of rules for making this judgment in all cases. Two criteria that

1 apply in these guidelines are that the underlying scientific principle has been generally accepted
2 within the scientific community and that supportive experiments are available that test the
3 application of the principle to the agent under review. For example, mutagenicity through
4 reactivity with DNA has been generally accepted as a carcinogenic influence for many years.
5 This acceptance, together with evidence of such mutagenicity in experiments on an agent,
6 provides plausible and persuasive support for the inference that mutagenicity is a mode of action
7 for the agent.

8 Judgments about plausibility and persuasiveness of analyses vary according to the
9 scientific nature of the default. An analysis of data may replace a default or modify it. An
10 illustration of the former is development of EPA science policy on the issue of the relevance for
11 humans of male rat kidney neoplasia involving alpha 2u globulin (U.S. EPA, 1991b). The 1991
12 EPA policy gives guidance on the kind of experimental findings that demonstrate whether the
13 alpha 2u globulin mechanism is present and responsible for carcinogenicity in a particular case.
14 Before this policy guidance was issued, the default assumption was that neoplasia in question
15 was relevant to humans and indicated the potential for hazard to humans. A substantial body of
16 data was developed by public and private research groups as a foundation for the view that the
17 alpha 2u globulin-induced response was not relevant to humans. These studies first addressed
18 the alpha 2u globulin mechanism in the rat and whether this mechanism has a counterpart in the
19 human being, both were large research efforts. The resulting data presented difficulties; some
20 reviewers were concerned that the mechanism in the rat appeared to be understood only in
21 outline, not in detail, and felt that the data were insufficient to show the lack of a counterpart
22 mechanism in humans. It was particularly difficult to support a negative such as the
23 nonexistence of a mechanism in humans because so little is known about what the mechanisms
24 are in humans. Despite these concerns, in its 1991 policy guidance, EPA concluded that the
25 alpha 2u globulin-induced response in rats should be regarded as not relevant to humans (i.e., as
26 not indicating human hazard).

27 One lesson in the development and peer review of this policy is that if the default
28 concerns an inherently complex biological question, large amounts of work will be required to
29 replace the default. A second is that addressing a negative is difficult. A third is that "proof" in
30 the strict sense of having laid all reasonable doubt to rest is not required. Instead, an alternative

1 may displace a default when it is generally accepted in peer review as the most reasonable
2 judgment. The issue of relevance may not always be so difficult. It would be an experimentally
3 easier task, for example, to determine whether carcinogenesis in an animal species is due to a
4 metabolite of the agent in question that is not produced in humans.

5 When scientific processes are understood but case-specific data are missing, defaults can
6 be constructed to be modified by experimental data, even if data do not suffice to replace them
7 entirely. For example, the approaches adopted in these guidelines for scaling dose from
8 experimental animals to humans are constructed to be either modified or replaced as data
9 become available on toxicokinetic parameters for the particular agent being assessed. Similarly,
10 the selection of an approach or approaches for dose response assessment is based on a series of
11 decisions that consider the nature and adequacy of available data in choosing among alternative
12 modeling and default approaches.

13 The 1994 NRC report notes (p. 6) that "[a]s scientific knowledge increases, the science
14 policy choices made by the Agency and Congress should have less impact on regulatory
15 decisionmaking. Better data and increased understanding of biological mechanisms should
16 enable risk assessments that are less dependent on conservative default assumptions and more
17 accurate as predictions of human risk." Undoubtedly, this is the trend as scientific understanding
18 increases. However, some gaps in knowledge and data will doubtless continue to be encountered
19 in assessment of even data-rich cases, and it will remain necessary for risk assessments to
20 continue using defaults within the framework set forth here.

21 22 **1.3.2. Major Defaults**

23 This discussion covers the major default assumptions commonly employed in a cancer
24 risk assessment and adopted in these guidelines. They are predominantly inferences necessary to
25 use data observed under empirical conditions to estimate events and outcomes under
26 environmental conditions. Several inferential issues arise when effects seen in a subpopulation
27 of humans or animals are used to qualitatively infer potential effects in the population of
28 environmentally exposed humans. Several more inferential issues arise in extrapolating the
29 exposure-effect relationship observed empirically to lower-exposure environmental conditions.

The following issues cover the major default areas. Typically, an issue has some subissues; they are introduced here, but are discussed in greater detail in subsequent sections.

- Ⓒ Is the presence or absence of effects observed in a human population predictive of effects in another exposed human population?
- Ⓒ Is the presence or absence of effects observed in an animal population predictive of effects in exposed humans?
- Ⓒ How do metabolic pathways relate across species?
- Ⓒ How do toxicokinetic processes relate across species?
- Ⓒ What is the correlation of the observed dose response relationship to the relationship at lower doses?

1.3.2.1. Is the Presence or Absence of Effects Observed in a Human Population Predictive of Effects in Another Exposed Human Population?

When cancer effects in exposed humans are attributed to exposure to an exogenous agent, the default assumption is that such data are predictive of cancer in any other exposed human population. Studies either attributing cancer effects in humans to exogenous agents or reporting no effects are often studies of occupationally exposed humans. By sex, age, and general health, workers are not representative of the general population exposed environmentally to the same agents. In such studies there is no opportunity to observe whether infants and children, males, or females who are under represented in the study, or people whose health is not good, would respond differently. Therefore, it is understood that this assumption could still underestimate the response of certain sensitive human subpopulations, i.e. biologically vulnerable parts of the population may be left out of risk assessments (NRC, 1993a, 1994). Consequently, this is a default that does not err on the side of public health conservatism, as the 1994 NRC report also recognizes.

On the one hand, if effects are seen in a worker population, this may be in fact indicative of heightened effects in sensitive subpopulations. There is not enough knowledge yet to form a basis for any generally applicable, qualitative inference to compensate for this knowledge gap. In these guidelines, this problem is left to case-by-case analysis, to be attended to as future research and information on particular agents allow. When information on a sensitive

subpopulation exists, it will be used. The topic of variability is addressed further in the discussion of quantitative default assumptions about dose response relationships below. On the other hand, *when cancer effects are not found in an exposed human population, this information by itself is not generally sufficient to conclude that the agent poses no carcinogenic hazard to this or other populations of potentially exposed humans.* This is because epidemiologic studies usually have low power to detect and attribute responses (section 2.2.1.). This may be particularly true when extrapolating null results from a healthy, worker population to other potentially sensitive exposed humans. Again, the problem is left to case-by-case analysis.

1.3.2.2. *Is the Presence or Absence of Effects Observed in an Animal Population Predictive of Effects in Exposed Humans?*

The default assumption is that positive effects in animal cancer studies indicate that the agent under study can have carcinogenic potential in humans. Thus, if no adequate human data are present, positive effects in animal cancer studies are a basis for assessing the carcinogenic hazard to humans. This assumption is a public health conservative policy, and it is both appropriate and necessary given that we do not test for carcinogenicity in humans. The assumption is supported by the fact that nearly all of the agents known to cause cancer in humans are carcinogenic in animals in tests with adequate protocols (IARC, 1994; Tomatis et al., 1989; Huff, 1994). Moreover, almost one-third of human carcinogens were identified subsequent to animal testing (Huff, 1993). Further support is provided by research on the molecular biology of cancer processes, which has shown that the mechanisms of control of cell growth and differentiation are remarkably homologous among species and highly conserved in evolution. Nevertheless, the same research tools that have enabled recognition of the nature and commonality of cancer processes at the molecular level also have the power to reveal differences and instances in which animal responses are not relevant to humans (Linjinsky, 1993; U.S. EPA, 1991b). Under these guidelines, available mode of action information is studied for its implications in both hazard and dose response assessment and its effect on default assumptions.

There may be instances in which the use of an animal model would identify a hazard in animals that is not truly a hazard in humans (e.g., the alpha-2u-globulin association with renal neoplasia in male rats (U.S. EPA, 1991b)). The extent to which animal studies may yield false

1 positive indications for humans is a matter of scientific debate. To demonstrate that a response
2 in animals is not relevant to any human situation, adequate data to assess the relevancy issue
3 must be available.

4 Animal studies are conducted at high doses in order to provide statistical power, the
5 highest dose being one that is minimally toxic (maximum tolerated dose). Consequently, the
6 question often arises whether a carcinogenic effect at the highest dose may be a consequence of
7 cell killing with compensatory cell replication or of general physiological disruption, rather than
8 inherent carcinogenicity of the tested agent. There is little doubt that this may happen in some
9 cases, but skepticism exists among some scientists that it is a pervasive problem (Ames and
10 Gold, 1990; Melnick et al., 1993a; Melnick et al., 1993b; Barrett, 1993). In light of this
11 question, *the default assumption is that effects seen at the highest dose tested are appropriate for*
12 *assessment, but it is necessary that the experimental conditions be scrutinized.* If adequate data
13 demonstrate that the effects are solely the result of excessive toxicity rather than carcinogenicity
14 of the tested agent per se, then the effects may be regarded as not appropriate to include in
15 assessment of the potential for human carcinogenicity of the agent. This is a matter of expert
16 judgment, considering all of the data available about the agent including effects in other toxicity
17 studies, structure-activity relationships, and effects on growth control and differentiation.

18 *When cancer effects are not found in well-conducted animal cancer studies in two or*
19 *more appropriate species and other information does not support the carcinogenic potential of*
20 *the agent, these data provide a basis for concluding that the agent is not likely to possess human*
21 *carcinogenic potential, in the absence of human data to the contrary.* This default assumption
22 about lack of cancer effects is not public health conservative. For instance, the tested animal
23 species may not be predictive of effects in humans; arsenic shows only minimal or no effect in
24 animals, while it is clearly positive in humans. (Other information, such as absence of mutagenic
25 activity or absence of carcinogenic activity among structural analogues, can increase the
26 confidence that negative results in animal studies indicate a lack of human hazard.) Also, it is
27 recognized that animal studies (and epidemiologic studies as well) have very low power to detect
28 cancer effects. Detection of a 10% tumor incidence is generally the limit of power with
29 currently conducted animal studies (with the exception of rare tumors that are virtually markers
30 for a particular agent, e.g., angiosarcoma caused by vinyl chloride).

1 Target organs of carcinogenesis for agents that cause cancer in both animals and humans
2 are most often concordant at one or more sites (Tomatis et al., 1989; Huff, 1994). However,
3 concordance by site is not uniform. *The default assumption is that target organ concordance is*
4 *not a prerequisite for evaluating the implications of animal study results for humans.* This is a
5 public health conservative science policy. The mechanisms of control of cell growth and
6 differentiation are concordant among species, but there are marked differences among species in
7 the way control is managed in various tissues. For example, in humans, mutation of the tumor
8 suppressor gene p53 is one of the most frequently observed genetic changes in tumors. This
9 tumor suppressor is also observed to be operating in some rodent tissues, but other growth
10 control mechanisms predominate in rodents. Thus, an animal response may be due to changes in
11 a control that are relevant to humans, but appear in animals in a different way. However, it is
12 appropriate under these guidelines to consider the influences of route of exposure, metabolism,
13 and, particularly, hormonal modes of action that may either support or not support target organ
14 concordance between animals and humans. When data allow, these influences are considered in
15 deciding whether the default remains appropriate in individual instances (NRC, 1994, p. 121).
16 An exception to the basic default of not assuming site concordance exists in the context of
17 toxicokinetic modeling. Site concordance is inherently assumed when these models are used to
18 estimate delivered dose in humans based on animal data.

19 As in the approach of the National Toxicology Program and the International Agency
20 for Research on Cancer, *the default is to include benign tumors observed in animal studies in the*
21 *assessment of animal tumor incidence if they have the capacity to progress to the malignancies*
22 *with which they are associated.* This treats the benign and malignant tumors as representative
23 of related responses to the test agent, which is scientifically appropriate. This is a science policy
24 decision that is somewhat more conservative of public health than not including benign tumors
25 in the assessment. Nonetheless, in assessing findings from animal studies, a greater proportion
26 of malignancy is weighed more heavily than a response with a greater proportion of benign
27 tumors. Greater frequency of malignancy of a particular tumor type in comparison with other
28 tumor responses observed in an animal study is also a factor to be considered in selecting the
29 response to be used in dose response assessment.

Benign tumors that are not observed to progress to malignancy are assessed on a case-by-case basis. There is a range of possibilities for their overall significance. They may deserve attention because they are serious health problems even though they are not malignant; for instance, benign tumors may be a health risk because of their effect on the function of a target tissue such as the brain. They may be significant indicators of the need for further testing of an agent if they are observed in a short term test protocol, or such an observation may add to the overall weight of evidence if the same agent causes malignancies in a long term study. Knowledge of the mode of action associated with a benign tumor response may aid in the interpretation of other tumor responses associated with the same agent. In other cases, observation of a benign tumor response alone may have no significant health hazard implications when other sources of evidence show no suggestion of carcinogenicity.

1.3.2.3. How Do Metabolic Pathways Relate Across Species?

The default assumption is that there is a similarity of the basic pathways of metabolism and the occurrence of metabolites in tissues in regard to the species-to-species extrapolation of cancer hazard and risk. If comparative metabolism studies were to show no similarity between the tested species and humans and a metabolite(s) were the active form, there would be less support for an inference that the animal response(s) relates to humans. In other cases, parameters of metabolism may vary quantitatively between species; this becomes part of deciding on an appropriate human equivalent dose based on animal studies, optimally in the context of a toxicokinetic model.

1.3.2.4. How Do Toxicokinetic Processes Relate Across Species?

A major issue is how to estimate human equivalent doses in extrapolating from animal studies. As a default for oral exposure, a human equivalent dose is estimated from data on another species by an adjustment of animal oral dose by a scaling factor of body weight to the 0.75 power. This adjustment factor is used because it represents scaling of metabolic rate across animals of different size. Because the factor adjusts for a parameter that can be improved on and brought into more sophisticated toxicokinetic modeling, when such data become available, the default assumption of 0.75 power can be refined or replaced.

1 *For inhalation exposure, a human equivalent dose is estimated by default methodologies*
2 *that provide estimates of lung deposition and of internal dose.* The methodologies can be refined
3 to more sophisticated forms with data on toxicokinetic and metabolic parameters of the specific
4 agent. This default assumption, like the one with oral exposure, is selected in part because it
5 lays a foundation for incorporating better data. The use of information to improve dose
6 estimation from applied, to internal, to delivered dose is encouraged, including use of
7 toxicokinetic modeling instead of any default, where data are available. Health conservatism is
8 not an element in choosing the default.

9 For a route-to-route of exposure extrapolation, *the default assumption is that an agent*
10 *that causes internal tumors by one route of exposure will be carcinogenic by another route if it is*
11 *absorbed by the second route to give an internal dose.* This is a qualitative assumption and is
12 considered to be public health conservative. The rationale is that for internal tumors an internal
13 dose is significant no matter what the route of exposure. Additionally, the metabolism of the
14 agent will be qualitatively the same for an internal dose. The issue of quantitative extrapolation
15 of the dose-response relationship from one route to another is addressed case by case.
16 Quantitative extrapolation is complicated by considerations such as first-pass metabolism, but is
17 approachable with empirical data. Adequate data are necessary to demonstrate that an agent will
18 act differently by one route versus another route of exposure.

19 20 **1.3.2.5. What Is the Correlation of the Observed Dose Response Relationship to the** 21 **Relationship at Lower Doses?**

22 The overriding preferred approach is to use a biologically based or case-specific model
23 for both the observed range and extrapolation below that range when there are sufficient data.
24 While biologically based models are still under development, it is likely that they will be used
25 more frequently in the future. *The default procedure for the observed range of data, when the*
26 *preferred approach cannot be used, is to use a curve-fitting model.*

27 In the absence of data supporting a biologically based or case-specific model for
28 extrapolation outside of the observed range, the choice of approach is based on the view of mode
29 of action of the agent arrived at in the hazard assessment. *A linear default approach is used*
30 *when the mode of action information is supportive of linearity or, alternatively, is insufficient to*

1 *support a nonlinear mode of action.* The linear approach is used when a view of the mode of
2 action indicates a linear response, for example, when a conclusion is made that an agent
3 directly causes alterations in DNA, a kind of interaction that not only theoretically requires one
4 reaction, but also is likely to be additive to ongoing, spontaneous gene mutation. Other kinds of
5 activity may have linear implications, e.g., linear rate-limiting steps, that support a linear
6 procedure also. The linear approach is to draw a straight line between a point of departure from
7 observed data, generally, as a default, the LED₁₀, and the origin (zero dose, zero response).
8 Other points of departure may be more appropriate for certain data sets; these may be used
9 instead of the LED₁₀. This approach is generally considered to be public health conservative.
10 The LED₁₀ is the lower 95% limit on a dose that is estimated to cause a 10% response. This
11 level is chosen to account (conservatively) for experimental variability. Additionally, it is
12 chosen because it rewards experiments with better designs in regard to number of doses and dose
13 spacing, since these generally will have narrower confidence limits. It is also an appropriate
14 representative of the lower end of the observed range because the limit of detection of studies of
15 tumor effect is about 10%.

16 The linear default is thought to generally produce an upper bound on potential risk at low
17 doses, e.g., a 1/100,000 to 1/1,000,000 risk; the straight line approach gives numerical results
18 about the same as a linearized multistage procedure (Krewski et al., 1984). This upper bound is
19 thought to cover the range of human variability although, in some cases, it may not completely
20 do so (Bois et al., 1995). The EPA considers the linear default to be inherently conservative of
21 public health, without addition of another factor for human variability. In any case, the size of
22 such a factor would be hard to determine since a good empirical basis on which to construct an
23 estimate does not currently exist. The question of what may be the actual variability in human
24 sensitivity is one that the 1994 NRC report discussed as did the 1993 NRC report on pesticides
25 in children and infants. The NRC has recommended research on the question, and the EPA and
26 other agencies have begun such research.

27 *When adequate data on mode of action show that linearity is not the most reasonable*
28 *working judgment and provide sufficient evidence to support a nonlinear mode of action, the*
29 *default changes to a different approach-- a margin of exposure analysis--which assumes that*
30 *nonlinearity is more reasonable.* The departure point is again generally the LED₁₀. A margin of

1 exposure analysis compares the LED₁₀ with the dose associated with the environmental
2 exposure(s) of interest by computing the ratio between the two.

3 The purpose of a margin of exposure analysis is to provide the risk manager with all
4 available information on how much reduction in risk may be associated with reduction in
5 exposure from the point of departure. This is to support the risk manager's decision as to what
6 constitutes an acceptable margin of exposure, given requirements of the statute under which the
7 decision is being made. There are several factors to be considered. (For perspective, keep in
8 mind that a sufficient basis to support this nonlinear procedure often will include data on
9 responses that are precursors to tumor effects. This means that the point of departure may well
10 be from these biological response data rather than tumor incidence data, e.g., hormone levels,
11 mitogenic effects.) One factor to consider is the slope of the dose response curve at the point of
12 departure. A steeper slope implies an apparent greater reduction in risk as exposure decreases.
13 This may support a smaller margin of exposure. Conversely, a shallow slope may support use of
14 a greater margin of exposure. A second factor is the nature of the response used in the
15 assessment--A precursor effect or frank toxicity or tumor response. The latter two may support
16 a greater margin of exposure. A third factor is the nature and extent of human variability in
17 sensitivity to the phenomenon. A fourth factor is the agent's persistence in the body. Greater
18 variability or persistence argue for greater margins of exposure. A fifth factor is human
19 sensitivity to the phenomenon as compared with experimental animals. The size of the margin
20 of exposure that is acceptable would increase or decrease as this factor increases or decreases. If
21 human variability cannot be estimated based on data, it should be considered to be at least 10-
22 fold. Similarly, if comparison of species sensitivities cannot be estimated from available data,
23 humans can be considered to be 10-fold more sensitive. If it is found that humans are less
24 sensitive than animals a factor that is a fraction no smaller than 1/10 may be assumed. The 10-
25 fold factors are moderately conservative, traditional ones used for decades in the assessment of
26 toxicological effects. It should not be assumed that the numerical factors are the sole components
27 for determination of an acceptable margin of exposure. Each case calls for individual judgment.
28 It should be noted that for cancer assessment the margin of exposure analysis begins from a
29 point of departure that is adjusted for toxicokinetic differences between species to give a human
30 equivalent dose. Since the traditional factor for interspecies difference is thought to contain a

1 measure for toxicokinetics as well as sensitivity to effect, the result of beginning with a human
2 equivalent dose is to add some conservatism. The ultimate judgment whether a particular
3 margin of exposure is acceptable is a risk management decision under applicable law, rather than
4 being inherent in the risk assessment. Nonetheless, the risk assessor is responsible for providing
5 scientific rationale to support the the decision.

6 *When the mode of action information indicates that the dose response may be adequately*
7 *described by both a linear and a nonlinear approach, then the default is to present both the*
8 *linear and margin of exposure analyses.* An assessment may use both linear and nonlinear
9 approaches either for responses that are thought to result from different modes of action or for
10 presenting considerations for a response that appears to be very different at high and low doses
11 due to influence of separate modes of action. Also, separate approaches may be used for
12 different induced responses (i.e. tumor types) from the same agent. These would also be carried
13 forward and presented in the assessment.

14 Figure 1-1 presents the decision points in deciding on a dose response approach or approaches.
15

Data to Support:					
Biologically Based or Case-Specific Model	yes	no	no	no	no
Linearity		yes	no	yes	no
Nonlinearity		no	yes	yes	no
Extrapolation Used:	model	default--linear	default--nonlinear	default--linear and nonlinear	default--linear

Figure 1-1. Decisions on dose response assessment approaches for the range of extrapolation.

1 *A default assumption is made that cumulative dose received over a lifetime, expressed as*
2 *a lifetime average daily dose, is an appropriate measure of dose.* This assumes that a high dose
3 of such an agent received over a shorter period of time is equivalent to a low dose spread over a
4 lifetime. This is thought to be a relatively public health conservative assumption and has
5 empirical support (Monro, 1992). An example of effects of short-term, high exposure that
6 results in subsequent cancer development is treatment of cancer patients with certain
7 chemotherapeutic agents. An example of cancer from long-term exposure to an agent of
8 relatively low potency is smoking. Whether the cumulative dose measure is exactly the correct
9 measure in both such instances is not certain and should be assessed case by case and altered
10 when data are available to support another approach. Other measures of dose that consider dose
11 rate and duration are appropriate, e.g., when an agent acts by causing cell toxicity or hormone
12 disruption. In these cases both agent concentration and duration are likely to be important,
13 because such effects are generally observed to be reversible at cessation of short-term exposure.

15 **1.4. CHARACTERIZATIONS**

16 The risk characterization process first summarizes findings on hazard, dose response, and
17 exposure characterizations, then develops an integrative analysis of the whole risk case. It ends
18 in a nontechnical Risk Characterization Summary. The Risk Characterization Summary is a
19 presentation for risk managers who may or may not be familiar with the scientific details of
20 cancer assessment. It also provides information for other interested readers. The initial steps in
21 the risk characterization process are to make building blocks in the form of characterizations of
22 the assessments of hazard, dose response, and exposure. The individual assessments and
23 characterizations are then integrated to arrive at risk estimates for exposure scenarios of interest.
24 There are two reasons for individually characterizing the hazard, dose response, and exposure
25 assessments. One is that they are often done by different people than those who do the
26 integrative analyses. The second is that there is very often a lapse of time between the conduct
27 of hazard and dose response analyses and the conduct of exposure assessment and integrative
28 analysis. Thus, it is necessary to capture characterizations of assessments as the assessments are
29 done to avoid the need to go back and reconstruct them. Figure 1-2 shows the relationships of
30 analyses. The figure does not necessarily correspond to the number of documents involved;

1 there may be one or several. "Integrative analysis" is a generic term. At EPA, the documents of
2 various programs that contain integrative analyses have other names such as the "Staff Paper"
3 that discusses air quality criteria issues. In the following sections, the elements of this figure are
4 discussed.
5

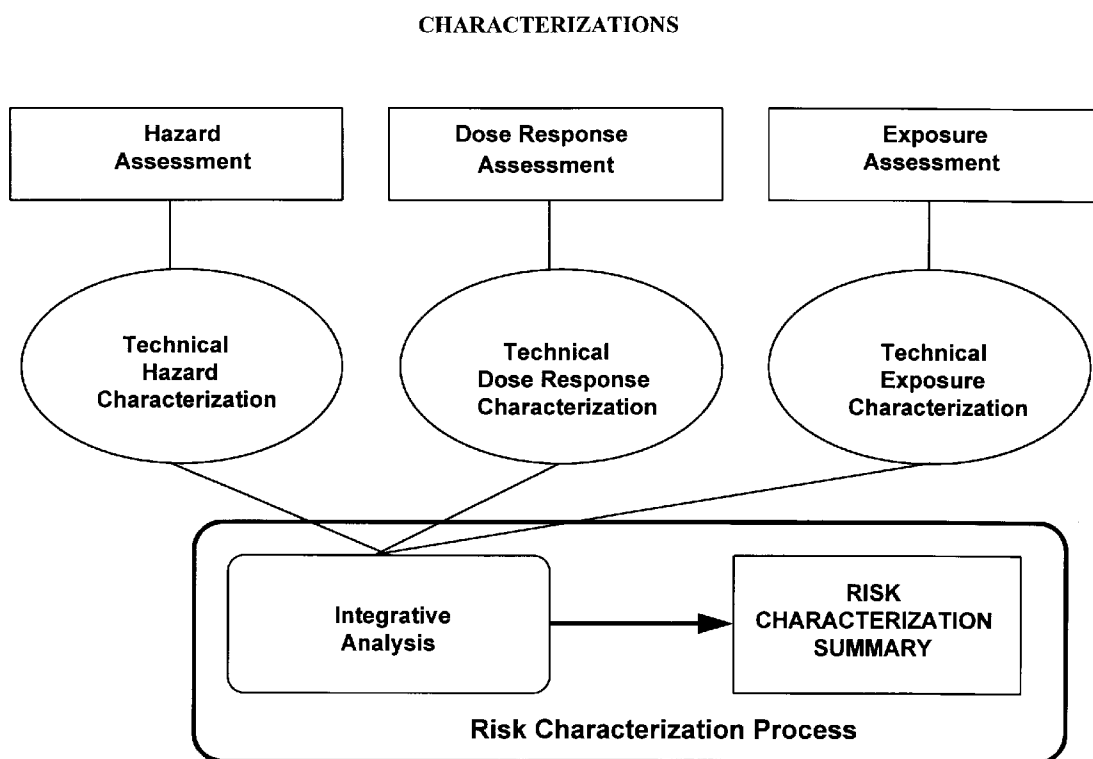


Figure 1-2. Risk Characterization

2. HAZARD ASSESSMENT

2.1. OVERVIEW OF HAZARD ASSESSMENT AND CHARACTERIZATION

2.1.1. Analyses of Data

The purpose of hazard assessment is to review and evaluate data pertinent to two questions: (1) whether an agent may pose a carcinogenic hazard to human beings and (2) under what circumstances an identified hazard may be expressed (NRC, 1994, p. 142). Hazard assessment is composed of analyses of a variety of data that may range from observations of tumor responses to analysis of structure-activity relationships. The purpose of the assessment is not simply to assemble these separate evaluations; its purpose is to construct a total case analysis examining the biological story the data reveal as a whole about carcinogenic effects, mode of action, and implications of these for human hazard and dose response evaluation. Weight of evidence conclusions come from the combined strength and coherence of inferences appropriately drawn from all of the available evidence. To the extent that data permit, hazard assessment addresses the mode of action question as both an initial step in considering appropriate approaches to dose response assessment and as a part of identifying human hazard potential.

The topics in this section include analysis of tumor data, both animal and human, and analysis of other key information about properties and effects that relate to carcinogenic potential. The section addresses how information can be used to evaluate potential modes of action. It also provides guidance on performing a weight of evidence evaluation.

2.1.2. Cross-Cutting Topics for Data Integration

Two topics are included in the analysis of each kind of available data: first, gathering information from available data about the conditions of expression of hazard and second, gathering perspectives on the agent's potential mode of action.

2.1.2.1. *Conditions of Expression*

Information on the significance of the route of exposure may be available from human or animal studies on the agent itself or on structural analogues. This information may be found in

1 studies of the agent or analogue for toxicological endpoints other than cancer under acute or
2 subchronic or chronic exposure regimens. Studies of metabolism or toxicokinetics of the agent
3 similarly may provide pertinent data.

4 Each kind of data is also examined for information on conditions that affect expression
5 of carcinogenic effect such as presence or absence of metabolic pathways. If carcinogenicity is
6 secondary to another toxic effect, the physiological or tissue changes that mark the other toxicity
7 are examined. Comparison of metabolic processes and toxicity processes in humans and animals
8 also bears on the relevance of animal responses to human hazard. Included in the examination
9 are the questions of the potential range of human variability and whether any special sensitivity
10 may occur because of age, sex, preexisting disease, or other condition.

11 12 **2.1.2.2. *Mode of Action***

13 Information on an agent's potential mode(s) of action is important in considering the
14 relevance of animal effects to assessment of human hazard. It also plays an important role in
15 selecting dose response approach(es), which are generally either biologically based models or
16 case-specific models incorporating mode of action data or default procedures based on more
17 limited data that support inferences about the likely shape of the dose response curve.

18 Each kind of data may provide some insight about mode of action and insights are
19 gathered from each to be considered together as discussed in section 2.4. In Appendix C, is a
20 background discussion of some of the development of views about carcinogenic processes.

21 22 **2.1.3. *Presentation of Results***

23 Presentation of the results of hazard assessment follows Agency guidance as discussed in
24 section 2.7. The results are presented in a technical hazard characterization that serves as a
25 support to later risk characterization. It includes:

- 26 • a summary of the evaluations of hazard data,
- 27 • the rationales for its conclusions, and
- 28 • an explanation of the significant strengths or limitations of the conclusions.

29 Another presentation feature is the use of a weight of evidence narrative that includes
30 both a conclusion about the weight of evidence of carcinogenic potential and a summary of the

1 data on which the conclusion rests. This narrative is a brief summary that replaces the
2 alphanumerical classification system used in EPA's previous guidelines.

3 4 **2.2. ANALYSIS OF TUMOR DATA**

5 Evidence of carcinogenicity comes from finding tumor increases in humans or laboratory
6 animals exposed to a given agent, or from finding tumors following exposure to structural
7 analogues to the compound under review. The significance of observed or anticipated tumor
8 effects is evaluated in reference to all of the other key data on the agent. This section contains
9 guidance for analyzing human and animal studies to decide whether there is an association
10 between exposure to an agent or a structural analogue and occurrence of tumors. Note that the
11 use of the term "tumor" here is generic, meaning malignant neoplasms or a combination of
12 malignant and corresponding benign neoplasms.

13 Observation of only benign neoplasias may or may not have significance. Benign tumors
14 that are not observed to progress to malignancy are assessed on a case-by-case basis. There is a
15 range of possibilities for their overall significance. They may deserve attention because they are
16 serious health problems even though they are not malignant; for instance, benign tumors may be
17 a health risk because of their effect on the function of a target tissue such as the brain. They
18 may be significant indicators of the need for further testing of an agent if they are observed in a
19 short term test protocol, or such an observation may add to the overall weight of evidence if the
20 same agent causes malignancies in a long term study. Knowledge of the mode of action
21 associated with a benign tumor response may aid in the interpretation of other tumor responses
22 associated with the same agent. In other cases, observation of a benign tumor response alone
23 may have no significant health hazard implications when other sources of evidence show no
24 suggestion of carcinogenicity.

25 26 **2.2.1. Human Data**

27 Human data may come from epidemiologic studies or case reports. Epidemiology is the
28 study of the distributions and causes of disease within human populations. The goals of cancer
29 epidemiology are to identify differences in cancer risk between different groups in a population
30 or between different populations, and then to determine the extent to which these differences in

1 risk can be attributed causally to specific exposures to exogenous or endogenous factors.
2 Epidemiologic data are extremely useful in risk assessment because they provide direct evidence
3 that a substance produces cancer in humans, thereby avoiding the problem of species to species
4 inference. Thus, when available human data are extensive and of good quality, they are
5 generally preferable over animal data and should be given greater weight in hazard
6 characterization and dose response assessment, although both are utilized.

7 Null results from a single epidemiologic study cannot prove the absence of carcinogenic
8 effects because they can arise either from being truly negative or from inadequate statistical
9 power, inadequate design, imprecise estimates, or confounding factors. However, null results
10 from a well-designed and well-conducted epidemiologic study that contains usable exposure data
11 can help to define upper limits for the estimated dose of concern for human exposure if the
12 overall weight of the evidence indicates that the agent is potentially carcinogenic in humans.

13 Epidemiology can also complement experimental evidence in corroborating or clarifying
14 the carcinogenic potential of the agent in question. For example, observations from
15 epidemiologic studies that elevated cancer incidence occurs at sites corresponding to those at
16 which laboratory animals experience increased tumor incidence can strengthen the weight of
17 evidence of human carcinogenicity. On the other hand, strong nonpositive epidemiologic data
18 alone or in conjunction with compelling mechanistic information can lend support to a
19 conclusion that animal responses may not be predictive of a human response. Furthermore, the
20 advent of biochemical or molecular epidemiology may help improve understanding of the
21 mechanisms of human carcinogenesis.

22 23 **2.2.1.1. *Types of Studies***

24 The major types of cancer epidemiologic studies are analytical epidemiologic studies and
25 descriptive or correlation epidemiologic studies. Each study type has well-known strengths and
26 weaknesses that affect interpretation of study results as summarized below (Kelsey et al., 1986;
27 Lilienfeld and Lilienfeld, 1979; Mausner and Kramer, 1985; Rothman, 1986).

28 Analytical epidemiologic studies are most useful for identifying an association between
29 human exposure and adverse health effects. Analytical study designs include case-control
30 studies and cohort studies. In case-control studies, groups of individuals with (cases) and

without (controls) a particular disease are identified and compared to determine differences in exposure. In cohort studies, a group of "exposed" and "nonexposed" individuals are identified and studied over time to determine differences in disease occurrence. Cohort studies can either be performed prospectively or retrospectively from historical records.

Descriptive or correlation epidemiologic studies (sometimes called ecological studies) examine differences in disease rates among populations in relation to age, gender, race, and differences in temporal or environmental conditions. In general, these studies can only identify patterns or trends in disease occurrence over time or in different geographical locations but cannot ascertain the causal agent or degree of exposure. These studies, however, are often very useful for generating hypotheses for further research.

Biochemical or molecular epidemiologic studies are studies in which laboratory methods are incorporated in analytical investigations. The application of techniques for measuring cellular and molecular alterations due to exposure to specific environmental agents may allow conclusions to be drawn about the mechanisms of carcinogenesis. The use of biological biomarkers in epidemiology may improve assessment of exposure and internal dose.

Case reports describe a particular effect in an individual or group of individuals who were exposed to a substance. These reports are often anecdotal or highly selected in nature and are of limited use for hazard assessment. However, reports of cancer cases can identify associations particularly when there are unique features such as an association with an uncommon tumor (e.g., vinyl chloride and angiosarcoma or diethylstilbestrol and clear-cell carcinoma of the vagina).

2.2.1.2. *Criteria for Assessing Adequacy of Epidemiologic Studies*

Criteria for assessing the adequacy of epidemiologic studies are well recognized. Characteristics that are desirable in these studies include (1) clear articulation of study objectives or hypothesis, (2) proper selection and characterization of the exposed and control groups, (3) adequate characterization of exposure, (4) sufficient length of follow-up for disease occurrence, (5) valid ascertainment of the causes of cancer morbidity and mortality, (6) proper consideration of bias and confounding factors, (7) adequate sample size to detect an effect, (8) clear, well-documented, and appropriate methodology for data collection and analysis, (9) adequate

1 response rate and methodology for handling missing data, and (10) complete and clear
2 documentation of results. Ideally, these conditions should be satisfied, where appropriate, but
3 rarely can a study meet all of them. No single criterion determines the overall adequacy of a
4 study. The following discussions highlight the major factors included in an analysis of
5 epidemiologic studies.

6 7 *Population Issues*

8 The ideal comparison would be between two populations that differ only in exposure to
9 the agent in question. Because this is seldom the case, it is important to identify sources of bias
10 inherent in a study's design or data collection methods. Bias can arise from several sources,
11 including noncomparability between populations of factors such as general health (McMichael,
12 1976), diet, lifestyle, or geographic location; differences in the way case and control individuals
13 recall past events; differences in data collection that result in unequal ascertainment of health
14 effects in the populations; and unequal follow-up of individuals. Both acceptance of studies for
15 assessment and judgment of their strengths or weaknesses depend on identifying their sources of
16 bias and the effects on study results.

17 18 *Exposure Issues*

19 For epidemiologic data to be useful in determining whether there is an association
20 between health effects and exposure to an agent, there must be adequate characterization of
21 exposure information. In general, greater weight should be given to studies with more precise
22 and specific exposure estimates.

23 Questions to address about exposure are: What can one reliably conclude about the level,
24 duration, route, and frequency of exposure of individuals in one population as compared with
25 another? How sensitive are study results to uncertainties in these parameters?

26 Actual exposure measurements are not available for many retrospective studies.
27 Therefore, surrogates are often used to reconstruct exposure parameters when historical
28 measurements are not available. These may involve attributing exposures to job classifications
29 in a workplace or to broader occupational or geographic groupings. Use of surrogates carries a
30 potential for misclassification in that individuals may be placed in the incorrect exposure group.

1 Misclassification generally leads to reduced ability of a study to detect differences between study
2 and referent populations.

3 When either current or historical monitoring data are available, the exposure evaluation
4 includes consideration of the error bounds of the monitoring and analytic methods and whether
5 the data are from routine or accidental exposures. The potentials for misclassification and
6 measurement errors are amenable to both qualitative and quantitative analysis. These are
7 essential analyses for judging a study's results because exposure estimation is the most critical
8 part of a retrospective study.

9 Biological markers potentially offer excellent measures of exposure (Hulka and
10 Margolin, 1992; Peto and Darby, 1994). Validated markers of exposure such as alkylated
11 hemoglobin from exposure to ethylene oxide (van Sittert et al., 1985) or urinary arsenic
12 (Enterline et al., 1987) can greatly improve estimates of dose. Markers closely identified with
13 effects promise to greatly increase the ability of studies to distinguish real effects from bias at
14 low levels of relative risk between populations (Taylor et al., 1994; Biggs et al., 1993) and to
15 resolve problems of confounding risk factors.

16 17 *Confounding Factors*

18 Because epidemiologic studies are mostly observational, it is not possible to guarantee
19 the control of confounding variables, which may affect the study outcome. A confounding
20 variable is a risk factor, independent of the putative agent, that is distributed unequally among
21 the exposed and unexposed populations (e.g., smoking habits, lifestyle). Adjustment for
22 possible confounding factors can occur either in the design of the study (e.g., matching on
23 critical factors) or in the statistical analysis of the results. The influence of a potential
24 confounding factor is limited by the effect of the exposure of interest. For example, a twofold
25 effect of an exposure requires that the confounder effect be at least as big. The latter may not be
26 possible due to the presentation of the data or because needed information was not collected
27 during the study. In this case, indirect comparisons may be possible. For example, in the
28 absence of data on smoking status among individuals in the study population, an examination of
29 the possible contribution of cigarette smoking to increased lung cancer risk may be based on
30 information from other sources such as the American Cancer Society's longitudinal studies

(Hammand, 1966; Garfinkel and Silverberg, 1991). The effectiveness of adjustments contributes to the ability to draw inferences from a study.

Different studies involving exposure to an agent may have different confounding factors. If consistent increases in cancer risk are observed across a collection of studies with different confounding factors, the inference that the agent under investigation was the etiologic factor is strengthened, even though complete adjustment for confounding factors cannot be made and no single study supports a strong inference.

It also may be the case that the agent of interest is a risk factor in conjunction with another agent. This relationship may be revealed in a collection of studies such as in the case of asbestos exposure and smoking.

Sensitivity

Sensitivity, or the ability of a study to detect real effects, is a function of several factors. Greater size of the study population(s) (sample size) increases sensitivity, as does greater exposure (levels and duration) of the population members. Because of the often long latency period in cancer development, sensitivity also depends on whether adequate time has elapsed since exposure began for effects to occur. A unique feature that can be ascribed to the effects of a particular agent (such as a tumor type that is seen only rarely in the absence of the agent) can increase sensitivity by permitting separation of bias and confounding factors from real effects. Similarly, a biomarker particular to the agent can permit these distinctions. Statistical reanalyses of data, particularly an examination of different exposure indices, can give insight on potential exposure-response relationships. These are all factors to explore in statistical analysis of the data.

Statistical Considerations

The analysis applies appropriate statistical methods to ascertain whether or not there is any significant association between exposure and effects. A description of the method or methods should include the reasons for their selection. Statistical analyses of the potential effects of bias or confounding factors are part of addressing the significance of an association, or lack of one, and whether a study is able to detect any effect.

The analysis augments examination of the results for the whole population with exploration of the results for groups with comparatively greater exposure or time since first exposure. This may support identifying an association or establishing a dose response trend. When studies show no association, such exploration may apply to determining an upper limit on potential human risk for consideration alongside results of animal tumor effects studies.

Combining Statistical Evidence Across Studies

Meta-analysis is a means of comparing and synthesizing studies dealing with similar health effects and risk factors. It is intended to introduce consistency and comprehensiveness into what otherwise might be a more subjective review of the literature. When utilized appropriately, meta-analysis can enhance understanding of associations between sources and their effects that may not be apparent from examination of epidemiologic studies individually. Whether to conduct a meta-analysis depends on several issues. These include the importance of formally examining sources of heterogeneity, the refinement of the estimate of the magnitude of an effect, and the need for information beyond that provided by individual studies or a narrative review. Meta-analysis may not be useful in some circumstances. These include when the relationship between exposure and disease is obvious without a more formal analysis, when there are only a few studies of the key health outcomes, when there is insufficient information from available studies related to disease, risk estimate, or exposure classification, or when there are substantial confounding or other biases that cannot be adjusted for in the analysis (Blair et al., 1995; Greenland, 1987; Peto, 1992).

2.2.1.3. Criteria for Causality

A causal interpretation is enhanced for studies to the extent that they meet the criteria described below. None of the criteria is conclusive by itself, and the only criterion that is essential is the temporal relationship. These criteria are modeled after those developed by Bradford Hill in the examination of cigarette smoking and lung cancer (Rothman, 1986) and they need to be interpreted in the light of all other information on the agent being assessed.

- **Temporal relationship**: The development of cancers require certain latency periods, and while latency periods vary, existence of such periods is generally acknowledged.

Thus, the disease has to occur within a biologically reasonable time after initial exposure. This feature must be present if causality is to be considered.

- Consistency: Associations occur in several independent studies of a similar exposure in different populations, or associations occur consistently for different subgroups in the same study. This feature usually constitutes strong evidence for a causal interpretation when the same bias or confounding is not also duplicated across studies.
- Magnitude of the association: A causal relationship is more credible when the risk estimate is large and precise (narrow confidence intervals).
- Biological gradient: The risk ratio (i.e., the ratio of the risk of disease or death among the exposed to the risk of the unexposed) increases with increasing exposure or dose. A strong dose response relationship across several categories of exposure, latency, and duration is supportive for causality given that confounding is unlikely to be correlated with exposure. The absence of a dose response relationship, however, is not by itself evidence against a causal relationship.
- Specificity of the association: The likelihood of a causal interpretation is increased if an exposure produces a specific effect (one or more tumor types also found in other studies) or if a given effect has a unique exposure.
- Biological plausibility: The association makes sense in terms of biological knowledge. Information is considered from animal toxicology, toxicokinetics, structure-activity relationship analysis, and short-term studies of the agent's influence on events in the carcinogenic process considered.
- Coherence: The cause-and-effect interpretation is in logical agreement with what is known about the natural history and biology of the disease, i.e., the entire body of knowledge about the agent.

2.2.1.4. Assessment of Evidence of Carcinogenicity from Human Data

In the evaluation of carcinogenicity based on epidemiologic studies, it is necessary to critically evaluate each study for the confidence in findings and conclusions as discussed under section 2.2.1.2. All studies that are properly conducted, whether yielding positive or null results,

1 or even suggesting protective carcinogenic effects, should be considered in assessing the totality
2 of the human evidence. Although a single study may be indicative of a cause-effect relationship,
3 confidence in inferring a causal relationship is increased when several independent studies are
4 concordant in showing the association, when the association is strong, and when other criteria
5 for causality are also met. Conclusions about the overall evidence for carcinogenicity from
6 available studies in humans should be summarized along with a discussion of strengths or
7 limitations of the conclusions.

8 9 **2.2.2. Animal Data**

10 Various kinds of whole animal test systems are currently used or are under development
11 for evaluating potential carcinogenicity. Cancer studies involving chronic exposure for most of
12 the life span of an animal are generally accepted for evaluation of tumor effects (Tomatis et al.,
13 1989; Rall, 1991; Allen et al., 1988; but see Ames and Gold, 1990). Other studies of special
14 design are useful for observing formation of preneoplastic lesions or tumors or investigating
15 specific modes of action.

16 17 **2.2.2.1. Long-Term Carcinogenicity Studies**

18 The objective of long-term carcinogenesis bioassays is to determine the carcinogenic
19 potential and dose response relationships of the test agent. Long-term rodent studies are
20 designed to examine the production of tumors as well as preneoplastic lesions and other
21 indications of chronic toxicity that may provide evidence of treatment-related effects and
22 insights into the way the test agent produces tumors. Current standardized long-term studies in
23 rodents test at least 50 animals per sex per dose group in each of three treatment groups and in a
24 concurrent control group, usually for 18 to 24 months, depending on the rodent species tested
25 (OECD, 1981; U.S. EPA, 1983a; U.S. EPA, 1983b; U.S. EPA, 1983c). The high dose in long-
26 term studies is generally selected to provide the maximum ability to detect treatment-related
27 carcinogenic effects while not compromising the outcome of the study due to excessive toxicity
28 or inducing inappropriate toxicokinetics (e.g., overwhelming detoxification or absorption
29 mechanisms). The purpose of two or more lower doses is to provide some information on the

1 shape of the dose response curve. Similar protocols have been and continue to be used by many
2 laboratories worldwide.

3 All available studies of tumor effects in whole animals are considered, at least
4 preliminarily. The analysis discards studies judged to be wholly inadequate in protocol, conduct,
5 or results. Criteria for the technical adequacy of animal carcinogenicity studies have been
6 published and should be used as guidance to judge the acceptability of individual studies (NTP,
7 1984; OSTP, 1985). Care is taken to include studies that provide some evidence bearing on
8 carcinogenicity or help interpret effects noted in other studies even if they have some limitations
9 of protocol or conduct. Such limited, but not wholly inadequate, studies can contribute as their
10 deficiencies permit. The findings of long-term rodent bioassays are always interpreted in
11 conjunction with results of prechronic studies along with toxicokinetic and metabolism studies
12 and other pertinent information, if available. Evaluation of tumor effects requires consideration
13 of both biological and statistical significance of the findings (Haseman, 1984, 1985, 1990,
14 1995). The following sections highlight the major issues in the evaluation of long-term
15 carcinogenicity studies.

16 17 *Dosing Issues*

18 In order to obtain the most relevant information from a long-term carcinogenicity study,
19 it is important to require maximization of exposure to the test material. At the same time, there
20 is a need for caution in using excessive high dose levels that would confound the interpretation
21 of study results to humans. The high dose is conventionally defined as a dose that produces
22 some toxic effects without either unduly affecting mortality from effects other than cancer or
23 producing significant adverse effects on the nutrition and health of the test animals (OECD,
24 1981; NRC, 1993b). It should be noted that practical upper limits have been established to avoid
25 the use of excessive high doses in long-term carcinogenicity studies (e.g., 5% of the test
26 substance in the feed for dietary studies [OECD, 1981]).

27 Evaluating the appropriateness of the high dose in carcinogenicity studies is based on
28 scientific judgment using all available relevant information. In general, if the test agent does not
29 appear to cause any specific target organ toxicity or perturbation of physiological function, an
30 adequate high dose would be a dose that causes no more than 10% reduction of body weight

gain over the life span of the animals. On the other hand, significant increases in mortality from effects other than cancer is accepted as clear evidence of frank toxicity, which indicates that an adequate high dose may have been exceeded. Other signs of treatment-related toxicity that may indicate that an adequate high dose has been exceeded include the following: (a) reduction of body weight gain of 10% or greater, (b) significant increases in abnormal behavioral and clinical signs, (c) significant changes in hematology or clinical chemistry, (d) saturation of absorption and detoxification mechanisms, or (e) marked changes in organ weight, morphology, and histopathology.

For dietary studies, weight gain reductions should be evaluated as to whether there is a palatability problem or an issue with food efficiency; certainly, the latter is a toxic manifestation. In the case of inhalation studies with respirable particles, evidence of impairment of normal clearance of particles from the lung should be considered along with other signs of toxicity to the respiratory airways to determine whether the high exposure concentration has been appropriately selected. For dermal studies, evidence of skin irritation may indicate that an adequate high dose has been reached.

Interpretation of carcinogenicity study results is profoundly affected by exposure conditions, especially by inappropriate dose selection. This is particularly important in studies that are nonpositive for carcinogenicity, since failure to reach a sufficient dose reduces the sensitivity of a study. A lack of tumorigenic responses at exposure levels that cause significant impairment of animal survival may also not be acceptable as negative findings because of the reduced sensitivity of the study. On the other hand, overt toxicity or inappropriate toxicokinetics due to excessive high doses may result in tumor effects that are secondary to the toxicity rather than directly attributable to the agent.

There are several possible outcomes regarding the study interpretation of the significance and relevance of tumorigenic effects associated with exposure or dose levels below, at, or above an adequate high dose. General guidance is given here that should not be taken as prescriptive; for each case, the information at hand is evaluated and a rationale should be given for the position taken.

- Adequate high dose: If an adequate high dose has been utilized, tumor effects are judged positive or negative depending on the presence or absence of significant tumor incidence increases, respectively.
- Excessive high dose: If toxicity or mortality is excessive at the high dose, interpretation depends on the finding of tumors or not.
 - (a) Studies that show tumor effects only at excessive doses may be compromised and may or may not carry weight, depending on the interpretation in the context of other study results and other lines of evidence. Results of such studies, however, are generally not considered suitable for risk extrapolation.
 - (b) Studies that show tumors at lower doses, even though the high dose is excessive and may be discounted, should be evaluated on their own merits.
 - (c) If a study does not show an increase in tumor incidence at a toxic high dose and appropriately spaced lower doses are used without such toxicity or tumors, the study is generally judged as negative for carcinogenicity.
- Inadequate high dose: Studies of inadequate sensitivity where an adequate high dose has not been reached may be used to bound the dose range where carcinogenic effects might be expected.

Statistical Considerations

The main aim of statistical evaluation is to determine whether exposure to the test agent is associated with an increase of tumor development. Statistical analysis of a long-term study should be performed for each tumor type separately. The incidence of benign and malignant lesions of the same cell type, usually within a single tissue or organ, are considered separately and are combined when scientifically defensible (McConnell et al., 1986).

Trend tests and pairwise comparison tests are the recommended tests for determining whether chance, rather than a treatment-related effect, is a plausible explanation for an apparent increase in tumor incidence. A trend test such as the Cochran-Armitage test (Snedecor and Cochran, 1967) asks whether the results in all dose groups together increase as dose increases. A pairwise comparison test such as the Fisher exact test (Fisher, 1932) asks whether an incidence in one dose group is increased over the control group. By convention, for both tests a

1 statistically significant comparison is one for which $p < 0.05$ that the increased incidence is due
2 to chance. Significance in either kind of test is sufficient to reject the hypothesis that chance
3 accounts for the result. A statistically significant response may or may not be biologically
4 significant and vice versa. The selection of a significance level is a policy choice based on a
5 trade-off between the risks of false positives and false negatives. A significance level of greater
6 or less than 5% is examined to see if it confirms other scientific information. When the
7 assessment departs from a simple 5% level, this should be highlighted in the risk
8 characterization. A two-tailed test or a one-tailed test can be used. In either case a rationale is
9 provided.

10 Considerations of multiple comparisons should also be taken into account. Haseman
11 (1983) analyzes typical animal bioassays testing both sexes of two species and concludes that,
12 because of multiple comparisons, a single tumor increase for a species-sex-site combination that
13 is statistically significant at the 1% level for common tumors or 5% for rare tumors corresponds
14 to a 7-8% significance level for the study as a whole. Therefore, animal bioassays presenting
15 only one significant result that falls short of the 1% level for a common tumor may be treated
16 with caution.

18 *Concurrent and Historical Controls*

19 The standard for determining statistical significance of tumor incidence comes from a
20 comparison of tumors in dosed animals as compared with concurrent control animals.
21 Additional insights about both statistical and biological significance can come from an
22 examination of historical control data (Tarone, 1982; Haseman, 1995). Historical control data
23 can add to the analysis particularly by enabling identification of uncommon tumor types or high
24 spontaneous incidence of a tumor in a given animal strain. Identification of common or
25 uncommon situations prompts further thought about the meaning of the response in the current
26 study in context with other observations in animal studies and with other evidence about the
27 carcinogenic potential of the agent. These other sources of information may reinforce or weaken
28 the significance given to the response in the hazard assessment. Caution should be exercised in
29 simply looking at the ranges of historical responses because the range ignores differences in
30 survival of animals among studies and is related to the number of studies in the database.

1 In analyzing results for uncommon tumors in a treated group that are not statistically
2 significant in comparison to concurrent controls, the analyst can use the experience of historical
3 controls to conclude that the result is in fact unlikely to be due to chance. In analyzing results
4 for common tumors, a different set of considerations comes into play. Generally speaking,
5 statistically significant increases in tumors should not be discounted simply because incidence
6 rates in the treated groups are within the range of historical controls or because incidence rates in
7 the concurrent controls are somewhat lower than average. Random assignment of animals to
8 groups and proper statistical procedures provide assurance that statistically significant results are
9 unlikely to be due to chance alone. However, caution should be used in interpreting results that
10 are barely statistically significant or in which incidence rates in concurrent controls are unusually
11 low in comparison with historical controls.

12 In cases where there may be reason to discount the biological relevance to humans of
13 increases in common animal tumors, such considerations should be weighed on their own merits
14 and clearly distinguished from statistical concerns.

15 When historical control data are used, the discussion needs to address several issues that
16 affect comparability of historical and concurrent control data. Among these issues are the
17 following: genetic drift in the laboratory strains; differences in pathology examination at
18 different times and in different laboratories (e.g., in criteria for evaluating lesions; variations in
19 the techniques for preparation or reading of tissue samples among laboratories); comparability of
20 animals from different suppliers. The most relevant historical data come from the same
21 laboratory and same supplier, gathered within 2 or 3 years one way or the other of the study
22 under review; other data should be used only with extreme caution.

24 *Assessment of Evidence of Carcinogenicity from Long-Term Animal Studies*

25 In general, observation of tumor effects under different circumstances lends support to
26 the significance of the findings for animal carcinogenicity. Significance is a function of the
27 number of factors present, and for a factor such as malignancy, the severity of the observed
28 pathology. The following observations add significance to the tumor findings:

- 29 ● uncommon tumor types
- 30 ● tumors at multiple sites

- tumors by more than one route of administration
- tumors in multiple species, strains, or both sexes
- progression of lesions from preneoplastic to benign to malignant
- reduced latency of neoplastic lesions
- metastases
- unusual magnitude of tumor response
- proportion of malignant tumors
- dose-related increases

These guidelines adopt the science policy position that tumor findings in animals indicate that an agent may produce such effects in humans. Moreover, the absence of tumor findings in well-conducted, long-term animal studies in at least two species provides reasonable assurance that an agent may not be a carcinogenic concern for humans. Each of these is a default assumption that may be adopted, when appropriate, after evaluation of tumor data and other key evidence.

Site concordance of tumor effects between animals and humans is an issue to be considered in each case. Thus far, there is evidence that growth control mechanisms at the level of the cell are homologous among mammals, but there is no evidence that these mechanisms are site concordant. Moreover, agents observed to produce tumors in both humans and animals have produced tumors either at the same (e.g., vinyl chloride) or different sites (e.g., benzene) (NRC, 1994). Hence, site concordance is not assumed a priori. On the other hand, certain processes with consequences for particular tissue sites (e.g., disruption of thyroid function) may lead to an anticipation of site concordance.

2.2.2.2. Other Studies

Various intermediate-term studies often use protocols that screen for carcinogenic or preneoplastic effects, sometimes in a single tissue. Some involve the development of various proliferative lesions, like foci of alteration in the liver (Goldsworthy et al., 1986). Others use tumor endpoints, like the induction of lung adenomas in the sensitive strain A mouse (Maronpot et al., 1986) or tumor induction in initiation-promotion studies using various organs such as the bladder, intestine, liver, lung, mammary gland, and thyroid (Ito et al., 1992). In these tests, the

1 selected tissue is, in a sense, the test system rather than the whole animal. Important information
2 concerning the steps in the carcinogenic process and mode of action can be obtained from
3 "start/stop" experiments. In these protocols, an agent is given for a period of time to induce
4 particular lesions or effects, then stopped to evaluate the progression or reversibility of processes
5 (Todd, 1986; Marsman and Popp, 1994).

6 Assays in genetically engineered rodents may provide insight into the chemical and gene
7 interactions involved in carcinogenesis (Tennant et al., 1995a). These mechanistically based
8 approaches involve activated oncogenes that are introduced (transgenic) or tumor suppressor
9 genes that are deleted (knocked-out). If appropriate genes are selected, not only may these
10 systems provide information on mechanisms, but the rodents typically show tumor development
11 earlier than the standard bioassay. Transgenic mutagenesis assays also represent a mechanistic
12 approach for assessing the mutagenic properties of agents as well as developing quantitative
13 linkages between exposure, internal dose, and mutation related to tumor induction (Morrison and
14 Ashby, 1994; Sisk et al., 1994; Hayward et al., 1995). These systems use a stable genomic
15 integration of a lambda shuttle vector that carries a *lacI* target gene and a *lacZ* reporter gene.

16 The support that these studies give to a determination of carcinogenicity rests on their
17 contribution to the consistency of other evidence about an agent. For instance, benzoyl peroxide
18 has promoter activity on the skin, but the overall evidence may be less supportive (Kraus et al.,
19 1995). These studies also may contribute information about mode of action. One needs to
20 recognize the limitations of these experimental protocols such as short duration, limited
21 histology, lack of complete development of tumors, or experimental manipulation of the
22 carcinogenic process that may limit their contribution to the overall assessment. Generally, their
23 results are appropriate as aids in the assessment for interpreting other toxicological evidence
24 (e.g., rodent chronic bioassays), especially regarding potential modes of action. With sufficient
25 validation, these studies may partially or wholly replace chronic bioassays in the future (Tennant
26 et al., 1995).

27 28 **2.2.3. Structural Analogue Data**

29 For some chemical classes, there is significant information available on the
30 carcinogenicity of analogues, largely in rodent bioassays. Analogue effects are instructive in

investigating carcinogenic potential of an agent as well as identifying potential target organs, exposures associated with effects, and potential functional class effects or modes of action. All appropriate studies are included and analyzed, whether indicative of a positive effect or not. Evaluation includes tests in various animal species, strains, and sexes; with different routes of administration; and at various doses, as data are available. Confidence in conclusions is a function of how similar the analogues are to the agent under review in structure, metabolism, and biological activity. This confidence needs to be considered to ensure a balanced position.

2.3. ANALYSIS OF OTHER KEY DATA

The physical, chemical, and structural properties of an agent, as well as data on endpoints that are thought to be critical elements of the carcinogenic process, provide valuable insights into the likelihood of human cancer risk. The following sections provide guidance for analyses of these data.

2.3.1. Physicochemical Properties

Physicochemical properties affect an agent's absorption, tissue distribution (bioavailability), biotransformation, and degradation in the body and are important determinants of hazard potential (and dose response analysis). Properties to analyze include, but are not limited to, the following: molecular weight, size, and shape; valence state; physical state (gas, liquid, solid); water or lipid solubility, which can influence retention and tissue distribution; and potential for chemical degradation or stabilization in the body.

An agent's potential for chemical reaction with cellular components, particularly with DNA and proteins, is also important. The agent's molecular size and shape, electrophilicity, and charge distribution are considered in order to decide whether they would facilitate such reactions.

2.3.2. Structure-Activity Relationships

Structure-activity relationship (SAR) analyses and models can be used to predict molecular properties, surrogate biological endpoints, and carcinogenicity. Overall, these

analyses provide valuable initial information on agents, which may strengthen or weaken the concern for an agent's carcinogenic potential.

Currently, SAR analysis is useful for chemicals and metabolites that are believed to initiate carcinogenesis through covalent interaction with DNA (i.e., DNA-reactive, mutagenic, electrophilic, or proelectrophilic chemicals) (Ashby and Tennant, 1991). For organic chemicals, the predictive capability of SAR analysis combined with other toxicity information has been demonstrated (Ashby and Tennant, 1994). The following parameters are useful in comparing an agent to its structural analogues and congeners that produce tumors and affect related biological processes such as receptor binding and activation, mutagenicity, and general toxicity (Woo and Arcos, 1989):

- nature and reactivity of the electrophilic moiety or moieties present,
- potential to form electrophilic reactive intermediate(s) through chemical, photochemical, or metabolic activation,
- contribution of the carrier molecule to which the electrophilic moiety(ies) is attached,
- physicochemical properties (e.g., physical state, solubility, octanol-water partition coefficient, half-life in aqueous solution),
- structural and substructural features (e.g., electronic, steric, molecular geometric),
- metabolic pattern (e.g., metabolic pathways and activation and detoxification ratio), and
- possible exposure route(s) of the agent.

Suitable SAR analysis of non-DNA-reactive chemicals and of DNA-reactive chemicals that do not appear to bind covalently to DNA requires knowledge or postulation of the probable mode(s) of action of closely related carcinogenic structural analogues (e.g., receptor-mediated, cytotoxicity-related). Examination of the physicochemical and biochemical properties of the agent may then provide the rest of the information needed in order to make an assessment of the likelihood of the agent's activity by that mode of action.

2.3.3. Comparative Metabolism and Toxicokinetics

Studies of the absorption, distribution, biotransformation, and excretion of agents permit comparisons among species to assist in determining the implications of animal responses for

human hazard assessment, supporting identification of active metabolites, identifying changes in distribution and metabolic pathway or pathways over a dose range, and making comparisons among different routes of exposure.

If extensive data are available (e.g., blood/tissue partition coefficients and pertinent physiological parameters of the species of interest), physiologically based pharmacokinetic models can be constructed to assist in a determination of tissue dosimetry, species-to-species extrapolation of dose, and route-to-route extrapolation (Connolly and Andersen, 1991; see section 3.2.2). If it is not contrary to available data, it is assumed as a default that toxicokinetic and metabolic processes are qualitatively comparable between species. Discussion of the defaults regarding quantitative comparison and their modifications appears in section 3.

The qualitative question of whether an agent is absorbed by a particular route of exposure is important for weight of evidence classification discussed in section 2.7.1. Decisions whether route of exposure is a limiting factor on expression of any hazard, in that absorption does not occur by a route, are based on studies in which effects of the agent, or its structural analogues, have been observed by different routes, on physical-chemical properties, or on toxicokinetics studies.

Adequate metabolism and pharmacokinetic data can be applied toward the following as data permit. Confidence in conclusions is enhanced when in vivo data are available.

- Identifying metabolites and reactive intermediates of metabolism and determining whether one or more of these intermediates are likely to be responsible for the observed effects. This information on the reactive intermediates will appropriately focus SAR analysis, analysis of potential modes of action, and estimation of internal dose in dose response assessment (D'Souza et al., 1987; Krewski et al., 1987).
- Identifying and comparing the relative activities of metabolic pathways in animals with those in humans. This analysis can provide insights for extrapolating results of animal studies to humans.
- Describing anticipated distribution within the body and possibly identifying target organs. Use of water solubility, molecular weight, and structure analysis can support qualitative inferences about anticipated distribution and excretion. In addition, describing whether the agent or metabolite of concern will be excreted rapidly or

slowly or will be stored in a particular tissue or tissues to be mobilized later can identify issues in comparing species and formulating dose response assessment approaches.

- Identifying changes in toxicokinetics and metabolic pathways with increases in dose. These changes may result in important differences in disposition of the agent or its generation of active forms of the agent between high and low dose levels. These studies play an important role in providing a rationale for dose selection in carcinogenicity studies.
- Determining bioavailability via different routes of exposure by analyzing uptake processes under various exposure conditions. This analysis supports identification of hazards for untested routes. In addition, use of physicochemical data (e.g., octanol-water partition coefficient information) can support an inference about the likelihood of dermal absorption (Flynn, 1990).

In all of these areas, attempts are made to clarify and describe as much as possible the variability to be expected because of differences in species, sex, age, and route of exposure. The analysis takes into account the presence of subpopulations of individuals who are particularly vulnerable to the effects of an agent because of toxicokinetic or metabolic differences (genetically or environmentally determined) (Bois et al., 1995).

2.3.4. Toxicological and Clinical Findings

Toxicological findings in experimental animals and clinical observations in humans are an important resource to the cancer hazard assessment. Such findings provide information on physiological effects, effects on enzymes, hormones, and other important macromolecules as well as on target organs for toxicity. Given that the cancer process represents defects in terminal differentiation, growth control, and cell death, developmental studies of agents may provide an understanding of the activity of an agent that carries over to cancer assessment. Toxicity studies in animals by different routes of administration support comparison of absorption and metabolism by those routes. Data on human variability in standard clinical tests may provide insight into the range of human sensitivity and common mechanisms to agents that affect the tested parameters.

2.3.5. Mode of Action-Related Endpoints and Short-Term Tests

A myriad of biochemical and biological endpoints relevant to the carcinogenic process provide important information in determining whether a cancer hazard exists and include, but are not limited to, mutagenesis, inhibition of gap junctional intercellular communication, increased cell proliferation, inhibition of programmed cell death, receptor activation, and immunosuppression. These precursor effects are discussed below.

2.3.5.1. *Direct DNA Effects*

Because cancer is the result of multiple genetic defects in genes controlling proliferation and tissue homeostasis (Vogelstein et al., 1988), the ability of an agent to affect DNA is of obvious importance. It is well known that many carcinogens are electrophiles that interact directly with DNA, resulting in DNA damage and adducts, and subsequent mutations (referred to in these guidelines as direct DNA effects) that are thought to contribute to the carcinogenic process (Shelby and Zeiger, 1990; Tinwell and Ashby, 1991). Thus, studies of these phenomena continue to be important in the assessment of cancer hazard. The EPA has published testing guidelines for detecting the ability of agents to affect DNA or chromosomes (EPA, 1991a). Information on agents that induce mutations in animal germ cells also deserves attention; several human carcinogens have been shown to be positive in rodent tests for the induction of genetic damage in both somatic and germ cells (Shelby, 1995).

2.3.5.2. *Secondary DNA Effects*

Similarly of interest are secondary mechanisms that either increase mutation rates or the number of dividing cells. An increase in mutations might be due to cytotoxic exposures causing regenerative proliferation or mitogenic influences, either of which could result in clonal expansion of initiated cells (Cohen and Ellwein, 1990). An agent might interfere with the enzymes involved in DNA repair and recombination (Barrett and Lee, 1992). Also, programmed cell death (apoptosis) can potentially be blocked by an agent, thereby permitting replication of damaged cells. For example, peroxisome proliferators may act by suppressing apoptosis pathways (Shulte-Hermann et al., 1993; Bayly et al., 1994). An agent may also generate reactive oxygen species that produce oxidative damage to DNA and other important

1 macromolecules that become important elements of the carcinogenic process (Kehrer, 1993;
2 Clayson et al., 1994; Chang et al., 1988). Damage to certain critical DNA repair genes or other
3 genes (e.g., the p53 gene) may result in genomic instability, which predisposes cells to further
4 genetic alterations and increases the probability of neoplastic progression independent of any
5 exogenous agent (Harris and Hollstein, 1993; Levine, 1994).

6 The loss or gain of chromosomes (i.e., aneuploidy) is an effect that can result in genomic
7 instability (Fearon and Vogelstein, 1990; Cavenee et al., 1986). Although the relationship
8 between induced aneuploidy and carcinogenesis is not completely established, several
9 carcinogens have been shown to induce aneuploidy (Gibson et al., 1995; Barrett, 1992). Agents
10 that cause aneuploidy interfere with the normal process of chromosome segregation and lead to
11 chromosomal losses, gains, or aberrations by interacting with the proteins (e.g., microtubules)
12 needed for chromosome movement.

14 **2.3.5.3. *Nonmutagenic and Other Effects***

15 A failure to detect DNA damage and mutation induction in several test systems suggests
16 that a carcinogenic agent may act by another mode of action.

17 It is possible for an agent to alter gene expression (transcriptional, translational, or post-
18 translational modifications) by means not involving mutations (Barrett, 1995). For example,
19 perturbation of DNA methylation patterns may cause effects that contribute to carcinogenesis
20 (Jones, 1986; Goodman and Counts, 1993; Holliday, 1987). Overexpression of genes by
21 amplification has been observed in certain tumors (Vainio et al., 1992). Other mechanisms may
22 involve cellular reprogramming through hormonal mechanisms or receptor-mediated
23 mechanisms (Ashby et al., 1994; Barrett, 1992).

24 Gap-junctional intercellular communication is widely believed to play a role in tissue and
25 organ development and in the maintenance of a normal cellular phenotype within tissues. A
26 growing body of evidence suggests that chemical interference with gap-junctional intercellular
27 communication is a contributing factor in tumor development; many carcinogens have been
28 shown to inhibit this communication. Thus, such information may provide useful mechanistic
29 data in evaluating cancer hazard (Swierenga and Yamasaki, 1992; Yamasaki, 1995).

Both cell death and cell proliferation are mandatory for the maintenance of homeostasis in normal tissue. The balance between the two directly affects the survival and growth of initiated cells, as well as preneoplastic and tumor cell populations (i.e., increase in cell proliferation or decrease in cell death) (Bellamy et al., 1995; Cohen and Ellwein, 1990, 1991; Cohen et al., 1991). In studies of proliferative effects, distinctions should be made between mitogenesis and regenerative proliferation (Cohen and Ellwein, 1990, 1991; Cohen et al., 1991). In applying information from studies on cell proliferation and apoptosis to risk assessment, it is important to identify the tissues and target cells involved, to measure effects in both normal and neoplastic tissue, to distinguish between apoptosis and necrosis, and to determine the dose that affects these processes.

2.3.5.4. Criteria for Judging Mode of Action

Criteria that are applicable for judging the adequacy of mechanistically based data include the following:

- mechanistic relevance of the data to carcinogenicity,
- number of studies of each endpoint,
- consistency of results in different test systems and different species,
- similar dose response relationships for tumor and mode of action-related effects,
- tests conducted in accordance with generally accepted protocols, and
- degree of consensus and general acceptance among scientists regarding interpretation of the significance and specificity of the tests.

Although important information can be gained from in vitro test systems, a higher level of confidence is generally given to data that are derived from in vivo systems, particularly those results that show a site concordance with the tumor data.

2.4. BIOMARKER INFORMATION

Various endpoints can serve as biological markers of events in biological systems or samples. In some cases, these molecular or cellular effects (e.g., DNA or protein adducts, mutation, chromosomal aberrations, levels of thyroid stimulating hormone) can be measured in

1 blood, body fluids, cells and tissues to serve as biomarkers of exposure in both animals and
2 humans (Callemén et al., 1978; Birner et al., 1990). As such, they can do the following:

- 3 ● act as an internal surrogate measure of chemical dose, representing as appropriate,
4 either recent (e.g., serum concentration) or accumulated (e.g., hemoglobin adducts)
5 exposure,
- 6 ● help identify doses at which elements of the carcinogenic process are operating,
- 7 ● aid in interspecies extrapolations when data are available from both experimental
8 animal and human cells, and
- 9 ● under certain circumstances, provide insights into the possible shape of the dose
10 response curve below levels where tumor incidences are observed (e.g., Choy, 1993).

11 Genetic and other findings (like changes in proto-oncogenes and tumor suppressor genes
12 in preneoplastic and neoplastic tissue or possibly measures of endocrine disruption) can indicate
13 the potential for disease and as such serve as biomarkers of effect. They, too, can be used in
14 different ways:

- 15 ● The spectrum of genetic changes in proliferative lesions and tumors following
16 chemical administration to experimental animals can be determined and compared
17 with those in spontaneous tumors in control animals, in animals exposed to other
18 agents of varying structural and functional activities, and in persons exposed to the
19 agent under study.
- 20 ● They may provide a linkage to tumor response.
- 21 ● They may help to identify subpopulations of individuals who may be at an elevated
22 risk for cancer, e.g., cytochrome P450 2D6/debrisoquine sensitivity for lung cancer
23 (Caporaso et al., 1989) or inherited colon cancer syndromes (Kinzler et al., 1991;
24 Peltomäki et al., 1993).
- 25 ● As with biomarkers of exposure, it may be justified in some cases to use these
26 endpoints for dose response assessment or to provide insight into the potential shape
27 of the dose response curve at doses below those at which tumors are induced
28 experimentally.

29 In applying biomarker data to cancer assessment (particularly assessments based on
30 epidemiologic data), one should consider the following:

- routes of exposure
- exposure to mixtures
- time after exposure
- sensitivity and specificity of biomarkers
- dose response relationships.

2.5. MODE OF ACTION--IMPLICATIONS FOR HAZARD CHARACTERIZATION AND DOSE RESPONSE

The interaction of the biology of the organism and the chemical properties of the agent determine whether there is an adverse effect. Thus, mode of action analysis is based on physical, chemical, and biological information that helps to explain critical events in an agent's influence on development of tumors. The entire range of information developed in the assessment is reviewed to arrive at a reasoned judgment. An agent may work by more than one mode of action both at different sites and at the same tumor site. It is felt that at least some information bearing on mode of action (e.g., SAR, screening tests for mutagenicity) is present for most agents undergoing assessment of carcinogenicity, even though certainty about exact molecular mechanisms may be rare.

Inputs to mode of action analysis include tumor data in humans, animals, and among structural analogues as well as the other key data. The more complete the data package and generic knowledge about a given mode of action, the more confidence one has and the more one can replace or refine default science policy positions with relevant information. Making reasoned judgments is generally based on a data-rich source of chemical, chemical class, and tumor type-specific information. Many times there will be conflicting data and gaps in the information base; one must carefully evaluate these uncertainties before reaching any conclusion.

Some of the questions that need to be addressed include the following:

- Has a body of data been developed on the agent that fits with a generally accepted mode of action?
- Has the mode of action been published and gained general scientific acceptance through peer-reviewed research or is it still speculative?

- Is the mode of action consistent with generally agreed-upon principles and understanding of carcinogenesis?
- Is the mode of action reasonably anticipated or assumed, in the absence of specific data, to operate in humans? How is this question influenced by information on comparative uptake, metabolism, and excretion patterns across animals and humans?
- Do humans appear to be more or less sensitive to the mode of action than are animals?
- Does the agent affect DNA, directly or indirectly?
- Are there important determinants in carcinogenicity other than effects on DNA, such as changes in cell proliferation, apoptosis, gene expression, immune surveillance, or other influences?

In making decisions about potential modes of action and the relevance of animal tumor findings to humans (Ashby et al., 1990), very often the results of chronic animal studies may give important clues. Some of the important factors to review include the following:

- tumor types, e.g., those responsive to endocrine influence, those produced by reactive carcinogens (Ashby and Tennant, 1991),
- number of tumor sites, sexes, studies, and species affected or unaffected (Tennant, 1993),
- influence of route of exposure; spectrum of tumors; local or systemic sites,
- target organ or system toxicity, e.g., urinary chemical changes associated with stone formation, effects on immune surveillance,
- presence of proliferative lesions, e.g., hepatic foci, hyperplasias,
- progression of lesions from preneoplastic to benign to malignant with dose and time,
- ratio of malignant to benign tumors as a function of dose and time,
- time of appearance of tumors after commencing exposure,
- tumors invading locally, metastasizing, producing death,
- tumors at sites in laboratory animals with high or low spontaneous historical incidence,
- biomarkers in tumor cells, both induced and spontaneous, e.g., DNA or protein adducts, mutation spectra, chromosome changes, oncogene activation, and

- shape of the dose response in the range of tumor observation, e.g., linear vs. profound change in slope.

Some of the myriad of ways that information from chronic animal studies influences mode of action judgments include the following. Multisite and multispecies tumor effects are often associated with mutagenic agents. Tumors restricted to one sex/species may suggest an influence restricted to gender, strain, or species. Late onset of tumors that are primarily benign or are at sites with a high historical background incidence or show reversal of lesions on cessation of exposure may point to a growth-promoting mode of action. The possibility that an agent may act differently in different tissues or have more than one mode of action in a single tissue must also be kept in mind.

Simple knowledge of sites of tumor increase in rodent studies can give preliminary clues as to mode of action. Experience at the National Toxicology Program (NTP) indicates that substances that are DNA reactive and produce gene mutations may be unique in producing tumors in certain anatomical sites, while tumors at other sites may arise from both mutagenic or nonmutagenic influences (Ashby and Tennant, 1991; Huff et al., 1991).

Effects on tumor sites in rodents and other mode of action information has been explored for certain agents (Alison et al., 1994; Clayson, 1989; ECETOC, 1991; MacDonald et al., 1994; McClain, 1994; Tischer et al., 1991; ILSI, 1995; Cohen and Ellwein, 1991; FASEB, 1994; Havu et al., 1990; U.S. EPA, 1991; Li et al., 1987; Grasso and Hinton, 1991; Larson et al., 1994; IARC, 1990; Jack et al., 1983; Stitzel et al., 1989; Ingram and Grasso, 1991; Bus and Popp, 1987; Prahalada et al., 1994; Yamada et al., 1994; Hill et al., 1989; Burek et al., 1988).

The selection of a dose response extrapolation procedure for cancer risk estimation considers mode of action information. When information is extensive and there is considerable certainty in a given mode of action, a biologically based or case-specific model that incorporates data on processes involved is preferred. Obviously, use of such a model requires the existence of substantial data on component parameters of the mode of action, and judgments on its applicability must be made on a case-by-case basis.

In the absence of information to develop a biologically based or case-specific model, understanding of mode of action should be employed to the extent possible in deciding upon one of three science policy defaults: low-dose linear extrapolation, nonlinear, and both procedures.

1 The overall choice of the default(s) depends upon weighing the various inputs and deciding
2 which best reflect the mode of action understanding. A rationale accompanies whichever default
3 or defaults are chosen.

4 A default assumption of linearity is appropriate when the evidence supports a mode of
5 action of gene mutation due to DNA reactivity or supports another mode of action that is
6 anticipated to be linear. Other elements of empirical data may also support an inference of
7 linearity, e.g., the background of human exposure to an agent might be such that added human
8 exposure is on the linear part of a dose response curve that is sublinear overall. The default
9 assumption of linearity is also appropriate as the ultimate default when evidence shows no DNA
10 reactivity or other support for linearity, but neither is it sufficient evidence of a nonlinear mode
11 of action to support a nonlinear procedure.

12 A default assumption of nonlinearity is appropriate when there is no evidence for
13 linearity and sufficient evidence to support an assumption of nonlinearity and a nonlinear
14 procedure. The mode of action may lead to a dose response relationship that is nonlinear, with
15 response falling much more quickly than linearly with dose, or being most influenced by
16 individual differences in sensitivity. Alternatively, the mode of action may theoretically have a
17 threshold, e.g., the carcinogenicity may be a secondary effect of toxicity that is itself a threshold
18 phenomenon.

19 Both linear and nonlinear procedures may be used in particular cases. If a mode of
20 action analysis finds substantial support for differing modes of action for different tumor sites,
21 an appropriate procedure is used for each. Both procedures may also be appropriate to discuss
22 implications of complex dose response relationships. For example, if it is apparent that an agent
23 is both DNA reactive and is highly active as a promotor at high doses, and there are insufficient
24 data for modeling, both linear and nonlinear default procedures may be needed to decouple and
25 consider the contribution of both phenomena.

2.6. WEIGHT OF EVIDENCE EVALUATION FOR POTENTIAL HUMAN CARCINOGENICITY

A weight of evidence evaluation is a collective evaluation of all pertinent information so that the full impact of biological plausibility and coherence are adequately considered. Identification and characterization of human carcinogenicity is based on human and experimental data, the nature, advantages and limitations of which have been discussed in the preceding sections.

The subsequent sections outline: (1) the basics of weighing individual lines of evidence and combining the entire body of evidence to make an informed judgment, (2) classification descriptors of cancer hazard, and (3) some case study examples to illustrate how the principles of guidance can be applied to arrive at a classification.

2.6.1. Weight of Evidence Analysis

Judgment about the weight of evidence involves considerations of the quality and adequacy of data and consistency of responses induced by the agent in question. The weight of evidence judgment requires combined input of relevant disciplines. Initial views of one kind of evidence may change significantly when other information is brought to the interpretation. For example, a positive animal carcinogenicity finding may be diminished by other key data; a weak association in epidemiologic studies may be bolstered by consideration of other key data and animal findings. Factors typically considered are illustrated in figures below. Generally, no single weighing factor on either side determines the overall weight. The factors are not scored mechanically by adding pluses and minuses; they are judged in combination.

Human Evidence

Analyzing the contribution of evidence from a body of human data requires examining available studies and weighing them in the context of well-accepted criteria for causation (see section 2.2.1). A judgment is made about how closely they satisfy these criteria, individually and jointly, and how far they deviate from them. Existence of temporal relationships, consistent results in independent studies, strong association, reliable exposure data, presence of dose-

1 related responses, freedom from biases and confounding factors, and high level of statistical
2 significance are among the factors leading to increased confidence in a conclusion of causality.

3 Generally, the weight of human evidence increases with the number of adequate studies
4 that show comparable results on populations exposed to the same agent under different
5 conditions. The analysis takes into account all studies of high quality, whether showing positive
6 associations or null results, or even protective effects. In weighing positive studies against null
7 studies, possible reasons for inconsistent results should be sought, and results of studies that are
8 judged to be of high quality are given more weight than those from studies judged to be
9 methodologically less sound. See figure 2-1.

1
2

Human Evidence Factors

Increase Weight	Decrease Weight
Number of independent studies with consistent results	Few studies Equally well designed and conducted studies with null results
Most causal criteria satisfied:	
Temporal relationship	
Strong association	
Reliable exposure data	Few causal criteria satisfied
Dose response relationship	
Freedom from bias and confounding	
Biological plausibility	
High statistical significance	

Figure 2-1. Factors for Weighing Human Evidence

Generally, no single factor is determinative. For example, the strength of association is one of the causal criteria. A strong association (i.e., a large relative risk) is more likely to indicate causality than a weak association. However, finding of a large excess risk in a single study must be balanced against the lack of consistency as reflected by null results from other equally well designed and well conducted studies. In this situation, the positive association of a single study may either suggest the presence of chance, bias or confounding, or reflect different exposure conditions. On the other hand, evidence of weak but consistent associations across several studies suggests either causality or the same confounder may be operating in all of these studies.

Animal Evidence

Evidence from long-term or other carcinogenicity studies in laboratory animals constitutes the second major class of information bearing on carcinogenicity. See figure 2-2. As discussed in section 2.2.2., each relevant study must be reviewed and evaluated as to its adequacy of design and conduct as well as the statistical significance and biological relevance of its findings. Factors that usually increase confidence in the predictivity of animal findings are those of (1) multiplicity of observations in independent studies; (2) severity of lesions, latency, and lesion progression; (3) consistency in observations.

Animal Evidence Factors	
Increase Weight	Decrease Weight
Number of independent studies with consistent results	Single study Inconsistent results
Same site across species, structural analogues	
Multiple observations Species Sites Sexes	Single site/species/sex
Severity and progression of lesions Early in life tumors/malignancy Dose response relationships Lesion progression Uncommon tumor	Benign tumors only High background of incidence tumors
Route of administration like human exposure	Route of administration unlike human exposure

Figure 2-2. Factors for Weighing Animal Evidence

Other Key Evidence

Additional information bearing on the qualitative assessment of carcinogenic potential may be gained from comparative pharmacokinetic and metabolism studies, genetic toxicity studies, SAR analysis, and other studies of an agent's properties. See figure 2-3. Information from these studies helps to elucidate potential modes of action and biological fate and disposition. The knowledge gained supports interpretation of cancer studies in humans and animals and provides a separate source of information about carcinogenic potential.

Other Key Evidence Factors

Increase Weight	Decrease Weight
A rich set of other key data are available	Few or poor data
Physicochemical data	or
Data indicate reactivity with macromolecules	Inadequate data necessitate use of default assumptions
Structure activity relationships support hazard potential	or
Comparable metabolism and toxicokinetics between species	Data show that animal findings are not relevant to humans
Toxicological and human clinical data support tumor findings	
Biomarker data support attribution of effects to agent	
Mode of action data support causal interpretation of human evidence or relevance of animal evidence	

a

Figure 2-3. Factors for Weighing Other Key Evidence

Totality of Evidence

In reaching a view of the entire weight of evidence, all data and inferences are merged. Figure 2-4 indicates the generalities. In fact, possible weights of evidence span a broad continuum that cannot be capsulized. Most of the time the data in various lines of evidence fall in the middle of the weights represented in the four figures in this section.

Totality of Evidence Factors

Increase Weight	Decrease Weight
Evidence of human causality	Data not available or do not show causality
Evidence of animal effects relevant to humans	Data not available or not relevant
Coherent inferences	Conflicting data
Comparable metabolism and toxicokinetics between species	Metabolism and toxicokinetics not comparable
Mode of action comparable across species	Mode of action not comparable across species

Figure 2-4. Factors for Weighing Totality of Evidence

The following section and the weight of evidence narrative discussed in 2.7.2. provide a way to state a conclusion and capture this complexity in a consistent way.

2.6.2. Descriptors for Classifying Weight of Evidence

Hazard classification uses three categories of descriptors for human carcinogenic potential: "known/likely," "cannot be determined," and "not likely." Each category has associated subdescriptors to further define the conclusion. The descriptors are not meant to replace an explanation of the nuances of the biological evidence, but rather to summarize it. Each category spans a wide variety of potential data sets and weights of evidence. There will always be gray areas, gradations, and borderline cases. That is why the descriptors are presented only in the context of a weight of evidence narrative whose format is given in section 2.7.2. Using them within a narrative preserves and presents the complexity that is an essential part of the hazard classification. Applying a descriptor is a matter of judgment and cannot be reduced to a formula. Risk managers should consider the entire range of information included in the narrative rather than focusing simply on the descriptor.

A single agent may be categorized in more than one way if, for instance, the agent is likely to be carcinogenic by one route of exposure but not by another (section 2.3.3).

The descriptors and subdescriptors are standardized and are to be used consistently from case to case. The discussions below explain descriptors and subdescriptors which appear in italics, and along with Appendix A and section 2.6.3, illustrate their use.

"Known/Likely"

This category of descriptors is appropriate when the available tumor effects and other key data are adequate to convincingly demonstrate carcinogenic potential for humans; it includes:

- C agents *known* to be carcinogenic in humans based on either epidemiologic evidence or a combination of epidemiologic and experimental evidence, demonstrating causality between human exposure and cancer,

1 C agents that should be treated *as if* they were *known* human carcinogens, based on a
2 combination of epidemiologic data showing a plausible causal association (not
3 demonstrating it definitively) and strong experimental evidence.

4 C agents that are *likely* to produce cancer in humans due to the production or anticipated
5 production of tumors by modes of action that are relevant or assumed to be relevant
6 to human carcinogenicity.

7 Modifying descriptors for particularly high or low ranking in the "known/likely" group can be
8 applied based on scientific judgment and experience and are as follows:

9 C agents that are *likely* to produce cancer in humans based on data that are at the *high*
10 *end* of the weights of evidence typical of this group,

11 C agents that are *likely* to produce cancer in humans based on data that are at the *low*
12 *end* of the weights of evidence typical of this group.

13 14 ***"Cannot Be Determined"***

15 This category of descriptors is appropriate when available tumor effects or other key data
16 are suggestive or conflicting or limited in quantity and, thus, are not adequate to convincingly
17 demonstrate carcinogenic potential for humans. In general, further agent specific and generic
18 research and testing are needed to be able to describe human carcinogenic potential. The
19 descriptor *cannot be determined* is used with a subdescriptor that captures the rationale:

20 C agents whose carcinogenic potential *cannot be determined*, but for which there is
21 *suggestive* evidence that raises concern for carcinogenic effects,

22 C agents whose carcinogenic potential *cannot be determined* because the existing
23 evidence is composed of *conflicting data* (e.g., some evidence is suggestive of
24 carcinogenic effects, but other equally pertinent evidence does not confirm any
25 concern),

26 C agents whose carcinogenic potential *cannot be determined* because there are
27 *inadequate data* to perform an assessment,

28 C agents whose carcinogenic potential *cannot be determined* because *no data* are
29 available to perform an assessment.

"Not Likely"

This is the appropriate descriptor when experimental evidence is satisfactory for deciding that there is no basis for human hazard concern, as follows (in the absence of human data suggesting a potential for cancer effects):

- C agents *not likely* to be carcinogenic to humans because they have been evaluated in at least two well conducted studies in two appropriate animal species without demonstrating carcinogenic effects,
- C agents *not likely* to be carcinogenic to humans because they have been appropriately evaluated in animals and show only carcinogenic effects that have been shown not to be relevant to humans (e.g., showing only effects in the male rat kidney due to accumulation of alpha_{2u}-globulin),
- C agents *not likely* to be carcinogenic to humans when carcinogenicity is dose or route dependent. For instance, *not likely* below a certain dose range (categorized as *likely* above that range) or *not likely* by a certain route of exposure (may be categorized as *likely* by another route of exposure). To qualify, agents will have been appropriately evaluated in animal studies and the only effects show a dose range or route limitation or a route limitation is otherwise shown by empirical data.
- C agents not likely to be carcinogenic to humans based on extensive human experience that demonstrates lack of effect (e.g., phenobarbital).

2.6.3. Case Study Examples

This section provides examples of substances that fit the three broad categories described above. These examples are based on available information about real substances and are selected to illustrate the principles for weight-of-evidence evaluation and the application of the classification scheme.

These case studies show the interplay of differing lines of evidence in making a conclusion. Some particularly illustrate the role that “other key data” can play in conclusions.

1 ***Example 1: "Known Human Carcinogen"--Route-Dependent/Linear Extrapolation***

2 Human Data

3 Substance 1 is an aluminosilicate mineral that exists in nature with a fibrous habit.
4 Several descriptive epidemiologic studies have demonstrated very high mortality from malignant
5 mesothelioma, mainly of the pleura, in three villages in Turkey, where there was a
6 contamination of this mineral and where exposure had occurred from birth. Both sexes were
7 equally affected and at an unusually young age.

8
9 Animal Data

10 Substance 1 has been studied in a single long-term inhalation study in rats at one
11 exposure concentration that showed an extremely high incidence of pleural mesothelioma (98%
12 in treated animals versus 0% in concurrent controls). This is a rare malignant tumor in the rat
13 and the onset of tumors occurred at a very early age (as early as 1 year of age). Several studies
14 involving injection into the body cavities of rats or mice (i.e., pleural or peritoneal cavities) also
15 produced high incidences of pleural or peritoneal mesotheliomas. No information is available on
16 the carcinogenic potential of substance 1 in laboratory animals via oral and dermal exposures.

17
18 Other Key Data

19 Information on the physical and chemical properties of substance 1 indicates that it is
20 highly respirable to humans and laboratory rodents. It is highly insoluble and is not likely to be
21 readily degraded in biological fluid.

22 No information is available on the deposition, translocation, retention, lung clearance,
23 and excretion of the substance after inhalation exposure or ingestion. Lung burden studies have
24 shown the presence of elevated levels of the substance in lung tissue samples of human cases of
25 pleural mesotheliomas from contaminated villages compared with control villages.

26 No data are available on genetic or related effects in humans. The substance has been
27 shown to induce unscheduled DNA synthesis in human cells in vitro and transformation and
28 unscheduled DNA synthesis in mouse cells.

29 The mechanisms by which this substance causes cancer in humans and animals are not
30 understood, but appear to be related to its unique physical, chemical, and surface properties. Its

1 fiber morphology is similar to a known group of naturally occurring silicate minerals that have
2 been known to cause respiratory cancers (including pleural mesothelioma) from inhalation
3 exposure and genetic changes in humans.
4

5 Evaluation

6 Human evidence is judged to establish a causal link between exposure to substance 1 and
7 human cancer. Even though the human evidence does not satisfy all criteria for causality, this
8 judgment is based on a number of unusual observations: large magnitude of the association,
9 specificity of the association, demonstration of environmental exposure, biological plausibility,
10 and coherence based on the entire body of knowledge of the etiology of mesothelioma.

11 Animal evidence demonstrates a causal relationship between exposure and cancer in
12 laboratory animals. Although available data are not optimal in terms of design (e.g., the use of
13 single dose, one sex only), the judgment is based on the unusual findings from the only
14 inhalation experiment in rats (i.e., induction of an uncommon tumor, an extremely high
15 incidence of malignant neoplasms, and onset of tumors at an early age). Additional evidence is
16 provided by consistent results from several injection studies showing an induction of the same
17 tumors by different modes of administration in more than one species.

18 Other key data, while limited, support the human and animal evidence of carcinogenicity.
19 It can be inferred from human and animal data that this substance is readily deposited in the
20 respiratory airways and deep lung and is retained for extended periods of time since first
21 exposure. Information on related fibrous substances indicates that the modes of action are likely
22 mediated by the physical and chemical characteristics of the substance (e.g., fiber shape, high
23 aspect ratio, a high degree of insolubility in lung tissues).

24 Insufficient data are available to evaluate the human carcinogenic potential of substance
25 1 by oral exposure. Even though there is no information on its carcinogenic potential via dermal
26 uptake, it is not expected to pose a carcinogenic hazard to humans by that route because it is
27 very insoluble and is not likely to penetrate the skin.
28

Conclusion

It is concluded that substance 1 is a *known human carcinogen by inhalation exposure*. The weight of evidence of human carcinogenicity is based on (a) exceptionally increased incidence of malignant mesothelioma in epidemiologic studies of environmentally exposed human populations; (b) significantly increased incidence of malignant mesothelioma in a single inhalation study in rats and in several injection studies in rats and mice; and (c) supporting information on related fibrous substances that are known to cause cancer via inhalation and genetic damage in exposed mammalian and human mesothelial cells. The human carcinogenic potential of substance 1 via oral exposure cannot be determined on the basis of insufficient data. It is not likely to pose a carcinogenic hazard to humans via dermal uptake because it is not anticipated to penetrate the skin.

The mode of action of this substance is not understood. In addition to this uncertainty, dose response information is lacking for both human and animal data. Epidemiologic studies contain observations of significant excess cancer risks at relatively low levels of environmental exposure. The use of *linear* extrapolation in a dose response relationship assessment is appropriate as a default since mode of action data are not available.

Example 2: "As If Known Human Carcinogen"--Any Exposure Conditions/Linear Extrapolation

Human Data

Substance 2 is an alkene oxide. Several cohort studies of workers using substance 2 as a sterilant have been conducted. In the largest and most informative study, mortality from lymphatic and hematopoietic cancer was marginally elevated, but a significant trend was found, especially for lymphatic leukemia and non-Hodgkin's lymphoma, in relation to estimated cumulative exposure to the substance. Nonsignificant excesses of lymphatic and hematopoietic cancer were found in three other smaller studies of sterilization personnel.

In one cohort study of chemical workers exposed to substance 2 and other agents, mortality rate from lymphatic and hematopoietic cancer was elevated, but the excess was confined to a small subgroup with only occasional low-level exposure to substance 2. Six other studies of chemical workers are considered more limited due to a smaller number of deaths.

Four studies found an excess of lymphatic and hematopoietic cancer (which were significant in two); no increase in mortality rate was observed in the other two studies.

Animal Data

Substance 2 was studied in an oral gavage study in rats. Treatment of substance 2 resulted in a dose-dependent increased incidence in forestomach tumors that were mainly squamous-cell carcinomas.

Substance 2 was also studied in two inhalation studies in mice and two inhalation studies in rats. In the first mouse study, dose-dependent increases in combined benign and malignant tumors at several tissue sites were induced in mice of both sexes (lung tumors and tumors of the Harderian gland in each sex, and uterine adenocarcinomas, mammary carcinomas, and malignant lymphomas in females). In a second study--a screening study for pulmonary tumors in mice--inhalation exposure to substance 2 resulted in a dose-dependent increase in lung tumors. In the two inhalation studies in rats, increased incidences of mononuclear-cell leukemia and brain tumors were induced in exposed animals of each sex; increased incidences of peritoneal tumors in the region of the testis and subcutaneous fibromas were induced in exposed male rats.

Substance 2 induced local sarcomas in mice following subcutaneous injection. No tumors were found in a limited skin painting study in mice.

Other Key Data

Substance 2 is a flammable gas at room temperature. The gaseous form is readily taken up in humans and rats, and in aqueous solution it can penetrate human skin. Studies in rats indicate that, once absorbed, substance 2 is uniformly distributed throughout the body. It is eliminated metabolically by hydrolysis and by conjugation with glutathione. The ability to form glutathione conjugate varies across animal species, with the rat being most active, followed by mice and rabbits.

Substance 2 is a directly acting alkylating agent. It has been shown to form adducts with hemoglobin in both humans and animals and with DNA in animals. The increased frequency of hemoglobin adducts, which have been used as markers of internal dose, has been found to correlate with the level and cumulative exposure to substance 2. Significant increases in

1 chromosomal aberrations and sister chromatid exchanges in peripheral lymphocytes and
2 induction of micronuclei in the bone marrow cells have been observed in exposed workers.

3 Substance 2 also induced chromosomal aberrations and sister chromatid exchanges in
4 peripheral lymphocytes of monkeys exposed in vivo. It also induced gene mutation, specific
5 locus mutation, sister chromatid exchanges, chromosomal aberrations, micronuclei, dominant
6 lethal mutations, and heritable translocation in rodents exposed in vivo. In human cells in vitro,
7 it induced sister chromatid exchanges, chromosomal aberrations, and unscheduled DNA
8 synthesis. Similar genetic and related effects were observed in rodent cells in vitro and in
9 nonmammalian systems.

10 11 Evaluation

12 Available epidemiologic studies, taken together, suggest that a causal association
13 between exposure to substance 2 and elevated risk of cancer is plausible. This judgment is based
14 on small but consistent excesses of lymphatic and hematopoietic cancer in the studies of
15 sterilization workers. Interpretation of studies of chemical workers is difficult because of
16 possible confounding exposures. Nevertheless, findings of elevated risks of cancer at similar
17 sites in chemical workers support the findings in studies of sterilization workers. Additional
18 support is provided by observations of DNA damage in the same tissue in which elevated cancer
19 was seen in exposed workers.

20 Extensive evidence indicates that substance 2 is carcinogenic to laboratory animals.
21 Positive results were consistently observed in all well-designed and well-conducted studies.
22 Substance 2 causes dose-related increased incidences of tumors at multiple tissue sites in rats and
23 mice of both sexes by two routes of exposure (oral and inhalation). The only dermal study that
24 yielded a nonpositive finding is considered of limited quality.

25 Other key data significantly add support to the potential carcinogenicity of substance 2.
26 There is strong evidence of heritable mutations of exposed rodents and mutagenicity and
27 clastogenicity both in vivo and in vitro. These findings are reinforced by observations of similar
28 genetic damage in exposed workers. Additional support is based on SAR analysis that indicates
29 that substance 2 is a highly DNA-reactive agent. Structurally related chemicals, i.e., low-
30 molecular-weight epoxides, also exhibit carcinogenic effects in laboratory animals.

1 Conclusion

2 Substance 2 should be considered *as if it were a known human carcinogen by all routes*
3 *of exposure*. The weight of evidence of human carcinogenicity is based on (a) consistent
4 evidence of carcinogenicity in rats and mice by oral and inhalation exposure; (b) epidemiologic
5 evidence suggestive of a causal association between exposure and elevated risk of lymphatic and
6 hematopoietic cancer; (c) evidence of genetic damage in blood lymphocytes and bone marrow
7 cells of exposed workers; (d) mutagenic effects in numerous in vivo and in vitro test systems; (e)
8 membership in a class of DNA-reactive compounds that have been shown to cause carcinogenic
9 and mutagenic effects in animals; and (f) ability to be absorbed by all routes of exposure,
10 followed by rapid distribution throughout the body.

11 Although the exact mechanisms of carcinogenic action of substance 2 are not completely
12 understood, available data strongly indicate a mutagenic mode of action. Linear extrapolation
13 should be assumed in dose response assessment.

14
15 ***Example 3: "Likely Human Carcinogen"--Any Exposure Conditions/Linear Extrapolation***

16 Human Data

17 Substance 3 is a brominated alkane. Three studies have investigated the cancer mortality
18 of workers exposed to this substance. No statistically significant increase in cancer at any site
19 was found in a study of production workers exposed to substance 3 and several other chemicals.
20 Elevated cancer mortality was reported in a much smaller study of production workers. An
21 excess of lymphoma was reported in grain workers who may have had exposure to substance 3
22 and other chemical compounds. These studies are considered inadequate due to their small
23 cohort size; lack of, or poorly characterized, exposure concentrations; or concurrent exposure of
24 the cohort to other potential or known carcinogens.

25
26 Animal Data

27 The potential carcinogenicity of substance 3 has been extensively studied in an oral
28 gavage study in rats and mice of both sexes, two inhalation studies of rats of different strains of
29 both sexes, an inhalation study in mice of both sexes, and a skin painting study in female mice.

1 In the oral study, increased incidences of squamous-cell carcinoma of the forestomach
2 were found in rats and mice of both sexes. Additionally, there were increased incidences of liver
3 carcinomas in female rats, hemangiosarcomas in male rats, and alveolar/bronchiolar adenoma of
4 the lung of male and female mice. Excessive toxicity and mortality were observed in the rat
5 study, especially in the high-dose groups, which resulted in early termination of study, and
6 similar time-weighted average doses for the high- and low-treatment groups.

7 In the first inhalation study in rats and mice, increased incidences of carcinomas and
8 adenocarcinomas of the nasal cavity and hemangiosarcoma of the spleen were found in exposed
9 animals of each species of both sexes. Treated female rats also showed increased incidences of
10 alveolar/bronchiolar carcinoma of the lung and mammary gland fibroadenomas. Treated male
11 rats showed an increased incidence of peritoneal mesothelioma. In the second inhalation study
12 in rats (single exposure only), significantly increased incidences of hemangiosarcoma of the
13 spleen and adrenal gland tumors were seen in exposed animals of both sexes. Additionally,
14 increased incidences of subcutaneous mesenchymal tumors and mammary gland tumors were
15 induced in exposed male and female rats, respectively.

16 Lifetime dermal application of substance 3 to female mice resulted in significantly
17 increased incidences of skin papillomas and lung tumors.

18 Several chemicals structurally related to substance 3 are also carcinogenic in rodents.
19 The spectrum of tumor responses induced by related substances was similar to those seen with
20 substance 3 (e.g., forestomach, mammary gland, lung tumors).

21 Other Key Data

22 Substance 3 exists as a liquid at room temperature and is readily absorbed by ingestion,
23 inhalation, and dermal contact. It is widely distributed in the body and is eliminated in the urine
24 mainly as metabolites (e.g., glutathione conjugate).

25 Substance 3 is not itself DNA-reactive, but is biotransformed to reactive metabolites as
26 inferred by findings of its covalent binding to DNA and induction of DNA strand breaks, both in
27 vivo and in vitro. Substance 3 has been shown to induce sister chromatid exchanges, mutations,
28 and unscheduled DNA synthesis in human and rodent cells in vitro. Reverse and forward
29 mutations have been consistently produced in bacterial assays and in vitro assays using
30

eukaryotic cells. Substance 3, however, did not induce dominant lethal mutations in mice or rats, or chromosomal aberrations or micronuclei in bone marrow cells of mice treated in vivo.

Evaluation

Available epidemiologic data are considered inadequate for an evaluation of a causal association of exposure to the substance and excess of cancer mortality due to major study limitations.

There is extensive evidence that substance 3 is carcinogenic in laboratory animals. Increased incidences of tumors at multiple sites have been observed in multiple studies in two species of both sexes with different routes of exposure. It induces tumors both at the site of entry (e.g., nasal tumors via inhalation, forestomach tumors by ingestion, skin tumor with dermal exposure) and at distal sites (e.g., mammary gland tumors). Additionally, it induced tumors at the same sites in both species and sexes via different routes of exposure (e.g., lung tumors). With the exception of the oral study in which the employed doses caused excessive toxicity and mortality, the other studies are considered adequately designed and well conducted. Overall, given the magnitude and extent of animal carcinogenic responses to substance 3, coupled with similar responses to structurally related substances, these animal findings are judged to be highly relevant and predictive of human responses.

Other key data, while not very extensive, are judged to be supportive of carcinogenic potential. Substance 3 has consistently been shown to be mutagenic in mammalian cells, including human cells, and nonmammalian cells; thus, mutation is likely a mode of action for its carcinogenic activity. However, the possible involvement of other modes of action has not been fully investigated. Furthermore, induction of genetic changes from in vivo exposure to substance 3 has not been demonstrated.

Conclusion

Substance 3 is *likely to be a human carcinogen by any route of exposure*. In comparison with other agents designated as likely human carcinogens, the overall weight of evidence for substance 3 puts it at the *high end* of the grouping.

1 The weight of evidence of human carcinogenicity is based on animal evidence and other
2 key evidence. Human data are inadequate for an evaluation of human carcinogenicity. The
3 overall weight of evidence is based on (a) extensive animal evidence showing induction of
4 increases of tumors at multiple sites in both sexes of two rodent species via three routes of
5 administration relevant to human exposure; (b) tumor data of structural analogues exhibiting
6 similar patterns of tumors in treated rodents; (c) in vitro evidence for mutagenic effects in
7 mammalian cells and nonmammalian systems; and (d) its ability to be absorbed by all routes of
8 exposure followed by rapid distribution throughout the body.

9 Some uncertainties are associated with the mechanisms of carcinogenicity of substance
10 3. Although there is considerable evidence indicating that mutagenic events could account for
11 carcinogenic effects, there is still a lack of adequate information on the mutagenicity of
12 substance 3 in vivo in animals or humans. Moreover, alternative modes of action have not been
13 explored. Nonetheless, available data indicate a likely mutagenic mode of action. Linear
14 extrapolation should be assumed in dose response assessment.

15
16 ***Example 4: "Likely Human Carcinogen"--All Routes/Linear and Nonlinear Extrapolation***

17 **Human Data**

18 Substance 4 is a chlorinated alkene solvent. Several cohort studies of dry cleaning and
19 laundry workers exposed to substance 4 and other solvents reported significant excesses of
20 mortality due to cancers of the lung, cervix, esophagus, kidney, bladder, lymphatic and
21 hematopoietic system, colon, or skin. No significant cancer risks were observed in a subcohort
22 of one these investigations of dry cleaning workers exposed mainly to substance 4. Possible
23 confounding factors such as smoking, alcohol consumption, or low socioeconomic status were
24 not considered in the analyses of these studies.

25 A large case-control study of bladder cancer did not show any clear association with dry
26 cleaning. Several case-control studies of liver cancer identified an increased risk of liver cancer
27 with occupational exposure to organic solvents. The specific solvents to which workers were
28 exposed and exposure levels were not identified.

1 Animal Data

2 The potential carcinogenicity of substance 4 has been investigated in two long-term
3 studies in rats and mice of both sexes by oral administration and inhalation.

4 Significant increases in hepatocellular carcinomas were induced in mice of both sexes
5 treated with substance 4 by oral gavage. No increases in tumor incidence were observed in
6 treated rats. Limitations in both experiments included control groups smaller than treated
7 groups, numerous dose adjustments during the study, and early mortality due to treatment-
8 related nephropathy.

9 In the inhalation study, there were significantly increased incidences of hepatocellular
10 adenoma and carcinoma in exposed mice of both sexes. In rats of both sexes, there were
11 marginally significant increased incidences of mononuclear cell leukemia (MCL) when
12 compared with concurrent controls. The incidences of MCL in control animals, however, were
13 higher than historical controls from the conducting laboratory. The tumor finding was also
14 judged to be biologically significant because the time to onset of tumor was decreased and the
15 disease was more severe in treated than in control animals. Low incidences of renal tubular cell
16 adenomas or adenocarcinomas were also observed in exposed male rats. The tumor incidences
17 were not statistically significant but there was a significant trend.

18
19 Other Key Data

20 Substance 4 has been shown to be readily and rapidly absorbed by inhalation and
21 ingestion in humans and laboratory animals. Absorption by dermal exposure is slow and
22 limited. Once absorbed, substance 4 is primarily distributed to and accumulated in adipose
23 tissue and the brain, kidney, and liver. A large percentage of substance 4 is eliminated
24 unchanged in exhaled air, with urinary excretion of metabolites comprising a much smaller
25 percentage. The absorption and distribution profiles of substance 4 are similar across species
26 including humans.

27 Two major metabolites (trichloroacetic acid (TCA), and trichloroethanol), which are
28 formed by a P-450-dependent mixed-function oxidase enzyme system, have been identified in
29 all studied species, including humans. There is suggestive evidence for the formation of an
30 epoxide intermediate based on the detection of two other metabolites (oxalic acid and

1 trichloroacetyl amide). In addition to oxidative metabolism, substance 4 also undergoes
2 conjugation with glutathione. Further metabolism by renal beta-lyases could lead to two minor
3 active metabolites (trichlorovinyl thiol and dichlorothiokente).

4 Toxicokinetic studies have shown that the enzymes responsible for the metabolism of
5 substance 4 can be saturated at high exposures. The glutathione pathway was found to be a
6 minor pathway at low doses, but more prevalent following saturation of the cytochrome P-450
7 pathway. Comparative in vitro studies indicate that mice have the greater capacity to metabolize
8 to TCA than rats and humans. Inhalation studies also indicate saturation of oxidative
9 metabolism of substance 4, which occurs at higher dose levels in mice than in rats and humans.
10 Based on these findings, it has been postulated that the species differences in the carcinogenicity
11 of substance 4 between rats and mice may be related to the differences in the metabolism to
12 TCA and glutathione conjugates.

13 Substance 4 is a member of the class of chlorinated organics that often cause liver and
14 kidney toxicity and carcinogenesis in rodents. Like many chlorinated organics, substance 4 itself
15 does not appear to be mutagenic. Substance 4 was generally negative in in vitro bacterial
16 systems and in vivo mammalian systems. However, a minor metabolite formed in the kidney by
17 the glutathione conjugation pathway has been found to be a strong mutagen.

18 The mechanisms of induced carcinogenic effects of substance 4 in rats and mice are not
19 completely understood. It has been postulated that mouse liver carcinogenesis is related to liver
20 peroxisomal proliferation and toxicity of the metabolite TCA. Information on whether or not
21 TCA induces peroxisomal proliferation in humans is not definitive. The induced renal tumors in
22 male rats may be related either to kidney toxicity or the activity of a mutagenic metabolite. The
23 mechanisms of increases in MCL in rats are not known.

24 25 Evaluation

26 Available epidemiologic studies, taken together, provide suggestive evidence of a
27 possible causal association between exposure to substance 4 and cancer incidence in the laundry
28 and dry cleaning industries. This is based on consistent findings of elevated cancer risks in
29 several studies of different populations of dry cleaning and laundry workers. However, each
30 individual study is compromised by a number of study deficiencies including small numbers of

1 cancers, confounding exposure to other solvents, and poor exposure characterization. Others
2 may interpret these findings collectively as inconclusive.

3 There is considerable evidence that substance 4 is carcinogenic to laboratory animals. It
4 induces tumors in mice of both sexes by oral and inhalation exposure and in rats of both sexes
5 via inhalation. However, due to incomplete understanding of the mode of mechanism of action,
6 the predictivity of animal responses to humans is uncertain.

7 Animal data of structurally related compounds showing common target organs of toxicity
8 and carcinogenic effects (but lack of mutagenic effects) provide additional support for the
9 carcinogenicity of substance 4. Comparative toxicokinetic and metabolism information indicates
10 that the mouse may be more susceptible to liver carcinogenesis than rats and humans. This may
11 indicate differences of the degree and extent of carcinogenic responses, but does not detract from
12 the qualitative weight of evidence of human carcinogenicity. The toxicokinetic information also
13 indicates that oral and inhalation are the major routes of human exposure.

14 15 Conclusion

16 Substance 4 is *likely to be carcinogenic to humans by all routes of exposure*. The weight
17 of evidence of human carcinogenicity is based on: (a) demonstrated evidence of carcinogenicity
18 in two rodent species of both sexes via two relevant routes of human exposure; (b) the
19 substance's similarity in structure to other chlorinated organics that are known to cause liver and
20 kidney toxicity and carcinogenesis in rodents; (c) suggestive evidence of a possible association
21 between exposure to the substance in the laundry and dry cleaning industries and increased
22 cancer incidence; and (d) human and animal data indicating that the substance is absorbed by all
23 routes of exposure.

24 In comparison with other agents designated as likely carcinogens, the overall weight of
25 evidence places it the lower end of the grouping. This is because there is a lack of good
26 evidence that observed excess cancer risk in exposed workers is due solely to substance 4.
27 Moreover, there is considerable scientific uncertainty about the human significance of certain
28 rodent tumors associated with substance 4 and related compounds. In this case, the human
29 relevance of the animal evidence of carcinogenicity relies on the default assumption.

Overall, there is not enough evidence to give high confidence in a conclusion about any single mode of action; it appears that more than one is plausible in different rodent tissues. Nevertheless, the lack of mutagenicity of substance 4 and its general growth-promoting effect on high background tumors as well as its toxicity toward mouse liver and rat kidney tissue support the view that the predominant mode is growth-promoting rather than mutagenic. A mutagenic contribution to carcinogenicity due to a metabolite cannot be ruled out. The dose response assessment should, therefore, adopt both default approaches, nonlinear and linear extrapolations. The latter approach is very conservative since it likely overestimates risk at low doses in this case, and is primarily useful for screening analyses.

***Example 5: "Likely/Not Likely Human Carcinogen"--Range of Dose
Limited, Margin-of-Exposure Extrapolation***

Human Data

Substance 5 is a metal-conjugated phosphonate. No human tumor or toxicity data exist on this chemical.

Animal Data

Substance 5 caused a statistically significant increase in the incidence of urinary bladder tumors in male, but not female, rats at 30,000 ppm (3%) in the diet in a long-term study. Some of these animals had accompanying urinary tract stones and toxicity. No bladder tumors or adverse urinary tract effects were seen in two lower dose groups (2,000 and 8,000 ppm) in the same study. A chronic dietary study in mice at doses comparable to those in the rat study showed no tumor response or urinary tract effects. A 2-year study in dogs at doses up to 40,000 ppm showed no adverse urinary tract effects.

Other Key Data

Subchronic dosing of rats confirmed that there was profound development of stones in the male bladder at doses comparable to those causing cancer in the chronic study, but not at lower doses. Sloughing of the epithelium of the urinary tract accompanied the stones.

1 There was a lack of mutagenicity relevant to carcinogenicity. In addition, there is
2 nothing about the chemical structure of substance 5 to indicate DNA-reactivity or
3 carcinogenicity.

4 Substance 5 is composed of a metal, ethanol, and a simple phosphorus-oxygen-
5 containing component. The metal is not absorbed from the gut, whereas the other two
6 components are absorbed. At high doses, ethanol is metabolized to carbon dioxide, which makes
7 the urine more acidic; the phosphorus level in the blood is increased and calcium in the urine is
8 increased. Chronic testing of the phosphorus-oxygen-containing component alone in rats did not
9 show any tumors or adverse effects on the urinary tract.

10 Because substance 5 is a metal complex, it is not likely to be readily absorbed from the
11 skin.

12 13 Evaluation

14 Substance 5 produced cancer of the bladder and urinary tract toxicity in male, but not
15 female rats and mice, and dogs failed to show the toxicity noted in male rats. The mode of
16 action developed from the other key data to account for the toxicity and tumors in the male rats
17 is the production of bladder stones. At high but not lower subchronic doses in the male rat,
18 substance 5 leads to elevated blood phosphorus levels; the body responds by releasing excess
19 calcium into the urine. The calcium and phosphorus combine in the urine and precipitate into
20 multiple stones in the bladder. The stones are very irritating to the bladder; the bladder lining is
21 eroded, and cell proliferation occurs to compensate for the loss of the lining. Cell layers pile up,
22 and finally, tumors develop. Stone formation does not involve the chemical per se but is
23 secondary to the effects of its constituents on the blood and, ultimately, the urine. Bladder
24 stones, regardless of their cause, commonly produce bladder tumors in rodents, especially the
25 male rat.

26 27 Conclusion

28 Substance 5, a metal aliphatic phosphonate, is *likely* to be carcinogenic to humans only
29 under high-exposure conditions following *oral and inhalation exposure* that lead to bladder
30 stone formation, but is *not likely* to be carcinogenic under low-exposure conditions. It is *not*

likely to be a human carcinogen via the dermal route, given that the compound is a metal conjugate that is readily ionized and its dermal absorption is not anticipated. The weight of evidence is based on (a) bladder tumors only in male rats; (b) the absence of tumors at any other site in rats or mice; (c) the formation of calcium-phosphorus-containing bladder stones in male rats at high, but not low, exposures that erode bladder epithelium and result in profound increases in cell proliferation and cancer; and (d) the absence of structural alerts or mutagenic activity.

There is a strong mode of action basis for the requirements of (a) high doses of substance 5, (b) which lead to excess calcium and increased acidity in the urine, (c) which result in the precipitation of stones and (d) the necessity of stones for toxic effects and tumor hazard potential. Lower doses fail to perturb urinary constituents, lead to stones, produce toxicity, or give rise to tumors. Therefore, dose response assessment should assume nonlinearity.

A major uncertainty is whether the profound effects of substance 5 may be unique to the rat. Even if substance 5 produced stones in humans, there is only limited evidence that humans with bladder stones develop cancer. Most often human bladder stones are either passed in the urine or lead to symptoms resulting in their removal. However, since one cannot totally dismiss the male rat findings, some hazard potential may exist in humans following intense exposures. Only fundamental research could illuminate this uncertainty.

Example 6: "Cannot Be Determined"--Suggestive Evidence

Human Data

Substance 6 is an unsaturated aldehyde. In a cohort study of workers in a chemical plant exposed to a mixture of chemicals with substance 6 as a minor component, an elevated risk of cancer than was expected was reported. This study is considered inadequate because of multiple exposures, small cohort, and poor exposure characterization.

Animal Data

Substance 6 was tested for potential carcinogenicity in a drinking water study in rats, an inhalation study in hamsters, and a skin painting study in mice. No significant increases in tumors were observed in male rats treated with substance 6 at three dose levels in drinking

1 water. However, a significant increase of adrenal cortical adenomas was found in the only
2 treated female dose group administered a dose equivalent to the high dose of males. This study
3 used a small number of animals (20 per dose group).

4 No significant finding was detected in the inhalation study in hamsters. This study is
5 inadequate due to the use of too few animals, short duration of exposure, and inappropriate dose
6 selection (use of a single exposure that was excessively toxic as reflected by high mortality).

7 No increase in tumors was induced in the skin painting study in mice. This study is of
8 inadequate design for carcinogenicity evaluation because of several deficiencies: small number
9 of animals, short duration of exposure, lack of reporting about the sex and age of animals, and
10 purity of test material.

11 Substance 6 is structurally related to lowmolecularweight aldehydes that generally
12 exhibit carcinogenic effects in the respiratory tracts of laboratory animals via inhalation
13 exposure. Three skin painting studies in mice and two subcutaneous injection studies of rats and
14 mice were conducted to evaluate the carcinogenic potential of a possible metabolite of substance
15 6 (identified in vitro). Increased incidences of either benign or combined benign and malignant
16 skin tumors were found in the dermal studies. In the injection studies of rats and mice, increased
17 incidences of local sarcomas or squamous cell carcinoma were found at the sites of injection.
18 All of these studies are limited by the small number of test animals, the lack of characterization
19 of test material, and the use of single doses.

20 21 Other Key Data

22 Substance 6 is a flammable liquid at room temperature. Limited information on its
23 toxicokinetics indicates that it can be absorbed by all routes of exposure. It is eliminated in the
24 urine mainly as glutathione conjugates. Substance 6 is metabolized in vitro by rat liver and lung
25 microsomal preparations to a dihydroxylated aldehyde.

26 No data were available on the genetic and related effects of substance 6 in humans. It did
27 not induce dominant lethal mutations in mice. It induced sister chromatid exchanges in rodent
28 cells in vitro. The mutagenicity of substance 6 is equivocal in bacteria. It did not induce DNA
29 damage or mutations in fungi.

Evaluation

Available human data are judged inadequate for an evaluation of any causal relationship between exposure to substance 6 and human cancer.

The carcinogenic potential of substance 6 has not been adequately studied in laboratory animals due to serious deficiencies in study design, especially the inhalation and dermal studies. There is some evidence of carcinogenicity in the drinking water study in female rats. However, the significance and predictivity of that study to human response are uncertain since the finding is limited to occurrence of benign tumors, one sex, and at the high dose only. Additional suggestion for animal carcinogenicity comes from observation that a possible metabolite is carcinogenic at the site of administration. This metabolite, however, has not been studied in vivo. Overall, the animal evidence is judged to be suggestive for human carcinogenicity.

Other key data, taken together, do not add significantly to the overall weight of evidence of carcinogenicity. SAR analysis indicates that substance 6 would be DNA-reactive. However, mutagenicity data are inconclusive. Limited in vivo data do not support a mutagenic effect. While there is some evidence of DNA damage in rodent cells in vitro, there is either equivocal or no evidence of mutagenicity in nonmammalian systems.

Conclusion

The human carcinogenicity potential of substance 6 cannot be determined on the basis of available information. Both human and animal data are judged inadequate for an evaluation. There is evidence suggestive of potential carcinogenicity on the basis of limited animal findings and SAR considerations. Data are not sufficient to judge whether there is a mode of carcinogenic action. Additional studies are needed for a full evaluation of the potential carcinogenicity of substance 6. Hence, dose response assessment is not appropriate.

Example 7: "Not Likely Human Carcinogen"--Appropriately Studied Chemical in Animals Without Tumor Effects

Human Data

Substance 7, a plant extract, has not been studied for its toxic or carcinogenic potential in humans.

Animal Data

Substance 7 has been studied in four chronic studies in three rodent species. In a feeding study in rats, males showed a nonsignificant increase in benign tumors of the parathyroid gland in the high-dose group, where the incidence in concurrent controls greatly exceeded the historical control range. Females demonstrated a significant increase in various subcutaneous tumors in the low-dose group, but findings were not confirmed in the high-dose group, and there was no dose response relationship. These effects were considered as not adding to the evidence of carcinogenicity. No tumor increases were noted in a second adequate feeding study in male and female rats. In a mouse feeding study, no tumor increases were noted in dosed animals. There was some question as to the adequacy of the dosing; however it was noted that in the mouse 90-d subchronic study, a dose of twice the high dose in the chronic study led to significant decrements in body weight. In a hamster study there were no significant increases in tumors at any site. No structural analogues of substance 7 have been tested for cancer.

Other Key Data

There are no structural alerts that would suggest that substance 7 is a DNA-reactive compound. It is negative for gene mutations in bacteria and yeast, but positive in cultured mouse cells. Tests for structural chromosome aberrations in cultured mammalian cells and in rats are negative; however, the animals were not tested at sufficiently high doses. Substance 7 binds to proteins of the cell division spindle; therefore, there is some likelihood for producing numerical chromosome aberrations, an endpoint that is sometimes noted in cancers. In sum, there is limited and conflicting information concerning the mutagenic potential of the agent.

The compound is absorbed via oral and inhalation exposure but only poorly via the skin.

Evaluation

The only indication of a carcinogenic effect comes from the finding of benign tumors in male rats in a single study. There is no confirmation of a carcinogenic potential from dosed females in that study, in males and females in a second rat study, or from mouse and hamster studies.

There is no structural indication that substance 7 is DNA-reactive, there is inconsistent evidence of gene mutations, and chromosome aberration testing is negative. The agent binds to cell division spindle proteins and may have the capacity to induce numerical chromosome anomalies. Further information on gene mutations and in vivo structural and numerical chromosome aberrations may be warranted.

Conclusion

Substance 7 is *not likely* to be carcinogenic to humans via all relevant routes of exposure. This weight of evidence judgment is largely based on the absence of significant tumor increases in chronic rodent studies. Adequate cancer studies in rats, mice, and hamsters fail to show any carcinogenic effect; a second rat study showed an increase in benign tumors at a site in dosed males, but not females.

2.7. PRESENTATION OF RESULTS

The results of the hazard assessment are presented in the form of an overall technical hazard characterization. Additionally, a weight of evidence narrative is used when the conclusion as to carcinogenic potential needs to be presented separately from the overall characterization.

2.7.1. Technical Hazard Characterization

The hazard characterization has two functions. First, it presents results of the hazard assessment and an explanation of how the weight of evidence conclusion was reached. It explains the potential for human hazard, anticipated attributes of its expression, and mode of action considerations for dose response. Second, it contains the information needed for eventual incorporation into a risk characterization consistent with EPA guidance on risk characterization (U.S. EPA, 1995).

The characterization qualitatively describes the conditions under which the agent's effects may be expressed in human beings. These qualitative hazard conditions are ones that are observable in the toxicity data without having done either quantitative dose response or exposure assessment. The description includes how expression is affected by route of exposure and dose levels and durations of exposure.

The discussion of limitations of dose as a qualitative aspect of hazard addresses the question of whether reaching a certain dose range appears to be a precondition for a hazard to be expressed; for example, when carcinogenic effects are secondary to another toxic effect that appears only when a certain dose level is reached. The assumption is made that an agent that causes internal tumors by one route of exposure will be carcinogenic by another route, if it is absorbed by the second route to give an internal dose. Conversely, if there is a route of exposure by which the agent is not absorbed (does not cross an absorption barrier; e.g., the exchange boundaries of skin, lung, and digestive tract through uptake processes) to any significant degree, hazard is not anticipated by that route. An exception to the latter statement would be when the site of contact is also the target tissue of carcinogenicity. Duration of exposure may be a precondition for hazard if, for example, the mode of action requires cytotoxicity or a physiologic change, or is mitogenicity, for which exposure must be sustained for a period of time before

1 effects occur. The characterization could note that one would not anticipate a hazard from
2 isolated, acute exposures. The above conditions are qualitative ones regarding preconditions for
3 effects, not issues of relative absorption or potency at different dose levels. The latter are dealt
4 with under dose response assessment (section 3), and their implications can only be assessed
5 after human exposure data are applied in the characterization of risk.

6 The characterization describes conclusions about mode of action information and its
7 support for recommending dose response approaches.

8 The hazard characterization routinely includes the following in support of risk
9 characterization:

- 10 ● a summary of results of the assessment,
- 11 ● identification of the kinds of data available to support conclusions and explanation of
12 how the data fit together, highlighting the quality of the data in each line of evidence,
13 e.g., tumor effects, short-term studies, structure-activity relationships), and
14 highlighting the coherence of inferences from the different kinds of data,
- 15 ● strengths and limitations (uncertainties) of the data and assessment, including
16 identification of default assumptions invoked in the face of missing or inadequate
17 data,
- 18 ● identification of alternative interpretations of data that are considered equally
19 plausible,
- 20 ● identification of any subpopulations believed to be more susceptible to the hazard
21 than the general population,
- 22 ● conclusions about the agent's mode of action and recommended dose response
23 approaches,
- 24 ● significant issues regarding interpretation of data that arose in the assessment.

25 Typical ones may include:

- 26 -- determining causality in human studies,
- 27 -- dosing (MTD), background tumor rates, relevance of animal tumors to humans,
- 28 -- weighing studies with positive and null results, considering the influence of
29 other available kinds of evidence,

-- drawing conclusions based on mode of action data versus using a default assumption about the mode of action.

2.7.2. Weight of Evidence Narrative

The weight of evidence narrative summarizes the results of hazard assessment employing the descriptors defined in section 2.6.1. The narrative (about two pages in length) explains an agent's human carcinogenic potential and the conditions of its expression. If data do not allow a conclusion as to carcinogenicity, the narrative explains the basis of this determination. An example narrative appears below. More examples appear in Appendix A.

The items regularly included in a narrative are:

- name of agent and Chemical Abstracts Services number, if available,
- conclusions (by route of exposure) about human carcinogenicity, using a standard descriptor from section 2.6.1,
- summary of human and animal tumor data on the agent or its structural analogues, their relevance, and biological plausibility,
- other key data (e.g., structure-activity data, toxicokinetics and metabolism, short-term studies, other relevant toxicity or clinical data),
- discussion of possible mode(s) of action and appropriate dose response approach(es),
- conditions of expression of carcinogenicity, including route, duration, and magnitude of exposure.

Example Narrative

Aromatic Compound

CAS# XXX

CANCER HAZARD SUMMARY

Aromatic compound (AR) is *known* to be carcinogenic to humans by all routes of exposure.

The weight of evidence of human carcinogenicity is based on (a) consistent evidence of elevated leukemia incidence in studies of exposed workers and significant increases of genetic damage in bone marrow cells and blood lymphocytes of exposed workers; (b) significantly

1 increased incidence of cancer in both sexes of several strains of rats and mice; (c) genetic
2 damage in bone marrow cells of exposed rodents and effects on intracellular signals that control
3 cell growth.

4 AR is readily absorbed by all routes of exposure and rapidly distributed throughout the
5 body. The mode of action of AR is not understood. A dose response assessment that assumes
6 linearity of the relationship is recommended as a default.

8 **SUPPORTING INFORMATION**

9 Data include numerous human epidemiologic and biomonitoring studies, long-term
10 bioassays, and other data on effects of AR on genetic material and cell growth processes. The
11 key epidemiologic studies and animal studies are well conducted and reliable. The other data are
12 generally of good quality also.

14 **Human Effects**

15 Numerous epidemiologic and case studies have reported an increased incidence or a
16 causal relationship associating exposure to AR and leukemia. Among the studies are five for
17 which the design and performance as well as follow-up are considered adequate to demonstrate
18 the causal relationship. Biomonitoring studies of exposed workers have found dose-related
19 increases in chromosomal aberrations in bone marrow cells and blood lymphocytes.

21 **Animal Effects**

22 AR caused increased incidence of tumors in various tissues in both sexes of several rat
23 and mouse strains. AR also caused chromosomal aberrations in rabbits, mice, and rats--as it
24 does in humans.

26 **Other Key Data**

27 AR itself is not DNA-reactive and is not mutagenic in an array of test systems both in
28 vitro and in vivo. Metabolism of AR yields several metabolites that have been separately
29 studied for effects on carcinogenic processes. Some have mutagenic activity in test systems and
30 some have other effects on cell growth controls inside cells.

MODE OF ACTION

No rodent tumor precisely matches human leukemia in pathology. The closest parallel is a mouse cancer of blood-forming tissue. Studies of the effects of AR at the cell level in this model system are ongoing. As yet, the mode of action of AR is unclear, but most likely the carcinogenic activity is associated with one or a combination of its metabolites. It is appropriate to apply a linear approach to the dose response assessment pending a better understanding because: (a) genetic damage is a typical effect of AR exposure in mammals and (b) metabolites of AR produce mutagenic effects in addition to their other effects on cell growth controls; AR is a multitissue carcinogen in mammals suggesting that it is affecting a common controlling mechanism of cell growth.

3. DOSE RESPONSE ASSESSMENT

Dose response assessment first addresses the relationship of dose² to the degree of response observed in an experiment or human study. When environmental exposures are outside of the range of observation, extrapolations are necessary in order to estimate or characterize the dose relationship (ILSI, 1995). In general, three extrapolations may be made: from high to low doses, from animal to human responses, and from one route of exposure to another.

The dose response assessment proceeds in two parts. The first is assessment of the data in the range of empirical observation. This is followed by extrapolations either by modeling, if there are sufficient data to support a model, or by a default procedure based as much as possible on information about the agent's mode of action. The following discussion covers the assessment of observed data and extrapolation procedures, followed by sections on analysis of response data and analysis of dose data. The final section discusses dose response characterization.

3.1. DOSE RESPONSE RELATIONSHIP

In the discussion that follows, reference to “response” data includes measures of tumorigenicity as well as other responses related to carcinogenicity. The other responses may include effects such as changes in DNA, chromosomes, or other key macromolecules, effects on growth signal transduction, induction of physiological or hormonal changes, effects on cell proliferation, or other effects that play a role in the process. Responses other than tumorigenicity may be considered part of the observed range in order either to extend the tumor dose response analysis or to inform it. The nontumor response or responses also may be used in lieu of tumor data if they are considered to be a more informative representation of the carcinogenic process for an agent (see section 3.2).

For this discussion, “exposure” means contact of an agent with the outer boundary of an organism. “Applied dose” means the amount of an agent presented to an absorption barrier and available for absorption. “Internal dose” means the amount crossing an absorption barrier (e.g., the exchange boundaries of skin, lung, and digestive tract) through uptake processes. “Delivered dose” for an organ or cell means the amount available for interaction with that organ or cell (U.S. EPA, 1992a).

3.1.1. Analysis in the Range of Observation

Biologically Based and Case-Specific Models

A biologically based model is one whose parameters are calculated independently of curve-fitting of tumor data. If data are sufficient to support a biologically based model specific to the agent and the purpose of the assessment is such as to justify investing resources supporting use, this is the first choice for both the observed tumor and related response data and for extrapolation below the range of observed data in either animal or human studies. Examples are the two-stage models of initiation plus clonal expansion and progression developed by Moolgavkar and Knudson (1981) and Chen and Farland (1991). Such models require extensive data to build the form of the model as well as to estimate how well it conforms with the observed carcinogenicity data. Theoretical estimates of process parameters, such as cell proliferation rates, are not used to enable application of such a model (Portier, 1987).

Similarly preferred as a first choice are dose response models based on general concepts of mode of action and data on the agent. For a case-specific model, model parameters and data are obtained from studies on the agent.

In most cases, a biologically based or case-specific model will not be practicable, either because the necessary data do not exist or the decisions that the assessment are to support do not justify or permit, the time and resources required. In these cases, the analysis proceeds using curve-fitting models followed by default procedures for extrapolation, based, to the extent possible, on mode of action and other biological information about the agent. These methods and assumptions are described below.

Curve-Fitting and Point of Departure for Extrapolation

Curve-fitting models are used that are appropriate to the kind of response data in the observed range. Any of several models can be used; e.g., the models developed for benchmark dose estimation for noncancer endpoints may be applied (Barnes et al., 1995). For some data sets, particularly those with extreme curvature, the impact of model selection can be significant. In these cases, the choice is rationalized on biological grounds as possible. In other cases, the nature of the data or the way it is reported will suggest other types of models; for instance, when

1 longitudinal data on tumor development are available, time to tumor or survival models may be
2 necessary and appropriate to fit the data.

3 A point of departure for extrapolation is estimated. This is a point that is either a data
4 point or an estimated point that can be considered to be in the range of observation, without any
5 significant extrapolation. The LED₁₀--the lower 95% confidence limit on a dose associated with
6 10% extra risk--is such a point and is the standard point of departure, adopted as a matter of
7 science policy to remain as consistent and comparable from case to case as possible.³ It is also a
8 comparison point for noncancer endpoints (U.S. EPA, 1991f). The central estimate of the ED₁₀
9 also may be appropriate for use in relative hazard and potency ranking.

10 For some data sets, a choice of point of departure other than the LED₁₀ may be
11 appropriate. For example, if the observed response is below the LED₁₀, then a lower point may
12 be a better choice. Moreover, some forms of data may not be amenable to curve-fitting
13 estimation, but to estimation of a "low-" or "no-observable-adverse-effect level" (LOAEL,
14 NOAEL) instead, e.g., certain continuous data.

15 The rationale supporting the use of the LED₁₀ is that it a 10% response is at or just below
16 the limit of sensitivity of discerning a significant difference in most long-term rodent studies.
17 The lower confidence limit on dose is used to appropriately account for experimental uncertainty
18 (Barnes et al., 1995) and for consistency with the "benchmark dose" approach for noncancer
19 assessment; it does not provide information about human variability. In laboratory studies of
20 cancer or noncancer endpoints, the level of dose at which increased incidence of effects can be
21 detected, as compared to controls, is a function of the size of the sample (e.g., number of
22 animals), dose spacing, and other design aspects. In noncancer assessment, the dose at which
23 significant effects are not observed is traditionally termed the NOAEL. This is not, in fact, a
24 level of zero effect. The NOAEL in most study protocols is about the same as an LED₅ or
25 LED₁₀--the lower 95% confidence limit on a dose associated with a 5% or 10% increased effect
26 (Faustman et al., 1994; Haseman, 1983). Adopting parallel points of departure for cancer and
27 noncancer assessment is intended to make discussion and comparison of the two kinds of

It is appropriate to report the central estimate of the ED₁₀, the upper and lower 95% confidence limits, and a graphical representation of model fit.

assessment more comparable because of their similar science and science policy bases and similar analytic approaches.

Analysis of human studies in the observed range is designed case by case, depending on the type of study and how dose and response are measured in the study. In some cases the agent may have discernible interactive effects with another agent (e.g., asbestos and smoking), making possible estimation of contribution of the agent and others as risk factors. Also, in some cases, estimation of population risk in addition, or in lieu of, individual risk may be appropriate.

3.1.2. Analysis in the Range of Extrapolation

Extrapolation to lower doses is usually necessary, and in the absence of a biologically based or case-specific model, is based on one of the three default procedures described below. The Agency has adopted these three procedures as a matter of science policy based on current hypotheses of the likely shapes of dose response curves for differing modes of action. The choice of the procedure to be used in an individual case is a judgment based on the agent's modes of action.

Linear

A default assumption of linearity is appropriate when the evidence supports a mode of action of gene mutation due to DNA reactivity or supports another mode of action that is anticipated to be linear. Other elements of empirical support may also support an inference of linearity, e.g., the background of human exposure to an agent might be such that added human exposure is on the linear part of a dose response curve that is sublinear overall. The default assumption of linearity is also appropriate as the ultimate science policy default when evidence shows no DNA reactivity or other support for linearity, but neither does it show sufficient evidence of a nonlinear mode of action to support a nonlinear procedure.

For linear extrapolation, a straight line is drawn from the point of departure to the origin-zero dose, zero response (Flamm and Winbush, 1984; Gaylor and Kodell, 1980; Krewski et al., 1984). This approach is generally conservative of public health, in the absence of information about the extent of human variability in sensitivity to effects. When a linear extrapolation procedure is used, the risk characterization summary displays the degree of extrapolation that is

1 being made from empirical data and discusses its implications for the interpretation of the
2 resulting quantitative risk estimates.

4 *Nonlinear*

5 A default assumption of nonlinearity is appropriate when there is no evidence for
6 linearity and sufficient evidence to support an assumption of nonlinearity. The mode of action
7 may lead to a dose response relationship that is nonlinear, with response falling much more
8 quickly than linearly with dose, or being most influenced by individual differences in sensitivity.
9 Alternatively, the mode of action may theoretically have a threshold, e.g., the carcinogenicity
10 may be a secondary effect of toxicity or of an induced physiological change (see example 5,
11 section 2.6.3) that is itself a threshold phenomenon.

12 As a matter of science policy under this analysis, nonlinear probability functions are not
13 fitted to the response data to extrapolate quantitative low-dose risk estimates because different
14 models can lead to a very wide range of results, and there is currently no basis, generally, to
15 choose among them. Sufficient information to choose leads to a biologically based or case-
16 specific model. In cases of nonlinearity, the risk is not extrapolated as a probability of an effect
17 at low doses. A margin of exposure analysis is used, as described below, to evaluate concern for
18 levels of exposure. The margin of exposure is the LED_{10} or other point of departure divided by
19 the environmental exposure of interest. The EPA does not generally try to distinguish between
20 modes of action that might imply a "true threshold" from others with a nonlinear dose response
21 relationship. Except in unusual cases where extensive information is available, it is not possible
22 to distinguish between these empirically.

23 The environmental exposures of interest, for which margins of exposure are estimated,
24 may be actual or projected future levels. The risk manager decides whether a given margin of
25 exposure is acceptable under applicable management policy criteria. The risk assessment
26 provides supporting information to assist the decisionmaker.

1 The EPA often conducts margin of exposure analyses to accompany estimates of
2 reference doses or concentrations (RfD, RfC) for noncancer endpoints.⁴ The procedure for a
3 margin of exposure analysis for a response related to carcinogenicity is operationally analogous,
4 the difference being that a threshold of cancer response is not necessarily presumed. If, in a
5 particular case, the evidence indicates a threshold, as in the case of carcinogenicity being
6 secondary to another toxicity that has a threshold, the margin of exposure analysis for the
7 toxicity is the same as is done for a noncancer endpoint, and an RfD or RfC for that toxicity also
8 may be estimated and considered in cancer assessment.

9 The analogy between margin of exposure analysis for noncancer and cancer responses
10 begins with the analogy of points of departure; for both it is an effect level, either LED₁₀ or other
11 point (presented as a human equivalent dose or concentration), as data support. For cancer
12 responses, when animal data are used, the point of departure is a human equivalent dose or
13 concentration arrived at by interspecies dose adjustment or toxicokinetic analysis. It is likely
14 that many of the margin of exposure analyses for cancer will be for responses other than tumor
15 incidence. This is because the impetus for considering a carcinogenic agent to have a nonlinear
16 dose response will be a conclusion that there is sufficient evidence to support that view, and this
17 evidence will often be information about a response that is a precursor to tumors.

18 To support a risk manager's consideration of the margin of exposure, information is
19 provided in a risk assessment about current understanding of the phenomena that may be
20 occurring as dose (exposure) decreases substantially below the observed data. The goal is to
21 provide as much information as possible about the risk reduction that accompanies lowering of
22 exposure. To this end, some important points to address include:

- 23 ● the slope of the observed dose response relationship at the point of departure and its
24 uncertainties and implications for risk reduction associated with exposure reduction (a
25 shallow slope suggests less reduction than a steep slope),

An RfD or RfC is an estimate with uncertainty spanning perhaps an order of magnitude of daily exposure to the human population (including sensitive subgroups) that is anticipated to be without appreciable deleterious effects during a lifetime. It is arrived at by dividing empirical data on effects by uncertainty factors that consider inter- and intraspecies variability, extent of data on all important chronic exposure toxicity endpoints, and availability of chronic as opposed to subchronic data.

- the nature of the response used for the dose response assessment,
- the nature and extent of human variability in sensitivity to the phenomena involved,
- persistence of the agent in the body,
- human sensitivity to the phenomena as compared with experimental animals.

As a default assumption for two of these points, a factor of no less than 10-fold each may be employed to account for human variability and for interspecies differences in sensitivity when humans may be more sensitive than animals. When humans are found to be less sensitive than animals, a default factor of no smaller than a 1/10 fraction may be employed to account for this. If any information about human variability or interspecies differences is available, it is used instead of the default or to modify it as appropriate. In the case of analysis based on human studies, obviously, interspecies differences are not a factor. It should be noted that the dose response relationship and inter- or intraspecies variability in sensitivity are independent. That is, reduction of dose reduces risk; it does not change variability. To support consideration of acceptability of a margin of exposure by the risk manager, the assessment considers all of the hazard and dose response factors together; hence, the factors for inter- and intraspecies differences alone are not to be considered a default number for an acceptable margin of exposure. (See Section 1.3.2.5.)

It is appropriate to provide a graphical representation of the data and dose response modeling in the observed range, also showing exposure levels of interest to the decisionmaker. (See figure 3-1.) In order to provide a frame of reference, by way of comparison, a straight line extrapolation may be displayed to show what risk levels would be associated with decreasing dose, if the dose response were linear. If this is done, the

1 Insert Figure 3-1 here.

2

3 **Note: Figure 3-1 is in a separate file. It can be viewed, but cannot be printed unless a high quality printer is used, such as an**
4 **HP LaserJet 4si or better. If you need a paper copy, please phone the contact listed at the beginning of this notice.**

5

1 clear accompanying message is that, in this case of nonlinearity, the response falls
2 disproportionately with decreasing dose.

3 4 *Linear and Nonlinear*

5 Both linear and nonlinear procedures may be used in particular cases. If a mode of
6 action analysis finds substantial support for differing modes of action for different tumor sites,
7 an appropriate procedure is used for each. Both procedures may also be appropriate to discuss
8 implications of complex dose response relationships. For example, if it is apparent that an agent
9 is both DNA reactive and is highly active as a promotor at high doses, and there are insufficient
10 data for modeling, both linear and nonlinear default procedures may be needed to decouple and
11 consider the contribution of both phenomena.

12 13 **3.1.3. Use of Toxicity Equivalence Factors and Relative Potency Estimates**

14 A toxicity equivalence factor (TEF) procedure is one used to derive quantitative dose
15 response estimates for agents that are members of a category or class of agents. TEFs are based
16 on shared characteristics that can be used to order the class members by carcinogenic potency
17 when cancer bioassay data are inadequate for this purpose (U.S. EPA, 1991c). The ordering is
18 by reference to the characteristics and potency of a well-studied member or members of the
19 class. Other class members are indexed to the reference agent(s) by one or more shared
20 characteristics to generate their TEFs. The TEFs are usually indexed at increments of a factor of
21 10. Very good data may permit a smaller increment to be used. Shared characteristics that may
22 be used are, for example, receptor-binding characteristics, results of assays of biological activity
23 related to carcinogenicity, or structure-activity relationships.

24 TEFs are generated and used for the limited purpose of assessment of agents or mixtures
25 of agents in environmental media when better data are not available. When better data become
26 available for an agent, its TEF should be replaced or revised. Criteria for constructing TEFs are
27 given in U.S. EPA (1991b). The criteria call for data that are adequate to support summing
28 doses of the agents in mixtures. To date, adequate data to support use of TEF's has been found
29 in only one class of compounds (dioxins) (U.S. EPA, 1989a).

1 Relative potencies can be similarly derived and used for agents with carcinogenicity or
2 other supporting data. These are conceptually similar to TEFs, but they are less firmly based in
3 science and do not have the same level of data to support them. They are used only when there
4 is no better alternative.

5 The uncertainties associated with both TEFs and relative potencies are explained
6 whenever they are used.

7 8 **3.2. RESPONSE DATA**

9 Response data for analysis include tumor incidence data from human or animal studies as
10 well as data on other responses as they relate to an agent's carcinogenicity, such as effects on
11 growth control processes or cell macromolecules or other toxic effects. Tumor incidence data
12 are ordinarily the basis of dose response assessment, but other response data can augment such
13 assessment or provide separate assessments of carcinogenicity or other important effects.

14 Data on carcinogenic processes underlying tumor effects may be used to support
15 biologically based or case-specific models. Other options for such data exist. If confidence is
16 high in the linkage of a precursor effect and the tumor effect, the assessment of tumor incidence
17 may be extended to lower dose levels by linking it to the assessment of the precursor effect
18 (Swenberg et al., 1987). Even if a quantitative link is not appropriate, the assessment for a
19 precursor effect may provide a view of the likely shape of the dose response curve for tumor
20 incidence below the range of tumor observation (Cohen and Ellwein, 1990; Choy, 1993). If
21 responses other than tumor incidence are regarded as better representations of the
22 carcinogenicity of the agent, they may be used in lieu of tumor responses. For example, if it is
23 concluded that the carcinogenic effect is secondary to another toxic effect, the dose response for
24 the other effect will likely be more pertinent for risk assessment. As another example, if
25 disruption of hormone activity is the key mode of action of an agent, data on hormone activity
26 may be used in lieu of tumor incidence data.

27 If adequate positive human epidemiologic response data are available, they provide an
28 advantageous basis for analysis since concerns about interspecies extrapolation do not arise.
29 Adequacy of human exposure data for quantification is an important consideration in deciding
30 whether epidemiologic data are the best basis for analysis in a particular case. If adequate

exposure data exist in a well-designed and well-conducted epidemiologic study that detects no effects, it may be possible to obtain an upper-bound estimate of the potential human risk to provide a check on plausibility of available estimates based on animal tumor or other responses, e.g., do confidence limits on one overlap the point estimate of the other?

When animal studies are used, response data from a species that responds most like humans should be used if information to this effect exists. If this is unknown and an agent has been tested in several experiments involving different animal species, strains, and sexes at several doses and different routes of exposure, all of the data sets are considered and compared, and a judgment is made as to the data to be used to best represent the observed data and important biological features such as mode of action. Appropriate options for presenting results include:

- use of a single data set,
- combining data from different experiments (Stiteler et al., 1993; Vater et al., 1993),
- showing a range of results from more than one data set,
- showing results from analysis of more than one statistically significant tumor response based on differing modes of action,
- representing total response in a single experiment by combining animals with statistically significant tumors at more than one site, or
- a combination of these options.

The approach judged to best represent the data is presented with the rationale for the judgment, including the biological and statistical considerations involved. The following are some points to consider:

- quality of study protocol and execution,
- proportion of malignant neoplasms,
- latency of onset of neoplasia,
- number of data points to define the relationship of dose and response,
- background incidence in test animal,
- differences in range of response among species, sexes, strains,
- most sensitive responding species, and
- availability of data on related precursor events to tumor development.

Analyses of carcinogenic effects other than tumor incidence are similarly presented and evaluated for their contribution to a best judgment on how to represent the biological data for dose response assessment.

3.3. DOSE DATA

Whether animal experiments or epidemiologic studies are the sources of data, questions need to be addressed in arriving at an appropriate measure of dose for the anticipated environmental exposure. Among these are:

- whether the dose is expressed as an environmental concentration, applied dose, or delivered dose to the target organ,
- whether the dose is expressed in terms of a parent compound, one or more metabolites, or both,
- the impact of dose patterns and timing where significant,
- conversion from animal to human doses, where animal data are used, and
- the conversion metric between routes of exposure where necessary and appropriate.

In practice, there may be little or no information on the concentration or identity of the active form at a target; being able to compare the applied and delivered doses between routes and species is the ideal, but is rarely attained. Even so, the objective is to use available data to obtain as close to a measure of internal or delivered dose as possible.

The following discussion assumes that the analyst will have data of varying detail in different cases about toxicokinetics and metabolism. Discussed below are approaches to basic data that are most frequently available, as well as approaches and judgments for improving the analysis based on additional data. The estimation of dose in human studies is tailored to the form of dose data available.

3.3.1. Interspecies Adjustment of Dose

When adequate data are available, the doses used in animal studies can be adjusted to equivalent human doses using toxicokinetic information on the particular agent. The methods used should be tailored to the nature of the data on a case-by-case basis. In rare cases, it may also be possible to make adjustments based on toxicodynamic considerations. In most cases,

1 however, there are insufficient data available to compare dose between species. In these cases,
2 the estimate of human equivalent dose is based on science policy default assumptions. The
3 defaults described below are modified or replaced whenever better comparative data on
4 toxicokinetic or metabolic relationships are available. The availability and discussion of the
5 latter also may permit reduction or discussion of uncertainty in the analysis.

6 For oral exposure, the default assumption is that delivered doses are related to applied
7 dose by a power of body weight. This assumption rests on the similarities of mammalian
8 anatomy, physiology, and biochemistry generally observed across species. This assumption is
9 more appropriate at low applied dose concentrations where sources of nonlinearity, such as
10 saturation or induction of enzyme activity, are less likely to occur. To derive an equivalent
11 human oral dose from animal data, the default procedure is to scale daily applied doses
12 experienced for a lifetime in proportion to body weight raised to the 0.75 power ($W^{0.75}$).
13 Equating exposure concentrations in parts per million units for food or water is an alternative
14 version of the same default procedure because daily intakes of these are in proportion to $W^{0.75}$.
15 The rationale for this factor rests on the empirical observation that rates of physiological
16 processes consistently tend to maintain proportionality with $W^{0.75}$. A more extensive discussion
17 of the rationale and data supporting the Agency's adoption of this scaling factor is in U.S. EPA,
18 1992b. Information such as blood levels or exposure biomarkers or other data that are available
19 for interspecies comparison are used to improve the analysis when possible.

20 The default procedure to derive an human equivalent concentration of inhaled particles
21 and gases is described in U.S. EPA (1994) and Jarabek (1995a,b). The methodology estimates
22 respiratory deposition of inhaled particles and gases and provides methods for estimating
23 internal doses of gases with different absorption characteristics. The method is able to
24 incorporate additional toxicokinetics and metabolism to improve the analysis if such data are
25 available.

26 27 **3.3.2. Toxicokinetic Analyses**

28 Physiologically based mathematical models are potentially the most comprehensive way
29 to account for toxicokinetic processes affecting dose. Models build on physiological

1 compartmental modeling and attempt to incorporate the dynamics of tissue perfusion and the
2 kinetics of enzymes involved in metabolism of an administered compound.

3 A comprehensive model requires the availability of empirical data on the carcinogenic
4 activity contributed by parent compound and metabolite or metabolites and data by which to
5 compare kinetics of metabolism and elimination between species. A discussion of issues of
6 confidence accompanies presentation of model results (Monro, 1992). This includes
7 considerations of model validation and sensitivity analysis that stress the predictive performance
8 of the model. When a delivered dose measure is used in animal to human extrapolation of dose
9 response data, the assessment should discuss the confidence in the assumption that the
10 toxicodynamics of the target tissue(s) will be the same in both species. Toxicokinetic data can
11 improve dose response assessment by accounting for sources of change in proportionality of
12 applied to internal or delivered dose at various levels of applied dose. Many of the sources of
13 potential nonlinearity involve saturation or induction of enzymatic processes at high doses. An
14 analysis that accounts for nonlinearity (for instance, due to enzyme saturation kinetics) can assist
15 in avoiding overestimation or underestimation of low dose response otherwise resulting from
16 extrapolation from a sublinear or supralinear part of the experimental dose response curve
17 (Gillette, 1983). Toxicokinetic processes tend to become linear at low doses, an expectation that
18 is more robust than low-dose linearity of response (Hattis, 1990). Accounting for toxicokinetic
19 nonlinearities allows better description of the shape of the curve at relatively high levels of dose
20 in the range of observation, but cannot determine linearity or nonlinearity of response at low
21 dose levels (Lutz, 1990a; Swenberg et al., 1987).

22 Toxicokinetic modeling results may be presented as the preferred method of estimating
23 human equivalent dose or in parallel discussion with default assumptions depending on relative
24 confidence in the modeling.

26 **3.3.3. Route-to-Route Extrapolation**

27 Judgments frequently need to be made about the carcinogenicity of an agent through a
28 route of exposure different than the one in the underlying studies. For example, exposures of
29 interest may be through inhalation of an agent tested primarily through animal feeding studies or

1 through ingestion of an agent that showed positive results in human occupational studies from
2 inhalation exposure.

3 Route-to-route extrapolation has both qualitative and quantitative aspects. For the
4 qualitative aspect, the assessor weighs the degree to which positive results through one route of
5 exposure in human or animal studies support a judgment that similar results would have been
6 observed in appropriate studies using the route of exposure of interest. In general, confidence in
7 making such a judgment is strengthened when the tumor effects are observed at a site distant
8 from the portal of entry and when absorption through the route of exposure of interest is similar
9 to absorption via the tested routes. In the absence of contrary data, the qualitative default
10 assumption is that, if the agent is absorbed by a route to give an internal dose, it may be
11 carcinogenic by that route. (See section 2.7.1.)

12 When a qualitative extrapolation can be supported, quantitative extrapolation may still be
13 problematic in the absence of adequate data. The differences in biological processes among
14 routes of exposure (oral, inhalation, dermal) can be great because of, for example, first-pass
15 effects and differing results from different exposure patterns. There is no generally applicable
16 method for accounting for these differences in uptake processes in quantitative route-to-route
17 extrapolation of dose response data in the absence of good data on the agent of interest.
18 Therefore, route-to-route extrapolation of dose data relies on a case-by-case analysis of available
19 data. When good data on the agent itself are limited, an extrapolation analysis can be based on
20 expectations from physical and chemical properties of the agent, properties and route-specific
21 data on structurally analogous compounds, or in vitro or in vivo uptake data on the agent.
22 Route-to-route uptake models may be applied if model parameters are suitable for the compound
23 of interest. Such models are currently considered interim methods; further model development
24 and validation is awaiting the development of more extensive data (see generally, Gerrity and
25 Henry, 1990). For screening or hazard ranking, route-to-route extrapolation may be based on
26 assumed quantitative comparability as a default, as long as it is reasonable to assume absorption
27 by compared routes. When route-to-route extrapolation is used, the assessor's degree of
28 confidence in both the qualitative and quantitative extrapolation should be discussed in the
29 assessment and highlighted in the dose response characterization.

3.3.4. Dose Averaging

The cumulative dose received over a lifetime, expressed as lifetime average daily dose, is generally considered an appropriate default measure of exposure to a carcinogen (Monro, 1992). The assumption is made that a high dose of a carcinogen received over a short period of time is equivalent to a corresponding low dose spread over a lifetime. While this is a reasonable default assumption based on theoretical considerations, departures from it are expected. Another approach is needed in some cases, such as when dose-rate effects are noted (e.g., formaldehyde). Cumulative dose may be replaced, as appropriate and justified by the data, with other dose measures. In such cases, modifications to the default assumption are made to take account of these effects; the rationale for the selected approach is explained.

In cases where a mode of action or other feature of the biology has been identified that has special dose implications for sensitive subpopulations (e.g., differential effects by sex or disproportionate impacts of early-life exposure), these are explained and are recorded to guide exposure assessment and risk characterization. Special problems arise when the human exposure situation of concern suggests exposure regimens (e.g., route and dosing schedule) that are substantially different from those used in the relevant animal studies. These issues are explored and pointed out for attention in the exposure assessment and risk characterization.

3.4. DISCUSSION OF UNCERTAINTIES

The exploration of significant uncertainties in data for dose and response and in extrapolation procedures is part of the assessment. The presentation distinguishes between model uncertainty and parameter uncertainty. Model uncertainty is an uncertainty about a basic biological question. For example, a default, linear dose response extrapolation may have been made based on tumor and other key evidence supporting the view that the model for an agent's mode of action is a DNA-reactive process. Discussion of the confidence in the extrapolation is appropriately done qualitatively or by showing results for alternatives that are equally plausible. It is not useful, for example, to conduct quantitative uncertainty analysis running multiple forms of linear models. This would obviate the function of the policy default.

Parameter uncertainties deal with numbers representing statistical or analytical measures of variance or error in data or estimates. Uncertainties in parameters are described

quantitatively, if practicable, through sensitivity analysis and statistical uncertainty analysis. With the recent expansion of readily available computing capacity, computer methods are being adapted to create simulated biological data that are comparable with observed information. These simulations can be used for sensitivity analysis, for example, to analyze how small, plausible variations in the observed data could affect dose response estimates. These simulations can also provide information about experimental uncertainty in dose response estimates, including a distribution of estimates that are compatible with the observed data. Because these simulations are based on the observed data, they cannot assist in evaluating the extent to which the observed data as a whole are idiosyncratic rather than typical of the true situation. If quantitative analysis is not possible, significant parameter uncertainties are described qualitatively. In either case, the discussion highlights uncertainties that are specific to the agent being assessed, as distinct from those that are generic to most assessments.

Estimation of the applied dose in a human study has numerous uncertainties such as the exposure fluctuations that humans experience compared with the controlled exposures received by animals on test. In a prospective cohort study, there is opportunity to monitor exposure and human activity patterns for a period of time that supports estimation of applied dose (U.S. EPA, 1992a). In a retrospective study, exposure may be based on monitoring data but is often based on human activity patterns and levels reconstructed from historical data, contemporary data, or a combination of the two. Such reconstruction is accompanied by analysis of uncertainties considered with sensitivity analysis in the estimation of dose (Wyzga, 1988; U.S. EPA, 1986a). These uncertainties can also be assessed for any confounding factor for which a quantitative adjustment of dose response data is made (U.S. EPA, 1984).

3.5. TECHNICAL DOSE RESPONSE CHARACTERIZATION

As with hazard characterization, the dose response characterization serves the dual purposes of presenting a technical characterization of the assessment results and supporting the risk characterization.

The characterization presents the results of analyses of dose data, of response data, and of dose response. When alternative approaches are plausible and persuasive in selecting dose data, response data, or extrapolation procedures, the characterization follows the alternative paths of

analysis and presents the results. The discussion covers the question of whether any should be preferred over others because it (or they) better represents the available data or corresponds to the view of the mechanism of action developed in the hazard assessment. The results for different tumor types by sex and species are provided along with the one(s) preferred. Similarly, results for responses other than tumor incidence are shown if appropriate.

Numerical dose response estimates are presented to one significant figure. Numbers are qualified as to whether they represent central tendency or upper bounds and whether the method used is inherently more likely to overestimate or underestimate (Krewski et al., 1984).

In cases where a mode of action or other feature of the biology has been identified that has special implications for early-life exposure, differential effects by sex, or other concerns for sensitive subpopulations, these are explained. Similarly, any expectations that high dose-rate exposures may alter the risk picture for some portion of the population are described. These and other perspectives are recorded to guide exposure assessment and risk characterization. Whether the lifetime average daily dose or another measure of dose should be considered for differing exposure scenarios is discussed.

Uncertainty analyses, qualitative or quantitative if possible, are highlighted in the characterization.

The dose response characterization routinely includes the following, as appropriate for the data available:

- identification of the kinds of data available for analysis of dose and response and for dose response assessment,
- results of assessment as above,
- explanation of analyses in terms of quality of data available,
- selection of study/response and dose metric for assessment,
- discussion of implications of variability in human susceptibility, including for susceptible subpopulation,
- applicability of results to varying exposure scenarios--issues of route of exposure, dose rate, frequency, and duration,
- discussion of strengths and limitations (uncertainties) of the data and analyses that are quantitative as well as qualitative, and

- 1 ● special issues of interpretation of data, such as:
- 2 -- selecting dose data, response data, and dose response approach(es),
- 3 -- use of meta-analysis,
- 4 -- uncertainty and quantitative uncertainty analysis.
- 5

4. TECHNICAL EXPOSURE CHARACTERIZATION

Guidelines for exposure assessment of carcinogenic and other agents are published (U.S. EPA, 1992a) and are used in conjunction with these cancer risk assessment guidelines. Presentation of exposure descriptors is a subject of discussion in EPA risk characterization guidance (U.S. EPA, 1995). The exposure characterization is a technical characterization that presents the assessment results and supports risk characterization.

The characterization provides a statement of purpose, scope, level of detail, and approach used in the assessment, identifying the exposure scenario(s) covered. It estimates the distribution of exposures among members of the exposed population as the data permit. It identifies and compares the contribution of different sources and routes and pathways of exposure. Estimates of the magnitude, duration, and frequency of exposure are included as available monitoring or modeling results or other reasonable methods permit. The strengths and limitations (uncertainties) of the data and methods of estimation are made clear.

The exposure characterization routinely includes the following, as appropriate and possible for the data available:

- identification of the kinds of data available,
- results of assessment as above,
- explanation of analyses in terms of quality of data available,
- uncertainty analyses as discussed in Exposure Assessment Guidelines, distinguishing uncertainty from variability, and
- explanation of derivation of estimators of "high end" or central tendency of exposure and their appropriate use.

5. RISK CHARACTERIZATION

5.1. PURPOSE

The risk characterization process includes an integrative analysis followed by a presentation in a Risk Characterization Summary, of the major results of the risk assessment. The Risk Characterization Summary is a nontechnical discussion that minimizes the use of technical terms. It is an appraisal of the science that supports the risk manager in making public health decisions, as do other decisionmaking analyses of economic, social, or technology issues. It also serves the needs of other interested readers. The summary is an information resource for preparation of risk communication information, but being somewhat technical, is not itself the usual vehicle for communication with every audience.

The integrative analysis brings together the assessments and characterizations of hazard, dose response, and exposure to make risk estimates for the exposure scenarios of interest. This analysis is generally much more extensive than the Risk Characterization Summary. It may be peer-reviewed or subject to public comment along with the summary in preparation for an Agency decision. The integrative analysis may be titled differently by different EPA programs (e.g., "Staff Paper" for criteria air pollutants), but it typically will identify exposure scenarios of interest in a decisionmaking and present risk analyses associated with them. Some of the analyses may concern scenarios in several media, others may examine, for example, only drinking water risks. It also may be the document that contains quantitative analyses of uncertainty.

The values supported by a risk characterization throughout the process are *transparency* in environmental decisionmaking, *clarity* in communication, *consistency* in core assumptions and science policies from case to case, and *reasonableness*. While it is appropriate to err on the side of protection of health and the environment in the face of scientific uncertainty, common sense and reasonable application of assumptions and policies are essential to avoid unrealistic estimates of risk (U.S. EPA, 1995). Both integrative analyses and the Risk Characterization Summary present an integrated and balanced picture of the analysis of the hazard, dose response, and exposure. The risk analyst should provide summaries of the evidence and results and describe the quality of available data and the degree of confidence to be placed in the risk

estimates. Important features include the constraints of available data and the state of knowledge, significant scientific issues, and significant science and science policy choices that were made when alternative interpretations of data existed (U.S. EPA, 1995). Choices made about using default assumptions or data in the assessment are explicitly discussed in the course of analysis, and if a choice is a significant issue, it is highlighted in the summary.

5.2. APPLICATION

Risk characterization is a necessary part of generating any Agency report on risk, whether the report is preliminary to support allocation of resources toward further study or comprehensive to support regulatory decisions. In the former case, the detail and sophistication of the characterization are appropriately small in scale; in the latter case, appropriately extensive. Even if a document covers only parts of a risk assessment (hazard and dose response analyses for instance), the results of these are characterized.

Risk assessment is an iterative process that grows in depth and scope in stages from screening for priority-making, to preliminary estimation, to fuller examination in support of complex regulatory decisionmaking. Default assumptions are used at every stage because no database is ever complete, but they are predominant at screening stages and are used less as more data are gathered and incorporated at later stages. Various provisions in EPA-administered statutes require decisions based on findings that represent all stages of iteration. There are close to 30 provisions within the major statutes that require decisions based on risk, hazard, or exposure assessment. For example, Agency review of premanufacture notices under section 5 of the Toxic Substances Control Act relies on screening analyses, while requirements for industry testing under section 4 of that Act rely on preliminary analyses of risk or simply of exposure. At the other extreme, air quality criteria under the Clean Air Act rest on a rich data collection required by statute to undergo reassessment every few years. There are provisions that require ranking of hazards of numerous pollutants--by its nature a screening level of analysis--and other provisions that require a full assessment of risk. Given this range in the scope and depth of analyses, not all risk characterizations can or should be equal in coverage or depth. The risk assessor must carefully decide which issues in a particular assessment are important to present, choosing those that are noteworthy in their impact on results. For example, health effect

assessments typically rely on animal data since human data are rarely available. The objective of characterization of the use of animal data is not to recount generic issues about interpreting and using animal data. Agency guidance documents cover these. Instead, the objective is to call out any significant issues that arose within the particular assessment being characterized and inform the reader about significant uncertainties that affect conclusions.

5.3. PRESENTATION OF RISK CHARACTERIZATION SUMMARY

The presentation is a nontechnical discussion of important conclusions, issues, and uncertainties that uses the hazard, dose-response, exposure, and integrative analyses for technical support. The primary technical supports within the risk assessment are the hazard characterization, dose response characterization, and exposure characterization described in this guideline. The risk characterization is derived from these. The presentation should fulfill the aims outlined in the purpose section above.

5.4. CONTENT OF RISK CHARACTERIZATION SUMMARY

Specific guidance on hazard, dose response, and exposure characterization appears in previous sections. Overall, the risk characterization routinely includes the following, capturing the important items covered in hazard, dose response, and exposure characterization.

- primary conclusions about hazard, dose response, and exposure, including equally plausible alternatives,
- nature of key supporting information and analytic methods,
- risk estimates and their attendant uncertainties, including key uses of default assumptions when data are missing or uncertain,
- statement of the extent of extrapolation of risk estimates from observed data to exposure levels of interest (i.e., margin of exposure) and its implications for certainty or uncertainty in qualifying risk,
- significant strengths and limitations of the data and analyses, including any major peer reviewers' issues,
- appropriate comparison with similar EPA risk analyses or common risks with which people may be familiar, and

- 1
 - comparison with assessment of the same problem by another organization.

1 **APPENDIX A**

2
3 This appendix contains several general illustrations of weight of evidence narratives. In
4 addition, after narrative #5 is an example of a briefing summary format.

5
6 **NARRATIVE #1**

7 **Chlorinated Alkene**

8 **CAS# XXX**

9 **CANCER HAZARD SUMMARY**

10 Chlorinated alkene (cl-alkene) is *likely to be carcinogenic to humans by all routes of*
11 *exposure*. The weight of evidence of human carcinogenicity of cl-alkene is based on (a)
12 findings of carcinogenicity in rats and mice of both sexes by oral and inhalation exposures; (b)
13 its similarity in structure to other chlorinated organics that are known to cause liver and kidney
14 damage, and liver and kidney tumors in rats and mice; (c) suggestive evidence of a possible
15 association between cl-alkene exposure of workers in the laundry and dry cleaning industries and
16 increased cancer risk in a number of organ systems; and (d) human and animal data indicating
17 that cl-alkene is absorbed by all routes of exposure.

18 In comparison with other agents designated as likely carcinogens, the overall weight of
19 evidence for cl-alkene places it at the *low end* of the grouping. This is because one cannot
20 attribute observed excess cancer risk in exposed workers solely to cl-alkene. Moreover, there is
21 considerable scientific uncertainty about the human significance and relevance of certain rodent
22 tumors associated with exposure to cl-alkene and other chlorinated organics, but insufficient
23 evidence about mode of action for the animal tumors. Hence, the human relevance of the animal
24 evidence of carcinogenicity relies on a default assumption of relevance.

25 There is no clear evidence about the mode of action for each tumor type induced in rats
26 and mice. Available evidence suggests that cl-alkene induces cancer mainly by promoting cell
27 growth rather than via direct mutagenic action, although a mutagenic mode of action for rat
28 kidney tumors cannot be ruled out. The dose response assessment should, therefore, adopt *both*
29 *default approaches, nonlinear and linear*. It is recognized that the latter approach likely

overestimates risk at low doses if the mode of action is primarily growth-promoting. This approach, however, may be useful for screening analyses.

SUPPORTING INFORMATION

Human Data

A number of epidemiologic studies of dry cleaning and laundry workers that have reported elevated incidences of lung, cervix, esophagus, kidney, blood and lymphoid cancers. Many of these studies are confounded by co-exposure to other petroleum solvents, making them limited for determining whether the observed increased cancer risks are causally related to cl-alkene. The only investigation of dry cleaning workers with no known exposure to other chemicals did not evaluate other confounding factors such as smoking, alcohol consumption, and low socioeconomic status to exclude the possible contribution of these factors to cancer risks.

Animal Data

The carcinogenic potential of cl-alkene has been adequately investigated in two chronic studies in two rodent species, the first study by gavage and the second study by inhalation. Cl-alkene is carcinogenic in the liver in both sexes of mice when tested by either route of exposure. It causes marginally increased incidences of mononuclear cell leukemia (MCL) in both sexes of rats and low incidences of a rare kidney tumor in male rats by inhalation. No increases in tumor incidence were found in rats treated with cl-alkene by gavage. This rat study was considered limited because of high mortality of the animals.

Although cl-alkene causes increased incidences of tumors at multiple sites in two rodent species, controversy surrounds each of the tumor endpoints concerning their relevance and/or significance to humans (see discussion under Mode of Action).

Other Key Data

Cl-alkene is a member of a class of chlorinated organics that often cause liver and kidney toxicity and carcinogenesis in rodents. Like many chlorinated hydrocarbons, cl-alkene itself is tested negative in a battery of standard genotoxicity tests using bacterial and mammalian cells systems including human lymphocytes and fibroblast cells. There is evidence, however, that a

1 minor metabolite generated by an enzyme found in rat kidney tissue is mutagenic. This kidney
2 metabolite has been hypothesized to be related to the development of kidney tumors in the male
3 rat. This metabolic pathway appears to be operative in the human kidney.

4 Human data indicate that cl-alkene is readily absorbed via inhalation but to a much lesser
5 extent by skin contact. Animal data show that cl-alkene is absorbed well by the oral route.

7 **MODE OF ACTION**

8 The mechanisms of cl-alkene-induced mouse liver tumors are not completely understood.
9 One mechanism has been hypothesized to be mediated by a genotoxic epoxide metabolite
10 generated by enzymes found in the mouse liver, but there is a lack of direct evidence in support
11 of this mechanism. A more plausible mechanism that still needs to be further defined is related
12 to liver peroxisomal proliferation and toxicity by TCA (trichloroacetic acid), a major metabolite
13 of cl-alkene. However, there are no definitive data indicating that TCA induces peroxisomal
14 proliferation in humans.

15 The mechanisms by which cl-alkene induces kidney tumors in male rats are even less
16 well understood. The rat kidney response may be related to either kidney toxicity or the activity
17 of a mutagenic metabolite of the parent compound.

18 The human relevance of cl-alkene-induced MCL in rats is unclear. The biological
19 significance of marginally increased incidences of MCL has been questioned by some, since this
20 tumor occurs spontaneously in the tested rat strain at very high background rates. On the other
21 hand, it has been considered by others to be a true finding because there was a decreased time to
22 onset of the disease and the disease was more severe in treated as compared with untreated
23 control animals. The exact mechanism by which cl-alkene increases incidences of MCL in rats
24 is not known.

25 Overall, there is not enough evidence to give high confidence in a conclusion about any
26 single mode of action; it would appear that more than a single mode operates in different rodent
27 tissues. The apparent lack of mutagenicity of cl-alkene itself and its general growth-promoting
28 effect on high background tumors as well as its toxicity toward mouse liver and rat kidney tissue
29 support the view that its predominant mode of action is cell growth promoting rather than

1 mutagenic. A mutagenic contribution to the renal carcinogenicity due to a metabolite cannot be
2 entirely ruled out.

3 4 **NARRATIVE #2**

5 **Unsaturated Aldehyde**

6 **CAS# XXX**

7 **CANCER HAZARD SUMMARY**

8 The potential human hazard of unsaturated aldehyde (UA) *cannot be determined*, but
9 there are *suggestive* data for carcinogenicity.

10 The evidence on carcinogenicity consists of (a) data from an oral animal study showing a
11 response only at the highest dose in female rats, with no response in males and (b) the fact that
12 other low-molecular-weight aldehydes have shown tumorigenicity in the respiratory tract after
13 inhalation. The one study of UA effects by the inhalation route was not adequately performed.
14 The available evidence is too limited to describe human carcinogenicity potential or support dose
15 response assessment.

16 17 **SUPPORTING INFORMATION**

18 **Human Data**

19 An elevated incidence of cancer was reported in a cohort of workers in a chemical plant
20 who were exposed to a mixture of chemicals including UA as a minor component. The study is
21 considered inadequate because of the small size of the cohort studied and the lack of adequate
22 exposure data.

23 24 **Animal Data**

25 In a long-term drinking water study in rats, an increased incidence of adrenal cortical
26 adenomas was found in the highest-dosed females. No other significant finding was made. The
27 oral rat study was well conducted by a standard protocol. In a 1-year study in hamsters at one
28 inhalation dose, no tumors were seen. This study was inadequate due to high mortality and
29 consequent short duration. The chemical is very irritating and is a respiratory toxicant in
30 mammals. The animal data are too limited for conclusions to be drawn.

1 **Structural Analogue Data**

2 UA's structural analogues, formaldehyde and acetaldehyde, both have carcinogenic
3 effects on the rat respiratory tract.

4
5 **Other Key Data**

6 The weight of results of mutagenicity tests in bacteria, fungi, fruit flies, and mice result
7 in an overall conclusion of not mutagenic; UA is lethal to bacteria to a degree that makes testing
8 difficult and test results difficult to interpret. The chemical is readily absorbed by all routes.

9
10 **MODE OF ACTION**

11 Data are not sufficient to judge whether there is a carcinogenic mode of action.

12
13 **NARRATIVE #3**

14 **Alkene Oxide**

15 **CAS# XXX**

16 **CANCER HAZARD SUMMARY**

17 Alkene oxide (AO) should be dealt with *as if* it were a *known* human carcinogen by *all*
18 *routes of exposure*. Several studies in workers, when considered together, suggest an elevated
19 risk of leukemia and lymphoma after long-term exposure to AO, even though no single study
20 conclusively demonstrates that AO caused the cancer. In addition, animal cancer and
21 mutagenicity studies as well as short-term tests of mutagenicity have strongly consistent results
22 that support a level of concern equal to having conclusive human studies.

23 The weight of evidence of human carcinogenicity is based on (a) consistent evidence of
24 carcinogenicity of AO in rats and mice by both oral and inhalation exposure; (b) studies in
25 workers that taken together suggest elevated risk of leukemia and lymphoma to workers exposed
26 to AO and show genetic damage in blood lymphocytes in exposed workers; (c) mutagenic
27 effects in numerous test systems and heritable gene mutations in animals; and (d) membership in
28 a class of DNA-reactive compounds that are regularly observed to cause cancer in animals.

29 Due to its ready absorption by all routes of exposure and rapid distribution throughout
30 the body, AO is expected to pose a risk by any route of exposure. The strong evidence of a

1 mutagenic mode of action supports dose response assessment that assumes *linearity* of the
2 relationship.

4 **SUPPORTING INFORMATION**

5 **Human Data**

6 Elevated risks of lymphatic cancer and cancer of blood-forming tissue have been
7 reported in exposed workers in several studies. The interpretation of the studies separately is
8 complicated by exposures to other agents in each so there is no single study that demonstrates
9 that AO caused the effects; nevertheless, several of the studies together are considered
10 suggestive of AO carcinogenicity because they consistently show cancer elevation in the same
11 tissues. Biomonitoring studies of exposed workers find DNA damage in blood lymphocytes and
12 the degree of DNA damage correlates with the level and duration of AO exposure. Finding this
13 damage in the same tissue in which elevated cancer was seen in workers adds further weight to
14 the positive suggestion from the worker cancer studies. The human data are from well-
15 conducted studies.

17 **Animal Data**

18 AO causes cancer in multiple tissue sites in rats and mice of both sexes by oral and
19 inhalation exposure. The database is more extensive than usual and the studies are good. The
20 observation of multisite, multispecies carcinogenic activity by an agent is considered to be very
21 strong evidence and is often the case with highly mutagenic agents. There are also good studies
22 showing that AO causes heritable germ cell mutations in mice after inhalation exposure--a
23 property that is very highly correlated with carcinogenicity.

25 **Structural Analogue Data**

26 Organic epoxides are commonly found to have carcinogenic effects in animals,
27 particularly the low-molecular-weight ones.

1 **Other Key Data**

2 The structure and DNA reactivity of AO support potential carcinogenicity. Both
3 properties are highly correlated with carcinogenicity. Positive mutagenicity tests *in vitro* and *in*
4 *vivo* add to this support and are reinforced by observation of similar genetic damage in exposed
5 workers.

6 AO is experimentally observed to be readily absorbed by all routes and rapidly
7 distributed through the body.

8
9 **MODE OF ACTION**

10 All of the available data are strongly supportive of a mutagenic mode of action, with a
11 particular human target in lymphatic and blood-forming tissue. The current scientific consensus
12 is that there is virtually complete correspondence between ability of an agent to cause heritable
13 germ cell mutations, as AO does, and carcinogenicity. All of this points to a mutagenic mode of
14 action and supports assuming linearity of the dose response relationship.

15
16 **NARRATIVE #4**

17 **Bis-benzenamine**

18 **CAS# XXX**

19 **CANCER HAZARD SUMMARY**

20 This chemical is *likely* to be carcinogenic to humans by *all routes* of exposure. Its
21 carcinogenic potential is indicated by (a) tumor and toxicity studies on structural analogues,
22 which demonstrate the ability of the chemical to produce thyroid follicular cell tumors in rats
23 and hepatocellular tumors in mice following ingestion and (b) metabolism and hormonal
24 information on the chemical and its analogues, which contributes to a working mode of action
25 and associates findings in animals with those in exposed humans. In comparison with other
26 agents designated as likely carcinogens, the overall weight of evidence for this chemical places it
27 at the *lower end* of the grouping. This is because there is a lack of tumor response data on this
28 agent itself.

29 Biological information on the compound is contradictory in terms of how to quantitate
30 potential cancer risks. The information on disruption on thyroid-pituitary status argues for using

1 a margin of exposure evaluation. However, the chemical is an aromatic amine, a class of agents
2 that are DNA-reactive and induce gene mutation and chromosome aberrations, which argues for
3 low-dose linearity. Additionally, there is a lack of mode of action information on the mouse
4 liver tumors produced by the structural analogues, also pointing toward a low-dose linear default
5 approach. In recognition of these uncertainties, it is recommended to quantitate tumors using
6 *both nonlinear* (to place a lower bound on the risks) *and linear* (to place an upper bound on the
7 risks) *default approaches*. Given the absence of tumor response data on the chemical per se, it is
8 recommended that tumor data on close analogues be used to possibly develop toxicity equivalent
9 factors or relative potencies.

10 Overall, this chemical is an inferential case for potential human carcinogenicity. The
11 uncertainties associated with this assessment include (1) the lack of carcinogenicity studies on
12 the chemical, (2) the use of tumor data on structural analogues, (3) the lack of definitive
13 information on the relevance of thyroid-pituitary imbalance for human carcinogenicity, and (4)
14 the different potential mechanisms that may influence tumor development and potential risks.

16 **SUPPORTING INFORMATION**

17 **Human Data**

18 Worker exposure has not been well characterized or quantified, but recent medical
19 monitoring of workers exposed over a period of several years has uncovered alterations in
20 thyroid-pituitary hormones (a decrease in T3 and T4 and an increase in TSH) and symptoms of
21 hypothyroidism. A urinary metabolite of the chemical has been monitored in workers, with
22 changes in thyroid and pituitary hormones noted, and the changes were similar to those seen in
23 an animal study.

25 **Animal Data**

26 The concentration of the urinary metabolite in rats receiving the chemical for 28 days
27 was within twofold of that in exposed workers, a finding associated with comparable changes in
28 thyroid hormones and TSH levels. In addition, the dose of the chemical given to rats in this
29 study was essentially the same as that of an analogue that had produced thyroid and pituitary
30 tumors in rats. The human thyroid responds in the same way as the rodent thyroid following

1 short-term, limited exposure. Although it is not well established that thyroid-pituitary imbalance
2 leads to cancer in humans as it does in rodents, information in animals and in exposed humans
3 suggests similar mechanisms of disrupting thyroid-pituitary function and the potential role of
4 altered TSH levels in leading to thyroid carcinogenesis.

5 6 **Structural Analogue Data**

7 This chemical is an aromatic amine, a member of a class of chemicals that has regularly
8 produced carcinogenic effects in rodents and gene and structural chromosome aberrations in
9 short-term tests. Some aromatic amines have produced cancer in humans.

10 Close structural analogues produce thyroid follicular cell tumors in rats and
11 hepatocellular tumors in mice following ingestion. The thyroid tumors are associated with
12 known perturbations in thyroid-pituitary functioning. These compounds inhibit the use of iodide
13 by the thyroid gland, apparently due to inhibition of the enzyme that synthesizes the thyroid
14 hormones (T3, T4). Accordingly, blood levels of thyroid hormones decrease, which induce the
15 pituitary gland to produce more TSH, a hormone that stimulates the thyroid to produce more of
16 its hormones. The thyroid gland becomes larger due to increases in the size of individual cells
17 and their proliferation and upon chronic administration, tumors develop. Thus, thyroid tumor
18 development is significantly influenced by disruption in the thyroid-pituitary axis.

19 20 **Other Key Data**

21 The chemical can be absorbed by the oral, inhalation, and dermal routes of exposure.
22

23 **MODE OF ACTION**

24 Data on the chemical and on structural analogues indicate the potential association of
25 carcinogenesis with perturbation of thyroid-pituitary homeostasis. Structural analogues are
26 genotoxic, thus raising the possibility of different mechanisms by which this chemical may
27 influence tumor development.
28

1 **NARRATIVE #5**

2 **Brominated Alkane (BA)**

3 **CAS# XXX**

4
5 **CANCER HAZARD SUMMARY**

6 Brominated alkane (BA) is *likely* to be a human carcinogen by *all routes* of exposure.
7 The weight of evidence for human carcinogenicity is at the *high end* of agents in the "likely"
8 group. Findings are based on very extensive and significant experimental findings that include
9 (a) tumors at multiple sites in both sexes of two rodent species via three routes of administration
10 relevant to human exposure, (b) close structural analogues that produce a spectrum of tumors
11 like BA, (c) significant evidence for the production of reactive BA metabolites that readily bind
12 to DNA and produce gene mutations in many systems including cultured mammalian and human
13 cells, and (d) two null and one positive epidemiologic study; in the positive study, there may
14 have been exposure to BA. These findings support a decision that BA might produce cancer in
15 exposed humans. In comparison to other agents considered likely human carcinogens, the
16 overall weight of evidence for BA puts it near the top of the grouping. Given the agent's
17 mutagenicity, which can influence the carcinogenic process, a linear dose-response extrapolation
18 is recommended.

19 Uncertainties include the lack of adequate information on the mutagenicity of BA in
20 mammals or humans *in vivo*, although such effects would be expected.

21
22 **SUPPORTING INFORMATION**

23 **Human Data**

24 The information on the carcinogenicity of BA from human studies is inadequate. Two
25 studies of production workers have not shown significant increases in cancer from exposure to
26 BA and other chemicals. An increase in lymphatic cancer was reported in a mortality study of
27 grain elevator workers who may have been exposed to BA (and other chemicals).

28
29 **Animal Data**

1 BA produced tumors in four chronic rodents studies. Tumor increases were noted in
2 males and females of rats and mice following oral dermal and inhalation exposure (rat--oral and
3 two inhalation, mouse--oral and dermal). It produces tumors both at the site of application (e.g.,
4 skin with dermal exposure) and at sites distal to the portal of entry into the body (e.g., mammary
5 gland) following exposure from each route. Tumors at the same site were noted in both sexes of
6 a species (blood vessel), both species (forestomach) and via different routes of administration
7 (lung). Some tumors developed after very short latency, metastasized extensively, and produced
8 death, an uncommon findings in rodents. The rodent studies were well designed and conducted
9 except for the oral studies, in which the doses employed caused excessive toxicity and mortality.
10 However, given the other rodent findings, lower doses would also be anticipated to be
11 carcinogenic.

12 13 **Structural Analogue Data**

14 Several chemicals structurally related to BA are also carcinogenic in rodents. Among
15 four that are closest in structure, tumors like those seen for BA were often noted (e.g.,
16 forestomach, mammary, lung), which helps to confirm the findings for BA itself. In sum, all of
17 the tumor findings help to establish animal carcinogenicity and support potential human
18 carcinogenicity for BA.

19 20 **Other Key Data**

21 BA itself is not reactive, but from its structure it was expected to be metabolized to
22 reactive forms. Extensive metabolism studies have confirmed this presumption and have
23 demonstrated metabolites that bind to DNA and cause breaks in the DNA chain. These lesions
24 are readily converted to gene mutations in bacteria, fungi, higher plants, insects and mammalian
25 and human cells in culture. There are only a limited number of reports on the induction of
26 chromosome aberrations in mammals and humans; thus far they are negative.

27 28 **MODE OF ACTION**

29 Human carcinogens often produce cancer in multiple sites of multiple animal species and
30 both sexes and are mutagenic in multiple test systems. BA satisfies these findings. It produces

cancer in males and females of rats and mice. It produces gene mutations in cells across all life forms--plants, bacteria and animals--including mammals and humans. Given the mutagenicity of BA exposure and the multiplicity and short latency of BA tumor induction, it is reasonable to use a linear approach for cancer dose-response extrapolation.

BRIEFING SUMMARY

<u>Route(s)</u>	<u>Class</u>	<u>Designation or rationale</u>	<u>Dose Response</u>
all	likely	high end	default-linear

Basis for classification/dose response

- Human data:** Two studies of production workers show no increase in cancer (one had a small sample size; the other had mixed chemical exposures). An increase in lymphatic cancer is seen among grain elevator workers who may have been exposed to other chemicals.
- Animal data:** BA produces tumors at multiple sites in male and female rats and mice following oral, dermal, and inhalation exposure. Tumors are seen at the site of administration and distally and are often consistent across sex, species, and route of administration; some develop early, metastasize, and cause death.
- Structural analogue data:** Close analogues produce some of the same tumors as are seen with BA.
- Other key data:** BA is metabolized to a reactive chemical that binds DNA and produces gene mutations in essentially every test system including cultured human cells.
- Mode of action:** Like most known human carcinogens, BA is mutagenic in most test systems.
- Hazard classification/uncertainties:** There is a rich database on BA demonstrating its potential ability to cause tumors in humans, including (a) multiple animal tumors, (b) by appropriate routes of exposure, (c) a mode of action relevant to human carcinogenicity, and (d) some information in humans. Together they lead to a designation near the high end of the *likely* human carcinogen class.

- 1 **7. Dose response:** Given the anticipated mode of action, a linear default dose response
- 2 relationship should be assumed.
- 3

APPENDIX B

This appendix contains responses to the National Academy of Sciences National Research Council report *Science and Judgment in Risk Assessment* (NRC, 1994).

Recommendations of the National Academy of Sciences National Research Council

In 1994, the National Academy of Sciences published a report *Science and Judgment in Risk Assessment*. The 1994 report was written by a Committee on Risk Assessment of Hazardous Air Pollutants formed under the Academy's Board on Environmental Studies and Toxicology, Commission on Life Sciences, National Research Council. The report was called for under Section 112(o)(1)(A,B) of the Clean Air Act Amendments of 1990, which provided for the EPA to arrange for the Academy to review:

- risk assessment methodology used by the EPA to determine the carcinogenic risk associated with exposure to hazardous air pollutants from source categories and subcategories subject to the requirements of this section and
- improvements in such methodology.

Under Section 112(o)(2)(A,B), the Academy was to consider the following in its review:

- the techniques used for estimating and describing the carcinogenic potency to humans of hazardous air pollutants and
- the techniques used for estimating exposure to hazardous air pollutants (for hypothetical and actual maximally exposed individuals as well as other exposed individuals).

To the extent practicable, the Academy was also to review methods of assessing adverse human health effects other than cancer for which safe thresholds of exposure may not exist [Section 112(o)(3)]. The Congress further provided that the EPA Administrator should consider, but need not adopt, the recommendations in the report and the views of the EPA Science Advisory Board with respect to the report. Prior to the promulgation of any standards under Section 112(f), the Administrator is to publish revised guidelines for carcinogenic risk assessment or a detailed explanation of the reasons that any recommendations contained in the report will not be implemented [Section 112(o)(6)].

1 The following discussion addresses the recommendations of the 1994 report that are
2 pertinent to the EPA cancer risk assessment guidelines. Guidelines for assessment of exposure,
3 of mixtures, and of other health effects are separate EPA publications. Many of the
4 recommendations were related to practices specific to the exposure assessment of hazardous air
5 pollutants, which are not covered in cancer assessment guidelines. Recommendations about
6 these other guidelines or practices are not addressed here.

7 8 **Hazard Classification**

9 The 1994 report contains the following recommendation about classifying cancer hazard:

- 10 • The EPA should develop a two-part scheme for classifying evidence on
11 carcinogenicity that would incorporate both a simple classification and a narrative
12 evaluation. At a minimum, both parts should include the strength (quality) of the
13 evidence, the relevance of the animal model and results to humans, and the relevance
14 of the experimental exposures (route, dose, timing, and duration) to those likely to be
15 encountered by humans.

16 The report also presented a possible matrix of 24 boxes that would array weights of evidence
17 against low, medium, or high relevance, resulting in 24 codes for expressing the weight and
18 relevance.

19 These guidelines adopt a set of descriptors and subdescriptors of weight of evidence in
20 three categories: “known/likely,” “cannot be determined,” and “not likely,” and a narrative for
21 presentation of the weight of evidence findings. The descriptors are used within the narrative.
22 There is no matrix of alphanumerical weight of evidence boxes.

23 The issue of an animal model that is not relevant to humans has been dealt with by not
24 including an irrelevant response in the weighing of evidence, rather than by creating a weight of
25 evidence then appending a discounting factor as the NRC scheme would do. The issue is more
26 complex than the NRC matrix makes apparent. Often the question of relevance of the animal
27 model applies to a single tumor response, but one encounters situations in which there are more
28 tumor responses in animals than the questioned one. Dealing with this complexity is more
29 straightforward if it is done during the weighing of evidence rather than after as in the NRC

1 scheme. Moreover, the same experimental data are involved in deciding on the weight of
2 evidence and the relevance of a response. It would be awkward to go over the same data twice.

3 In recommending that the relevance of circumstances of human exposure also be taken
4 into account, the NRC appears to assume that all of the actual conditions of human exposure will
5 be known when the classification is done. This is not the case. More often than not, the hazard
6 assessment is applied to assessment of risks associated with exposure to different media or
7 environments at different times. In some cases, there is no priority to obtaining exposure data
8 until the hazard assessment has been done. The approach of these guidelines is to characterize
9 hazards as to whether their expression is intrinsically limited by route of exposure or by reaching
10 a particular dose range based strictly on toxicological and other biological features of the agent.
11 Both the use of descriptors and the narrative specifically capture this information. Other aspects
12 of appropriate application of the hazard and dose response assessment to particular human
13 exposure scenarios are dealt with in the characterization of the dose response assessment, e.g.,
14 the applicability of the dose response assessment to scenarios with differing frequencies and
15 durations.

16 The NRC scheme apparently intended that the evidence would be weighed, then given a
17 low, medium, or high code for some combination of relevance of the animal response, route of
18 exposure, timing, duration, or frequency. The 24 codes contain none of this specific
19 information, and in fact, do not communicate what the conclusion is about. To make the codes
20 communicate the information apparently intended would require some multiple of the 24 in the
21 NRC scheme. As the number of codes increases, their utility for communication decreases.

22 Another reason for declining to use codes is that they tend to become outdated as
23 research reveals new information that was not contemplated when they were adopted. This has
24 been the case with the classification system under the EPA, 1986 guidelines.

25 Even though these guidelines do not adopt a matrix of codes, the method they provide of
26 using descriptors and narratives captures the information the NRC recommended as the most
27 important, and in the EPA's view, in a more transparent manner.

Dose Response

The 1994 report contains the following recommendations about dose response issues:

- EPA should continue to explore, and when scientifically appropriate, incorporate toxicokinetic models of the link between exposure and biologically effective dose (i.e., dose reaching the target tissue).
- Despite the advantages of developing consistent risk assessments between agencies by using common assumptions (e.g., replacing surface area with body weight to the 0.75 power), EPA should indicate other methods, if any, that would be more accurate.
- EPA should continue to use the linearized multistage model as a default option but should develop criteria for determining when information is sufficient to use an alternative extrapolation model.
- EPA should continue to use as one of its risk characterization metrics upper-bound potency estimates of the probability of developing cancer due to lifetime exposure. Whenever possible, this metric should be supplemented with other descriptions of cancer potency that might more adequately reflect the uncertainty associated with the estimates.
- EPA should adopt a default assumption for differences in susceptibility among humans in estimating individual risks.
- In the analysis of animal bioassay data on the occurrence of multiple tumor types, the cancer potencies should be estimated for each relevant tumor type that is related to exposure and the individual potencies should be summed for those tumors.

The use of toxicokinetic models is encouraged in these guidelines with discussion of appropriate considerations for their use. When there are questions as to whether such a model is more accurate in a particular case than the default method for estimating the human equivalent dose, both alternatives may be used. It should be noted that the default method for inhalation exposure is a toxicokinetic model.

The rationale for adopting the oral scaling factor of body weight to the 0.75 power has been discussed above in the explanation of major defaults. The empirical basis is further explored in U.S. EPA, 1992b. The more accurate approach is to use a toxicokinetic model when data become available or to modify the default when data are available as encouraged under

1 these guidelines. As the U.S. EPA, 1992b discussion explores in depth, data on the differences
2 among animals in response to toxic agents are basically consistent with using a power of 1.0,
3 0.75, or 0.66. The Federal agencies chose the power of 0.75 for the scientific reasons given in
4 the previous discussion of major defaults; these were not addressed specifically in the NRC
5 report. It was also considered appropriate, as a matter of policy, for the agencies to agree on one
6 factor. Again, the default for inhalation exposure is a model that is constructed to become better
7 as more agent-specific data become available.

8 The EPA proposes not to use a computer model such as the linearized multistage model
9 as a default for extrapolation below the observed range. The reason is that the basis for default
10 extrapolation is a theoretical projection of the likely shape of the curve considering mode of
11 action. For this purpose, a computer model looks more sophisticated than a straight line
12 extrapolation, but is not. The extrapolation will be by straight line as explained in the
13 explanation of major defaults. This was also recommended by workshop reviewers of a
14 previous draft of these guidelines (U.S. EPA, 1994b). In addition, a margin of exposure analysis
15 is proposed to be used in cases in which the curve is thought to be nonlinear, based on mode of
16 action. In both cases, the observed range of data will be modeled by curve fitting in the absence
17 of supporting data for a biologically based or case-specific model.

18 The result of using straight line extrapolation is thought to be an upper bound on low-
19 dose potency to the human population in most cases, but as discussed in the major defaults
20 section, it may not always be. Exploration and discussion of uncertainty of parameters in curve-
21 fitting a model of the observed data or in using a biologically based or case-specific model is
22 called for in the dose response assessment and characterization sections of these guidelines.

23 The issue of a default assumption for human differences in susceptibility has been
24 addressed under the major defaults discussion in section 1.3 with respect to margin of exposure
25 analysis. The EPA has considered but decided not to adopt a quantitative default factor for
26 human differences in susceptibility when a linear extrapolation is used. In general, the EPA
27 believes that the linear extrapolation is sufficiently conservative to protect public health. Linear
28 approaches (both LMS and straight line extrapolation) from animal data are consistent with
29 linear extrapolation on the same agents from human data (Goodman and Wilson, 1991; Hoel and

Portier, 1994). If actual data on human variability in sensitivity are available they will, of course, be used.

In analyzing animal bioassay data on the occurrence of multiple tumor types, these guidelines outline a number of biological and other factors to consider. The objective is to use these factors to select response data (including nontumor data as appropriate) that best represent the biology observed. As stated in section 3 of the guidelines, appropriate options include use of a single data set, combining data from different experiments, showing a range of results from more than one data set, showing results from analysis of more than one tumor response based on differing modes of action, representing total response in a single experiment by combining animals with tumors, or a combination of these options. The approach judged to best represent the data is presented with the rationale for the judgment, including the biological and statistical considerations involved. The EPA has considered the approach of summing tumor incidences and decided not to adopt it. While multiple tumors may be independent, in the sense of not arising from metastases of a single malignancy, it is not clear that they can be assumed to represent different effects of the agent on cancer processes. In this connection, it is not clear that summing incidences provides a better representation of the underlying mode(s) of action of the agent than combining animals with tumors or using another of the several options noted above. Summing incidences would result in a higher risk estimate, a step that appears unnecessary without more reason.

Risk Characterization

- When EPA reports estimates of risk to decisionmakers and the public, it should present not only point estimates of risk, but also the sources and magnitudes of uncertainty associated with these estimates.
- Risk managers should be given characterizations of risk that are both qualitative and quantitative, i.e., both descriptive and mathematical.
- EPA should consider in its risk assessments the limits of scientific knowledge, the remaining uncertainties, and the desire to identify errors of either overestimation or underestimation.

1 In part as a response to these recommendations, the Administrator of EPA issued
2 guidelines for risk characterization and required implementation plans from all programs in EPA
3 (U.S. EPA, 1995). The Administrator's guidance is followed in these cancer guidelines. The
4 assessments of hazard, dose response, and exposure will all have accompanying technical
5 characterizations covering issues of strengths and limitations of data and current scientific
6 understanding, identification of defaults utilized in the face of gaps in the former, discussions of
7 controversial issues, and discussions of uncertainties in both their qualitative, and as practicable,
8 their quantitative aspects.

APPENDIX C

OVERVIEW OF CANCER PROCESSES

The following picture is changing as research reveals more about carcinogenic processes. Nevertheless, it is apparent that several general modes of action are being elucidated from direct reaction with DNA to hormonal or other growth-signaling processes. While the exact mechanism of action of an agent at the molecular level may not be clear from existing data, the available data will often provide support for deducing the general mode of action. Under these guidelines, using all of the available data to arrive at a view of the mode of action supports both characterization of human hazard potential and assessment of dose response relationships.

Cancers are diseases of somatic mutation affecting cell growth and differentiation. The genes that control cell growth, programmed cell death, and cell differentiation are critical to normal development of tissues from embryo to adult metazoan organisms. These genes continue to be critical to maintenance of form and function of tissues in the adult (e.g., Meyn, 1993) and changes in them are essential elements of carcinogenesis (Hsu et al., 1991; Kakizuka et al., 1991; Bottaro et al., 1991; Sidransky et al., 1991; Salomon et al., 1990; Srivastava et al., 1990). The genes involved are among the most highly conserved in evolution as evidenced by the great homology of many of them in DNA sequence and function in organisms as phylogenetically distant as worms, insects, and mammals (Auger et al., 1989a, b; Hollstein et al., 1991; Herschman, 1991; Strausfeld et al., 1991; Forsburg and Nurse, 1991).

Mutations affecting three general categories of genes have been implicated in carcinogenesis. Over 100 oncogenes have been found in human and animal tumors that act as dominant alleles, whereas there are about 10 known tumor suppressor genes that are recessive in action. The normal alleles of these genes are involved with control of cell division and differentiation; mutated alleles lead to a disruption in these functions. The third class are mutator genes that predispose the genome to enhanced mutagenic events that contribute further to the carcinogenic process.

Adult tissues, even those that are composed of rapidly replicating cells, maintain a constant size and cell number (Nunez et al., 1991) by balancing three cell fates: (1) continued replication, (2) differentiation to take on specialized functions, or (3) programmed cell death

(apoptosis) (Raff, 1992; Maller, 1991; Naeve et al., 1991; Schneider et al., 1991; Harris, 1990). Neoplastic growth through clonal expansion can result from somatic mutations that inactivate control over cell fate (Kakizuka et al., 1991; deThe et al., 1991; Sidransky et al., 1992; Nowell, 1976).

Cancers may also be thought of as diseases of the cell cycle. For example, genetic diseases that cause failure of cells to repair DNA damage prior to cell replication predispose people to cancer. These changes are also frequently found in tumor cells in sporadic cancers. These changes appear to be particularly involved at points in cell replication called "checkpoints" where DNA synthesis or mitosis is normally stopped until DNA damage is repaired or cell death induced (Tobey, 1975). A cell that bypasses a checkpoint may acquire a heritable growth advantage. Similar effects on the cell cycle occur when mitogens such as hormones or growth factors stimulate cell growth. Rapid replication in response to tissue injury may also lead to unrepaired DNA damage that is a risk factor for carcinogenesis.

Normally a cell's fate is determined by a timed sequence of biochemical signals. Signal transduction in the cell involves chemical signals that bind to receptors, generating further signals in a pathway whose target in many cases is control of transcription of a specific set of genes (Hunter, 1991; Cantley et al., 1991; Collum and Alt, 1990). Cells are subject to growth signals from the same and distant tissues, e.g., endocrine tissues (Schuller, 1991). In addition to hormones produced by endocrine tissues, numerous soluble polypeptide growth factors have been identified that control normal growth and differentiation (Cross and Dexter, 1991; Wellstein et al., 1990). The cells responsive to a particular growth factor are those that express transmembrane receptors that specifically bind the growth factor.

Solid tumors develop in stages operationally defined as initiation, promotion, and progression (see, for example, Pitot and Dragan, 1991). These terms, which were coined in the context of specific experimental designs, are used for convenience in discussing concepts, but they refer to complex events that are not completely understood. During initiation, the cell acquires a genetic change that confers a potential growth advantage. During promotion, clonal expansion of this altered cell occurs. Later, during progression, a series of genetic and other biological events both enhance the growth advantage of the cells and enlist normal host processes to support tumor development and cells develop the ability to invade locally and

1 metastasize distally, taking on the characteristics of malignancy. Many endogenous and
2 exogenous factors are known to participate in the process as a whole. These include specific
3 genetic predispositions or variations in ability to detoxify agents, medical history (Harris, 1989;
4 Nebreda et al., 1991), infections, exposure to chemicals or ionizing radiation, hormones and
5 growth factors, and immune suppression. Several such risk factors likely work together to cause
6 individual human cancers.

7 A cell that has been transformed, acquiring the potential to establish a line of cells that
8 grow to a tumor, will probably realize that potential only rarely. The process of tumorigenesis
9 in animals and humans is a multistep one (Bouk, 1990; Fearon and Vogelstein, 1990; Hunter,
10 1991; Kumar et al., 1990; Sukumar, 1989; Sukumar, 1990) and normal physiological processes
11 appear to be arrayed against uncontrolled growth of a transformed cell (Weinberg, 1989).
12 Powerful inhibition by signals from contact with neighboring normal cells is one known barrier
13 (Zhang et al., 1992). Another is the immune system (at least for viral infection). How a cell
14 with tumorigenic potential acquires additional properties that are necessary to enable it to
15 overcome these and other inhibitory processes is a subject of ongoing research. For known
16 human carcinogens studied thus far, there is an often decades-long latency between exposure to
17 carcinogenic agents and development of tumors (Fidler and Radinsky, 1990; Tanaka et al., 1991;
18 Thompson et al., 1989). This latency is also typical of tumor development in individuals with
19 genetic diseases that make them prone to cancer (Meyn, 1993; Srivastava et al., 1990).

20 The importance of genetic mutation in the carcinogenic process calls for special attention
21 to assessing agents that cause such mutations. Heritable genetic defects that predispose humans
22 to cancer are well known and the number of identified defects is growing. Examples include
23 xeroderma pigmentosum (DNA repair defect) and Li Fraumeni and retinoblastoma (both are
24 tumor suppressor gene mutations). Much of the screening and testing of agents for carcinogenic
25 potential has been driven by the idea of identifying this mode of action. Cognizance of and
26 emphasis on other modes of action such as ones that act at the level of growth signalling within
27 or between cells, through cell receptors, or that indirectly cause genetic change, comes from
28 more recent research. There are not yet standardized tests for many modes of action, but
29 pertinent information may be available in individual cases.

1 Agents of differing characteristics influence cancer development: inorganic and organic,
2 naturally occurring and synthetic, of inanimate or animate origin, endogenous or exogenous,
3 dietary and nondietary. The means by which these agents act to influence carcinogenesis are
4 variable, and reasoned hazard assessment requires consideration of the multiple ways that
5 chemicals influence cells in experimental systems and in humans. Agents exert mutagenic
6 effects either by interacting directly with DNA or by indirect means through intermediary
7 substances (e.g., reactive oxygen species) or processes. Most DNA-reactive chemicals are
8 electrophilic or can become electrophilic when metabolically activated. Electrophilic molecules
9 may bind covalently to DNA to form adducts, and this may lead to depurination,
10 depyrimidation, or produce DNA strand breaks; such lesions can be converted to mutations with
11 a round of DNA synthesis and cell division. Other DNA-interactive chemicals may cause the
12 same result by intercalation into the DNA helix. Still other chemicals may methylate DNA,
13 changing gene expression. Non-DNA-reactive chemicals produce genotoxic effects by many
14 different processes. They may affect spindle formation or chromosome proteins, interfere with
15 normal growth control mechanisms, or affect enzymes involved with ensuring the fidelity of
16 DNA synthesis (e.g., topoisomerase), recombination, or repair.

17 The "classical" chemical carcinogens in laboratory rodent studies are agents that
18 consistently produce gene mutations and structural chromosome aberrations in short-term tests.
19 A large database reveals that these mutagenic substances commonly produce tumors at multiple
20 sites and in multiple species (Ashby and Tennant, 1991). Most of the carcinogens identified in
21 human studies, aside from hormones, are also gene or structural chromosome mutagens (Tennant
22 and Ashby, 1991). Most of these compounds or their metabolites contain electrophilic moieties
23 that react with DNA.

24 Numerical chromosome aberrations, gene amplification, and the loss of gene
25 heterozygosity are also found in animal and human tumor cells and may arise from initiating
26 events or during progression. There is reason to believe that accumulation of additional genetic
27 changes is favored by selection in the evolution of tumor cells because they confer additional
28 growth advantages (Hartwell and Kastan, 1994). Exogenous agents may function at any stage of
29 carcinogenesis (Barrett, 1993). Some aberrations may arise as a consequence of genomic
30 instability arising from tumor suppressor gene mutation, e.g., p53 (Harris and Hollstein, 1993).

1 The frequent observation in tumor cells that both of a pair of homologous chromosomes have
2 identical mutation spectra in tumor suppressor genes suggests an ongoing, endogenous process
3 of gene conversion. Currently, there is a paucity of routine test methods to screen for events
4 such as gene conversion or gene amplification and knowledge regarding the ability of particular
5 agents of environmental interest to induce them is, for the most part, wanting. Work is under
6 way to characterize, measure, and evaluate their significance (Travis et al., 1991).

7 Several kinds of mechanistic studies aid in risk assessment. Comparison of DNA lesions
8 in tumor cells taken from humans with the lesions that a tumorigenic agent causes in
9 experimental systems can permit inferences about the association of exposure to the agent and an
10 observed human effect (Vahakangas et al., 1992; Hollstein et al., 1991; Hayward et al., 1991).
11 An agent that is observed to cause mutations experimentally may be inferred to have potential
12 for carcinogenic activity (U.S. EPA, 1991a). If such an agent is shown to be carcinogenic in
13 animals, the inference that its mode of action is through mutagenicity is strong. A carcinogenic
14 agent that is not mutagenic in experimental systems but is mitogenic or affects hormonal levels
15 or causes toxic injury followed by compensatory growth may be inferred to have effects on
16 growth signal transduction or to have secondary carcinogenic effects. The strength of these
17 inferences depends in each case on the nature and extent of all the available data.

18 Differing modes of action at the molecular level have different dose response
19 implications for the activity of agents. The carcinogenic activity of a direct-acting mutagen
20 should be a function of the probability of its reaching and reacting with DNA. The carcinogenic
21 activity of an agent that interferes at the level of signal pathways with many potential receptor
22 targets should be a function of multiple reactions. The carcinogenic activity of an agent that acts
23 by causing cell toxicity followed by compensatory growth should be a function of the toxicity.

REFERENCES

- Alison, R.H.; Capen, C.C.; Prentice, D.E. (1994) Neoplastic lesions of questionable significance to humans. *Toxicol. Pathol.* 22: 179-186.
- Allen, B.C.; Crump, K.S.; Shipp, A.M. (1988) Correlation between carcinogenic potency of chemicals in animals and humans. *Risk Anal.* 8: 531-544.
- Ames, B.N.; Gold, L.S. (1990) Too many rodent carcinogens: mitogenesis increases mutagenesis. *Science* 249: 970-971.
- Anderson, E.; Deisler, P.F.; McCallum, D.; St. Helaire, C.; Spitzer, H.L.; Strauss, H.; Wilson, J.D.; Zimmerman, R. (1993) Key issues in carcinogen risk assessment guidelines. Society for Risk Analysis.
- Ashby, J.; Tennant, R.W. (1991) Definitive relationships among chemical structure, carcinogenicity and mutagenicity for 301 chemicals tested by the U.S. NTP. *Mutat. Res.* 257: 229-306.
- Ashby, J.; Tennant, R.W. (1994) Prediction of rodent carcinogenicity for 44 chemicals: results. *Mutagenesis* 9: 7-15.
- Ashby, J.; Doerrler, N.G.; Flamm, F.G.; Harris, J.E.; Hughes, D.H.; Johannsen, F.R.; Lewis, S.C.; Krivanek, N.D.; McCarthy, J.F.; Moolenaar, R.J.; Raabe, G.K.; Reynolds, R.C.; Smith, J.M.; Stevens, J.T.; Teta, M.J.; Wilson, J.D. (1990) A scheme for classifying carcinogens. *Regul. Toxicol. Pharmacol.* 12: 270-295.
- Ashby, J.; Brady, A.; Elcombe, C.R.; Elliott, B.M.; Ishmael, J.; Odum, J.; Tugwood, D.; Keltle, S.; Purchase, I.F.H. (1994) Mechanistically based human hazard assessment of peroxisome proliferator-induced hepatocarcinogenesis. *Hum. Exper. Toxicol.* 13: 1-117.
- Auger, K.R.; Carpenter, C.L.; Cantley, L.C.; Varticovski, L. (1989a) Phosphatidylinositol 3-kinase and its novel product, phosphatidylinositol 3-phosphate, are present in *Saccharomyces cerevisiae*. *J. Biol. Chem.* 264: 20181-20184.
- Auger, K.R.; Sarunian, L.A.; Soltoff, S.P.; Libby, P.; Cantley, L.C. (1989b) PDGF-dependent tyrosine phosphorylation stimulates production of novel polyphosphoinositides in intact cells. *Cell* 57: 167-175.
- Barnes, D.G.; Daston, G.P.; Evans, J.S.; Jarabek, A.M.; Kavlock, R.J.; Kimmel, C.A.; Park, C.; Spitzer, H.L. (1995) Benchmark dose workshop: criteria for use of a benchmark dose to estimate a reference dose. *Regul. Toxicol. Pharmacol.* 21: 296-306.
- Barrett, J.C. (1992) Mechanisms of action of known human carcinogens. In: *Mechanisms of carcinogenesis in risk identification*. IARC Sci. Pubs. No. 116, Lyon, France: International Agency for Research on Cancer; 115-134.
- Barrett, J.C. (1993) Mechanisms of multistep carcinogenesis and carcinogen risk assessment. *Environ. Health Perspect.* 100: 9-20.
- Barrett, J.C. (1995) Role of mutagenesis and mitogenesis in carcinogenesis. *Environ. Mutagenesis*, in press.
- Barrett, J. C.; Lee, T. C. (1992) Mechanisms of arsenic-induced gene amplification. In: *Gene amplification in mammalian cells: A comprehensive guide* (ed. R. E. Kellems), Marcel Dekker, New York: 441-446.

- 1 Bayly, A.C.; Roberts, R.A.; Dive, C. (1994) Suppression of liver cell apoptosis in vitro by the nongenotoxic
2 hepatocarcinogen and peroxisome proliferator nafenopin. *J. Cell. Biol.* 125: 197-203.
- 3
- 4 Bellamy, C.O.C.; Malcomson, R.D.G.; Harrison, D.J.; Wyllie, A.H. (1995) Cell death in health and disease: The
5 biology and regulation of apoptosis. *Seminars in cancer biology, Apoptosis in oncogenesis and chemotherapy* 6:
6 3-16.
- 7
- 8 Bianchi, A.B.; Navone, N.M.; Alda, C.M.; Conti, C.J. (1991) Overlapping loss of heterozygosity by mitotic
9 recombination on more chromosome 7F1-ter in skin carcinogenesis. *Proc. Nat. Acad. Sci.* 88: 7590-7594.
- 10
- 11 Biggs, P.J.; Warren, W.; Venitt, S.; Stratton, M.R. (1993) Does a genotoxic carcinogen contribute to human
12 breast cancer? The value of mutational spectra in unraveling the etiology of cancer. *Mutagenesis* 8: 275-283.
- 13
- 14 Birner et al. (1990) Biomonitoring of aromatic amines. III: Hemoglobin binding and benzidine and some
15 benzidine congeners. *Arch. Toxicol.* 64(2): 97-102.
- 16
- 17 Blair, A.; Burg, J.; Foran, J.; Gibb, H.; Greenland, S.; Morris, R.; Raabe, G.; Savitz, D.; Teta, J.; Wartenberg, D.;
18 Wong, O.; Zimmerman, R. (1995) Guidelines for application of meta-analysis in environmental epidemiology.
19 *Regul. Toxicol. Pharmacol.* 22: 189-197.
- 20
- 21 Bois, F.Y.; Krowech, G.; Zeise, L. (1995) Modeling human interindividual variability in metabolism and risk: the
22 example of 4-aminobiphenyl. 15: 205-213.
- 23
- 24 Bottaro, D.P.; Rubin, J.S.; Faletto, D.L.; Chan, A.M.L.; Kmieck, T.E.; Vande Woude, G.F.; Aaronson, S.A.
25 (1991) Identification of the hepatocyte growth factor receptor as the *c-met* proto-oncogene product. *Science* 251:
26 802-804.
- 27
- 28 Bouck, N. (1990) Tumor angiogenesis: the role of oncogenes and tumor suppressor genes. *Cancer Cells* 2: 179-
29 183.
- 30
- 31 Burek, J.D.; Patrick, D.H.; Gerson, R.J. (1988) Weight-of-biological evidence approach for assessing
32 carcinogenicity. In: Grice, H.C.; Cimina, J.L., eds. *Carcinogenicity*. New York, NY: Springer Verlag; pp. 83-95.
- 33
- 34 Bus, J.S.; Popp, J.A. (1987) Perspectives on the mechanism of action of the splenic toxicity of aniline and
35 structurally related compounds. *Fd. Chem. Toxicol.* 25: 619-626.
- 36
- 37 Callemen, C.J.; Ehrenberg, L.; Jansson, B.; Osterman-Golkar, S.; Segerback, D.; Svensson, K.; Wachtmeister,
38 C.A. (1978) Monitoring and risk assessment by means of alkyl groups in hemoglobin in persons occupationally
39 exposed to ethylene oxide. *J. Environ. Pathol. Toxicol.* 2: 427-442.
- 40
- 41 Cantley, L.C.; Auger, K.R.; Carpenter, C.; Duckworth, B.; Graziani, A.; Kapeller, R.; Soltoff, S. (1991)
42 Oncogenes and signal transduction. *Cell* 64: 281-302.
- 43
- 44 Caporaso, N.; Hayes, R.B.; Dosemeci, M.; Hoover, R.; Ayesh, R.; Hetzel, M.; Idle, J. (1989) Lung cancer risk,
45 occupational exposure, and the debrisoquine metabolic phenotype. *Cancer Res.* 49: 3675-3679.
- 46
- 47 Cavenee, W. K.; Koufos, A.; Hansen, M. F. (1986) Recessive mutant genes predisposing to human cancer.
48 *Mutation Research* 168: 3-14.
- 49
- 50 Chang, C. C.; Jone, C.; Trosko, J. E.; Peterson, A. R.; Sevanian, A. (1988) Effect of cholesterol epoxides on the
51 inhibition of intercellular communication and on mutation induction in Chinese hamster V79 cells. *Mutation*
52 *Research* 206: 471-478.

- Chen, C.; Farland, W. (1991) Incorporating cell proliferation in quantitative cancer risk assessment: approaches, issues, and uncertainties. In: Butterworth, B.; Slaga, T.; Farland, W.; McClain, M., eds. Chemical induced cell proliferation: Implications for risk assessment. New York, NY: Wiley-Liss; pp. 481-499.
- Choy, W.N. (1993) A review of the dose-response induction of DNA adducts by aflatoxin B₂ and its implications to quantitative cancer-risk assessment. *Mutat. Res.* 296: 181-198.
- Clayson, D.B. (1989) Can a mechanistic rationale be provided for non-genotoxic carcinogens identified in rodent bioassays? *Mutat. Res.* 221: 53-67.
- Clayson, D.B.; Mehta, R.; Iverson, F. (1994) Oxidative DNA damage--The effects of certain genotoxic and operationally non-genotoxic carcinogens. *Mutat. Res.* 317: 25-42.
- Cogliano, V.J. (1986) The U.S. EPA's methodology for adjusting the reportable quantities of potential carcinogens. Proceedings of the 7th National Conference on Management of Uncontrollable Hazardous Wastes (Superfund '86). Washington, DC: Hazardous Wastes Control Institute, 182-185.
- Cohen, S.W.; Ellwein, L.B. (1990) Cell proliferation in carcinogenesis. *Science* 249: 1007-1011.
- Cohen, S.M.; Ellwein, L.B. (1991) Genetic errors, cell proliferation and carcinogenesis. *Cancer Res.* 51: 6493-6505.
- Cohen, S.M.; Purtilo, D.T.; Ellwein, L.B. (1991) Pivotal role of increased cell proliferation in human carcinogenesis. *Mod. Pathol.* 4: 371-375.
- Collum, R.G.; Alt, F.W. (1990) Are *myc* proteins transcription factors? *Cancer Cells* 2: 69-73.
- Connolly, R.B.; Andersen, M.E. (1991) Biologically based pharmacodynamic models: tools for toxicological research and risk assessment. *Ann. Rev. Pharmacol. Toxicol.* 31: 503-523.
- Cross, M.; Dexter, T. (1991) Growth factors in development, transformation, and tumorigenesis. *Cell* 64: 271-280.
- D'Souza, R.W.; Francis, W.R.; Bruce, R.D.; Andersen, M.E. (1987) Physiologically based pharmacokinetic model for ethylene chloride and its application in risk assessment. In: *Pharmacokinetics in risk assessment. Drinking Water and Health. Vol. 8.* Washington, DC: National Academy Press.
- deThe, H.; Lavau, C.; Marchio, A.; Chomienne, C.; Degos, L.; Dejean, A. (1991) The PML-RAR % fusion mRNA generated by the t (15;17) translocation in acute promyelocytic leukemia encodes a functionally altered RAR. *Cell* 66: 675-684.
- Enterline, P.E.; Henderson, V.L.; Marsh, G.M. (1987) Exposure to arsenic. *Amer. J. Epidemiol.* 125: 929-938.
- European Centre for Ecotoxicology and Toxicology of Chemicals (ECETOC). (1991) Early indicators of non-genotoxic carcinogenesis. ECETOC Monograph No. 16. Brussels: ECETOC. Printed in *Mutat. Res.* 248: 211-374.
- Faustman, E.M.; Allen, B.C.; Kavlock, R.J.; Kimmel, C.A. (1994) Dose-response assessment for developmental toxicity. *Fund. Appl. Toxicol.* 23: 478-486.
- Fearon, E.; Vogelstein, B. (1990) A genetic model for colorectal tumorigenesis. *Cell* 61: 959-967.

- 1 Federated Association of Societies of Experimental Biology (FASEB) (1994) Evaluation of evidence for the
2 carcinogenicity of butylated hydroxyanisole (BHA). Life Sciences Research Office, Bethesda, MD. Letter from
3 Hamilton Brown (FDA) to John Rice (FASEB), July 28, 1994. Letter from John Rice (FASEB) to Ed Arnold
4 (FDA), August 4, 1994.
- 5
- 6 Fidler, I.J.; Radinsky, R. (1990) Genetic control of cancer metastasis. J. Natl. Cancer Inst. 82: 166-168.
- 7
- 8 Fisher, R.A. (1950) Statistical methods for research workers. Edinburgh, Scotland: Oliver and Boyd.
- 9
- 10 Flamm, W.G.; Winbush, J.S. (1984) Role of mathematical models in assessment of risk and in attempts to define
11 management strategy. Fund. Appl. Toxicol. 4: S395-S401.
- 12
- 13 Flynn, G.L. (1990) Physicochemical determinants of skin absorption. In: Gerrity, T.R.; Henry, C.J., eds.
14 Principles of route to route extrapolation for risk assessment. New York, NY: Elsevier Science Publishing Co.;
15 pp. 93-127.
- 16
- 17 Forsburg, S.L.; Nurse, P. (1991) Identification of a G1-type cyclin *pug1+* in the fission yeast
18 *Schizosaccharomyces pombe*. Nature 351: 245-248.
- 19
- 20 Garfinkel, L.; Silverberg, E. (1991) Lung cancer and smoking trends in the United States over the past 25 years.
21 Cancer 41: 137-145.
- 22
- 23 Gaylor, D.W.; Kodell, R.L. (1980) Linear interpolation algorithm for low-dose risk assessment of toxic
24 substances. J. Environ. Pathol. Toxicol. 4: 305-312.
- 25
- 26 Gerrity, T.R.; Henry, C., eds. (1990) Principles of route to route extrapolation for risk assessment. New York,
27 NY: Elsevier Science Publishing Co.
- 28
- 29 Gibson, D.P.; Aardema, M.J.; Kerckaert, G.A.; Carr, G.J.; Brauninger, R.M.; LeBoeuf, R.A. (1995) Detection of
30 aneuploidy-inducing carcinogens in the Syrian hamster embryo (SHE) cell transformation assay. Mutat. Res.;
31 Genet. Toxicol. 343: 7-24.
- 32
- 33 Gillette, J.R. (1983) The use of pharmacokinetics in safety testing. In: Homburger, ed. Safety evaluation and
34 regulation of chemicals 2. 2nd Int. Conf., Cambridge, MA: Karger, Basel; pp. 125-133.
- 35
- 36 Goldsworthy, T.L.; Hanigan, M.H.; and Pitot, H.C. (1986) Models of hepatocarcinogenesis in the rat--contrasts
37 and comparisons. CRC Crit. Rev. Toxicol. 17: 61-89.
- 38
- 39 Goodman, G.; Wilson, R. (1991) Predicting the carcinogenicity of chemicals in humans from rodent bioassay
40 data. Environ. Health Perspect. 94: 195-218.
- 41
- 42 Goodman, J.I.; Counts, J.L. (1993) Hypomethylation of DNA: A possible nongenotoxic mechanism underlying
43 the role of cell proliferation in carcinogenesis. Environ. Health Perspect. 101 Supp. 5: 169-172.
- 44
- 45 Goodman, J.I.; Ward, J.M.; Popp, J.A.; Klaunig, J.E.; Fox, T.R. (1991) Mouse liver carcinogenesis: Mechanisms
46 and relevance. Fund. Appl. Toxicol. 17: 651-665.
- 47
- 48 Grasso, P.; Hinton, R.H. (1991) Evidence for and possible mechanisms of non-genotoxic carcinogenesis in rodent
49 liver. Mutat. Res. 248: 271-290.
- 50
- 51 Greenland, S. (1987) Quantitative methods in the review of epidemiologic literature. Epidemiol. Rev. 9: 1-29.
- 52

- 1 Hammand, E.C. (1966) Smoking in relation to the death rates of one million men and women. In: Haenxzel, W.,
2 ed. Epidemiological approaches to the study of cancer and other chronic diseases. National Cancer Institute
3 Monograph No. 19. Washington, DC.
- 4
- 5 Harris, C.C. (1989) Interindividual variation among humans in carcinogen metabolism, DNA adduct formation
6 and DNA repair. *Carcinogenesis* 10: 1563-1566.
- 7
- 8 Harris, C.C.; Hollstein, M. (1993) Clinical implications of the p53 tumor suppressor gene. *N. Engl. J. Med.* 329:
9 1318-1327.
- 10
- 11 Harris, H. (1990) The role of differentiation in the suppression of malignancy. *J. Cell Sci.* 97: 5-10.
- 12
- 13 Hartwell, L.H.; Kastan, M.B. (1994) Cell cycle control and cancer. *Science* 266: 1821-1828.
- 14
- 15 Haseman, J.K. (1983) Issues: a reexamination of false-positive rates for carcinogenesis studies. *Fund. Appl.*
16 *Toxicol.* 3: 334-339.
- 17
- 18 Haseman, J.K. (1984) Statistical issues in the design, analysis and interpretation of animal carcinogenicity
19 studies. *Environ. Health Perspect.* 58: 385-392.
- 20
- 21 Haseman, J.K. (1985) Issues in carcinogenicity testing: dose selection. *Fund. Appl. Toxicol.* 5: 66-78.
- 22
- 23 Haseman, J.K. (1990) Use of statistical decision rules for evaluating laboratory animal carcinogenicity studies.
24 *Fund. Appl. Toxicol.* 14: 637-648.
- 25
- 26 Haseman, J.K. (1995) Data analysis: Statistical analysis and use of historical control data. *Regul. Toxicol.*
27 *Pharmacol.* 21: 52-59.
- 28
- 29 Haseman, J.K.; Huff, J.; Boorman, G.A. (1984) Use of historical control data in carcinogenicity studies in
30 rodents. *Toxicol. Pathol.* 12: 126-135.
- 31
- 32 Hattis, D. (1990) Pharmacokinetic principles for dose-rate extrapolation of carcinogenic risk from genetically
33 active agents. *Risk Anal.* 10: 303-316.
- 34
- 35 Havu, N.; Mattsson, H.; Ekman, L.; Carlsson, E. (1990) Enterochromaffin-like cell carcinoids in the rat gastric
36 mucosa following long-term administration of ranitidine. *Digestion* 45: 189-195.
- 37
- 38 Hayward, N.K.; Walker, G.J.; Graham, W.; Cooksley, E. (1991) Hepatocellular carcinoma mutation. *Nature* 352:
39 764.
- 40
- 41 Hayward, J.J.; Shane, B.S.; Tindall, K.R.; Cunningham, M.L. (1995) Differential in vivo mutagenicity of the
42 carcinogen-noncarcinogen pair 2,4- and 2,6-diaminotoluene. *Carcinogenesis*. In press.
- 43
- 44 Herschman, H.R. (1991) Primary response genes induced by growth factors or promoters. *Ann. Rev. Biochem.*
45 60: 281-319.
- 46
- 47 Hill, R.N.; Erdreich, L.S.; Paynter, O.E.; Roberts, P.A.; Rosenthal, S.L.; Wilkinson, C.F. (1989) Thyroid
48 follicular cell carcinogenesis. *Fund. Appl. Toxicol.* 12: 629-697.
- 49
- 50 Hoel, D.G.; Portier, C.J. (1994) Nonlinearity of dose-response functions for carcinogenicity. *Environ. Health*
51 *Perspect.* 102 Suppl 1: 109-113.
- 52

- 1 Hoel, D.G.; Haseman, J.K.; Hogam, M.D.; Huff, J.; McConnell, E.E. (1988) The impact of toxicity on
2 carcinogenicity studies: Implications for risk assessment. *Carcinogenesis* 9: 2045-2052.
- 3
- 4 Holliday, R. (1987) DNA methylation and epigenetic defects in carcinogenesis. *Mutation Research* 181: 215-
5 217.
- 6
- 7 Hollstein, M.; Sidransky, D.; Vogelstein, B.; Harris, C.C. (1991) p53 mutations in human cancers. *Science* 253:
8 49-53.
- 9
- 10 Hsu, I.C.; Metcalf, R.A.; Sun, T.; Welsh, J.A.; Wang, N.J.; Harris, C.C. (1991) Mutational hotspot in human
11 hepatocellular carcinomas. *Nature* 350: 427-428.
- 12
- 13 Huff, J.E. (1993) Chemicals and cancer in humans: first evidence in experimental animals. *Environ. Health*
14 *Perspect.* 100: 201-210.
- 15
- 16 Huff, J.E. (1994) Chemicals causally associated with cancers in humans and laboratory animals. A perfect
17 concordance. In: *Carcinogenesis*. Waalkes, M.P.; Ward, J.M., eds., New York, NY: Raven Press; pp. 25-37.
- 18
- 19 Hulka, B.S.; Margolin, B.H. (1992) Methodological issues in epidemiologic studies using biological markers.
20 *Am. J. Epidemiol.* 135: 122-129.
- 21
- 22 Hunter, T. (1991) Cooperation between oncogenes. *Cell* 64: 249-270.
- 23
- 24 Ingram, A.J.; Grasso, P. (1991) Evidence for and possible mechanisms of non-genotoxic carcinogenesis in mouse
25 skin. *Mutat. Res.* 248: 333-340.
- 26
- 27 International Agency for Research on Cancer (IARC). (1990) Cyclosporin. IARC monographs on the evaluation of
28 carcinogenic risks to humans. Vol. 50. Lyon, France: IARC; pp. 77-114.
- 29
- 30 IARC. (1994) IARC monographs on the evaluation of carcinogenic risks to humans. Vol. 60, Some industrial
31 chemicals. Lyon, France: IARC; pp. 13-33.
- 32
- 33 International Life Science Institute (ILSI). (1995) The use of biological data in cancer risk assessment. In: Olin,
34 S.; Farland, W.; Park, C.; Rhomberg, L.; Scheuplein, R.; Starr, T.; Wilson, J., eds. *Low-dose extrapolation of*
35 *cancer risks: Issues and Perspectives*. Washington, DC: ILSI Press; pp. 45-60.
- 36
- 37 Ito, N.; Shirai, T.; and Hasegawa, R. (1992) Medium-term bioassays for carcinogens. in "Mechanisms of
38 Carcinogenesis in Risk Identifications" (eds., H. Vainio, P.N. Magee, D.B. McGregor and A.J. McMichael), Lyon,
39 International Agency for Research on Cancer, pp. 353-388.
- 40
- 41 Jack, D.; Poynter, D.; Spurling, N.W. (1983) Beta-adrenoreceptor stimulants and mesoovarian leiomyomas in the
42 rat. *Toxicology* 2: 315-320.
- 43
- 44 Jarabek, A.M. (1995a) The application of dosimetry models to identify key processes and parameters for default
45 dose-response assessment approaches. *Toxicol. Lett.* 79:171-184.
- 46
- 47 Jarabek, A.M. (1995b) Interspecies extrapolation based on mechanistic determinants of chemical disposition.
48 *Human Eco. Risk Assess.* 1(5):641-662.
- 49
- 50 Jones, P.A. (1986) DNA methylation and cancer. *Cancer Res.* 46: 461-466.
- 51
- 52 Kehrer, J.P. (1993) Free radicals as mediators of tissue injury and disease. *Crit. Rev. Toxicol.* 23: 21-48.

- 1 Kelsey, J.L.; Thompson, W.D.; Evans, A.S. (1986) Methods in observational epidemiology. New York, NY:
2 Oxford University Press.
- 3
- 4 Kinzler, K.W.; Nilbert, M.C.; Su, L.-K.; Vogelstein, B.; Bryan, T.M.; Levy, D.B.; Smith, K.J.; Preisinger, A.C.;
5 Hedge, P.; McKechnie, D.; Finnear, R.; Markham, A.; Groffen, J.; Boguski, M.S.; Altschul, S.J.; Horii, A.;
6 Ando, H.; Miyoshi, Y.; Miki, Y.; Nishisho, I.; Nakamura, Y. (1991) Identification of FAP locus genes from
7 chromosome 5q21. *Science* 253: 661-665.
- 8
- 9 Kodell, R.L.; Park, C.N. (1995) Linear extrapolation in cancer risk assessment. ILSI Risk Science Institute:
10 Washington, D.C. In press.
- 11
- 12 Kraus, A.L.; Munro, I.C.; Orr, J.C.; Binder, R.L.; LeBoeuf, R.A.; Williams, G.M. (1995) Benzoyl peroxide: An
13 integrated human safety assessment for carcinogenicity. *Regul. Toxicol. Pharmacol.* 21: 87-107.
- 14
- 15 Krewski, D.; Brown, C.; Murdoch, D. (1984) Determining "safe" levels of exposure: Safety factors of
16 mathematical models. *Fund. Appl. Toxicol.* 4: S383-S394.
- 17
- 18 Krewski, D.; Murdoch, D.J.; Withey, J.R. (1987) The application of pharmacokinetic data in carcinogenic risk
19 assessment. In: *Pharmacokinetics in risk assessment. Drinking water and health.* Vol. 8. Washington, DC:
20 National Academy Press; pp. 441-468.
- 21
- 22 Kripke, M.L. (1988) Immunoregulation of carcinogenesis: Past, present, and future. *J. Natl. Cancer Inst.* 80: 722-
23 727.
- 24
- 25 Kumar, R.; Sukumar, S.; Barbacid, M. (1990) Activation of *ras* oncogenes preceding the onset of neoplasia.
26 *Science* 248: 1101-1104.
- 27
- 28 Larson, J.L.; Wolf, D.C.; Butterworth, B.E. (1994) Induced cytotoxicity and cell proliferation in the
29 hepatocarcinogenicity of chloroform in female B6C3F1 mice: comparison of administration by gavage in corn oil
30 vs. ad libitum in drinking water. *Fundam. Appl. Toxicol.* 22: 90-102.
- 31
- 32 Levine, A.M. (1993) AIDs-related malignancies: The emerging epidemic. *J. Natl. Cancer Inst.* 85: 1382-1397.
- 33
- 34 Levine, P.H.; Hoover, R.N., eds. (1992) The emerging epidemic of non-Hodgkin's lymphoma: Current knowledge
35 regarding etiological factors. *Cancer Res.* 52: 5426s-5574s.
- 36
- 37 Levine, A. J.; Momand, J.; Finlay, C. A. (1991) The p53 tumour suppressor gene. *Nature* 351: 453-456.
- 38
- 39 Levine, A.J.; Perry, M.E.; Chang, A., et al. (1994) The 1993 Walter Hubert Lecture: The role of the p53 tumor-
40 suppressor gene in tumorigenesis. *Brit. J. Cancer* 69: 409-416.
- 41
- 42 Li, J.L.; Okada, S.; Hamazaki, S.; Ebina, Y.; Midorikawa, O. (1987) Subacute nephrotoxicity and induction of
43 renal cell carcinoma in mice treated with ferric nitrilotriacetate. *Cancer Res.* 47: 1867-1869.
- 44
- 45 Lijinsky, W. (1993) Species differences in carcinogenesis. *In Vivo* 7: 65-72.
- 46
- 47 Lilienfeld, A.M.; Lilienfeld, D. (1979) Foundations of epidemiology, 2nd ed. New York, NY: Oxford University
48 Press.
- 49
- 50 Loeb, L.A. (1991) Mutator phenotype may be required for multistage carcinogenesis. *Cancer Res.* 51: 3075-3079.
- 51

- 1 Lutz, W.K. (1990a) Endogenous genotoxic agents and processes as a basis of spontaneous carcinogenesis. *Mutat.*
2 *Res.* 238: 287-295.
- 3
- 4 Lutz, W.K. (1990b) Dose-response relationship and low dose extrapolation in chemical carcinogenesis.
5 *Carcinogenesis* 11: 1243-1247.
- 6
- 7 MacDonald, J.S.; Lankas, G.R.; Morrissey, R.E. (1994) Toxicokinetic and mechanistic considerations in the
8 interpretaion of the rodent bioassay. *Toxicol. Pathol.* 22: 124-140.
- 9
- 10 Maller, J.L. (1991) Mitotic control. *Curr. Opin. Cell Biol.* 3: 269-275.
- 11
- 12 Maronpot, R.R.; Shimkin, M.B.; Witschi, H.P.; Smith, L.H.; and Cline, J.M. (1986) Strain A mouse pulmonary
13 tumor test results for chemicals previously tested in National Cancer Institute carcinogenicity test. *J. Natl. Cancer*
14 *Inst.* 76: 1101-1112.
- 15
- 16 Marsman, D.S.; Popp, J.A. (1994) Biological potential of basophilic hepatocellular foci and hepatic adenoma
17 induced by the peroxisome proliferator, Wy-14,643. *Carcinogenesis* 15: 111-117.
- 18
- 19 Mausner, J.S.; Kramer, S. (1985) *Epidemiology*, 2nd ed. Philadelphia: W.B. Saunders Company.
- 20
- 21 McClain, R.M. (1994) Mechanistic considerations in the regulation and classification of chemical carcinogens.
22 In: Kotsonis, F.N.; Mackey, M.; Hjelle, J., eds. *Nutritional toxicology*. New York, NY: Raven Press; pp. 273-
23 304.
- 24
- 25 McConnell, E.E.; Solleveld, H.A.; Swenberg, J.A.; Boorman, G.A. (1986) Guidelines for combining neoplasms
26 for evaluation of rodent carcinogenesis studies. *J. Natl. Cancer Inst.* 76: 283-289.
- 27
- 28 McMichael, A.J. (1976) Standardized mortality ratios and the "healthy worker effect": scratching beneath the
29 surface. *J. Occup. Med.* 18: 165-168.
- 30
- 31 Melnick, R.L.; Huff, J.E.; Barrett, J.C.; Maronpot, R.R.; Lucier, G.; Portier, C.J. (1993a) Cell proliferation and
32 chemical carcinogenesis: A symposium overview. *Molecular Carcinogenesis* 7: 135-138.
- 33
- 34 Melnick, R.L.; Huff, J.E.; Barrett, J.C.; Maronpot, R.R.; Lucier, G.; Portier, C.J. (1993b) Cell proliferation and
35 chemical carcinogenesis. *Molecular Carcinogeneis* 7: 135-138.
- 36
- 37 Meyn, M.S. (1993) High spontaneous intrachromosomal recombination rates in ataxia-telangiectasia. *Science*
38 260: 1327-1330.
- 39
- 40 Modrich, P. (1994) Mismatch repair, genetic stability, and cancer. *Science* 266: 1959-1960.
- 41
- 42 Monro, A. (1992) What is an appropriate measure of exposure when testing drugs for carcinogenicity in rodents?
43 *Toxicol. Appl. Pharmacol.* 112: 171-181.
- 44
- 45 Moolgavkar, S.H.; Knudson, A.G. (1981) Mutation and cancer: A model for human carcinogenesis. *J. Natl.*
46 *Cancer Inst.* 66: 1037-1052.
- 47
- 48 Morrison, V.; Ashby, J. (1994) A preliminary evaluation of the performance of the mutaTM mouse (lacZ) and Big
49 BlueTM (lacI) transgenic mouse mutation assays. *Mutagenesis* 9: 367-375.
- 50
- 51 Naeve, G.S.; Sharma, A.; Lee, A.S. (1991) Temporal events regulating the early phases of the mammalian cell
52 cycle. *Curr. Opin. Cell Biol.* 3: 261-268.

- 1 National Research Council (NRC). (1983) Risk assessment in the federal government: Managing the process.
2 Committee on the Institutional Means for Assessment of Risks to Public Health, Commission on Life Sciences,
3 NRC. Washington, DC: National Academy Press.
4
- 5 NRC. (1993a) Pesticides in the diets of infants and children. Committee on Pesticides in the Diets of Infants and
6 Children, Commission on Life Sciences, NRC. Washington, DC: National Academy Press.
7
- 8 NRC. (1993b) Issues in risk assessment. Committee on Risk Assessment Methodology, Commission on Life
9 Sciences, NRC. Washington, DC: National Academy Press.
10
- 11 NRC. (1994) Science and judgment in risk assessment. Committee on Risk Assessment of Hazardous Air
12 Pollutants, Commission on Life Sciences, NRC. Washington, DC: National Academy Press.
13
- 14 National Toxicology Program (NTP). (1984) Report of the Ad Hoc Panel on Chemical Carcinogenesis Testing
15 and Evaluation of the National Toxicology Program, Board of Scientific Counselors. Washington, DC: U.S.
16 Government Printing Office. 1984-421-132: 4726.
17
- 18 Nebreda, A.R.; Martin-Zanca, D.; Kaplan, D.R.; Parada, L.F.; Santos, E. (1991) Induction by NGF of meiotic
19 maturation of *xenopus* oocytes expressing the *trk* proto-oncogene product. *Science* 252: 558-561.
20
- 21 Nowell, P. (1976) The clonal evolution of tumor cell populations. *Science* 194: 23-28.
22
- 23 Nunez, G.; Hockenberry, D.; McDonnell, J.; Sorenson, C.M.; Korsmeyer, S.J. (1991) Bcl-2 maintains B cell
24 memory. *Nature* 353: 71-72.
25
- 26 Office of Science and Technology Policy (OSTP). (1985) Chemical carcinogens: Review of the science and its
27 associated principles. *Federal Register* 50: 10372-10442.
28
- 29 Organization for Economic Cooperation and Development (OECD). (1981) Guidelines for testing of chemicals.
30 Carcinogenicity studies. No. 451. Paris, France.
31
- 32 Peltomäki, P.; Aaltonen, L.A.; Sisonen, P.; Pylkkänen, L.; Mecklin, J.-P.; Järvinen, H.; Green, J.S.; Jass, J.R.;
33 Weber, J.L.; Leach, F.S.; Petersen, G.M.; Hamilton, S.R.; de la Chapelle, A.; Vogelstein, B. (1993) Genetic
34 mapping of a locus predisposing human colorectal cancer. *Science* 260: 810-812.
35
- 36 Peto, J. (1992) Meta-analysis of epidemiological studies of carcinogenesis. In: *Mechanisms of carcinogenesis in*
37 *risk assessment*. IARC Sci. Pubs. No. 116, Lyon, France: IARC; pp. 571-577.
38
- 39 Peto, J.; Darby, S. (1994) Radon risk reassessed. *Nature* 368: 97-98.
40
- 41 Pitot, H.; Dragan, Y.P. (1991) Facts and theories concerning the mechanisms of carcinogenesis. *FASEB J.* 5:
42 2280-2286.
43
- 44 Portier, C. (1987) Statistical properties of a two-stage model of carcinogenesis. *Environ. Health Perspect.* 76:
45 125-131.
46
- 47 Prahalada, S.; Majka, J.A.; Soper, K.A.; Nett, T.M.; Bagdon, W.J.; Peter, C.P.; Burek, J.D.; MacDonald, J.S.;
48 van Zwieten, M.J. (1994) Leydig cell hyperplasia and adenomas in mice treated with finasteride, a 5 α -reductase
49 inhibitor: A possible mechanism. *Fund. Appl. Toxicol.* 22: 211-219.
50
- 51 Raff, M.C. (1992) Social controls on cell survival and cell death. *Nature* 356: 397-400.
52

- 1 Rall, D.P. (1991) Carcinogens and human health: Part 2. *Science* 251: 10-11.
- 2
- 3 Rothman, K.T. (1986) *Modern epidemiology*. Boston: Little, Brown and Company.
- 4
- 5 Salomon, D.S.; Kim, N.; Saeki, T.; Ciardiello, F. (1990) Transforming growth factor " --an oncodevelopmental
- 6 growth factor. *Cancer Cells* 2: 389-397.
- 7
- 8 Schneider, C.; Gustincich, S.; DelSal, G. (1991) The complexity of cell proliferation control in mammalian cells.
- 9 *Curr. Opin. Cell Biol.* 3: 276-281.
- 10
- 11 Schuller, H.M. (1991) Receptor-mediated mitogenic signals and lung cancer. *Cancer Cells* 3: 496-503.
- 12
- 13 Schulte-Hermann, R.; Bursch, W.; Kraupp-Grasl, B.; Oberhammer, F.; Wagner, A.; Jirtle, R. (1993) Cell
- 14 proliferation and apoptosis in normal liver and preneoplastic foci. *Environ. Health Perspect.* 101 (Supp. 5): 87-90.
- 15
- 16 Shelby, M.D.; Zeiger, E. (1990) Activity of human carcinogens in the *Salmonella* and rodent bone-marrow
- 17 cytogenetics tests. *Mutat. Res.* 234: 257-261.
- 18
- 19 Shelby, M.D. (1994) Human germ cell mutations. *Environ. Molec. Mutagen.* 23 (Supp. 24): 30-34.
- 20
- 21 Sidransky, D.; Von Eschenbach, A.; Tsai, Y.C.; Jones, P.; Summerhayes, I.; Marshall, F.; Paul, M.; Green, P.;
- 22 Hamilton, P.F.; Vogelstein, B. (1991) Identification of p53 gene mutations in bladder cancers and urine samples.
- 23 *Science* 252: 706-710.
- 24
- 25 Sidransky, D.; Mikkelsen, T.; Schwechheimer, K.; Rosenblum, M.L.; Cavanee, W.; Vogelstein, B. (1992) Clonal
- 26 expansion of p53 mutant cells is associated with brain tumor progression. *Nature* 355: 846-847.
- 27
- 28 Sisk, S.C.; Pluta, L.J.; Bond, J.A.; Recio, L. (1994) Molecular analysis of lacI mutants from bone marrow of
- 29 B6C3F1 transgenic mice following inhalation exposure to 1,3-butadiene. *Carcinogenesis* 15(3): 471-477.
- 30
- 31 Snedecor, G.W.; Cochran, W.G. (1978) *Statistical methods*, Sixth ed. Ames, Iowa: Iowa State University Press;
- 32 593 pp.
- 33
- 34 Srivastava, S.; Zou, Z.; Pirolo, K.; Blattner, W.; Chang, E. (1990) Germ-line transmission of a mutated p53 gene
- 35 in a cancer-prone family with Li-Fraumeni syndrome. *Nature* 348(6303): 747-749.
- 36
- 37 Stewart, B.W. (1994) Mechanisms of apoptosis: Integration of genetic, biochemical, and cellular indicators. *J.*
- 38 *Natl. Cancer Inst.* 86: 1286-1296.
- 39
- 40 Stiteler, W.H.; Knauf, L.A.; Hertzberg, R.C.; Schoeny, R.S. (1993) A statistical test of compatibility of data sets
- 41 to a common dose-response model. *Reg. Tox. Pharmacol.* 18: 392-402.
- 42
- 43 Stitzel, K.A.; McConnell, R.F.; Dierckman, T.A. (1989) Effects of nitrofurantoin on the primary and secondary
- 44 reproductive organs of female B6C3F1 mice. *Toxicol. Pathol.* 17: 774-781.
- 45
- 46 Strausfeld, U.; Labbe, J.C.; Fesquet, D.; Cavadore, J.C.; Dicard, A.; Sadhu, K.; Russell, P.; Dor'ee, M. (1991)
- 47 Identification of a G1-type cyclin puc1+ in the fission yeast *Schizosaccharomyces pombe*. *Nature* 351: 242-245.
- 48
- 49 Sukumar, S. (1989) *ras* oncogenes in chemical carcinogenesis. *Curr. Top. Microbiol. Immunol.* 148: 93-114.
- 50
- 51 Sukumar, S. (1990) An experimental analysis of cancer: Role of *ras* oncogenes in multistep carcinogenesis.
- 52 *Cancer Cells* 2: 199-204.

- 1 Swenberg, J.A.; Richardson, F.C.; Boucheron, J.A.; Deal, F.H.; Belinsky, S.A.; Charbonneau, M.; Short, B.G.
2 (1987) High to low dose extrapolation: Critical determinants involved in the dose-response of carcinogenic
3 substances. *Environ. Health Perspect.* 76: 57-63.
4
- 5 Swierenga, S.H.H.; Yamasaki, H. (1992) Performance of tests for cell transformation and gap junction
6 intercellular communication for detecting nongenotoxic carcinogenic activity. In: *Mechanisms of carcinogenesis*
7 *in risk identification*. IARC Sci. Pubs. No. 116, Lyon, France: International Agency for Research on Cancer; pp.
8 165-193.
9
- 10 Tanaka, K.; Oshimura, M.; Kikiuchi, R.; Seki, M.; Hayashi, T; Miyaki, M. (1991) Suppression of tumorigenicity
11 in human colon carcinoma cells by introduction of normal chromosome 5 or 18. *Nature* 349: 340-342.
12
- 13 Tarone, R.E. (1982) The use of historical control information in testing for a trend in proportions. *Biometrics* 38:
14 215-220.
15
- 16 Taylor, J.H.; Watson, M.A.; Devereux, T.R.; Michels, R.Y.; Saccomanno, G.; Anderson, M. (1994) *Lancet* 343:
17 86-87.
18
- 19 Tennant, R.W. (1993) Stratification of rodent carcinogenicity bioassay results to reflect relative human hazard.
20 *Mutat. Res.* 286: 111-118.
21
- 22 Tennant, R.W.; Ashby, J. (1991) Classification according to chemical structure, mutagenicity to *Salmonella* and
23 level of carcinogenicity of a further 39 chemicals tested for carcinogenicity by the U.S. National Toxicology
24 Program. *Mutat. Res.* 257: 209-277.
25
- 26 Tennant, R.W.; Elwell, M.R.; Spalding, J.W.; Griesemer, R.A. (1991) Evidence that toxic injury is not always
27 associated with induction of chemical carcinogenesis. *Molec. Carcinogen.* 4: 420-440.
28
- 29 Tennant, R.W.; French, J.E.; Spalding, J.W. (1995) Identifying chemical carcinogens and assessing potential risk
30 in short-term bioassays using transgenic mouse models. *Environ. Health Perspect.* 103:942-950.
31
- 32 Thompson, T.C.; Southgate, J.; Kitchener, G.; Land, H. (1989) Multistage carcinogenesis induced by *ras* and *myc*
33 oncogenes in a reconstituted organ. *Cell* 56: 917-3183.
34
- 35 Tinwell, H.; Ashby, J. (1991) Activity of the human carcinogen MeCCNU in the mouse bone marrow
36 micronucleus test. *Environ. Molec. Mutagen.* 17: 152-154.
37
- 38 Tischler, A.S.; McClain, R.M.; Childers, H.; Downing, J. (1991) Neurogenic signals regulate chromaffin cell
39 proliferation and mediate the mitogenic effect of reserpine in the adult rat adrenal medulla. *Lab. Invest.* 65: 374-
40 376.
41
- 42 Tobey, R.A. (1975) Different drugs arrest cells at a number of distinct stages in G2. *Nature* 254: 245-247.
43
- 44 Todd, G.C. (1986) Induction of reversibility of thyroid proliferative changes in rats given an antithyroid
45 compound. *Vet. Pathol.* 23: 110-117.
46
- 47 Tomatis, L.; Aitio, A.; Wilbourn, J.; Shuker, L. (1989) Human carcinogens so far identified. *Jpn. J. Cancer Res.*
48 80: 795-807.
49
- 50 Travis, C.C.; McClain, T.W.; Birkner, P.D. (1991) Diethylnitrosamine-induced hepatocarcinogenesis in rats: A
51 theoretical study. *Toxicol. Appl. Pharmacol.* 109: 289-309.
52

1 U.S. Environmental Protection Agency. (1983a) Good laboratory practices standards--toxicology testing. Federal
2 Register 48: 53922.
3
4 U.S. Environmental Protection Agency. (1983b) Hazard evaluations: Humans and domestic animals. Subdivision
5 F. Available from: NTIS, Springfield, VA; PB 83-153916.
6
7 U.S. Environmental Protection Agency. (1983c) Health effects test guidelines. Available from: NTIS,
8 Springfield, VA; PB 83-232984.
9
10 U.S. Environmental Protection Agency. (1984) Estimation of the public health risk from exposure to gasoline
11 vapor via the gasoline marketing system. Office of Health and Environmental Assessment, Washington, DC.
12
13 U.S. Environmental Protection Agency. (1986a) Health assessment document for beryllium. Office of Health and
14 Environmental Assessment, Washington, DC.
15
16 U.S. Environmental Protection Agency. (1986b) Guidelines for carcinogen risk assessment. Federal Register
17 51(185):33992-34003.
18
19 U.S. Environmental Protection Agency. (1989a) Interim procedures for estimating risks associated with
20 exposures to mixtures of chlorinated dibenzo-*p*-dioxins and -dibenzofurans (CDDs and CDFs) and 1989 update.
21 Risk Assessment Forum, Washington, DC. EPA/625/3-89/016.
22
23 U.S. Environmental Protection Agency. (1989b) Workshop on EPA guidelines for carcinogen risk assessment.
24 Risk Assessment Forum, Washington, DC. EPA/625/3-89/015.
25
26 U.S. Environmental Protection Agency. (1989c) Workshop on EPA guidelines for carcinogen risk assessment:
27 use of human evidence. Risk Assessment Forum, Washington, DC. EPA/625/3-90/017.
28
29 U.S. Environmental Protection Agency. (1991a) Pesticide assessment guidelines: Subdivision F, hazard
30 evaluation, human and domestic animals. Series 84, Mutagenicity, Addendum 9. Office of Pesticide Programs,
31 Washington, DC. PB91-158394, 540/09-91-122.
32
33 U.S. Environmental Protection Agency. (1991b) Alpha-2u-globulin: Association with chemically induced renal
34 toxicity and neoplasia in the male rat. Risk Assessment Forum, Washington, DC. EPA/625/3-91/019F.
35
36 U.S. Environmental Protection Agency. (1991c) Workshop report on toxicity equivalency factors for
37 polychlorinated biphenyl congeners. Risk Assessment Forum, Washington, DC. EPA/625/3-91/020.
38
39 U.S. Environmental Protection Agency. (1991f) Guidelines for developmental toxicity risk assessment. Federal
40 Register 56(234): 63798-63826.
41
42 U.S. Environmental Protection Agency. (1992a) Guidelines for exposure assessment. Federal Register 57(104):
43 22888-22938.
44
45 U.S. Environmental Protection Agency. (1992b) Draft report: A cross-species scaling factor for carcinogen risk
46 assessment based on equivalence of mg/kg^{3/4}/day. Federal Register 57(109): 24152-24173.
47
48 U.S. Environmental Protection Agency. (1992c) Health assessment for 2,3,7,8-tetrachlorodibenzo-*p*-dioxin
49 (TCDD) and related compounds (Chapters 1 through 8). Workshop Review Drafts. EPA/600/AP-92/001a through
50 001h.
51

U.S. Environmental Protection Agency. (1994) Methods for derivation of inhalation reference concentrations and application of inhalation dosimetry. Office of Health and Environmental Assessment, Environmental Criteria and Assessment Office, Research Triangle Park, NC. EPA/600/8-90/066F.

U.S. Environmental Protection Agency. (1994a) Estimating exposure to dioxin-like compounds. Office of Health and Environmental Assessment, Office of Research and Development, Washington, DC. External Review Draft, 3 vol. EPA/600/6-88/005Ca, Cb, Cc.

U.S. Environmental Protection Agency. (1994b) Report on the workshop on cancer risk assessment guidelines issues. Office of Research and Development, Risk Assessment Forum, Washington, DC. EPA/630/R-94/005a.

U.S. Environmental Protection Agency. (1995) Policy for risk characterization. Memorandum of Carol M. Browner, Administrator, March 21, 1995, Washington, D.C.

U.S. Food and Drug Administration (1987) Sponsored compounds in food-producing animals; criteria and procedures for evaluating the safety of carcinogenic residues. final rule. 21 CFR Parts 70, 500, 514 and 571.

Vahakangas, K.H.; Samet, J.M.; Metcalf, R.A.; Welsh, J.A.; Bennett, W.P.; Lane, D.P.; Harris, C.C. (1992) Mutation of p53 and *ras* genes in radon-associated lung cancer from uranium miners. *Lancet* 339: 576-578.

Vainio, H.; Magee, P.; McGregor, D.; McMichael, A.J. (1992) Mechanisms of carcinogenesis in risk identification. IARC Sci. Pubs. No. 116. Lyon, France: IARC.

Van Sittert, N.J.; De Jong, G.; Clare, M.G.; Davies, R.; Dean, B.J.; Wren, L.R.; Wright, A.S. (1985) Cytogenetic, immunological, and hematological effects in workers in an ethylene oxide manufacturing plant. *Br. J. Indust. Med.* 42:19-26.

Vater, S.T.; McGinnis, P.M.; Schoeny, R.S.; Velazquez, S. (1993) Biological considerations for combining carcinogenicity data for quantitative risk assessment. *Reg. Toxicol. Pharmacol.* 18: 403-418.

Vogelstein, B.; Fearon, E. R.; Hamilton, S. R.; Kern, S. E.; Presinger, A. C.; Leppert, M.; Nakamura, Y.; White, R.; Smits, A. M. M.; Bos, J. L. (1988) Genetic alterations during colorectal-tumor development. *New England Journal of Medicine* 319: 525-532.

Weinberg, R.A. (1989) Oncogenes, antioncogenes, and the molecular bases of multistep carcinogenesis. *Cancer Res.* 49: 3713-3721.

Wellstein, A.; Lupu, R.; Zugmaier, G.; Flamm, S.L.; Cheville, A.L.; Bovi, P.D.; Basicico, C.; Lippman, M.E.; Kern, F.G. (1990) Autocrine growth stimulation by secreted Kaposi fibroblast growth factor but not by endogenous basic fibroblast growth factor. *Cell Growth Differ.* 1: 63-71.

Woo, Y.T.; Arcos, J.C. (1989) Role of structure-activity relationship analysis in evaluation of pesticides for potential carcinogenicity. In: Ragsdale, N.N.; Menzer, R.E., eds. *Carcinogenicity and pesticides: Principles, issues, and relationship*. ACS Symposium Series No. 414. San Diego: Academic Press; pp. 175-200.

Wyzga, R.E. (1988) The role of epidemiology in risk assessments of carcinogens. *Adv. Mod. Environ. Toxicol.* 15: 189-208.

Yamada, T.; Nakamura, J.; Murakami, M.; Okuno, Y.; Hosokawa, S.; Matsuo, M.; Yamada, H. (1994) The correlation of serum luteinizing hormone levels with the induction of Leydig cell tumors in rats by oxolinic acid. *Toxicol. Appl. Pharmacol.* 129: 146-154.

1 Yamasaki, H. (1990) Gap junctional intercellular communication and carcinogenesis. *Carcinogenesis* 11: 1051-
2 1058.

3
4 Yamasaki, H. (1995) Non-genotoxic mechanisms of carcinogenesis: Studies of cell transformation and gap
5 junctional intercellular communication. *Toxicol. Lett.* 77: 55-61.

6
7 Zhang, K.; Papageorge, A.G.; Lowry, D.R. (1992) Mechanistic aspects of signalling through ras in NIH 3T3
8 cells. *Science* 257: 671-674.

9
10
11 Billing Code: 6560-50-P
12
13