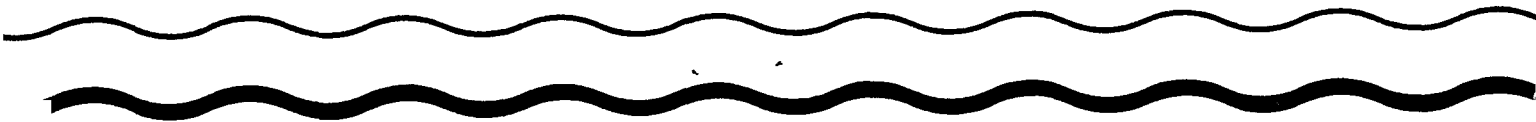
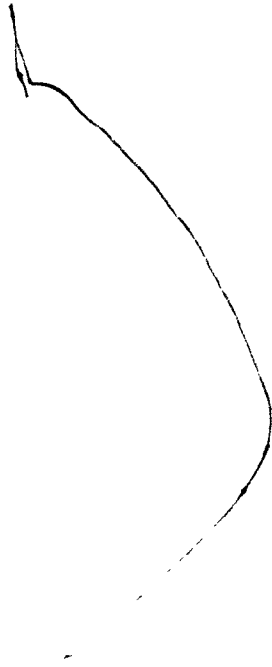




Ambient Water Quality Criteria for Polynuclear Aromatic Hydrocarbons



AMBIENT WATER QUALITY CRITERIA FOR
POLYNUCLEAR AROMATIC HYDROCARBONS

Prepared By
U.S. ENVIRONMENTAL PROTECTION AGENCY

Office of Water Regulations and Standards
Criteria and Standards Division
Washington, D.C.

Office of Research and Development
Environmental Criteria and Assessment Office
Cincinnati, Ohio

Carcinogen Assessment Group
Washington, D.C.

Environmental Research Laboratories
Corvallis, Oregon
Duluth, Minnesota
Gulf Breeze, Florida
Narragansett, Rhode Island

U.S. Environmental Protection Agency
Region 5, Library (PL-12J)
77 West Jackson Boulevard, 12th Floor
Chicago, IL 60604-3590

DISCLAIMER

This report has been reviewed by the Environmental Criteria and Assessment Office, U.S. Environmental Protection Agency, and approved for publication. Mention of trade names or commercial products does not constitute endorsement or recommendation for use.

AVAILABILITY NOTICE

This document is available to the public through the National Technical Information Service, (NTIS), Springfield, Virginia 22161.

FOREWORD

Section 304 (a)(1) of the Clean Water Act of 1977 (P.L. 95-217), requires the Administrator of the Environmental Protection Agency to publish criteria for water quality accurately reflecting the latest scientific knowledge on the kind and extent of all identifiable effects on health and welfare which may be expected from the presence of pollutants in any body of water, including ground water. Proposed water quality criteria for the 65 toxic pollutants listed under section 307 (a)(1) of the Clean Water Act were developed and a notice of their availability was published for public comment on March 15, 1979 (44 FR 15926), July 25, 1979 (44 FR 43660), and October 1, 1979 (44 FR 56628). This document is a revision of those proposed criteria based upon a consideration of comments received from other Federal Agencies, State agencies, special interest groups, and individual scientists. The criteria contained in this document replace any previously published EPA criteria for the 65 pollutants. This criterion document is also published in satisfaction of paragraph 11 of the Settlement Agreement in Natural Resources Defense Council, et. al. vs. Train, 8 ERC 2120 (D.D.C. 1976), modified, 12 ERC 1833 (D.D.C. 1979).

The term "water quality criteria" is used in two sections of the Clean Water Act, section 304 (a)(1) and section 303 (c)(2). The term has a different program impact in each section. In section 304, the term represents a non-regulatory, scientific assessment of ecological effects. The criteria presented in this publication are such scientific assessments. Such water quality criteria associated with specific stream uses when adopted as State water quality standards under section 303 become enforceable maximum acceptable levels of a pollutant in ambient waters. The water quality criteria adopted in the State water quality standards could have the same numerical limits as the criteria developed under section 304. However, in many situations States may want to adjust water quality criteria developed under section 304 to reflect local environmental conditions and human exposure patterns before incorporation into water quality standards. It is not until their adoption as part of the State water quality standards that the criteria become regulatory.

Guidelines to assist the States in the modification of criteria presented in this document, in the development of water quality standards, and in other water-related programs of this Agency, are being developed by EPA.

STEVEN SCHATZOW
Deputy Assistant Administrator
Office of Water Regulations and Standards

ACKNOWLEDGEMENTS

Aquatic Life Toxicology:

William A. Brungs, ERL-Narragansett
U.S. Environmental Protection Agency

David J. Hansen, ERL-Gulf Breeze
U.S. Environmental Protection Agency

Mammalian Toxicology and Human Health Effects:

Joseph Santodonato (author)
Syracuse Research Corporation

Julian Andelman
University of Pittsburgh

Debdas Mukerjee (doc. mgr.) ECAO-Cin
U.S. Environmental Protection Agency

Edmund LaVoie
American Health Foundation

Bonnie Smith (doc. mgr.) ECAO-Cin
U.S. Environmental Protection Agency

S.D. Lee, ECAO-Cin
U.S. Environmental Protection Agency

Steven D. Lutkenhoff, ECAO-Cin
U.S. Environmental Protection Agency

Leo Newland
Texas Christian University

James Selkirk
Oakridge National Laboratory

Jerry F. Stara, ECAO-Cin
U.S. Environmental Protection Agency

Yin Tak Woo

Roy E. Albert, CAG*
U.S. Environmental Protection Agency

Technical Support Services Staff: D.J. Reisman, M.A. Garlough, B.L. Zwyer,
P.A. Daunt, K.S. Edwards, T.A. Scandura, A.T. Pressiey, C.A. Cooper,
M.M. Denessen.

Clerical Staff: C.A. Haynes, S.J. Faehr, L.A. Wade, D. Jones, B.J. Bordicks,
B.J. Quesnell, P. Gray, R. Rubinstein.

*CAG Participating Members: Elizabeth L. Anderson, Larry Anderson, Ralph Annicar,
Steven Bayard, David L. Bayliss, Chao W. Chen, John R. Fowle III, Bernard Haberman,
Charalingayya Hiremath, Chang S. Lao, Robert McGaughy, Jeffrey Rosenblatt,
Dharm V. Singh, and Todd W. Thorslund.

TABLE OF CONTENTS

	<u>Page</u>
Criteria Summary	
Chemical Abbreviations Used Within This Document	A-1
Introduction	A-2
Aquatic Life Toxicology	B-1
Introduction	B-1
Effects	B-1
Criteria	B-2
References	B-5
Mammalian Toxicology and Human Health Effects	C-1
Exposure	C-1
Ingestion from Water	C-1
Ingestion from Food	C-10
Inhalation	C-28
Dermal	C-37
Pharmacokinetics	C-37
Absorption	C-37
Distribution	C-38
Metabolism	C-39
Excretion	C-49
Effects	C-50
Acute, Subacute, and Chronic Toxicity	C-50
Synergism and/or Antagonism	C-57
Teratogenicity	C-63
Mutagenicity	C-64
Carcinogenicity	C-72
Criterion Formulation	C-108
Existing Guidelines and Standards	C-108
Current Levels of Exposure	C-109
Special Groups at Risk	C-111
Basis and Derivation of Criterion	C-117
References	C-124
Appendix	C-128

CRITERIA DOCUMENT
POLYNUCLEAR AROMATIC HYDROCARBONS

CRITERIA

Aquatic Life

The limited freshwater data base available for polynuclear aromatic hydrocarbons, mostly from short-term bioconcentration studies with two compounds, does not permit a statement concerning acute or chronic toxicity.

The available data for polynuclear aromatic hydrocarbons indicate that acute toxicity to saltwater aquatic life occurs at concentrations as low as 300 $\mu\text{g/l}$ and would occur at lower concentrations among species that are more sensitive than those tested. No data are available concerning the chronic toxicity of polynuclear aromatic hydrocarbons to sensitive saltwater aquatic life.

Human Health

For the maximum protection of human health from the potential carcinogenic effects due to exposure of polynuclear aromatic hydrocarbons through ingestion of contaminated water and contaminated aquatic organisms, the ambient water concentration should be zero based on the non-threshold assumption for this chemical. However, zero level may not be attainable at the present time. Therefore, the levels which may result in incremental increase of cancer risk over the lifetime are estimated at 10^{-5} , 10^{-6} , and 10^{-7} . The corresponding recommended criteria are 28.0 ng/l, 2.8 ng/l, and 0.28 ng/l, respectively. If the above estimates are made for consumption of aquatic organisms only, excluding consumption of water, the levels are 311.0 ng/l, 31.1 ng/l, and 3.11 ng/l, respectively.

CHEMICAL ABBREVIATIONS USED WITHIN THIS DOCUMENT

<u>Abbreviation</u>	<u>Chemical</u>
A:	anthracene
ANT:	anthranthrene
BaA:	benz[a]anthracene
BaP:	benz[a]pyrene
BbFL:	benzo[b]fluoranthene
BeP:	benzo[e]pyrene
BjFL:	benzo[j]fluoranthene
BkFL:	benzo[k]fluoranthene
BPR:	benzo[g,h,i]perylene
CH:	chrysene
CR:	coronene
DBA:	dibenz[a,h]anthracene
DMBA:	7,12-dimethylbenz[a]anthracene
F:	fluorene
FL:	fluoranthene
IP:	indeno[1,2,3-cd]pyrene
MCA:	3-methylcholanthrene
NA:	naphthalene
P:	pyrene
PA:	phenanthrene
PAH:	polynuclear aromatic hydrocarbons
PR:	perylene

INTRODUCTION

Polynuclear aromatic hydrocarbons (PAH) are a diverse class of compounds consisting of substituted and unsubstituted polycyclic and heterocyclic aromatic rings. PAH are formed as a result of incomplete combustion of organic compounds with insufficient oxygen. This leads to the formation of C-H free radicals which can polymerize to form various PAH. Among these PAH are compounds such as benzo[a]pyrene, and benz[a]anthracene.

PAH are present in the environment from both natural and anthropogenic sources. As a group, they are widely distributed in the environment, having been detected in animal and plant tissue, sediments, soils, air, and surface water (Radding et al. 1976); Shackelford and Keith (1976) report that PAH have been detected in surface waters, finished drinking water, industrial effluents, ambient river water, well water, and ground water.

PAH will adsorb strongly onto suspended particulates and biota and that their transport will be determined largely by the hydrogeologic condition of the aquatic system. PAH dissolved in the water column will probably undergo direct photolysis at a rapid rate. The ultimate fate of those which accumulate in the sediment is believed to be biodegradation and biotransformation by benthic organisms (U.S. EPA, 1979).

The general physical properties of acenaphthylene and fluorene are as follows:

	<u>Acenaphthylene</u>	<u>Fluorene</u>
Molecular weight	152.21 ^a	116.15 ^a
Melting point	92°C ^a	116-117°C ^a
Vapor pressure (20°C)	10 ⁻³ to 10 ⁻² torr ^b	10 ⁻³ to 10 ⁻² torr ^b
Solubility in water (25°C)	3.93 mg/l ^c	1.98 mg/l ^c 1.69 mg/l ^d
Log octanol/water partition coefficients	4.07 ^d	4.18 ^d

a) Weast, 1977.

b) Estimated, based on data for structurally similar compounds.

c) Mackay and Shiu, 1977.

d) Calculated as per Leo, et al. 1971.

e) May and Wasik, 1978.

The general physical properties of anthracene and phenanthrene are as follows:

	<u>Anthracene</u>	<u>Phenanthrene</u>
Molecular weight	178.23 ^f	178.23 ^f
Melting point	216°C ^f	101°C
Vapor Pressure (20°C)	1.95x10 ⁻⁴ torr ^f	6.8x10 ⁻⁴ torr ^f
Solubility in water (25°C)	0.045 mg/l ^h 0.073 mg/l ^g	1.00 mg/l ^h 1.29 mg/l ^g
Log octanol/water partition coefficients	4.45 ^f	4.46 ^f

f) Radding, et al. 1976.

g) Mackay and Shiu, 1977.

h) May and Wasik, 1978.

The physical properties of polycyclic aromatic hydrocarbons are as follows:

	<u>Benzo[a] Anthracene</u>	<u>Benzo[b] Fluoranthene</u>	<u>Benzo[k] Fluornathene</u>	<u>Chrysene</u>	<u>Pyrene</u>
Molecular weight	228.28 ⁱ	252.32 ^l	252.32 ^o	228.28 ^k	202 ^o
Melting point	155-157°C ⁱ	167-168°C ^l	217°C ^o	256°C ^k	150°C ^p
Vapor Pressure (20°C)	5x10 ⁻⁹ torr ⁱ	10 ⁻¹¹ -10 ⁻⁶ torr ^m	9.59x10 ⁻¹¹ torr ^k	10 ⁻¹¹ -10 ⁻⁶ torr ^m	6.85x10 ⁻⁷ torr ^k
Solubility in water (25°C)	0.014 mg/l ^j .009 mg/l ^q	NA	NA	0.002 mg/l ^j 0.002 mg/l ^q	0.14 mg/l ^j 0.132 mg/l ^q
Log Octanol/ Partition Coefficient	5.61 ^k	6.57 ⁿ	6.84 ⁿ	5.61 ^k	5.32 ⁿ

i) Smith et al. 1978.

j) Mackay and Shiu, 1977.

k) Radding et al. 1976.

l) IARC, 1973.

m) Estimated based on data for sturcturally similar compounds.

n) Calculated as per Leo et al. 1971.

o) Weast, 1977.

p) Cleland and Kingsbury, 1977.

q) May and Wasik, 1978.

NA = No data found.

The general physical properties of the polycyclic aromatic hydrocarbons having 5 or more aromatic rings which are discussed in this chapter are shown below.

	<u>Benzo[g,h,i] perylene</u>	<u>Benzo[a] pyrene</u>	<u>Dibenzo[a,h] anthracene</u>	<u>Indeno[1,2,3-cd] pyrene</u>
Molecular weight	276 ^r	252 ^s	278.36 ^r	276.34 ^r
Melting point	222°C ^t	179°C ^s	270°C ^r	162.5-164°C ^r
Vapor pressure (torr)	~10-10 ^u , ^y	5x10 ⁻⁹ ^{s,v}	~10-10 ^u , ^y	~10-10 ^u , ^y
Solubility in water (25°C)	0.00026 mg/l ^w	0.0038 mg/l ^w	0.0005 mg/l ^x	NA
Log octanol/water partition coefficient	7.23 ^z	6.04 ^{aa}	5.97 ^z	7.66 ^z

r) Weast, 1977.

s) Smith et al. 1978.

t) Cleland and Kingsbury, 1977.

u) 20°C

v) 25°C

w) Mackay and Shiu, 1977.

x) Davis, et al. 1942.

y) Estimated, based on data for structurally similar compounds.

z) Calculated according to Leo, et al. 1971.

aa) Radding, et al. 1976.

NA = No data found.

REFERENCES

- Cleland, J.G. and G.L. Kingsbury. 1977. Multimedia environmental goals for environmental assessment, Vol. II. MEG charts and background information. U.S. Environ. Prot. Agency, (Office of Research and Develop.), Washington, D.C. EPA-600/7-77-136b. p. 451.
- Davis, W.W., et al. 1942. Solubility of carcinogenic and related hydrocarbons in water. Jour. Am. Chem. Soc. 64: 108.
- International Agency for Research on Cancer. 1973. IARC monographs on the evaluation of carcinogenic risk of the chemical to man. Certain polycyclic aromatic hydrocarbons and heterocyclic compounds Vol. 3. Benzo[b]fluoranthene. p. 69. IARC, Lyon. p. 271.
- Leo, A., et al. 1971. Partition coefficients and their uses. Chem. Rev. 71: 525.
- Mackay, D. and W.Y. Shiu. 1977. Aqueous solubility of polynuclear aromatic hydrocarbons. Chem. Eng. Data. 22: 399.
- Mackay, D. and A.W. Wolkoff. 1973. Rate of evaporation of low-solubility contaminants from water bodies to atmosphere. Environ. Sci. Technol. 8: 611.
- May, W.E. and S.P. Wasik. 1978. Determination of the solubility behavior of some polyaromatic hydrocarbons in water. Anal. Chem. 50(7): 997.

May, W.E. and S.P. Wasik. 1978. Determination of the aqueous solubility of polynuclear aromatic hydrocarbons by a coupled chromatographic technique. Anal. Chem. 50(11): 997

Radding, S.B., et al. 1976. The environmental fate of selected polynuclear aromatic hydrocarbons. U.S. Environ. Prot. Agency, (Office of Toxic Sub.), Washington, D.C. EPA-560/5-75-009. p. 122.

Shackelford, W. and L.H. Keith. 1976. Frequency of organic compounds identified in water. U.S. Environ. Prot. Agency, (Office of Research and Development), Athens, Georgia. EPA-600/4-76-062. p. 618.

Smith, J.H., et al. 1978. Environmental pathways of selected chemicals in freshwater systems; Part II: Laboratory Studies. U.S. Environ. Prot. Agency, Athens, Georgia. EPA-600/7-78-074. p. 432.

U.S. EPA. 1979. Water-related environmental fate of 129 priority pollutants. EPA-440/4-79-0296.

Weast, R.C. (ed.) 1977. Handbook of Chemistry and Physics, 58th edition. CRC Press Inc., Cleveland, Ohio. p. 4, p. 2398.

INTRODUCTION

No standard freshwater toxicity tests have been reported for any polynuclear aromatic hydrocarbon not already discussed in criterion documents on specific compounds (e.g., fluoranthene and acenaphthene). There are some data for bioconcentration during tests with model ecosystems or for short periods of time.

As was true for freshwater organisms, no standard toxicity tests with saltwater organisms have been conducted with any polynuclear aromatic hydrocarbon. There are a variety of data for bioconcentration during short exposures.

EFFECTS

Miscellaneous

Lu, et al. (1977) conducted studies with benzo[a]pyrene in a terrestrial-aquatic model ecosystem and observed bioconcentration factors after 3 days ranging from 930 for the mosquitofish to 134,248 for Daphnia pulex (Table 1). Bioconcentration factors for Daphnia magna and Hexagenia sp. for a shorter time were 200 to 3,500 (Table 1).

The bioconcentration factors for polynuclear aromatic hydrocarbons by saltwater species are lower than those observed with freshwater organisms but may be due to the short exposure periods (Table 1). A polychaete worm was exposed to various crude oil fractions and 96-hour LC₅₀ values were between 300 and 1,000 ug/l (Neff, et al. 1976a).

*The reader is referred to the Guidelines for Deriving Water Quality Criteria for the Protection of Aquatic Life and Its Uses in order to better understand the following discussion and recommendation. The following tables contain the appropriate data that were found in the literature, and at the bottom of each table are calculations for deriving various measures of toxicity as described in the Guidelines.

CRITERIA

The limited freshwater data base available for polynuclear aromatic hydrocarbons, mostly from short-term bioconcentration studies with two compounds, does not permit a statement concerning acute or chronic toxicity.

The available data for polynuclear aromatic hydrocarbons indicate that acute toxicity to saltwater aquatic life occurs at concentrations as low as 300 $\mu\text{g/l}$ and would occur at lower concentrations among species that are more sensitive than those tested. No data are available concerning the chronic toxicity of polynuclear aromatic hydrocarbons to sensitive saltwater aquatic life.

Table 1. Other data for polynuclear aromatic hydrocarbons

<u>Species</u>	<u>Chemical</u>	<u>Duration</u>	<u>Effect</u>	<u>Result (µg/l)</u>	<u>Reference</u>
<u>FRESHWATER SPECIES</u>					
<u>Alga,</u> <u>Oedogonium cardiacum</u>	Benz[a]pyrene	3 days	Model ecosystem, bioconcentration factor = 5,258	-	Lu, et al. 1977
<u>Snail,</u> <u>Physa sp.</u>	Benz[a]pyrene	3 days	Model ecosystem, bioconcentration factor = 82,231	-	Lu, et al. 1977
<u>Cladoceran,</u> <u>Daphnia pulex</u>	Benz[a]pyrene	3 days	Model ecosystem, bioconcentration factor = 134,248	-	Lu, et al. 1977
<u>Mosquito,</u> <u>Culex pipiens</u> <u>quinquefasciatus</u>	Benz[a]pyrene	3 days	Model ecosystem, bioconcentration factor = 11,536	-	Lu, et al. 1977
<u>Mosquitofish,</u> <u>Gambusia affinis</u>	Benz[a]pyrene	3 days	Model ecosystem, bioconcentration factor = 930	-	Lu, et al. 1977
<u>Protozoa,</u> <u>Paramecium caudatum</u>	Anthracene	60 min	90% lethal photo- dynamic response	0.1	Epstein, 1963
<u>Cladoceran,</u> <u>Daphnia magna</u>	Anthracene	1 hr	Bioconcentration factor = 200	-	Herbes, 1976
<u>Cladoceran,</u> <u>Daphnia pulex</u>	Anthracene	24 hrs	Bioconcentration factor = 760	-	Herbes & Risl, 1978
<u>Mayfly,</u> <u>Hexagenia sp.</u>	Anthracene	28 hrs	Bioconcentration factor = 3,500	-	Herbes, 1976
<u>Bluegill,</u> <u>Lepomis macrochirus</u>	Benzo-(a)-anthracene	6 mos	87% mortality	1,000	Brown, et al. 1975
<u>SALTWATER SPECIES</u>					
<u>Eastern oyster,</u> <u>Crassostrea virginica</u>	Benz[a]pyrene	14 days	Bioconcentration factor = 242	-	Couch, et al. In press

Table 1. (Continued)

<u>Species</u>	<u>Chemical</u>	<u>Duration</u>	<u>Effect</u>	<u>Result ($\mu\text{g/l}$)</u>	<u>Reference</u>
Clam, <u>Rangia cuneata</u>	Benzolalpyrene	24 hrs	Bioconcentration factor = 8.66	-	Neff, et al. 1976a
Clam, <u>Rangia cuneata</u>	Benzolalpyrene	24 hrs	Bioconcentration factor = 236	-	Neff, et al. 1976b
Clam, <u>Rangia cuneata</u>	Chrysene	24 hrs	Bioconcentration factor = 8.2	-	Neff, et al. 1976a
Mudsucker, <u>Gillichthys mirabilis</u>	Benzolalpyrene (edible tissue)	96 hrs	Bioconcentration factor = 0.048	-	Lee, et al. 1972
Tidepool sculpin, <u>Oligocottus maculosus</u>	Benzolalpyrene (edible tissue)	1 hr	Bioconcentration factor = 0.13	-	Lee, et al. 1972
Sand dab, <u>Citharichthys stigmacus</u>	Benzolalpyrene (edible tissue)	1 hr	Bioconcentration factor = 0.02	-	Lee, et al. 1972
Polychaete worm, <u>Neanthes arenaceodentata</u>	Crude oil extract (fluorene)	96 hrs	LC50	1,000	Neff, et al. 1976a
Polychaete worm, <u>Neanthes arenaceodentata</u>	Crude oil fraction (phenanthrene)	96 hrs	LC50	600	Neff, et al. 1976a
Polychaete worm, <u>Neanthes arenaceodentata</u>	Crude oil fraction (1-methyl- phenanthrene)	96 hrs	LC50	300	Neff, et al. 1976a

REFERENCES

- Brown, E.R., et al. 1975. Tumors in fish caught in polluted waters: Possible explanations. Comparative Leukemia Res. 1973, Leukemogenesis. Univ. Tokyo Press/Karger, Basel. p. 47.
- Couch, J.A., et al. The American Oyster as an Indicator of Carcinogens in the Aquatic Environment. In: Pathobiology of Environmental Pollutants - Animal Models and Wildlife as Monitors. Storrs, Conn. National Academy of Sciences. (In press)
- Epstein, S.S., et al. 1963. The photodynamic effect of the carcinogen, 3,4-benzopyrene, on Paramecium caudatum. Cancer Res. 23: 35.
- Herbes, S.E. 1976. Transport and Bioaccumulation of Polycyclic Aromatic Hydrocarbons (PAH) in Aquatic Systems. In: Coal Technology Program Quarterly Progress Report for the Period Ending December 31, 1975. Oak Ridge National Lab., Oak Ridge, Tennessee. ORNL-5120. p. 65.
- Herbes, S.E. and G.F. Risi. 1978. Metabolic alteration and excretion of anthracene by Daphnia pulex. Bull. Environ. Contam. Toxicol. 19: 147.
- Lee, R.G., et al. 1972. Uptake, metabolism and discharge of polycyclic aromatic hydrocarbons by marine fish. Mar. Biol. 17: 201.

Lu, P., et al. 1977. The environmental fate of three carcinogens; benzo-(a)-pyrene, benzidine, and vinyl chloride evaluated in laboratory model ecosystems. Arch. Environ. Contam. Toxicol. 6: 129.

Neff, J.M., et al. 1976a. Effects of Petroleum on Survival, Respiration and Growth of Marine Animals. In: Sources, Effects and Sinks of Hydrocarbons in the Aquatic Environment. Proc. of a Symposium, American University, Washington, D.C. Amer. Inst. of Biol. Sci.

Neff, J.M., et al. 1976b. Accumulation and release of petroleum-derived aromatic hydrocarbons by four species of marine animals. Mar. Biol. 38: 279.

Mammalian Toxicology and Human Health Effects

EXPOSURE

Ingestion from Water

The uptake of polynuclear aromatic hydrocarbons (PAH) in humans from water occurs through the consumption of drinking water. In the United States, the sources of drinking water are ground waters and surface waters, such as lakes and rivers. Although a small amount of PAH originates from natural or endogenous sources, the predominant sources of PAH in surface waters are man-made. The discharges of raw and industrial sewage, atmospheric fallout and precipitation, road runoff, and leaching from polluted soils, all of which contain substantial PAH concentrations (Andelman and Suess, 1970), contribute to the PAH contamination in surface waters. Other than leaching from soils, the only source of PAH in ground water is of endogenous origin. Borneff (1977) estimated that low-level contaminated river and lake waters contain five times higher PAH concentrations than ground water, whereas in medium-level polluted river and lake waters this value may be 10 to 20 times higher. The concentration of PAH in ground water obtained by various authors is given in Table 1.

The PAH level in surface waters was determined by a number of German, English, and Russian workers. In all of these methods, the PAH were solvent extracted from the water, subjected to clean-up procedures and analyzed either by TLC-spectrofluorometry or by u.v.-spectrophotometry. These values are presented in Table 2.

Keegan (1971) analyzed the PAH content in three relatively unpolluted U.S. river waters by removing the PAH from water by sol-

TABLE 1

PAH Concentration in Ground Water

Source	Concentration, $\mu\text{g/l}$			Reference
	BaP	Carcinogenic PAH	Total PAH	
G. Finthen, Germany,		0.002		Borneff, 1964
Mainz, Germany		0.005		Borneff, 1964
Unspecified locations in Germany	0.0004	0.003	0.04	Borneff and Kunte, 1964
Average of 12 German ground waters*			0.06	Borneff and Kunte, 1969
Champaign, Ill.*	N.D. ^a	0.003	0.007	Basu and Saxena, 1977
Elkhart, Ind.*	0.004	0.004	0.02	Basu and Saxena, 1977
Fairborn, O.*	0.0003	0.0008	0.003	Basu and Saxena, 1977

*These are results of 6 specified PAH

^aN.D.: not detected

TABLE 2

Concentration of PAH in Surface Waters

Source	Concentration, $\mu\text{g}/\text{l}$			Reference
	BaP	Carcinogenic PAH	Total PAH	
Rhine River at Mainz	0.08	0.49	1.12	Borneff and Kunte, 1964
River Main at Seligenstadt	0.0024	0.155	0.48	Borneff and Kunte, 1964
River Danube at Ulm	0.0006	0.067	0.24	Borneff and Kunte, 1964
River Gersprenz at Munster	0.0096	0.047	0.14	Borneff and Kunte, 1964
River Aach at Stockach	0.017	0.95	2.5	Borneff and Kunte, 1965
River Schussen	0.01	0.20	1.0	Borneff and Kunte, 1965
River Plyussa: at Shale-oil effluent discharge site 3,500 m downstream	12			Dikun and Makhinenko, 1963
at Navy water intake	1			Dikun and Makhinenko, 1963
	0.1			Dikun and Makhinenko, 1963
A river: 15 m below coke by-product discharge site	8-12			Fedorenko, 1964
500 m downstream	2-3			Fedorenko, 1964
Thames River at Kew Bridge	0.13	0.18	0.50	Harrison, et al. 1975
at Albert Bridge	0.16	0.27	0.69	Harrison, et al. 1975
at Tower Bridge	0.35	0.56	1.33	Harrison, et al. 1975

vent extraction. The extract was subjected to clean-up and the PAH were analyzed by TLC-spectrofluorometry. Only samples from the Oyster River showed detectable amounts of four PAH. No PAH could be detected in the other two water samples from the Cocheco and Winnepesaukee Rivers.

The PAH levels in surface waters used as raw water sources for drinking water, and the effects of treatments of these waters on PAH levels, are shown in Table 3.

According to Borneff (1977), in surface waters, one-third of the total PAH is bound to larger suspended particles, a third is bound to finely dispersed particles, and the last third is present in dissolved form. The particle-bound portion of PAH can be removed by sedimentation, flocculation, and filtration processes. The remaining one-third dissolved PAH usually requires oxidation for partial removal/transformation. The use of Cl_2 , ClO_2 , O_3 , and uv light for this purpose has been studied. According to Borneff (1977), 50 to 60 percent of BaP can be removed by chlorination of water. However, the total PAH is reduced to a smaller degree by chlorination. ClO_2 on the other hand, reduces BaP concentration by 90 percent. But at BaP concentrations lower than 10 ppt, ClO_2 no longer functions as an oxidant for the transformation of BaP. The transformation of PAH is faster with O_3 , but the use of O_3 requires intensified prepurification to prevent oxidation of other chemicals. Filtration with activated carbon has been suggested by Borneff (1977) as the best method for PAH removal/transformation during water treatment. The reduction of BaP concentration with activated carbon was 99 percent efficient in actual field tests

TABLE 3

Concentrations of PAH in Raw and Treated Surface Water
used as Drinking Water Sources

Source	Treatment	Concentration, $\mu\text{g/l}$			Reference
		BaP	Carcinogenic PAH	Total PAH	
River Rhine	Untreated	0.082	0.485	1.11	Borneff and Kunte, 1964
River Rhine	Bank and activated carbon filtered	0.0005	0.015	0.13	Borneff and Kunte, 1964
Lake Constance	Untreated	0.0013	0.030	0.065	Borneff and Kunte, 1964
Lake Constance ^a	Rapid sand filtration chlorination	0.0017	0.017	0.053	Borneff and Kunte, 1964
English River	Untreated	0.06 ^b	0.37 ^c	0.73 ^b	Harrison, et al. 1976
English River	Filtration and chlorination	0.009	0.051 ^c	0.24	Harrison, et al. 1976
Monongahela River at Pittsburgh	Untreated	0.04	0.14	0.60	Basu and Saxena, 1978
same as above	Treated ^d	0.0004	0.002	0.003	Basu and Saxena, 1978
Ohio River at Huntington, W. Va.	Untreated	0.006	0.020	0.058	Basu and Saxena, 1978
same as above	Treated ^d	0.0005	0.002	0.007	Basu and Saxena, 1978
Ohio River at Wheeling, W. Va.	Untreated	0.21	0.57	1.59	Basu and Saxena, 1977
same as above	Treated ^d	0.002	0.011	0.14	Basu and Saxena, 1977
Delawater River at Philadelphia	Untreated	0.04	0.16	0.35	Basu and Saxena, 1978
same as above	Treated ^d	0.0003	0.002	0.015	Basu and Saxena, 1978
Lake Winnebago at Appleton, Wis.	Untreated	0.0006	0.002	0.007	Basu and Saxena, 1977
same as above	Treated ^d	0.0004	0.002	0.006	Basu and Saxena, 1977

^aThese are average of five determinations with the exclusion of a sixth high value.

^bThese values are estimates on the basis of average PAH adsorption in reservoir.

^cThese values may be a little higher due to the inability of separation of all the carcinogenic from non-carcinogenic PAH.

^dThe treatment included flocculation, activated carbon addition, filtration, pH control, chlorination and fluoridation.

(Borneff, 1977). With the exception of Appleton, Wis. drinking water, this finding of Borneff (1977) has been validated by the work of Basu and Saxena (1977, 1978), who demonstrated an 88 to 100 percent reduction of PAH in U.S. drinking waters by the use of activated carbon. In the case of Appleton, Wis. water, the initial PAH level in raw water was very low. Therefore, it can be concluded that below a certain minimum concentration, activated carbon may not be very effective for PAH removal/transformation.

As some derivatives of BaP and other PAH are formed during the disinfection of water with oxidizing agents and u.v. radiation, it is of interest to examine briefly the carcinogenicity of such derivatives. With the exception of alkylated derivatives, most BaP derivatives at best have only weak carcinogenic activity (Butenandt and Dannenberg, 1956). However, 10-chloro-compounds do cause tumors (Andelman and Suess, 1970). The quinones, some of which are also formed during chlorination (Andelman and Suess, 1970) do not produce tumors (Butenandt and Dannenberg, 1956), and may, in fact, inhibit the activity of other carcinogens (Buu-Hoi, 1959). The possibility of transformation of PAH into other carcinogenic compounds during water treatment processes is an area which remains largely unexplored.

The PAH content in U.S. drinking waters was analyzed by Basu and Saxena (1977, 1978). Six representative PAH recommended by the World Health Organization (WHO, 1970), as the measure of PAH contamination in drinking water, were monitored in this study (BbFL was replaced by BjFL) and the average concentration of PAH was found to be 13.5 ng/l. The U.S. EPA also conducted the National

Organic Monitoring Survey (NOMS) (U.S. EPA, 1977) to determine the frequency of occurrence and the levels of PAH in U.S. drinking water supplies. Of the 110 water samples analyzed, none showed any PAH other than fluoranthene. Seventeen out of 110 samples analyzed showed positive fluoranthene values with an average of 20 ng/l concentration. It should be mentioned that the detection limit of PAH in this study was as high as 50 ng/l. The PAH levels in various drinking waters are shown in Table 4.

Finished waters from various treatment sites are transported to the consumers through a variety of pipelines. Borneff (1977) reported a 10-fold increase in PAH concentration from beginning to end of a water supply pipe that resulted from the paint used on the water pipes. Leaching of PAH from the coating materials used on the pipes could possibly cause an increase in their concentration in the water reaching consumers. In other instances, PAH could be adsorbed from the water onto the surface of the pipes causing a decrease in their concentration. In the United States, two kinds of pipes are commonly used as distribution lines for transporting treated waters. These are cast/ductile iron, asbestos/cement pipes, and a combination of these. The effect of contact with these pipes on the quality of drinking water in terms of PAH concentration was studied by Basu and Saxena (1977). Because of the intermixing of the pipes, it is difficult to draw definite conclusions from their results. However, it seems likely that in instances where an enhancement of PAH concentration was observed, the tar/asphalt coating of the pipes was responsible for the increase. Cement-coated pipes, on the other hand, produced lower PAH concentrations, possibly due to adsorption of PAH from the water.

TABLE 4
PAH Levels in a Few Drinking Waters

Source	Concentration, ng/l			Reference
	BaP	Carcinogenic PAH	Total PAH	
Mixed tap water at Mainz, Germany			7.0	Borneff, 1964
Water at: ^a				
Syracuse, N.Y.	0.3	0.3	1.1	Basu and Saxena, 1978
Buffalo, N.Y.	0.2	0.2	0.9	Basu and Saxena, 1978
New York, N.Y.	0.5	3.9	6.4	Basu and Saxena, 1978
Lake George, N.Y.	0.3	1.5	4.2	Basu and Saxena, 1978
Endicott, N.Y.	0.2	1.1	8.3	Basu and Saxena, 1978
Hammondsport, N.Y.	0.3	1.5	3.5	Basu and Saxena, 1978
Pittsburgh, Pa.	0.4	1.9	2.8	Basu and Saxena, 1978
Philadelphia, Pa.	0.3	2.0	14.9	Basu and Saxena, 1978
Huntington, W. Va.	0.5	2.0	7.1	Basu and Saxena, 1978
Wheeling, W. Va.	2.1	11.3	138.5	Basu and Saxena, 1977
New Orleans, La.	1.6	1.6	2.2	Basu and Saxena, 1978
Appleton, Wis.	0.4	2.4	6.1	Basu and Saxena, 1977
Champaign, Ill,	N.D. ^b	1.2	2.8	Basu and Saxena, 1977
Fairborn, Ohio	0.1	0.8	2.9	Basu and Saxena, 1977
Elkhart, Ind.	N.D. ^b	0.3	0.3	Basu and Saxena, 1977

^aOnly the six WHO (1970) - recommended PAH were analyzed, with the exception that BbFL replaced B_jF_l. PAH were concentrated by passing 60 liters of drinking water through polyurethane foams. The eluate from the foams was subjected to cleanup and analyzed for PAH by TLC-spectrofluorometry.

^bN.D.: not detected.

There are very few epidemiological studies concerning the correlation between cancer and drinking water. It was, nevertheless, noted that four London boroughs, supplied largely by well water, had lower cancer mortalities than most of the other boroughs, which were supplied with surface water (Stocks, 1947). Another study concluded that the highest cancer death rates occurred in communities supplied by river water, followed by communities supplied by well water, and health water (Diehl and Tromp, 1953; Tromp, 1955). However, none of these studies attempted to correlate cancer morbidity with concentrations of PAH. Finally, it should be noted that one epidemiological study of the incidence of gastric cancer concluded that social factors and the kinds of soils present reduced the correlations otherwise obtained with the type of domestic water supply (Wynne-Griffith and Davies, 1954; Davies and Wynne-Griffith, 1954).

Although the levels of PAH detected in U.S. drinking waters are well below the WHO (1970) recommended limit of 200 parts per trillion (ppt), the health hazards associated with repeated exposure (more effective than an equivalent single dose (Payne and Hueper, 1960) of carcinogens through drinking water should not be underestimated. Shabad and Il'nitskii (1970) stated that the amount of carcinogenic PAH consumed by man from water is typically only 0.1 percent of the amount he consumes from foods. If the total PAH uptake from food is taken as 4.15 mg/year (Borneff, 1977), the human uptake of PAH from drinking water should not exceed 4 μ g/year. Assuming the PAH concentration value of 13.5 ng/l in U.S. drinking water (Basu and Saxena, 1977,1978), and a daily consump-

tion of 2.5 liters of drinking water, the yearly intake of PAH from U.S. drinking water would be 12.3 μg or 0.3 percent of the total food intake. Nevertheless, the accumulation of PAH in edible aquatic organisms through polluted surface waters can greatly increase their amount in foods, including fish, some mollusks, and edible algae (Andelman and Snodgrass, 1974). The use of contaminated water for irrigation can also spread PAH into other vegetable foodstuffs (Shabad and Il'nitskii, 1970). Therefore, it is important to monitor the PAH levels in surface waters not used as drinking water sources as well as drinking waters, in order to estimate accurately the human intake of PAH.

Ingestion from Food

PAH formed through both natural and man made sources can enter the food chain of man. There is considerable disagreement, however, concerning the contribution of each of these sources to the total PAH contamination in foods. From their work with marine algae and fishes obtained from polluted and unpolluted sources, Harrison, et al. (1975) concluded that endogenous synthesis may be the important factor for PAH contamination in these species. Others, however, believe that the endogenous formation of PAH occurs to such a limited extent that it is completely masked by the accumulation of PAH from the environment (Payer, et al. 1975). The latter conclusion was verified by Shabad and Smirnov (1972). It has been demonstrated by these authors that plants near an airport contained 10 to 20 times more BaP than areas remote from the runway. The results of Dunn and Stich (1976) indicated a correlation between the PAH level in mussels with industrial, urban, and recre-

ational activity. The highest occurrence of BaP in marine organisms in the areas adjacent to the sea lanes tends to support the view that exogenous sources are the predominant factor for PAH contamination in foods.

The primary routes of entry for PAH in foods are surface adsorption and biological accumulation from the environment (Binet and Malet, 1964). The adsorption of PAH from the soil by various plant roots and translocation to the shoots is well documented (Lo and Sandi, 1978). Similarly, the absorption of PAH by other marine organisms has been demonstrated by Lee, et al. (1972). Oysters and clams collected from moderately polluted waters also concentrate PAH via absorption (Cahnmann and Kuratsune, 1957; Guerrero, et al. 1976). The waxy surface of some plant leaves and fruits can concentrate PAH through surface adsorption (Hetteche, 1971; Kolar, et al. 1975). Kolar, et al. (1975) have shown that the concentration of BaP in vegetation is proportional to the exposure time during the growing season (bioaccumulation through adsorption) and the structure of the surface of the plant (surface adsorption). The above-ground parts of the vegetables contain more BaP than underground parts. Only about 10 percent of the externally deposited BaP in lettuce, kale, spinach, leeks, and tomatoes can be removed by cold water washing (Kolar, et al. 1975).

Food additives and food packaging materials such as paraffin waxes containing PAH, contribute to the enhancement of PAH levels in processed foods. For example, Swallow (1976) found that paraffin wax wrapping for food contained PaA, CH, BeP, and BaP at levels of 29 ppb, 2 ppb, 0-48 ppb, and 2 ppb, respectively. Certainly,

some of these PAH in the packing material can diffuse into the food. Hexane, a commercial solvent used to extract edible vegetable oils, is also a source of PAH contamination. PAH present in food-grade carbon blacks used for food processing can be transported to the food products. Curing smoke and other pyrolysis products used during cooking add to the level of PAH in food. However, in raw foods which require cooking, the largest source of PAH contamination originates from the cooking process itself.

In order to summarize the available data on PAH levels, various foods have been categorized following the pattern of USDA-FDA for total diet samples (Martin and Duggan, 1968). These are shown in table form later in the text. It should be recognized that the data presented in the tables are neither exhaustive nor absolute. Not all the PAH detected by the various authors are listed in these tables. Only the most frequently detected PAH are listed. The concentration values given in these tables are subject to considerable variation. The PAH concentrations in uncooked foods largely depend on the source of food. For example: vegetables, fruits, and fishes obtained from a polluted environment can be expected to contain higher concentrations of PAH. Therefore, the PAH content is subject to regional variation. In the case of raw foods which require cooking, the method of cooking is largely responsible for the PAH content in the food and is subject to regional or even personal variation. Therefore, the frequency of occurrence of PAH in a particular food is dependent on a number of factors. The results presented in Tables 5 and 6 represent only the values where the sample showed detectable levels of PAH.

TABLE 5

PAH Concentrations (ppb) in a few Vegetable Oils and Margarine

	A	PA	FL	P	BaA	BeP	BaP	BPR	CH
Corn ^a				3.1	0.8	0.7	0.7	0.6	
Coconut ^b	36	51	18.0	15.0	2.0	2.0			12
Margarine ^c					1.4- 29.5	0.5- 1.2	0.2- 6.8		
Sunflower ^c					13 ^d	4.0	8.0	4.0	
Soybean ^a			1.3	1.6	0.9	1.6	1.4	1.0	
Olive ^a			3.2	2.6	1.0	0.4	0.5	0.9	
Peanut ^a			3.3	2.9	1.1		0.6	0.9	

^aHoward, et al. 1966c^bBiernoth and Rost, 1967^cSwallow, 1976^dThis value represents concentration of BaA and CH

TABLE 6

PAH Concentrations (ppb) in Smoked and Nonsmoked Fish

Fish	F	A	PA	FL	P	BaA	BeP	BaP	PR	BPR
Smoked eel ^a	9.0	4.0	37.0	4.0	6.0		t ^b	1.0		
Smoked lumpfish ^a	5.0	t	10.0	2.0	1.0	t	t	0		
Smoked trout ^a	67.0	26.0	52.0	12.0	5.0		t	t		
Smoked herring ^b				3.0	2.2					
Smoked herring ^b (dried)				1.8	1.8	1.7	1.2	1.0		1.0
Smoked salmon ^b				3.2	2.0	0.5	0.4			
Smoked sturgeon ^b				2.4	4.4			0.8		
Smoked whitefish ^b				4.6	4.0			4.3		
Smoked whiting ^c					0.5			6.6	0.7	2.4
Smoked redfish ^d		1.5	4.1	4.0	3.0		0.3	0.3		
Smoked cod ^d					0.6			4.0	0.4	2.2
Electric smoked mackerel ^d	2.6	1.9	9.0	5.2	3.6	1.2	0.5	0.2	t	0.2
Gas smoked mackerel ^d	8.2	2.3	11.0	2.6	4.0	0.6	0.2	0.3	t	0.3
Non-smoked haddock ^b				1.6	0.8					
Non-smoked herring ^b (salted)				0.8	1.0					
Non-smoked salmon				1.8	1.4					

^aThorsteinsson, 1969; Dungal, 1961^bHoward, et al. 1966a^cMalanoski, et al. 1968^dMasuda and Kuratsune, 1971

t = trace

It has been claimed by Zitko (1975) that PAH are not bioaccumulated along the food chain. However, Bjørseth (1978) demonstrated that both common and horse mussels bioaccumulated PAH, although not to the same degree. Dunn and Stitch (1976) have shown that mussels cannot metabolize BaP upon their removal from water. In water, mussels released 79 percent of naphthalene in three days, with a half-life of 1.3 days. The BaP released from both clams and mussels in water takes place with a half-life of two to five weeks (Dunn and Stitch, 1976).

The human intake of PAH through the digestive system has been estimated by Borneff (1977). According to this estimate the human intake of PAH per year is about 3 to 4 mg from fruits, vegetables, and bread, 0.1 mg from vegetable fats and oils, and about 0.05 mg from smoked meat or fish and drinking water.

Vegetable Fats, Oils, and Shortening: Several PAH have been found in edible oils by European workers (Howard and Fazio, 1969). The PAH levels in a few vegetable oils and margarine are presented in Table 5. PAH other than those shown in Table 5 have been reported in these oils (Swallow, 1976). Since the concentration of PAH in vegetable oils depends on the nature of refinement of the crude oil (Grimmer and Hildebrandt, 1967), one can expect variations in their concentrations. Heating of the oils also leads to a slight increase in PAH concentrations. For example, Lijinsky and Shubik (1965b) did not detect any PAH in uncooked Wesson and Crisco oil. However, oil used previously for deep-frying of food showed 1.4 ppb BaP, 12 ppb FL, and 6 ppb pyrenes (Lijinsky and Ross, 1967; Malanowski, et al. 1968).

Swallow (1976) determined the level of PAH in butter and found the concentration of BaA + CH, BaP, IP + DBA, and BPR to be 1 ppb. In a total diet study with a composite sample containing the fats, oils, and shortening, Howard, et al. (1968) found less than 0.5 ppb of seven PAH. However, Borneff (1977) estimated that the human intake of PAH from vegetable fats and oils amounted to 0.1 mg/year.

Fish and Other Marine Foods: Raw fish from unpolluted waters usually do not contain detectable amounts of PAH, but smoked or cooked fish contain varying levels of PAH. In addition to the origin of the fish, (polluted or unpolluted water), the amount of PAH in smoked fish depends on various parameters, such as type of smoke, temperature of combustion, and degree of smoking (Draudt, 1963).

The skin of fish apparently serves as a barrier to the migration of PAH into the body tissues. This was postulated by Malanowski, et al. (1968) from their observations that the BaP level in the skin was much higher than in the interior of cooked fish.

The PAH levels in various smoked and unsmoked fish are shown in Table 6. In addition to the fish presented in this table, various other marine organisms had been tested for PAH content. For example, cooked squid and prawns had BaP concentrations of 1.04 ppb and 0.08 ppb, respectively (Shiraishi, et al. 1975). Various other edible marine organisms were investigated and found to contain PAH. Swallow (1976) analyzed smoked oysters and determined the levels of BaA + Ch, BbFL + BkFL + BjFL, IP + DBA and BPR to be 19 ppb, 8 ppb, 9 ppb, 7 ppb, and 3 ppb, respectively. Cooked scallops were found to contain 9.9 ppb BaP (Shiraishi, et al. 1975). Shiraishi, et al.

(1973) detected 0 to 31.3 ppb BaP in various Japanese seaweeds. However, no BaP was detected in crab (Shiraishi, et al. 1975). The absence of BaP in crab is corroborated by the work of Lee, et al. (1976), who found no evidence of PAH storage by any of the crab tissues.

A bioconcentration factor (BCF) relates the concentration of a chemical in aquatic animals to the concentration in the water in which they live. The steady-state BCFs for a lipid-soluble compound in the tissues of various aquatic animals seem to be proportional to the percent lipid in the tissue. Thus the per capita ingestion of a lipid-soluble chemical can be estimated from the per capita consumption of fish and shellfish, the weighted average percent lipids of consumed fish and shellfish, and a steady-state BCF for the chemical.

Data from a recent survey on fish and shellfish consumption in the United States were analyzed by SRI International (U.S. EPA, 1980). These data were used to estimate that the per capita consumption of freshwater and estuarine fish and shellfish in the United States is 6.5 g/day (Stephan, 1980). In addition, these data were used with data on the fat content of the edible portion of the same species to estimate that the weighted average percent lipids for consumed freshwater and estuarine fish and shellfish is 3.0 percent.

No measured steady-state BCF is available for any of the following compounds, but the equation " $\text{Log BCF} = (0.85 \text{ Log P}) - 0.70$ " can be used (Veith, et al. 1979) to estimate the steady-state BCF for aquatic organisms that contain about 7.6 percent lipids (Veith,

1980) from the octanol/water partition coefficient (P). The log P values were obtained from Hansch and Leo (1979) or were calculated by the method described therein. The adjustment factor of $3.0/7.6 = 0.395$ is used to adjust the estimated BCF from the 7.6 percent lipids on which the equation is based to the 3.0 percent lipids that is the weighted average for consumed fish and shellfish in order to obtain the weighted average bioconcentration factor for the edible portion of all freshwater and estuarine aquatic organisms consumed by Americans (Table 7). Caution must be exercised in application of common practice in obtaining the BCF as described here, because the ecological impact of PAH is not well understood at this time. Numerous studies show that despite their high lipid solubility, PAH show little tendency for bioaccumulation in the fatty tissues of animals or man (Lee, et al. 1972; Ahokas, et al. 1975; Graf, et al. 1975). This observation is not unexpected, in light of convincing evidence to show that PAH are rapidly and extensively metabolized. Since only low levels of PAH are detected in plants and lower organisms, (Radding, et al. 1976), transfer of PAH through the food chain does not seem likely. The direct impact of PAH on plants, animals, or the ecological balance of nature is difficult to evaluate, since few data are available which suggest that adverse effects may occur. Thus it is appropriate in the case of PAH to use the octanol-water partition coefficient for estimation of the BCF. Instead a more realistic value of 30, based on the work of Lu, et al. (1977) in fish, is recommended for criteria derivation.

TABLE 7

Calculated Bioconcentration Factors of PAH
Based upon the Octanol/Water Partition Coefficient

Chemical	Log P	Estimated Steady State BCF	Weighted Average BCF
Acenaphthalene	3.74	301	119
Anthracene	4.45	1,210	478
Benzo(a) pyrene	6.06	28,200	11,100
Phenanthrene	4.46	1,230	486
3-Methylcholanthrene	6.97	168,000	66,400
1-Methylphenanthrene	5.00	3,550	1,400
Dibenzofuran	4.12	634	250
Fluoranthene	4.90	2,920	1,150
Fluorene	4.18	713	282
Benz(a)anthracene	5.61	11,700	4,620
1,12-Benzoperylene	6.51	68,200	26,900
Benzo(K) fluoranthene	6.06	28,200	11,100
Benzo(B) fluoranthene	6.06	28,200	11,100
Chrysene	5.61	11,700	4,620
Dibenzo(A,H)anthracene	6.77	113,000	4,460
2,3-Phenylene pyrene	6.51	68,200	26,900
Pyrene	4.88	2,800	1,110
Dibenz(A,H)acridine	5.73	14,800	5,850

Meat and Meat Products: Raw meat does not normally contain PAH, but smoked or cooked meat may contain varying amounts of PAH (Lo and Sandi, 1978). Table 8 shows the concentration of PAH detected in a few meats and meat products. The higher concentration of PAH in charcoal broiled ribs (containing more fats) than in charcoal broiled steaks tends to support the idea that the most likely source of PAH is the melted fat. These fats drip on the heat source and are pyrolyzed. The PAH compounds in the smoke are then deposited on the meat as the smoke rises (Lijinsky and Shubik, 1965a). Many factors, such as degree of smoking, and the temperature of combustion affect the composition and concentration of PAH in cooked meat (Howard, et al. 1966a). In addition to the pyrolysis of fats, incomplete combustion of charcoal can also contribute to the PAH content in broiled meat. Thus, the source of heat used for cooking is responsible for the PAH concentration in cooked meats. These effects are indicated in Table 9.

In North America, except for smoked ham, most smoked meats contained much less carcinogenic PAH than European samples (Howard, et al. 1966a,b). The high incidence of stomach carcinoma in Iceland has been explained by the high concentration of BaP in smoked trout and mutton which are consumed in large quantities in the area (Bailey and Dungal, 1958). On the other hand, very low concentrations of PAH in Norwegian bologna sausages (see Table 8) are probably indicative of the tradition of light smoking of food in Norway (Frethein, 1976).

About 60 to 75 percent of the BaP in smoked food has been found to be in the superficial layer of meat (Thorsteinsson, 1969). This

TABLE 8
PAH Concentrations (ppb) in a Few Smoked Meat and Meat Products

Meat	A	PA	FL	P	BaA	BeP	BaP	PR	BPR	CH
Charcoal broiled steaks ^a		21.0	43.0	35.0	1.4	5.5	5.8	0.9	6.7	0.6
Barbecued ribs ^a	7.1	58.0	49.0	42.0	3.6	7.5	10.5	1.5	4.7	2.2
Smoked beef (chipped) ^b			0.6	0.5	0.4					
Smoked ham ^b			14.0	11.2	2.8	1.2	3.2		1.4	
Smoked pork (roll) ^b			3.1	2.5						
Smoked frankfurters ^b			6.4	3.8	1.5	2.0				
Barbecued beef ^e			2.0	3.2	13.2	1.7	3.5		4.3	9.6
Smoked hot sausages ^c				1.5	0.5		0.4			1.0
Smoked mutton ^d	13.0	104.0	18.0	8.0	2.0	5.0	t ^h			
Smoked mutton sausages ^d	2.0	17.0	6.0	2.0	0.5	t	t			
Smoked bologna ^e					0.04- 0.55	5.0	0.04- 0.08 2.0	0.04- 0.07	0.04- 0.20	0.15- 1.20 ⁱ
Smoked salami ^f	0.7	D ^g	5.6	5.2	0.6	0.2	0.8	3.2	D	1.2
Smoked Mortadella ^f	2.6	D	22.0	15.0	2.8	1.8	0.7	0.1	0.4	3.4
Heavily smoked bacon ^f	20.0	D	35.0	27.0	29.0	D	3.6	0.9	3.0	D

C-21

^aLijinsky and Shubik, 1965a
^bHoward, et al. 1966a,b; Panalaks, 1976
^cMalanoski, et al. 1968
^dThorsteinsson, 1969
^eFretheim, 1976; Panalaks, 1976

^fLo and Sandi, 1978
^gD = detected
^ht = trace
ⁱcompound unseparated

TABLE 9

Effect of Different Cooking Variables on the Concentration of PAH (ppb) in Cooked Meat

Meat	Effect	FL	P	BaA	BeP	BaP	BPR	CH	CR
Charcoal broiled									
hamburger ^a	Fat Content								
Fat, ^c hot ^d		13.3	7.7	2.7		2.6	14.9	1.7	1.0
Lean, ^e hot		0.3	1.6				0.9		
No-drip pan		0.2	0.1				t	t	
Charcoal broiled									
hamburger ^a	Heating temperature								
Lean, ^f hot		0.3	1.6				0.9	0.3	
Lean, cool		1.3	0.6						
Broiled T-bone									
steak ^a	Heat source								
Charcoal, hot		19.8	19.1	31.0	17.6	50.4	12.4	25.4	8.0
Flame, hot		19.0	20.0	3.9	5.7	4.4	6.2	2.0	9.0
Smoked ham ^b									
Light	Degree of Smoke	4.0-	2.0-	0.5-	0-2.0	3.0-	0-1.4	0-3.0	
		14.0	11.0	3.0		4.0			
Heavy		48.0-	35.0-	6.0-	4.0-	3.8-	2.5-	12.0-	
		156.0	161.0	33.0	26.0	55.0	25.0	66.0	

^aLijinsky and Ross, 1967^bFilipovic and Toth, 1971; Toth and Blass, 1972^cFat: 21% fat^dHot: 7 cm. from heat source^eLean: 7% fat^fCool: 25 cm from heat source

low penetration has also been noted by Rhee and Bratzier (1970), who observed that in smoked bologna sausages, the BaP is located within 1.5 mm from the surface. Cellulose casings can be used as a more effective barrier to BaP permeation during smoking of frankfurters than animal casing (Simon, et al. 1969).

In addition to meat and meat products, liquid smoke flavorings used during the cooking of meat have been found to contain a variety of PAH. Lijinsky and Shubik (1965b) have detected BaP, FL, P, BPR, BaA, and CH in liquid smoke at concentrations of 1 ppb, 16 ppb, 7 ppb, 1 ppb, 12 ppb, and 6 ppb, respectively. In liquid hickory smoke flavoring, Youngblood and Blumer (1975) found the total concentration of PAH as 9,400 ppm. The high level of PAH present in the resinous condensate in liquid smoke flavoring indicates the importance of its efficient removal from the aqueous flavoring prior to its use in foodstuffs (White, et al. 1971).

Vegetables, Fruits, Grains and Cereal Products, Sugar and Adjuncts, and Beverages: Various European and Japanese workers have reported the presence of BaP and other PAH in these products; their results are summarized in Table 10. Studies in this field in North America are lacking. Test results indicate that surface adsorption and root uptake are the principal modes of PAH accumulation in vegetables (Binet and Mallet, 1964). The frizzly leaf of kale, for example, has a large surface area and holds dust particularly well. PAH are adsorbed by the wax layer and protected against solar reactions (Hetteche, 1971). In kale, Hetteche (1971) found the concentration of PAH to be the following: PA, 70-586 ppb; A, 2.4-97.5 ppb; P, 36.2-510 ppb; FL, 53.6-1,196 ppb; BaA, 11.2-230 ppb; CH,

TABLE 10

BaP Content in Fruits and Other Foods

Fruits	Concentration (ppb)	Comments	References
Apple	0.02		Shiraishi, et al. 1975
Apple	8.3	Polluted environment	Kolar, et al. 1975
Banana	0.02		Shiraishi, et al. 1975
Banana peel	0.03		Shiraishi, et al. 1975
Grape	0.2	Polluted environment	Kolar, et al. 1975
Grape	0.02		Shiraishi, et al. 1975
Japanese pear	0.05		Shiraishi, et al. 1975
Pear	1.9	Polluted environment	Kolar, et al. 1975
Persimmon	0.02		Shiraishi, et al. 1975
Pineapple	0.02		Shiraishi, et al. 1975
Plums	0.04		Shiraishi, et al. 1975
Plums	29.7	Polluted environment	Kolar, et al. 1975
Dried prunes	0.2 to 1.5		IARC, 1973
Mandarin orange	0.03		Shiraishi, et al. 1975
Orange peel	0.15		Shiraishi, et al. 1975
Strawberry	N.D. ^a		Shiraishi, et al. 1975
Pumpkin	N.D. to trace		Shiraishi, et al. 1974

Table 10 (cont.)

Grains & Cereal Products			
Product	Concentration (ppb)	Comments	References
Wheat grain	0.1	Polluted environment	Kolar, et al. 1975
Wheat sprouts	60.0		Siddiqui and Wagner, 1972
Cereals	0.2 to 4.1		IARC, 1973
Barley	0.3	Polluted environment	Kolar, et al. 1975
Oats	0.2	Polluted environment	Kolar, et al. 1975
Polished rice	N.D. ^a		Shiraishi, et al. 1973
Rye seedling	10.0 to 20.0	8 other PAH identified	Graf and Nowak, 1966
Lentil seedlings	10.0 to 20.0	8 other PAH identified	Graf and Nowak, 1966
Sesame seeds	N.D.		Shiraishi, et al. 1973
Sugar and Adjuncts			
Product	Concentration (ppb)	Comments	References
Charred biscuits	11.0-72.0		Kuratsune, 1956
Caramel	N.D. ^a		Shiraishi, et al. 1973
Chocolate	0.2-1.7	4 other PAH quantified	Fabian, 1965

Table 10 (cont.)

Vegetables			
Vegetable	Concentration (ppb)	Comments	References
Parsley leaf and stem	24.3	Polluted environment	Kolar, et al. 1975
Red clover	7.5	Polluted environment	Kolar, et al. 1975
Mushroom	7.0	Polluted environment	Kolar, et al. 1975
Lettuce	8.6	Polluted environment	Kolar, et al. 1975
Lettuce	N.D.		Shiraishi, et al. 1974
Spinach	6.2	Polluted environment	Kolar, et al. 1975
Spinach	1.3		Shiraishi, et al. 1973
Spinach	7.4		IARC, 1973
Radish leaves	5.3	Polluted environment	Kolar, et al. 1975
Radish roots	1.2	Polluted environment	Kolar, et al. 1975
Radish roots	N.D. ^a		Shiraishi, et al. 1974
Tomatoes	0.1	Polluted environment	Kolar, et al. 1975
Tomatoes	0.2		IARC, 1973
Cabbage	12.3 to 20.9	Polluted environment	Kolar, et al. 1975
Cabbage	N.D.		Shiraishi, et al. 1974
Chinese cabbage	0.05		Shiraishi, et al. 1974
Potatoes	N.D. to 0.01		Shiraishi, et al. 1974
Potatoes	0.2	Polluted environment	Kolar, et al. 1975
Sweet potatoes	N.D.		Shiraishi, et al. 1974
Sweet pepper	N.D.		Shiraishi, et al. 1974
Cauliflower	5.1	Polluted environment	Kolar, et al. 1975
Bean paste	N.D.		Shiraishi, et al. 1973
Kidney bean	N.D.		Shiraishi, et al. 1973
Carrot	N.D. to 0.02		Shiraishi, et al. 1973
Cucumber	N.D.		Shiraishi, et al. 1973
Eggplant	N.D.		Shiraishi, et al. 1973
Onion bulb	N.D. to 0.01		Shiraishi, et al. 1974
Onion greens	0.01		Shiraishi, et al. 1974

Table 10 (cont.)

Beverages			
Beverage	Concentration (ppb)	Comments	References
Dark rum	1.0		Swallow, 1976
Whiskey	0.04	3 quinolines detected	IARC, 1973; Nishimura and Masuda, 1971
Tea leaves	3.9 to 21.3		IARC, 1973
Black tea aroma ^b		7 quinolines detected	Vitzthum, et al. 1975
Roasted coffee (moderate dark)	N.D.		Kuratsune and Hueper, 1960
Roasted coffee (darkest)	N.D. to 4.0		Kuratsune and Hueper, 1958, 1960
Coffee soots ^c	200.0-440.0		Kuratsune and Hueper, 1958

^aN.D. = Not detected

^bThis is the volatile components of black tea.

^cThese are the soots generated during direct and indirect roasting of coffee beans.

28.6-395 ppb; BeP, 3.8-67.2 ppb; BaP, 0.9-48.6 ppb; PR, N.D.-7 ppb; BPR, 1.2-46.4 ppb; and CR 0.1-7.2 ppb.

The concentration of BaP in vegetables is directly proportional to exposure time during the growing season and structure of the surface of the plant. The above-ground parts contain more BaP than underground parts. Washings with cold water do not remove more than 10 percent of the BaP (Kolar, et al. 1975). Fruits grown in polluted environments show a high degree of PAH contamination mainly through adsorption on the waxy surface.

In smoked Gouda cheese, Panalaks (1976) found 0.5 ppb BaP and Howard, et al. (1966a) found 2.8 ppb FL and 2.6 ppb P. The unsmoked cheese contained lower levels of PAH. Grimmer (1974) analyzed baker's yeasts and determined the level of PAH. The values are shown in Table 11.

Inhalation

A variety of PAH have been detected in ambient air in the United States and elsewhere in the world. Because of its carcinogenic properties, BaP has been most extensively monitored and has frequently been used as an indicator of ambient PAH. The presumed correlation between the concentration of BaP and other PAH, however, does not always exist. For example, a study by Kertesz-Saringer and Morlin (1975) found little or no relationship between BaP and other PAH in Budapest air. Gordon (1976) and Gordon and Bryan (1973) came to a similar conclusion from their work with ambient Los Angeles air.

The concentration and the nature of PAH in ambient air are dependent on a number of factors. In general, the PAH concentra-

TABLE 11
PAH Concentrations (ppb) in a Variety of Baker's Yeast^{a,b}

PAH	French	German	Scottish	Russian
PA	17.8-34.60	67.0	1,620	7.2
A	2.6-13.6	4.8-10.2	567	4.7
P	11.6-19.6	11.5-35.0	327	16.9
FL	18.5-21.2	17.2-66.8	93	32.1
BaA	9.8-23.3	2.5-15.8	203	10.8
CH	8.1-13.4	4.2-14.0	50	11.1
BeP	8.0-10.6	3.1-14.3	40.4	8.7
BaP	8.0-12.2	1.8-13.2	6.2	0.5
PR	0.9-1.2	N.D]-0.5	16.7	6.0

^aSource: Grimmer, 1974

^bThis is baker's yeast as opposed to dietary or brewer's yeast.

tion is lowest during the summer months and highest during the winter, (Sawicki, et al. 1962) probably due to commercial and residential heating during winter (U.S. EPA, 1974). However, there are some exceptions. Cleveland, Ohio, for instance, does not follow the high winter-low summer pattern (U.S. EPA, 1974). It has been suggested that this may be due to significant industrial emissions that are uniform throughout the year (U.S. EPA, 1974).

The nature and relative amounts of individual PAH in ambient air are also dependent on the source of these compounds. Thus, the content of PAH sampled in an industrial area is a composite of the emissions from various industrial and transportation sources within the area. For example, Gordon (1976), from his study of the relative PAH concentration pattern for different areas in Los Angeles, found a correlation between coronene concentration and automobile emissions. Similarly, Greinke and Lewis (1975) had demonstrated that emissions from coke ovens contain lower amounts of certain methyl-substituted PAH than emissions from petroleum pitch volatiles. Bartle, et al. (1974) also used a PAH profiling technique for the identification of air pollution sources, such as coal burning, vehicular emissions, and oil and gas burning.

Meteorological factors have a dominant effect on PAH concentrations. For example, Lunde and Bjørseth (1977) demonstrated that under favorable wind conditions PAH from downtown London could be transported to Norway. The tendency of atmospheric inversion to increase the PAH levels in urban areas has also been shown (Hoffmann and Wynder, 1977).

The annual average ambient BaP concentrations for different U.S. urban and rural locations during the period 1966-70 have been compiled by U.S. EPA report (Santodonato, et al. 1978). The average BaP concentrations in U.S. urban and rural areas obtained from this U.S. EPA study are shown in Table 12.

An interesting trend has developed from the National Air Surveillance Network (NASN) monitored BaP values listed in Table 12. As can be seen, the average BaP concentrations in urban areas decreased from 3.2 ng/m³ in 1966 to 2.1 ng/m³ in 1970, approximately a 30 percent decrease. The decrease is more dramatic (i.e., > 80 percent) between the period 1966 to 1976. Even the concentrations in rural areas indicate a downward trend. This decline in BaP concentration is believed to be due primarily to decreases in coal consumption for commercial and residential heating, improved disposal of solid wastes, and restrictions on open burning (Faoro and Manning, 1978). A further observation that can be made from Table 12 is the 5- to 10-fold difference in BaP concentration between urban and rural locations.

The NASN study did not include the determination of concentrations of other PAH. The summer and winter averages of ambient PAH concentrations for seven urban locations were determined by Sawicki, et al. (1962). The averages of summer and winter data from this work are presented in Table 13.

The average of total PAH concentrations for all cities listed in Table 13 is 46.4 ng/m³. However, these values were obtained from ambient air sampled in 1958-59 and probably have decreased during subsequent years. If an 80 percent decrease of total PAH

TABLE 12
 Average BaP Concentrations (ng/m³) in U.S. Urban
 and Rural Areas During 1966-76^a

Period	1966	1970	1976
Urban	3.2	2.1	0.5
Rural	0.4	0.2	0.1 ^b

^aSource: Santodonato, et al. 1978

^bThis value is the average of two rural locations.

TABLE 13

Summer-Winter Average of Ambient PAH Concentrations (ng/m³)
in the Air of Selected Cities^a

City	BPR	BaP	BeP	BkFL	P	CR	PR	A	Total
Atlanta	7.0	4.5	3.1	3.7	3.4	3.4	0.8	0.4	26.3
Birmingham	13.2	15.7	8.0	8.8	9.6	3.0	3.8	1.3	63.4
Detroit	21.3	18.5	14.2	12.5	19.4	4.1	3.9	1.2	95.1
Los Angeles	10.2	2.9	4.4	3.1	3.2	7.1	0.8	0.1	31.8
Nashville	10.2	13.2	7.6	8.0	15.3	3.0	2.3	1.0	60.6
New Orleans	6.0	3.1	4.8	2.9	1.3	14.8	0.6	0.1	33.6
San Francisco	5.1	1.3	1.7	1.0	1.0	3.3	0.2	0.1	13.7

^aSource: Sawicki, et al. 1962

concentration is assumed (as in the case of BaP), the present ambient PAH concentration in the U.S. urban areas can be extrapolated as 9.3 ng/m^3 . Although the concentration of BaP and some other PAH might have decreased in past decades, the concentration of corenene and some other PAH may not have maintained the same trend. This could be due to the higher number of automobiles in current use. Therefore, this 80 percent decrease figure may or may not be valid for all PAH.

The concentrations of PAH in recent years in individual U.S. cities have been determined by a number of authors. The lowest and highest values of these determinations published during the period 1971-77 are shown in Table 14.

The exact amount of human PAH intake from all modes is difficult to determine because of the different modes of inhalation due to smoking, occupational exposure, or exposure to ambient air. Considering only exposure to ambient air, one needs an average PAH concentration in air in order to determine the PAH intake through inhalation. In the absence of national average data for PAH equivalent to NASN data on national average BaP levels, the yearly average data for Los Angeles are used for the derivation of PAH intake due to inhalation. These values are given in Table 15.

It can be seen from Table 15 that the yearly intake of total PAH, carcinogenic PAH, and BaP through inhalation is $39.8 \text{ } \mu\text{g}$, $9.9 \text{ } \mu\text{g}$, and $1.9 \text{ } \mu\text{g}$, respectively. It should be recognized that these data are based on the average ambient air concentration of one city and probably will not reflect the true U.S. average. It is noteworthy, however, that the total ambient PAH concentration of 10.9

TABLE 14

PAH Concentration Range in U.S. Cities Determined
by Various Authors in Recent Years

Compound	Concentration, Range, ng/m ³	Reference
NA	0.052 - 0.350	Krstulovic, et al. 1977
A	0.068 - 0.278 ^a	Lunde and Bjørseth, 1977
BaA	0.18 - 4.6	Fox and Staley, 1976; Gordon, 1976
PA	0.011 - 0.340	Krstulovic, et al. 1977
FL	0.10 - 4.1	Fox and Staley, 1976; Hoffman and Wynder, 1977
BbFL	0.1 - 1.6	Gordon and Bryan, 1973
BjFL	0.01 - 0.8	Gordon and Bryan, 1973
BkFL	0.03 - 1.3	Gordon and Bryan, 1973
P	0.18 - 5.2	Fox and Staley, 1976; Gordon and Bryan, 1973
BaP	0.13 - 3.2	Colucci and Begeman, 1971; Fox and Staley, 1976
BeP	0.9 - 4.6	Gordon, 1976; Fox and Staley, 1976
IP	0.03 - 1.34	Gordon, 1976; Gordon and Bryan, 1973
CH	0.6 - 4.8	Gordon, 1976; Fox and Staley, 1976
PR	0.01 - 1.2	Gordon and Bryan, 1973
BPR	0.2 - 9.2	Gordon and Bryan, 1973
CR	0.2 - 6.4	Gordon and Bryan, 1973

^aThis Norwegian value is included because no recent U.S. data are available.

TABLE 15

Average Ambient PAH Concentration in U.S. and
Daily Intake of PAH Through Inhalation^a

PAH	BaP	Carcinogenic PAH ^b	Total PAH
Ambient conc., ng/m ³	0.5	2.7	10.9
Inhalation intake/day, ng ^c	5.0	27.0	109.0

^aThese values are based on the study of Gordon, 1976.

^bCarcinogenicity of PAH are derived from NAS, 1972.

^cThese values are based on 10 m³ inhalation of air/day.

ng/m³ derived in this work is very close to the earlier extrapolated value of 9.3 ng/m³.

Dermal

No direct information is available on the importance of dermal absorption in total human exposure to PAH. PAH can be absorbed across the skin by animals. For those humans exposed to only ambient levels of PAH, dermal absorption is not likely to be a significant route of entry.

PHARMACOKINETICS

There are no data available concerning the pharmacokinetics of PAH in humans. Nevertheless, it is possible to make limited assumptions based on the results of animal studies conducted with several PAH, particularly BaP. The metabolism of PAH in human and animal tissues has been especially well-studied, and has contributed significantly to an understanding of the mechanisms of PAH-induced cancer.

Absorption

The demonstrated toxicity of PAH by oral and dermal administration (Smyth, et al. 1962) indicates that they are capable of passage across epithelial membranes. The high lipid solubility of compounds in this class supports this observation. Animal studies with structurally-related PAH such as benzo(a)pyrene (BaP), chrysene, 7,12-dimethylbenz(a)anthracene (DMBA), benz(a)anthracene, and 3-methylcholanthrene (MCA) confirmed that intestinal transport readily occurs, primarily by passive diffusion (Rees, et al. 1971). In addition, there is ample evidence to indicate that benzo(a)pyrene, and presumably other PAH, are easily absorbed through the lungs (Kotin, et al. 1969; Vainio, et al. 1976).

Distribution

The tissue distribution and accumulation of PAH have not been studied in humans. It is known, however, that several PAH (e.g., benzo(a)pyrene, 7,12-dimethylbenz(a)anthracene, 3-methylcholanthrene, phenanthrene) become localized in a wide variety of body tissues following their absorption in experimental rodents (Kotin, et al. 1969; Bock and Dao, 1961; Dao, et al. 1959; Flesher, 1967). Relative to other tissues, PAH localize primarily in body fat and fatty tissues (e.g., breast) (Schlede, et al. 1970a,b; Bock and Dao, 1961).

Disappearance of BaP from the blood and liver of rats following a single intravenous injection was very rapid (Schlede, et al. 1970a). The concentration of BaP in the blood one minute after a 10 µg injection was 193 ± 29 ng/ml; after five minutes concentration of BaP in the blood was 31 ± 1 ng/ml. Similarly, in the liver, the half-time for BaP disappearance was about ten minutes. In both blood and liver, however, the initial rapid elimination phase was followed by a slower disappearance phase, lasting six hours or more. In the same experiment, disappearance of BaP from the brain was slower than from blood or liver, and the concentration of BaP in fat increased during the six-hour observation period. Schlede, et al. (1970a) concluded that a rapid equilibrium occurs for BaP between blood and liver, and that rapid disappearance from the blood is due to both metabolism and distribution into tissues. This contention is supported by data (Schlede, et al. 1970b) showing that pretreatment with BaP (which induces microsomal enzyme activity) accelerates both the rate of BaP disappearance from all

tissues and the excretion of BaP metabolites into the bile. The ability of BaP to stimulate its own metabolism may have important implications for human situations, where lifelong exposure to PAH is known to occur.

With certain PAH, passage into the fetus following intragastric or intravenous administration to pregnant rats has been variable (Shendrikova and Aleksandrov, 1974).

Metabolism

In the past, the relative lack of chemical reactivity for tumorigenic PAH has been puzzling in light of their dramatic biological effects. Early attempts to explain the carcinogenicity of various PAH utilized physico-chemical calculations (Pullman and Pullman, 1955). These early hypotheses were based on the assumption that those regions of the molecule favoring substitution or addition reactions would preferentially react with critical cellular target sites to initiate a carcinogenic transformation. This concept, however, did not prove successful for PAH.

More recently it was learned that PAH are metabolized via enzyme-mediated oxidative mechanisms to form reactive electrophiles (Lehr, et al. 1978). For many of the PAH, certain "bioactivated" metabolites are formed having the capability for covalent interaction with cellular constituents (i.e., RNA, DNA, proteins) and ultimately leading to tumor formation (see Effects section).

The obligatory involvement of metabolic activation for the expression of PAH-induced carcinogenesis has prompted the investigation of PAH metabolism in numerous animal models and human tissues. From these studies has emerged an understanding of the gen-

eral mechanisms involved in PAH biotransformation. It is now known that PAH are metabolized by the cytochrome P-450-dependent microsomal mixed-function oxidase (MFO) system, often designated aryl hydrocarbon hydroxylase (Conney, 1967; Marquardt, 1976; Sims, 1976; Gelboin, et al. 1972). The activity of this enzyme system is readily inducible by exposure to chemicals and is found in most mammalian tissues, although predominantly in the liver (Bast, et al. 1976; Chuang, et al. 1977; Andrews, et al. 1976; Cohn, et al. 1977; Wiebel, et al. 1975; Grundin, et al. 1973; Zampaglione and Mannerling, 1973). The MFO system is involved in the metabolism of endogenous substrates (e.g., steroids) and the detoxification of many xenobiotics. Paradoxically, however, the MFO system also catalyzes the formation of reactive epoxide metabolites from certain PAH, possibly leading to carcinogenesis in experimental mammals (Sims and Grover, 1974; Selkirk, et al. 1971, 1975a; Sims, 1976; Thakker, et al. 1977; Levin, et al. 1977a; Lehr, et al. 1978; see Effects section). A second microsomal enzyme, epoxide hydrase, converts epoxide metabolites of PAH to vicinal glycols, a process which may also play a critical role in carcinogenic bioactivation. Figure 1 presents a schematic representation of the various enzymes involved in activation and detoxification pathways for BaP. At present this also appears to be representative of the general mechanism for PAH metabolism.

A discussion of the metabolism of PAH in mammalian species, including man, is best approached by examining in detail the chemical fate of the most representative and well-studied compound in the PAH class, namely BaP. The metabolism of BaP has been exten-

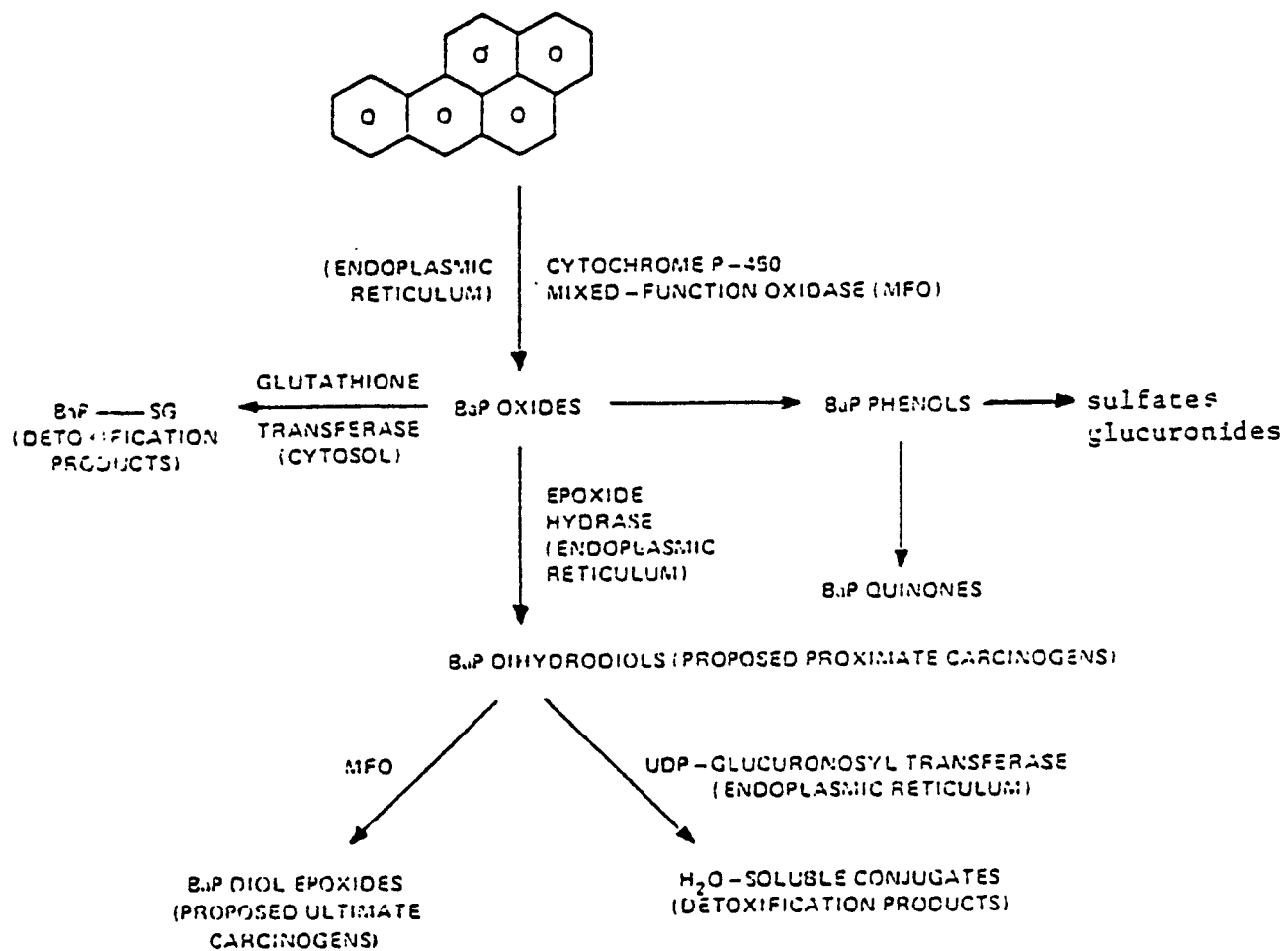


FIGURE 1
 Enzymatic Pathways Involved in the Activation
 and Detoxification of BaP

sively studied in rodents, and the results of these investigations provide useful data which can be directly compared to and contrasted with the results of more limited studies employing human cells and tissues. Therefore, separate discussions are based upon the available experimental evidence regarding PAH metabolism in general, and BaP metabolism in particular, in both animals and man.

Metabolism of PAH in Animals: The metabolites of PAH produced by microsomal enzymes in mammals can arbitrarily be divided into two groups on the basis of solubility. In one group are those metabolites which can be extracted from an aqueous incubation mixture by an organic solvent. This group consists of ring-hydroxylated products such as phenols and dihydrodiols (Selkirk, et al. 1974; Sims, 1970), and hydroxymethyl derivatives of those PAH having aliphatic side chains, such as 7,12-dimethylbenz(a)anthracene (Boylard and Sims, 1967) and 3-methylcholanthrene (Stoming, et al. 1977; Thakker, et al. 1978). In addition to the hydroxylated metabolites are quinones, produced both enzymatically by microsomes and non-enzymatically by air oxidation of phenols. Labile metabolic intermediates such as epoxides can also be found in this fraction (Selkirk, et al. 1971, 1975a,b; Sims and Grover, 1974; Yang, et al. 1978).

In the second group of PAH metabolites are the water soluble products remaining after extraction with an organic solvent. Many of these derivatives are formed by reaction (conjugation) of hydroxylated PAH metabolites with glutathione, glucuronic acid, and sulfate. Enzyme systems involved in the formation of water-soluble metabolites include glutathione S-transferase, UDP-glucuronosyl

transferase, and sulfotransferases (Bend, et al. 1976; Jerina and Daly, 1974; Sims and Grover, 1974). Conjugation reactions are believed to represent detoxification mechanisms only, although this group of derivatives has not been rigorously studied.

The metabolite profile of BaP which has recently been expanded and clarified by the use of high pressure liquid chromatography is depicted in Figure 2. This composite diagram shows three groups of positional isomers, three dihydrodiols, three quinones, and several phenols. The major BaP metabolites found in microsomal incubations are 3-hydroxy-BaP, 1-hydroxy-BaP, 7-hydroxy-BaP, and 9-hydroxy-BaP. The BaP-4,5-epoxide has been isolated and identified as a precursor of the BaP-4,5-dihydrodiol. Other studies indicate that epoxides are the precursors of the 7,8-dihydrodiol and 9,10-dihydrodiol as well. Considerable evidence has recently become available which implicates the diol epoxide, $7\beta,8\gamma$ -dihydro-7,8-dihydroxybenzo(a)pyrene-9,10 α -oxide, as an ultimate carcinogen derived from BaP (Jerina, et al. 1976; Kapitulnik, et al. 1977b, 1978a,b; Levin, et al. 1976a,b; Yang, et al. 1978).

Since the resonance properties of PAH make ring openings difficult, enzymatic attack in the microsomes functions to open double bonds and add an oxygen-containing moiety, such as a hydroxyl group, to give it more solubility in aqueous media (e.g., urine) and thus facilitate removal from the body. In the formation of metabolic intermediates by oxidation mechanisms, relatively stable PAH are converted to unstable products (i.e., epoxides). Thus, nucleophilic attack of this reactive intermediate, through the formation of a transient carbonium ion, would be greatly enhanced.

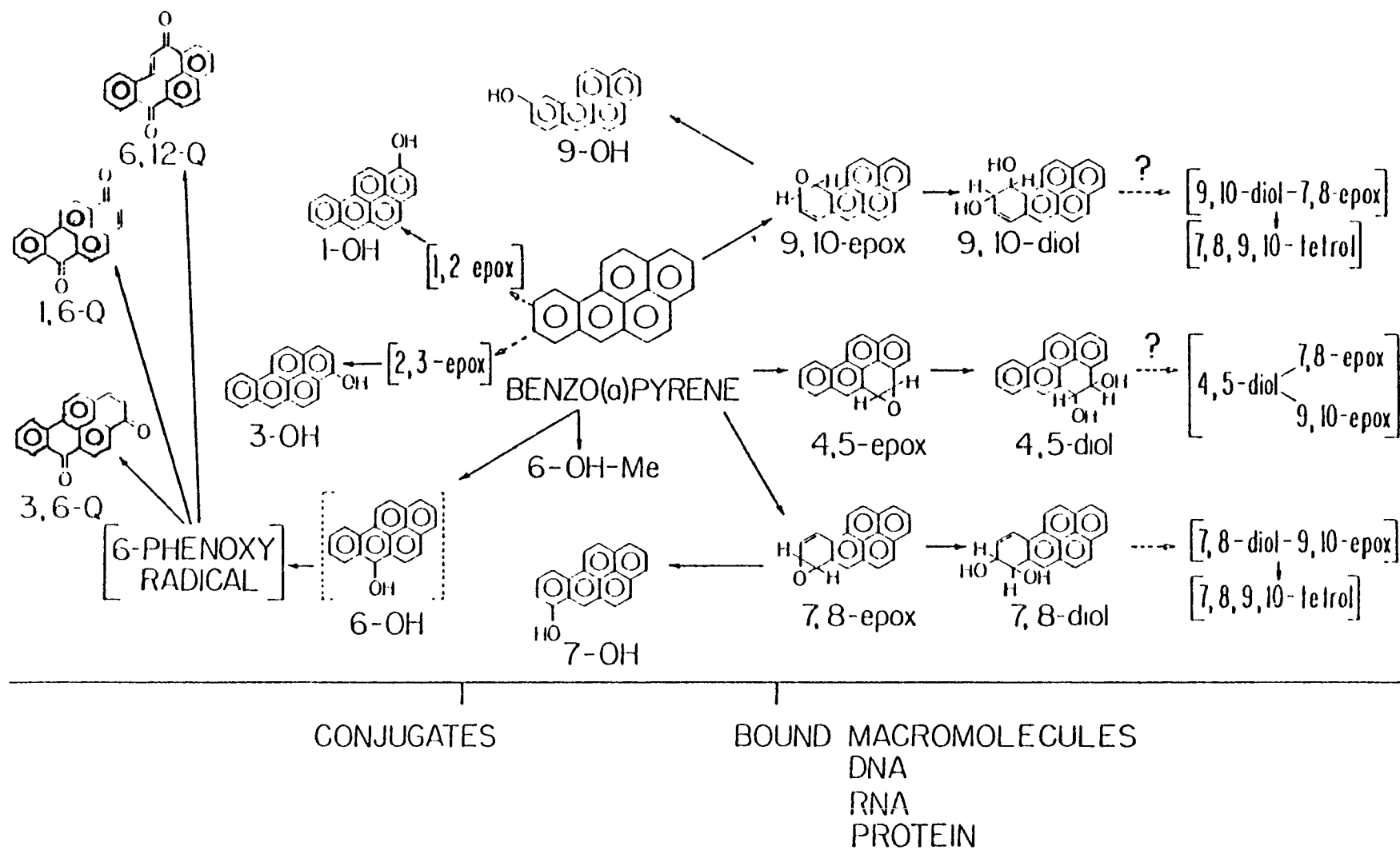


FIGURE 2

Metabolites of Benzo(a)pyrene

Arylations of this type are common to many classes of carcinogenic chemicals. Therefore, the microsomal cytochrome P-450-containing MFO system and epoxide hydrase play a critical role in both the metabolic activation and detoxification of many PAH.

Various forms of liver microsomal cytochrome P-450 can be isolated from animals treated with different enzyme inducers (Wiebel, et al. 1973; Nebert and Felton, 1976; Conney, et al. 1977a,b; Lu, et al. 1978). Moreover, the metabolite profiles of BaP can be qualitatively altered depending on the type of cytochrome P-450 present in the incubation mixture (Lu, et al. 1976; Wiebel, et al. 1975). This observation has important implications in considering the carcinogenic action of certain PAH toward tissues from animals of different species, sex, age, nutritional status, and exposure to enzyme-inducing chemicals. Limited evidence is also available indicating that multiple forms of epoxide hydrase exist among animal species, which may also influence the pattern of PAH metabolism with respect to carcinogenic bioactivation (Lu, et al. 1978).

Comparative Metabolism of PAH in Animals and Man: An important consideration in evaluating the health hazards of PAH is whether metabolism in various animal tissues and species is indicative of the pattern of PAH metabolism in the target organs of humans. Moreover, it is essential to determine whether differences occur in the metabolism of PAH by: (a) different tissues in the same animal; and (b) different animals of the same species.

Numerous studies have shown that the qualitative and quantitative differences exist in the metabolism of BaP by different tissues and animal species (Sims, 1976; Leber, et al. 1976; Wang, et

al. 1976; Pelkonen, 1976; Kimura, et al. 1977; Selkirk, et al. 1976). For the most part, however, interspecies extrapolations of qualitative patterns of PAH metabolism appears to be a valid practice. On the other hand, marked differences in patterns of tissue-specific metabolism may prevent the reliable extrapolation of data from hepatic to extrahepatic (i.e., target organ) tissues. These differences may also exist in human tissues (Conney, et al. 1976).

Freudenthal, et al. (1978) recently examined the metabolism of BaP by lung microsomes isolated from the rat, Rhesus monkey, and man. Metabolite profiles obtained by high pressure liquid chromatography are shown in Figure 3. Their results confirmed previous observations regarding the existence of considerable individual variation in BaP metabolism among samples from the same species. In addition, it was apparent that qualitative and quantitative interspecies variation also existed (Table 16). Nevertheless, the qualitative differences between man and the other animal species were by no means dramatic, and probably do not compromise the validity of extrapolations concerning PAH metabolism.

The metabolite pattern obtained for BaP in human lymphocytes is similar to that obtained with human liver microsomes (Selkirk, et al. 1975b), and human lymphocytes (Booth, et al. 1974). However, in cultured human bronchus (24 hrs) and pulmonary alveolar macrophages an absence of phenols (i.e., 3-hydroxy-BaP) and paucity of quinones were observed (Autrup, et al. 1978). Instead, a relative abundance of the trans-7,8-diol metabolite of BaP was demonstrated. This result is noteworthy in light of the possibility that the 7,8-diol is capable of further oxidative metabolism to an ulti-

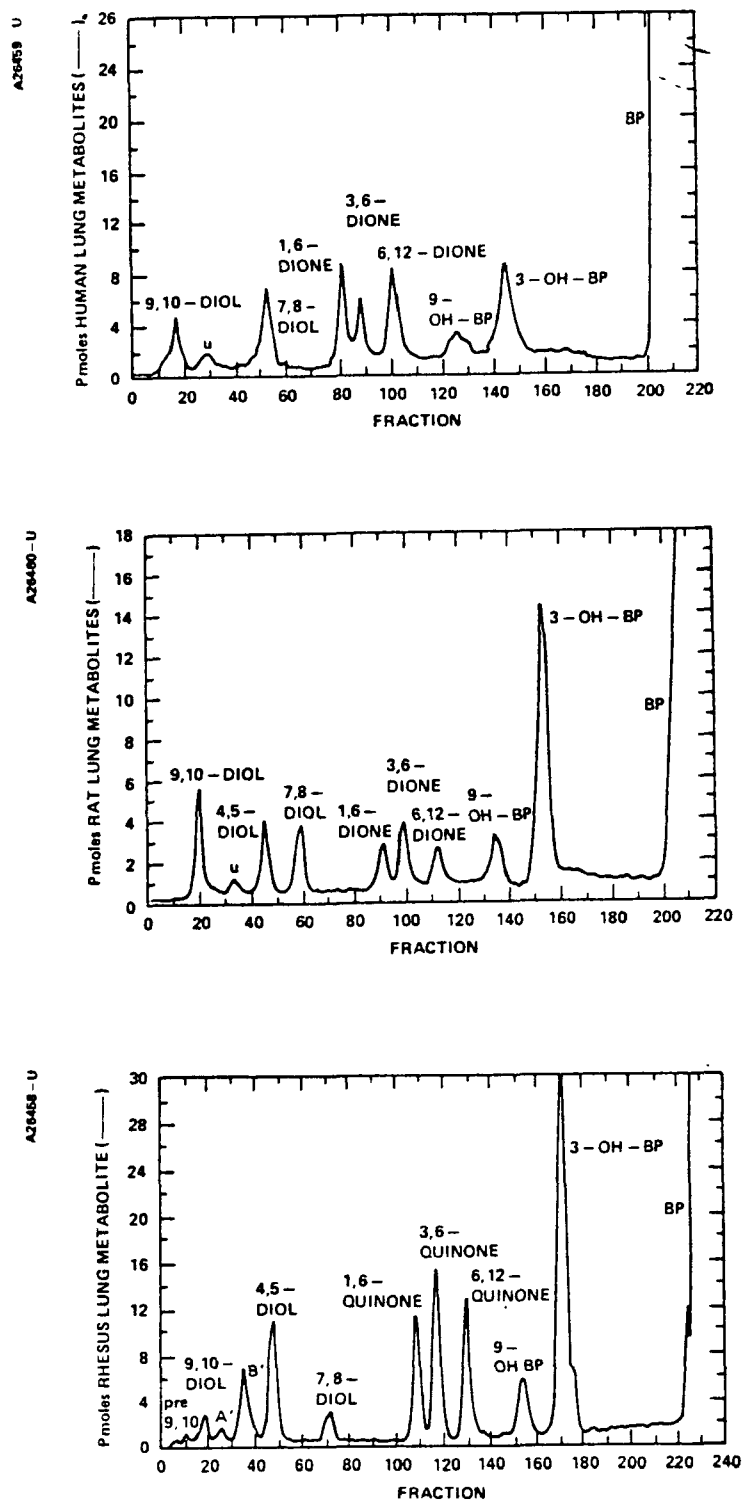


FIGURE 3

Comparative Metabolism of Benzo(a)pyrene by Lung Microsomes from Rat, Rhesus Monkey, and Human

Source: Freudenthal, et al. 1978

TABLE 16

Metabolite Percentages of BP Metabolites from Rat, Rhesus, and Human Lung Microsomal Assays*

Metabolite	Metabolite percentages (pmoles metabolite/pmoles total metabolites x 100)									
	Rat ^a			Rhesus ^b			Man ^{b,c}			
	1	2	3	1	2	3	1	2	3	4
Pre-9,10					3.0	5.3				
9,10-Diol	9.7	6.3	9.6	2.7	4.6	2.6		7.1	6.0	
A				1.5						
U (B)	4.4	3.4	2.9	6.9		7.7	8.9	3.9	7.5	30.0
4,5-Diol	8.3	9.2	8.3	9.0	9.2	7.7	4.1			
7,8-Diol	5.3	5.2	8.0	4.2	8.6	5.1		15.0	13.3	9.9
1,6-Dione	4.4	7.5	8.3	11.4	14.8	12.8	24.9	11.6	12.6	4.4
3,6-Dione	7.8	8.0	9.9	14.5	16.0	20.5	22.5	13.8	19.2	8.5
6,12-Dione	6.8	8.6	8.6	11.8	8.0	15.3	22.5	18.3	27.4	15.7
9-OH	12.6	11.5	3.5	7.3			5.7	6.2		8.5
3-OH	40.8	40.2	41.1	30.8	35.9	23.1	11.4	24.0	13.9	22.9

^aLungs of 5 rats pooled for each group.^bDeterminations made on lung samples from separate individuals.^cWith the exception of subject 4, activity determinations were made using microsomes which had been stored at -84°C.^dThe structural characteristics of unknown, U, may differ between species.

*Source: Freudenthal, et al. 1978

mate carcinogenic form of BaP. It is not known whether a longer incubation period would have changed the pattern of metabolite formation.

Excretion

There is no direct information available concerning the excretion of PAH in man. Limited inferences can be drawn from animal studies with PAH, however.

As long ago as 1936, researchers recognized that various PAH were excreted primarily through the hepatobiliary system and the feces (Peacock, 1936; Chalmers and Kirby, 1940). However, the rate of disappearance of various PAH from the body, and the principal routes of excretion are influenced both by structure of the parent compound and the route of administration (Heidelberger and Weiss, 1951; Aitio, 1974a,b). Moreover, the rate of disappearance of a PAH [i.e., benzo(a)pyrene] from body tissues can be markedly stimulated by prior treatment with inducers of microsomal enzymes [e.g., benzo(a)pyrene, 7,12-dimethylbenz(a)anthracene, 3-methylcholanthrene, chrysene] (Schlede, et al. 1970a,b). Likewise, it has been shown that inhibitors of microsomal enzyme activity, such as parathion and paraoxon, can decrease the rate of BaP metabolism in certain animal tissues (Weber, et al. 1974). From the available evidence concerning excretion of PAH in animals, it is apparent that extensive bioaccumulation is not likely to occur.

EFFECTS

Acute, Subacute, and Chronic Toxicity

The potential for PAH to induce malignant transformation dominates the consideration given to health hazards resulting from exposure. This is because overt signs of toxicity are usually not produced by many PAH until the dose is sufficient to produce a high tumor incidence. Although the emphasis on carcinogenicity is certainly justified when dealing with public health issues concerning PAH, one must recognize that nonneoplastic lesions may also result from environmental and occupational contact. Such effects can be seen with low doses of carcinogenic PAH and with those compounds which possess no tumorigenic activity. Numerous PAH have demonstrated carcinogenic activity when administered to laboratory animals by various routes of administration. However, since many PAH have not been tested for biologic activity it is not possible to list all carcinogenic PAH. A summary of those PAH which are thus far known to be carcinogenic in animals is provided in surveys published by the U.S. Public Health Service (Hartwell and Shubik, 1951; Shubik and Hartwell, 1957, 1969; Tracor Jitco, Inc., 1974, 1976). Since only a small percentage of PAH compounds are known to be carcinogenic, measurements of total PAH (i.e., the sum of all multiple fused-ring hydrocarbons having no heteroatoms) cannot be equated with carcinogenic risk. When the term "total PAH" is used it is necessary in each case to specify the compounds being considered.

As long ago as 1937, investigators knew that carcinogenic PAH, produced systemic toxicity as manifested by an inhibition of body

growth in rats and mice (Haddow, et al. 1937). Tissue damage resulting from the administration of various PAH to experimental animals is often widespread and severe, although selective organ destruction may occur (e.g., adrenal necrosis, lymphoid tissue damage). Few investigators, however, have attempted to ascertain the molecular mechanism of PAH-induced cytotoxicity. Nevertheless, current opinion favors the concept that normally proliferating tissues (intestinal epithelium, bone marrow, lymphoid organs, testis) are preferred targets for PAH, and this susceptibility may be due to a specific attack on DNA of cells in the S phase of the mitotic cycle (Philips, et al. 1973). Additional factors which may have an important bearing on the adverse effects resulting from PAH exposure are primary and secondary alterations in enzyme activity and immunologic competence. Moreover, these toxicant-induced changes may play an important role in the eventual induction of neoplasia.

Target organs for the toxic action of PAH are diverse, due partly to extensive distribution in the body and also to the selective attack by these chemicals on proliferating cells. Damage to the hematopoietic and lymphoid systems in experimental animals is a particularly common observation. Yasuhira (1964) described severe degeneration of the thymus and marked reduction in weight of the spleen and mesenteric lymph nodes of CF_1 Swiss and C57BL mice given a single intraperitoneal injection of MCA (0.3 to 1.0 mg) between 12 hours and 9 days after birth. Degeneration of young cells in the bone marrow and retardation of thyroid gland development were also noted. Newborn mice were highly susceptible to the toxic effects of MCA, with many animals dying from acute or chronic wasting dis-

ease following treatment. Among surviving CF₁ mice, numerous thymomas eventually developed; none were evident, however, in C57BL mice despite serious thymic damage.

DMBA is well-known for its effects on the bone marrow and lymphoid tissues. With single feedings (112 or 133 mg/kg body weight) to female Sprague-Dawley rats, age 50 days, DMBA induced pancytopenia by causing a severe depression of hematopoietic and lymphoid precursors (Cawein and Sydnor, 1968). Maturation arrest occurred at the proerythroblast levels; no injury to the stem cells or the formed elements in the peripheral blood was evident. The fact that only the more rapidly proliferating hematopoietic elements were vulnerable to attack by DMBA led the authors to suggest that inhibition of DNA replication may be involved in the toxicologic response.

Philips and coworkers (1973) provided strong support for the argument that DMBA-induced cytotoxicity is mediated via an interaction with DNA. Female Sprague-Dawley rats receiving 300 mg/kg of body weight of DMBA orally and male rats receiving an intravenous injection of 50 mg/kg of body weight of DMBA displayed injury to the intestinal epithelium, extreme atrophy of the hematopoietic elements, shrinkage of lymphoid organs, agranulocytosis, lymphopenia, and progressive anemia. Mortality among rats receiving DMBA by gastric intubation (females) was about 65 percent. In rats given 50 mg/kg of body weight of DMBA intravenously, incorporation of ¹⁴C-labeled thymidine into DNA of small and large intestine, spleen, bone marrow, cervical lymph nodes, thymus, and testis was significantly inhibited. This inhibition was as high as 90 percent

in several organs at six hours which indicated a strong inhibition of DNA synthesis. Consequently, the authors postulated that DNA in S phase cells is particularly susceptible to DMBA attack. This phenomenon probably applies for other carcinogenic PAH as well.

Another lesion, characteristic of that produced by X-rays, is the severe testicular damage induced by DMBA in rats (Ford and Huggins, 1963). Single intravenous injections of DMBA (0.5 to 2.0 mg) given to adolescent (25 days of age) rats caused transient degenerative changes in the testis which were most evident 38 to 40 days after treatment. Essentially the same effects were produced in adult rats, age 60 days, given DMBA orally (20 mg) and intravenously (5 mg). Lesions of the testes were highly specific and involved destruction of spermatogonia and resting spermatocytes, both of which are the only testicular cells actively synthesizing DNA. Neither the remaining germinal cells nor the interstitial cells were damaged by DMBA. Surprisingly, no testicular damage was produced by single feedings of BaP (100 mg), MCA (105 mg), or 2-acetaminophenanthrene (40 mg).

It is well known that the application of carcinogenic polycyclic hydrocarbons to mouse skin leads to the destruction of sebaceous glands, hyperplasia, hyperkeratosis, and even ulceration (Bock, 1964). Sebaceous glands are the skin structures most sensitive to polycyclic hydrocarbons, and assay methods for detection of carcinogens have been based on this effect. Although a good correlation can be obtained between carcinogenic activity and sebaceous gland suppression for many PAH [e.g., MCA, DMBA, BaP, DBA, benz(a)-anthracene], such an effect is neither necessary nor sufficient for

carcinogenesis. However, workers exposed to PAH-containing materials such as coal tar, mineral oil, and petroleum waxes are known to show chronic dermatitis, hyperkeratoses, etc. (Hueper, 1963; NAS, 1972), though the possible significance of these skin disorders to human cancer is not known.

In female animals, ovotoxicity has been reported to result from the administration of PAH. DMBA was shown to cause the destruction of small oocytes and to reduce the numbers of growing and large oocytes after oral administration to mice (Kraup, 1970). More recently a report was published that destruction of primordial oocytes in mice by injection of MCA was correlated with the genetic capability for PAH-induced increases in ovarian aryl hydrocarbon hydroxylase activity (Mattison and Thorgeirsson, 1977). Thus, the ovarian metabolism of PAH and ovotoxicity are apparently linked.

A toxic reaction which is apparently unique to DMBA is the selective destruction of the adrenal cortex and induction of adrenal apoplexy in rats (Boyland, et al. 1965). Adrenal apoplexy, increased adrenal gland weight, and increased adrenal hemoglobin content were induced in female Sprague-Dawley rats by a single intragastric dose of 30 mg DMBA. The same amount of adrenal damage could be produced by a 5 mg dose of the principal DMBA oxidative metabolite, 7-hydroxymethyl-12-methylbenz(a)anthracene. Other DMBA metabolites produced no adrenal damage, thus indicating that a specific reactive intermediate may be responsible for this phenomenon.

Repeated injections of benz(a)anthracene derivatives to mice and rats have produced gross changes in the lymphoid tissues.

Hoch-Ligeti (1941) administered DBA, benz(a)anthracene, and anthracene to mice in weekly subcutaneous injections for 40 weeks. Analysis of lymph glands removed at weekly intervals showed an increase of reticulum (stem) cells and an accumulation of iron in all treated groups of animals. Lymphoid cells were reduced and lymph sinuses dilated in all groups, although these effects were more common in mice receiving DBA. The weights of the spleens in mice treated with DBA were significantly reduced in comparison to controls and those animals receiving benz(a)anthracene or anthracene.

A more detailed study on the effects of repeated administration of DBA on lymph nodes of male rats was reported in 1944 (Lasnitzki and Woodhouse, 1944). Subcutaneous injections given five times weekly for several weeks caused normal lymph nodes to undergo hemolymphatic changes. These changes are characterized by the presence of extravascular red blood cells in the lymph spaces and the presence of large pigmented cells. These changes were not observed by Hoch-Ligeti (1941) in mice, but could be produced in rats by BaP and MCA in addition to DBA. The noncarcinogen, anthracene, on the other hand, did not produce as dramatic a change in the lymph nodes of rats.

In light of the concern over PAH-induced neoplasms of the respiratory tract, an understanding of early pathological alterations and preneoplastic lesions in this tissue has particular significance.

In a study conducted by Reznik-Schuller and Mohr (1974), BaP-induced damage to the bronchial epithelium of Syrian golden hamsters was examined in detail using tissue sections. Animals were

treated intratracheally with 0.63 mg BaP (total dose) dispersed in a solution of saline, dodecylsulfate, Tris-HCl, and EDTA once weekly for life. Animals were serially sacrificed at weekly intervals following the first month of treatment, and sections of the bronchi were examined microscopically. In the first animals sacrificed, minimal focal cell proliferation in the area of the basement membrane was evident in the bronchial epithelium. By 7 weeks, cytoplasmic vacuolization of both goblet and ciliated cells had occurred. Epithelial and basal cell proliferation continued for several weeks and led to the formation of three-to four-layered hyperplastic regions by the 11th week. Epithelial cells began to penetrate through the basement membrane by the 12th week, and within two or more weeks the bronchial epithelium began to grow continuously into the surrounding lung tissues. Microscopic bronchogenic adenomata had developed by the 20th week. These tumors consisted primarily of ciliated cells and goblet cells, with only a few basal cells present. The apparently small amount of basal cell proliferation may have been the reason why squamous metaplasia was not observed by the time the experiment had ended after 21 weeks. Squamous metaplasia and keratinization were found in the trachea, but not in the bronchi, after 21 weeks of treatment. Although these investigators found no increase in the number of alveolar macrophages, others have reported numerous alveolar macrophage responses in BaP-treated hamsters as well as focal areas of accumulated macrophages containing a yellow pigment having unknown biological significance (Henry, et al. 1973; Saffiotti, et al. 1968).

Epithelial proliferation and cell hyperplasia in the absence of necrosis and/or marked inflammation is a common observation in the tracheobronchial mucosa of animals directly exposed to carcinogenic PAH. This phenomenon was shown with repeated exposures of DMBA, BaP, and dibenzo(a,i)pyrene in hamsters (Reznik-Schuller and Mohr, 1974; Saffiotti, et al. 1968; Stenbäck and Sellakumar, 1974a,b).

Numerous investigators have demonstrated that carcinogenic PAH can produce an immunosuppressive effect. This effect was first observed by Malmgren, et al. (1952) using high doses of MCA and DB(a,h)A in mice. Subsequent studies established that single carcinogenic doses of MCA, DMBA, and BaP caused a prolonged depression of the immune response to sheep red blood cells (Stjernswärd, 1966, 1969). Noncarcinogenic hydrocarbons such as benzo(e)pyrene and anthracene reportedly had no immunosuppressive activity. In a recent review on immunosuppression and chemical carcinogenesis, substantial evidence was presented to indicate that the degree of immunosuppression was correlated with carcinogenic potency for PAH (Baldwin, 1973). Both cell-mediated and humoral immune reactions are affected by PAH.

Synergism and/or Antagonism

It is well-known that the development of PAH-induced tumors in epithelial and non-epithelial tissues can be altered by: (1) components in the diet, (2) inducers and inhibitors of microsomal enzymes, (3) other co-administered noncarcinogenic or weakly carcinogenic chemicals, and (4) the vehicle used to deliver a carcinogenic PAH to experimental animals. These factors tend to compli-

cate the extrapolation of animal dose-response data to human situations. On the other hand, these observations in animals reinforce the belief that similar interactions occur with regard to the action of PAH in humans.

Early studies conducted by Falk and coworkers (1964) indicated that the carcinogenic effect of BaP on subcutaneous injection in mice could be markedly inhibited by the simultaneous administration of various noncarcinogenic PAH. Similarly, they showed that neutral extracts of particulate air pollutant fractions also produced inhibitory effects on BaP-induced tumorigenesis. However, when Pfeiffer (1973, 1977) conducted similar studies with BaP and DBA in the presence of 10 noncarcinogenic PAH, no inhibitory effect was evident. Moreover, an increased tumor yield resulted from injection of mixtures containing increasing amounts of the components. This effect, however, was less dramatic than if BaP were administered alone, and it paralleled the dose-response curve for DBA acting singly.

Many studies on cocarcinogenesis have been concerned with the identification of tumor accelerating substances present in cigarette smoke. These compounds are generally tested for cocarcinogenic activity by repeated application to mouse skin together with low doses of BaP. A positive response would be obtained in cases where the tumor yield of the combination exceeds that produced by either agent alone at the same doses. Van Duuren and coworkers (1973, 1976) established that a pronounced cocarcinogenic effect could be obtained with catechol and the noncarcinogens, pyrene, BeP, and benzo(g,h,i)perylene. Doses of 12, 15, 21, and 2,000 μg

of these compounds, respectively, were applied three times a week for 52 weeks to female ICR/Ha Swiss mice. Each animal also received 5 ug of BaP in 0.1 ml acetone with each dose of test substance. Although phenol has been regarded as a tumor-promotor in the two-stage carcinogenesis system (Van Duuren, 1976), this compound has a slight inhibitory effect on BaP carcinogenesis when administered in combination. These results, therefore, indicated that tumor-promoters and cocarcinogens may not have the same mode of action, and that the two terms should not be used interchangeably. Other PAH (e.g., fluoranthene, pyrene, pyrogallol) also possess cocarcinogenic activity but have no tumor-promoting activity (Van Duuren, 1976). Additional studies by Schmeltz, et al. (1978) established that most of the naphthalenes found in cigarette smoke (250 ug, three times a week) have an inhibitory effect on skin tumorigenesis as induced by BaP (3 ug, three times a week). On the other hand, several of the alkylnaphthalenes tested (dimethyl-, trimethyl-, tetramethyl-) enhanced the carcinogenic activity of BaP on mouse skin.

Numerous investigators have shown that antioxidants are effective inhibitors of PAH-induced tumor development. This action has been demonstrated with selenium (Shamberger, 1970; Shamberger and Rudolph, 1966; Riley, 1969), dl- α -tocopherol (vitamin E) (Shamberger, 1970; Shamberger and Rudolph, 1966), and ascorbic acid (Shamberger, 1972) in mice treated with DMBA and croton oil. The carcinogenic action of MCA has been reduced by tocopherol-rich diets in rats and mice (Jaffe, 1946; Haber and Wissler, 1962). The antioxidant food additives butylated hydroxytoluene (BHT), ethoxy-

quin, and butylated hydroxyanisole (BHA) have inhibited lung, breast, and gastric tumor formation induced in rats and mice by various carcinogens in the diet (Wattenberg, 1972, 1973; Wattenberg, et al. 1976). The sulfur-containing antioxidants (disulfuram, dimethyldithiocarbamate, and benzyl thiocyanate) inhibited DMBA-induced mammary cancer in rats when they were added to the diet; in the mouse, disulfuram prevented the formation of forestomach tumors induced by BaP in the diet, but had no effect on BaP-induced pulmonary adenoma (Wattenberg, 1974). The agricultural herbicide, maleic hydrazide, and its precursor, maleic anhydride, can inhibit the initiating activity of DMBA in the mouse skin two-stage carcinogenesis system (Akin, 1976).

Rahimtula and coworkers (1977) examined the abilities of several antioxidants to affect BaP hydroxylation by rat liver microsomal mixed-function oxidases. Their results indicated that antioxidants can markedly inhibit BaP hydroxylation by an apparently direct action on microsomal oxidation mechanisms. Furthermore, all of the antioxidants tested reduced the bacterial mutagenicity of BaP in the presence of rat liver microsomes and cofactors. The authors suggested that antioxidants may exert their protective effect in vivo by inhibiting the formation of carcinogenic intermediates from PAH. This conclusion, however, seems to conflict with data indicating that inducers of increased BaP hydroxylase activity can also inhibit tumor formation (Wattenberg and Leong, 1970). However, flavones are also inhibitors of BaP metabolism in vitro, thereby indicating that their specific effects depend upon how and where they are used. These investigators found that sev-

eral synthetic and naturally occurring flavones when incorporated in the diet (3 to 5 mg/g) or applied to the skin caused a profound increase in BaP hydroxylase activity in the small intestine and skin, respectively. In addition, pulmonary adenoma formation resulting from oral administration of BaP was totally prevented, and skin tumors initiated by BaP application to mice were significantly reduced (> 50 percent) by treatment with the synthetic flavone, β -naphthoflavone. Pulmonary tumor formation was also reduced 50 percent by incorporation of the naturally occurring flavone, quercetin pentamethyl ether, into the diet. Sullivan and coworkers (1978) recently demonstrated that BHA, BHT, phenothiazine, phenothiazine methosulfate, and ethoxyquin all can reduce the quantitative yield of BaP metabolites in incubations with rat liver microsomes. The possibility that only specific components of the drug metabolizing enzyme system may be induced by antioxidants has not been fully explored.

In addition to flavones, other naturally occurring compounds have exhibited protective effects against PAH-induced tumor formation. Retinoids have clearly been shown to play a role in reducing carcinogen-induced tumors (Nettesheim, et al. 1975; Cone and Nettesheim, 1973; Chu and Malmgren, 1965; Smith, et al. 1975). Nettesheim and Williams (1976) recently examined whether inadequate vitamin A consumption may predispose individuals to carcinogenesis, or whether increased vitamin A intake exerts a protective effect against neoplasia. They found that a diet deficient in vitamin A increased the formation of MCA-induced metaplastic lung nodules in female Fisher 344 rats, even though adequate amounts of the vitamin

were stored in the liver. On the other hand, moderate amounts of the vitamin A added to the diet markedly reduced the development of MCA-induced lesions of the lung. High doses of the vitamin given intragastrically provided no additional protection, however.

Further studies on naturally occurring antineoplastic compounds were recently reported by Wattenberg (1977). Benzyl isothiocyanate and phenethyl isothiocyanate, both found in cruciferous plants such as cabbage, brussel sprouts, cauliflower, etc., inhibited DMBA-induced mammary cancer in Sprague-Dawley rats. When added to the diet together with DMBA, these compounds inhibited the development of forestomach tumors and pulmonary adenomas in female ICR/Ha mice. Similar anticarcinogenic actions were obtained when BaP was incorporated into the diet. These results lead to interesting speculation regarding the role and importance of diet in human susceptibility to environmental carcinogens. In cases where dietary constituents can alter the metabolism of xenobiotics such as PAH, then the anticarcinogenic effect may result from an alteration of steady state levels of activated versus detoxified metabolites.

Studies have shown that not only can specific substances in the diet affect the response to carcinogens, but decreased protein content in the diet may also decrease the activation of carcinogens (Czygan, et al. 1974). The feeding of protein-deficient diets to male mice decreased liver weights and reduced cytochrome P-450 content in the total liver. Diets deficient in both protein and choline produced even further reductions in liver weight and cytochrome P-450 content. Liver microsomes isolated from these animals

displayed a decreased ability to activate dimethylnitrosamine to a mutagen in the Ames Salmonella test system, which paralleled the reduction in cytochrome P-450 content produced by the diet. Conversely, the inactivation of the direct-acting (ultimate) carcinogen, N-methyl-N'-nitro-N-nitrosoguanidine, was reduced in liver microsomes from mice receiving a protein-deficient diet.

In humans fed charcoal-broiled beef, the metabolism of the drug phenacetin was enhanced; in pregnant rats a similar diet stimulated the activity of AHH in the placenta and liver (Conney, et al. 1977a,b). Further studies showed that high-protein diets enhanced the metabolism of antipyrone and theophylline in man, while a high-carbohydrate diet depressed the rate of metabolism of these drugs. Additional agents in man's environment which inhibit AHH activity include certain organophosphate pesticides, piperonyl butoxide, carbon tetrachloride, ozone, carbon monoxide, nickel carbonyl, and nickel, tin, cobalt, and other metals (Conney, et al. 1977a,b).

Teratogenicity

No information is available concerning the possible teratogenic effects of PAH in man. Furthermore, only limited data are available regarding the teratogenic effects of PAH in experimental animals.

BaP had little effect on fertility or the developing embryo in several mammalian and nonmammalian species (Rigdon and Rennels, 1964; Rigdon and Neal, 1965). On the other hand, DMBA and its hydroxymethyl derivatives apparently are teratogenic in the rat (Currie, et al. 1970; Bird, et al. 1970). However, DMBA is not generally regarded as an environmental contaminant.

Mutagenicity

No reliable way presently exists to measure whether PAH may induce heritable mutations in humans. However, the concept that carcinogenesis is an expression of an alteration in the genetic material of a cell (i.e., somatic mutation) implies that a formal relationship exists between mutagenesis and carcinogenesis (Nery, 1976; Miller, 1978). The results obtained with several in vitro mutagenesis test systems, particularly the Ames Salmonella typhimurium assay, support the belief that most carcinogenic chemicals are mutagenic as well. For PAH, the Ames assay has been very effective in detecting those parent structures and their biotransformation products which possess carcinogenic activity (McCann, et al. 1975; Teranishi, et al. 1975; McCann and Ames, 1976; Sugimura, et al. 1976; Wislocki, et al. 1976b; Wood, et al. 1976a; Tokiwa, et al. 1977; Brookes, 1977). The Ames assay, however, may not be 100 percent effective in detecting all PAH carcinogens, nor does the assay provide a reliable quantitative measure of carcinogenic potency or tumor-initiating activity.

The availability of Salmonella typhimurium strains for the detection of chemically induced mutations and the use of microsomal preparation to provide metabolic activation has made possible an investigation of the mechanisms of PAH-induced mutagenesis. In particular, an exhaustive survey of the mutagenicity of all the possible oxidative metabolites of BaP has helped to confirm the belief that diol epoxide intermediates are the ultimate mutagens/carcinogens derived from PAH (Jerina, et al. 1976; Wood, et al. 1976a,b; Wislocki, et al. 1976a,b; Thakker, et al. 1976; Levin, et al. 1977a,b). These results are summarized in Table 17.

TABLE 17

Comparison of Inherent Mutagenic Activity of Thirty BaP Derivatives in Salmonella typhimurium TA98 and in Chinese Hamster V79 Cells^{a,b}

Compound ^c	Relative % Activity	
	Strain TA98	V79
Diol epoxide-1	100	40
Diol epoxide-2	35	100
H ₄ 9,10-epoxide	95	40
H ₄ 7,8-epoxide	10	0.2
BaP 4,5-oxide	20	1
BaP 7,8-oxide	1	<0.1
BaP 9,10-oxide	1	<0.1
BaP 11,12-oxide	0.5	1
6-HOBaP	5	0.3
12-HOBaP	1.5	<0.1
1-HOBaP	0.5	0.1
3-HOBaP	0.5	<0.1
2-, 4-, 5-, 7-, 8-, 9-, 10-, 11-HOBaP	<0.1	<0.1
BaP 1,6-, 3,6-, 6,12-, 4,5-, 11,12-quinone	<0.1	<0.1
BaP 4,5-, 7,8-, 9,10-, 11,12-dihydrodiol	<0.1	<0.1
BaP	<0.1	<0.1

^aSource: Jerina, et al. 1976

^bThe relative percent mutagenic activities are approximations since the data were compiled from several separate studies conducted at different times. In some experiments, BaP 7,8-dihydrodiol was 0.1 to 0.4% as active as diol epoxide-2 in V79 cells.

^cAbbreviations used: BaP, benzo(a)pyrene; 1-HOBaP, 1-hydroxybenzo(a)pyrene; 2- to 12-HOBaP, other BaP phenols; BaP 1,6-quinone, benzo(a)pyrene 1,6-quinone; BaP 3,6-quinone, BaP 4,5-quinone, BaP 6,12-quinone, and BaP 11,12-quinone, other BaP quinones; BaP 4,5-dihydrodiol, trans-4,5-dihydroxy-4,5-dihydrobenzo(a)pyrene; BaP 7,8-, 9,10- and 11,12-dihydrodiol, other dihydrodiols of BaP; BaP 4,5-oxide, benzo(a)pyrene 4,5-oxide; BaP 7,8-, 9,10-, and 11,12-oxide, other BaP oxides; diol epoxide-1(+)-7 β ,8 α -dihydroxy-9 β ,10 β -epoxy-7,8,9,10-tetrahydro BaP; diol epoxide-2, (+)-7 β ,8 α -dihydroxy-9 α ,10 α -epoxy-7,8,9,10-tetrahydro BaP; H₄ 9,10-epoxide, 9,10-epoxy-7,8,9,10-tetrahydro BaP; H₄ 7,8-epoxide, 7,9-epoxy-7,8,9,10-tetrahydro BaP.

Further examination of the mutagenic activity of PAH and their derivatives has been conducted in mammalian cell culture systems. These systems operate with concentrations of test compounds which are lower than those used in the Ames assay. This work has been conducted primarily with Chinese hamster cell lines, either V79 cells derived from male lung tissue or CHO cells derived from the ovary. These cells, however, do not possess a microsomal enzyme system and thus co-cultivation with lethally irradiated rodent embryo cells which retain metabolic activity is required for testing of PAH.

Using this system, Huberman and Sachs (1974, 1976) demonstrated that a number of carcinogenic PAH produced forward mutations involving three genetic markers: (1) ouabain resistance; (2) temperature sensitivity; and (3) 8-azaguanine resistance. Noncarcinogenic PAH such as BeP, phenanthrene, and pyrene were not mutagenic. In addition, studies by Huberman indicated that a correlation could be shown between the degree of carcinogenicity and the frequency of induced somatic mutations (Huberman, et al. 1977). The demonstration that covalent binding of carcinogenic PAH with DNA of V79 cells was the same as occurs in vivo further strengthened the argument that genetic interaction (i.e., somatic mutation or gene depression) may be involved in tumor formation (Newbold, et al. 1977).

The use of Chinese hamster V79 cells to test the mutagenicity of BaP metabolites has contributed significantly to an understanding of the molecular action of PAH (Huberman, et al. 1976a,b, 1977; Malaveille, et al. 1975; Newbold and Brooks, 1976; Jerina, et al.

1976). Comparison of the mutagenic activities of the optically pure (+) and (-)-enantiomers of BaP 7,8-dihydrodiol revealed that, in the presence of a metabolic activating system, the (-)trans, 7,8-dihydrodiol was the most active mutagen (Huberman, et al. 1977). These results are consistent with the fact that the (-)trans 7,8-dihydrodiol is the only BaP enantiomer by rat liver microsomes (Yang, et al. 1977), and that it is highly carcinogenic to newborn mice (Kapitulnik, et al. 1978a,b). Because the (-)trans 7,8-dihydrodiol had no mutagenic activity in the absence of enzymes required for PAH metabolism, it was apparent that the BaP 7,8-diol-9,10-epoxide, which is derived from this intermediate, is an ultimate mutagen/carcinogen. Studies by Wood, et al. (1977a) on the mutagenicity to V79 cells by the four optically pure enantiomers of the BaP 7,8-diol-9-10-epoxides supported this belief. None of the triols and tetrols which are derived from BaP diol epoxides were mutagenic to V79 cells, and thus represent probable detoxification products (Huberman, et al. 1977).

The current belief that neoplastic transformation may arise from a chemically induced somatic mutation was made even more convincing by the recent studies of Huberman and coworkers (1976b). They demonstrated for the first time that BaP and BaP 7,8-dihydrodiol can induce both neoplastic transformation and mutagenesis (ouabain resistance) in the same culture of normal diploid hamster embryo cells. The concentrations for transformation and mutagenesis were the same, and showed a dose-response effect in both transformation and ouabain resistance for BaP 7,8-dihydrodiol.

In further adaptation of the cell-mediated mutagenesis system, V79 cells are metabolically activated by rat liver homogenates containing microsomes and cofactors (Krahn and Heidelberger, 1977). The mutagenic activity of BaP, MCA, DMBA, and benz(a)anthracene in this system showed a limited correlation with their respective carcinogenic potencies. It should be noted, however, that the selection of a particular activating system (i.e., microsomes v. feeder cells) may have a significant influence on the test results.

The analysis of chromosomal aberrations and sister chromatid exchanges (SCEs) is often recommended as a screening technique for potential mutagens and carcinogens. Several investigators have examined the effects of PAH on the chromosomes of mammalian cells. Early studies indicated that variations in chromosome number and structure may accompany tumors induced by BaP, MCA, and DMBA in the rat, mouse, and hamster (Kato, et al. 1975). However, in cultured human leukocytes exposed to DMBA, chromosome damage was not the same as that produced in hamster cells. Although it is argued that chromosome changes in PAH-induced tumors are all specific (Levan and Levan, 1975; Ahlstrom, 1974), others (Popescu, et al. 1976; Nery, 1976) claim that detectable chromosome changes are not specific for the carcinogenic agent nor are they a prerequisite for neoplastic growth. Moreover, an increased rate of SCEs can be produced by BaP in cultured human lymphocytes (Rüdiger, et al. 1976; Schönwald, et al. 1977) but this increase is not correlated with different rates of BaP metabolism (Rüdiger, et al. 1976), a surprising result in light of the known importance of metabolic activation for BaP mutagenicity. BaP-induced SCEs rates did not differ

between lymphocytes taken from normal humans and those from patients with lung cancer (Schönwald, et al. 1977). In recent studies with cultured Chinese hamster cells exposed to DMBA, BaP, and MCA, none of the chemicals produced chromosome breaks and only DMBA could successfully induce SCEs (Abe and Sasaki, 1977). Although it cannot be denied that PAH cause chromosome damage, it is not clear whether this effect may represent an epigenetic phenomenon which is merely secondary to mutagenesis and neoplastic transformation. Furthermore, in cases where a chemically induced mutation is "silent" (i.e., neutral amino acid substitution), there is no reason to believe that detectable chromosome damage should occur.

In recent comparisons of three cytogenetic tests, (1) induction of chromosome aberrations, (2) induction of micronuclei, and (3) in vivo induction of sister chromatid exchanges, the last test proved to be the most sensitive with carcinogenic polycyclic hydrocarbons (Bayer, 1978). Since positive results were also obtained with phenanthrene, the usefulness of sister chromatid exchange as a screening technique for carcinogen detection is limited. BaP was positive in the sister chromatid exchange test, weakly active in the chromosome aberration test, and negative in the micro-nucleus test. On the other hand, DMBA was clearly positive in all three tests. The conclusion was that cytological tests do not provide reliable correlations with all carcinogens tested and thus cannot be used alone in mutagenicity/carcinogenicity evaluations.

Damage to the genome resulting from chemical insult can theoretically also be detected by examining DNA repair (Stich and Laishes, 1973). The suggestion that DNA repair is applicable as a

screening procedure for evaluating potential chemical mutagens is based on the assumption that the level of DNA repair synthesis in a cell reflects the extent of DNA damage produced by a chemical. Indeed, unscheduled incorporation of ³H-thymidine into nuclear DNA of normal human cells exposed to epoxides of benz(a)anthracene and MCA has been observed (Stich and Laishes, 1973). However, since a metabolic activation system was not present in this system, the parent hydrocarbons showed no activity. More recent studies confirmed that K-region epoxides of BaP, DMBA, and DBahA caused DNA damage in human skin fibroblasts which was repaired with the same system used for repairing lesions induced by ultraviolet radiation (Maher, et al. 1977). As would be expected, the parent hydrocarbons exerted no effect. More important, results were obtained which indicated that the DNA repair process itself does not induce mutations, but rather that mutagenesis occurs before the DNA lesion can be excised.

DNA repair synthesis in human fibroblasts (Regan, et al. 1978; Stich, et al. 1975,1976; San and Stich, 1975), rat liver cells (Williams, 1976), and Chinese hamster V79 cells (Swenberg, et al. 1976) has been successfully used for the detection of chemical carcinogens, including numerous PAH. However, the percentage of carcinogens giving positive results for DNA repair is considerably less than in the cell transformation or microbial mutagenesis assays. Nevertheless, tests with human skin fibroblasts showed that DNA repair synthesis results from exposure to BaP 7,8-diol-9,10-epoxides, whereas BaP 4,5-, 9,10-, and 11,12-oxides did not produce DNA damage which was repairable by the ultraviolet excision

repair system (Regan, et al. 1978). These results support the concept that diol epoxide metabolites of PAH are ultimate mutagens.

Tumors induced in vivo by PAH are commonly associated with chromosome abnormalities in the neoplastic cells. In particular, sarcomas induced by DMBA, MCA, and BaP in the rat display karyotype variations which were reportedly nonrandom and distinctly different from sarcomas induced by Rous sarcoma virus (Levan and Levan, 1975; Mitelman, et al. 1972). The chromosome patterns of DMBA-induced sarcomas were found to be identical with those observed in rat leukemias (Mitelman and Levan, 1972) and in primary carcinomas of the auricular skin (Ahlstrom, 1974) induced by DMBA.

Considerable evidence is also available to indicate that chromosome alterations in PAH-induced tumors in vivo are not consistent either in frequency or in pattern. DMBA-induced tumors (fibrosarcoma, squamous carcinoma, lymphosarcoma) of the uterine cervix in ICR mice revealed various karyotypic profiles (Joneja and Coulson, 1973; Joneja, et al. 1971). These tumors displayed diploid, aneuploid, tetraploid, and octaploid chromosome constitutions. Tumors induced in mice with MCA and dibenzo(a,i)pyrene also showed a wide variation in chromosome constitution (Biedler, et al. 1961; Hellstrom, 1959). Mice treated with 30 μ g DMBA, a dose sufficient to produce a 100 percent incidence of thymic lymphomas, did not reveal an excess of chromosome abnormalities in bone marrow or thymus (Ottonen and Ball, 1973). Even at higher doses (60 μ g DMBA), the incidence of abnormal chromosomes did not significantly differ from controls. Subcutaneous tumors in Syrian hamsters induced by single injections of BaP (0.1 μ g) or DMBA (0.1 mg), and cultured cell

populations derived from these tumors, failed to reveal common karyotypic changes (DiPaolo, et al. 1971a,b). Tumor cells had aneuploid, diploid, and hypotetraploid chromosome constitutions; further karyotype rearrangements occurred with subsequent growth in vitro.

In humans, the presence of the "Philadelphia" chromosome in myeloid leukemia appears to be the only example of a human chromosome abnormality which is tumor-specific (Nowell and Hungerford, 1960). In PAH-induced experimental tumors, lymphatic leukemia in mice produced by DMBA also displays consistent chromosome abnormalities (Joneja and Coulson, 1973). Beyond this common feature, convincing data have not been presented to indicate that somatic cells exposed to PAH may suffer characteristic or reproducible damage to the genome. Instead, random karyotypic mutants of transformed cells are thought to be selected in response to growth pressures in the host environment (e.g., tissue necrosis, infection, anoxia, lack of nutrition) (Joneja and Coulson, 1973).

Carcinogenicity

Animal data: Numerous polycyclic aromatic compounds are distinctive in their ability to produce tumors in skin and most epithelial tissues of practically all species tested. Malignancies are often induced by acute exposures to microgram quantities of PAH. Latency periods can be short (four to eight weeks) and the tumors produced may resemble human carcinomas. Carcinogenesis studies involving PAH have historically involved primarily effects on the skin or lungs. In addition, subcutaneous or intramuscular injections are frequently employed to produce sarcomas at the in-

jection site. Ingestion has not been a preferred route of administration for the bioassay of PAH (Santodonato, et al. 1980).

Concern over potential human cancer risk posed by PAH present in the atmosphere stems from studies demonstrating that crude extracts of airborne particulate matter can be carcinogenic to animals (Hoffmann and Wynder, 1976; Wynder and Hoffman, 1965; Hueper, et al. 1962; Kotin, et al. 1954). Fractions soluble in benzene or benzene-methanol produced tumors in mice by skin painting or subcutaneous injection. Both the aromatic and oxygenated neutral subfractions were active as complete carcinogens, and indicated the presence of numerous carcinogenic materials, including non-PAH. Since the carcinogenicity of the total organic particulates and aromatic neutral subfractions could be explained only partly by the presence of BaP, its usefulness as a measure of carcinogenic risk from air pollution may be limited.

From investigations in which polycyclic carcinogens were painted on the skin of mice has emerged the two-stage theory of skin carcinogenesis (Berneblum, 1941; Van Duuren, 1969, 1976). The first stage, initiation, results from the ability of a carcinogen to effect a permanent change within a cell or cell population following a single application. The measure of carcinogenic potency is often regarded as the capacity for tumor initiation. However, some weak or inactive complete carcinogens can be active as tumor initiators (e.g., dibenz(a,c)anthracene, 1-methylchrysene, benz(a)anthracene). The second stage, promotion, is a prolonged process which does not necessarily require the presence of a carcinogen, but nevertheless a chemical stimulus must be supplied (e.g.,

by croton oil). A complete carcinogen is one which, if applied in sufficient quantity, can supply both initiating and promoting stimuli (e.g., DMBA, BaP). The formation of skin tumors by polycyclic hydrocarbons may also be influenced by inhibitors and accelerators (cocarcinogens), thus complicating the interpretation of experimental data.

The tumorigenic effects of PAH when applied to the skin of animals have been known for decades. Iball (1939) collected the results of a series of experiments to arrive at a method for comparing the carcinogenic potencies of various polycyclic aromatic chemicals. His results, presented in Table 18, express tumorigenic potency in mouse skin as the ratio of percent tumor incidence to the average latency period. This expression, commonly referred to as the Iball index, is still used as a means of comparing the relative activity of carcinogens. An important data compilation on agents tested for carcinogenicity has more recently been published by the U.S. Public Health Service (Publication No. 149) which lists the results of tests on hundreds of chemicals in numerous animals including rodent, avian, and amphibian species.

Experimental models for respiratory carcinogenesis have major limitations in that the delivery of carcinogens to the tracheobronchial tree in measured amounts and their adequate retention at the target tissue are poorly controlled. Therefore, the conduct of dose-response studies on lung tumor induction has been seriously hampered. Moreover, the possible relevance of the two-stage theory of carcinogenesis to lung cancer has not been clearly established. Many of the bioassay data on PAH-induced lung cancer have been de-

TABLE 18

Carcinogenic Compounds in Descending Order of Potency*

Compound	Number of Mice Alive when First Tumor	Number of Tumors	Percentage of Tumors (A)	Papilloma	Epithelioma	Average Latent Period(B)	Index (A/B x 100)
1. 7,12-Dimethylbenz(a)anthracene	20	13	65	6	7	43	151
2. 3-Methylcholanthrene (a)	18	18	100	1	17	99	101
3. 3-Methylcholanthrene (b)	8	5	62.5	0	5	151	41
4. 3-Methylcholanthrene (a and b added together)	26	23	88.5	1	22	109	80
5. Benzo(a)pyrene (from pitch)	10	10	100	2	8	127	79
6. Benzo(a)pyrene (synthetic)	9	7	78	2	5	109	72
7. Benzo(a)pyrene (5 and 6 added together)	19	17	89.5	4	13	119	75
8. Cholanthrene	49	28	57	5	23	112	51
9. 5,6-Cyclopenteno-benz(a)anthracene	14	13	93	1	12	194	48
10. 2-Methyl-benzo(c)phenanthrene	16	12	75	5	7	155	48
11. 10-Methyl-benz(a)anthracene	18	12	66.5	2	10	147	45
12. 5,6-Dimethyl-benz(a)anthracene	19	16	84	0	16	220	38
13. 6-Isopropyl-benz(a)anthracene	15	11	73.5	1	10	204	36
14. Dibenz(c,g)carbazole	19	9	47.5	4	5	143	33
15. Dibenz(a,h)pyrene	17	10	59	0	10	205	29
16. 5-Methyl-benz(a)anthracene	8	7	87.5	2	5	317	28
17. 5-Ethyl-benz(a)anthracene	9	7	77.5	2	5	285	27
18. Dibenz(a,h)anthracene	65	41	63	8	33	239	26
19. Benzo(c)phenanthrene	18	12	67	5	7	387	17
20. Dibenz(a,g)carbazole	9	4	44.5	1	3	263	17
21. 5-n-Propyl-benz(a)anthracene	20	6	30	3	3	192	16
22. Dibenz(c,h)acridine	28	11	39.3	2	9	357	11
23. 3-Methyl-dibenz(a,h)anthracene	25	7	28	1	6	325	9
24. Dibenz(a,h)acridine	25	6	24	2	4	350	7
Totals		305		60	245		

*Source: Iball, 1939

rived from animal model systems employing various modes of administration (inhalation, intratracheal instillation, intravenous injection), and the use of carrier particles (e.g., ferric oxide) for the delivery of the carcinogen to the bronchial epithelium. Thus, the results obtained from these studies cannot always be directly compared. The most commonly employed method for the study of PAH-induced lung cancer involves intratracheal instillation of test material in the Syrian golden hamster.

Following the identification of the first carcinogenic hydrocarbon from soot (BaP) an intensive effort was mounted to isolate the various active components of carcinogenic tars (IARC, 1973). From the earliest studies conducted, the realization emerged that carcinogenic PAH are structurally derived from the simple angular phenanthrene nucleus (Arcos and Argus, 1974). However, unsubstituted PAH with less than four condensed rings that have been tested have not shown tumorigenic activity. Furthermore, of the six possible arrangements with four benzene rings, only two of these compounds are active: benzo(c)phenanthrene and benz(a)anthracene. The unsubstituted penta- and hexacyclic aromatic hydrocarbons are clearly the most potent of the series. These include BaP, DBaA, dibenzo(a,h)pyrene, dibenzo(a,i)pyrene, dibenzo(a,l)pyrene, dibenzo(a,e)pyrene, benzo(b)fluoranthene, and benzo(j)fluoranthene. Somewhat less potent as carcinogens are the dibenzanthracenes and dibenzophenanthrenes. Only a few heptacyclic hydrocarbons show carcinogenic activity. These include phenanthro(2',3':3,4')pyrene, peropyrene, and dibenzo(h,rst)pentaphene. Beyond seven unsubstituted aromatic rings, there are very few known carcinogenic

hydrocarbons. However, many physico-chemical and enzymatic parameters must be dealt with in respect to carcinogenic PAH. Factors such as solubility and intracellular localization to achieve metabolic activation are likely to be important determinants of the true carcinogenicity of a particular PAH.

Among the unsubstituted polycyclic hydrocarbons containing a nonaromatic ring, a number of active carcinogens are known. The most prominent examples of this type of compound are cholanthrene, 11,12-ace-benz(a)anthracene, 8,9-cyclopentanobenz(a)anthracene, 6,7-ace-benz(a)anthracene, acenaphthanthracene, 1,2,5,6-tetrahydrobenzo(j)cyclopent(f,g)aceanthrylene, and "angular" steranthrene. All of these compounds retain an intact conjugated phenanthrene segment.

The addition of alkyl substituents in certain positions in the ring system of a fully aromatic hydrocarbon will often confer carcinogenic activity or dramatically enhance existing carcinogenic potency. In this regard, Arcos and Argus (1974) noted that monomethyl substitution of benz(a)anthracene can lead to strong carcinogenicity in mice, with potency depending on the position of substitution in the decreasing order, 7 > 6 > 8 ~ 12 > 9. A further enhancement of carcinogenic activity is produced by appropriate dimethyl substitution of benz(a)anthracene. Active compounds are produced by 6,8-dimethyl-, 8,9-dimethyl-, 8,12-dimethyl-, 7,8-dimethyl-, and 7,12-dimethyl-substitution. The latter compound is among the most potent PAH carcinogens known, although it has not been shown as a product of fossil fuel pyrolysis. Methyl substitution in the angular ring of benz(a)anthracene, however, tends to

deactivate the molecule, although 4,5-dimethylbenz(a)anthracene may be an exception. Carcinogenic trimethyl- and tetramethylbenz(a)anthracenes are known, and their relative potencies are comparable to the parent 7,12-DMBA. In general, free radical synthesis of polycyclic hydrocarbons by pyrolysis does not favor alkyl side chain formation.

Alkyl substitution of partially aromatic condensed ring systems may also add considerable carcinogenic activity. The best example of this type of activation is 3-methylcholanthrene, a highly potent carcinogen.

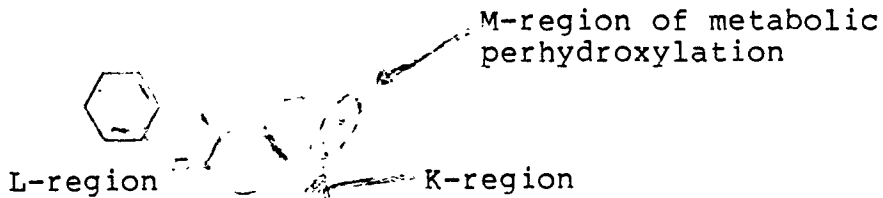
With alkyl substituents longer than methyl, carcinogenicity tends to decrease, possibly due to a decrease in transport through cell membranes. However, different positions in the benz(a)anthracene molecule will vary with respect to the effect of n-alkyl substitution on carcinogenicity. Benz(a)anthracene is especially sensitive to decreased carcinogenicity caused by the addition of bulky substituents at the 7-position, and is indicative of a once widely-held view for most polycyclics that high reactivity of the meso-phenanthrenic region (now called the "K-region") was a critical determinant for carcinogenicity. Current studies show that the K-region is not involved in critical binding to DNA.

Partial hydrogenation of the polycyclic aromatic skeleton can generally be expected to decrease carcinogenic potency. This was shown with various hydrogenated derivatives of BaP, benz(a)anthracene, and MCA. On the other hand, the carcinogenicity of DBahA, dibenzo(a,i)pyrene, and dibenzo(a,h)pyrene is not significantly altered by meso-hydrogenation. This may be due to the fact that

extensive resonance capability is preserved. Moreover, 5,6-dihydro-DBahA actually displayed a fourfold increase in carcinogenicity in comparison to the parent hydrocarbon (Arcos and Argus, 1974), possibly due to the hydrophilicity and ease of intracellular transport of its dihydrodiol derivative.

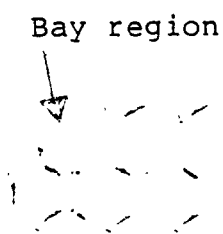
For many years, investigators have sought a common molecular feature among PAH carcinogens which would serve to explain their biological activity. The "electronic theory of carcinogenesis" has relied upon an analysis of the influence of electron density at specific molecular regions to explain unique reactivity with cellular constituents. A basic assumption arising from the work of the Pullmans and others (Pullman and Pullman, 1955) was that a meso-phenanthrenic region ("K'-region") of high π -electron density and with a propensity for addition reactions was a critical structural feature for polycyclic carcinogens. In expanding this hypothesis, further biological significance was attributed to the concomitant presence of a rather unreactive meso-anthracenic region ("L-region") for high carcinogenicity. In addition, a region of comparatively low reactivity which characteristically undergoes metabolic perhydroxylation (corresponding to the 3,4-positions of benz(a)anthracene) has been designated the M-region. According to the theory, only binding of the K'-region to critical cellular sites would cause tumor formation; protein binding at the L-region causes no tumorigenic effect, while inactivation is produced by metabolic

perhydroxylation in the M'-region. The three regions of reactivity are readily distinguished in the benz(a)anthracene skeleton:

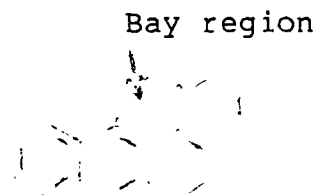


The electronic K'-L theory of carcinogenic reactivity has encountered numerous inconsistencies, primarily because these relationships were derived from physico-chemical properties of the parent hydrocarbon and gave no consideration to the biological effects of activated metabolites.

Advances in recent years have focused attention on the potential reactivity of diol epoxide metabolites of PAH, and their ease of conversion to triol carbonium ions. Under the assumption that diol epoxides, which are more readily converted to carbonium ions, will be better alkylating agents to produce carcinogenesis and mutagenesis, the "bay region" theory has been proposed (Lehr, et al. 1978; Wood, et al. 1977b). Examples of a "bay region" in a polycyclic hydrocarbon are the regions between the 10 and 11 positions of BaP and the 1 and 12 positions of benz(a)anthracene:



Benzo(a)pyrene



Benz(a)anthracene

The theory predicts that diol epoxides in which the oxirane oxygen forms part of a "bay region" (e.g., BaP 7,8-diol-9,10-epoxide) will be more reactive and hence more carcinogenic than diol epoxides in which the oxirane oxygen is not situated in a "bay region."

Experimentally, the "bay region" diol epoxides of benz(a)anthracene, BaP, and chrysene were more mutagenic in vitro and/or tumorigenic than other diol epoxide metabolites, their precursor dihydrodiols, the parent hydrocarbons, or other oxidative metabolites. Moreover, quantum mechanical calculations were in accord with the concept that reactivity at the "bay region" is highest for all the diol epoxides derived from polycyclic hydrocarbons.

The bay region concept has received enough confirmation to lead to suggestions that an analysis of theoretical reactivity in this manner may be useful in screening PAH as potential carcinogens (Smith, et al. 1978). Among several indices of theoretical reactivity examined, the presence of a bay region for a series of PAH displayed a high degree of correlation with positive carcinogenic activity (Table 19).

The carcinogenic activity of BaP has been studied extensively in various animal model systems. In recent years, research on BaP has been expanded to include an examination of the tumorigenic activity of various BaP metabolites. These efforts were directed at the objective of identifying a BaP derivative which acts as the principal ultimate carcinogen resulting from metabolic activation (Levin, et al. 1976a,b, 1977a,b; Slaga, et al. 1976, 1977; Kapitulnik, et al. 1976a,b; Wislocki, et al. 1977; Conney, et al. 1977a,b).

Studies on the activity of BaP and its derivatives as complete carcinogens on mouse skin (Table 20) and as tumor initiators (Table 21) revealed that marked differences in tumorigenic potency exist. The apparent lack of activity for the BaP 7,8-diol-9,10-epoxides,

TABLE 19

Reactivity Indices for Polycyclic Hydrocarbons*

Compound	K-region?	L-region?	Bay region	Carcinogenicity Index	
				Arcos and Argus (1974)	Jerina, et al. (1972)
Naphthalene	-	-	-	0	-
Anthracene	-	+	-	0	-
Tetracene	-	+	-	0	-
Pentacene	-	+	-	0	-
Hexacene	-	+	-		?
BA	+	+	+	5	+
Benzo(a) tetracene	+	+	+		-
Phenanthrene	+	-	+	0	-
Benzo(c) phenanthrene	+	-	+	4	+
Chrysene	+	-	+	3	+
Benzo(b) chrysene	+	+	+		-
Picene	+	-	+	0	-
Triphenylene	-	-	+	0	-
Benzo(g) chrysene	+	-	+	17	++
Dibenz(a,c) anthracene	-	+	+	3	+
Dibenz(a,j) anthracene	+	+	+	4	+
Dibenz(a,h) anthracene	+	+	+	26	++
Naphtho(2,3-b) pyrene	+	+	a	27	++
Benzo(a) pyrene	+	-	+	73	++++
Benzo(e) pyrene	+	-	+	2	+ ^b
Dibenzo(a,l) pyrene	+	-	+	33	++ ^b
Dibenzo(a,i) pyrene	+	-	+	74	++++
Dibenzo(a,e) pyrene	+	-	+	50	+++
Dibenzo(a,h) pyrene	+	-	+	70	++++
Tribenzo(a,e,i) pyrene	-	-	+	16	++

*Source: Smith, et al. 1978

^aThis compound does not strictly possess a bay region but does contain a "pseudo" bay region.

^bJerina, et al. (1972) have assigned this as ++++.

TABLE 20

Skin Tumors in Mice Treated with Benzo(a)pyrene and Derivatives

Treatment ^a	Total No. Animals	Dose, μ moles	Mice with Tumors, %	Total No. Skin Tumors ^b	Reference
BaP	25	0.4	100	32	Wislocki, et al. 1977
BaP	30	0.4	100	34	Wislocki, et al. 1977
BaP	26	0.4	92	34	Albert, et al. 1978
BaP	30	0.15	100	40	Levin, et al. 1976a,b
BaP	27	0.1	96	28	Wislocki, et al. 1977
BaP	30	0.1	38	13	Levin, et al. 1977a,b
BaP	30	0.1	50	15	Levin, et al. 1977a,b
BaP	30	0.1	91	24	Levin, et al. 1977a,b
BaP	30	0.05	59	20	Levin, et al. 1977a,b
BaP	30	0.025	7	2	Levin, et al. 1977a,b
BaP	30	0.02	4	1	Levin, et al. 1977a,b
BaP	30	0.02	0	0	Levin, et al. 1977a,b
1-HOBaP	25	0.4	0	0	Wislocki, et al. 1977
2-HOBaP	29	0.4	100	37	Wislocki, et al. 1977
3-HOBaP	29	0.4	0	0	Wislocki, et al. 1977
4-HOBaP ^C	26	0.4	0	0	Albert, et al. 1978
5-HOBaP ^C	26	0.4	0	0	Albert, et al. 1978
6-HOBaP ^C	28	0.4	0	0	Albert, et al. 1978
7-HOBaP ^C	30	0.4	0	0	Albert, et al. 1978
8-HOBaP ^C	27	0.4	0	0	Albert, et al. 1978
9-HOBaP ^C	26	0.4	0	0	Albert, et al. 1978
10-HOBaP ^C	28	0.4	0	0	Albert, et al. 1978
11-HOBaP	28	0.4	14	4	Wislocki, et al. 1977
12-HOBaP	23	0.4	0	0	Wislocki, et al. 1977
BaP 4,5-oxide	30-39	0.4	4	1	Levin, et al. 1976a
BaP 4,5-oxide	30-39	0.1	6	2	Levin, et al. 1976a
BaP 7,8-oxide	30-39	0.4	94	37	Levin, et al. 1976a
BaP 7,8-oxide	30	0.3	53	16	Levin, et al. 1976a
BaP 7,8-oxide	30	0.15	18	5	Levin, et al. 1976a
BaP 7,8-oxide	30-39	0.1	9	3	Levin, et al. 1976a
BaP 9,10-oxide	30-39	0.4	0	0	Levin, et al. 1976a
BaP 11,12-oxide	28	0.4	0	0	Wislocki, et al. 1977
BaP 11,12-oxide	17	0.1	0	0	Wislocki, et al. 1977

TABLE 20 (cont.)

Treatment ^a	Total No. Animals	Dose, μ moles	Mice with Tumors, %	Total No. Skin Tumors ^b	Reference
BaP 7,8-dihydro-diol	30	0.3	100	42	Levin, et al. 1976b
BaP 7,8-dihydro-diol	30	0.15	100	40	Levin, et al. 1976b
BaP 7,8-dihydro-diol	30	0.1	92	28	Levin, et al. 1976a
BaP 7,8-dihydro-diol	30	0.05	76	24	Levin, et al. 1976a
BaP 7,8-dihydro-diol	30	0.025	7	2	Levin, et al. 1976a
(+)-7 β ,8 α -Di-hydroxy-9 β ,10 β -epoxy-7,8,9,10-tetrahydrobenzo(a)pyrene (diol epoxide 1)	30	0.4	0	0	Levin, et al. 1976a
diol epoxide 1	30	0.1	0	0	Levin, et al. 1976a
diol epoxide 1	30	0.02	0	0	Levin, et al. 1976a
(+)-7 β ,8 α -Di-hydroxy-9 α ,10 α -epoxy-7,8,9,10-tetrahydrobenzo(a)pyrene (diol epoxide 2)	30	0.4	13	3	Levin, et al. 1976a
diol epoxide 2	30	0.1	7	2	Levin, et al. 1976a
diol epoxide 2	30	0.02	0	0	Levin, et al. 1976a

^aFemale C57BL/6J mice were treated with BaP or BaP derivatives (0.02-0.4 μ mole) once every 2 weeks for 60 weeks by topical application to the shaved skin of the back.

^bSkin tumors consisted mostly of squamous cell carcinomas; other skin tumors were fibro-sarcomas, papillomas, and keratocanthomas.

^cMice were treated once every 2 weeks for 56 weeks.

TABLE 21
Summary of the Skin Tumor Initiation Activities of Benzo(a)pyrene and its Metabolites^a

Initiator	No. Mice	Dose, hmoles	Weeks of Promotion	Mice with Tumors, %	Papillomas/ Mouse	Reference
BaP	30	200	23	94	4.8	Slaga, et al. 1976
BaP	30	200	30	92	5.3	Slaga, et al. 1977
BaP	30	200	21	77	2.6	Levin, et al. 1977b
BaP 4,5-epoxide	30	200	23	20	0.2	Slaga, et al. 1976
BaP 7,8-epoxide	29	200	23	81	1.9	Slaga, et al. 1976
BaP 9,10-epoxide	29	200	30	15	0.15	Slaga, et al. 1977
BaP 11,12-epoxide	30	200	30	38	0.45	Slaga, et al. 1977
BaP 7 β ,8 α -diol-9 α ,10 α -epoxide	29	200	30	69	1.5	Slaga, et al. 1977
BaP 7 β ,8 α -diol-9 β ,10 β -epoxide	28	200	30	7	0.07	Slaga, et al. 1977
BaP 7,8-dihydrodiol ^b	29	200	30	86	5.0	Slaga, et al. 1977
(-)-BaP 7,8-dihydrodiol ^b	30	100	21	77	3.8	Levin, et al. 1977b
(+)-BaP 7,8-dihydrodiol ^b	30	100	21	23	0.43	Levin, et al. 1977b

^aFemale CD-1 mice were treated with a single dose of initiator dissolved in acetone, acetone: NH₄OH (1,000;1), or dimethyl sulfoxide:acetone (1;3) and followed 1 week later by twice-weekly applications of 10 μ g of TPA.

^bPromotion was by twice-weekly applications of 16 hmoles of TPA beginning 11 days after treatment with initiator.

despite their exceptional mutagenicity, may be due to poor skin penetration of adult mouse skin because of high chemical reactivity. Indeed, as a carcinogen in newborn mice the (-) enantiomer of BaP, 7,8-dihydrodiol, and the 7,8-diol-9,10-epoxide derived therefrom are far more active than the parent hydrocarbon (Kapitulnik, et al. 1977a,b,c,d, 1978a,b). These studies on the newborn mouse clearly indicate the role of a BaP 7,8-diol-9,10-epoxide as an ultimate carcinogenic metabolite of BaP.

Further dose-response information on the sarcomagenic activity of BaP by subcutaneous injection to rats and mice is summarized in Table 22.

Temporal relationships for the development of BaP-induced skin cancers in mice have been examined by Albert, et al. (1978). Their results showed that increasing weekly doses of BaP caused a shortening of the latency period for carcinoma formation. Furthermore, it was determined that the development of papillomas as a precursor lesion to carcinoma formation occurred only at higher BaP doses (e.g., 32 and 64 µg/week). At the lower dose levels (8 and 16 µg/week), carcinomas appeared de novo without precursor papilloma formation.

The carcinogenicity of BaP by oral intake has not been studied as thoroughly as for other routes of administration. Nevertheless, tumors of various sites result when BaP is administered orally to rodents (Table 23).

With oral, intratracheal, and intravenous routes of administration, BaP is less effective than other PAH (e.g., DMBA, MCA, dibenz(a,h)anthracene) in producing carcinomas. On the other hand,

TABLE 22
Induction of Sarcoma by Benzo(a)pyrene

Species	No. and (Sex)	Total Dose μ moles	Animals with Sarcoma, %	Average Latency, Days	Reference
Rat (Sprague-Dawley)	13 (female)	6.0 ^a	100	101 \pm 2.7	Flesher, et al. 1976
Mouse	14 (male)	7.1 ^b	93	129	Buu-Hoi, 1964
Mouse	16 (female)	7.1 ^b	50	160	Buu-Hoi, 1964
Mouse	9 (?)	15.9 ^c	66.6	112	Gottschalk, 1942
Mouse	10 (?)	5.0 ^c	70	122	Gottschalk, 1942
Mouse	12 (?)	0.5 ^c	66.6	155	Gottschalk, 1942
Mouse	15 (?)	0.002 ^c	0	N.A. ^d	Gottschalk, 1942

^aAdministered as 0.2 μ mole dissolved in 0.1 ml sesame oil by subcutaneous injection on alternate days for 30 doses beginning at 30 days of age.

^bAdministered as three injections of 2.4 μ moles each, given at 1 month intervals.

^cAdministered as a single injection under the skin of the abdomen, dissolved in 0.5 ml of neutral olive oil.

^dNot applicable.

TABLE 23

Carcinogenicity of Benzo(a)pyrene by Oral Administration to Various Mammals*

Compound	Species	Dose	Route of Administration	Effects
BaP	Mouse	0.2 mg in PEG ^a	Intragastric	14 tumors of the forestomach in 5 animals out of 11
	Mouse (age 17-116 days)	50-250 ppm	Dietary (110-197 days)	>70% incidence of stomach tumors at 50-250 ppm for 197 days; no tumors with diets containing up to 30 ppm for 110 days
	Mouse	250 ppm	Dietary	100% stomach tumor incidence when diet was fed for 30 days; 5-7 days of feeding, 30-40%; 2 to 4 days of feeding, 10 percent; 1 day of feeding, 0 percent
	Mouse (age 18-30 days)	250 ppm	Dietary (140 days)	Leukemias, lung adenomas, and stomach tumors produced
	Rat (Sprague-Dawley; age 105 days)	2.5 mg per day	Oral	Papillomas developed in the esophagus and forestomach in 3 out of 40 animals
	Hamster	2-5 mg bi-weekly	Intragastric	5 stomach papillomas in 67 animals treated for 1-5 months; 7 papillomas and 2 carcinomas in 18 animals treated for 6-9 months; 5 papillomas in 8 animals treated for 10-11 months
	Hamster	500 ppm	Dietary (4 days per week for up to 14 mo.)	12 tumors (2 esophagus, 8 forestomach, 2 intestinal) in 8 animals

^aPolyethylene glycol

*Source: IARC, 1973

BaP has remarkable potency for the induction of skin tumors in mice. Therefore, caution must be exercised in considering the carcinogenicity of PAH as a class, and in extrapolating data derived from studies with BaP to the effects of PAH mixtures.

An examination of comparative carcinogenicities within the same tumor model system can provide valuable insight concerning relative risks of various PAH. By single intravenous injection of about 0.25 mg of aqueous dispersions of PAH to mice, a direct comparison of carcinogenic potency was possible (Table 24). In this test system, MCA displayed the greatest lung tumor-forming capability; dibenz(a,h)anthracene followed closely in activity with BaP being considerably less potent.

Intratracheal instillation of PAH to Syrian golden hamsters has been widely utilized for the conduct of studies on pulmonary carcinogenesis (Saffiotti, et al. 1968,1972; Henry, et al. 1975). Several studies are summarized in Table 25 and indicate that: (1) dose-response relationships are clearly evident, and (2) the co-administration of carrier particles such as Fe_2O_3 (i.e., with BaP) can markedly increase tumor incidence, depending on the conditions of the experiment and physical characteristics of the particle. Since environmental exposures to PAH occur in conjunction with particulate material in air, this effect may be particularly relevant to human situation.

In addition to the hamster model system, respiratory tract tumors have been readily induced by PAH in rats and mice. The results of several representative studies are summarized in Table 26.

TABLE 24

Comparative Carcinogenicity of Polycyclic Hydrocarbons and Related Compounds
Measured by Induction of Lung Tumors (LT)^{a, b}

Compound	Dose, μmoles/kg	Mice with LT/ No. of Mice	Mean No. LT/mouse	μMoles/kg for 1 LT Response
3-Methylcholanthrene, 0.1 mg	15	15/15	11	0.9
3-Methylcholanthrene, 0.5 mg	74	6/6	47	
Dibenz(a,h)anthracene	36	10/10	31	1.0
7H-Dibenzo(c,g)carbazole	38	12/12	5.7	6.0
Benzo(a)pyrene	40	10/10	3.7	9.5
Dibenz(a,j)aceanthrylene	33	9/10	2.7	14
Dibenz(a,h)acridine	36	11/12	2.0	18
8-Methylbenzo(c)phenanthrene	42	6/11	0.7	--
7-Methylbenzo(a)pyrene	38	5/10	0.6	--
5-Methoxy-7-propylbenz(a)anthracene	33	1/10	0.1	--
Benz(a)anthracene	44	2/11	0.2	--
Untreated controls	--	4/19	0.2	--

^aSource: Shimkin and Stoner, 1975

^bStrain A mice, 8-12 weeks old, received single intravenous injection of 0.24 mg of methylcholanthrene in aqueous dispersion and were killed 20 weeks later.

TABLE 25

Induction of Respiratory Tract Tumors in Syrian Golden Hamsters by Intratracheal Instillation of PAH

Compound	No. Animals	Total Dose, mg	Respiratory Tumor Incidence, Percent	Reference
BaP	30	3.25 ^a	10	Feron, et al. 1973
BaP	30	6.5 ^a	13	Feron, et al. 1973
BaP	30	13	30	Feron, et al. 1973
BaP	29	26 ^a	86	Feron, et al. 1973
BaP	28	52 ^a	93	Feron, et al. 1973
BaP	48	30 ^b	15	Sellakumar, et al. 1976
BaP and Fe ₂ O ₃	48	30 ^b	71	Sellakumar, et al. 1976
BaP and Fe ₂ O ₃ , coated	49	26.1 ^c	73	Henry, et al. 1975
BaP and Fe ₂ O ₃ , ground	49	27.4 ^c	84	Henry, et al. 1975
BaP and Fe ₂ O ₃ , mixed	43	26.3 ^c	12	Henry, et al. 1975
BaP and gelatin	46	26.4 ^c	17	Henry, et al. 1975
BaP and Fe ₂ O ₃	28 (male), 29 (female)	60 ^d	60.7 (male), 58.6 (female)	Saffioti, et al. 1972
BaP and Fe ₂ O ₃	33 (male), 34 (female)	30 ^d	66.7 (male), 58.8 (female)	Saffioti, et al. 1972
BaP and Fe ₂ O ₃	33 (male), 30 (female)	15 ^d	30.3 (male), 30.0 (female)	Saffioti, et al. 1972
BaP and Fe ₂ O ₃	47 (male), 41 (female)	7.5 ^d	12.8 (male), 9.8 (female)	Saffioti, et al. 1972

C-91

TABLE 25 (cont.)

Compound	No. Animals	Total Dose, mg	Respiratory Tumor Incidence, Percent	Reference
BaP	32 (male)	30 ^e	42.3	Kobayashi, 1975
BaP	28 (female)	30 ^e	57.7	Kobayashi, 1975
DB(a,i)P	48	12 ^f	75	Stenback and Sellakumar, 1974a
DB(a,i)P	48	8.5 ^g	64.6	Stenback and Sellakumar, 1974a
DMBA and Fe ₂ O ₃	46	1.2 ^h	43.5	Stenback and Sellakumar, 1974b
DMBA and Fe ₂ O ₃	28	0.85 ⁱ	46.4	Stenback and Sellakumar, 1974b

^aAnimals treated once weekly for 52 weeks with BaP suspended in 0.9% NaCl solution.

^b3 mg BaP administered once weekly for 10 weeks.

^cAnimals received 30 weekly intratracheal instillations.

^dAnimals received 30 weekly instillations of BaP mixed with equal amounts of Fe₂O₃ and suspended in 0.2 ml saline.

^eAnimals received 30 weekly intratracheal instillations of BaP suspended in 0.9% NaCl.

^fAnimals received 12 weekly intratracheal instillations of 1 mg DB(a,i)P suspended in distilled water.

^gAnimals received 17 weekly intratracheal instillations of 0.5 mg DB(a,i)P suspended in distilled water.

^hAnimals received 100 µg DMBA and 100 µg Fe₂O₃ intratracheally once a week for 12 weeks in saline suspensions.

ⁱAnimals received 50 µg DMBA and 50 µg Fe₂O₃ intratracheally once a week for 17 weeks in saline suspensions.

TABLE 26

Induction of Respiratory Tract Tumors in Rats and Mice

Compound	Organism	No. Animals	Total Dose, mg	Route of Administration	Tumor Incidence, %	Reference
DMBA and Indian ink	Rat (Wistar and random-bred)	34	2.5 ^a	Intratracheal instillation	17.6	Pylev, 1962
DMBA and Indian ink	Rat (Wistar and random-bred)	56	6 ^b	Intratracheal instillation	35.7	Pylev, 1962
DMBA and Indian ink	Rat (Wistar and random-bred)	61	10 ^c	Intratracheal instillation	26.2	Pylev, 1962
DB(a,h)A	Mouse (DBA/2)	14 (male) 13 (female)	236 (male) ^d 179 (female) ^d	Oral	100 (male) ^e 77 (female) ^e	Snell and Stewart, 1962
MCA	Rat (Osborne-Mendel)	100	0.005 ^f	Pulmonary injection	1 ^g	Hirano, et al. 1974
MCA	Rat (Osborne-Mendel)	100	0.05 ^f	Pulmonary injection	13 ^g	Hirano, et al. 1974
MCA	Rat (Osborne-Mendel)	100	0.10 ^f	Pulmonary injection	27 ^g	Hirano, et al. 1974

TABLE 26 (cont.)

Compound	Organism	No. Animals	Total Dose, mg	Route of Administration	Tumor Incidence, %	Reference
MCA	Rat (Osborne-Mendel)	100	0.20 ^f	Pulmonary injection	47 ^g	Hirano, et al. 1974
MCA	Rat (Osborne-Mendel)	100	0.30 ^f	Pulmonary injection	40 ^g	Hirano, et al. 1974
MCA	Rat (Osborne-Mendel)	100	0.40 ^f	Pulmonary injection	51 ^g	Hirano, et al. 1974
MCA	Rat (Osborne-Mendel)	100	0.50 ^f	Pulmonary injection	45 ^g	Hirano, et al. 1974

^aAdministered as a single dose with 0.2 mg of Indian ink in 0.2 ml of a colloid protein solution.

^bAdministered as three 2 mg doses at monthly intervals with 0.2 mg of Indian ink in 0.2 ml of a colloid protein solution.

^cAdministered as five 2 mg doses at monthly intervals with 0.2 mg of Indian ink in 0.2 ml of a colloid protein solution.

^dAdministered as an aqueous-olive oil emulsion of DB(a,h)A given in place of drinking water for 237 to 279 days.

^eTumors were alveologenic carcinomas, a 100% incidence of pulmonary adenomatosis was also observed.

^fAdministered as a single MCA-containing beeswax pellet placed directly into the lower peripheral segment of the left lung.

^gOvert squamous cell carcinoma.

The published literature regarding chemical carcinogenesis in cell cultures is vast, despite the fact that systematic studies were not begun until the early 1960's due to the lack of a reproducible transformation assay. Berwald and Sachs (1963) first demonstrated that polycyclic hydrocarbons (MCA, BaP) could cause the direct malignant transformation of hamster embryo cells in culture. Transformed colonies have growth characteristics visually distinct from normal colonies and are readily seen above a background of normal cells. This assay can therefore be easily used as a screen to compare carcinogenic activity of suspect compounds. A common feature of these, and nearly all, transformed cells is that they give rise to fibrosarcomas upon inoculation into immunosuppressed animals. In addition to hamster embryo cells, malignant transformation has been demonstrated in organ cultures, liver cell cultures, fibroblastic cells derived from mouse ventral prostate, 3TC cell lines derived from mouse embryo cells, and various types of epithelial cells from humans and other animals (Heidelberger, 1973, 1975; Heidelberger and Boshell, 1975).

Early reports by Berwald and Sachs (1965) and Dipaolo and Donovan (1967) described alterations in hamster embryo cells induced by BaP, DMBA, and MCA which could be used as indicators of a change from normal to neoplastic state. The compounds were applied to cells in culture either dissolved in paraffin and impregnated on filter disks or as a colloidal suspension in growth medium. Following marked cytotoxicity, foci of transformed cells developed which displayed chromosomal abnormalities and the ability to grow indefinitely in culture. In addition, these transformed mass cul-

tures, when transplanted to four- to six-week-old hamsters, continued to grow and form tumors. A good correlation was obtained between in vitro carcinogenicity of a polycyclic hydrocarbon and the number of transformed clones they produced. The maximum rate of cell transformation in these studies was 25.6 percent in surviving cells, obtained by treatment with 10 µg/ml of BaP for six days. BaP treatment at 1 µg/ml for six days produced 19.9 percent transformation in surviving cells. Further data indicating the activity of several polycyclic carcinogens and their derivatives are summarized in Table 27. The K-region epoxides of DBahA and MCA are more active in the production of malignant transformation in hamster embryo cells than the parent hydrocarbons or the corresponding K-region phenols (Grover, et al. 1971; Huberman, et al. 1972). Although these results confirm the view that metabolism is necessary for carcinogenic activity, they conflict with data generated in vivo which indicate that K-region epoxides of polycyclic carcinogens are less active than the parent compound in various species. A possible reason for the lack of correlation is the relative instability of K-region epoxides as compared to the parent hydrocarbon when applied to the skin. It is likely that in vivo far less of the reactive K-region epoxide can survive passage through the skin to reach the basal cell layer. Furthermore, it has become apparent that the non-K-region diol-epoxide is likely to be the ultimate carcinogenic metabolite for most PAH. Several investigators have also made it evident that the toxicity and transforming activity of PAH are dissociable and occur by different processes (Landolph, et al. 1976; DiPaolo, et al. 1971a,b), with the toxicity being due to

TABLE 27

Hamster Embryo Cell Transformation Produced by Several Polycyclic Hydrocarbons and Their Derivatives

	Concentration, µg/ml	Total No. Colonies	Cloning Efficiency, %	No. Transformed Colonies	Transformation, %	Reference
DB(a,h)A ^a	2.5	760	4.2	4	0.5	Huberman, et al. 1972
	5	690	3.8	4	0.7	Huberman, et al. 1972
	10	790	4.4	7	0.9	Huberman, et al. 1972
DB(a,h)A ^b	2.5	1,341	13.4	3	0.2	Grover, et al. 1971
	5.0	1,363	14.0	11	0.8	Grover, et al. 1971
	10	1,365	14.5	7	0.5	Grover, et al. 1971
DB(a,h)A5,6-epoxide ^a	2.5	598	3.3	3	0.5	Huberman, et al. 1972
	5	601	3.3	12	2.0	Huberman, et al. 1972
	7.5	395	2.5	31	7.8	Huberman, et al. 1972
	10	350	1.9	14	4.0	Huberman, et al. 1972
DB(a,h)A5,6-epoxide ^b	2.5	895	10.1	7	0.8	Grover, et al. 1971
	5.0	866	9.3	20	2.3	Grover, et al. 1971
	7.5	817	9.3	22	2.7	Grover, et al. 1971
	10	707	7.7	30	4.2	Grover, et al. 1971
MCA ^c	2.5	404	10.1	9	2.2	Huberman, et al. 1972
	5	370	9.2	10	2.7	Huberman, et al. 1972
	7.5	349	8.7	15	4.3	Huberman, et al. 1972
MCA ^d	2.5	664	9.6	20	3.46	DiPaolo, et al. 1971a,b
MCA epoxide ^c	3.5	364	2.4	13	3.6	Huberman, et al. 1972
	5	245	1.5	8	3.3	Huberman, et al. 1972
	7	103	0.7	17	16.5	Huberman, et al. 1972
BaP ^d	1	1,016	8.46	25	2.46	DiPaolo, et al. 1971a
	5	394	7.17	21	5.33	DiPaolo, et al. 1971a

^a7-day treatment of cells seeded on a feeder layer.

^b7-8 day treatment of cells.

^c4-hour treatment of cells seeded in conditioned medium.

^d8-day treatment of cells.

random alkylation of nucleophilic regions within the cell. However, when hamster embryo cells are pretreated with weak chemical carcinogens which can induce microsomal enzyme activity [e.g., benz(a)anthracene, methyl methanesulfonate, ethyl methanesulfonate] before the addition of a potent carcinogen (e.g., MCA, BaP, DMBA), transformation may be considerably enhanced (DiPaolo, et al. 1971a,b, 1974).

As a prescreen for chemical carcinogens, cell transformation in vitro may be one of the most sensitive techniques available. Pienta and coworkers (1977) reported that 90 percent (54/60) of the carcinogens they tested transformed hamster embryo cells in vitro, whereas none of the noncarcinogens tested showed any activity. Moreover, many of the carcinogens which have not been shown to be mutagenic toward S. typhimurium in vitro (e.g., chrysene) were capable of transforming the hamster cells. It is noteworthy, however, that large differences exist in dosage requirements for transformation among those various test systems. Calculations have been made which show that a battery of tests using S. typhimurium (Ames assay), polymerase A-deficient E. coli, and hamster embryo cell transformation is capable of detecting nearly all carcinogens tested, both PAH and non-PAH types.

The alteration of microsomal enzyme activity either in vitro or in vivo is known to have a marked effect on the carcinogenic response to PAH. Nesnow and Heidelberger (1976) reported that in 10T_{1/2}CL8 cells, a line of contact-sensitive C3H mouse embryo fibroblasts, transformation in culture was altered by chemical modifiers of microsomal enzymes. Pretreatment of 10T_{1/2}CL8 cells with

benz(a)anthracene, a microsomal enzyme inducer, caused a doubling in MCA-mediated transformation. Similarly, treatment with inhibitors of epoxide hydrase (e.g., cyclohexene oxide; styrene oxide; 1,2,3,4-tetra-hydronaphthalene-1,2-oxide) caused an increase in transformation over that obtained with MCA treatment alone. Thus, treatments which can induce epoxide-forming enzymes and/or lower the activity of epoxide-degrading enzymes seemed to enhance the degree of transformation in cultured cells by altering steady-state levels of oncogenic epoxides.

Chen and Heidelberger (1969a,b) developed a system using C3H mouse ventral prostate cells to examine transformation by carcinogenic hydrocarbons under conditions in which no spontaneous malignant transformation occurred. Cells treated with MCA (1 µg/ml) for six days in culture produced fibrosarcomas in 100 percent of mice into which they were subcutaneously injected. When treated for only one day with MCA at the single cell stage, transformed foci were found in all clones grown to confluency. A good quantitative correlation was obtained between the in vivo oncogenic activity of eight hydrocarbons (including BaP, MCA, DMBA, and DBahA) and the number of transformed colonies produced in this system. In contrast to the enhanced transforming ability of K-region epoxides relative to the parent hydrocarbon in hamster embryo cells, the K-region epoxide derived from DMBA was less active and the K-region epoxides from MCA, DBahA, and benz(a)anthracene were more active than the parent compound in mouse prostate cells (Marquardt, et al. 1972, 1974). Moreover, the epoxide derived from DMBA was more toxic than DMBA itself. The anomalous behavior of DMBA may have

been due, however, to a decreased intracellular half-life of the epoxide because of its greater chemical reactivity.

Attempts to transform human cells in culture with PAH (e.g., BaP, MCA, DMBA) have generally met with failure (Leith and Hayflick, 1974). However, Rhim and coworkers (1975) reported that a human osteosarcoma clonal cell line could be further transformed in vitro with DMBA. Morphologic alterations and abnormal growth patterns became evident in cells treated with DMBA at 2.5 and 1.0 $\mu\text{g}/\text{ml}$ in the fifth subculture 52 to 57 days after exposure. One of the altered cell lines obtained from the 1 $\mu\text{g}/\text{ml}$ treatment was tumorigenic in nude mice by subcutaneous and intracerebral injection. Interpretation of the significance of these results is made difficult by the fact that an aneuploid sarcomatous cell line had to be employed in order to demonstrate successful transformation.

The use of organ cultures for the assessment of chemical carcinogenicity suffers from the lack of reliable biochemical and morphological parameters for measuring early neoplastic changes. Nevertheless, pioneering work in the application of organ culture to chemical carcinogenesis was performed by Lasnitzki (1963). Microgram quantities of MCA added to organ cultures of rat and mouse prostate fragments caused extensive hyperplasia and squamous metaplasia. However, these preneoplastic morphological effects are generally not associated with subsequent tumor development when carcinogen-treated pieces of tissue are implanted into host animals (Heidelberger, 1973). Limited success has been achieved with organ cultures of rat tracheas, which showed characteristic morphologic alterations when treated with DMBA, BaP, and MCA (Heidelberger,

1973). In addition, Crocker (1970) has exposed respiratory epithelia from the hamster, rat, dog, and monkey to BaP at 7 to 15 $\mu\text{g}/\text{ml}$ and observed occasional squamous metaplasia. More commonly, pleomorphic cells in a dysplastic epithelium were evident as a result of the treatment. Rat tracheas maintained in organ culture have been suggested as a useful system for the predictive screening of potential carcinogens (Lindsay, et al. 1974).

A unique organ culture technique has recently been reported in which BaP (4 or 12 mg) was administered to pregnant mice (strain A and C57Bl), and lung tissue of their 19- to 20-day-old embryos was subsequently explanted in culture (Shabad, et al. 1974). A transplacental influence of BaP was manifested as a proliferative stimulus in embryonic lung tissue. Hyperplasia arising in the bronchial epithelium led to the development of adenomas in a large percentage of the explants.

In the environment, man is unlikely to come in contact with only a single PAH, regardless of the route of exposure. Instead, PAH occur as complex mixtures in all environmental media. Despite this generally accepted fact, very few studies have been conducted on the carcinogenicity of defined PAH mixtures.

Among the most relevant studies conducted on the effects of PAH mixtures were those concerned with the carcinogenic components of automotive engine exhaust. Pfeiffer (1973, 1977) treated groups of 100 female NMRI mice with single subcutaneous injections of a mixture containing 10 noncarcinogenic PAH, in addition to BaP and/or dibenz(a,h)anthracene. The treatment combinations and dosages are summarized in Table 28. As the results depicted in

TABLE 28

Classification of Test Groups*

A	Dose (μg)	Substance	B	Dose (μg)	Substance		
A ₁	3.12	benzo(a) pyrene	B ₁	2.35	dibenz(a,h) anthracene		
A ₂	6.25		B ₂	4.7			
A ₃	12.5		B ₃	9.3			
A ₄	25.0		B ₄	18.7			
A ₅	50.0		B ₅	37.5			
A ₆	100.0		B ₆	75.0			
C	Substance	C ₁ dose (μg)	C ₂ dose (μg)	C ₃ dose (μg)	C ₄ dose (μg)	C ₅ dose (μg)	C ₆ dose (μg)
C-102	benzo(e) pyrene	2.15	4.3	8.75	17.5	35.5	70.0
	benzo(a) anthracene	3.125	6.25	12.5	25.0	50.0	100.0
	phenanthrene	125.0	250.0	500.0	1,000.0	2,000.0	4,000.0
	anthracene	31.25	62.5	125.0	250.0	500.0	1,000.0
	pyrene	65.1	131.2	262.5	525.0	1,050.0	2,100.0
	fluoranthene	28.1	56.25	112.5	225.0	450.0	900.0
	chrysene	3.125	6.25	12.5	25.0	50.0	100.0
	perylene	0.2	0.4	0.87	1.75	3.5	7.0
	benzo(ghi)perylene	12.8	25.6	51.25	102.5	205.0	410.0
	coronene	3.125	6.25	12.5	25.0	50.0	100.0
D			E				
D ₁	A ₁ + B ₁		E ₁	C ₁ + D ₁			
D ₂	A ₂ + B ₂		E ₂	C ₂ + D ₂			
D ₃	A ₃ + B ₃		E ₃	C ₃ + D ₃			
D ₄	A ₄ + B ₄		E ₄	C ₄ + D ₄			
D ₅	A ₅ + B ₅		E ₅	C ₅ + D ₅			
D ₆	A ₆ + B ₆		E ₆	C ₆ + D ₆			

*Source: Pfeiffer, 1977

Table 29 indicate, increases in tumor incidence could be attributed to the presence of increased amounts of BaP and of dibenz(a,h)anthracene. It is noteworthy that, at the lower dosages, dibenz(a,h)anthracene was more effective in producing tumors at the injection site than was BaP. Moreover, no effect of the 10 noncarcinogens on tumorigenic response was evident. Probit analysis of tumor incidence data indicated that the tumorigenic response from application of all 12 PAH was attributable solely to dibenz(a,h)anthracene.

Similar studies intended to reveal carcinogenic interactions among PAH found in automobile exhaust were conducted by Schmahl, et al. (1977). Eleven PAH were selected for their experiments, and various combinations were applied to the skin of NMRI mice in a proportion based on their respective weights in automobile exhaust (Table 30). Animals received twice weekly treatments for life (or until a carcinoma developed). Their results (Table 31) indicated that a mixture of carcinogenic PAH was more effective than BaP alone, and that the whole mixture (carcinogenic plus noncarcinogenic PAH) was not significantly more effective than the carcinogenic PAH group alone. Thus, the carcinogenic effects observed were solely attributable to the carcinogenic components of the mixture.

Human data: Although exposure to PAH occurs predominantly by direct ingestion (i.e., in food and in drinking water) there are no studies to document the possible carcinogenic risk to humans by this route of exposure. It is known only that significant quantities of PAH can be ingested by humans, and that in animals such exposures are known to cause cancers at various sites in the body.

TABLE 29

Tumor Incidence Resulting, by the End of the 114th Week,
from a Single Subcutaneous Application of Test Substances*

<u>BaP Group (A)</u>		<u>DBA Group (B)</u>		<u>BaP + DAB Group (D)</u>	<u>10 PAH Group (C)</u>	<u>12 PAH Group (E)</u>
Dose (μg)	No. of Tumors	Dose (μg)	No. of Tumors	No. of Tumors	No. of Tumors	No. of Tumors
3.12	9	2.35	37	48	6	41
6.25	35	4.7	39	44	8	55
12.5	51	9.3	44	61	6	61
25.0	57	18.7	56	68	4	72
50.0	77	37.5	65	69	13	68
100.0	83	75.0	69	79	5	82

*Source: Pfeiffer, 1977

TABLE 30

Doses (μg) Applied in Dermal Administration
Experiments, in Relation to Benzo(a)pyrene*

<u>Controls</u>				
Acetone	as solvent			
Benzo(a)pyrene	1.0	1.7	3.0	
<u>C PAH</u>				
Benzo(a)pyrene	1.0	1.7	3.0	
Dibenz(a,h)anthracene	0.7	1.2	2.1	
Benzo(a)anthracene	1.4	2.4	4.2	
Benzo(b)fluoranthene	<u>0.9</u>	<u>1.5</u>	<u>2.7</u>	
total	4.0	6.8	12.0	
<u>NC PAH</u>				
(Benzo(a)pyrene	1.0	3.0	9.0	27.0)
Phenanthrene	27.0	81.0	243.0	729.0
Anthracene	8.5	25.5	76.5	229.5
Fluoranthene	10.8	32.4	97.2	291.6
Pyrene	13.8	41.4	124.2	372.6
Chrysene	1.2	3.6	10.8	32.4
Benzo(e)pyrene	0.6	1.8	5.4	16.2
Benzo(ghi)perylene	<u>3.1</u>	<u>9.3</u>	<u>27.9</u>	<u>83.7</u>
total	65.0	195.0	585.0	1,755.0
<u>C PAH + NC PAH</u>				
(Benzo(a)pyrene	1.0	1.7	3.0)	
Total C PAH	4.0	6.8	12.0	
Total NC PAH	<u>65.0</u>	<u>110.5</u>	<u>195.0</u>	
Total C PAH + NC PAH	69.0	117.3	207.0	
Relation of C PAH:NC PAH is constantly 1:16.25				

*Source: Schmahl, et al. 1977

TABLE 31

Findings at the Site of Application of PAH to Mouse Skin*

Application	Single Dose µg	Initial No. of Animals	Effective No. of Animals	Histological Diagnosis at the Site of Application							
				Negative Abs. %		Papilloma Abs. %		Carcinoma Abs. %		Sarcoma Abs. %	
Solvent	-	100	81	80	99	-	-	-	-	1	1
BaP	1.0	100	77	66	86	1	1	10	13	-	-
BaP	1.7	100	88	63	72	-	-	25	28	-	-
BaP	3.0	100	81	36	44	2	3	43	53	-	-
C PAH	4.0	100	81	52	64	4	5	25	31	-	-
C PAH	6.8	100	88	31	35	3	3	53	60	1	1
C PAH	12.0	100	90	25	28	1	1	63	70	1	1
NC PAH	65.0	100	85	84	99	-	-	1	1	-	-
NC PAH	195.0	100	84	84	100	-	-	-	-	-	-
NC PAH	585.0	100	88	87	99	-	-	1	1	-	-
NC PAH	1,755.0	100	86	70	81	-	-	15	17	1	1
C PAH + NC PAH	69.0	100	89	43	48	1	1	44	49	1	1
C PAH + NC PAH	117.3	100	93	36	39	2	2	54	58	1	1
C PAH + NC PAH	207.0	100	93	28	30	1	1	64	69	-	-

^aThe decimal points have been rounded off; therefore, the sum of % values will not always be equivalent to 100%.

*Source: Schmahl, et al. 1977

Convincing evidence from air pollution studies indicates an excess of lung cancer mortality among workers exposed to large amounts of PAH-containing materials such as coal gas, tars, soot, and coke-oven emissions (Kennaway, 1925; Kennaway and Kennaway, 1936, 1947; Henry, et al. 1931; Kuroda, 1937; Reid and Buck, 1956; Doll, 1952; Doll, et al. 1965, 1972; Redmond, et al. 1972, 1976; Mazumdar, et al. 1975; Hammond, et al. 1976; Kawai, et al. 1967). However, no definite proof exists that the PAH present in these materials are responsible for the cancers observed. Nevertheless, our understanding of the characteristics of PAH-induced tumors in animals, and their close resemblance to human carcinomas of the same target organs, strongly suggests that PAH pose a carcinogenic threat to man, regardless of the route of exposure (Santodonato, et al. 1980).

The magnitude of the carcinogenic risk of PAH to man remains obscure in the community setting. Ambient levels of PAH in air are much lower than are encountered in occupational situations, and populations exposed are much more heterogeneous with regard to age, sex, and health status. However, the current state of knowledge regarding chemical carcinogenesis would lead to the conclusion that the number of cancers produced is directly proportional to the dose received by any route. One must assume, therefore, that the small amounts of PAH present in the environment (air, food, and water) under ambient conditions contribute in some degree to the observed incidence of lung cancer in most populations.

CRITERION FORMULATION

Existing Guidelines and Standards

There have been few attempts to develop exposure standards for PAHs, either individually or as a class. In the occupational setting, a Federal standard has been promulgated for coke oven emissions, based primarily on the presumed effects of the carcinogenic PAH contained in the mixture as measured by the benzene soluble fraction of total particulate matter. Similarly, the American Conference of Governmental Industrial Hygienists recommends a workplace exposure limit for coal tar pitch volatiles, based on the benzene-soluble fraction containing carcinogenic PAH. The National Institute for Occupational Safety and Health has also recommended a workplace standard for coal tar products (coal tar, creosote, and coal tar pitch), based on measurements of the cyclohexane extractable fraction. These standards are summarized below:

<u>Substance</u>	<u>Exposure Limit</u>	<u>Agency</u>
Coke Oven Emissions	150 $\mu\text{g}/\text{m}^3$, 8-hr. time-weighted average	U.S. Occupational Safety and Health Administration
Coal Tar Products	0.1 mg/m^3 , 10-hr. time-weighted average	U.S. National Institute for Occupational Safety and Health
Coal Tar Pitch of Volatiles	0.2 mg/m^3 (benzene soluble fraction) 8-hr. time-weighted average	American Conference of Governmental Industrial Hygienists

A drinking water standard for PAH as a class has been developed. The 1970 World Health Organization European Standards for Drinking Water recommends a concentration of PAH not to exceed 0.2 $\mu\text{g}/\text{l}$. This recommended standard is based on the composite analysis

of six PAH in drinking water: 1) fluoranthene, (2) benzo(a)pyrene, (3) benzo(g,h,i) perylene, (4) benzo(b)fluoranthene, (5) benzo(k)-fluoranthene, and (6) indeno(1,3,-cd)pyrene.

The designation of these six PAH for analytical monitoring of drinking water was not made on the basis of potential health effects or bioassay data on these compounds (Borneff and Kunte, 1969). Thus, it should not be assumed that these six compounds have special significance in determining the likelihood of adverse health effects resulting from absorption of any particular PAH. They are, instead, considered to be useful indicators for the presence of PAH pollutants. Borneff and Kunte (1969) found that PAH were present in ground water at concentrations up to 50 ng/l, and in drinking water at concentrations up to 100 ng/l. Based on these data they suggested that water containing more than 200 ng/l should be rejected. However, as data from a number of U.S. cities indicate (see Exposure section), levels of PAH in raw and finished waters are typically much less than the 0.2 µg/l criterion.

Current Levels of Exposure

This report presents considerable data which may be used to calculate an estimate of human exposure to PAH by all routes of entry to the body. However, quantitative estimates of human exposure to PAH require numerous assumptions concerning principal routes of exposure, extent of absorption, conformity of human lifestyle, and lack of geographic-, sex-, and age-specific variables. Nevertheless, by working with estimates developed for PAH as a class, it is possible through certain extrapolations to arrive at an admittedly crude estimate of PAH exposure.

Unfortunately, there are no environmental monitoring data available for most of the PAH which are specified under the Consent Decree in NRDC v. Train. By far the most widely monitored PAH in the environment is BaP; data on BaP levels in food, air, and water are often used as a measure of total PAH. Among the PAH routinely monitored in water, four compounds are included in the Consent Decree list: BaP, IP, BbFL, and BjFL. In addition, levels of FL and BPR have been routinely determined in water, as recommended by the World Health Organization.

The reported estimated average concentrations of BaP, carcinogenic PAH (BaP, BjFL, and IP), and total PAH in drinking water are 0.55 ng/l, 2.1 ng/l, and 13.5 ng/l, respectively (see Exposure section; Basu and Saxena, 1977). Thus, assuming that a human consumes 2 liters of water per day, the daily intake of PAH via drinking water would be:

$$0.55 \text{ ng/l} \times 2 \text{ liters/day} = 1.1 \text{ ng/day (BaP)}$$

$$2.1 \text{ ng/l} \times 2 \text{ liters/day} = 4.2 \text{ ng/day (carcinogenic PAH)}$$

$$13.5 \text{ ng/l} \times 2 \text{ liters/day} = 27.0 \text{ ng/day (total PAH)}$$

Borneff (1977) estimates that the daily dietary intake of PAH is about 8 to 11 $\mu\text{g/day}$. As a check on this estimate, PAH intake may be calculated based on reported concentrations in various foods (see Exposure section) and the per capita estimates of food consumption by the International Commission on Radiological Protection (1974). Taking a range of 1.0 to 10.0 ppb as a typical concentration for PAH in various foods, and 1,600 g/day as the total daily food consumption by man from all types of foods (i.e., fruits, vegetables, cereals, dairy products, etc.), the intake of PAH from

the diet would be in the range of 1.6 to 16.0 $\mu\text{g}/\text{day}$. An estimate of BaP ingestion from the diet may be similarly derived. Using 0.1 to 1.0 ppb as the range of BaP concentration in various foods, total daily BaP intake would be 0.16 to 1.6 $\mu\text{g}/\text{day}$.

Ambient air is reported to contain average levels of 0.5 ng/m^3 , 2.0 ng/m^3 , and 10.9 ng/m^3 for BaP, carcinogenic PAH, and total PAH, respectively (see Exposure section, Table 15). Taking the range of 15 m^3 to 23 m^3 as the average amount of air inhaled by a human each day results in an estimated intake of 0.005 to 0.0115 ng/day , 0.03 to 0.046 ng/day , and 0.164 to 0.251 ng/day for BaP, carcinogenic PAH, and total PAH, respectively.

In summary, a crude estimate of total daily exposure to PAH would be as shown in Table 32.

Two important factors are not taken into account in this estimate. First, it is known that tobacco smoking can contribute greatly to PAH exposure in man. Exposure to BaP from smoking one pack of cigarettes per day was shown to be 0.4 $\mu\text{g}/\text{day}$ (NAS, 1972). Second, the possibility for dermal absorption of PAH is assumed to contribute only a negligible amount to the total exposure. Only in certain occupational situations is dermal exposure expected to be quantitatively important.

Special Groups at Risk

An area of considerable uncertainty with regard to the carcinogenic hazard of PAH to man involves the relationship between aryl hydrocarbon hydroxylase (AHH) activity and cancer risk. Genetic variation in AHH inducibility has been implicated as a determining factor for susceptibility to lung and laryngeal cancer (Kellerman,

TABLE 32

Estimate of Human Exposure to PAH from Various Media

Source	Estimated Exposure		
	BaP	Carcinogenic PAH ^a	Total PAH
Water	0.0011 µg/day	0.0042 ug/day	0.027 ug/day
Food	0.160-1.6 ug/day		1.600-0.251 ug/day
Air	0.005-0.0115 ug/day	0.03-0.046 ug/day	0.164-0.251 ug/day
Total	0.166-1.6 ug/day		1.6-16 ug/day

^aTotal of BaP, B_jFL, and IP; no data are available for food.

et al. 1973a,b). It was suggested that the extent of AHH inducibility in lymphocytes was correlated with increasing susceptibility to lung cancer formation.

Paigen, et al. (1978) have examined the question of genetic susceptibility to cancer, and concluded that epidemiologic evidence supports this hypothesis. Moreover, they were able to show that AHH inducibility in lymphocytes segregates in the human population as a genetic trait. However, their studies failed to find a correlation between this inducibility and presumed cancer susceptibility, either among healthy relatives of cancer patients or in patients who had their cancer surgically removed. It is noteworthy that previous investigations on AHH inducibility were conducted in persons with active cancer.

Recent studies with other human tissues (liver and placenta) have provided important new data concerning the carcinogen-metabolizing capacity of man and its implications for cancer susceptibility. Conney, et al. (1976) examined individual differences in the metabolism of drugs and carcinogens in human tissues, and have identified drugs which may serve as model substrates to provide an indirect index of carcinogen metabolism for man. The rates for antiprene, hexobarbital, and zoxazolamine hydroxylation in human autopsy livers were highly, but not perfectly, correlated with the rates of BaP metabolism. In human placenta, an almost perfect correlation was found between zoxazolamine hydroxylase activity and BaP hydroxylase activity (Kapitulnik, et al. 1976a). Thus, metabolism of BaP and zoxazolamine by human placenta occurs by the same enzyme system(s) or by different enzyme systems under the same reg-

ulatory control (Kapitulnik, et al. 1977a). BaP and zoxazolamine hydroxylase activities were also shown to be significantly enhanced in placentas obtained from women who smoked cigarettes.

The lack of perfect correlations for the hepatic metabolism of BaP and certain drugs in many subjects indicated the presence of several monooxygenases in human liver which catalyze the oxidative metabolism of these compounds. Furthermore, large inter-individual differences exist in the capacity of humans to metabolize foreign chemicals both in vitro and in vivo. Further studies showed that 7,8-benzoflavone markedly stimulated the hydroxylation of BaP, antipyrene, and zoxazolamine in human liver samples, but with a wide variation in magnitude among different samples. These results suggested the presence of multiple monooxygenases or cytochrome P-450 in the different liver samples (Kapitulnik, et al. 1977b). Moreover, 7,8-benzoflavone did not affect the hydroxylation of coumarin or hexobarbital, thereby indicating the existence of different monooxygenases for metabolism of these substrates.

Multiple forms of cytochrome P-450 have been shown in the livers of rats, rabbits, and mice, but not thus far in humans (Kapitulnik, et al. 1977a). More important, however, MCA is a potent inducer of BaP hydroxylase activity in rats but does not stimulate antipyrene hydroxylase, clearly suggesting that metabolism of PAH in rodents may be regulated by different enzyme systems than in humans (Kapitulnik, et al. 1977a).

In contrast to the apparent multiplicity of cytochrome P-450 dependent enzyme systems for the oxidative metabolism of PAH in man, a single epoxide hydrase with broad substrate specificity may

be present in human liver (Conney, et al. 1976; Kapitulnik, et al. 1977c). Because the hydration of arene oxides may lead to the formation of dihydrodiol carcinogen precursors, the capacity of different humans to metabolize epoxides may affect cancer susceptibility. It is not known, however, if enhanced dihydrodiol formation would increase cancer risk or decrease cancer risk.

Thomson and Slaga (1976) did not obtain a correlation of AHH induction with skin-tumor-inducing ability in mice for a series of unsubstituted hydrocarbons. Nevertheless, the highest AHH enzyme activity was found in the epidermal layer of the skin, which is the major point of contact with many environmental chemicals. These results may be interpreted to indicate that a chemical carcinogen may not necessarily induce its own bioactivation, but instead can be transformed into a reactive intermediate by virtue of increased AHH activity stimulated by other noncarcinogenic compounds.

Due consideration must also be given to the fact that, in addition to the initiation of resting cells by a chemical carcinogen, a promotion phase involving cell proliferation is also involved in skin carcinogenesis (Yuspa, et al. 1976). Therefore, although certain aromatic hydrocarbons are effective enzyme inducers, their bioactivated metabolites may function only as initiators having no promoting ability. A potent complete carcinogen, however, will be transformed not only into a powerful tumor initiator but will also be able to interact with cellular membranes, alter genetic expression, and ultimately cause irreversible cell proliferation. These observations raise certain doubts concerning the validity and/or reliability of equating enzyme inducibility with

carcinogenic potential for chemical agents. Further reinforcement of this opinion has been provided by Shulte-Hermann (1977) who showed that cell proliferation is not a direct result of enzyme induction, even though both processes are normally coupled.

The further possibility that the genetics of AHH inducibility is organ-dependent rather than strain-dependent in animals has important implications for evaluating susceptibility to PAH-induced cancers (Kouri, et al. 1976). Most significant is the demonstration that pulmonary AHH may be inducible in all strains of mice, regardless of the inducibility of hepatic AHH. Since the respiratory epithelium represents a primary portal of entry for PAH, AHH activity which is induced in this tissue may bear importantly on susceptibility to malignancy.

Enzyme induction by PAH is not limited to AHH. Owens (1977) recently demonstrated that MCA can induce hepatic UDP-glucuronosyltransferase activity in certain inbred strains of mice. This enzyme catalyzes the conjugation and excretion of PAH substrates after they have first been oxygenated by AHH. The induction of this transferase activity and that of AHH was apparently regulated by a single genetic locus. However, transferase inducibility does not depend on AHH levels, but rather is stoichiometrically related to the concentration of a specific and common cytosolic receptor regulating both enzyme induction processes. Owens further demonstrated that AHH activity can be fully induced in certain mouse strains (e.g., by 2,3,7,8-tetrachlorodibenzo-p-dioxin) without greatly enhancing the transferase activity. Earlier studies had established that chrysene and chlorpromazine were potent inducers

of AHH activity while having little effect on transferase activity (Aitio, 1974a,b). Subsequent exposure to carcinogenic PAH (i.e., MCA) could lead to maximal oxidative metabolism but little transferase-catalyzed removal of metabolites by glucuronic acid conjugation. This situation would be exacerbated by the fact that metabolites of MCA are incapable of further inducing the transferase activity. This effect may have considerable toxicologic significance in that the highly reactive epoxides of PAH formed by the action of AHH under these circumstances may not be adequately removed by glucuronidation. Thus, one must consider the total exposure of all environmental agents and their possible effect on critical enzymatic processes before attempting to assess the toxicologic impact of exposure to a specific PAH. In summary, there is a need to further explore the relative effects of enzyme induction on the metabolic activation of chemicals to toxic products, versus metabolism of chemicals via detoxification pathways, when considering the possibility of special groups at risk.

Basis and Derivation of Criterion

The presently available data base is inadequate to support the derivation of individual criteria for each of the PAH as specified under the Consent Decree. This problem arises primarily from the diversity of test systems and bioassay conditions employed for determining carcinogenic potential of individual PAH in experimental animals. Furthermore, it is not possible to estimate the intake via water of individual PAH, except for those compounds which have been selected by the World Health Organization for environmental monitoring. Therefore, an approach to criterion development is

adopted in this report with the objective of deriving criteria for individual carcinogenic PAH, which will lead to effective control of PAH as a class. This approach is attractive in that it recognizes the fact that environmental exposures to PAH invariably occur by contact with complex, undefined, PAH mixtures.

The attempt to develop a drinking water criterion for PAH as a class is hindered by several gaps in the scientific data base:

- (1) The PAH class is composed of numerous compounds having diverse biological effects and varying carcinogenic potential. A "representative" PAH mixture, has not been defined.
- (2) The common practice of using data derived from studies with BaP to make generalizations concerning the effects of environmental PAH may not be scientifically sound.
- (3) No chronic animal toxicity studies involving oral exposure to PAH mixtures exist.
- (4) No direct human data concerning the effects of exposure to defined PAH mixtures exist.

However, assuming that the development of a criterion must proceed despite these obstacles, certain approaches may be taken to circumvent deficiencies in the data base. The choice of an appropriate animal bioassay from which to derive data for application to the human cancer risk assessment should be guided by several considerations. Primary emphasis must be placed on appropriate animal studies which: (1) include sufficient numbers of animals for statistically reliable results; (2) involve long-term low-level exposures to PAH; (3) include a proper control group; and (4) achieve positive dose-related carcinogenic response.

Because there are no studies available regarding chronic oral exposure to PAH mixtures, it is necessary to derive a criterion based upon data involving exposure to a single compound. Two studies can be selected, one involving BaP ingestion (Neal and Rigdon, 1967) and one involving DBA ingestion (Snell and Stewart, 1962). Both compounds are recognized as animal carcinogens, and both are known to be environmental contaminants to which humans are exposed.

Presently, there is no way to quantitate the potential human health risks incurred by the interaction of PAH, either among themselves or with other agents (e.g., tumor initiators, promoters, inhibitors) in the environment. In addition, it is known that PAH commonly produce tumors at the site of contact (i.e., forestomach tumors by oral exposure to BaP; lung tumors by intratracheal administration; skin tumors by dermal application). Thus, consideration of the extent of absorption may not always be necessary in the case of carcinogenic PAH, and will in fact result in underestimation of actual risk if only distant target sites are considered. Calculation of the water quality criterion based upon bioassay data for BaP is presented in the Appendix.

The water quality criterion for BaP derived using the linearized multistage model, as described in the Human Health Methodology Appendices to the October 1980 Federal Register notice which announced the availability of this document, is 28 ng/l. For the sake of comparison, a water quality criterion for DBA was calculated using the procedure developed by Mantel and Bryan (1961). As opposed to the linearized multistage model, which is logistic and defines acceptable risk as 1/100,000, the Mantel and Bryan (1961)

model is probabilistic and defines acceptable risk as 1/100,000,000. Furthermore, the Mantel and Bryan model (1961) is concerned with the maximum tumor incidence in treated animals at the 99 percent confidence level versus the 95 percent confidence level in the linearized multistage model. Using the Mantel and Bryan (1961) approach with DBA, the resultant water quality criterion is 13.3 ng/l.

Under the Consent Decree in NRDC v. Train, criteria are to state "recommended maximum permissible concentrations (including where appropriate, zero) consistent with the protection of aquatic organisms, human health, and recreational activities." BaP is a known animal carcinogen. Because there is no recognized safe concentration for a human carcinogen, the recommended concentration in water for maximum protection of human health is zero.

Because attaining a zero concentration level may be infeasible in some cases and in order to assist the Agency and states in the possible future development of water quality regulations, the concentrations of BaP corresponding to several incremental lifetime cancer risk levels have been estimated. A cancer risk level provides an estimate of the additional incidence of cancer that may be expected in an exposed population. A risk of 10^{-5} for example, indicates a probability of one additional case of cancer for every 100,000 people exposed, a risk of 10^{-6} indicates one additional case of cancer for every million people exposed, and so forth.

PAH are widely distributed in the environment as evidenced by their detection in sediments, soils, air, surface waters, and plant and animal tissues. The ecological impact of these chemicals, how-

ever, is uncertain. Numerous studies show that despite their high lipid solubility, PAH show little tendency for bioconcentration in the fatty tissues of animals or man. This observation is not unexpected, in light of convincing evidence to show that PAH are rapidly and extensively metabolized.

Lu, et al. (1977) have published the only available study regarding the bioconcentration and biomagnification of a PAH in model ecosystem environments. They reported that the bioconcentration of BaP, expressed as concentration in mosquitofish/concentration in water was zero. This was apparently due to the fact that the fish metabolized the BaP about as rapidly as it was absorbed. On the other hand, in a 33-day terrestrial-aquatic model ecosystem study, BaP showed a small degree of biomagnification which probably resulted from food chain transfer. In this case the biomagnification factor for mosquitofish was 30. Based on the results of Lu, et al. (1977) a bioconcentration (BCF) factor of 30 was employed for the purpose of calculating a water quality criterion.

In the Federal Register notice of availability of draft ambient water quality criteria, EPA stated that it is considering setting criteria for BaP at an interim target risk level of 10^{-5} , 10^{-6} , or 10^{-7} as shown in the following table.

<u>Exposure Assumptions</u> (per day)	<u>BaP</u>			
	<u>Risk Levels and Corresponding Criteria (1)</u> ng/l			
	<u>0</u>	<u>10⁻⁷</u>	<u>10⁻⁶</u>	<u>10⁻⁵</u>
2 liters of drinking water and consumption of 6.5 grams fish and shellfish (2)	0	0.28	2.8	28.0
Consumption of fish and shellfish only.		3.11	31.1	311.0

- (1) Calculated by applying a linearized multistage model as previously discussed. Appropriate bioassay data used in the calculation of the model are presented in the Appendix. Since the extrapolation model is linear at low doses, the additional lifetime risk is directly proportional to the water concentration. Therefore, water concentrations corresponding to other risk levels can be derived by multiplying or dividing one of the risk levels and corresponding water concentrations shown in the table by factors such as 10, 100, 1,000, and so forth.
- (2) Approximately 9 percent of the PAH exposure, assumed to be BaP, results from the consumption of aquatic organisms which exhibit an average bioconcentration potential of 30-fold based on the work of Lu, et al. (1977). The remaining 91 percent of PAH exposure results from drinking water.

Concentration levels were derived assuming a lifetime exposure to various amounts of PAH (1) occurring from the consumption of both drinking water and aquatic life grown in water containing the corresponding PAH concentrations and, (2) occurring solely from the consumption of aquatic life grown in the waters containing the cor-

responding PAH concentrations. Because data indicating other sources of exposure and the concentration to total body burden are inadequate for quantitative use, the criterion reflects the increment to risks associated with ambient water exposure only.

REFERENCES

- Abe, S. and M. Sasaki. 1977. Studies on chromosomal aberrations and sister chromatid exchanges induced by chemicals. Proc. Japan Acad. 53: 46.
- Ahlstrom, U. 1974. Chromosomes of primary carcinomas induced by 7,12-dimethylbenz(a)-anthracene in the rat. Hereditas. 78: 235.
- Ahokas, J.T., et al. 1975. Metabolism of polycyclic hydrocarbons by a highly active aryl hydrocarbon hydroxylase system in the liver of a trout species. Biochem. Biophys. Res. Comm. 63: 635.
- Aitio, A. 1974a. Different elimination and effect on mixed function oxidase of 20-methylcholanthrene after intragastric and intraperitoneal administration. Res. Comm. Chem. Pathol. Pharmacol. 9: 701.
- Aitio, A. 1974b. Effect of chrysene and carbon tetrachloride administration on rat hepatic microsomal monooxygenase and UDP glucuronosyltransferase activity. FEBS Lett. 42: 46.
- Akin, F.J. 1976. Anti-tumorigenic effect of maleic hydrazide on mouse skin. Jour. Agric. Food Chem. 24: 672.

Albert, R., et al. 1978. Temporal aspects of tumorigenic response to individual and mixed carcinogens. Comprehensive Prog. Rep. Inst. Environ. Med., New York Univ. Med. Center, New York.

Andelman, J.B. and J.E. Snodgrass. 1974. Incidence and Significance of Polynuclear Aromatic Hydrocarbons in the Water Environment. In: CR, Critical Reviews in Environmental Control. p. 69.

Andelman, J.B. and M.J. Suess. 1970. Polynuclear aromatic hydrocarbons in the water environment. Bull. World Health Organ. 43: 479.

Andrews, L.S., et al. 1976. Characterization and induction of aryl hydrocarbon [benzo(a)pyrene] hydroxylase in rabbit bone marrow. Res. Comm. Chem. Pathol. Pharmacol. 15: 319.

Arcos, J.S. and M.F. Argus. 1974. Chemical Induction of Cancer. Vol. IIA. Academic Press, New York.

Autrup, H., et al. 1978. Metabolism of (³H)benzo(a)pyrene by cultured human bronchus and cultured human pulmonary alveolar macrophages. Lab. Inv. 38: 217.

Bailey, E.J. and N. Dungal. 1958. Polycyclic hydrocarbons in Iceland smoked food. Br. Jour. Cancer. 12: 348.

Baldwin, R.W. 1973. Immunological Aspects of Chemical Carcinogenesis. In: G. Klein and S. Weinhouse (eds.), Advances in Cancer Research. Academic Press, New York, London. 18: 1.

Bartle, K.D., et al. 1974. High-resolution GLC profiles of urban air pollutant polynuclear aromatic hydrocarbons. Int. Jour. Environ. Anal. Chem. 3: 349.

Bast, R.C., Jr., et al. 1976. Development of an assay for aryl hydrocarbon [benzo(a)pyrene] hydroxylase in human peripheral blood monocytes. Cancer Res. 36: 1967.

Basu, D.K. and J. Saxena. 1977. Analysis of raw and drinking water samples for polynuclear aromatic hydrocarbons. EPA P.O. No. CA-7-2999-A, and CA-8-2275-B. U.S. Environ. Prot. Agency, HERL, Cincinnati, Ohio.

Basu, D.K. and J. Saxena. 1978. Polynuclear aromatic hydrocarbons in selected U.S. drinking waters and their raw water sources. Environ. Sci. Technol. 12: 795.

Bayer, U. 1978. In vivo Induction of Sister Chromatid Exchanges by Three Polyaromatic Hydrocarbons. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: 2nd. Int. Symp. on Analysis, Chemistry, and Biology (Carcinogenesis - A Comprehensive Survey: Vol. 3). Raven Press, New York. 3: 101.

Bend, J.R., et al. 1976. Hepatic and Extrahepatic Glutathione S-transferase Activity Toward Several Arene Oxides and Epoxides in the Rat. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: Chemistry, Metabolism, and Carcinogenesis (Carcinogenesis - A Comprehensive Survey: Vol. 1). Raven Press, New York. 1: 63.

Berneblum, I. 1941. The cocarcinogenic action of croton resin. Cancer Res. 1: 44.

Berwald, Y. and L. Sachs. 1963. In vitro transformation with chemical carcinogens. Nature. 200: 1182.

Berwald, Y. and L. Sachs. 1965. In vitro transformation of normal cells to tumor cells by carcinogenic hydrocarbons. Jour. Natl. Cancer Inst. 35: 641.

Biedler, J.L., et al. 1961. Chromosome lesions associated with carcinogen-induced tumors in mice. Nature. 192: 286.

Biernoth, G. and H.E. Rost. 1967. The occurrence of PAH in coconut oil and their removal. Chem. Ind. 45: 2002.

Biernoth, G. and H.E. Rost. 1968. The occurrence of PAH in edible oils and their removal. Arch. Hyg. 152: 238. (Berl.)

Binet, L. and L. Mallet. 1964. Diffusion of PAH in the living environment. *Gaz. Hop. (Paris)* 135: 1142. (Included in *Chem. Abstr.* 60: 2282c).

Bird, C.C., et al. 1970. Protection from the embryopathic effects of 7-hydroxymethyl-12-methylbenz(a)anthracene by 2-methyl-1,2-bis-(3-pyridyl)-1-propanone(metopirone ciba) and -diethylamino-ethyl-diphenyl-n-propyl acetate (SKR 525-A). *Br. Jour. Cancer* 24: 548.

Bjørseth, A. 1978. Analysis of Polycyclic Aromatic Hydrocarbons in Environmental Samples by Glass Capillary Gas Chromatography. In: R.I. Freudenthal and P.W. Jones (eds.), *Polynuclear Aromatic Hydrocarbons: 2nd. Int. Symp. on Analysis, Chemistry, and Biology (Carcinogenesis - A Comprehensive Survey: Vol. 3)*. Raven Press, New York. 3: 75.

Bock, F.G. 1964. Early effects of hydrocarbons on mammalian skin. *Progr. Exp. Tumor Res.* 4: 126.

Bock, F.G. and T.L. Dao. 1961. Factors affecting the polynuclear hydrocarbon level in rat mammary glands. *Cancer Res.* 21: 1024.

Booth, J., et al. 1974. The metabolism of polycyclic hydrocarbons by cultured human lymphocytes. *FEBS Lett.* 43: 341.

Borneff, J. 1964. Carcinogenic substances in water and soil. Part XV: Interim results of the former investigations. Arch. Hyg. 148: 1. (Berl.)

Borneff, J. 1977. Fate of carcinogens in aquatic environment. Prepublication copy received from author.

Borneff, J. and H. Kunte. 1964. Carcinogenic substances in water and soil. XVI: Evidence of PAH in water samples through direct extraction. Arch. Hyg. Bakt. 148: 585.

Borneff, J. and H. Kunte. 1965. Carcinogenic substances in water and soil. XVII: About the origin and evaluation of PAH in water. Arch. Hyg. Bakt. 149: 226.

Borneff, J. and H. Kunte. 1969. Carcinogenic substances in water and soil. XXVI: A routine method for the determination of PAH in water. Arch. Hyg. (Berl) 153: 220.

Boyland, E. and P. Sims. 1967. The carcinogenic activities in mice of compounds related to benz(a)anthracene. Int. Jour. Cancer. 2: 500.

Boyland, E., et al. 1965. Induction of adrenal damage and cancer with metabolites of 7,12-dimethylbenz(a)anthracene. Nature. 207: 816.

Brookes, P. 1977. Mutagenicity of polycyclic aromatic hydrocarbons. *Mutat. Res.* 39: 257.

Butenandt, A. and H. Dannenberg. 1956. The Biochemistry of Tumors. In: F. Buchner, et al. (ed.), Springer-Verlag, Handbuch der Allgemeinen Pathologie. 6: 107. (Berlin)

Buu-Hoi, N.P. 1959. Carcinogenic Materials. In: K.F. Bauer (ed.), Georg Thieme Verlag. Medizinische Grundlagen Forschung. Stuttgart. 2: 465.

Buu-Hoi, N.P. 1964. New developments in chemical carcinogenesis by polycyclic hydrocarbons and related heterocycles: A review. *Cancer Res.* 24: 1511.

Cahnmann, H.J. and M. Kuratsune. 1957. Determination of polycyclic aromatic hydrocarbons in oysters collected in polluted water. *Anal. Chem.* 29: 1312.

Cawein, M.J. and K.L. Sydnor. 1968. Suppression of cellular activity in the reticuloendothelial system of the rat by 7,12-dimethylbenz(a)anthracene. *Cancer Res.* 28: 320.

Chalmers, J.G. and A.H.M. Kirby. 1940. The elimination of 3,4-benzpyrene from the animal body after subcutaneous injection. I. Unchanged benzpyrene. *Biochem. Jour.* 34: 1191.

Chen, T.T. and C. Heidelberger. 1969a. In vitro malignant transformation of cells derived from mouse prostate in the presence of 3-methylcholanthrene. Jour. Natl. Cancer Inst. 42: 915.

Chen, T.T. and C. Heidelberger. 1969b. Quantitative studies on the malignant transformation of mouse prostate cells by carcinogenic hydrocarbons in vitro. Int. Jour. Cancer. 4: 166.

Chu, E.W. and R.A. Malmgren. 1965. An inhibitory effect of vitamin A on the induction of tumors in the forestomach and cervix in the Syrian hamster by carcinogenic polycyclic hydrocarbons. Cancer Res. 25: 885.

Chuang, A.H.L., et al. 1977. Aryl hydrocarbon hydroxylase in mouse mammary gland: in vitro study using mammary cell lines. Chem. Biol. Inter. 17: 9.

Cohn, J.A., et al. 1977. On the occurrence of cytochrome P-450 and aryl hydrocarbon hydroxylase activity in rat brain. Jour. Exp. Med. 145: 1607.

Colucci, J.M. and C.R. Begeman. 1971. Polynuclear Aromatic Hydrocarbons and Other Pollutants in Los Angeles Air. In: Proc. Int. Clean Air Cong. Academic Press. 2: 28.

Cone, M.V. and P. Nettesheim. 1973. Effects of vitamin A on 3-methylcholanthrene-induced squamous metaplasia and early tumors in the respiratory tract of rats. Jour. Natl. Cancer Inst. 50: 1599.

Conney, A.H. 1967. Pharmacological implications of microsomal enzyme induction. Pharmacol. Rev. 19: 317.

Conney, A.H., et al. 1976. Use of Drugs in the Evaluation of Carcinogen Metabolism in Man. In: R. Montesano and L. Tomatis (eds.), Screening Tests in Chemical Carcinogenesis. IARC Publ. No. 12. Lyon, France. p. 319.

Conney, A.H., et al. 1977a. Metabolism and Biological Activity of Benzo(a)pyrene and its Metabolic Products. In: D.J. Jallow, et al. (eds.), Biological Reactive Intermediates. Plenum Press.

Conney, A.H., et al. 1977b. Regulation of drug metabolism in man by environmental chemicals and diet. Fed. Proc. 36: 1647.

Cordle, F., et al. 1978. Human exposure to polychlorinated biphenyls and polybrominated biphenyls. Environ. Health Perspect. 24: 157.

Cornfield, J. 1977. Carcinogenic risk assessment. Science. 198: 693.

Crocker, T.T. 1970. Effect of Benzo(a)pyrene on Hamster, Rat, Dog, and Monkey Respiratory Epithelia in Organ Culture. In: Proc. Biol. Div., Oak Ridge Natl. Lab. Conf. in Gatlinburg, Tenn. Oct. 8-11, 1968. AEC Symp. Series 18, Oak Ridge, Tennessee. Div. Tech. Info., U.S. Atomic Energy Comm.

Currie, A.R., et al. 1970. Embryopathic effects of 7,12-dimethylbenz(a)anthracene and its hydroxymethyl derivatives in the Sprague-Dawley rat. *Nature*. 226: 911.

Czygan, P., et al. 1974. The effect of dietary protein deficiency on the ability of isolated hepatic microsomes to alter the mutagenicity of a primary and a secondary carcinogen. *Cancer Res.* 34: 119.

Dao, T.L., et al. 1959. Level of 3-methylcholanthrene in mammary glands of rats after intragastric instillation of carcinogen. *Proc. Soc. Exptl. Biol. Med.* 102: 635.

Davies, R.I. and G. Wynne-Griffith. 1954. Cancer and soils in the country of Anglesey. *Br. Jour. Cancer.* 8: 56.

Diehl, J.S. and S.W. Tromp. 1953. First report on the geographical and geological distribution of carcinogens in the Netherlands. Leiden Foundation for the Study of Psychophysics.

Dikun, P.P. and A.I. Makhinenko. 1963. Detection of BP in the schistose plant resins, in its effluents and in water basins after discharge of effluents. Gig. i. Sanit. 28: 10.

DiPaolo, J.A. and P.J. Donovan. 1967. Properties of Syrian hamster cells transformed in the presence of carcinogenic hydrocarbons. Experi. Cell Res. 48: 261.

DiPaolo, J.A., et al. 1971a. Transformation of hamster cells in vitro by polycyclic hydrocarbons without cytotoxicity. Proc. Natl. Acad. Sci. 68: 2958.

DiPaolo, J.A., et al. 1971b. Characteristics of primary tumors induced by carcinogenic polycyclic hydrocarbons in Syrian hamsters. Jour. Natl. Cancer Inst. 46: 171.

DiPaolo, J.A., et al. 1974. Enhancement by alkylating agents of chemical carcinogen transformation of hamster cells in culture. Chem. Biol. Inter. 9: 351.

Doll, R. 1952. The causes of death among gas workers with special reference to cancer of the lung. Br. Jour. Ind. Med. 9: 180.

Doll, R., et al. 1965. Mortality of gas workers with special reference to cancers of the lung and bladder, chronic bronchitis, and pneumoconiosis. Br. Jour. Ind. Med. 22: 1.

Doll, R., et al. 1972. Mortality of gas workers - final report of a prospective study. Br. Jour. Ind. Med. 29: 394.

Draudt, H.N. 1963. The meat smoking process: A review. Food Technol. 17: 85.

Dungal, N. 1961. Can smoked food be carcinogenic? Acta Unio Intern. Contra. Cancrum. 17: 365.

Dunn, B.P. and H.F. Stich. 1976. Release of the carcinogen benzo(a)pyrene from environmentally contaminated mussels. Bull. Environ. Contam. Toxicol. 15: 398.

Fabian, B. 1965. Carcinogenic substances in edible fat and oil. Part VI: Further investigations on margarine and chocolate. Arch. Hyg. 153: 21. (Berl.)

Falk, H.L., et al. 1964. Inhibition of carcinogenesis. The effect of polycyclic hydrocarbons and related compounds. Arch. Environ. Health. 9: 169.

Faoro, R.B. and J.A. Manning. 1978. Trends in benzo(a)pyrene.

Fedorenko, Z.P. 1964. The effect of biochemical treatment of wastewater of a by-product coke plant on the BP content. Gig. i. Sanit. 29: 17.

Feron, V.J., et al. 1973. Dose-response correlation for the induction of respiratory-tract tumors in Syrian golden hamsters by intratracheal instillations of benzo(a)pyrene. Eur. Jour. Cancer 9: 387.

Filipovic, J. and L. Toth. 1971. Polycyclische Kohlenwasserstoffe in Geraeucherten Jugoslawischen Fleischwaren. Fleischwirtschaft. 51: 1323.

Flesher, J.S. 1967. Distribution of radioactivity in the tissues of rats after oral administration of 7,12-dimethylbenz(a)anthracene-³H. Biochem. Pharmacol. 16: 1821.

Flesher, J.S., et al. 1976. Oncogenicity of K-region epoxides of benzo(a)pyrene and 7,12-dimethylbenz(a)anthracene. Int. Jour. Cancer. 18: 351.

Ford, E. and C. Huggins. 1963. Selective destruction in testis induced by 7,12-dimethylbenz(a)anthracene. Jour. Exp. Med. 118: 27.

Fox, M.A. and S.W. Staley. 1976. Determination of polycyclic aromatic hydrocarbons in atmospheric particulate matter by high pressure liquid chromatography coupled with fluorescence techniques. Anal. Chem. 48: 992.

Frethein, K. 1976. Carcinogenic polycyclic aromatic hydrocarbons in Norwegian smoked meat. Jour. Agric. Food Chem. 24: 976.

Freudenthal, R.I., et al. 1978. A Comparison of the Metabolites of Benzo(a)pyrene by Lung Mixed Function Oxidase from Rat, Rhesus, and Humans. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: 2nd. Int. Symp. on Analysis, Chemistry, and Biology (Carcinogenesis - A Comprehensive Survey: Vol. 3). Raven Press, New York. 3: 313.

Gelboin, H.V., et al. 1972. Microsomal Hydroxylases: Studies on the Mechanism of Induction and their Role in Polycyclic Hydrocarbon Action. In: Collection of Papers Presented at the Ann. Symp. on Fundamental Cancer Res., Ser. 24. p. 214.

Gordon, R.J. 1976. Distribution of airborne polycyclic aromatic hydrocarbons throughout Los Angeles. Environ. Sci. Technol. 10: 370.

Gordon, R.J. and R.J. Bryan. 1973. Patterns of airborne polynuclear hydrocarbon concentrations at four Los Angeles sites. Environ. Sci. Technol. 7: 1050.

Gottschalk, R.G. 1942. Quantitative studies on tumor production in mice by benzpyrene. Proc. Soc. Exp. Biol. Med. 50: 369.

Graf, W. and W. Nowak. 1966. Promotion of growth in lower and higher plants by carcinogenic polycyclic aromatics. Arch. Hyg. Bakt. 150: 513.

Graf, W., et al. 1975. Levels of carcinogenic polycyclic aromatic hydrocarbons in human and animal tissues. 3rd communication. Zbl. Bakt. Hyg. I. Abst. Orig. B. 161: 85.

Greinke, R.A. and I.C. Lewis. 1975. Development of a gas chromatographic - Ultraviolet absorption spectrometric method for monitoring petroleum pitch volatiles in the environment. Anal. Chem. 47: 2151.

Grimmer, G. 1974. Detection and occurrence of polycyclic hydrocarbons in yeast cultured on mineral oils. Dtsch. Lebensm. Rundsch. 70: 394.

Grimmer, G. and A. Hildebrandt. 1967. Content of polycyclic hydrocarbons in crude vegetable oils. Chem. Ind. p. 2,000.

Grover, P.L., et al. 1971. In vitro transformation of rodent cells by K-region derivatives on polycyclic hydrocarbons. Proc. Natl. Acad. Sci. 68: 1098.

Grundin, R., et al. 1973. Induction of microsomal aryl hydrocarbon (3,4-benzo(a)pyrene) hydroxylase and cytochrome P-450 in rat cortex. I. Characteristics of the hydroxylase system. Arch. Biochem. Biophys. 158: 544.

Guerrero, H., et al. 1976. High-pressure liquid chromatography of benzo(a)pyrene and benzo(g,h,i)perylene in oil-contaminated shellfish. Jour. Assoc. Off. Anal. Chem. 59: 989.

Haber, S.L. and R.W. Wissler. 1962. Effects of vitamin E on carcinogenicity of methylcholanthrene. Proc. Soc. Exp. Biol. Med. 111: 774.

Haddow, A., et al. 1937. The influence of certain carcinogenic and other hydrocarbons on body growth in the rat. Proc. R. Soc. B. 122: 477.

Hammond, E.C., et al. 1976. Inhalation of benzpyrene and cancer in man. Ann. N.Y. Acad. Sci. 271: 116.

Hansch, C. and A.J. Leo. 1979. Substituent Constants for Correlation Analysis in Chemistry and Biology. Wiley-Interscience, New York.

Harrison, R.M., et al. 1975. Polynuclear aromatic hydrocarbons in raw, potable, and waste waters. Water Res. 9: 331.

Harrison, R.M., et al. 1976. Effect of water chlorination upon levels of some polynuclear aromatic hydrocarbons in water. *Environ. Sci. Technol.* 12: 1151.

Hartwell, J.L. and P. Shubik. 1951. Survey of compounds which have been treated for carcinogenic activity. *Pub. Health Ser. Publ.* 149 (2nd ed.). U.S. Govern. Print. Off., Washington, D.C.

Hecht, S.S., et al. 1976. On the Structure and Carcinogenicity of the Methylchrysenes. In: R.I. Freudentahl and P.W. Jones (eds.), *Polynuclear Aromatic Hydrocarbons: Chemistry, Metabolism, and Carcinogenesis (Carcinogenesis - A Comprehensive Survey: Vol. 1)*. Raven Press, New York 1: 325.

Heidelberger, C. 1973. Chemical oncogenesis in culture. *Adv. Cancer Res.* 18: 317.

Heidelberger, C. 1975. Chemical carcinogenesis. *Ann. Rev. Biochem.* 44: 79.

Heidelberger, C. and P.R. Boshell. 1975. Chemical oncogenesis in cultures. *Gann Monogr. Cancer Res.* 17: 39.

Heidelberger, C. and S.M. Weiss. 1951. The distribution of radioactivity in mice following administration of 3,4-benzpyrene-5C¹⁴ and 1,2,5,6-dibenzanthracene-9,10-C¹⁴. *Cancer Res.* 11: 885.

Hellstrom, K.E. 1959. Chromosome studies on primary methylcholanthrene-induced sarcomas in the mouse. Jour. Natl. Cancer Inst. 23: 1019.

Henry, M.C., et al. 1973. Respiratory tract tumors in hamsters induced by benzo(a)pyrene. Cancer Res. 33: 1585.

Henry, M.C., et al. 1975. Importance of physical properties of benzo(a)pyrene-ferric oxide mixtures in lung tumor induction. Cancer Res. 35: 207.

Henry, S.A., et al. 1931. The incidence of cancer of the bladder and prostate in certain occupations. Jour. Hyg. 31: 125.

Hetteche, H.O. 1971. Plant waxes as collectors of PCAH in the air of polluted areas. Staub. 31: 72.

Hirano, T., et al. 1974. Measurement of epidermoid carcinoma development induced in the lungs of rats by 3-methylcholanthrene containing beeswax pellets. Jour. Natl. Cancer Inst. 53: 1209.

Hoch-Ligeti, C. 1941. Studies on the changes in the lymphoid tissues of mice treated with carcinogenic and non-carcinogenic hydrocarbons. Cancer Res. 1: 484.

Hoffman, D. and E. Wynder. 1976. Respiratory Carcinogenesis. In: C.E. Searle (ed.), Chemical Carcinogens. ACS Monogr. 173. Am. Chem. Soc. Washington, D.C.

Hoffman, D. and E.L. Wynder. 1977. Organic Particulate Pollutants - Chemical Analysis and Bioassays for Carcinogenicity. In: Stern (ed.), Air Pollution, 3rd ed. Academic Press, New York. 2: 361.

Howard, J.W. and T. Fazio. 1969. A review of polycyclic aromatic hydrocarbons in foods. Jour. Agric. Food Chem. 17: 527.

Howard, J.W., et al. 1966a. Extraction and estimation of PAH in smoked foods. Part I. General Method. Jour. Assoc. Off. Anal. Chem. 49: 595.

Howard, J.W., et al. 1966b. Extraction and estimation of polycyclic aromatic hydrocarbons in smoked foods. II. Benzo(a)pyrene. Jour. Assoc. Off. Anal. Chem. 49: 611.

Howard, J.W., et al. 1966c. Extraction and estimation of polycyclic aromatic hydrocarbons in vegetable oils. Jour. Assoc. Off. Anal. Chem. 49: 1236.

Howard, J.W., et al. 1968. Extraction and estimation of polycyclic aromatic hydrocarbons in total diet composites. Jour. Assoc. Off. Anal. Chem. 51: 122.

Huberman, E. and L. Sachs. 1974. Cell-mediated mutagenesis of mammalian cells with chemical carcinogens. Int. Jour. Cancer. 13: 326.

Huberman, E. and L. Sachs. 1976. Mutability of different genetic loci in mammalian cells by metabolically activated carcinogenic polycyclic hydrocarbons. Proc. Natl. Acad. Sci. 73: 188.

Huberman, E., et al. 1972. Transformation of hamster embryo cells by epoxides and other derivatives of polycyclic hydrocarbons. Cancer Res. 32: 1391.

Huberman, E., et al. 1976a. Identification of mutagenic metabolites of benzo(a)pyrene in mammalian cells. Proc. Natl. Acad. Sci. 73: 607.

Huberman, E., et al. 1976b. Mutagenesis and transformation of normal cells by chemical carcinogens. Nature. 264: 360.

Huberman, E., et al. 1977. Mutagenicity to mammalian cells in culture by (+) and (-) trans-7,8-dihydroxy-7,8-dihydro-benzo(a)pyrenes and the hydrolysis and reduction products of two stereo-isomeric benzo(a)pyrene 7,8-diol-9,10-epoxides. Cancer Lett. 4: 35.

Hueper, W.C. 1963. Chemically induced skin cancers in man. Natl. Cancer Inst. Monogr. 10: 377.

Hueper, W.C., et al. 1962. Carcinogenic bioassays on air pollutants. Arch. Pathol. 74: 89.

Iball, J. 1939. The relative potency of carcinogenic compounds. Am. Jour. Cancer. 35: 188.

International Agency for Research on Cancer. 1973. Certain Polycyclic Aromatic Hydrocarbons and Heterocyclic Compounds. Monographs on the evaluation of carcinogenic risk of the chemical to man. Vol. 3. Lyon, France.

International Commission on Radiological Protection. 1974. No. 23, Report of the Task Group on Reference Man. Pergamon Press, New York.

Jaffe, W. 1946. The influence of wheat germ oil on the production of tumors in rats by methylcholanthrene. *Exp. Med. Surg.* 4: 278.

Jerina, D.M. and J.W. Daly. 1974. Arene oxides: A new aspect of drug metabolism. *Science.* 185: 573.

Jerina, D.M., et al. 1972. Bay Region Epoxides of Dihydrodiols. A Concept which Explains the Mutagenic and Carcinogenic Activity of Benzo(a)pyrene and Benzo(a)anthracene. In: *Origins of Human Cancer.* Cold Spring Harbor Lab., Cold Spring Harbor, New York.

Jerina, D.M., et al. 1976. Mutagenicity of Benzo(a)pyrene Derivatives and the Description of a Quantum Mechanical Model which Predicts the Ease of Carbonium Ion Formation from Diol Epoxides. In vitro Metabolic Activation in Mutagenesis Testing. In: F.J. de Serres, et al. (eds.), Elsevier/North Holland Biomedical Press, Amsterdam. p. 159.

Joneja, M.G. and D.B. Coulson. 1973. Histopathology and cytogenetics of tumors induced by the application of 7,12-dimethylbenz(a)anthracene (DMBA) in mouse cervix. Eur. Jour. Cancer. 9: 367.

Joneja, M.G., et al. 1971. Cytogenetic studies on two types of 7,12-dimethylbenz(a)anthracene (DMBA) induced malignant tumors of mice. Anat. Rec. 969: 350.

Kapitulnik, J., et al. 1976a. Comparison of the hydroxylation of zoxazolamine and benzo(a)pyrene in human placenta: Effect of cigarette smoking. Clin. Pharmacol. Therap. 20: 557.

Kapitulnik, J., et al. 1976b. Lack of carcinogenicity of 4-,5-,6-, 7-,8-,9-, and 10-hydroxybenzo(a)pyrene on mouse skin. Cancer Res. 36: 3625.

Kapitulnik, J., et al. 1977a. Hydration of arene and alkene oxides by epoxide hydrase in human liver microsomes. Clin. Pharmacol. Therap. 21: 158.

Kapitulnik, J., et al. 1977b. Benzo(a)pyrene 7,8-dihydrodiol is more carcinogenic than benzo(a)pyrene in newborn mice. Nature. 266: 378.

Kapitulnik, J., et al. 1977c. Activation of monooxygenases in human liver by 7,8-benzoflavone. Clin. Pharmacol. Therap. 22: 475.

Kapitulnik, J., et al. 1977d. Comparative metabolism of benzo(a)-pyrene and drugs in human liver. Clin. Pharmacol. Therap. 21: 166.

Kapitulnik, J., et al. 1978a. Marked differences in the carcinogenic activity of optically pure (+) and (-)-trans-7,8-dihydroxy-7,8-dihydrobenzo(a)pyrene in newborn mice. Cancer Res. 38: 2661.

Kapitulnik, J., et al. 1978b. Tumorigenicity studies with diol-epoxides of benzo(a)pyrene which indicate that (+)-trans-7 β ,8 α -dihydroxy-9 α ,10 α -epoxy-7,8,9,10-tetrahydrobenzo(a)pyrene is an ultimate carcinogen in newborn mice. Cancer Res. 38: 354.

Kato, T., et al. 1975. Studies on experimental formation of ovarian tumors - especially, the discussion of the developing process of ovarian tumors following an application of DMBA. Kurume Med. Jour. 22: 169.

Kawai, M., et al. 1967. Epidemiologic study of occupational lung cancer. Arch. Environ. Health. 14: 859.

Keegan, R.E. 1971. The trace fluorometric determination of polynuclear aromatic hydrocarbons in natural water, Ph.D. Thesis, University of New Hampshire. Available from University Microfilms, Ann Arbor, Michigan.

- Kellerman, G., et al. 1973a. Aryl hydroxylase inducibility and bronchogenic carcinoma. *New England Jour. Med.* 289: 934.
- Kellerman, G., et al. 1973b. Genetic variation of aryl hydrocarbon hydroxylase in human lymphocytes. *Am. Jour. Human Genet.* 25: 327.
- Kennaway, E.L. 1925. The anatomical distribution of the occupational cancers. *Jour. Ind. Hyg.* 7: 69.
- Kennaway, E.L. and N.M. Kennaway. 1936. A study of the incidence of cancer of the lung and larynx. *Jour. Hyg.* 36: 236.
- Kennaway, E.L. and N.M. Kennaway. 1947. A further study of the incidence of cancer of the lung and larynx. *Br. Jour. Cancer.* 1: 260.
- Kertesz-Saringer, M. and Z. Morlin. 1975. On the occurrence of polycyclic aromatic hydrocarbons in the urban area of Budapest. *Atmos. Environ.* 9: 831.
- Kimura, T., et al. 1977. Differences in benzo(a)pyrene metabolism between lung and liver homogenates. *Biochem. Pharmacol.* 26: 671.
- Kobayashi, N. 1975. Production of respiratory tract tumors in hamsters by benzo(a)pyrene. *Gann.* 66: 311.

Kolar, L.R., et al. 1975. Contamination of soil, agricultural crops, and vegetables by 3,4-benzopyrene in the vicinity of CESKA BUDE JOVICE. Cesu Hyg. 20: 135.

Kotin, P., et al. 1954. Aromatic hydrocarbons. I. Presence in the Los Angeles atmosphere and the carcinogenicity of atmospheric extracts. Arch. Ind. Hyg. 9: 153.

Kotin, P., et al. 1969. Distribution, retention, and elimination of C¹⁴-3,4-benzopyrene after administration to mice and rats. Jour. Natl. Cancer Inst. 23: 541.

Kouri, R.E., et al. 1976. Studies on pulmonary aryl hydrocarbon hydroxylase activity in inbred strains of mice. Chem. Biol. Inter. 13: 317.

Krahn, D.B. and C. Heidelberger. 1977. Liver homogenate-mediated mutagenesis in Chinese hamster V79 cells by polycyclic aromatic hydrocarbons and aflatoxins. Mutat. Res. 46: 27.

Kraup, T. 1970. Oocyte survival in the mouse ovary after treatment with 9,10-dimethyl-1,2-benz(a)anthracene. Jour. Endocrinol. 46: 483.

Krstulovic, A.M., et al. 1977. Distribution of some atmospheric polynuclear aromatic hydrocarbons. Am. Lab. p. 11.

Kuratsune, M. 1956. Benzo(a)pyrene content in certain pyrogenic materials. Jour. Natl. Cancer Inst. 16: 1485.

Kuratsune, M. and W.C. Hueper. 1958. Polycyclic aromatic hydrocarbons in coffee soots. Jour. Natl. Cancer Inst. 20: 37.

Kuratsune, M. and W.C. Hueper. 1960. Polycyclic aromatic hydrocarbons in roasted coffee. Jour. Natl. Cancer Inst. 24: 463.

Kuroda, S. 1937. Occupational pulmonary cancer of generator gas workers. Ind. Med. Surg. 6: 304.

Landolph, J.R., et al. 1976. Quantitative studies of the toxicity of benzo(a)pyrene to a mouse liver epithelial cell strain in culture. Cancer Res. 26: 4143.

Lasnitzki, A. and D.L. Woodhouse. 1944. The effect of 1,2,5,6-dibenzanthracene on the lymph nodes of the rat. Jour. Anat. 78: 121.

Lasnitzki, I. 1963. Growth pattern of the mouse prostate gland in organ culture and its response to sex hormones, vitamin A, and 3-methylcholanthrene. Natl. Cancer Inst. Monogr. 12: 318.

Leber, P., et al. 1976. A Comparison of Benzo(a)pyrene Metabolism by Primates, Rats, and Miniature Swine. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: Chemistry, Metabolism and Carcinogenesis (Carcinogenesis - A Comprehensive Survey: Vol. 1). Raven Press, New York. 1: 35.

Lee, R.F., et al. 1972. Uptake, metabolism and discharge of polycyclic aromatic hydrocarbons by marine fish. *Mar. Biol.* 17: 201.

Lee, R.F., et al. 1976. Fate of petroleum hydrocarbons taken up from food and water by the blue crab, Calinectes Sapidus. *Marine Biol.* 37: 363.

Lehr, R.E., et al. 1978. The Bay Region Theory of Polycyclic Aromatic Hydrocarbon-induced Carcinogenicity. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: 2nd. Int. Symp. on Analysis, Chemistry, and Biology (Carcinogenesis - A Comprehensive Survey: Vol. 3). Raven Press, New York. 3: 231.

Leith, R.S. and L. Hayflick. 1974. Efforts to transform cultured normal human cells with polycyclic aromatic hydrocarbons. *Proc. Am. Assoc. Cancer Res.* 15: 86.

Levan, G. and A. Levan. 1975. Specific chromosome changes in malignancy: Studies in rat sarcomas induced by two polycyclic hydrocarbons. *Hereditas.* 79: 161.

Levin, W., et al. 1976a. Carcinogenicity of benzo(a)pyrene 4,5-, 7,8-, and 9,10-oxides on mouse skin. Proc. Natl. Acad. Sci. 73: 243.

Levin, W., et al. 1976b. (+)-Trans-7,8-dihydroxy-7,8-dihydrobenzo(a)pyrene: A potent skin carcinogen when applied topically to mice. Proc. Natl. Acad. Sci. 73: 3867.

Levin, W., et al. 1977a. Role of Purified Cytochrome P-448 and Epoxide Hydrase in the Activation and Detoxification of Benzo(a)pyrene. In: D.M. Jerina (ed.), ACS Symp. Ser. No. 44, Drug Metabolism Concepts. p. 99.

Levin, W., et al. 1977b. Marked differences in the tumor-initiating activity of optically pure (+)-and (-)-trans-7,8-dihydroxy-7,8-dihydrobenzo(a)pyrene on mouse skin. Cancer Res. 37: 2721.

Lijinsky, W. and A.E. Ross. 1967. Production of carcinogenic polynuclear hydrocarbons in the cooking of food. Food Cosmet. Toxicol. 5: 343.

Lijinsky, W. and P. Shubik. 1965a. The detection of polycyclic aromatic hydrocarbons in liquid smoke and some foods. Toxicol. Appl. Pharmacol. 7: 337.

Lijinsky, W. and P. Shubik. 1965b. PH carcinogens in cooked meat and smoked food. Ind. Med. Surg. 34: 152.

Lindsay, D.W., et al. 1974. The Bioassay of Carcinogenesis: Effects on the Epithelial Cell Complement of Rat Trachea Maintained in vitro. In: Experimental Lung Cancer: Carcinogenesis and Bioassays. Int. Symp. p. 521.

Lo, M. and E. Sandi. 1978. Polycyclic Aromatic Hydrocarbons (polynuclears) in Foods. In: F.A. Gunther and J.D. Gunther (eds.), Residue Reviews. Springer-Verlag. 69: 34.

Lu, A.Y.H., et al. 1976. Metabolism of Benzo(a)pyridine by Purified Liver Microsomal Cytochrome P-448 and Epoxide Hydrase. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: Chemistry, Metabolism, and Carcinogenesis (Carcinogenesis - A Comprehensive Survey: Vol. 1). Raven Press, New York. 1: 115.

Lu, A.Y.H., et al. 1978. Enzymological Properties of Purified Liver Microsomal Cytochrome P-450 System and Epoxide Hydrase. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: 2nd. Int. Symp. on Analysis, Chemistry, and Biology (Carcinogenesis - A Comprehensive Survey: Vol. 3). Raven Press, New York. 3: 243.

Lu, P.Y., et al. 1977. The environmental fate of three carcinogens: Benzo(a)pyrene, benzidine, and vinyl chloride evaluated in laboratory model ecosystems. Arch. Environ. Contam. Toxicol. 6: 129.

Lunde, G. and Bjørseth. 1977. Polycyclic aromatic hydrocarbons in long-range transported aerosols. *Nature*. 268: 518.

Maher, V.M., et al. 1977. Effect of DNA repair on the cytotoxicity and mutagenicity of polycyclic hydrocarbon derivatives in normal and xeroderma pigmentosum human fibroblasts. *Mutat. Res.* 43: 117.

Malanoski, A.J., et al. 1968. Survey of polycyclic aromatic hydrocarbons in smoked foods. *Jour. Assoc. Off. Anal. Chem.* 51: 114.

Malaveille, C., et al. 1975. Mutagenicity of non-K-region diols and diol-epoxides of benz(a)anthracene and benzo(a)pyrene in S. typhimurium TA 100. *Biochem. Biophys. Res. Comm.* 66: 693.

Malmgren, R.A., et al. 1952. Reduced antibody titres in mice treated with carcinogenic and cancer chemotherapeutic agents. *Proc. Soc. Exp. Biol. Med.* 79: 484.

Mantel, N. and R.W. Bryan. 1961. Safety testing of carcinogenic agents. *Jour. Natl. Cancer Inst.* 27: 455.

Marquardt, H. 1976. Microsomal Metabolism of Chemical Carcinogens in Animals and Man. In: R. Montesano and L. Tomatis (eds.), *Screening Tests in Chemical Carcinogenesis*. Intl. Agency Res. Cancer. IARC Publ. No 12. Lyon, France. p. 309.

Marquardt, H., et al. 1972. Malignant transformation of cells derived from mouse prostate by epoxides and other derivatives of polycyclic hydrocarbons. *Cancer Res.* 32: 716.

Marquardt, H., et al. 1974. Malignant transformation in vitro of mouse fibroblasts by 7,12-dimethylbenz(a)anthracene and 7-hydroxymethylbenz(a)anthracene and by their K-region derivatives. *Int. Jour. Cancer.* 13: 304.

Martin, R.J. and R.E. Duggan. 1968. Pesticide residues in total diet samples (III). *Pestic. Monitor. Jour.* 1: 111.

Masuda, Y. and M. Kuratsune. 1971. Polycyclic aromatic hydrocarbons in smoked fish. *Katsuobuski, GANN.* 62: 27.

Mattison, D.R. and S.S. Thorgeirsson. 1977. Ovarian metabolism of polycyclic aromatic hydrocarbons and associated ovotoxicity in the mouse. *Gynecol. Invest.* 8: 11.

Mazumdar, S., et al. 1975. An epidemiological study of exposure to coal tar pitch volatiles among coke oven workers. *Air Pollut. Control Assoc. Jour.* 25: 382.

McCann, J. and B.N. Ames. 1976. Detection of carcinogens as mutagens in the Salmonella/microsome test: Assay of 300 chemicals: Discussion. *Proc. Natl. Acad. Sci.* 73: 950.

McCann, J., et al. 1975. Detection of carcinogens as mutagens in the Salmonella/microsome test: Assay of 300 chemicals. Proc. Natl. Acad. Sci. 72: 5135.

Miller, E.C. 1978. Some current perspectives on chemical carcinogenesis in humans and experimental animals: Presidential address. Cancer Res. 38: 1479.

Mitelman, F. and G. Levin. 1972. The chromosomes of primary 7,12-dimethyl(a)anthracene-induced rat sarcomas. Hereditas. 71: 325.

Mitelman, F., et al. 1972. Chromosomes of six primary sarcomas induced in the Chinese hamster by 7,12-dimethylbenz(a)anthracene. Hereditas. 72: 311.

National Academy of Sciences. 1972. Biological effects of atmospheric pollutants: Particulate polycyclic organic matter. Washington, D.C.

Neal, J. and R.H. Rigdon. 1967. Gastric tumors in mice fed benzo(a)pyrene: A quantitative study. Texas Rep. Biol. Med. 25: 553.

Nebert, D.W. and J.S. Felton. 1976. Importance of genetic factors influencing the metabolism of foreign compounds. Fed. Proc. 35: 1133.

Nery, R. 1976. Carcinogenic mechanisms: A critical review and a suggestion that oncogenesis may be adaptive ontogenesis. Chem. Biol. Inter. 12: 145.

Nesnow, S. and C. Heidelberger. 1976. The effect of modifiers of microsomal enzymes on chemical oncogenesis in cultures of C3H mouse cell lines. Cancer Res. 36: 1801.

Nettesheim, P. and M.L. Williams. 1976. The influence of vitamin A on the susceptibility of the rat lung to 3-methylcholanthrene. Int. Jour. Cancer. 17: 351.

Nettesheim, P., et al. 1975. Effect of vitamin A on lung tumor induction in rats. Proc. Am. Assoc. Cancer Res. 16: 54.

Newbold, R.F. and P. Brooks. 1976. Exceptional mutagenicity of a benzo(a)pyrene diol epoxide in cultured mammalian cells. Nature. 261: 52.

Newbold, R.F., et al. 1977. Cell-mediated mutagenesis in cultured Chinese hamster cells by carcinogenic polycyclic hydrocarbons: Nature and extent of the associated hydrocarbon-DNA reaction. Mutat. Res. 43: 101.

Nishimura, K. and M. Masuda. 1971. Minor constituents of whisky fusel oils. I. Basic, phenolic and lactonic compounds. Jour. Food Sci. 36: 819.

Nowell, P.C. and D.A. Hungerford. 1960. Chromosome studies in normal and leukemic human leucocytes. Jour. Natl. Cancer Inst. 25: 85.

Ottonen, P.O. and J.K. Ball. 1973. Lack of correlation between gross chromosome abnormalities and carcinogenesis with 7,12-dimethylbenz(a)anthracene. Jour. Natl. Cancer Inst. 50: 497.

Owens, I.S. 1977. Genetic regulation of UDP-glucuronosyltransferase induction by polycyclic aromatic compounds in mice. Jour. Biol. Chem. 252: 2827.

Paigen, B., et al. 1978. Human Aryl Hydrocarbon Hydroxylase and Cancer Risk. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: 2nd. Int. Symp. on Analysis, Chemistry, and Biology (Carcinogenesis - A Comprehensive Survey: Vol. 3). Raven Press, New York. 3: 429.

Panalaks, T. 1976. Determination and identification of polycyclic aromatic hydrocarbons in smoked and charcoal-broiled food products by high pressure liquid chromatography. Jour. Environ. Sci. Health. 11: 299.

Payer, H.D., et al. 1975. Accumulation of polycyclic aromatic hydrocarbons in cultivated microalge. Naturwiss. 62: 536.

Payne, W.W. and W.C. Hueper. 1960. The carcinogenic effects of single and repeated doses of BP. Am. Ind. Hyg. Assoc. Jour. 21: 350.

Peacock, P.R. 1936. Evidence regarding the mechanism of elimination of 1,2-benzpyrene, 1,2,5,6-dibenzanthracene, and anthracene from the blood-stream of injected animals. Br. Jour. Exptl. Pathol. 17: 164.

Pelkonen, O. 1976. Metabolism of Benzo(a)pyrene in Human Adult and Fetal Tissues. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: Chemistry, Metabolism, and Carcinogenesis (Carcinogenesis - A Comprehensive Survey: Vol. 1). Raven Press, New York. 1: 9.

Pfeiffer, E.H. 1973. Investigations on the carcinogenic burden by air pollution in man. VII. Studies on the oncogenetic interaction of polycyclic aromatic hydrocarbons. Abl. Bakt. Hyg., I. Abt. Orig. B. 158: 69.

Pfeiffer, E.H. 1977. Oncogenic Interaction of Carcinogenic and Noncarcinogenic Polycyclic Aromatic Hydrocarbons in Mice. In: V. Mohr, et al. (eds.), Air Pollution and Cancer in Man. Intl. Agency Res. Cancer. Sci. Publ. No. 16. p. 69.

Philips, F.S., et al. 1973. In vivo Cytotoxicity of Polycyclic Hydrocarbons. In: Pharmacology and the Future of Man. Proc. 5th Intl. Congr. Pharmacol., 1972, San Francisco. 2: 75.

Pienta, R.J., et al. 1977. III. Morphological transformation of early passage golden Syrian hamster embryo cells derived from cryo-preserved primary cultures as a reliable in vitro bioassay for identifying diverse carcinogens. Intl. Jour. Cancer. 19: 642.

Popescu, N.C., et al. 1976. Chromosome patterns (G and C bands) of in vitro chemical carcinogen-transformed guinea pig cells. Cancer Res. 36: 1404.

Pullman, A. and B. Pullman. 1955. Electronic structure and carcinogenic activity of aromatic molecules - new developments. Adv. Cancer Res. 3: 117.

Pylev, L.N. 1962. Induction of lung cancer in rats by intratracheal insufflation of cancerogenic hydrocarbons. Acta Un. Int. Cancer. 19: 688.

Radding, S.G., et al. 1976. The environmental fate of selected polynuclear aromatic hydrocarbons. Prepared by Stanford Research Institute, Menlo Park, California, under Contract No. 68-01-2681. Publ. No. EPA-560/5-750-009. U.S. Environ. Prot. Agency, Washington, D.C.

Rahimtula, A.D., et al. 1977. The effects of antioxidants of the metabolism and mutagenicity of benzo(a)pyrene in vitro. Biochem. Jour. 164: 473.

Redmond, C.K., et al. 1972. Long term mortality study of steelworkers. Jour. Occup. Med. 14: 621.

Redmond, C.K., et al. 1976. Cancer experience among coke by-product workers. Ann. N.Y. Acad. Sci. p. 102.

Rees, E.O., et al. 1971. A study of the mechanism of intestinal absorption of benzo(a)pyrene. Biochem. Biophys. Act. 225: 96.

Regan, J.D., et al. 1978. Repair of DNA damage by mutagenic metabolites of benzo(a)pyrene in human cells. Chem. Biol. Inter. 20: 279.

Reid, D.D. and C. Buck. 1956. Cancer in coking plant workers. Br. Jour. Ind. Med. 13: 265.

Reznik-Schuller, H. and U. Mohr. 1974. Investigations on the carcinogenic burden by air pollution in man. IX. Early pathological alterations of the bronchial epithelium in Syrian golden hamsters after intratracheal instillation of benzo(a)pyrene. Zbl. Bakt. Hyg., I. Abt. Orig. B. 159: 493.

Rhee, K.S. and L.J. Bratzler. 1970. Benzo(a)pyrene in smoked meat products. Jour. Food Sci. 35: 146.

Rhim, J.S., et al. 1975. Transformation of human osteosarcoma cells by a chemical carcinogen. Jour. Natl. Cancer Inst. 55: 1291.

Rigdon, R.H. and J. Neal. 1965. Effects of feeding benzo(a)pyrene on fertility, embryos, and young mice. Jour. Natl. Cancer Inst. 34: 297.

Rigdon, R.H. and E.G. Rennels. 1964. Effect of feeding benzpyrene on reproduction in the rat. Experientia. 20: 1291.

Riley, J.F. 1969. Mast cells. Co-carcinogenesis and anti-carcinogenesis in the skin of mice. Experientia. 4: 1237.

Rüdiger, H., et al. 1976. Benzpyrene induces sister chromatid exchanges in cultured human lymphocytes. Nature. 262: 290.

Saffiotti, U., et al. 1968. A method for the experimental induction of bronchogenic carcinoma. Cancer Res. 28: 104.

Saffiotti, U., et al. 1972. Respiratory tract carcinogenesis induced in hamsters by different dose levels of benzo(a)pyrene and ferric oxide. Jour. Natl. Cancer Inst. 49: 1199.

San, R.H.C. and H.F. Stich. 1975. DNA repair synthesis of cultured human cells as a rapid bioassay for chemical carcinogens. Int. Jour. Cancer. 16: 284.

Santodonato, J., et al. 1978. Health assessment document for polycyclic organic matters. U.S. Environ. Prot. Agency, Washington, D.C.

Santodonato, J., et al. 1980. Multimedia health assessment document for polycyclic organic matter. Jour. Environ. Pathol. Toxicol. (In press).

Sawicki, E. 1962. Analysis of airborne particulate hydrocarbons: Their relative proportions as affected by different types of pollution. Natl. Cancer Inst. Monogr. No. 9: 201.

Sawicki, E., et al. 1962. Polynuclear aromatic hydrocarbon composition of the atmosphere in some large American cities. Am. Ind. Hyg. Assoc. Jour. 23: 137.

Schlede, E., et al. 1970a. Effect of enzyme induction on the metabolism and tissue distribution of benzo(a)pyrene. Cancer Res. 30: 2893.

Schlede, E., et al. 1970b. Stimulatory effect of benzo(a)pyrene and phenobarbital pretreatment on the biliary excretion of benzo(a)pyrene metabolites in the rat. Cancer Res. 30: 2898.

Schmahl, D., et al. 1977. Syncarcinogenic Action of Polycyclic Hydrocarbons in Automobile Exhaust Gas Condensates. In: V. Mohr, et al. (eds.), Air Pollution and Cancer in Man. Intl. Agency Res. Cancer Sci. Publ. No. 16. p. 53.

Schmeltz, I., et al. 1978. Bioassays of Naphthalene and Lkyl-naphthalenes for Co-carcinogenic Activity. Relation to Tobacco Carcinogenesis. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: 2nd. Int. Symp. on Analysis, Chemistry, and Biology (Carcinogenesis - A Comprehensive Survey: Vol. 3). Raven Press, New York. 3: 47.

Schönwald, A.D., et al. 1977. Benzpyrene-induced sister chromatid exchanges in lymphocytes of patients with lung cancer. Human Genet. 36: 361.

Selkirk, J.K., et al. 1971. An epoxide is an intermediate in the microsomal metabolism of the chemical carcinogen, dibenz(a,h)-anthracene. Biochem. Biophys. Res. Comm. 43: 1010.

Selkirk, J.K., et al. 1974. High-pressure liquid chromatographic analysis of benzo(a)pyrene metabolism and covalent binding and the mechanism of action of 7,8-benzoflavone and 1,2-epoxy-3,3,3-trichloropropane. Cancer Res. 34: 3474.

Selkirk, J.K., et al. 1975a. Isolation by high-pressure liquid chromatography and characterization of benzo(a)pyrene-4,5-epoxide as a metabolite of benzo(a)pyrene. Arch. Biochem. Biophys. 168: 322.

Selkirk, J.K., et al. 1975b. In vitro metabolism of benzo(a)pyrene by human liver microsomes and lymphocytes. Cancer Res. 35: 3651.

Selkirk, J.K., et al. 1976. Analysis of Benzo(a)pyrene Metabolism in Human Liver and Lymphocytes and Kinetic Analysis of Benzo(a)pyrene in Rat Liver Microsomes. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: Chemistry, Metabolism, and Carcinogenesis (Carcinogenesis - A Comprehensive Survey: Vol. 1). Raven Press, New York. 1: 153.

Sellakumar, A., et al. 1976. Effects of different dusts of respiratory carcinogenesis in hamsters induced by benzo(a)pyrene and diethylnitrosamine. Eur. Jour. Cancer. 12: 313.

Shabad, L.M. and A.P. Il'nitskii. 1970. Perspective on the problem of carcinogenic pollution in water bodies. Gig. Sanit. 35: 84; Hyg. Sanit. 35: 268. (Russian, English Transt.).

Shabad, L.M. and G.A. Smirnov. 1972. Aircraft engines as a source of carcinogenic pollution of the environment (benzo(a)pyrene studies). Atmos. Environ. 6: 153.

Shabad, L.M., et al. 1974. Transplacental and direct action of benzo(a)pyrene studied in organ cultures of embryonic lung tissue. Neoplasma. 22: 113.

Shamberger, R.J. 1970. Relationship of selenium to cancer. I. Inhibitory effect of selenium on carcinogenesis. Jour. Natl. Cancer Inst. 44: 931.

Shamberger, R.J. 1972. Increase of peroxidation in carcinogenesis. Jour. Natl. Cancer Inst. 48: 1491.

Shamberger, R.J. and G. Rudolph. 1966. Protection against cocarcinogenesis by antioxidants. Experientia. 22: 116.

Shendrikova, I.A. and V.A. Aleksandrov. 1974. Comparative characteristics of penetration of polycyclic hydrocarbons through the placenta into the fetus in rats. Byull. Eksperiment. Biol. i Medit. 77: 169.

Shimkin, M.B. and G.D. Stoner. 1975. Lung Tumors in Mice: Application to Carcinogenesis Bioassay. In: G. Klein and S. Weinhouse (eds.), Advances in Cancer Research. Raven Press, New York. 12: 1.

Shiraishi, Y., et al. 1973. Determination of polycyclic aromatic hydrocarbons in foods. II. 3,4-Benzopyrene in Japanese foods. Jour. Food Hyg. Soc. Japan, Shokuhin Eiseigaku Zasshi. 14: 173.

Shiraishi, Y., et al. 1974. Determination of polycyclic aromatic hydrocarbons in foods. III. 3,4-benzopyrene in vegetables. Jour. Food Hyg. Soc. Japan. 15: 18.

Shiraishi, Y., et al. 1975. Determination of polycyclic aromatic hydrocarbons in foods. IV. 3,4-benzopyrene in fish and shellfish. Jour. Food Hyg. Soc. Japan, Shokuhin Eiseigaku Zasshi. 16: 178.

Shubik, P. and J.L. Hartwell. 1957. Survey of compounds which have been tested for carcinogenic activity. Supplement 1. Pub. Health Serv. Publ. 149-1. U.S. Govern. Print. Off., Washington, D.C. p. 388.

Shubik, P. and J.L. Hartwell. 1969. Survey of compounds which have been tested for carcinogenic activity. Supplement 2. Pub. Health Serv. Publ. 149-2. U.S. Govern. Print. Off., Washington, D.C. p. 655.

Shulte-Herman, R. 1977. Stimulation of Liver Growth and Mixed-function Oxidase by alpha-Hexachlorocyclohexane: Separation of Inductive Pathways. In: V. Ulrich (ed.), Microsomes and drug oxidations. Pergamon Press, New York.

Siddiqui, I. and K.H. Wagner. 1972. Determination of 3,4-benzopyrene and 3,4-benzo-fluoranthene in rain water, ground water, and wheat. Chemosphere. 1: 83.

Simon, S., et al. 1969. Effect of cellulose casing on absorption of polycyclic hydrocarbons in wood smoke by absorbents. Jour. Agric. Food Chem. 17: 1128.

Sims, P. 1970. Qualitative and quantitative studies on the metabolism of a series of aromatic hydrocarbons by rat-liver preparations. Biochem. Pharmacol. 19: 795.

Sims, P. 1976. The Metabolism of Polycyclic Hydrocarbons to Dihydrodiols and Diol Epoxides by Human and Animal Tissues. In: R. Montesano, et al. (eds.), Screening Tests in Chemical Carcinogenesis. IARC Publ. No. 12. Lyon, France. p. 211.

Sims, P. and P.L. Grover. 1974. Epoxides in polycyclic aromatic hydrocarbon metabolism and carcinogenesis. Adv. Cancer Res. 20: 165.

Slaga, T.J., et al. 1976. Skin tumor initiating ability of benzo(a)pyrene 4,5-7,8- and 7,8-diol-9,10-epoxides and 7,8-diol. Cancer Lett. 2: 115.

Slaga, T.J., et al. 1977. Comparison of the tumor-initiating activities of benzo(a)pyrene arene oxides and dio-epoxides. Cancer Res. 37: 4130.

Smith, D.M., et al. 1975. Vitamin A and benzo(a)pyrene carcinogenesis in the respiratory tract of hamsters fed a semi-synthetic diet. Cancer Res. 35: 1483.

Smith, I.A., et al. 1978. Relationships between carcinogenicity and theoretical reactivity indices in polycyclic aromatic hydrocarbons. Cancer Res. 38: 2968.

Smyth, H.F., et al. 1962. Range-finding toxicity data: List VI. Am. Ind. Hyg. Assoc. Jour. 23: 95.

Snell, K.C. and H.L. Stewart. 1962. Induction of pulmonary adenomatosis in DBA/2 mice by the oral administration of dibenz(a,h)-anthracene. Acta. Un. Int. Canc. 19: 692.

Stenbäck, F. and A. Sellakumar. 1974a. Lung tumor induction by dibenzo(a,i)pyrene in the Syrian golden hamster. Z. Krebsforsch. 82: 175.

Stenbäck, F. and A. Sellakumar. 1974b. Squamous metaplasia and respiratory tumors induced by intratracheal instillations of 7,12-dimethylbenz(a)anthracene in Syrian golden hamsters. Eur. Jour. Cancer. 10: 483.

Stephan, C.E. 1980. Memorandum to J. Stara. U.S. EPA. July 3.

Stich, H.F. and B.A. Laishes. 1973. DNA repair and chemical carcinogens. In: H.L. Ioachim, (ed.). Pathobiology Ann. 3: 341.

Stich, H.F., et al. 1975. The search for relevant short term bioassays for chemical carcinogens: The tribulation of modern Sisyphus. Can. Jour. Genet. Cytol. 17: 471.

Stich, H.F., et al. 1976. DNA Fragmentation and DNA Repair as on in vitro and in vivo Assay for Chemical Procarcinogens, Carcinogens, and Carcinogenic Nitrosation Products. In: R. Montesano, et al. (eds.), IARC Scien. Publ. No. 12, Screening Tests in Chemical Carcinogenesis. Lyon, France. p. 15.

Stjernsward, J. 1966. The effect of non-carcinogenic and carcinogenic hydrocarbons on antibody-forming cells measured at the cellular level in vitro. Jour. Natl. Cancer Inst. 36: 1189.

Stjernsward, J. 1969. Immunosuppression by carcinogens. Anti-biot. Chemother. 15: 213.

Stocks, P. 1947. Regional and local differences in cancer death rates. Studies on medical and population subjects, No. 1. Gen. Regis. Off., London.

Stoming, T.A., et al. 1977. The metabolism of 3-methyl-cholanthrene by rat liver microsomes - A reinvestigation. Biochem. Biophys. Res. Comm. 79: 461.

Sugimura, T., et al. 1976. Overlapping of Carcinogens and Mutagens. In: P.N. Magee (ed.), Fundamentals in Cancer Prevention. Univ. of Tokyo Press, Tokyo/Univ. Park Press, Baltimore. p. 191.

Sullivan, P.D., et al. 1978. Effect of Antioxidants on Benzo(a)-pyrene Free Radicals. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: 2nd Int. Symp. on Analysis, Chemistry, and Biology (Carcinogenesis - A Comprehensive Survey: Vol. 3). Raven Press, New York. 3: 1

Swallow, W.H. 1976. Survey of polycyclic aromatic hydrocarbons in selected foods and food additives available in New Zealand. New Zealand Jour. Sci. 19: 407.

Swenberg, J.A., et al. 1976. In vitro DNA damage/alkaline elution assay for predicting carcinogenic potential. Biochem. Biophys. Res. Comm. 72: 738.

Teranishi, K., et al. 1975. Quantitative relationship between carcinogenicity and mutagenicity of polyaromatic hydrocarbons in Salmonella typhimurium mutants. Mutat. Res. 31: 97.

Thakker, D.R., et al. 1976. Metabolism of benzo(a)pyrene: Conversion of (+)-trans-7,9-dihydroxy-7,8-dihydrobenzo(a)pyrene to highly mutagenic 7,8-diol-9,10-epoxides. Proc. Natl. Acad. Sci. 73: 3381.

Thakker, D.R., et al. 1977. Metabolism of benzo(a)pyrene. VI. Stereo-selective metabolism of benzo(a)pyrene and benzo(a)pyrene 7,8-dihydrodiol to diol epoxides. Chem. Biol. Inter. 16: 281.

Thakker, D.R., et al. 1978. Metabolism of 3-methylcholanthrene by Rat Liver Microsomes and a Highly Purified Monooxygenase System with and without Epoxide Hydrase. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: 2nd. Int. Symp. on Analysis, Chemistry, and Biology (Carcinogenesis - A Comprehensive Survey: Vol. 3). Raven Press, New York. 3: 253.

Thomson, S. and T.J. Slaga. 1976. Mouse epidermal aryl hydrocarbon hydroxylase. Jour. Invest. Dermatol. 66: 108.

Thorsteinsson, T. 1969. Polycyclic hydrocarbons in commercially and home-smoked food in Iceland. *Cancer*. 23: 455.

Tokiwa, H., et al. 1977. Detection of mutagenic activity in particulate air pollutants. *Mutat. Res.* 48: 237.

Toth, L. and W. Blass. 1972. Einfluss der Raeuchertechnologie auf den Gehalt von Geraeucherten Fleischwaren an Cancerogenen Kohlenwasserstoffen. *Fleischwirt.* 21: 1121.

Tracor Jitco, Inc. 1974. Survey of compounds which have been tested for carcinogenic activity (1970-1971). *Pub. Health Serv. Publ.* 149. U.S. Govern. Print. Off., Washington, D.C. p. 1667.

4

Tracor Jitco, Inc. 1976. Survey of compounds which have been tested for carcinogenic activity (1972-1973). *Pub. Health Serv. Publ.* 149. U.S. Govern. Print. Off., Washington, D.C. p. 1638.

Tromp, S.W. 1955. Possible effects of geophysical and geochemical factors on development and geographic distribution of cancer. *Schweiz. Z. Path.* 18: 929.

U.S. EPA. 1974. Special report: Trends in concentrations of benzene-soluble suspended particulate fraction and benzo(a)pyrene. *Publ. No.* EPA-450/2-74-022, Res. Triangle Park, North Carolina.

U.S. EPA. 1975. Scientific and technical assessment report on particulate polycyclic organic matter (PPOM), Publ. No. EPA-600/6-75-001, Washington, D.C.

U.S. EPA. 1977. National Organic Monitoring Survey, Tech. Support Div., Off. Water Supply, Cincinnati, Ohio.

U.S. EPA. 1980. Seafood consumption data analysis. Stanford Research Institute International, Menlo Park, California. Final report, Task 11, Contract No. 68-01-3887.

Vainio, H., et al. 1976. The fate of intratracheally installed benzo(a)pyrene in the isolated perfused rat lung of both control and 20-methylcholanthrene pretreated rats. Res. Comm. Chem. Pathol. Pharmacol. 13: 259.

Van Duuren, B.L. 1969. Tumor-promoting agents in two-stage carcinogenesis. Prog. Exp. Tumor Res. 11: 31.

Van Duuren, B.L. 1976. Tumor-promoting and Co-carcinogenic Agents in Chemical Carcinogenesis. In: C.E. Searle (ed.), Chemical Carcinogens. ACS Monogr. 172. Am. Chem. Soc., Washington, D.C. p. 24.

Van Duuren, B.L. and B.M. Goldschmidt. 1976. Co-carcinogenic and tumor-promoting agents in tobacco carcinogenesis. Jour. Natl. Cancer Inst. 56: 1237.

Van Duuren, B.L., et al. 1973. Brief communications: Co-carcinogenic agents in tobacco carcinogenesis. Jour. Natl. Cancer Inst. 51: 703.

Veith, G.D. 1980. Memorandum to C.E. Stephan. U.S. EPA. April 14.

Veith, G.D., et al. 1979. Measuring and estimating the bioconcentration factors of chemicals in fish. Jour. Fish. Res. Board Can. 36: 1040.

Vitzthum, O.G., et al. 1975. New volatile constituents of black tea aroma. Jour. Agric. Food Chem. 23: 999.

Wang, I.Y., et al. 1976. Enzyme Induction and the Difference in the Metabolite Patterns of Benzo(a)pyrene Produced by Various Strains of Mice. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: Chemistry, Metabolism, and Carcinogenesis (Carcinogenesis - A Comprehensive Survey: Vol. 1). Raven Press, New York. 1: 77.

Wattenberg, L.W. 1972. Inhibition of carcinogenic and toxic effects of polycyclic hydrocarbons by phenolic antioxidants. Jour. Natl. Cancer Inst. 48: 1425.

Wattenberg, L.W. 1973. Inhibition of chemical carcinogen-induced pulmonary neoplasia by butylated hydroxyanisole. Jour. Natl. Cancer Inst. 50: 1541.

Wattenberg, L.W. 1974. Inhibition of carcinogenic and toxic effects of polycyclic hydrocarbons by several sulfur-containing compounds. Jour. Natl. Cancer. Inst. 52: 1583.

Wattenberg, L.W. 1977. Inhibition of carcinogenic effects of polycyclic hydrocarbons by benzyl isothiocyanate and related compounds. Jour. Natl. Cancer Inst. 58: 395.

Wattenberg, L.W. and J.L. Leong. 1970. Inhibition of the carcinogenic action of benzo(a)pyrene by flavones. Cancer Res. 30: 1922.

Wattenberg, L.W., et al. 1976. Effects of antioxidants on metabolism of aromatic polycyclic hydrocarbons. Adv. Enzyme Regul. 14: 313.

Weber, R.P., et al. 1974. Effect of the organophosphate insecticide parathion and its active metabolite paraoxon on the metabolism of benzo(a)pyrene in the rat. Cancer Res. 34: 947.

Welch, R.M., et al. 1972. Effect of enzyme induction on the metabolism of benzo(a)pyrene and 3'-methyl-4-monomethyl-amino-azobenzene in the pregnant and fetal rat. Cancer Res. 32: 973.

White, R.H., et al. 1971. Determination of polycyclic aromatic hydrocarbons in liquid smoke flavors. Jour. Agric. Food Chem. 19: 143.

- Wiebel, F.J., et al. 1973. Aryl hydrocarbon [benzo(a)pyrene] hydroxylase: Inducible in extrahepatic tissues of mouse strains not inducible in liver. Arch. Biochem. Biophys. 154: 292.
- Wiebel, F.J., et al. 1975. Aryl hydrocarbon [benzo(a)pyrene] hydroxylase: A mixed-function oxygenase in mouse skin. Jour. Invest. Dermatol. 64: 184.
- Williams, G.M. 1976. Carcinogen-induced DNA repair in primary rat liver cells cultures; a possible screen for chemical carcinogens. Cancer Lett. 1: 231.
- Wislocki, P.G., et al. 1976a. High mutagenicity and toxicity of a diol epoxide derived from benzo(a)pyrene. Biochem. Biophys. Res. Comm. 68: 1006.
- Wislocki, P.G., et al. 1976b. Mutagenicity and cytotoxicity of benzo(a)pyrene arene oxides, phenols, quinones, and dihydrodiols in bacterial and mammalian cells. Cancer Res. 36: 3350.
- Wislocki, P.G., et al. 1977. High carcinogenicity of 2-hydroxy-benzo(a)pyrene on mouse skin. Cancer Res. 37: 2608.
- World Health Organization. 1970. European Standards for Drinking Water, 2nd ed., Revised, Geneva.

- Wood, A.W., et al. 1976a. Mutagenicity and cytotoxicity of benzo(a)pyrene benzo-ring epoxides. *Cancer Res.* 36: 3358.
- Wood, A.W., et al. 1976b. Metabolism of benzo(a)pyrene and benzo(a)pyrene derivatives to mutagenic products by highly purified hepatic microsomal enzymes. *Jour. Biol. Chem.* 251916: 4882.
- Wood, A.W., et al. 1977a. Differences in mutagenicity of the optical enantiomers of the diastereomeric benzo(a)pyrene 7,8-diol-9,10-epoxides. *Biochem. Biophys. Res. Comm.* 77: 1389.
- Wood, A.W., et al. 1977b. High mutagenicity of metabolically activated chrysene 1,2-dihydrodiol: Evidence for bay region activation of chrysene. *Biochem. Biophys. Res. Comm.* 78: 847.
- Wynder, E. and D. Hoffman. 1965. Some laboratory and epidemiological aspects of air pollution carcinogenesis. *Jour. Air Pollut. Contr. Assoc.* 15: 155.
- Wynne-Griffith, G. and R.I. Davies. 1954. Cancer and soils in the County of Anglesey - A revised method of comparison. *Br. Jour. Cancer.* 8: 594.
- Yang, S.K., et al. 1977. Metabolic activation of benzo(a)pyrene and binding to DNA in cultured human bronchus. *Cancer Res.* 37: 1210.

- Yang, S.K., et al. 1978. Benzo(a)pyrene Metabolism: Mechanism in the Formation of Epoxides, Phenols, Dihydrodiols, and the 7,8-diol-9,10-epoxides. In: R.I. Freudenthal and P.W. Jones (eds.), Polynuclear Aromatic Hydrocarbons: 2nd. Int. Symp. on Analysis, Chemistry, and Biology (Carcinogenesis - A Comprehensive Survey: Vol. 3). Raven Press, New York. 3: 285.
- Yasuhira, K. 1964. Damage to the thymus and other lymphoid tissues from 3-methyl-cholanthrene, and subsequent thymoma production, in mice. *Cancer Res.* 24: 558.
- Youngblood, W.W. and M. Blumer. 1975. Polycyclic aromatic hydrocarbons in the environment: Homologous series in soils and recent marine sediments. *Geochem. Cosmochim. Acta.* 39: 1303.
- Yuspa, S.H., et al. 1976. Cutaneous carcinogenesis: Past, present, and future. *Jour. Invest. Dermatol.* 67: 199.
- Zampaglione, N.G. and G.J. Mannering. 1973. Properties of benzpyrene hydroxylase in the liver, intestinal mucosa and adrenal of untreated and 3-methyl-cholanthrene treated rats. *Jour. Pharmacol. Exp. Ther.* 185: 676.
- Zitko, V. 1975. Aromatic hydrocarbons in aquatic fauna. *Bull. Environ. Contam. Toxicol.* 14: 621.

APPENDIX

Summary and Conclusion Regarding the Carcinogenicity of Polynuclear Aromatic Hydrocarbons (PAH)

Polynuclear aromatic hydrocarbons (PAH) comprise a diverse class of compounds consisting of substituted and unsubstituted polycyclic and heterocyclic aromatic rings. They are formed as a result of incomplete combustion of organic compounds and appear in food as well as ambient air and water.

Numerous studies of workers exposed to coal gas, coal tars, and coke oven emissions, all of which have large amounts of PAH, have demonstrated a positive association between the exposures and lung cancer.

Several PAH are well-known animal carcinogens, others are not carcinogenic alone but can enhance or inhibit the response of the carcinogenic PAH and some induce no tumors in experimental animals. Most of the information about the combined carcinogenic effects of several PAH come from skin painting and subcutaneous injection experiments in mice whereas oral administration, intratracheal instillation, and inhalation have been shown to induce carcinogenic responses to single compounds. In one subcutaneous injection study in mice it was shown that a combination of several noncarcinogenic PAH compounds, mixed according to the proportion occurring in auto exhaust, does not enhance or inhibit the action of two potent PAH carcinogens, benzo(a)pyrene (BaP) and dibenz(a,h)anthracene.

The mutagenicity of PAH in the Salmonella typhimurium assay correlates well with their carcinogenicity in animal systems. PAH compounds have damaged chromosomes in cytogenetic tests, have

induced mutations in mammalian cell culture systems and have induced DNA repair synthesis in human fibroblast cultures.

The water quality criterion for carcinogenic PAH compounds is based on the assumption that each compound is as potent as BaP and that the carcinogenic effect of the compounds is proportional to the sum of their concentrations. Based on an oral feeding study of BaP in mice, the concentration of BaP estimated to result in a lifetime risk of 10^{-5} is 28 ng/l. Therefore, with the assumption above, the sum of the concentrations of all carcinogenic PAH compounds should be less than 28 ng/l in order to keep the lifetime cancer risk below 10^{-5} .

Summary of Pertinent Data

The water quality criterion for BaP is based on the experiment reported by Neal and Rigdon (1967), in which benzo(a)pyrene at doses ranging between 1 and 250 ppm in the diet was fed to strain CFW mice for approximately 110 days. Stomach tumors, which were mostly squamous cell papillomas but some carcinomas, appeared with an incidence statistically higher than controls at several doses. The extrapolation was based on the following parameters:

<u>Dose</u> (mg/kg/day)	<u>Incidence^a</u> (No. responding/No. tested)
0.0	0/289
0.13	0/25
1.3	0/24
2.6	1/23
3.9	0/37
5.2	1/40
5.85	4/40
6.5	24/34
13.0	19/23
32.5	66/73

le = 110 days
 Le = 183 days
 L = 630 days

w = 0.034 kg
 R = 30 l/kg

With these parameters, the carcinogenic potency factor for humans, q_1^* , is $11.53 \text{ (mg/kg/day)}^{-1}$. The result is that the water concentration of BaP should be less than 28 ng/l in order to keep the individual lifetime risk below 10^{-5} . It is recognized that numerous carcinogenic PAH other than BaP are found in water. However, there is probably little need to derive criteria for all such PAH, since efforts to reduce BaP levels to within acceptable limits will result in the reduction of all PAH.

^aThe incidences at the highest three doses were not used in the extrapolation due to lack of fit of the multistage model. See the Human Health Methodology Appendices to the October 1980 Federal Register notice which announced the availability of this document for a discussion on the fit of data to the multistage model.