## IX. APPENDIX III

#### BIOLOGIC METHOD FOR SAMPLING

#### AND ANALYSIS OF BENZENE

The recommended biologic method for urinalysis is derived from Sherwood and Carter. [102] It has been designed to determine the concentration of phenol and its conjugates, sulfate and glucuronide, in urine. It also determines orthocresol and meta- and paracresols. Urine is hydrolyzed with perchloric acid at 95 C, and the phenols and cresols are extracted with isopropyl ether and determined by gas chromatography.

## Collection of Urine Samples

"Spot" urine specimens of about 100 ml are collected as close to the end of the working day as possible. If any worker's urine phenol level exceeds 75 mg/liter, procedures are instituted immediately to determine the cause of the elevated urine phenol levels and to reduce benzene exposure to the worker. Weekly specimens are collected as described above until 3 consecutive weekly determinations indicate that urinary phenol levels are below 75 mg/liter.

After thoroughly washing their hands with soap and water, workers shall collect urine samples from single voidings in clean, dry specimen containers having tight closures and at least a 120-ml capacity. Collection containers may be glass, waxcoated paper, or other disposable types if desired. Following collection of urine specimens, 1 ml of a 10% copper sulfate solution is added to each sample as a preservative, and samples are ammediately stored under refrigeration, preferably at 0-4 C.

Refrigerated specimens will remain stable for approximately 90 days. If shipment of samples is necessary to perform analyses, the most rapid method available shall be employed utilizing acceptable packing procedures as specified by the carrier. Proper identification of each specimen shall include as a minimum, the worker's name, date, and time of collection.

## **Analytical**

#### (a) Principle of the Method

Urine samples are treated with perchloric acid at 95 C to hydrolyze the phenol conjugates, phenyl sulfate, and phenyl glucuronide, formed as detoxification products following benzene absorption. The total phenol is extracted with disopropyl ether and the phenol concentration is determined by gas chromatography analysis of the disopropyl ether extract.

## (b) Apparatus

(1) Gas chromatograph with a flame ionization detector and equipped with a 5-foot x 3/16-inch column packed with 2 w/w polyethylene glycol adipate on universal 'B' support. Operating conditions are as follows:

Column temperature 150 C

Detector temperature 200 C

Injection port tempera-

ture 200 C

Carrier gas Nitrogen

Carrier gas flowrate 60 ml/min

- (2) Water bath
- (3) Glass-stoppered, 10-ml volumetric flasks

- (4) 1-ml, 2-ml, and 5-ml volumetric pipets
- (5)  $5-\mu 1$  syringe

#### (c) Reagents

- (1) Phenol
- (2) Perchloric acid
- (3) Diisopropyl ether
- (4) Distilled water

#### (d) Procedure

(1) Hydrolysis of Phenol Conjugates

Pipet 5 ml of urine into a 10-ml, glass-stoppered, volumetric flask. Add perchloric acid, mix by swirling, and transfer the lightly stoppered flask to a water bath at 95 C. After 2 hours, remove the flask from the water bath and allow to cool at room temperature.

(2) Diisopropyl ether extraction of phenol and cresols.

Pipet 1 ml of diisopropyl ether into the flask and adjust the volume to 10 ml with distilled water. Shake vigorously for 1 minute to extract the phenol and cresols. Allow the aqueous and ether layers to separate.

(3) Gas chromatographic analysis for phenol

Inject 5  $\mu$ l of the disopropyl ether layer into the gas chromatograph and record the attenuation and area of the phenol peak. Under the conditions described, phenol is eluted in 100 seconds, o-cresol in 130 seconds, and m- and p- cresols in 320 seconds.

#### (e) Standards Preparation

A 50 mg/liter standard aqueous solution of phenol is prepared. A 5-ml aliquot of the standard solution is then subjected to the hydrolysis, extraction, and gas chromatographic analysis procedures described under Procedure above.

#### (f) Calculations

Determine the phenol concentration in the urine by comparing the gas chromatographic peak area of the sample with that of the 50 mg/liter standard and adjust the value to a specific gravity of 1.024.

## (g) Specific Gravity Correction

Due to the magnitude of correction which is required, samples having uncorrected specific gravities less than 1.010 shall be rejected and another sample shall be obtained.

Based on a survey of a large population in the United States in connection with urinary lead excretion, Levine and Fahy [139] found the mean specific gravity to be 1.024. Many investigators throughout the world now use this figure. Buchwald [130] in 1964 determined the mean specific gravity for residents in the United Kingdom to be 1.016, a value now frequently used for Northern Europeans. The importance of specific gravity adjustments can be seen in that a specific gravity of 1.016 will give results having two-thirds the value of those corrected to 1.024. It is important, therefore, that a value be chosen for standardization; since greater acceptance seems to be for 1.024, this value has been selected for adjustment of urinary concentrations of benzene recommended for biological monitoring.

corrected concentration = 
$$\frac{\text{observed concentration x 24}}{\text{last 2 digits of sp gr (eg, 1.021)}}$$

#### X. APPENDIX IV

#### SPECIAL MEDICAL CONSIDERATIONS

The literature on the subject of benzene intoxication, both acute and chronic, has been reviewed elsewhere in this document. Levels of exposure permitted in the standards set by this document have been shown to reduce the danger of acute intoxications to a minimum. [1,24,23] Barring accidental exposure, the need for constant monitoring for signs and symptoms of acute intoxication is unnecessary. The toxic effects of chronic low level exposures are not as well documented and, as has been discussed, exposures to 40 ppm have caused hematologic changes in animals. [66] The need for constant and complete monitoring of the organ systems known to be affected by chronic benzene exposure is, therefore, prudent and necessary.

The hematologic system is especially singled out by benzene's toxic effects. There is no agreement in the literature as to which parameter of hematologic function is the first indicator of early benzene intoxication. Monitoring a number of components, therefore, becomes necessary.

The life span of the erythrocyte has been calculated by various methods to be approximately 120 days. [140] This means that if erythrocyte production were to stop suddenly, as in the development of aplastic anemia, 0.83% of the red cell mass would be lost daily. In the asymptomatic individual exposed to very low concentrations of benzene, measurements of the red cell mass could safely be done every 3 months. In workers exposed to higher concentrations, the risk of developing aplastic anemia increases, and more frequent determinations become necessary. In the event of red

cell agenesis, 2 weeks would be a sufficient time to reduce the red cell mass by 12%. A longer delay in discovering this condition would be deleterious to the prognosis; thus, monitoring the red cell mass in individuals with higher levels of exposure to benzene should be done at intervals not exceeding 2 weeks. Macrocytosis has also been stated to be the second most frequent toxic effect of benzene on the bone marrow [140]; therefore, bone marrow monitoring for macrocytosis by the measurement of appropriate corpuscular indices at the most frequent practical period is indicated.

No such simple means for estimating the decay of the white blood cell mass in the case of WBC agenesis is available because, to date, the life span of neutrophils has not been measured successfully, despite estimates of less than 12 days. [140] It is difficult to rationally set a maximum period beyond which it would be dangerous to delay measurement. Quarterly intervals in exposed individuals are felt to be maximum intervals prudent in this situation, reflecting the expense and difficulty of the differential WBC count, but measurement at shorter intervals is desirable where practical.

The life span of platelets has been variously estimated as from 9-12 days. These data are imprecise because of the difficulty inherent in the measurements. For those individuals exposed to greater than the maximum suggested TWA, a bimonthly measurement would seem sufficient to find a marked platelet reduction by estimation of platelets from a smear of peripheral blood. This finding might precede symptoms. However, by the time the abnormality is sufficiently advanced, the worker may already be complaining of symptoms caused by a decreased clotting function; therefore,

no test more frequently than quarterly is recommended for a platelet determination.

Increased turnover of erythrocytes, probably through hemolysis, has been reported. [140,141] Counts of reticulocytes (immature, still nucleate red blood cells) give a rough estimate of the rapidity of erythrocyte turnover. Obtaining this value on a quarterly basis is suggested in workers having exposures from 1-10 ppm of benzene and annually in others. Hemolysis is discovered early by laboratory estimation of the breakdown products of hemoglobin, of which bilirubin is the easiest to measure. Again, the frequency of the determination is predicated upon the level of individual exposure.

## Normal Hematologic Values

The generally accepted ranges of normal for the hematologic tests discussed in the body of this document are presented in Table XII-14 and are derived from values reported by Conn. [142] It should be noted that these values do not represent a definition of normal, but are only a rough guideline. Interpretation of laboratory results should be made on the basis of that laboratory's established normal range for the procedure as performed there. The values listed in Table XII-14 are applicable only to adults.

#### XI. APPENDIX V.

#### MATERIAL SAFETY DATA SHEET

The following items of information which are applicable to a specific product or material containing benzene shall be provided in the appropriate section of the Material Safety Data Sheet or approved form. If a specific item of information is inapplicable, the initials "n.a." (not applicable) should be inserted.

- (a) Section I. Source and Nomenclature,
- (1) The name, address, and telephone number of the manufacturer or supplier of the product.
- (2) The trade name and synonyms for a mixture of chemicals, a basic structural material, or for a process material; and the trade name and synonyms, chemical name and synonyms, chemical family, and formula for a single chemical.
  - (b) Section II. Hazardous Ingredients.
- (1) Chemical or widely recognized common name of all hazardous ingredients.
- (2) The approximate percentage by weight or volume (indicate basis) which each hazardous ingredient of the mixture bears to the whole mixture. This may be indicated as a range or maximum amount, ie, 10-20 by volume; 10% maximum by weight.
- (3) Basis for toxicity for each hazardous material such as an established standard in appropriate units.

(c) Section III. Physical Data.

Physical properties of the total product including boiling point and melting point in degrees Fahrenheit; vapor pressure in millimeters of mercury; vapor density of gas or vapor (air=1); solubility in water, in parts/hundred parts of water by weight; specific gravity (water=1); volatility, indicate if by weight or volume, at 70 degrees Fahrenheit; evaporation rate for liquids (indicate whether butyl acetate or ether=1); and appearance and odor.

(d) Section IV. Fire and Explosion Hazard Data.

Fire and explosion hazard data about a single chemical or a mixture of chemicals, including flash point, in degrees Fahrenheit; flammable limits in percent by volume in air; suitable extinguishing media or agents; special fire fighting procedures; and unusual fire and explosion hazard information.

(e) Section V. Health Hazard Data.

Toxic level for total compound or mixture, effects of exposure, and emergency and first-aid procedures.

(f) Section VI. Reactivity Data.

Chemical stability, incompatibility, hazardous decomposition products, and hazardous polymerization.

(g) Section VII. Spill or Leak Procedures.

Detailed procedures to be followed with emphasis on precautions to be taken in cleaning up and safe disposal of materials leaked or spilled. This includes proper labeling and disposal of containers holding residues, contaminated absorbents, etc.

(h) Section VIII. Special Protection Information.

Requirements for personal protective equipment, such as respirators, eye protection, clothing, and ventilation, such as local exhaust (at site of product use or application), general, or other special types.

(i) Section IX. Special Precautions.

Any other general precautionary information.

## U.S. DEPARTMENT OF LABOR Occupational Safety and Health Administration

Form Approved OMB No. 44-R1387

# MATERIAL SAFETY DATA SHEET

Required under USDL Safety and Health Regulations for Ship Repairing,

Shipbuilding, a	nd Sh	nipbreakin	g (29 CFR 1915, 1916, 1917)						
		SECT	ION I	······································					
MANUFACTURER'S NAME				EPHONE NO.					
ADDRESS (Number, Street, City, State, and ZIP Co	de)								
	· · · -		TRADE NAME AND SYNONY	MS					
CHEMICAL FAMILY			FORMULA						
SECTION	11 -	HAZAF	RDOUS INGREDIENTS	**************************************					
PAINTS, PRESERVATIVES, & SOLVENTS	%	TLV (Units)	ALLOYS AND METALLIC COATING	GS %	TLV (Units)				
PIGMENTS		(0.1110)	BASE METAL						
CATALYST			ALLOYS						
VEHICLE	EHICLE METALLIC COATINGS								
SOLVENTS			FILLER METAL PLUS COATING OR CORE FLUX						
ADDITIVES			OTHERS						
OTHERS									
HAZARDOUS MIXTURE	S OF	OTHER LI	QUIDS, SOLIDS, OR GASES	%	TLV (Units)				
	CTIO	N III - I	PHYSICAL DATA						
BOILING POINT (°F.)			SPECIFIC GRAVITY (H <sub>2</sub> O=1)						
VAPOR PRESSURE (mm Hg.)	$\perp$		PERCENT, VOLATILE BY VOLUME (%)						
VAPOR DENSITY (AIR=1)	<u> </u>		(=1)						
SOLUBILITY IN WATER									
APPEARANCE AND ODOR			······································	·					
SECTION IV -	FIR	E AND	EXPLOSION HAZARD DATA	<del> </del>					
ADDRESS (Number, Street, City, State, and ZIP Code)  CHEMICAL NAME AND SYNONYMS  CHEMICAL FAMILY  SECTION II - HAZARDOUS INGREDIENTS  PAINTS, PRESERVATIVES, & SOLVENTS		Uel							
EXTINGUISHING MEDIA									
SPECIAL FIRE FIGHTING PROCEDURES		****			·———				
UNUSUAL FIRE AND EXPLOSION HAZARDS									
<u> </u>									

		SECTION	NV - HEAI	LTH HAZARD D	ATA	
THRESHOLD LIM	IIT VALUE					
EFFECTS OF OVE	EREXPOSU	RE				
EMERGENCY AN	D FIRST AI	D PROCEDURES				
	,					
	-					
STABILITY		SECTI		EACTIVITY DAT	ГА '	
STABILITY	UNSTA	ABLE	CONDITION	IS TO AVOID		
INCOMPATABILI	STABL	į				- · · ·
		·				
HAZARDOUS DE	COMPOSITI	ON PRODUCTS	· · · · · · · · · · · · · · · · · · ·	T 444.414.414.4		
HAZARDOUS POLYMERIZATIO	i	MAY OCCUR		CONDITIONS TO A	AVOID	
. Julian in writers I I'c		WILL NOT OCCUR				
		SECTION V	II - SPILL	OR LEAK PROC	EDURES	
STEPS TO BE TA	KEN IN CA	SE MATERIAL IS RE	ELEASED OR S	SPILLED		
	<del></del>					<del></del>
					NUCL.	
WASTE DISPOSA	L METHOD	)				
		· · · · · · · · · · · · · · · · · · ·				
		· · · · · · · · · · · · · · · · · · ·				*···
		SECTION VIII -	SPECIAL F	PROTECTION IN	FORMATION	
RESPIRATORY F	PROTECTIO	N (Specify type)				
VENTILATION	LOCA	L EXHAUST		•	SPECIAL	
	MECH	ANICAL (General)			OTHER	···
PROTECTIVE GL	OVES			EYE PROTECTION		
OTHER PROTEC	TIVE EQUI	PMENT				<del></del>
						<del></del>
		SECTIO	N IX - SPE	CIAL PRECAUT	IONS	
PRECAUTIONS 1	TO BE TAK	EN IN HANDLING A	ND STORING			
OTHER PRECAU	ITIONS	•	<del> </del>			
					-	
	<del></del>					

PAGE (2)

GPO 934-110

Form OSHA-20 Rev. May 72

## TABLE XII-1

# SIGNIFICANT PHYSICAL PROPERTIES OF BENZENE

Formula	С6н6
Formula Weight	78.1
Boiling Point	80.1 C (176 F) at 760 mm Hg
Melting Point	5.5 C (42 F)
Specific Gravity	0.8790 g/ml at $\frac{20 \text{ C } (68 \text{ F})}{4 \text{ C } (39.2 \text{ F})}$
Solubility	0.06% in water, mixes freely with alcohol, ether and most organic solvents.
Explosive Range for Vapor	1.4 - 7.1% by volume in air
Flash Point	-12 to -10 C (10.4-14 F)
Ignition Temperature	490 C (914 F)
Vapor Density	2.7 (Air = 1.0)

Derived from references 7 and 8

## TABLE XII-2

## BENZENE REACTIONS OF COMMERCIAL IMPORTANCE

1.	Halogenation and subsequent hydrolysis to produce phenol:
	C6H6 + C12 (Metallic Iron Catalyst) C6H5C1 + HC1 chlorobenzene
	C6H5Cl + NaOH (6-8% aqueous solution) 360 C 4500 lb/sq in
	C6H5ONa + HC1 C6H5OH + NaCl phenol
2.	Hydrogenation of benzene to produce cyclohexane:
	C6H6 + 3H2 (Metallic Nickel Catalyst) C6H12 150-200 C, 25 atm cyclohexane
3.	Friedel-Crafts reaction of benzene and ethylene to pro-
	duce ethyl benzene which is then dehydrogenated to yield styrene:
	C6H6 + C2H4 (Phosphoric Acid Catalyst) C6H5C2H5 ethyl benzene
	C6H5C2H5 (Cr2O3 . A12O3 Catalyst) C6H5CH=CH2 + H2 styrene
Fro	m Chemical Economics Handbook [3]

TABLE XII-3

SUMMARY OF BLOOD FINDINGS
ON EXAMINATION OF WORKERS
EXPOSED TO BENZENE

		Loca1	_	e benzene ir, ppm	Blood fin	Blood findings		
Group	Room	ventila- tion	Summer	Winter	Number of persons examined	Number positive		
I-A								
Small amount of benzene;	150B	-	100		9	2		
no local ventilation;	60	_	150		1	0		
low benzene content in air.	27A	***	110		2	1		
I-B								
Small amount of benzene;	27B	_	700		2	0		
no local ventilation; high	59	_	150	210	9	1		
benzene content in air.	61A	_	130	210	12	6		
	61B	-	1,360	580	1	1		
II-A				1				
Large amount of benzene;	78A	+	70	90	0			
local ventilation; low	150A	+	90		1	1		
benzene content in air.	75B	+	100		3	1		
II-B								
Large amount of benzene;	91	+	180	400	5	*0		
local ventilation; high	50B	+		430	3	1		
benzene content in air.	50A	+ -		500	4	1		
	75A	+	130	330	10	1		
III								
Large amount of benzene; no			340		1	0		
local ventilation; high	23	<del>-</del> .			6	2		
benzene content in air.	83	_	620		9	6		
	95	-	1,800		3	2		
Total					81	26		

<sup>\*3</sup> clinical cases, 1 fatal, since tests were made. From Greenburg [19]

TABLE XII-4

DETAILED BLOOD COUNTS ON 13 WORKERS
EXHIBITING THE PICTURE OF EARLY BENZOL POISONING

Plant Code No.	Нb	RBC	WBC	Poly	Lym- phocytes	Large Mono- nuclears	Eosin	Trans
				%	%	%	%	%
23	65	4,376,000	5,300	58	36	3.5	1.5	0.5
23	75	4,400,000	5,200	55	39	3.5	2.0	0.5
23			4,100					
23			4,800					
27	55	4,304,000	4,667	55	36	5.0	1.0	2.0
59	70	5,424,000	6,140	47	47	3.5	0.5	1.0
61	85		4,450					
61	50		4,000					
	40	1,736,000	3,000					
61	75		2,850					
	80	1,736,000	4,200					
61	23	800,000	3,000					
83	27	1,055,000	1,450	58	36	5.0	1.0	0.0
	41							
	30	2,100,000	2,100					
	29	1,365,000	2,200	44	49	6.0	1.0	0.0
95	55	3,193,000	3,100	50	39	1.5	7.0	1.5
95	70	4,968,000	3,600	47	41	0.5	8.0	3.0
Normal		5,000,000						
male	90-110	5,500,000	7,500	65-70	30	1-2	1-2	2-4
Normal		4,500,000						
female	50-100	5,000,000	7,500	65-70	30	1-2	1-2	2-4

From Greenburg [19]

TABLE XII-5

INCIDENCE OF SIGNIFICANT ABNORMALITIES
IN CASES COMPLETELY STUDIED, BY DIAGNOSIS

Test	Criteria	of Abnormality	Se <b>v</b> Cas	ere es	Early Cases		Negative Cases	
			No.	*	No.	%	No.	%
RBC	Less than	4.5 million	15	68.2	31	72.1	-	
Mean corp volume	More than	94 cu µm	14	63.6	25	58.1	9	24.3
Platelets	Less than	100,000	18	81.3	14	32.6		
Hemoglobin	Less than	13.0 gm/100 cc	8	36.4	11	25.6		
WBC	Less than	5,000	19	86.5	13	30.2		<u></u> -
Numbe	er of cases	examined		22		43		37

From Greenburg et al [17]

TABLE XII-6

COMBINATIONS OF TESTS WHICH WOULD REVEAL A HIGH PROPORTION
OF INDIVIDUALS SHOWING THE BENZENE EFFECT, ACCORDING TO
POSITIVE CASES WITH COMPLETE BLOOD STUDIES\*

Combined Tests		soning Revealed
	•	t Combinations
	No.	<b>%</b>
MCV + RBC	61	82.4
MCV + WBC	59	79.7
MCV + Hb	59	79.7
MCV + Platelets	57	77.0
RBC + Platelets	56	75.7
RBC + WBC	54	73.0
RBC + Hb	51	68.9
MCV + RBC + WBC + Platelets	72	97.3
MCV + RBC + WBC	69	93.2
MCV + RBC + Platelets	66	89.2
MCV + RBC + Hb	65	87.8
Single Tests		
MCV	48	64.9
RBC	47	63.5
Platelets	31	41.9
WBC	30	40.5
НЬ	30	40.5
Total positive cases having		
complete blood studies	74	100.0

<sup>\*</sup>Includes 9 cases with macrocytosis as the only blood abnormality.
From Greenburg et al [17]

TABLE XII-7 PRESUMPTIVE DURATION OF CONTACT AND INTERVAL BETWEEN LAST CONTACT AND DEATH OR BIOPSY IN CHRONIC BENZENE POISONING

Case	Sex	Age	Age Industry Duration of Contact		Interval Since Last Contact			
1	M	22	Rubber factory	6 months	9 months (N)			
2	M	54	Artificial leather	7 years	1 month (N)			
3	F	20	Rubber cement	8 months	1 month (N)			
4	M	25	Artificial leather	3 years	1 month (N)			
5	M	46	Cobbler*	years	l month (N)			
6	F	44	Rubber factory	4 years	6 months (N)			
7	M	48	Artificial leather	12 years	5 months (B)			
8	M	45	Artificial leather	1 1/2 years	4 months (N)			
9	M	45	Artificial leather	3 years	1 1/2 years (N)			
10	M	43	Artificial leather	years	? (N)			
11	F	18	Rubber factory	7 months	1 month (N)			
12	M	54	Artificial leather	3 years	3 months (N)			
13	M	51	Cobbler*	2 years	? (N)			
14	F	63	Telephone operator**	5 years	3 months (N)			
15	M	28	Artificial leather	4 years	6 years (N)			
16	M	57	Artificial leather	l year	2 years (A) (B)			
17	M	57	Artificial leather	5 years	5 months (A) (B)			
18	M	41	Furniture finisher***	years	2 1/2 months (N)			
19	M	12	Schoolboy***	?	2 months (A) (B)			

From Mallory et al [22]

<sup>(</sup>N) Necropsy, (B) Biopsy, (A) Alive. \*Used benzene as solvent for rubber cement.

<sup>\*\*</sup>Used solvent containing 50% benzene for eradicating names on switchboard.

<sup>\*\*\*</sup>Used paint remover containing benzene.

TABLE XII-8

AIR ANALYSES AT A BENZENE COATING PLANT

	Benzene Vapor ppm					
Location	De- cem- ber 1938	July 1946**	Au- gust 1946	Aver- age		
Coating Room-Machine						
No. 1	60*	70*	50*	60*		
Coating Room-Average	45	40	40	40		
Coating Room-Maximum	60	70	55	60		
Mixing Room-Average	80	80		80		

<sup>\*</sup>Exposure of deceased worker or successor.

<sup>\*\*</sup>Analysis by an insurance company.

Derived from Hardy and Elkins [57]

TABLE XII-9

SUMMARY OF ENVIRONMENTAL BENZENE LEVELS AND URINARY PHENOL EXCRETIONS FOR WORKERS IN A RUBBER COATING PLANT USING NAPHTHA SOLVENTS (3-7.5% by Volume)

W	kr J	lob	Date Empl Began	Age When Hired	Urine Phenol mg/l	5/25/60 Equiv Air Level ppm	Actl Air	Urine Phenol mg/1	7/14/60 Equiv Air Level	Actl Air Anal ppm	Urine Phenol mg/1	1/13/61 Equiv Air Level	Actl Air Anal ppm	Urine Phenol mg/l	9/6/61 Equiv Air Level	Actl Air Anal ppm
	A Spre	ader	7/55	55	106	10	5,11,	158	19	7,25	250	29	20,25	<del>-</del>		19,36,
	В		9/44	17	114	13	12,27	75	10	(16.0)	160	19	(22.5)	130	13	25
	С		6/57	24	68	10	(13.8)*	_	_	, ,	250	29	, ,	162	19	(26.3)
	D		8/51	47	111	13		_	_		330	38		200	25	•
	E		12/55	34	270	31			_		-	_		200	. 25	
	F		2/46	34	_	-		-	_		350	41		260	31	
,	G		7/60	33	-	-		-	-		-	-		255	31	
]	H Satu	rator	8/57	20	570	74	68	-	-	57	700	95**	90	295	35	22,23
:	I Chur	ner	9/47	38	_	~		190	22	12,17	360	44		152	19	14,16,
	J		9/53	22	-	_		_	-	(14.5)		31		106	10	44
1	K		2/59	18	-	_		-	_	, ,	300	35		-	_	(24.7)
1	L		10/58	21	_	_			_		480	62		390	47	,

<sup>\*</sup> Mean

129

<sup>\*\*</sup> Extrapolated

From Pagnotto (written communication, 1972)

SUMMARY OF ENVIRONMENTAL BENZENE LEVELS AND
URINARY PHENOL EXCRETIONS FOR WORKERS IN A RUBBER
COATING PLANT USING NAPHTHA SOLVENTS (3-7.5% by Volume)
(Continued)

Work	er Job	Urine Phenol mg/l	8/16/62 Equiv Air Level ppm	Actual Air Analysis ppm	Urine Phenol mg/l	4/10/63 Equiv Air Level ppm	Actual Air Analysis ppm	Urine Phenol mg/1	2/12/63 Equiv Air Level ppm	Actual Air Analysis ppm	Years Expos
A	Spreader			12,20,	195	25	35,10,	133	16	17,23,	8
В		96	10	18,3,4	230	27	10,21,	193	25	17,30,	19
C		68	10	(11.4)*	145	16	14,17,	132	16	35,20	6
D		87	10		350	41	38,39,	232	29	(25.3)	12
E		85	10		280	33	25,29	152	19	<del>-</del>	8
F		268	31		370	44	(21.5)	119	13	-	17
G		130	13		435	56		165	19		3
H	Saturator	280	33	10,14 (12)	440	56	43,43, 33 (39.7)	260	31	38,82, 140 (86.7)	6
I	Churner	_			160	19	6	_	_		16
J		-	_		150	16		-	•		10
K		_	_		_	-		-	_		?
L		206	25		300	35		325	38		5

\*Mean

From Pagnotto (written communication, 1972)

TABLE XII-10

URINARY PHENOL LEVELS WITH CORRESPONDING
EQUIVALENT ENVIRONMENTAL BENZENE EXPOSURE LEVELS

Urine Phenol (mg/liter)	Approx. Av. Equiv. Benzene Air Level (ppm)	
100	10	
120	13	
140	16	
160	19	
180	22	
200	25	
220	27	
240	29	
260	31	
280	33	
300	35	
320	38	
340	41	
360	44	
380	47	
400	50	
420	53	
440	56	
460	59	
480	62	
500	65	
520	68	
540	71	
560	74	
580	77	
600	80	

From Pagnotto (written communication, 1972)

TABLE XII-11

SUMMARY OF HEMOGLOBIN LEVELS FOR WORKERS IN A RUBBER COATING PLANT USING NAPHTHA SOLVENTS (3-7% by Volume)

Worker	3/10/61	3/30/61	9/20/63	10/31/63
В	<del>,</del>	12.5		12.6
H		13.0	13.8	
J	13.4	12.8		
L	12.2	11.3	11.2	11.5
M	14.6			
N		12.7		
0		12.2		

From Pagnotto (written communication, 1972)

TABLE XII-12

COMPARISON OF BENZENE AIR LEVELS
FROM URINE PHENOL AND AIR SAMPLE DATA

Benzene in Air ppm

Occupation	Urine Pheno1* mg/liter	Estimated from Urine Phenols	Air Sampling Data (TWA)
Agitator operator	105	10	1.3
Agitator operator	107	10	10.7
Benzol loader	<65	<5	1.7
Benzol still operator	<65	<5	6.7
Benzol oil still operator	r <65	<5	0.8
Naphthalene operator	115	12	8.5
Analyst	105	10	2.4
Chemical observer	68	5	12.0
Foreman	<65	<b>&lt;</b> 5	none
Repairman	<65	<b>&lt;</b> 5 ′	2.6
Chemical observer	65	5	17.1
Chemical observer	112	11	12.2
Chemical observer	66	5	6.5
Control tester	66	5	14.6
Stillman	212	24	39.2
Chemist	157	17	8.8
Pumpman helper	302	36	55
Pumpman helper	84	7	9.5

From Bethlehem Steel data (written communication, 1972)

<sup>\*</sup>Values less than 65 mg/liter were not considered to differ significantly from that of an unexposed normal adult.

TABLE XII-13
BENZENE PLANT AIR LEVELS
ppm

Benzene in Air

Occupation	8-Hour TWA	Range
Agitator Operator	6.0	0.5 - 20
Benzol Loader & Loader Helper	4.0	0.5 - 15
Benzol Still Operator	4.0	1 - 15
Light Oil Still Operator	2.5	1 - 15
Naphthalene Operator	10	2 - 30
Analyst	10	2 - 30
Chemical Observer	10	4 - 50
Foreman	1.5	1 - 10

From Bethlehem Steel data (written communication, 1972)

# TABLE XII-14

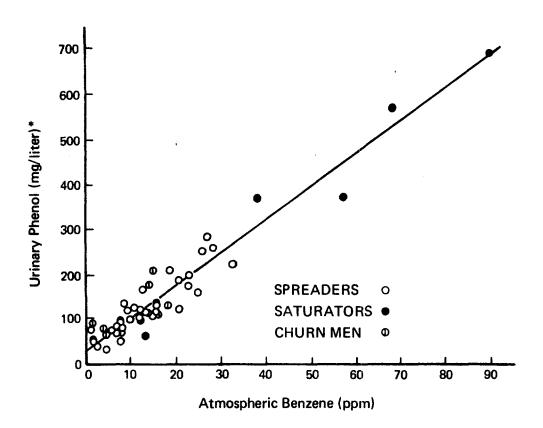
# NORMAL HEMATOLOGIC VALUES

# Cell Counts

Erythrocytes	Male Female	4.6-6.2 million/cu mm 4.2-5.4 million/cu mm		
Leukocytes	Total	5,000-10,000/cu mm		
	Differential			
	Myelocytes Immature polymorpho-	0%		
·	nuclears Segmented	3-5%		
	neutrophils	54-62%		
	Lymphocytes	25-33%		
	Monocytes	3-7%		
	Eosinophils	1-3%		
	Basophils	0-0.75%		
Platelets		150,000-350,000/cu mm		
Reticulocytes		0.5-1.5% of erythrocytes		
Corpuscular Values for Erythrocytes				
Mean Corpuscular Hemoglobin		27-31 picograms		
Mean Corpuscular Volume		82-92 cu micra		
Mean Corpuscular Hemo-				
globin Concent	ration	32–36%		
Hematocrit Mal	.e	40-54%		
Fem	ale	37-47%		
Hemoglobin Mal	.e	14.0-18.0 g%		
	ale	12.0-16.0 g%		
Serum Bilirubin Concentration				
Tot	:al	0.3-1.1 mg%		
Dir	ect	0.1-0.4 mg%		
Ind	lirect	0.2-0.7 mg%		
From Conn [142]				

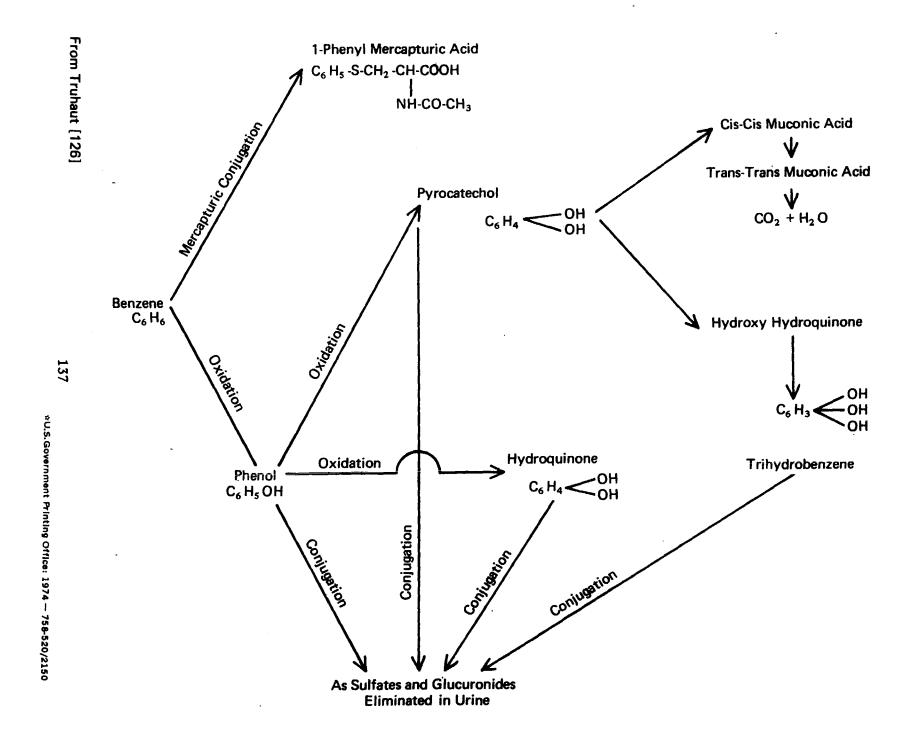
## FIGURE XII-1

# COMPARISON OF PHENOL IN URINE WITH BENZENE IN AIR



<sup>\*</sup>Represents both phenol and paracresol. Phenol alone would result in values lower than indicated.

Derived from Pagnotto [12]



SUGGESTED METABOLIC TRANSFORMATION OF BENZENE IN MAN

FIGURE XII-2