

ジフェニルアミンのラットを用いた  
経口投与によるがん原性試験（混餌試験）報告書

試験番号：0684

# APPENDICES

## APPENDICES

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APPENDIX 1-1

IDENTITY OF DIPHENYLAMINE IN THE 2-YEAR FEED STUDY

## IDENTITY OF DIPHENYLAMINE IN THE 2-YEAR FEED STUDY

Test Substance : Diphenylamine (Wako Pure Chemical Industries, Ltd.)

Lot No. : LTF7564

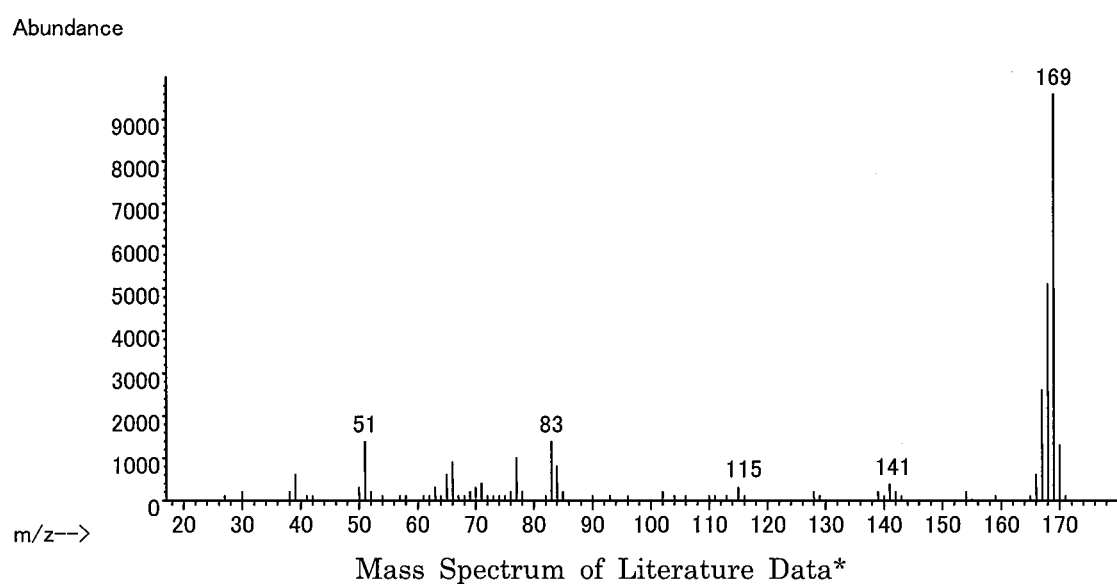
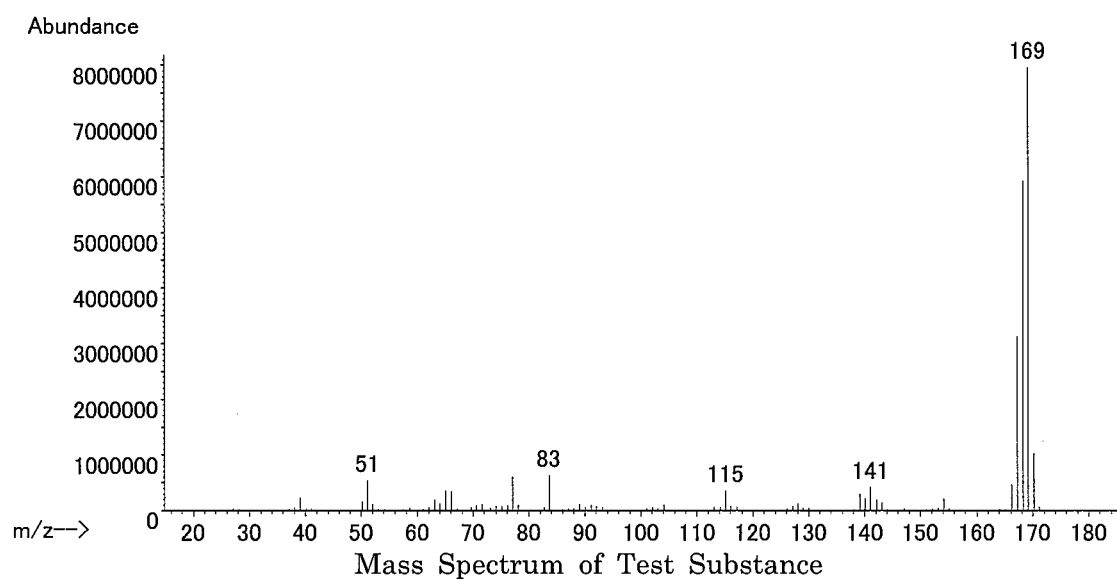
## 1. Spectral Data

Mass Spectrometry

Instrument : Agilent Technologies 5973N Mass Spectrometer

Ionization : EI (Electron Ionization)

Ionization Voltage : 70eV



Result: The mass spectrum was consistent with literature spectrum.

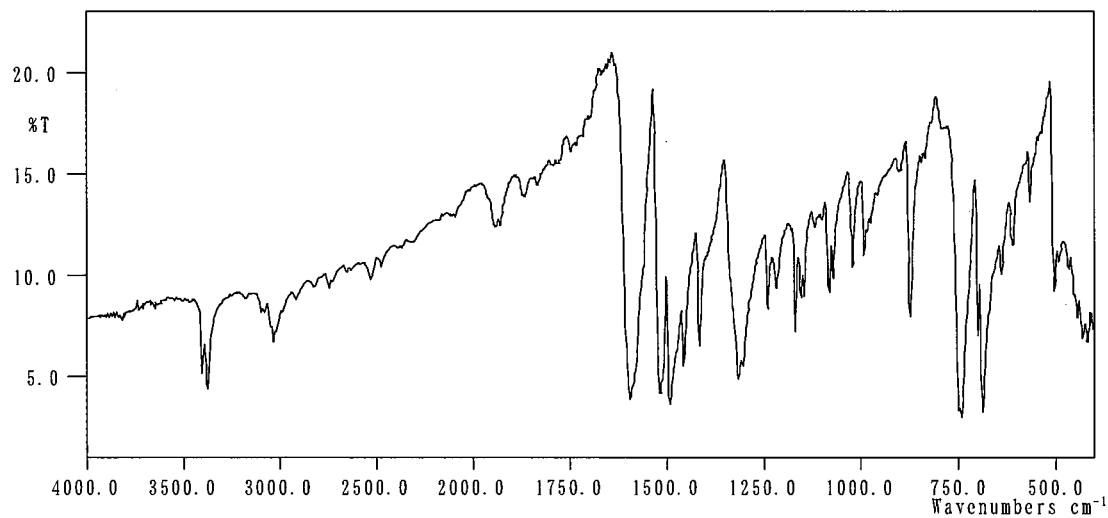
(\*McLafferty FW, ed. 1994. Wiley Registry of Mass Spectral Data. 6th ed. New York, NY:John Wiley and Sons.)

## Infrared Spectrometry

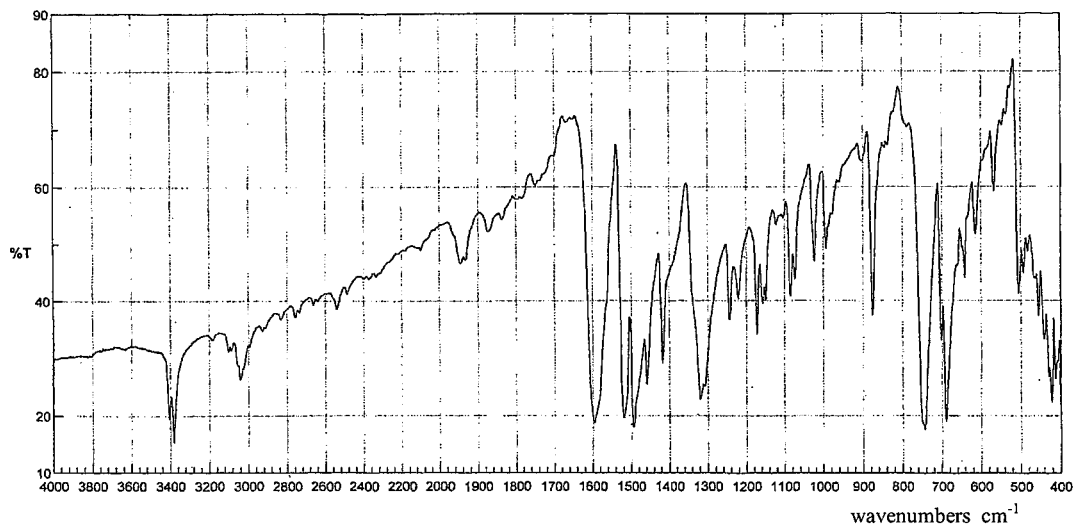
Instrument : Shimadzu FTIR-8200PC Infrared Spectrometer

Cell : KBr

Resolution : 2  $\text{cm}^{-1}$



Infrared Spectrum of Test Substance



Infrared Spectrum of Literature Data\*

Result: The infrared spectrum was consistent with literature spectrum.  
(\*Performed by Wako Pure Chemical Industries, Ltd.)

2. Conclusion: The test substance was identified as diphenylamine by mass spectrum and infrared spectrum.

## APPENDIX 1-2

### STABILITY OF DIPHENYLAMINE IN THE 2-YEAR FEED STUDY

## STABILITY OF DIPHENYLAMINE IN THE 2-YEAR FEED STUDY

Test Substance : Diphenylamine (Wako Pure Chemical Industries, Ltd.)

Lot No. : LTF7564

## 1. High Performance Liquid Chromatography

Instrument : Shimadzu LC-10 High Performance Liquid Chromatograph

Column : TSK-GEL ODS-80TM (4.6 mm  $\phi$   $\times$  15 cm)

Column Temperature: 40 °C

Flow Rate : 1 mL/min

Mobile Phase : Acetonitrile : Distilled Water = 70 : 30

Detector : UV (285 nm)

Injection Volume : 10  $\mu$ L

Date analyzed	Peak No.	Retention Time (min)	Area (%)
2007.07.03	1	3.869	100
2009.08.07	1	3.985	100

Result: High performance liquid chromatography indicated one major peak (peak No.1) analyzed on 2007.7.3 and one major peak (peak No.1) analyzed on 2009.8.7. No new trace impurity peak in the test substance analyzed on 2009.8.7 was detected.

2. Conclusion: The test substance was stable for the period that the test substance had been used for the study.

APPENDIX 1-3

CONCENTRATION OF DIPHENYLAMINE  
IN FORMULATED DIETS IN THE 2-YEAR FEED STUDY



## CONCENTRATION OF DIPHENYLAMINE IN FORMULATED DIETS IN THE 2-YEAR FEED STUDY

Analytical Method : The samples were analyzed by high performance liquid chromatography.

Instrument : Shimadzu LC-10 High Performance Liquid Chromatograph

Column : TSK-GEL ODS-80TM (4.6 mm  $\phi$   $\times$  15 cm)

Column Temperature: 40 °C

Flow Rate : 1 mL/min

Mobile Phase : Acetonitrile : Distilled Water = 70 : 30

Detector : UV (285 nm)

Injection Volume : 10  $\mu$ L

Date Analyzed	Target Concentration		
	250 <sup>a</sup>	1000	4000
2007.07.09	260 <sup>b</sup> (104) <sup>c</sup>	1020 (102)	4180 (105)
2007.09.24	261 (104)	969 ( 96.9)	3950 ( 98.8)
2007.12.17	243 ( 97.2)	983 ( 98.3)	4050 (101)
2008.03.10	253 (101)	958 ( 95.8)	3730 ( 93.3)
2008.06.02	248 ( 99.2)	958 ( 95.8)	3880 ( 97.0)
2008.08.25	259 (104)	965 ( 96.5)	3740 ( 93.5)
2008.11.17	266 (106)	1040 (104)	4170 (104)
2009.02.09	247 ( 98.8)	1040 (104)	4090 (102)
2009.05.04	240 ( 96.0)	1020 (102)	4150 (104)

<sup>a</sup> ppm

<sup>b</sup> ppm (Mean measured concentration.)

<sup>c</sup> % (Mean measured concentration/target concentration  $\times$  100.)

APPENDIX 1-4

HOMOGENEITY OF DIPHENYLAMINE  
IN FORMULATED DIETS IN THE 2-YEAR FEED STUDY

HOMOGENEITY OF DIPHENYLAMINE IN FORMULATED DIETS IN THE 2-YEAR  
FEED STUDY

Analytical Method : The samples were analyzed by high performance liquid chromatography.

Instrument : Shimadzu LC-10 High Performance Liquid Chromatograph

Column : TSK-GEL ODS-80TM (4.6 mm  $\phi$   $\times$  15 cm)

Column Temperature: 40 °C

Flow Rate : 1 mL/min

Mobile Phase : Acetonitrile : Distilled Water = 70 : 30

Detector : UV (285 nm)

Injection Volume : 10  $\mu$ L

	Target Concentration		
	250 <sup>a</sup>	1000	4000
Coefficient Variation	4.69 <sup>b</sup>	5.85	3.04

<sup>a</sup> ppm<sup>b</sup> % (n=7)

## APPENDIX 1-5

### STABILITY OF DIPHENYLAMINE IN FORMULATED DIETS

## STABILITY OF DIPHENYLAMINE IN FORMULATED DIETS

Analytical Method : The samples were analyzed by high performance liquid chromatography.

Instrument : Shimadzu LC-10 High Performance Liquid Chromatograph

Column : TSK-GEL ODS-80TM (4.6 mm  $\phi$   $\times$  15 cm)

Column Temperature: 40 ° C

Flow Rate : 1 mL/min

Mobile Phase : Acetonitrile : Distilled Water = 70 : 30

Detector : UV (285 nm)

Injection Volume : 10  $\mu$ L

Date Analyzed	Target Concentration	
	100 <sup>a</sup>	10000
2006.09.08	99.0 (100) <sup>b</sup>	10100 (100)
2006.09.16 <sup>c</sup>	92.7 ( 93.6)	9690 ( 95.9)
2006.09.16 <sup>d</sup>	100 (101)	10100 (100)

<sup>a</sup> ppm

<sup>b</sup> % (Percentage was based on the concentration at the date of preparation.)

<sup>c</sup> Animal room samples

<sup>d</sup> Cold storage samples

## APPENDIX 2

METHODS, UNITS AND DECIMAL PLACE FOR HEMATOLOGY

AND BIOCHEMISTRY IN THE 2-YEAR FEED

STUDY OF DIPHENYLAMINE

**METHODS, UNITS AND DECIMAL PLACE FOR HEMATOLOGY AND BIOCHEMISTRY  
IN THE 2-YEAR FEED STUDY OF DIPHENYLAMINE**

Item	Method	Unit	Decimal place
<b>Hematology</b>			
Red blood cell (RBC)	Light scattering method <sup>1)</sup>	$\times 10^6/\mu\text{L}$	2
Hemoglobin(Hgb)	Cyanmethemoglobin method <sup>1)</sup>	g/dL	1
Hematocrit(Hct)	Calculated as $\text{RBC} \times \text{MCV}/10$ <sup>1)</sup>	%	1
Mean corpuscular volume(MCV)	Light scattering method <sup>1)</sup>	fL	1
Mean corpuscular hemoglobin(MCH)	Calculated as $\text{Hgb}/\text{RBC} \times 10$ <sup>1)</sup>	pg	1
Mean corpuscular hemoglobin concentration (MCHC)	Calculated as $\text{Hgb}/\text{Hct} \times 100$ <sup>1)</sup>	g/dL	1
Platelet	Light scattering method <sup>1)</sup>	$\times 10^3/\mu\text{L}$	0
Reticulocyte	Light scattering method <sup>1)</sup>	%	1
Methemoglobin	Van Assendelft method <sup>2)</sup>	%	1
White blood cell(WBC)	Light scattering method <sup>1)</sup>	$\times 10^3/\mu\text{L}$	2
Differential WBC	Light scattering method <sup>1)</sup>	%	0
<b>Biochemistry</b>			
Total protein(TP)	Biuret method <sup>3)</sup>	g/dL	1
Albumin (Alb)	BCG method <sup>3)</sup>	g/dL	1
A/G ratio	Calculated as $\text{Alb}/(\text{TP} - \text{Alb})$ <sup>3)</sup>	—	1
T-bilirubin	Azobilirubin method <sup>3)</sup>	mg/dL	2
Glucose	GlcK·G-6-PDH method <sup>3)</sup>	mg/dL	0
T-cholesterol	CE·COD·POD method <sup>3)</sup>	mg/dL	0
Triglyceride	MGLP·GK·GPO·POD method <sup>3)</sup>	mg/dL	0
Phospholipid	PLD·ChOD·POD method <sup>3)</sup>	mg/dL	0
Aspartate aminotransferase (AST)	JSCC method <sup>3)</sup>	IU/L	0
Alanine aminotransferase (ALT)	JSCC method <sup>3)</sup>	IU/L	0
Lactate dehydrogenase (LDH)	JSCC method <sup>3)</sup>	IU/L	0
Alkaline phosphatase (ALP)	JSCC method <sup>3)</sup>	IU/L	0
$\gamma$ -Glutamyl transpeptidase ( $\gamma$ -GTP)	JSCC method <sup>3)</sup>	IU/L	0
Creatine kinase (CK)	JSCC method <sup>3)</sup>	IU/L	0
Urea nitrogen	Urease·GLDH method <sup>3)</sup>	mg/dL	1
Creatinine	Jaffé method <sup>3)</sup>	mg/dL	1
Sodium	Ion selective electrode method <sup>3)</sup>	mEq/L	0
Potassium	Ion selective electrode method <sup>3)</sup>	mEq/L	1
Chloride	Ion selective electrode method <sup>3)</sup>	mEq/L	0
Calcium	OCPC method <sup>3)</sup>	mg/dL	1
Inorganic phosphorus	PNP·XOD·POD method <sup>3)</sup>	mg/dL	1

1) Automatic blood cell analyzer (ADVIA120 : Siemens Healthcare Diagnostics Inc.)

2) Spectrophotometer (DU-530 : Beckman Coulter, Inc.)

3) Automatic analyzer (Hitachi 7080 : Hitachi, Ltd.)