

エチレングリコールモノエチルエーテルアセテートのラットを用いた吸入による2週間毒性試験報告書

試験番号：0735

# APPENDICES

## APPENDICES

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ETHYLENE GLYCOL MONOETHYL ETHER ACETATE

**APPENDIX 1 1**

**IDENTITY OF ETHYLENE GLYCOL  
MONOETHYL ETHER ACETATE  
IN THE 2-WEEK INHALATION STUDY**

# IDENTITY OF ETHYLENE GLYCOL MONOETHYL ETHER ACETATE IN THE 2-WEEK INHALATION STUDY

Test Substance : Ethylene glycol monoethyl ether acetate (Wako Pure Chemical Industries, Ltd.)

Lot No. : KWR2691

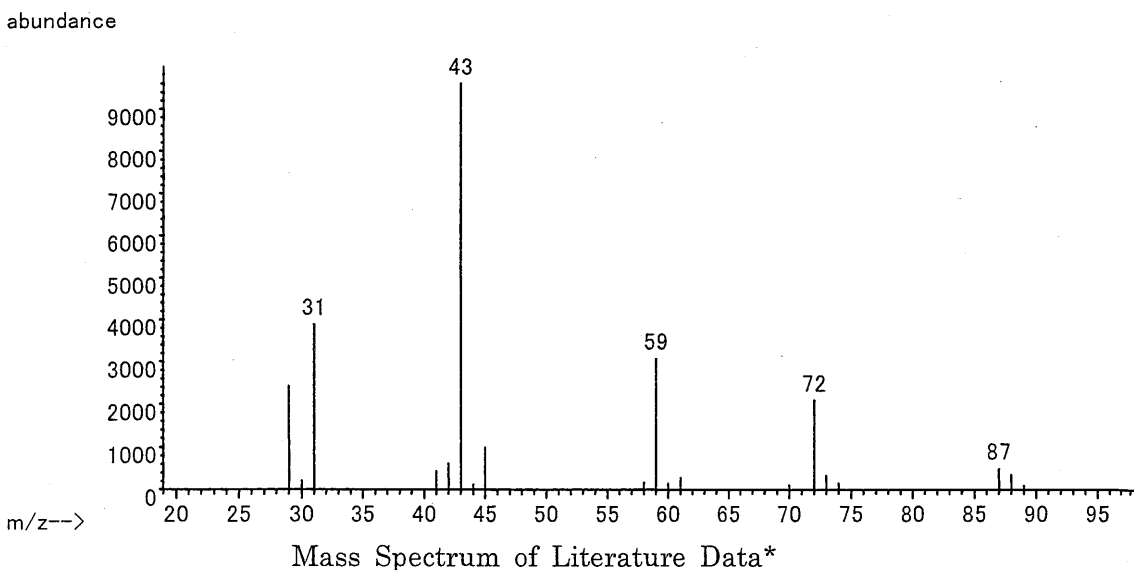
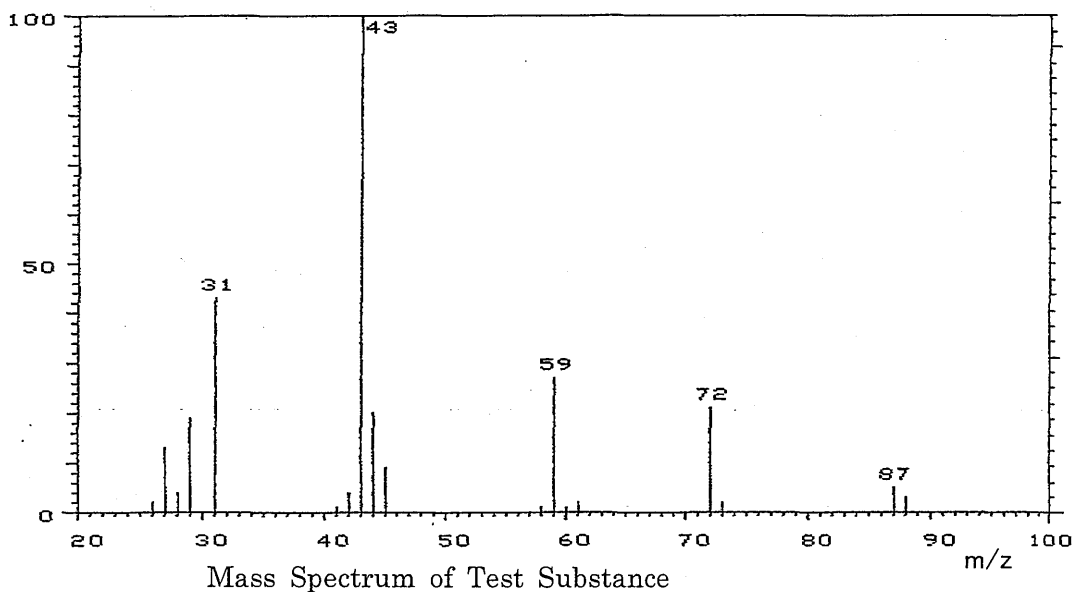
## 1. Spectral Data

### Mass Spectrometry

Instrument : Hitachi M-80B Mass Spectrometer

Ionization : EI (Electron Ionization)

Ionization Voltage : 70eV



Result: The mass spectrum was consistent with literature spectrum.

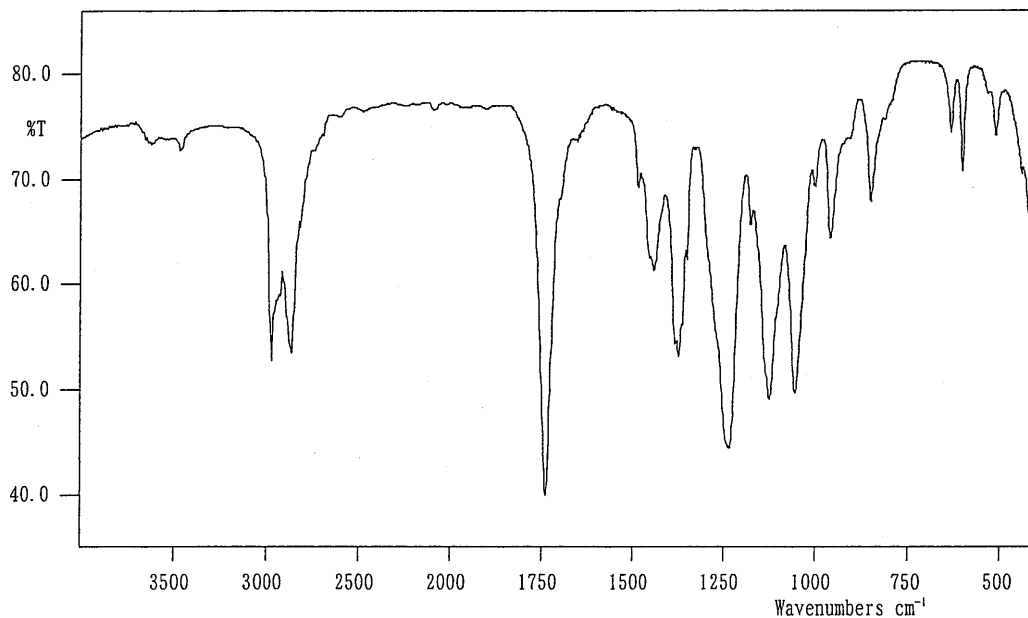
(\*McLafferty FW, ed. 1994. Wiley Registry of Mass Spectral Data. 6th ed. New York, NY:John Wiley and Sons.)

## Infrared Spectrometry

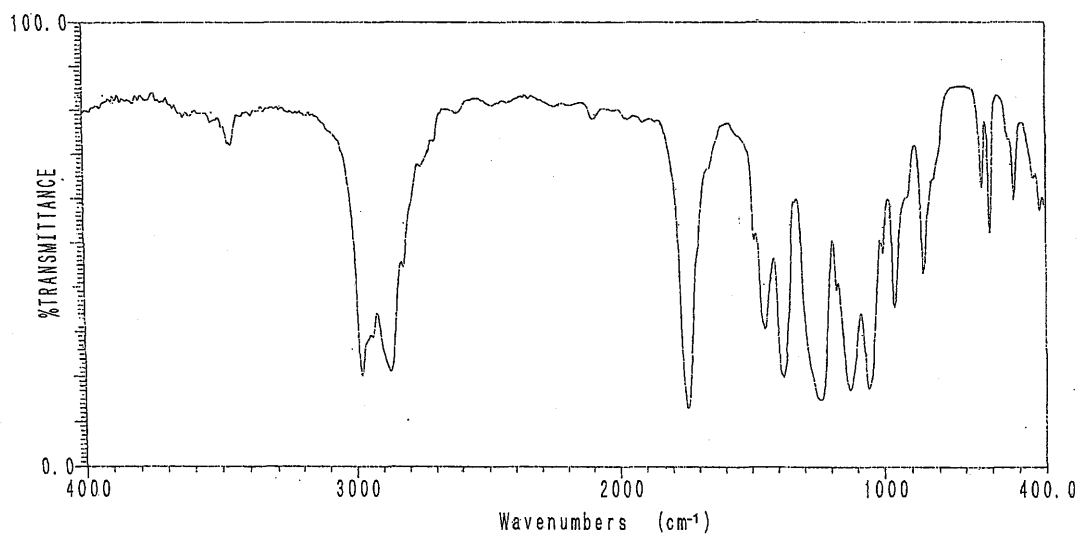
Instrument : Shimadzu FTIR-8200PC Infrared Spectrometer

Cell : KBr Liquid Cell

Resolution : 4  $\text{cm}^{-1}$



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 ethylene glycol monoethyl ether acetate by mass spectrum and infrared spectrum.

**APPENDIX 1 2**

**STABILITY OF ETHYLENE GLYCOL  
MONOETHYL ETHER ACETATE  
IN THE 2-WEEK INHALATION STUDY**

## STABILITY OF ETHYLENE GLYCOL MONOETHYL ETHER ACETATE IN THE 2-WEEK INHALATION STUDY

Test Substance : Ethylene glycol monoethyl ether acetate (Wako Pure Chemical Industries, Ltd.)

Lot No. : KWR2691

## 1. Gas Chromatography

Instrument : Agilent Technologies 5890A Gas Chromatograph

Column : INNOWAX ( 0.53 mm $\phi$   $\times$  60 m)

Column Temperature: 140 $^{\circ}$  C

Flow Rate : 10 mL/min

Detector : FID (Flame Ionization Detector)

Injection Volume : 1  $\mu$ L

Date (date analyzed)	Peak No.	Retention Time (min)	Area (%)
2009.02.26	1	2.885	100
2009.03.16	1	2.886	100

Result: Gas chromatography indicated one major peak (peak No.1) analyzed on 2009.2.26 and one major peak (peak No.1) analyzed on 2009.3.16. No new trace impurity peak in the test substance analyzed on 2009.3.16 was detected.

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

## **APPENDIX 2**

### **ENVIRONMENTAL CONDITIONS OF INHALATION CHAMBER IN THE 2-WEEK INHALATION STUDY OF ETHYLENE GLYCOL MONOETHYL ETHER ACETATE**



ENVIRONMENTAL CONDITIONS OF INHALATION CHAMBER IN THE 2-WEEK  
INHALATION STUDY OF ETHYLENE GLYCOL MONOETHYL ETHER ACETATE

Group Name	Temperature (°C) Mean ± S.D.	Humidity (%) Mean ± S.D.	Ventilation Rate (L/min) Mean ± S.D.	Air Change (time/h) Mean
Control	22.3 ± 0.1	56.2 ± 0.7	104.4 ± 0.2	12.0
50 ppm	21.9 ± 0.1	57.2 ± 1.0	104.3 ± 0.5	12.0
100 ppm	21.7 ± 0.1	57.2 ± 0.9	104.5 ± 0.3	12.1
200 ppm	21.7 ± 0.1	57.0 ± 1.9	104.4 ± 0.5	12.0
400 ppm	21.6 ± 0.1	55.5 ± 1.2	104.3 ± 0.4	12.0
800 ppm	21.8 ± 0.1	48.9 ± 3.4	104.1 ± 0.5	12.0

## **APPENDIX 3**

**METHODS, UNITS AND DECIMAL PLACE FOR  
HEMATOLOGY AND BIOCHEMISTRY  
IN THE 2-WEEK INHALATION STUDY OF  
ETHYLENE GLYCOL MONOETHYL ETHER ACETATE**

METHODS, UNITS AND DECIMAL PLACE FOR HEMATOLOGY AND BIOCHEMISTRY  
IN THE 2-WEEK INHALATION STUDY OF  
ETHYLENE GLYCOL MONOETHYL ETHER ACETATE

Item	Method	Unit	Decimal place
<b>Hematology</b>			
Red blood cell (RBC)	Light scattering method <sup>1)</sup>	$\times 10^9/\mu\text{L}$	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
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>2)</sup>	g/dL	1
Albumin (Alb)	BCG method <sup>2)</sup>	g/dL	1
A/G ratio	Calculated as $\text{Alb}/(\text{TP} - \text{Alb})$ <sup>2)</sup>	-	1
T-bilirubin	Azobilirubin method <sup>2)</sup>	mg/dL	2
Glucose	GlcK·G-6-PDH method <sup>2)</sup>	mg/dL	0
T-cholesterol	CE·COD·POD method <sup>2)</sup>	mg/dL	0
Triglyceride	MGLP·GK·GPO·POD method <sup>2)</sup>	mg/dL	0
Phospholipid	PLD·ChOD·POD method <sup>2)</sup>	mg/dL	0
Aspartate aminotransferase (AST)	JSCC method <sup>2)</sup>	IU/L	0
Alanine aminotransferase (ALT)	JSCC method <sup>2)</sup>	IU/L	0
Lactate dehydrogenase (LDH)	JSCC method <sup>2)</sup>	IU/L	0
Alkaline phosphatase (ALP)	JSCC method <sup>2)</sup>	IU/L	0
$\gamma$ -Glutamyl transpeptidase ( $\gamma$ -GTP)	JSCC method <sup>2)</sup>	IU/L	0
Creatine kinase (CK)	JSCC method <sup>2)</sup>	IU/L	0
Urea nitrogen	Urease·GLDH method <sup>2)</sup>	mg/dL	1
Creatinine	Jaffé method <sup>2)</sup>	mg/dL	1
Sodium	Ion selective electrode method <sup>2)</sup>	mEq/L	0
Potassium	Ion selective electrode method <sup>2)</sup>	mEq/L	1
Chloride	Ion selective electrode method <sup>2)</sup>	mEq/L	0
Calcium	OCPC method <sup>2)</sup>	mg/dL	1
Inorganic phosphorus	PNP·XOD·POD method <sup>2)</sup>	mg/dL	1

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

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