

Revision date: 06-Sep-2012 Version: 1.1 Page 1 of 9

## IDENTIFICATION OF THE SUBSTANCE/PREPARATION AND THE COMPANY/UNDERTAKING

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Contact E-Mail: pfizer-MSDS@pfizer.com

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International CHEMTREC (24 hours): +1-703-527-3887

Material Name: Methylprednisolone Acetate Injectable Suspension, Single-Dose Vial

Trade Name: DEPO-MEDROL Chemical Family: DEPO-MEDROL Glucocorticoid

Intended Use: Pharmaceutical product used as anti-inflammatory

## 2. HAZARDS IDENTIFICATION

Appearance: Clear, colorless solution

Signal Word: DANGER

Statement of Hazard: May damage the unborn child.

**Short Term:** May be harmful if absorbed through the skin.

Long Term: Repeat-dose studies in animals have shown a potential to cause adverse effects on

developing fetus and blood and blood forming organs

Known Clinical Effects: Adverse clinical reactions include the development of hypersensitivity and/or irritation leading

to rashes, itching, and burning. Clinical use has resulted in hormonal alterations. Clinical use

has resulted in changes in electrolytes and/or blood chemistry changes.

**EU Indication of danger:** Toxic to reproduction: Category 1

**EU Hazard Symbols:** 



**EU Risk Phrases:** 

R61 - May cause harm to the unborn child. Hazardous Substance. Non-Dangerous Goods.

Australian Hazard Classification (NOHSC):

Note:

This document has been prepared in accordance with standards for workplace safety, which require the inclusion of all known hazards of the active substance or its intermediates regardless of the potential risk. The precautionary statements and warnings included may not apply in all cases. Your needs may vary depending upon the potential for exposure in your workplace.

Material Name: Methylprednisolone Acetate Injectable Page 2 of 9

Suspension, Single-Dose Vial

Revision date: 06-Sep-2012 Version: 1.1

## 3. COMPOSITION/INFORMATION ON INGREDIENTS

## **Hazardous**

Ingredient	CAS Number	<b>EU EINECS/ELINCS List</b>	<b>EU Classification</b>	%
Methylprednisolone Acetate	53-36-1	200-171-3	T;48/22-R61	4-8
Myristyl-gamma-picolinium chloride	2748-88-1	220-387-1	Xn;R22	*
Sodium chloride	7647-14-5	231-598-3	Not Listed	*
Sodium hydroxide	1310-73-2	215-185-5	C;R35	**
Hydrochloric Acid	7647-01-0	231-595-7	C;R35 T;R23	**

Ingredient	CAS Number	<b>EU EINECS/ELINCS List</b>	<b>EU Classification</b>	%
Water for injection	7732-18-5	231-791-2	Not Listed	*
Polyethylene glycol	25322-68-3	Not Listed	Not Listed	*

Additional Information: \* Proprietary

\*\* to adjust pH

Ingredient(s) indicated as hazardous have been assessed under standards for workplace

safety.

For the full text of the R phrases mentioned in this Section, see Section 16

## 4. FIRST AID MEASURES

**Eye Contact:** Flush with water while holding eyelids open for at least 15 minutes. Seek medical attention

immediately.

Skin Contact: Remove contaminated clothing. Flush area with large amounts of water. Use soap. Seek

medical attention.

**Ingestion:** Never give anything by mouth to an unconscious person. Wash out mouth with water. Do not

induce vomiting unless directed by medical personnel. Seek medical attention immediately.

**Inhalation:** Remove to fresh air and keep patient at rest. Seek medical attention immediately.

Symptoms and Effects of Exposure: For information on potential signs and symptoms of exposure, See Section 2 - Hazards

Identification and/or Section 11 - Toxicological Information.

# 5. FIRE FIGHTING MEASURES

**Extinguishing Media:** Use carbon dioxide, dry chemical, or water spray.

Hazardous Combustion Products: May include oxides of carbon.

Fire Fighting Procedures: During all fire fighting activities, wear appropriate protective equipment, including self-

contained breathing apparatus.

Fire / Explosion Hazards: Fine particles (such as dust and mists) may fuel fires/explosions.

Material Name: Methylprednisolone Acetate Injectable Page 3 of 9

Suspension, Single-Dose Vial

Revision date: 06-Sep-2012 Version: 1.1

## 6. ACCIDENTAL RELEASE MEASURES

Health and Safety Precautions: Personnel involved in clean-up should wear appropriate personal protective equipment (see

Section 8). Minimize exposure.

Measures for Cleaning / Collecting: Contain the source of spill if it is safe to do so. Collect spill with absorbent material. Clean spill

area thoroughly.

**Measures for Environmental** 

**Protections:** 

Place waste in an appropriately labeled, sealed container for disposal. Care should be taken to

avoid environmental release.

**Additional Consideration for Large** 

Spills:

Non-essential personnel should be evacuated from affected area. Report emergency situations immediately. Clean up operations should only be undertaken by trained personnel.

## 7. HANDLING AND STORAGE

General Handling: Avoid breathing vapor or mist. Avoid contact with eyes, skin and clothing. When handling, use

appropriate personal protective equipment (see Section 8). Wash thoroughly after handling. Review and implement appropriate technical and procedural waste water and waste disposal measures to prevent occupational exposure or environmental releases. Potential points of process emissions of this material to the atmosphere should be controlled with dust collectors, HEPA filtration systems or other equivalent controls. Releases to the environment should be

avoided.

**Storage Conditions:** Store as directed by product packaging.

## 8. EXPOSURE CONTROLS / PERSONAL PROTECTION

Refer to available public information for specific member state Occupational Exposure Limits.

**Methylprednisolone Acetate** 

Pfizer OEL TWA-8 Hr: 4µg/m³, Skin

Sodium chloride

Latvia OEL - TWA 5 mg/m³
Lithuania OEL - TWA 5 mg/m³

Sodium hydroxide

**ACGIH Ceiling Threshold Limit:**  $2 \text{ mg/m}^3$ Australia PEAK 2 mg/m<sup>3</sup> 2 mg/m<sup>3</sup> **Austria OEL - MAKs Bulgaria OEL - TWA** 2.0 mg/m<sup>3</sup> Czech Republic OEL - TWA  $1 \text{ mg/m}^3$ **Estonia OEL - TWA**  $1 \text{ mg/m}^3$ France OEL - TWA 2 mg/m<sup>3</sup> **Greece OEL - TWA** 2 mg/m<sup>3</sup> 2 mg/m<sup>3</sup> **Hungary OEL - TWA** Japan - OELs - Ceilings  $2 \text{ mg/m}^3$ Latvia OEL - TWA  $0.5 \text{ mg/m}^{3}$ **OSHA - Final PELS - TWAs:**  $2 \text{ mg/m}^3$ Poland OEL - TWA 0.5 mg/m<sup>3</sup> 2 mg/m<sup>3</sup> Slovakia OEL - TWA 2 mg/m<sup>3</sup> Slovenia OEL - TWA  $1 \text{ mg/m}^3$ Sweden OEL - TWAs

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Material Name: Methylprednisolone Acetate Injectable Page 4 of 9

Suspension, Single-Dose Vial

Revision date: 06-Sep-2012 Version: 1.1

# 8. EXPOSURE CONTROLS / PERSONAL PROTECTION

Hydrochloric Acid	
ACGIH Ceiling Threshold Limit:	2 ppm
Australia PEAK	5 ppm
Additional Exit	7.5 mg/m <sup>3</sup>
Austria OEL - MAKs	5 ppm
, we want to be a second of the second of th	8 mg/m <sup>3</sup>
Belgium OEL - TWA	5 ppm
. <b>.</b> .	8 mg/m <sup>3</sup>
Bulgaria OEL - TWA	8.0 mg/m <sup>3</sup>
Cyprus OEL - TWA	5 ppm
•	8 mg/m <sup>3</sup>
Czech Republic OEL - TWA	8 mg/m <sup>3</sup>
Estonia OEL - TWA	5 ppm
	8 mg/m³
Germany - TRGS 900 - TWAs	2 ppm
	3 mg/m <sup>3</sup>
Germany (DFG) - MAK	2 ppm
	3.0 mg/m <sup>3</sup>
Greece OEL - TWA	5 ppm
U OFI TWA	7 mg/m <sup>3</sup>
Hungary OEL - TWA	8 mg/m <sup>3</sup>
Ireland OEL - TWAs	5 ppm 8 mg/m³
Holy OEL TWA	•
Italy OEL - TWA	5 ppm 8 mg/m³
Japan - OELs - Ceilings	5 ppm
Japan - OLLS - Cennigs	7.5 mg/m <sup>3</sup>
Latvia OEL - TWA	5 ppm
	8 mg/m <sup>3</sup>
Lithuania OEL - TWA	5 ppm
	8 mg/m <sup>3</sup>
Luxembourg OEL - TWA	5 ppm
	8 mg/m <sup>3</sup>
Malta OEL - TWA	5 ppm
	8 mg/m <sup>3</sup>
Netherlands OEL - TWA	8 mg/m <sup>3</sup>
Poland OEL - TWA	5 mg/m <sup>3</sup>
Romania OEL - TWA	5 ppm

Polyethylene glycol

Slovakia OEL - TWA

Slovenia OEL - TWA

Spain OEL - TWA

Austria OEL - MAKs1000 mg/m³Germany - TRGS 900 - TWAs1000 mg/m³

**Germany (DFG) - MAK** 1000 mg/m³ average molecular weight 200-600

Slovakia OEL - TWA 1000 mg/m³
Slovenia OEL - TWA 1000 mg/m³

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8 mg/m<sup>3</sup> 5 ppm

8.0 mg/m<sup>3</sup>

5 ppm 8 mg/m<sup>3</sup>

5 ppm 7.6 mg/m<sup>3</sup>

Material Name: Methylprednisolone Acetate Injectable Page 5 of 9

Suspension, Single-Dose Vial

Revision date: 06-Sep-2012 Version: 1.1

## 8. EXPOSURE CONTROLS / PERSONAL PROTECTION

Analytical Method:

Analytical method available for methylprednisolone. Contact Pfizer Inc for further information.

Engineering Controls:

Engineering controls should be used as the primary means to control exposures. Use process

Engineering controls should be used as the primary means to control exposures. Use process containment, local exhaust ventilation, or other engineering controls to maintain airborne levels

below recommended exposure limits.

Environmental Exposure Controls: Refer to specific Member State legislation for requirements under Community environmental

legislation.

Personal Protective Equipment: Refer to applicable national standards and regulations in the selection and use of personal

protective equipment (PPE).

Hands: Impervious gloves are recommended if skin contact with drug product is possible and for bulk

processing operations.

**Eyes:** Wear safety glasses or goggles if eye contact is possible.

**Skin:** Impervious protective clothing is recommended if skin contact with drug product is possible and

for bulk processing operations.

Respiratory protection: If the applicable Occupational Exposure Limit (OEL) is exceeded, wear an appropriate

respirator with a protection factor sufficient to control exposures to below the OEL.

## 9. PHYSICAL AND CHEMICAL PROPERTIES

Physical State:SolutionColor:ColorlessMolecular Formula:MixtureMolecular Weight:Mixture

**pH:** 3.5 to 7.0

Polymerization: Will not occur

### 10. STABILITY AND REACTIVITY

**Chemical Stability:** Stable under normal conditions of use.

**Conditions to Avoid:** Fine particles (such as dust and mists) may fuel fires/explosions. **Incompatible Materials:** As a precautionary measure, keep away from strong oxidizers

### 11. TOXICOLOGICAL INFORMATION

**General Information:** The information included in this section describes the potential hazards of the individual

ingredients. The information included in this section describes the potential hazards of various

forms of the active ingredient.

Acute Toxicity: (Species, Route, End Point, Dose)

Methylprednisolone

Rat Oral LD 50 > 2000 mg/kg Mouse Oral LD 50 450 mg/kg

Rat Intraperitoneal LD 50 1000 mg/kg Mouse Intraperitoneal LD 50 1409 mg/kg Rat Subcutaneous LD 50 >3000 mg/kg

**Methylprednisolone Acetate** 

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Material Name: Methylprednisolone Acetate Injectable Page 6 of 9

Suspension, Single-Dose Vial

Revision date: 06-Sep-2012 Version: 1.1

# 11. TOXICOLOGICAL INFORMATION

Rat Oral LD50 >10,000 mg/kg

Mouse Sub-tenon injection (eye) LD50 >1,409 mg/kg

Rat Subcutaneous LD50 265 mg/kg

#### Sodium hydroxide

Mouse IP LD50 40 mg/kg

#### Myristyl-gamma-picolinium chloride

Rat Oral LD 50 250 mg/kg

Rat Para-periosteal LD50 30 mg/kg
Rat Intraperitoneal LD50 7500 ug/kg
Rat Subcutaneous LD50 200 mg/kg

## Sodium chloride

Rat Oral LD50 3000 mg/kg Mouse Oral LD50 4000 mg/kg

Acute Toxicity Comments: A greater than symbol (>) indicates that the toxicity endpoint being tested was not achievable

at the highest dose used in the test.

## Irritation / Sensitization: (Study Type, Species, Severity)

#### Methylprednisolone

Skin Irritation Rabbit No effect Eve Irritation Rabbit No effect

Skin Sensitization - GPMT Guinea Pig No effect

#### Polyethylene glycol

Eye Irritation Rabbit Mild Skin Irritation Rabbit Mild

## **Methylprednisolone Acetate**

Eye Irritation Rabbit No effect Skin Irritation Rabbit No effect

#### Sodium hydroxide

Eye Irritation Rabbit Severe Skin Irritation Rabbit Severe

## **Hydrochloric Acid**

Skin Irritation Severe Eye Irritation Severe

## Sodium chloride

Eye Irritation Rabbit Moderate Skin Irritation Rabbit Mild

## Repeated Dose Toxicity: (Duration, Species, Route, Dose, End Point, Target Organ)

#### Methylprednisolone

42 Day(s) Dog Oral 167 μg/kg/day LOAEL Adrenal gland

6 Week(s) Rat Subcutaneous 500 μg/kg/day LOAEL None identified

14 Week(s) Rat Subcutaneous 0.4 µg/kg/day NOAEL Blood forming organs, Adrenal gland

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Material Name: Methylprednisolone Acetate Injectable Page 7 of 9

Suspension, Single-Dose Vial

Revision date: 06-Sep-2012 Version: 1.1

11. TOXICOLOGICAL INFORMATION

52 Week(s) Rat Subcutaneous 4 µg/kg/day NOAEL Blood forming organs Adrenal gland

Myristyl-gamma-picolinium chloride

60 Day(s) Rat Oral 2400 mg/kg Death

Reproduction & Development Toxicity: (Duration, Species, Route, Dose, End Point, Effect(s))

Methylprednisolone

Reproductive & Fertility Rat Subcutaneous 0.004 mg/kg/day Paternal toxicity NOAEL Reproductive & Fertility Rat Subcutaneous 0.02 mg/kg/day LOAEL Fetotoxicity

Embryo / Fetal Development Rat Subcutaneous 1.0 mg/kg/day LOAEL Fetotoxicity, Teratogenic

Embryo / Fetal Development Mouse Intramuscular 330 mg/kg/day LOAEL Teratogenic Embryo / Fetal Development Rabbit 0.1 mg/kg/day LOAEL Teratogenic Intramuscular

Genetic Toxicity: (Study Type, Cell Type/Organism, Result)

Methylprednisolone

**Bacterial Mutagenicity (Ames)** Salmonella Negative Unscheduled DNA Synthesis Rat Hepatocyte Negative

Mammalian Cell Mutagenicity Chinese Hamster Ovary (CHO) cells Negative

Direct DNA Interaction Negative

**Methylprednisolone Acetate** 

Not applicable Direct DNA Interaction Negative In Vitro Cytogenetics Not applicable Negative

Carcinogen Status: None of the components of this formulation are listed as a carcinogen by IARC, NTP or OSHA.

**Hydrochloric Acid** 

IARC: Group 3 (Not Classifiable)

12. ECOLOGICAL INFORMATION

Environmental properties have not been investigated. Releases to the environment should be **Environmental Overview:** 

avoided.

13. DISPOSAL CONSIDERATIONS

**Waste Treatment Methods:** Dispose of waste in accordance with all applicable laws and regulations. Member State

specific and Community specific provisions must be considered. Considering the relevant known environmental and human health hazards of the material, review and implement appropriate technical and procedural waste water and waste disposal measures to prevent occupational exposure and environmental release. It is recommended that waste minimization be practiced. The best available technology should be utilized to prevent environmental

releases. This may include destructive techniques for waste and wastewater.

14. TRANSPORT INFORMATION

The following refers to all modes of transportation unless specified below.

Not regulated for transport under USDOT, EUADR, IATA, or IMDG regulations.

Material Name: Methylprednisolone Acetate Injectable Page 8 of 9

Suspension, Single-Dose Vial

Revision date: 06-Sep-2012 Version: 1.1

# 15. REGULATORY INFORMATION

EU Symbol:

**EU Indication of danger:** Toxic to reproduction: Category 1

**EU Risk Phrases:** 

R61 - May cause harm to the unborn child.

**EU Safety Phrases:** 

S36/37 - Wear suitable protective clothing and gloves.

S53 - Avoid exposure - obtain special instructions before use.

**OSHA Label:** 

DANGER

May damage the unborn child.

Canada - WHMIS: Classifications

WHMIS hazard class:

Class D, Division 2, Subdivision A



**Methylprednisolone Acetate** 

Australia (AICS): Present EU EINECS/ELINCS List 200-171-3

Myristyl-gamma-picolinium chloride

Inventory - United States TSCA - Sect. 8(b)

Australia (AICS):

EU EINECS/ELINCS List

Present
220-387-1

Sodium chloride

Inventory - United States TSCA - Sect. 8(b)

Australia (AICS):

Present

EU EINECS/ELINCS List

231-598-3

Water for injection

Inventory - United States TSCA - Sect. 8(b)PresentAustralia (AICS):PresentREACH - Annex IV - Exemptions from thePresent

obligations of Register:

Material Name: Methylprednisolone Acetate Injectable Page 9 of 9

Suspension, Single-Dose Vial

Revision date: 06-Sep-2012 Version: 1.1

15. REGULATORY INFO	RMATION
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EU EINECS/ELINCS List 231-791-2

Sodium hydroxide

CERCLA/SARA Hazardous Substances
and their Reportable Quantities:
454 kg
Inventory - United States TSCA - Sect. 8(b)
Australia (AICS):
Present
Standard for the Uniform Scheduling
for Drugs and Poisons:
Schedule 5
EU EINECS/ELINCS List
1000 lb
Present
Stoel kg
Present
Schedule 5
Schedule 6
215-185-5

**Hydrochloric Acid** 

CERCLA/SARA 313 Emission reporting 1.0 %
CERCLA/SARA Hazardous Substances 5000 lb and their Reportable Quantities: 2270 kg
CERCLA/SARA - Section 302 Extremely Hazardous 500 lb

**TPQs** 

CERCLA/SARA - Section 302 Extremely Hazardous 5000 lb

**Substances EPCRA RQs** 

Inventory - United States TSCA - Sect. 8(b)PresentAustralia (AICS):PresentStandard for the Uniform SchedulingSchedule 5for Drugs and Poisons:Schedule 6EU EINECS/ELINCS List231-595-7

Polyethylene glycol

Inventory - United States TSCA - Sect. 8(b)

Australia (AICS):

Standard for the Uniform Scheduling

Present
Schedule 3

for Drugs and Poisons:

## 16. OTHER INFORMATION

#### Text of R phrases mentioned in Section 3

R61 - May cause harm to the unborn child. R20/22 - Harmful by inhalation and if swallowed.

R48/22 - Harmful: danger of serious damage to health by prolonged exposure if swallowed.

**Data Sources:** The data contained in this MSDS may have been gathered from confidential internal sources,

raw material suppliers, or from the published literature.

**Reasons for Revision:** Updated Section 2 - Hazard Identification.

Prepared by: Product Stewardship Hazard Communication

Pfizer Global Environment, Health, and Safety Operations

Pfizer Inc believes that the information contained in this Material Safety Data Sheet is accurate, and while it is provided in good faith, it is without warranty of any kind, expressed or implied. If data for a hazard are not included in this document there is no known information at this time.

**End of Safety Data Sheet** 

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