

Revision date: 26-Oct-2009

Version: 1.3

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IDENTIFICATION OF THE SUBSTANCE/PREPARATION AND THE COMPANY/UNDERTAKING 1.

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Material Name: Dinoprostone Endocervical Gel

Trade Name:	PREPIDIL; PROSTIN E2; MINPROSTIN
Chemical Family:	Mixture
Intended Use:	Pharmaceutical product used for smooth muscle stimulation

2. HAZARDS IDENTIFICATION

Appearance:	Colorless gel
Statement of Hazard:	Non-hazardous in accordance with international standards for workplace safety.
Long Term:	Repeat-dose studies in animals have shown a potential to cause adverse effects on the developing fetus.
Known Clinical Effects:	Clinical use of this drug has caused hot flashes diarrhea nausea vomiting May cause low blood pressure and dizziness. Uterine contractions, vaginal bleeding, and prevention/termination of pregnancy have been seen in women taking this drug.
EU Indication of danger:	Not classified

Australian Hazard Classification (NOHSC):	Non-Hazardous Substance. Non-Dangerous Goods.
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.

3. COMPOSITION/INFORMATION ON INGREDIENTS

Hazardous

Ingredient CAS Number EU EINECS/ELINCS List Classification	%
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3. COMPOSITION/INFORMATIO	ON ON INGREDIENTS	6		
Dinoprostone	363-24-6	206-656-6	Xn;R22	<0.1
Silica gel, amorphous	112926-00-8	Not listed	T;R60 Not Listed	*
Ingredient Triacetin	CAS Number 102-76-1	EU EINECS/ELINCS List 203-051-9	Classification Not Listed	<u>%</u> *
Additional Information:	* Proprietary Ingredient(s) indicated as hazardous have been assessed under standards for workplace safety.			
For the full text of the R phrases me 4. FIRST AID MEASURES	ntioned in this Section, se	ee Section 16		
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 cl	nemical, or water spray.		
Hazardous Combustion Products:	Carbon monoxide and car	bon dioxide		
Fire Fighting Procedures:	During all fire fighting activ contained breathing appar	vities, wear appropriate protec atus.	ctive equipment, inclu	ding self-
Fire / Explosion Hazards:	Fine particles (such as dust and mists) may fuel fires/explosions.			
6. ACCIDENTAL RELEASE ME	ASURES			
Health and Safety Precautions:	Personnel involved in clean-up should wear appropriate personal protective equipment (see Section 8). Minimize exposure.			
Measures for Cleaning / Collecting:	Use absorbent material to spill area thoroughly.	wipe up spill and place in a s	ealed container for di	sposal. Clean
Measures for Environmental Protections:	Place waste in an appropr avoid environmental relea	iately labeled, sealed contain se.	er for disposal. Care	should be taken to
Additional Consideration for Large Spills:		hould be evacuated from affe lean up operations should onl		

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7. HANDLING AND STORAGE

General Handling: Storage Conditions: 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. Store as directed by product packaging.

8. EXPOSURE CONTROLS / PERSONAL PROTECTION

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

Dinoprostone			
Pfizer OEL TWA-8 Hr:		0.5 μg/m³, Skin	
Silica gel, amorphous			
Australia TWA		10 mg/m ³	
Belgium OEL - TWA		Listed	
Finland OEL - TWA		Listed	
OSHA - Final PELs - Table Z-3	Mineral D:	- (80)/(% SiO2) mg/m ³ TWA TWA-20 mppcf	
Poland OEL - TWA		Listed	
Spain OEL - TWA		Listed	
The exposure limit(s) listed for solid co	mponents are only rel	evant if dust or mist may be generated.	
Analytical Method: Engineering Controls:	Analytical method available for Dinoprostone. Contact Pfizer Inc for further information. Engineering controls should be used as the primary means to control exposures. General room ventilation is adequate unless the process generates dust, mist or fumes. Keep airborne contamination levels below the exposure limits listed above in this section.		
Environmental Exposure Controls:	Refer to specific Me legislation.	mber State legislation for requirements under Community environmental	
Personal Protective Equipment:	Refer to applicable national standards and regulations in the selection and use of personal protective equipment (PPE).		
Hands:	Wear impervious glo	oves if skin contact is possible.	
Eyes:	Wear safety glasses or goggles if eye contact is possible.		
Skin:		hing when working with large quantities.	
Respiratory protection:		cupational Exposure Limit (OEL) is exceeded, wear an appropriate tection factor sufficient to control exposures to below the OEL.	

9. PHYSICAL AND CHEMICAL PROPERTIES

Physical State:GelColor:ColourlessMolecular Formula:MixtureMolecular Weight:Mixture

Polymerization:

Will not occur

10. STABILITY AND REACTIVITY

Stability: Conditions to Avoid: Incompatible Materials: Stable at normal conditions Fine particles (such as dust and mists) may fuel fires/explosions. 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.

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

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

Triacetin

Rat Oral LD 50 3000 mg/kg Mouse Oral LD 50 1100 mg/kg

Dinoprostone

RatOralLD 50500 mg/kgRatIntravenousLD 5059.5 mg/kgRatSubcutaneousLD 5031.6 mg/kgMouseOralLD 50750 mg/kgMouseIntravenousLD 5023.2 mg/kg

Dinoprostone

Skin Sensitization - GPMT Guinea Pig Negative

Reproduction & Developmental Toxicity: (Study Type, Species, Route, Dose, End Point, Effect(s))

Dinoprostone

Embryo / Fetal Development Oral 6 mg/kg LOAEL Fetotoxicity Mouse 6 mg/kg Embryo / Fetal Development Rat Oral LOAEL Fetotoxicitv Embryo / Fetal Development Rat Intraperitoneal 12.5 mg/kg/day LOEL Teratogenic

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

Dinoprostone

Bacterial Mutagenicity (Ames) Salmonella Negative Direct DNA Damage Negative Micronucleus Negative

Carcinogen Status:

Silica gel, amorphous IARC:

Group 3

12. ECOLOGICAL INFOR Environmental Overview:	MATION Environmental properties have not been investigated. Releases to the environment should be avoided.
13. DISPOSAL CONSIDE	RATIONS
Disposal Procedures:	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.

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14. TRANSPORT INFORMATION

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

15. REGULATORY INFORMATION

EU Indication of danger:

Not classified

OSHA Label:

Non-hazardous in accordance with international standards for workplace safety.

Canada - WHMIS: Classifications

WHMIS hazard class:

None required This product has been classified in accordance with the hazard criteria of the CPR and the MSDS contains all of the information required by the CPR.

Dinoprostone Standard for the Uniform Scheduling for Drugs and Poisons:	Schedule 4
EU EINECS/ELINCS List	206-656-6
Triacetin	
Inventory - United States TSCA - Sect. 8(b)	Listed
Australia (AICS):	Listed
EU EINECS/ELINCS List	203-051-9
Silica gel, amorphous Australia (AICS):	Listed

16. OTHER INFORMATION

Text of R phrases mentioned in Section 3

R22 - Harmful if swallowed. R60 - May impair fertility.

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Reasons for Revision:	Updated Section 1 - Identification of the Substance/Preparation and the Company/Undertaking. Updated Section 3 - Composition / Information on Ingredients.
Prepared by:	Toxicology and 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