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

Material Name: Phenytoin Sodium Solution for Injection

Trade Name: Dilantin®; Aurantin®; Epanutin®; Epamin®; Epelin®
Chemical Family: Mixture
Intended Use: Pharmaceutical product used for seizures and epilepsy.

2. COMPOSITION/INFORMATION ON INGREDIENTS

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>CAS Number</th>
<th>EU EINECS List</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phenytoin Sodium</td>
<td>630-93-3</td>
<td>211-148-2</td>
<td>5</td>
</tr>
<tr>
<td>Ethyl alcohol (ethanol)</td>
<td>64-17-5</td>
<td>200-578-6</td>
<td>9</td>
</tr>
<tr>
<td>Propylene glycol</td>
<td>57-55-6</td>
<td>200-338-0</td>
<td>*</td>
</tr>
<tr>
<td>SODIUM HYDROXIDE</td>
<td>1310-73-2</td>
<td>215-185-5</td>
<td>**</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>CAS Number</th>
<th>EU EINECS List</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Water for injection</td>
<td>7732-18-5</td>
<td>231-791-2</td>
<td>###</td>
</tr>
</tbody>
</table>

Additional Information:
* Proprietary
** to adjust pH
### as required
Ingredient(s) indicated as hazardous have been assessed under standards for workplace safety.

3. HAZARDS IDENTIFICATION

Appearance: Clear colorless solution.
Signal Word: DANGER

Statement of Hazard:
Harmful if swallowed.
Suspected of causing cancer.
Suspected of damaging the unborn child.
May cause damage to central nervous system through prolonged or repeated exposure.
Highly flammable liquid and vapor.
May cause eye irritation

Additional Hazard Information:
Short Term: Exposure to high concentrations may cause irritation, headache, drowsiness, and symptoms of alcohol intoxication.
**MATERIAL SAFETY DATA SHEET**

**Material Name:** Phenytoin Sodium Solution for Injection

**Revision date:** 18-Jan-2007

**Version:** 1.0

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**Long Term:**
Repeat-dose studies in animals have shown a potential to cause adverse effects on blood and blood forming organs, gastrointestinal system and liver. This product contains ethanol which can cause liver changes, central nervous system effects, and birth defects in the developing fetus.

**Known Clinical Effects:**
The most common adverse effects observed with clinical use of phenytoin are lack of appetite, headache, dizziness, transient nervousness, ataxia, slurred speech, decreased coordination, mental confusion, insomnia, and GI disturbances (nausea, vomiting, and constipation). IV administration has been associated with hypotension and CNS depression. Mild hypersensitivity reactions (skin rashes) are common. Effects on blood-forming organs and the liver have occurred rarely. Other less common effects include swollen lymph nodes, sore mouth and symptoms of dependence/withdrawal. There is an unconfirmed association between the use of anticonvulsants during pregnancy and an increased risk of birth defects. This material has been shown to be secreted in low concentrations in human breast milk.

**EU Indication of danger:**
- Carcinogenic: Category 3
- Toxic to Reproduction: Category 3
- F - Highly flammable

**EU Hazard Symbols:**

**EU Risk Phrases:**
- R40 - Limited evidence of a carcinogenic effect
- R63 - Possible risk of harm to the unborn child
- R11 - Highly flammable

**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.

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**4. FIRST AID MEASURES**

**Eye Contact:**
Immediately flush eyes with water for at least 15 minutes. If irritation occurs or persists, get medical attention.

**Skin Contact:**
Remove clothing and wash affected skin with soap and water. If irritation occurs or persists, get medical attention.

**Ingestion:**
Get medical attention. Do not induce vomiting unless directed by medical personnel. Never give anything by mouth to an unconscious person.

**Inhalation:**
Remove to fresh air. If not breathing, give artificial respiration. Get medical attention.

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**5. FIRE FIGHTING MEASURES**

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

**Hazardous Combustion Products:**
No data available

**Fire Fighting Procedures:**
Wear approved positive pressure, self-contained breathing apparatus and full protective turn out gear.
6. ACCIDENTAL RELEASE MEASURES

Health and Safety Precautions: Personnel involved in clean-up should wear appropriate personal protective equipment (see Section 8). Minimize exposure. Eliminate all sources of ignition and ventilate area using explosion-proof equipment.

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: Use with adequate ventilation. Avoid breathing vapor or mist. Flammable liquid and vapor-keep away from ignition sources and clean up spills promptly. Eliminate possible ignition sources (e.g., heat, sparks, flame, impact, friction, electricity), and follow appropriate grounding and bonding procedures. Avoid contact with eyes, skin, and clothing. Use appropriate personal protective equipment. Wash thoroughly after handling.

Storage Conditions: Protect from freezing. Protect from light.

Storage Temperature: Store at controlled room temperature 20-25°C (68-77°F)

8. EXPOSURE CONTROLS / PERSONAL PROTECTION

Phenytoin Sodium
Pfizer OEL TWA-8 Hr: 0.4 mg/m³

Ethyl alcohol (ethanol)
OSHA - Final PELS - TWAs: = 1000 ppm TWA
ACGIH Threshold Limit Value (TWA) = 1900 mg/m³ TWA
Australia TWA = 1000 ppm TWA
= 1880 mg/m³ TWA

Propylene glycol
Australia TWA = 10 mg/m³ TWA
= 150 ppm TWA
= 474 mg/m³ TWA

SODIUM HYDROXIDE
ACGIH Ceiling Threshold Limit: = 2 mg/m³ Ceiling
Australia PEAK = 2 mg/m³ Peak

The exposure limit(s) listed for solid components are only relevant if dust or mist may be generated.


Engineering Controls: Engineering controls should be used as the primary means to control exposures.

Personal Protective Equipment:
9. PHYSICAL AND CHEMICAL PROPERTIES:

<table>
<thead>
<tr>
<th>Physical State:</th>
<th>Solution</th>
</tr>
</thead>
<tbody>
<tr>
<td>Molecular Formula:</td>
<td>Mixture</td>
</tr>
<tr>
<td>Color:</td>
<td>Clear, colorless</td>
</tr>
<tr>
<td>Molecular Weight:</td>
<td>Mixture</td>
</tr>
</tbody>
</table>

10. STABILITY AND REACTIVITY

Stability: Stable under normal conditions of use.
Conditions to Avoid: Exposure to light and freezing. Prevent vapor accumulation. Keep away from heat, spark, flames and all other sources of ignition.
Incompatible Materials: Strong oxidizers
Polymerization: Will not occur

11. TOXICOLOGICAL INFORMATION

General Information: The information included in this section describes the potential hazards of the individual ingredients. The information in this section describes the hazards of various forms of the active ingredient.

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

**Propylene glycol**
- Mouse Oral LD50 22,000 mg/kg
- Rat Oral LD50 20,000 mg/kg
- Rabbit Dermal LD50 20,800 mg/kg

**Ethyl alcohol (ethanol)**
- Mouse Oral LD50 3450 mg/kg
- Rat Oral LD50 7060 mg/kg
- Rat Inhalation LC50 10h 20,000 ppm

**Phenytoin Sodium**
- Mouse Oral LD50 165 mg/kg
- Rat Oral LD50 1530 mg/kg
- Rat IV LD50 90 mg/kg
- Mouse IV LD 50 98 mg/kg

**Phenytoin**
- Mouse Oral LD50 150 mg/kg
- Rat Oral LD50 1635 mg/kg
- Rat Intravenous LD 50 96 mg/kg
- Rat IM LD 50 >337 mg/kg
- Rabbit Oral LD 50 >3000 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)

Propylene glycol
Skin Irritation Rabbit Mild
Eye Irritation Rabbit Mild

Ethyl alcohol (ethanol)
Eye Irritation Rabbit Severe

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

Phenytoin
2 Week(s) Rat Oral <3125 ppm/day NOEL Bone marrow
2 Week(s) Mouse Oral <125 ppm/day NOEL Central Nervous System
13 Week(s) Rat Oral 300 ppm/day NOEL None identified
13 Week(s) Mouse Oral 150 ppm/day NOEL Blood forming organs, Gastrointestinal system, Liver

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

Phenytoin
Embryo / Fetal Development Mouse Oral 75 mg/kg/day NOEL Maternal toxicity, Fetotoxicity, Teratogenic
Embryo / Fetal Development Mouse Oral 45 mg/kg/day NOEL Teratogenic
Embryo / Fetal Development Rabbit Oral 50 mg/kg/day NOEL Fetotoxicity, Teratogenic
Embryo / Fetal Development Monkey Oral 10 mg/kg/day NOEL Fetotoxicity, Teratogenic
Embryo / Fetal Development Mouse Subcutaneous <12.5 mg/kg/day NOEL Maternal Toxicity, Fetotoxicity, Teratogenic

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

Phenytoin
Bacterial Mutagenicity (Ames) Salmonella Negative
In Vitro Chromosome Aberration Chinese Hamster Ovary (CHO) cells Negative
In Vitro Chromosome Aberration Human Lymphocytes Negative
In Vivo Sister Chromatid Exchange Human Lymphocytes Positive
In Vivo Mitotic Spindle Assay Human Lymphocytes Negative

Carcinogenicity: (Duration, Species, Route, Dose, End Point, Effect(s))

Phenytoin
2 Year(s) Male Rat, in feed 50 mg/kg/day NOEL Benign neoplasms, Skin
2 Year(s) Mouse Oral, in feed 25 mg/kg/day NOEL Benign tumors, Liver
2 Year(s) Female Mouse Oral, in feed 60 ppm LOAEL Liver, neoplasms
2 Year(s) Female Rat Oral, in feed 240 ppm NOAEL Not carcinogenic

Carcinogen Status: See below

Phenytoin Sodium
IARC: Group 2B
OSHA: Present

Phenytoin
IARC: Group 2B
NTP: Reasonably Anticipated To Be A Carcinogen
12. ECOLOGICAL INFORMATION

Environmental Overview: The environmental characteristics of this mixture have not been fully evaluated. Releases to
the environment should be avoided. See aquatic toxicity data, below:

Aquatic Toxicity: (Species, Method, End Point, Duration, Result)

Ethyl alcohol (ethanol)
Rainbow Trout LC50/96h 12,900-15,300 mg/L

Phenytoin
Hyalella azteca (Freshwater Amphipod) OPPTS LC50 96 Hours 18 mg/L
Daphnia Magna (Water Flea) TAD EC50 48 Hours >39 mg/L
Pimephales promelas (Fathead Minnow) OPPTS LC50 96 Hours >23 mg/L

Aquatic Toxicity Comments: A greater than (>) symbol indicates that acute ecotoxicity was not observed at the maximum
solubility. Since the substance is insoluble in aqueous solutions above this concentration, an
acute ecotoxicity value (i.e. LC(EC)50) is not achievable.

13. DISPOSAL CONSIDERATIONS

Disposal Procedures: Dispose of waste in accordance with all applicable laws and regulations.

14. TRANSPORT INFORMATION

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

15. REGULATORY INFORMATION

EU Symbol: Xn F+
EU Indication of danger: Carcinogenic: Category 3
Toxic to Reproduction; Category 3
F - Highly flammable

EU Risk Phrases: R40 - Limited evidence of a carcinogenic effect
R63 - Possible risk of harm to the unborn child.
R11 - Highly flammable.

EU Safety Phrases: S36/37 - Wear suitable protective clothing and gloves.
S16 - Keep away from sources of ignition - No smoking.
OSHA Label:
DANGER
Harmful if swallowed.
Suspected of causing cancer.
Suspected of damaging the unborn child.
May cause damage to central nervous system through prolonged or repeated exposure.
Highly flammable liquid and vapor.
May cause eye irritation

Canada - WHMIS: Classifications

WHMIS hazard class:
D2a very toxic materials
D2b toxic materials

Phenytoin Sodium
California Proposition 65
carcinogen, initial date 1/1/88
Inventory - United States TSCA - Sect. 8(b) Present
Australia (AICS): Present
EU EINECS List 211-148-2

Ethyl alcohol (ethanol)
California Proposition 65
developmental toxicity, initial date 10/1/87 (when in alcoholic beverages)
Inventory - United States TSCA - Sect. 8(b) Present
Australia (AICS): Present
EU EINECS List 200-578-6

Propylene glycol
Inventory - United States TSCA - Sect. 8(b) Present
Australia (AICS): Present
EU EINECS List 200-338-0

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

Water for injection
Inventory - United States TSCA - Sect. 8(b) Present
Australia (AICS): Present
EU EINECS List 231-791-2

16. OTHER INFORMATION