

PRODUCT MONOGRAPH

Fr **DIPRIVAN[®] 1% w/v**

propofol injection

10 mg/mL

Intravenous Emulsion – Anaesthetic - Sedative

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PrDIPRIVAN[®]

Propofol injection 10 mg/mL

Intravenous Emulsion - Anaesthetic - Sedative

PART I: HEALTH PROFESSIONAL INFORMATION

SUMMARY PRODUCT INFORMATION

Route of Administration	Dosage Form / Strength	All Nonmedicinal Ingredients
Parenteral	Intravenous Emulsion/ 10 mg/ml	Disodium edetate, egg phosphatide, glycerol, sodium hydroxide, soybean oil and water for injection.

INDICATIONS AND CLINICAL USE

Adults (>18 years of age):

DIPRIVAN (propofol) is indicated for:

- Induction and maintenance of general anaesthesia
- Conscious sedation for surgical and diagnostic procedures
- Sedation during intensive care

DIPRIVAN is a short-acting i.v. general anaesthetic agent, that can be used for both induction and maintenance of anaesthesia as part of a balanced anaesthesia technique, including total intravenous anaesthesia (TIVA), for inpatient and outpatient surgery.

DIPRIVAN, when administered i.v. as directed, can be used to initiate and maintain sedation in conjunction with local/regional anaesthesia in **adult** patients undergoing surgical procedures. DIPRIVAN may also be used for sedation during diagnostic procedures in **adults** (see WARNINGS AND PRECAUTIONS, General).

DIPRIVAN should only be administered to intubated, mechanically ventilated, **adult** patients in the Intensive Care Unit (ICU) to provide continuous sedation and control of stress responses. In this setting, DIPRIVAN should be administered only by or under the supervision of persons trained in general anaesthesia or critical care medicine.

Geriatrics (> 65 years of age):

Elderly patients should be given reduced doses of propofol, commensurate with their age and physical condition (see WARNINGS AND PRECAUTIONS, Special Populations and DOSAGE AND ADMINISTRATION).

Pediatrics (≥ 3 years of age):

DIPRIVAN is **only** indicated for anaesthesia in children 3 years of age and older.

Pediatrics (≤ 18 years of age):

DIPRIVAN is not recommended for sedation or during surgical/diagnostic procedures in children under the age of 18, as safety and efficacy have not been established in this patient population. (see WARNINGS AND PRECAUTIONS, Special Populations and DOSAGE AND ADMINISTRATION).

CONTRAINDICATIONS

DIPRIVAN (propofol) is contraindicated:

- for the sedation of children 18 years or younger receiving intensive care (see DOSAGE AND ADMINISTRATION)
- when sedation or general anaesthesia are contraindicated
- in patients with a known allergy and/or hypersensitivity to DIPRIVAN or to lipid emulsions (see DOSAGE FORMS, COMPOSITION AND PACKAGING, Composition).

WARNINGS AND PRECAUTIONS

General

Strict aseptic techniques must always be maintained during handling as DIPRIVAN (propofol) is a single-use parenteral product, for use in an individual patient, and contains no antimicrobial preservatives. The vehicle is capable of supporting rapid growth of microorganism (see DOSAGE AND ADMINISTRATION). Failure to follow aseptic handling procedures may result in microbial contamination causing fever/infection/sepsis, which could lead to life-threatening illness.

For general anaesthesia or sedation for surgical/diagnostic procedures, DIPRIVAN should be administered only by persons trained in the administration of general

anaesthesia and not involved in the conduct of surgical/diagnostic procedures. Patients should be continuously monitored and facilities for maintenance of a patent airway, artificial ventilation, and oxygen enrichment and circulatory resuscitation must be immediately available.

For sedation of intubated, mechanically ventilated, adult patients in the Intensive Care Unit (ICU), DIPRIVAN should be administered only by persons trained in general anaesthesia or critical care medicine.

Propofol Infusion Syndrome (PRIS)

Use of DIPRIVAN Injectable Emulsion infusions for both adult and pediatric ICU sedation has been associated with a constellation of metabolic derangements and organ system failures, referred to as Propofol Infusion Syndrome, that have resulted in death.

The syndrome is characterized by severe metabolic acidosis, hyperkalemia, lipemia, rhabdomyolysis, hepatomegaly, cardiac and renal failure. The syndrome is most often associated with prolonged, high-dose infusions (> 5 mg/kg/h for > 48h) but has also been reported following large-dose, short-term infusions during surgical anesthesia. The following appear to be the major risk factors for the development of these events: decreased oxygen delivery to tissues; serious neurological injury and/or sepsis; high dosages of one or more of the following pharmacological agents - vasoconstrictors, steroids, inotropes and/or propofol.

In the elderly, debilitated and ASA III or IV patients, rapid (single or repeated) bolus administration should not be used during general anaesthesia or sedation in order to minimize undesirable cardiorespiratory depression including hypotension, apnea, airway obstruction and/or oxygen desaturation.

Extreme care should be used in administering DIPRIVAN in elderly, debilitated or other ASA III or IV patients.

Very rarely reports of metabolic acidosis, rhabdomyolysis, hyperkalaemia, and/or cardiac failure, in some cases with a fatal outcome, have been received concerning seriously ill patients receiving propofol for ICU sedation (see ADVERSE REACTIONS, Post-Market Adverse Drug Reactions). The following appear to be the major risk factors for the development of these events: decreased oxygen delivery to tissues; serious neurological injury and/or sepsis; high doses of one or more of the following pharmacological agents – vasoconstrictors, steroids, inotropes and/or propofol. All sedatives and therapeutic agents used in the ICU (including propofol) should be titrated to maintain optimal oxygen delivery and haemodynamic parameters.

Extreme care should be used in administering DIPRIVAN in patients with impaired left ventricular function because DIPRIVAN may produce a negative inotropic effect.

Extreme care should be used in administering DIPRIVAN in patients who are hypotensive, hypovolemic or in shock because DIPRIVAN may cause excessive arterial hypotension.

DIPRIVAN lacks vagolytic activity and has been associated with reports of bradycardia, (occasionally profound) and also asystole. The intravenous administration of an anticholinergic agent before induction, or during maintenance of anaesthesia should be considered, especially in situations where vagal tone is likely to predominate or when DIPRIVAN is used in conjunction with other agents likely to cause a bradycardia.

DIPRIVAN should not be co-administered through the same i.v. catheter with blood or plasma because compatibility has not been established. In vitro tests have shown that aggregates of the globular component of the emulsion vehicle have occurred with blood/plasma/serum from humans and animals. The clinical significance is not known.

The neuromuscular blocking agents, atracurium and mivacurium should not be given through the same i.v. line as DIPRIVAN without prior flushing.

The administration of DIPRIVAN should be initiated as a continuous infusion and changes in the rate of administration made slowly (>5 min) in order to minimize hypotension and avoid acute overdosage.

Since DIPRIVAN is formulated in an oil-water emulsion, patients should be monitored for lipemia. Administration of DIPRIVAN should be adjusted if fat is being inadequately cleared from the body. A reduction in the quantity of concurrently administered lipids is indicated to compensate for the amount of lipid infused as part of the DIPRIVAN formulation; 1.0 mL of DIPRIVAN contains approximately 0.1 g of fat (1.1 kcal).

In adults and children, attention should be paid to minimize pain on administration of propofol. Transient local pain during intravenous injection may be reduced by prior injection of i.v. lidocaine (1.0 mL of a 1% solution).

Cardiovascular

Patients should be monitored for early signs of significant hypotension and/or cardiovascular depression, which may be profound. These effects are responsive to discontinuation of DIPRIVAN, i.v. fluid administration, and/or vasopressor therapy.

Cardiac Anaesthesia

DIPRIVAN was evaluated in 328 patients undergoing coronary artery bypass graft (CABG). Of these patients 85% were males (mean age 61, range 32-83) and 15% were females (mean age 65, range 42-86).

The majority of patients undergoing CABG had good left ventricular function. Experience in patients with poor left ventricular function, as well as, in patients with hemodynamically significant valvular or congenital heart disease is limited.

Slower rates of administration should be utilized in premedicated patients, geriatric patients, patients with recent fluid shift, or patients who are hemodynamically unstable. Any fluid deficits should be corrected prior to administration of DIPRIVAN. In those patients where additional fluid therapy may be contraindicated, other measures, e.g. elevation of lower extremities, or use of pressor agents, may be useful to offset the hypotension which is associated with the induction of anaesthesia with DIPRIVAN.

Endocrine and Metabolism

DIPRIVAN should not be used for Intensive Care Unit (ICU) sedation in patients who have severely disordered fat metabolism because the vehicle of DIPRIVAN is similar to that of INTRALIPID 10%. The restrictions that apply to INTRALIPID 10% should also be considered when using DIPRIVAN in the ICU.

Hematologic

EDTA is a chelator of metal ions, including zinc. The need for supplemental zinc should be considered during prolonged administration of DIPRIVAN, particularly in patients who are predisposed to zinc deficiency, such as those with burns, diarrhoea and/or major sepsis.

Hepatic/Biliary/Pancreatic

The long-term administration of DIPRIVAN to patients with hepatic insufficiency has not been evaluated.

Neurologic

Neurosurgical Anaesthesia

When using DIPRIVAN in patients with increased intracranial pressure (ICP) or impaired cerebral circulation, significant decreases in mean arterial pressure should be avoided because of the resultant decreases in cerebral perfusion pressure. When increased ICP is suspected, hyperventilation and hypocarbia should accompany the administration of DIPRIVAN (see DOSAGE AND ADMINISTRATION).

Epilepsy: Since various manifestations of seizures have been reported during DIPRIVAN anaesthesia, special care should be taken when giving the drug to epileptic patients.

Locomotion and Coordination: Patients receiving DIPRIVAN on an outpatient basis should not engage in hazardous activities requiring complete mental alertness such as driving a motor vehicle or operating machinery until the effects of DIPRIVAN have completely subsided.

Peri-Operative Considerations

As with other sedative medications, there is wide interpatient variability in DIPRIVAN dosage requirements, and these requirements may change with time.

Patients who receive large doses of narcotics during surgery may require very small doses of DIPRIVAN for appropriate sedation.

When DIPRIVAN is administered as a sedative for surgical or diagnostic procedures, patients should be continuously monitored by persons not involved in the conduct of the surgical/diagnostic procedure. Oxygen supplementation should be immediately available and provided where clinically indicated; and oxygen saturation should be monitored in all patients. Patients should be continuously monitored for early signs of hypotension, apnea, airway obstruction and/or oxygen desaturation. These cardiorespiratory effects are more likely to occur following rapid initiation (loading) boluses or during supplemental maintenance boluses, especially in the elderly, debilitated and ASA III or IV patients.

Patients should be continuously monitored for early signs of significant hypotension and/or bradycardia. Treatment may include increasing the rate of intravenous fluid, elevation of lower extremities, use of pressor agents or administration of anticholinergic agents (e.g. atropine) or use of plasma volume expanders. Apnea often occurs during induction and may persist for more than 60 seconds. Ventilatory support may be required. Because DIPRIVAN is a lipid emulsion, caution should be exercised in patients with disorders of lipid metabolism such as primary hyperlipoproteinemia, diabetic hyperlipemia and pancreatitis.

As with other sedative agents, when DIPRIVAN is used for sedation during operative procedures, involuntary patient movements may occur. During procedures requiring immobility these movements may be hazardous to the operative site.

Abrupt discontinuation of DIPRIVAN infusion prior to weaning should be avoided since, due to the rapid clearance of DIPRIVAN, it may result in rapid awakening with associated anxiety, agitation and resistance to mechanical ventilation. Infusions of DIPRIVAN should be adjusted to maintain a light level of sedation throughout the weaning process.

Since DIPRIVAN is rarely used alone, an adequate period of evaluation of the awakened patient is indicated to ensure satisfactory recovery from general anaesthesia or sedation prior to discharge of the patient from the recovery room or to home. Very rarely the use of DIPRIVAN may be associated with the development of a period of post-operative unconsciousness, which may be accompanied by an increase in muscle tone. This may or may not be preceded by a period of wakefulness. Although recovery is spontaneous, appropriate care of an unconscious patient should be administered.

Renal

The long-term administration of DIPRIVAN to patients with renal failure has not been evaluated.

Special Populations

Pregnant Women: DIPRIVAN should not be used during pregnancy. DIPRIVAN has been used during termination of pregnancy in the first trimester. Teratology studies in rats and rabbits show some evidence of delayed ossification or abnormal cranial ossification, however such developmental delays are not considered indicative of a teratogenic effect. Reproductive

studies in rats suggest that administration of DIPRIVAN to the dam adversely affects perinatal survival of the offspring.

Labour and Delivery: DIPRIVAN should not be used in obstetrics including Caesarean section deliveries, because DIPRIVAN crosses the placenta and may be associated with neonatal depression.

Nursing Women: DIPRIVAN is not recommended for use in nursing women because preliminary findings indicate that it is excreted in human milk and the effects of oral absorption of small amounts of DIPRIVAN are not known.

Pediatrics (≤18 years of age): In the absence of sufficient clinical experience, DIPRIVAN is not recommended for anaesthesia in children less than 3 years of age (see INDICATIONS AND CLINICAL USE and DOSAGE AND ADMINISTRATION).

Geriatrics (> 65 years of age): Elderly patients may be more sensitive to the effects of DIPRIVAN; therefore, the dosage of DIPRIVAN should be reduced in these patients according to their condition and clinical response (see DETAILED PHARMACOLOGY, Pharmacokinetics and DOSAGE AND ADMINISTRATION).

ADVERSE REACTIONS

Adverse Drug Reaction Overview

Anaesthesia and Sedation For Surgical/Diagnostic Procedures

During induction of anaesthesia in clinical trials, hypotension and apnea occurred in the majority of patients. The incidence of apnea varied considerably, occurring in between 30 and 100% of patients depending upon premedication, speed of administration and dose (see ACTION AND CLINICAL PHARMACOLOGY). Decreases in systolic and diastolic pressures ranged between 10 and 28%, but were more profound in the elderly and in ASA III and IV patients. Excitatory phenomena occurred in up to 14% of adult patients and in 33 to 90% of pediatric patients; they consisted most frequently of spontaneous musculoskeletal movements and twitching and jerking of the hands, arms, feet or legs. Epileptiform movements including convulsions and opisthotonus have occurred rarely, but a causal relationship with DIPRIVAN (propofol) has not been established. Flushing and rash have occurred in 10 to 25% of pediatric patients. Local pain occurred during intravenous injection of DIPRIVAN at an incidence of 28% when veins of the dorsum of the hand were used and 5% when the larger veins of the forearm and the antecubital fossa were used. DIPRIVAN increased plasma glucose concentrations significantly, but no other significant changes in hematological or biochemical values were observed.

In the sedation clinical trials, the adverse reaction profile of DIPRIVAN was similar to that seen during anaesthesia. The most common adverse reactions included hypotension, nausea, pain and/or hotness at injection site and headache. Respiratory events included upper airway obstruction, apnea, hypoventilation, dyspnea and cough.

Rarely, clinical features of anaphylaxis, which may include angioedema, bronchospasm, erythema and hypotension, occur following DIPRIVAN administration.

Very rarely the use of DIPRIVAN may be associated with the development of a period of post-operative unconsciousness, which may be accompanied by an increase in muscle tone. This may or may not be preceded by a period of wakefulness.

There have been reports of post-operative fever.

Pulmonary oedema may be a potential side effect associated with the use of DIPRIVAN.

As with other anaesthetics, sexual disinhibition may occur during recovery.

Intensive Care Unit (ICU) Sedation - Adults

The most frequent adverse reactions during Intensive Care Unit (ICU) sedation were hypotension (31.5%), hypoxia (6.3%), and hyperlipemia (5.5%). In some patients, hypotension was severe. Other reactions considered severe were observed in single patients and included ventricular tachycardia, decreased cardiac output, decrease in vital capacity and negative inspiratory force, increase in triglycerides, and agitation. Two patients with head injury suffered renal failure with severe increases in BUN accompanied in one patient by an increase in creatinine.

There have been very rare reports of rhabdomyolysis when DIPRIVAN has been administered at doses greater than 4 mg/kg/hr for ICU sedation.

Very rarely pancreatitis has been observed following the use of DIPRIVAN for induction and maintenance of anaesthesia, and for intensive care sedation. A causal relationship has not been clearly established.

Clinical Trial Adverse Drug Reactions

Because clinical trials are conducted under very specific conditions the adverse reaction rates observed in the clinical trials may not reflect the rates observed in practice and should not be compared to the rates in the clinical trials of another drug. Adverse drug reaction information from clinical trials is useful for identifying drug-related adverse events and for approximating rates.

The following table compares the overall occurrence rates of adverse reactions in DIPRIVAN patients from non-ICU and ICU clinical trials where the rate of occurrence was greater than 1%. Major differences include lack of metabolic/nutritional (hyperlipemia) and respiratory events in the non-ICU group and lack of nausea, vomiting, headache, movement and injection site events in the ICU group.

Table 1 Non-ICU vs. ICU adverse events occurring in greater than 1% of DIPRIVAN patients.

Body System	Event	Non-ICU	ICU
Number of patients		2588	127
Cardiovascular	Hypotension	7.38%	31.50%
	Bradycardia	2.82%	3.94%
	Hypertension	2.82%	1.57%
	Arrhythmia	1.24%	0.79%
	Tachycardia	0.81%	3.15%
	Cardiovascular Disorder	0.23%	2.36%
	Hemorrhage	0.23%	1.57%
	Atrial Fibrillation	0.15%	1.57%
	Cardiac Arrest	0.12%	3.15%
	Ventricular Tachycardia	0.08%	1.57%
Digestive	Nausea	14.57%	0%
	Vomiting	8.31%	0%
	Abdominal Cramping	1.24%	0%
Nervous	Movement	4.44%	0%
	Headache	1.78%	0%
	Dizziness	1.70%	0%
	Twitching	1.47%	0%
	Agitation	0.19%	2.36
	Intracranial Hypertension	0%	3.94%
Metabolic/Nutritional	Hyperlipemia	0.08%	5.51%
	Acidosis	0.04%	1.57%
	Creatinine Increased	0%	2.36%
	BUN Increased	0%	1.57%
	Hyperglycemia	0%	1.57%
	Hypernatremia	0%	1.57%
	Hypokalemia	0%	1.57%
Respiratory	Dyspnea	0.43%	1.57%
	Hypoxia	0.08%	6.30%
	Acidosis	0%	1.57%
	Pneumothorax	0%	1.57%

Body System	Event	Non-ICU	ICU
Other	Injection Site:		
	Pain	8.11%	0%
	Burning/stinging	7.77%	0%
	Fever	1.89%	2.36%
	Hiccough	1.78%	0%
	Cough	1.55%	0%
	Rash	1.20%	1.57%
	Anemia	0.35%	1.57%
	Kidney Failure	0%	1.57%

Less Common Clinical Trial Adverse Drug Reactions ($\leq 1.0\%$) reported during anaesthesia and sedation for surgical/diagnostic procedures:

Cardiovascular System

Significant hypotension, premature atrial contractions, premature ventricular contractions, tachycardia, syncope, abnormal ECG, bigeminy, edema.

Respiratory System

Burning in throat, tachypnea, dyspnea, upper airway obstruction, wheezing, bronchospasm, laryngospasm, hypoventilation, hyperventilation, sneezing.

Excitatory

Hypertonia, dystonia, rigidity, tremor.

Central Nervous System

Confusion, dizziness, paresthesia, somnolence, shivering, abnormal dreams, agitation, delirium, euphoria, fatigue.

Injection Site

Phlebitis, thrombosis, hives/itching, redness/discolouration.

Digestive System

Hypersalivation, dry mouth.

Skin and Appendages

Flushing/rash (for incidence in children, see above), urticaria, pruritus.

Special Senses

Diplopia, amblyopia, tinnitus.

Musculoskeletal

Myalgia.

Urogenital

Urine retention, discolouration of urine.

Less Common Adverse Drug Reactions ($\leq 1\%$) reported during ICU sedation.

Cardiovascular

Arrhythmia, extrasystole, heart block, right heart failure, bigeminy, ventricular fibrillation, heart failure, myocardial infarction.

Respiratory

Lung function decreased, respiratory arrest.

Central Nervous System

Seizure, thinking abnormal, akathisia, chills, anxiety, confusion, hallucinations.

Digestive

Ileus, hepatomegaly.

Metabolic/Nutritional

Osmolality increased.

Urogenital

Green urine, urination disorder, oliguria.

Body as a Whole

Sepsis, trunk pain, whole body weakness.

Post-Market Adverse Drug Reactions

Clinical Trial Adverse Drug Reactions

A randomised, controlled, clinical trial that evaluated the safety and effectiveness of DIPRIVAN versus standard sedative agents (SSA) in pediatric ICU patients has been conducted. In that study, a total of 327 pediatric patients were randomised to receive either DIPRIVAN 2% (113 patients), DIPRIVAN 1% (109 patients), or an SSA (e.g. lorazepam, chloral hydrate, fentanyl, ketamine, morphine, or phenobarbital).

DIPRIVAN therapy was initiated at an infusion rate of 5.5 mg/kg/hr and titrated as needed to maintain sedation at a standardized level. The results of the study showed an increase in the number of deaths in patients treated with DIPRIVAN as compared to SSAs. A total of 25 patients died during the trial or within the 28-day follow-up period: 12 (11%) in the DIPRIVAN 2% treatment group, 9 (8%) in the DIPRIVAN 1% treatment group, and 4 (4%) in the SSA treatment group.

Spontaneous Reports and Publications

Propofol Infusion Syndrome (PRIS)

There are several publications identifying an association in adults between high infusion rates (greater than 5 mg/kg/h) of propofol for more than 48 hours in ICUs and a potentially fatal constellation of adverse events characterized by metabolic acidosis, rhabdomyolysis, hyperkalaemia, and cardiovascular collapse (see WARNINGS AND PRECAUTIONS).

The majority of the above-reported cases occurred in adults with head injury. These patients were treated with propofol at infusion rates greater than 5 mg/kg/h in an attempt to control intracranial hypertension. It is unclear at this time whether propofol at these high infusion rates can provide enhanced intracranial pressure reduction. A causal relationship between these adverse events and propofol and/or the lipid carrier cannot yet be established.

Similar findings were first reported in the literature in 1992 in children who received high doses of propofol in the ICU. Since the 1992 publication, several similar reports have been published, including an article that summarized 18 cases of children who received propofol infusions and suffered serious adverse events, including death.

Drug Abuse and Dependence

Rare cases of self administration of DIPRIVAN by health care professionals have been reported, including some fatalities.

DRUG INTERACTIONS

Overview

DIPRIVAN (propofol) has been used in association with spinal and epidural anaesthesia and with a range of premedicants, muscle relaxants, inhalational agents, analgesic agents and with local anaesthetic agents; no significant adverse interactions have been observed.

Drug-Food Interactions

Interactions of propofol with food have not been established.

Drug-Herb Interactions

Interactions of propofol with herbal products have not been established.

Drug-Laboratory Interactions

Interactions of propofol with laboratory tests have not been established.

Drug-Lifestyle Interactions

Interactions of propofol with lifestyle have not been established.

DOSAGE AND ADMINISTRATION

Dosing Considerations

General

Strict aseptic techniques must always be maintained during handling as DIPRIVAN (propofol)

is a single-use parenteral product, for use in an individual patient, and contains no antimicrobial preservatives. The vehicle is capable of supporting rapid growth of microorganisms. Failure to follow aseptic handling procedures may result in microbial contamination causing fever/infection/sepsis which could lead to life-threatening illness.

DIPRIVAN should be visually inspected for particulate matter, emulsion separation and/or discoloration prior to use. Do not use if any of these things are seen. If no signs of particulate matter, emulsion separation and/or discoloration are seen, shake gently before use.

Dosage and rate of administration should be individualized and titrated to the desired effect according to clinically relevant factors including preinduction and concomitant medications, age, ASA status and level of debilitation of the patient. In heavily premedicated patients, both the induction and maintenance doses should be reduced.

Recommended Dose and Dosage Adjustment

INDUCTION OF GENERAL ANAESTHESIA

As with most anaesthetic agents, the effects of DIPRIVAN may be potentiated in patients who have received intravenous sedative or narcotic premedications shortly prior to induction.

Adults (< 55 years of age):

Most *adult* patients under 55 years of age and classified ASA I and II are likely to require 2.0 to 2.5 mg/kg of DIPRIVAN for induction when unpremedicated or when premedicated with oral benzodiazepines or intramuscular narcotics. For induction, it is recommended that DIPRIVAN should be titrated (approximately 40 mg every 10 seconds by bolus injection or infusion) against the response of the patient until the clinical signs show the onset of general anaesthesia.

Geriatric, debilitated and adults ASA Classes III and IV

It is important to be familiar and experienced with the appropriate intravenous use of DIPRIVAN before treating *elderly, debilitated and/or adult patients in ASA Physical Status Classes III and IV*. These patients may be more sensitive to the effects of DIPRIVAN; therefore, the dosage of DIPRIVAN should be reduced in these patients by approximately 50% (20 mg every 10 seconds) according to their condition and clinical response. A rapid bolus should not be used as this will increase the likelihood of undesirable cardiorespiratory depression including hypotension, apnea, airway obstruction and/or oxygen desaturation (see WARNINGS AND PRECAUTIONS and DOSAGE AND ADMINISTRATION, Induction of General Anaesthesia - Dosage Guide).

During *cardiac anaesthesia*, a rapid bolus induction should be avoided. A slow rate of approximately 20 mg every 10 seconds until induction onset (0.5 to 1.5 mg/kg) should be used.

Pediatrics (3-18 years of age):

Most children over 8 years of age require approximately 2.5 mg/kg of DIPRIVAN for induction of anaesthesia. Children 3 to 8 years of age may require somewhat higher doses, however the dose should be titrated by administering DIPRIVAN slowly until the clinical signs show the onset of anaesthesia. Reduced dosage is recommended for children of ASA Classes III and IV.

Pediatrics < 3 years of age

DIPRIVAN is not recommended for induction of anaesthesia in children less than 3 years of age.

Dosage Guide for Induction of General Anaesthesia

Dosage should be individualized

<i>Adult Patients < 55 Years of Age</i>	Are likely to require 2.0 to 2.5 mg/kg (approximately 40 mg every 10 seconds until induction onset).
<i>Elderly, Debilitated and/or Adult ASA III or IV Patients</i>	Are likely to require 1.0 to 1.5 mg/kg (approximately 20 mg every 10 seconds until induction onset) but dose should be carefully titrated to effect.
<i>Cardiac Anaesthesia</i>	Patients are likely to require 0.5 to 1.5 mg/kg (approximately 20 mg every 10 seconds until induction onset).
<i>Neurosurgical Patients</i>	Are likely to require 1.0 to 2.0 mg/kg (approximately 20 mg every 10 seconds until induction onset).
<i>Pediatric Patients 3-8 and 8-18 Years of Age</i>	Children over 8 years of age require approximately 2.5 mg/kg. Children 3 to 8 years of age may require somewhat higher doses but doses should be titrated slowly to the desired effect. Reduced dosage is recommended for children of ASA Classes III and IV.

<i>Pediatric Patients <3 Years of Age</i>	In the absence of sufficient clinical experience, DIPRIVAN is not recommended for induction of anaesthesia in children less than 3 years of age (see INDICATIONS AND CLINICAL USE and WARNINGS AND PRECAUTIONS).
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MAINTENANCE OF GENERAL ANAESTHESIA

Anaesthesia can be maintained by administering DIPRIVAN by infusion or intermittent i.v. bolus injection. The patient's clinical response will determine the infusion rate or the amount and frequency of incremental injections.

When administering DIPRIVAN by infusion, drop counters, syringe pumps or volumetric pumps must be used to provide controlled infusion rates.

Continuous Infusion

DIPRIVAN 0.10 to 0.20 mg/kg/min (6 - 12 mg/kg/h) administered in a variable rate infusion with 60% - 70% nitrous oxide and oxygen provides anaesthesia for patients undergoing general surgery. Maintenance by infusion of DIPRIVAN should immediately follow the induction dose in order to provide satisfactory or continuous anaesthesia during the induction phase. During this initial period following the induction injection higher rates of infusion are generally required (0.15 - 0.20 mg/kg/min; 9 - 12 mg/kg/h) for the first 10 to 15 minutes. Infusion rates should subsequently be decreased by 30% - 50% during the first half-hour of maintenance. Changes in vital signs (increases in pulse rate, blood pressure, sweating and/or tearing) that indicate a response to surgical stimulation or lightening of anaesthesia may be controlled by the administration of DIPRIVAN 25 mg (2.5 mL) to 50 mg (5.0 mL) incremental boluses and/or by increasing the infusion rate. If vital sign changes are not controlled after a five minute period, other means such as a narcotic, barbiturate, vasodilator or inhalation agent therapy should be initiated to control these responses.

For minor surgical procedures (i.e. body surface) 60% to 70% nitrous oxide can be combined with a variable rate DIPRIVAN infusion to provide satisfactory anaesthesia. With more stimulating surgical procedures (i.e. intra-abdominal) supplementation with i.v. analgesic agents should be considered to provide a satisfactory anaesthetic and recovery profile. When supplementation with nitrous oxide is not provided, administration rate(s) of DIPRIVAN and/or opioids should be increased in order to provide adequate anaesthesia.

Infusion rates should always be titrated downward in the absence of clinical signs of light anaesthesia until a mild response to surgical stimulation is obtained in order to avoid administration of DIPRIVAN at rates higher than are clinically necessary. Generally, rates of 0.05 to 0.10 mg/kg/min should be achieved during maintenance in order to optimize recovery times.

During *cardiac anaesthesia*, when DIPRIVAN is used as the primary agent, maintenance infusion rates should not be less than 0.10 mg/kg/min and should be supplemented with analgesic levels of continuous opioid administration. When an opioid is used as the primary agent, DIPRIVAN maintenance rates should not be less than 0.05 mg/kg/min. Higher doses of DIPRIVAN will reduce the opioid requirements.

For *children*, the average rate of administration varies considerably but rates between 0.10 to 0.25 mg/kg/min (6 - 15 mg/kg/h) should achieve satisfactory anaesthesia. These infusion rates may be subsequently reduced depending on patient response and concurrent medication.

Intermittent Bolus

Increments of DIPRIVAN 25 mg (2.5 mL) to 50 mg (5.0 mL) may be administered with nitrous oxide in patients undergoing general surgery. The incremental boluses should be administered when changes in vital signs indicate a response to surgical stimulation or light anaesthesia.

DIPRIVAN has been used in conjunction with a wide variety of agents commonly used in anaesthesia such as atropine, scopolamine, glycopyrrolate, diazepam, depolarizing and nondepolarizing muscle relaxants, and narcotic analgesics, as well as with inhalational and regional anaesthetic agents. No pharmacological incompatibilities have been encountered.

Lower doses of DIPRIVAN may be required when used as an adjunct to regional anaesthesia.

Dosage Guide for Maintenance Of General Anaesthesia

Infusion: Variable rate infusion titrated to the desired clinical effect.

<i>Adult Patients < 55 Years of Age</i>	Generally, 0.10 to 0.20 mg/kg/min (6 to 12 mg/kg/h).
<i>Elderly, Debilitated and/or Adult ASA III or IV Patients</i>	Generally, 0.05 to 0.10 mg/kg/min (3 to 6 mg/kg/h).
<i>Cardiac Anaesthesia</i>	Most patients require: <ul style="list-style-type: none"> – Primary DIPRIVAN with Secondary Opioid - 0.10 to 0.15 mg/kg/min (6 to 9 mg/kg/h). – Low Dose DIPRIVAN with Primary Opioid - 0.05 to 0.10 mg/kg/min (3 to 6 mg/kg/h).
<i>Neurosurgical Patients</i>	Generally, 0.10 to 0.20 mg/kg/min (6 to 12 mg/kg/h).

<i>Pediatric Patients 3-18 Years of Age</i>	Generally, 0.10 to 0.25 mg/kg/min (6-15 mg/kg/h).
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Intermittent Bolus: Increments of 25 mg to 50 mg, as needed.

SURGICAL/DIAGNOSTIC SEDATION

Adults

When DIPRIVAN is administered for sedation, rates of administration should be individualized and titrated to clinical response. In most patients, the rates of DIPRIVAN administration will be approximately 25 to 30% of those used for maintenance of general anaesthesia.

During initiation of sedation, slow injection or slow infusion techniques are preferable over rapid bolus administration. During maintenance of sedation, a variable rate infusion is preferable over intermittent bolus dose administration.

Initiation of sedation

Slow injection: most adult patients will generally require 0.5 to 1.0 mg/kg administered over 3 to 5 minutes and titrated to clinical response.

In the elderly, debilitated, hypovolemic and ASA III or IV patients, the dosage of DIPRIVAN should be reduced to approximately 70 to 80% of the adult dosage and administered over 3 to 5 minutes.

Infusion: sedation may be initiated by infusing DIPRIVAN at 0.066 to 0.100 mg/kg/min (4.0 - 6.0 mg/kg/h) and titrating to the desired level of sedation while closely monitoring respiratory function.

Maintenance of sedation

Patients will generally require maintenance rates of 0.025 to 0.075 mg/kg/min (1.5 - 4.5 mg/kg/h) during the first 10 to 15 minutes of sedation maintenance.

Infusion rates should always be titrated downward in the absence of clinical signs of light sedation until mild responses to stimulation are obtained in order to avoid sedative administration of DIPRIVAN at rates higher than are clinically necessary.

In addition to the infusion, bolus administration of 10 to 15 mg may be necessary if a rapid increase in sedation depth is required.

In the elderly, debilitated, hypovolemic and ASA III or IV patients, the rate of administration and the dosage of DIPRIVAN should be reduced to approximately 70 to 80% of the adult

dosage according to their condition, responses, and changes in vital signs. Rapid (single or repeated) bolus dose administration should not be used for sedation in these patients (see WARNINGS AND PRECAUTIONS).

Dosage Guide for Surgical/Diagnostic Sedation

Dosage and rate should be individualized and titrated to the desired clinical effect.

<i>Adult Patients < 55 Years of Age</i>	Are likely to require 0.5 to 1.0 mg/kg over 3 to 5 min to initiate sedation, followed by 0.025 to 0.075 mg/kg/min (1.5 - 4.5 mg/kg/h) for continued sedation.
<i>Elderly, Debilitated, Hypovolemic and/or ASA III or IV patients</i>	The dosage and rate of administration may need to be reduced in these patients by approximately 20 to 30% (see previous section for details).
<i>Pediatric Patients <18 Years of Age</i>	DIPRIVAN is not recommended for sedation during surgical/diagnostic procedures in children under the age of 18, as safety and efficacy have not been established (see INDICATIONS and CLINICAL USE).

INITIATION AND MAINTENANCE OF ICU SEDATION IN INTUBATED, MECHANICALLY VENTILATED ADULT PATIENTS

DIPRIVAN should be individualized according to the patient's condition and response, blood lipid profile, and vital signs.

Adults

For intubated, mechanically ventilated, adult patients, Intensive Care Unit (ICU) sedation should be initiated slowly with a continuous infusion in order to titrate to desired clinical effect and minimize hypotension. When indicated, initiation of sedation should begin at 0.005 mg/kg/min (0.3 mg/kg/h). The infusion rate should be increased by increments of 0.005 to 0.010 mg/kg/min (0.3 - 0.6 mg/kg/h) until the desired level of sedation is achieved. A minimum period of 5 minutes between adjustments should be allowed for onset of peak drug effect.

Most adult patients require maintenance rates of 0.005 to 0.050 mg/kg/min (0.3 - 3.0 mg/kg/h). Administration of DIPRIVAN for ICU sedation in adult patients should not exceed 4 mg/kg/hour. Dosages of DIPRIVAN should be reduced in patients who have received large dosages of narcotics. As with other sedative medications, there is interpatient variability in dosage requirements and these requirements may change with time. (See DOSAGE AND

ADMINISTRATION, Initiation and Maintenance of ICU Sedation in Intubated, Mechanically Ventilated Adult Patients - Dosage Guide).

Bolus administration of 10 to 20 mg should only be used to rapidly increase sedation depth in patients where hypotension is not likely to occur. A rapid bolus should not be used, as this will increase the likelihood of hypotension. Patients with compromised myocardial function, intravascular volume depletion or abnormally low vascular tone (e.g. sepsis) may be more susceptible to hypotension.

Children Under 18 Years of Age

Propofol is contraindicated for the sedation of children 18 years or younger receiving intensive care.

Dosage Guide for Initiation and Maintenance of ICU Sedation in Intubated, Mechanically Ventilated Adult Patients

Dosage and rate of infusion should be individualized.

<i>Adult Patients</i>	<ul style="list-style-type: none">- For initiation, most patients require an infusion of 0.005 mg/kg/min (0.3 mg/kg/h) for at least 5 minutes. Subsequent increments of 0.005 to 0.010 mg/kg/min (0.3 - 0.6 mg/kg/h) over 5 to 10 minutes may be used until desired level of sedation is achieved. Administration of DIPRIVAN for ICU sedation in adult patients should not exceed 4 mg/kg/hour- For maintenance, most patients require 0.005 to 0.050 mg/kg/min (0.3 - 3.0 mg/kg/h).- The long-term administration of DIPRIVAN to patients with renal failure and/or hepatic insufficiency has not been evaluated.
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Handling and Administration

Handling Procedures

Parenteral drug products should be inspected visually for particulate matter and discoloration prior to administration whenever solution and container permit. Do not freeze.

DIPRIVAN should be visually inspected for particulate matter, emulsion separation and/or discoloration prior to use. Do not use if any of these changes are seen. If no signs of particulate matter, emulsion separation and/or discoloration are seen, shake gently before use.

Aseptic techniques must be applied to the handling of the drug. DIPRIVAN contains no antimicrobial preservatives and the vehicle supports growth of microorganisms. When DIPRIVAN is to be aspirated it should be drawn aseptically into a sterile syringe or intravenous administration set immediately after breaking the vial seal. Administration should commence without delay. Asepsis must be maintained for both DIPRIVAN and the infusion equipment throughout the infusion period. Any drugs or fluids added to the infusion line must be administered close to the cannula site. DIPRIVAN must not be administered via a microbiological filter.

DIPRIVAN is for single use in an individual patient only. If a vial is utilized for infusion, both the reservoir of DIPRIVAN and the infusion line must be discarded and replaced as appropriate at the end of the procedure or at 12 hours, whichever is sooner. (When using **DILUTED** DIPRIVAN see DOSAGE AND ADMINISTRATION, Reconstitution).

Administration into a Running I.V. Catheter

Compatibility of DIPRIVAN with the co-administration of blood/serum/plasma has not been established (see WARNINGS AND PRECAUTIONS, General). DIPRIVAN has been shown to be compatible with the following intravenous fluids when administered into a running i.v. catheter:

- 5% Dextrose Injection, USP
- Lactated Ringers Injection, USP
- Lactated Ringers and 5% Dextrose Injection
- 5% Dextrose and 0.45% Sodium Chloride Injection, USP
- 5% Dextrose and 0.2% Sodium Chloride Injection, USP

Since DIPRIVAN contains no preservative or bacteriostatic agents, any unused portions of DIPRIVAN or solutions containing DIPRIVAN should be discarded at the end of the surgical procedure.

Dilution Prior to Administration

When DIPRIVAN is diluted prior to administration, it should only be diluted with 5% Dextrose Injection, USP, and it should not be diluted to a concentration less than 2 mg/mL because it is an emulsion. Dilutions should be prepared aseptically immediately before administration and should not be used beyond 6 hours of preparation. In diluted form it has been shown to be more stable when in contact with glass than with plastic (95% potency after 2 hours of running infusion in plastic).

Pre-mixing with alfentanil

DIPRIVAN may be pre-mixed with alfentanil injection containing 500 µg/ml alfentanil in the ratio of 20:1 to 50:1 v/v. Mixtures should be prepared using sterile technique and used within 6 hours of preparation.

DIPRIVAN can be pre-mixed with alfentanil. DIPRIVAN should not be mixed with other therapeutic agents prior to administration.

The neuromuscular blocking agents, atracurium and mivacurium should not be given through the same i.v. line as DIPRIVAN without prior flushing.

OVERDOSAGE

For management of suspected drug overdose, contact your regional Poison Control Centre.

To date, there is no known case of acute overdose, and no specific information on emergency treatment of overdose is available. If accidental overdose occurs, DIPRIVAN (propofol) administration should be discontinued immediately. Overdosage is likely to cause cardiorespiratory depression. Respiratory depression should be treated by artificial ventilation with oxygen. Cardiovascular depression may require repositioning of the patient by raising the patient's legs, increasing the flow rate of intravenous fluids and if severe may require the administration of plasma volume expanders and/or pressor agents.

ACTION AND CLINICAL PHARMACOLOGY

Mechanism of Action

DIPRIVAN (propofol) is an intravenous hypnotic agent for use in the induction and maintenance of general anaesthesia or sedation.

The drug, an alkylphenol formulated in an oil-in-water emulsion, is chemically distinct from currently available intravenous anaesthetic agents. Intravenous injection of a therapeutic dose of propofol produces hypnosis rapidly and smoothly, usually within 40 seconds from the start of an injection (one arm-brain circulation time), although induction times >60 seconds have been observed.

Pharmacodynamics

Propofol induces anaesthesia in a dose-dependent manner. In unpremedicated, ASA I or II patients, propofol induced anaesthesia in 87% and 95% of patients at doses of 2.0 and 2.5 mg/kg, respectively. Elderly patients require lower doses; for unpremedicated patients older than 55 years of age, the mean dose requirement was 1.66 mg/kg. Premedication profoundly alters dose requirements; at 1.75 mg/kg, propofol induced anaesthesia in 65% of patients who had no premedication and in 85% and 100% of patients who received diazepam or papaveretum-hyoscine premedication, respectively.

During induction of anaesthesia, the hemodynamic effects of propofol vary. If spontaneous ventilation is maintained, the major cardiovascular effects are arterial hypotension (sometimes greater than a 30% decrease) with little or no change in heart rate and no appreciable decrease in cardiac output. If ventilation is assisted or controlled (positive pressure ventilation), the degree and incidence of decrease in cardiac output are accentuated. Maximal fall in blood pressure occurs within the first few minutes of the administration of a bolus dose. The fall in arterial pressure is greater under propofol anaesthesia than under anaesthesia induced by thiopental or methohexital. Increases in heart rate with propofol are generally less pronounced or absent after an induction dose, than after equivalent doses of these other two agents.

During maintenance of anaesthesia with propofol, systolic and diastolic blood pressures generally remain below pre-anaesthetic levels, although the depth of anaesthesia, the rate of maintenance infusion as well as stimulation from tracheal intubation and/or surgery may increase or decrease blood pressure. Heart rate may also vary as a function of these factors but will generally remain below pre-anaesthetic levels.

In the presence of a potent opioid (e.g. fentanyl), the blood pressure lowering effect of propofol is substantially increased. Fentanyl also decreases heart rate and this might lead to a significant decrease in cardiac output.

Age is highly correlated with the fall in blood pressure. In elderly subjects, both the incidence and degree of hypotension are greater than in younger subjects. Thus, a lower induction dose and a slower maintenance rate of administration should be used in the elderly (see DOSAGE AND ADMINISTRATION). Particular caution should be exercised in elderly patients with severe coronary and/or cerebral arteriosclerosis; reduction in perfusion pressure may impair adequate blood supply to these organs.

Insufficient data are available regarding the cardiovascular effects of propofol when used for induction and/or maintenance of anaesthesia or sedation in elderly, hypotensive, debilitated or other ASA III and IV patients. However, limited information suggests that these patients may have more profound cardiovascular responses. It is recommended that if propofol is used in these patients, a lower induction dose and a slower maintenance rate of administration of the drug be used (see WARNINGS AND PRECAUTIONS, General and DOSAGE AND ADMINISTRATION).

The first respiratory disturbance after a bolus dose of propofol is a profound fall in tidal volume leading to apnea in many patients. There has been no accompanying cough or hiccough and otherwise anaesthesia is smooth. However, there might be some difficulty in uptake of volatile agents if respiration is not assisted.

In unpremedicated, healthy patients, there is a steep dose-response relationship regarding apnea; 0% and 44% of patients had apnea after receiving 2.0 and 2.5 mg/kg of propofol, respectively. Fentanyl enhanced both the incidence and the onset of apnea and the episode lasted for >60 seconds in the majority of patients.

Opioid premedication - in the presence of hyoscine - affected respiratory function (rate of respiration and minute volume) substantially more than atropine premedication. Respiratory function was more depressed when these premedicants were combined with propofol than when they were combined with thiopental. Enhanced respiratory depression with propofol and an opioid have been observed in the postoperative period.

During maintenance, propofol (0.1 to 0.2 mg/kg/min; 6 - 12 mg/kg/h) caused a decrease in ventilation usually associated with an increase in carbon dioxide tension which may be marked depending upon the rate of administration and other concurrent medication (e.g. narcotics, sedatives, etc.). Propofol was not evaluated in patients with any respiratory dysfunction.

During sedation, attention must be given to the cardiorespiratory effects of DIPRIVAN. Hypotension, apnea, airway obstruction, and/or oxygen desaturation can occur, especially with a rapid bolus injection. During initiation of sedation, slow infusion or slow injection techniques are preferable over rapid bolus administration, and during maintenance of sedation, a variable rate infusion is preferable over intermittent bolus administration in order to minimize undesirable cardiorespiratory effects. In the elderly, debilitated and ASA III or IV patients, rapid (single or repeated) bolus dose administration should not be used for sedation (see WARNINGS AND PRECAUTIONS, General)

Clinical and preclinical studies suggest that propofol is rarely associated with elevation of plasma histamine levels and does not cause signs of histamine release.

Clinical and preclinical studies show that propofol does not suppress the adrenal response to ACTH.

Preliminary findings in patients with normal intraocular pressure indicate that propofol anaesthesia produces a decrease in intraocular pressure which may be associated with a concomitant decrease in systemic vascular resistance.

Propofol is devoid of analgesic or antanalgesic activity.

Pharmacokinetics

The pharmacokinetic profile of propofol can be described by a 3-compartment open model. After a single bolus dose, there is fast distribution from blood into tissues ($t_{1/2\alpha}$: 1.8 to 8.3 min), high metabolic clearance ($t_{1/2\beta}$: 34 to 66 min) and a terminal slow elimination from poorly perfused tissues ($t_{1/2\gamma}$: 184-480 min). With 12 and 24 hour samplings, $t_{1/2\gamma}$ values of 502 and 674 min, respectively, were observed.

When given by an infusion for up to two hours, the pharmacokinetics of propofol appear to be independent of dose (0.05 - 0.15 mg/kg/min; 3 - 9 mg/kg/h) and similar to i.v. bolus pharmacokinetics. Pharmacokinetics are linear over recommended infusion rates.

Propofol is highly protein-bound (97 - 99%); the degree of binding seems to be unrelated to either sex or age.

In the presence of DIPRIVAN, alfentanil concentrations were higher than expected based upon the rate of infusion. However, alfentanil did not affect the pharmacokinetics of DIPRIVAN (see DOSAGE AND ADMINISTRATION, Handling and Administration).

Pharmacokinetics in Adult Patients in Intensive Care Unit (ICU)

Regarding most parameters, the pharmacokinetics of propofol in these patients are similar to those of patients undergoing anaesthesia/sedation for short surgical procedures. However, the terminal half-life ($t_{1/2\gamma}$) is substantially prolonged after long-term infusion, reflecting extensive tissue distribution.

Distribution: Propofol has large volumes of distribution as would be expected with a highly lipophilic anaesthetic agent. The volume of central compartment (V_c) is between 21 and 56 L (0.35 - 0.93 L/kg based on a 60 kg patient), and the volume of distribution at steady state (V_{ss}) is between 171 and 364 L (2.85 - 6.07 L/kg). Values for volume of distribution during the terminal phase (V_d) are two to three times the corresponding V_{ss} values.

Metabolism: The termination of the anaesthetic or sedative effects of propofol after a single i.v. bolus or a maintenance infusion is due to extensive redistribution from the CNS to other tissues and high metabolic clearance, both of which will decrease blood concentrations. The mean propofol concentration at time of awakening is 1 $\mu\text{g/mL}$ (range : 0.74 to 2.2 $\mu\text{g/mL}$). Recovery from anaesthesia or sedation is rapid. When propofol is used for both induction (2.0 to 2.5 mg/kg) and maintenance (0.1 to 0.2 mg/kg/min) of anaesthesia, the majority of patients are generally awake, responsive to verbal command and oriented in approximately 7 to 8 minutes. Recovery from the effects of propofol occurs due to rapid metabolism and is not dependent on the terminal elimination half-life since the blood levels achieved in this phase are not clinically significant.

Excretion: A study in six subjects showed that 72% and 88% of the administered radio-labelled dose was recovered in the urine within 24 hours and 5 days, respectively. Less than 2% was excreted in the feces. Unchanged drug was less than 0.3%. Propofol is chiefly metabolized by conjugation in the liver to inactive metabolites which are excreted by the kidney. Propofol glucuronide accounts for about 50% of the administered dose. The remainder consists of the 1- and 4-glucuronide and 4-sulphate conjugates of 2,6-diisopropyl-1,4-quinol.

The total body clearance (Cl) of propofol ranges from 1.6 L/min to 2.3 L/min (0.026 - 0.038 L/min/kg based on a 60 kg patient). This clearance exceeds estimates of hepatic blood flow, suggesting possible extrahepatic metabolism.

Special Populations and Conditions

Pediatrics: The results were obtained in ASA I children, ranging in age from 3 to 10 years, who received a single bolus dose of propofol, 2.5 mg/kg. Propofol was rapidly distributed from blood into tissue ($t_{1/2\alpha}$: 1.5 - 4.1 min), metabolic clearance was high ($t_{1/2\beta}$: 9.3 - 56.1 min) and terminal elimination slow ($t_{1/2\gamma}$: 209 - 735 min). The volume of central compartment (V_c)

ranged between 0.53 - 0.72 L/kg, the volume of distribution at steady state (V_{ss}) was between 2.1 - 10.9 L/kg and clearance (Cl) ranged between 0.032 - 0.040 L/min/kg. The mean plasma concentration of propofol at awakening was 2.3 µg/mL.

Geriatrics: With increasing age, the dose of DIPRIVAN needed to achieve a defined anaesthetic endpoint (dose-requirement) decreases. Elderly patients had higher propofol blood concentrations at 2 minutes than young ones (6.07 versus 4.15 µg/mL), probably due to a significantly lower initial distribution volume (20 versus 26 L). The relatively high blood concentrations during the first few minutes can predispose elderly patients to cardiorespiratory effects including hypotension, apnea, airway obstruction and/or oxygen desaturation. The clearance of DIPRIVAN also decreased from a mean ± S.D. of 1.8 ± 0.4 L/min in young patients (18-35 years) to 1.4 ± 0.4 L/min in elderly patients (65-80 years). The reduced clearance could decrease maintenance propofol requirements and prolong recovery if inappropriate infusions are used. Obesity is associated with significantly larger volumes of distribution (399 L versus 153 L) and clearance rates (2.8 L/min versus 1.8 L/min) but there is no change in the elimination half-life.

Gender: The pharmacokinetics of DIPRIVAN do not appear to be altered by gender.

Hepatic Insufficiency: The pharmacokinetics of DIPRIVAN do not appear to be altered by chronic hepatic cirrhosis. The effects of acute hepatic failure on the pharmacokinetics of DIPRIVAN have not been studied.

Renal Insufficiency: In renal failure, the data is based on very limited findings. There was a trend towards longer half-lives, although the differences versus control patients did not reach statistical significance.

STORAGE AND STABILITY

Store between 2° and 25°C; do not freeze. The emulsion should be visually inspected for particulate matter, emulsion separation and/or discolouration prior to use. Do not use if any of these things are seen. If no signs of particulate matter, emulsion separation and/or discolouration are seen, shake gently before use. Any unused portions of DIPRIVAN (propofol) or solutions containing DIPRIVAN should be discarded at the end of the surgical procedure.

DOSAGE FORMS, COMPOSITION AND PACKAGING

Composition

DIPRIVAN (propofol) is a white, oil in water emulsion. Each mL contains 10 mg of propofol for i.v. administration. In addition to the active component, propofol, the formulation also contains soybean oil (100 mg/mL), glycerol (22.5 mg/mL), egg phosphatide (12 mg/mL) and disodium edetate (0.055 mg/mL)-and water for injection with sodium hydroxide to adjust pH. It is isotonic with a pH of 6.5-8.5.

Dosage Forms and Packaging

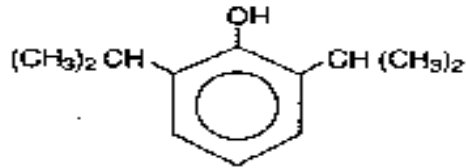
DIPRIVAN is available as DIPRIVAN 1% w/v in 20 mL, 50 mL and 100 mL glass vials for single infusion only. Each vial contains 10 mg/mL of propofol.

PART II: SCIENTIFIC INFORMATION

PHARMACEUTICAL INFORMATION

Drug Substance

Proper Name:	propofol
Chemical Name:	2,6-diisopropylphenol or 2,6-bis(1-methylethyl)phenol
Code Name:	
Molecular Formula and Molecular Mass:	C ₁₂ H ₁₈ O, 178.27
Structural Formula:	



Physicochemical Properties:

Colourless to pale straw-coloured liquid at room temperature. Practically insoluble in water. Completely miscible in all proportions with the following solvents at 20°C: acetone, 95% ethanol, chloroform, cyclohexane, diethyl ether, n-hexane, methanol, isooctane. pKa of 11.1 in water and a Melting Point of 18°C.

DETAILED PHARMACOLOGY

Propofol was administered as a 1% w/v aqueous emulsion containing 10% w/v soya bean oil, 1.2% w/v egg phosphatide and 2.25% w/v glycerol. Control animals received the vehicle. The drug was administered by the intravenous route unless otherwise indicated.

Anaesthetic Activity

Mice

The HD₅₀, i.e., the dose that abolished the righting reflex in 50% of the animals for 30 seconds or more, was 12.8 mg/kg. Induction times were significantly affected by the rate of injection. Namely, they were 4.6 and 12.6 seconds, respectively, after the injection of a 15 mg/kg dose over one or ten seconds. However, the rate of injection did not influence sleeping times or the duration of apnea to a significant degree. To evaluate the effect of repeated bolus injections, propofol was administered at a 25 mg/kg dose, which in previous experiments was shown to induce anaesthesia for approximately 4 minutes. When the mice regained their righting reflex, they were re-injected after an interval of 30 seconds. This procedure was repeated until the animals received a total of 10 injections. Sleeping times increased with subsequent injections.

The mean sleeping time was 4.6 minutes after the first injection and 19.4 minutes after the 8th injection indicating a slight cumulative effect.

Rats

In rats propofol produced a very steep dose-response curve, thus, only a range could be established for the HD₅₀ (5.0 to 7.5 mg/kg). In this species, the anaesthetic effect was sex dependent, female rats having significantly longer sleeping times than male rats. In a separate experiment, utilizing female rats, anaesthesia was induced by a 7.5-10 mg/kg bolus dose of propofol and then maintained by the infusion of a 50 mg/kg/h dose. Anaesthesia was maintained for either one or two hours. The experiment established that heart rate and rate of respiration remained stable over the two hour infusion and did not change vis à vis baseline. Recovery times were slightly longer after the two hour infusion but were significantly shorter in propofol-treated rats than in alphaxalone/alphadolone-treated animals.

Cardiovascular and Respiratory Effects

The experiments were conducted in mini-pigs. Propofol was evaluated at a 3.75 mg/kg dose injected over a 30 second interval.

All the changes mentioned were statistically significant. Maximal changes from baseline are indicated in brackets. Mean arterial pressure decreased (107 → 78 mmHg) and was accompanied by reflex tachycardia (105 → 189 beats/min). Rate of respiration became depressed (27.5 → 21.5 breaths/min) and was accompanied by decreased PO₂ (94.6 → 79.1 mmHg) and increased PCO₂ (40.4 → 44.3 mmHg). Total peripheral resistance decreased (2919 → 1902 dyne. sec. cm⁻⁵) while cardiac output increased (2.67 → 3.58 L/min). The hypotensive effect of propofol is probably due to reduced peripheral resistance.

Drug Interaction Studies

Preanaesthetic Medications

The experiments were conducted in mice. Diazepam, droperidol, promethazine, atropine, amylobarbitol or papaveretum were administered subcutaneously, 30 minutes before the induction of anaesthesia with propofol (18 mg/kg). At the doses tested, only diazepam (2 mg/kg) enhanced significantly the duration of sleep. Recovery times were prolonged significantly by both diazepam and atropine (1 mg/kg).

Drugs Used in Balanced Anaesthesia

The experiments were conducted in dogs. Anaesthesia was induced by propofol and maintained by halothane. In the first set of experiments, the dogs were premedicated with atropine and fentanyl, while in the second set of experiments premedication was not employed. The following changes were observed in the presence of atropine and fentanyl: the induction doses of propofol were smaller (4.3 versus 7.9 mg/kg) and the concentration of halothane lower (1% versus 2%). However, both tachycardia and apnea occurred. Recovery times were shorter, probably due to the lower dose of propofol.

The Effect of Propofol on Adrenocortical Function

The effect of propofol was compared to that of etomidate on ACTH-stimulated cortisol production in two in vitro models, namely in guinea pig and bovine adrenal cells. Propofol affected cortisol production only at the 10^{-4} M concentration, while etomidate exerted an activity at 10^{-7} M concentration.

In propofol-anaesthetized rats, 5 µg ACTH elicited serum corticosterone levels of 37.2 µg/100 mL while in etomidate-anaesthetized rats, the value was 8.5 µg/100 mL. Since basal corticosterone levels ranged between 5.6 and 10.4 µg/100 mL, ACTH did not elevate corticosterone levels in etomidate-anaesthetized rats. The experiments indicated that propofol is only a very weak inhibitor of adrenal steroidogenesis.

***In Vitro* Studies**

The aim of the experiments was to establish whether or not propofol exerts agonist or antagonist activity at various receptor sites. Only weak antagonist activities were detected at the β₁ adrenoceptor, muscarinic cholinergic and 5-HT₂ receptor sites. The pA₂ values were 5.23, 5.43 and 5.18, respectively. At the same receptor sites the pA₂ values for the standards were as follow: propranolol: 8.55, hyoscine: 9.38 and cyproheptadine: 8.2.

Behavioral Studies

The behaviour of mice was observed following the administration of 30, 100 and 300 mg/kg oral doses of propofol. None of the animals were anaesthetized by these oral doses. The lowest dose had no behavioral effects. The mid dose decreased locomotion. The highest dose produced sedation, ptosis, ataxia, slight tremors, hypothermia and decreased rate of respiration.

Miscellaneous Studies

Histamine Release

The administration of propofol, 7.5 mg/kg to dogs was not associated either with elevated plasma histamine levels or with clinical signs indicative of histamine release.

Hypersensitivity

Mini-pigs were anaesthetized with propofol, 2.5 mg/kg on two occasions at one week interval. No reactions, indicative of an anaphylactoid response were seen following the second injection.

Bronchomotor Tone

In the guinea-pig Konzett-Rossler technique, propofol (2.5 mg/kg) was devoid of both bronchoconstrictor and bronchodilator activity. The latter effect was tested against histamine-induced bronchoconstriction.

Blood Coagulation

ADP-induced platelet aggregation was similar in propofol (15 mg/kg) and saline-treated rats. Whole blood clotting times were similar in propofol (15 mg/kg) and saline-treated rats.

Renal Function

In rats, propofol, 15 mg/kg, had little effect on urine volume and urinary potassium and chloride levels. Sodium levels were slightly but significantly decreased (81% of control).

Cat Nictitating Membrane Preparation

Propofol, in a dose-range of 0.5 to 5.0 mg/kg, did not affect the contraction of the nictitating membrane, evoked by preganglionic stimulation of the cervical sympathetic nerve. The study indicates that propofol is devoid of ganglion blocking activity. Furthermore, propofol did not affect the pressor effect of norepinephrine, indicating a lack of effect on adrenoceptors.

Pharmacokinetics

Pharmacokinetic studies were carried out in male and female rats, male and female dogs and female rabbits. In all species, following a single intravenous dose, the pharmacokinetics fit a two-compartment open model with a very rapid distribution phase ($t_{1/2\alpha}$: 1.2-4.9 min) and a rapid elimination phase ($t_{1/2\beta}$: 15-27 min; [Table 2](#)). In rats, but not in dogs, a sex difference was observed regarding several pharmacokinetic parameters.

In rats, maximal mean propofol blood concentrations as well as AUC values and propofol blood concentrations at awakening were significantly higher in females. However, elimination half-lives were the same for the two sexes. Both propofol blood concentrations and AUC values increased in a dose-dependent manner. In contrast, waking blood concentrations were independent of the dose. In dogs, the pharmacokinetic parameters were determined either after a bolus injection of propofol (5 and 10 mg/kg) or in an infusion model where an initial bolus dose of 7.5 mg/kg was followed by an infusion at the rate of 0.5 mg/kg/min for 45 minutes (22.5 mg/kg). Steady-state blood concentrations (C_{ss}) in the infusion model were achieved within 25 minutes. The elimination half-life was significantly longer following the infusion than after the bolus doses ([Table 2](#)). In addition, total body clearance (TBCL) was significantly slower after the infusion (TBCL= 1.0 L/min) than after the 5 mg/kg (TBCL = 1.92 L/min) or 10 mg/kg (TBCL=2.12 L/min) bolus administrations. Waking propofol concentrations in the dog were ~ 1 µg/mL.

Table 2 Pharmacokinetic parameters following a single intravenous dose of propofol

Species	Dose (mg/kg)	Sex	Max propofol blood conc. (µg/mL)	AUC (µg.mL min)	$t_{1/2\beta}$ (min)	Sleeping time (min)	Waking propofol conc. (µg/mL)
Rat	5	M	0.57	13.7	23	Rats did not sleep	
		F	2.55	34.4	22		
		F (pregnant)	2.35	34.8	25		
	10	M	4.3	48.3	23	6.1	1.7
		F	11.3	87.9	18	7.9	2.8
	15	M	11.3	97.2	22	9.6	1.0
F		20.8	174.9	27	11.4	3.7	
Dog	5	M&F	2.35	40.3	16		
	10	M&F	4.31	71.4	21		
	30 (infusion)	M&F	C_{ss} : 6.5		33		
Rabbit	5	F	260	14.4	15		

Distribution

Tissue levels of total radioactivity and propofol were determined in rats following the administration of a 9.7 mg/kg intravenous dose of ¹⁴C-propofol. In all tissues assayed, other than fat, the highest concentration of radioactivity were detected five minutes after dosing and decreased thereafter. Maximal concentration in brown fat occurred at 10 minutes and in white fat at 30 and 60 minutes in males and females, respectively. This indicated that the distribution of propofol into fat occurs after five minutes.

Table 3 Concentrations of total radioactivity and propofol in selected tissues of rats 5 minutes after the intravenous administration of ¹⁴C-propofol

Tissue	Sex	Total radioactivity µg/equivalents/ mL	Propofol µg/mL of g	Propofol % of total radioactivity
Blood	M	5.18	1.47	28
	F	6.83	3.42	50
Brain	M	5.54	5.12	92
	F	9.87	9.16	93
Liver	M	32.77	1.58	5
	F	32.12	15.10	47

Tissue concentrations of total radioactivity were similar in male and female rats, except in the brain where radioactivity was significantly higher in the females (Table 3). The rate of decrease of radioactivity was greatest in the brain; by 30 minutes total radioactivity decreased to 19% and 15% of the 5 minute levels in males and females, respectively. The concentration of propofol in the blood, brain and liver was significantly higher in females. While propofol comprised >90% of the radioactivity in the brain of both sexes, in the blood and liver propofol concentrations were considerably lower and a sex difference was evident. In the liver, propofol levels were about ten times higher in female rats indicating initial differences in the rate of metabolism between the sexes.

Metabolism And Elimination

¹⁴C-propofol (10 mg/kg) was extensively metabolized and rapidly eliminated in the urine and feces of rats and dogs. In the rabbit, excretion occurred almost exclusively in the urine (Table 4).

Table 4 **The excretion of ¹⁴C-propofol**

Species	Sex	% dose			Total Recovery
		Urine	Feces	Bile	
Rat	M	60	31		92 ^a
	F	75	15		91 ^a
	F (pregnant)	77	16		95 ^a
	M ^b	13	1	78	95 ^c
	F ^b	15	1	53	82 ^c
Dog	M & F	60	29		90 ^d
Rabbit	M	95	2		93 ^e

a: includes dose found in ¹⁴CO₂ and carcass; 120-h collection

b: bile-duct cannulated rats

c: includes dose found in gastrointestinal tract and carcass; 24-h collection

d: 48-h collection

e: 24-h collection

In rats, the differences between the excretion data for the two sexes were statistically significant. Extensive biliary excretion and enterohepatic recirculation was observed in both sexes. In the urine, propofol was completely metabolized prior to elimination. In the feces, propofol comprised ~ 10% and 6% of the dose in male and female rats, respectively. The presence of propofol in the feces may be due to hydrolysis of propofol glucuronide. The radioactivity in the urine consisted of the 4-glucuronide and 4-sulphate conjugates of 2,6-diisopropyl 1,4 quinol and 4-sulphate of 2-(1-propionic acid)-6-isopropyl 1,4 quinol.

In dogs, the urine contained the 4-substituted glucuronic acid and sulphate conjugates of 2,6-diisopropyl 1,4-quinol and minor metabolites; unchanged propofol was <1%. The feces contained 2,6-diisopropyl 1,4-quinol and some uncharacterized polar metabolites. At 2 minutes, the blood concentration of radioactivity was 10.02 µg equivalents/mL, that of propofol was 2.7 µg/mL, constituting 26% of total radioactivity. At 2 hours, propofol comprised only ~ 1% of radioactivity.

In rabbits, urinary radioactivity consisted of the 4-glucuronide and 4-sulphate conjugates of propofol and 2,6-diisopropyl 1,4-quinol. Unchanged propofol was not detected. At 2 minutes, the blood concentration of radioactivity was ~ 30 µg equivalents/mL, that of propofol was 15.9 µg/mL, constituting 53% of total radioactivity. At 2 hours, propofol represented ~ 2% radioactivity.

Binding To Plasma Proteins

Propofol was 98% and 97% bound to plasma proteins in the dog and rat, respectively, over a concentration range of 0.1-20 µg/mL. In the rabbit, binding was concentration dependent; propofol binding decreased from 97% at 0.5 µg/mL to 95% at 50 µg/mL.

TOXICOLOGY

Acute Toxicity

Studies in Rats and Mice

Rats and mice of the Alderley Park Albino strain received graded intravenous or oral doses of propofol. At each dose level, six male and six female animals were used. The drug was available as an emulsion for the i.v. studies and as a solution in soya bean oil for the oral studies. At the doses used, all animals became anaesthetized. Several rats and mice, both in the i.v. and oral studies, regained consciousness and then became re-anaesthetized before fully recovering. The LD₅₀ values and observations are summarized in Table 5 below.

Table 5 LD₅₀ values and observations in Rats and Mice

Species	Route of admin	LD ₅₀ mg/kg (95% confidence limits)	Observations
Rats	i.v.	42 (38-46)	Death occurred within 5 minutes of dosing.
	oral	600 (540-660)	The majority of rats died 1 to 3 days after propofol administration. Following recovery from anaesthesia, several rats exhibited decreased activity, piloerection, hunched posture and tremors.
Mice	i.v.	53 (46-60)	Death occurred within 2 minutes of dosing and was due to respiratory depression.
	oral	1230 (1010 –1500)	The majority of mice died 1 to 2 days after propofol administration. During anaesthesia, both the rate and depth of respiration was decreased. Following recovery from anaesthesia, several mice exhibited locomotor incoordination and tremors.

Single-Dose Tolerance Study in Rabbits

Three male and three female Dutch rabbits received propofol, 15 mg/kg, by the intravenous route. The drug was given at a rate of 0.5 mg/kg/second. All rabbits became lightly

anaesthetized, with 6/6 rabbits retaining their pedal reflex and 2/6 rabbits retaining their palpebral reflex. Ten to 15 minutes after dosing, all rabbits recovered completely without any untoward effect.

Long-Term Toxicology

One-Month Toxicity Study in Rats

Five groups of albino rats were dosed daily for 28 days. Injections were given intravenously into the tail vein. Group I received saline, Group II the emulsion vehicle, Groups III, IV and V propofol at doses of 5, 10 and 15 mg/kg/day, respectively.

Propofol induced anaesthesia in a dose-dependent manner; at 5 mg/kg rats were not anaesthetized while the duration of anaesthesia was significantly longer at the 15 mg/kg than at the 10 mg/kg dose. With repeated administration, the duration of anaesthesia became prolonged and on Day 26, anaesthesia lasted significantly longer than on Day 1.

High dose male rats gained slightly but significantly less weight than control rats (131 versus 150 g). In female rats, weight gain was slightly less in all treated animals, however, the effect was not dose-related. Urine volume was significantly but not dose-dependently elevated on Day 26 in all propofol-treated rats. In female rats, relative kidney weights were significantly and dose-dependently elevated in all propofol-treated groups.

One-Month Toxicity Study in Dogs

Five groups of Beagle dogs were dosed intravenously over a 30-day period. Group I received saline, Group II the emulsion vehicle, and Groups III and IV propofol at doses of 5 and 10 mg/kg/day, respectively. Group V received propofol, 30 mg/kg 3 times weekly for a total of 13 doses.

Each dose consisted of a 7.5 mg/kg bolus dose and an infusion of 0.5 mg/kg/min for a total of 22.5 mg/kg.

Each group was comprised of 5 male and 5 female dogs. In addition, 3 dogs/sex were used to evaluate recovery in the control and high dose groups.

Propofol induced anaesthesia in a dose-dependent manner. With repeated administration, the duration of anaesthesia became prolonged and on Day 28, anaesthesia lasted significantly longer than on Day 1.

During the 30-day treatment period, Hb, RBC and PCV values declined below the normal range in a few animals. On Day 30, abnormally low values were recorded in 3/10 dogs in both Groups III and IV. (In both groups, the same three dogs were affected.) In Groups II and V, 1/16 dogs each showed similar changes.

REPRODUCTION AND TERATOLOGY

Fertility and Reproductive Performance in Rats

Three groups of 50 rats each were dosed intravenously with the vehicle or propofol at doses of 10 or 15 mg/kg/day for two weeks prior to mating, during the mating period to untreated males and up to Day 7 of gestation. Generally, reproductive studies require that treatment be continued during both gestation and lactation, thus, this study provides information about propofol's effect upon fertility but not necessarily upon reproduction.

Approximately half of the females of the F₀ generation were sacrificed on Day 21 of pregnancy. The remainder were allowed to litter and rear their offspring to weaning at Day 22 of lactation. At weaning, two females and one male were selected from each litter to form the F₁ generation. These animals were kept until sexually mature and then mated. As with the F₀ generation, approximately half the females were sacrificed on Day 21 of pregnancy and the remainder were allowed to litter and rear their young to weaning, when the F₁ dams and their pups (the F₂ generation) were sacrificed.

The administration of propofol was associated with the following changes:

In the F₀ generation, treated rats gained significantly less weight than controls prior to mating (9.7, -0.8 and 1.7 g in the control, low and high dose groups, respectively). However, weight gains between Days 7 and 16, or 1 to 21 of pregnancy, were similar in all three groups.

Gestation period was dose-dependently decreased. In the control, low and high dose groups 9.5, 16 and 33% of the rats, respectively delivered on Day 21, rather than Day 22.

Survival of the F₁ generation pups was lower in the treated groups. On Day 1, the number of alive pups was similar in all three groups. From Day 5 on, survival in treated groups was lower. Numerical values on Day 22 were as follows: 73, 49 and 52% of pups were alive in the control, low and high dose groups, respectively.

Pups which died, were subjected to necropsy. None showed soft tissue abnormalities, however, reduced vertebral ossification was present in 13, 38 and 40% of pups in the control, low and high dose groups, respectively.

Postimplantation loss (as a % of implants) in the F₁ generation was higher in rats born to high dose animals (2.3, 1.2 and 15.6% in control, low and high dose rats, respectively).

Teratology Study in Rats

Four groups of 40 mated female rats each were dosed intravenously with the vehicle or propofol at doses of 5, 10 or 15 mg/kg/day from Day 6 to Day 15 of pregnancy. The rats were sacrificed on Day 20 of pregnancy and the pups checked for internal and skeletal anomalies.

Maternal weight gain during Days 6 to 15 was significantly less in propofol-treated rats than in controls. The incidence of abnormal cranial ossification was higher in fetuses born to high dose dams than in control fetuses (19.9% versus 11.0%).

In rats, sacrificed on Day 15 of pregnancy, 10 minutes after the last dose, propofol was detected in maternal blood, amniotic fluid and the developing embryo. Drug concentrations increased linearly with increasing doses.

The study indicated that propofol is not teratogenic in rats at the doses studied.

Teratology Study in Rabbits

Four groups of 22 mated female rabbits each were dosed intravenously with the vehicle or propofol at doses of 5, 10 or 15 mg/kg/day from Day 6 to Day 18 of pregnancy. The rabbits were sacrificed on Day 28 of pregnancy.

Maternal weight gain during Days 6 to 18 was less in propofol-treated rabbits than in controls. Incomplete sternebral ossification increased dose-dependently in fetuses born to propofol-treated dams as compared to control fetuses.

Propofol was detected in maternal blood, amniotic fluid and embryonic tissue. Drug concentrations increased in a dose-dependent manner.

The study indicated that propofol is not teratogenic in rabbits at the doses studied.

Perinatal and Postnatal Study in Rats

Three groups of 22 rats each were dosed intravenously with the vehicle or propofol at doses of 10 to 15 mg/kg/day from Day 16 of gestation through Day 22 of lactation. The number of rats in whom treatment was completed was 18, 16 and 12, in the control, low and high dose groups, respectively. In the high dose group, four dams died during dosing, the cause of death might have been due to respiratory depression. In addition, mothers were sacrificed if the litters died. Maternal weight gain, during the last week of pregnancy, was significantly less in high dose rats than in control animals (47.1 versus 60.3 g). Litter survival on Day 22 was slightly but dose-dependently decreased; the percent of litters which survived was 65, 61 and 53% in the control, low and high dose groups, respectively.

Propofol did not affect the gestation period, maternal weight gain during lactation or the weight gain and developmental landmarks of the litter.

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PART III: CONSUMER INFORMATION

[Fr] DIPRIVAN[®] 1% w/v Propofol injection 10 mg/mL

This leaflet is part III of a three-part “Product Monograph” published when DIPRIVAN was approved for sale in Canada and is designed specifically for Consumers. This leaflet is a summary and will not tell you everything about DIPRIVAN. Contact your doctor or pharmacist if you have any questions about the drug.

ABOUT THIS MEDICATION

WHAT THE MEDICATION IS USED FOR:

DIPRIVAN (propofol) belongs to a group of medicines called general anaesthetics. It is used to induce or maintain anaesthesia.

WHAT IT DOES:

DIPRIVAN causes you to become unconscious (asleep) while surgical operations or other procedures are being conducted. In adults only, smaller doses can also be used in certain circumstances to sedate you (make you sleepy or make you go to sleep lightly).

The effect of DIPRIVAN starts rapidly, within approximately 30 seconds. The recovery period from anaesthesia or sedation is also usually rapid.

WHEN IT SHOULD NOT BE USED:

- In children under the age of 18 years for sedation during surgical/diagnostic procedures or in the intensive care unit.
- If you have ever received DIPRIVAN before and have experienced an allergic reaction to its use or if you know that you are allergic to any of the ingredients listed in this leaflet (see WHAT THE NONMEDICINAL INGREDIENTS ARE)

WHAT THE MEDICINAL INGREDIENT IS:

Propofol 10 mg/mL.

WHAT THE NONMEDICINAL INGREDIENTS ARE:

DIPRIVAN contains disodium edetate, egg phosphatide, glycerol, sodium hydroxide, soybean oil and water for injection.

WHAT DOSAGE FORMS IT COMES IN:

DIPRIVAN is available as DIPRIVAN 1% w/v in 20 mL, 50 mL and 100 mL glass vials for single infusion only.

WARNINGS AND PRECAUTIONS

You should talk to your doctor or anesthetist prior to surgery:

- About all health problems you have now or have had in the past.
- About other medicines you take, including ones you can buy without a prescription.
- If you have any other health problems such as problems with your heart, breathing, kidneys or liver or if you have been generally unwell for some time.
- If you have ever had an epileptic fit or convulsion.
- If you have ever been told that either you have very high fat levels in your blood or your body has problems in metabolising fat adequately.
- If you are pregnant, plan to become pregnant or are breastfeeding.
- If you are planning to drive or operate any tools or machinery on the day of surgery, because DIPRIVAN may temporarily interfere with your reactions and muscular coordination.

INTERACTIONS WITH THIS MEDICATION

Some drugs interact with DIPRIVAN. Tell your doctor about all prescription, over-the-counter and natural health products that you are using (see WARNINGS AND PRECAUTIONS above).

Usage of such medicines at the same time as DIPRIVAN may increase the risk of serious side effects.

PROPER USE OF THIS MEDICATION

HOW DIPRIVAN IS USED:

DIPRIVAN will be given to you by your anaesthetist or intensive care doctor.

Usual Dose

The dose given is decided by the doctor based on the clinical need and your physical condition.

When Receiving DIPRIVAN

DIPRIVAN will be given to you as an injection into a vein, usually in the back of the hand or in the forearm. Your anaesthetist may use a needle, or a fine plastic tube called a cannula. For long operations and for use in intensive care situations, an electric pump may be used to control the rate at which the injection is given.

After Receiving DIPRIVAN

You may feel some pain in the arm into which DIPRIVAN is given; this is normal. Your anaesthetist or intensive care doctor will closely control the amount of DIPRIVAN that is given to you. The amount will be adjusted according to how deeply your anaesthetist or intensive care doctor wishes you to be sedated or anaesthetised.

He/she will also take account of your age and physical fitness and adjust the amount accordingly.

Several different medicines may be needed in order to keep you asleep or sedated, free from pain, breathing in a healthy way and to keep your blood pressure steady. Your anaesthetist or intensive care doctor will decide which medicines to use as the need arises.

Serious adverse effects resulting from an overdose are extremely rare and need special treatment. The doctor is trained and equipped to handle such situations.

SIDE EFFECTS AND WHAT TO DO ABOUT THEM

Like all medicines, DIPRIVAN can have side effects. **Your anaesthetist or intensive care doctor will care for you if any of the following side effects or any other undesirable events occurs.**

Medicines affect different people in different ways. Just because side effects have occurred in some patients, does not mean that you will get them.

SERIOUS SIDE EFFECTS, HOW OFTEN THEY HAPPEN AND WHAT TO DO ABOUT THEM

DURING ANAESTHESIA

More than 10 of every 100 patients (very common) may experience:

- A feeling of pain near the injection site before the injection makes you unconscious

1 to 10 of every 100 patients (common) may experience:

- A fall in blood pressure
- A slower heart beat
- Changes in your breathing pattern

Less than 1 of every 1000 patients (rare) may experience:

- Some twitching and shaking

The majority of the above reactions are mild and transient in nature.

Less than 1 of every 10 000 patients (very rare) may experience:

- Allergic reactions
- Fluid in the lungs

AFTER ANAESTHESIA

1 to 10 of every 100 patients (common) may experience:

- Nausea and vomiting
- Headache

Less than 1 of every 100 patients (uncommon) may experience:

- Redness or soreness where the anaesthetic was given

Less than 1 of every 1000 patients (rare) may experience:

- Some twitching and shaking

Less than 1 of every 10 000 patients (very rare) may experience:

- Inflammation of the pancreas
- Fluid in the lungs
- Increased body temperature
- A feeling of sexual arousal
- Postoperative unconsciousness – recovery without complications has always occurred

DURING INTENSIVE CARE SEDATION (ADULTS ONLY)

1 to 10 of every 100 patients (common) may experience:

- A fall in blood pressure
- A slower heart beat

Less than 1 of every 100 patients (uncommon) may experience:

- Redness or soreness where the anaesthetic was given

Less than 1 of every 1000 patients (rare) may experience:

- Some twitching and shaking

Less than 1 of every 10 000 patients (very rare) may experience:

- Allergic reactions
- Increased body temperature
- Breakdown of muscle cells (rhabdomyolysis) when DIPRIVAN has been given in excess of the maximum recommended dose rate. A causal relationship with DIPRIVAN has not been established.
- Inflammation of the pancreas
- Discolouration of the urine

This is not a complete list of side effects. For any unexpected effects while or after taking DIPRIVAN contact your doctor or pharmacist.

REPORTING SUSPECTED SIDE EFFECTS

You can report any suspected adverse reactions associated with the use of health products to the Canada Vigilance Program by one of the following 3 ways:

- Report online at www.healthcanada.gc.ca/medeffect
- Call toll-free at 1-866-234-2345
- Complete a Canada Vigilance Reporting Form and:
 - Fax toll-free to 1-866-678-6789, or
 - Mail to: Canada Vigilance Program
Health Canada
Postal Locator 0701E
Ottawa, ON K1A 0K9

Postage paid labels, Canada Vigilance Reporting Form and the adverse reaction reporting guidelines are available on the MedEffect™ Canada Web site at www.healthcanada.gc.ca/medeffect.

NOTE: Should you require information related to the management of side effects, contact your health professional. The Canada Vigilance Program does not provide medical advice.

MORE INFORMATION

NOTE: This INFORMATION FOR THE CONSUMER leaflet provides you with the most current information at the time of printing.

The Consumer Information Leaflet plus the full Product Monograph, prepared for health professionals can be found at:

www.astrazeneca.ca,

or by contacting the sponsor, AstraZeneca Canada Inc. at:
Customer Inquiries – 1 (800) 668-6000,
Renseignements – 1 (800) 461-3787.

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