APOSTILLE

(Convention de La Haye du 5 octobre 1961)

1. Country: *United States of America*

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2. has been signed by Andrei Perlloni

3. acting in the capacity of Branch Chief, Drug Import Export Compliance Branch

4. bears the seal/stamp of U. S. Department of Health and Human Services

Certified

5. at Washington, D.C.

6. the twenty-fourth of November, 2017

7. by *Assistant Authentication Officer, United States Department of State*

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Matthews

Veda Matthews

United States Food and Drug Administration

Center for Drug Evaluation and Research

10903 New Hampshire Ave, Silver Spring, MD 20993, United States of America

CDERExportCertificateProgram@fda.hhs.gov - Telephone (301) 796-4950

Certificate of a Pharmaceutical Product - Approved Drug Product

Certificate Number: **3MPP-HK85**

Certificate Issue Date: **November 13, 2017**

Certificate Expiration Date: **November 12, 2019**

Importing Country: **CHILE**

Exporting Country: **UNITED STATES of AMERICA**

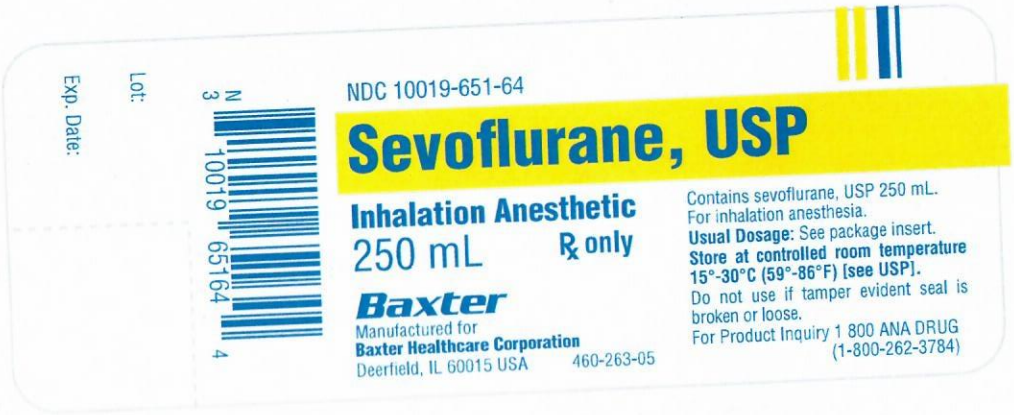
1.	Drug Trade Name, International or National non-proprietary name (as applicable) & dosage form: **SEVOFLURANE, USP, Inhalant**
1.1	Active Ingredient(s) and amount(s) per unit dose (complete quantitative composition is preferred): **sevoflurane, usp 250 ML**
1.2	Is this product licensed to be placed on the market for use in the exporting country? **Yes**
1.3	Is this product actually on the market in the exporting country? **Yes**
2.A.1	Product license number & date of issue: **075895 07/02/2002**
2.A.2	Product license holder name & address: **Baxter Healthcare Corporation, 1 Baxter Parkway, Deerfield, IL 60015 United States of America**
2.A.3	Status of Product license holder: **Manufacturer**
2.A.3.1	Manufacturer name & address: **Baxter Healthcare Corporation, Route 3 Km 144.2, Guayama, PUERTO RICO 00784**
2.A.4	Is a summary basis for approval appended? **Yes**
2.A.5	Is the attached product information, complete and consonant with the license? **Yes**
2.A.6	Applicant name & address for certificate (if different from the license holder): **Baxter Healthcare Corporation, 32650 N Wilson Rd, Round Lake, IL 60073 United States of America**
2.B.4	Remarks: **Manufacturing Facility: Baxter Healthcare Corporation,Route 3 Km 144.2, Guayama, Puerto Rico 00784(USA). Marketing Authorization Holder:Baxter Healthcare Corporation, 1 Baxter Parkway Deerfield IL 60015 USA**
3.	Does the certifying authority arrange for periodic inspection of the manufacturing plant in which the dosage form is produced? **Yes**
3.1	Periodicity of routine inspections (years): **Pursuant to section 510(h)(3) of the Federal Food, Drug & Cosmetic Act, Inspections will occur in accordance with a risk-based schedule**
3.2	Has the manufacture of this type of dosage form been inspected? **Yes**
3.3	Do the facilities and operations conform to GMPs as recommended by the WHO? (GMPs including 21 Code of Federal Regulations parts 210, 211, or ICH Q7A): **Yes, at time of inspection, site complies with FDA cGMP**
3.4	Does the information submitted by the applicant satisfy the certifying authority on all aspects of the manufacture of the product undertaken by another party? **Yes**

Andrei Perlloni

Andrei Perlloni, Branch Chief
Drug Import Export Compliance Branch
Division of Imports, Exports & Recalls
Office of Drug Security, Integrity & Respons

conforms to the format recommended by the World Health Organization format revised October 1, 1997. Website: www.who.int

3MPP-HK85



460-263-05 Label, Sevoflurane, USP, 250 mL Bottle
Size: 2" x 5 3/8" Corner: 1/4"
USA
Submission - 1 4-May-2011

PMS 108

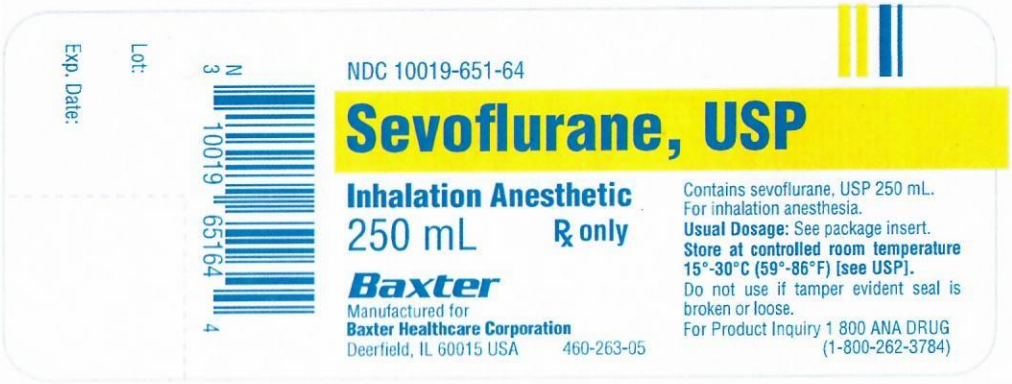
PMS 287

Bar Code: UPC Code

Multi-Color Pharmacode (5): Prints in PMS 108 and 287.

Guide Lines: DO NOT PRINT.

Dotted Lines: Indicates Unvarnished Imprint Area. DO NOT PRINT.



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3MPP-HK85

071975755

Sevoflurane, USP

Volatile Liquid for Inhalation

DESCRIPTION

Sevoflurane, USP volatile liquid for inhalation, a nonflammable and nonexplosive liquid administered by vaporization, is a halogenated general inhalation anesthetic drug. Sevoflurane, USP is fluoromethyl 2,2,2-trifluoro-1-(trifluoromethyl) ethyl ether and its structural formula is:

Sevoflurane, USP, Physical Constants are:

Molecular weight	200.05
Boiling point at 760 mm Hg	58.6°C
Specific gravity at 20°C	1.520 - 1.525
Vapor pressure in mm Hg	157 mm Hg at 20°C
	197 mm Hg at 25°C
	317 mm Hg at 36°C

Distribution Partition Coefficients at 37°C:

Blood/Gas	0.63 - 0.69
Water/Gas	0.36
Olive Oil/Gas	47 - 54
Brain/Gas	1.15

Mean Component/Gas Partition Coefficients at 25°C for Polymers Used Commonly in Medical Applications:

Conductive rubber	14.0
Butyl rubber	7.7
Polyvinylchloride	17.4
Polyethylene	1.3

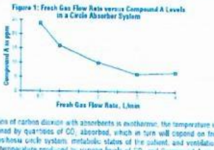
Sevoflurane, USP is nonflammable and nonexplosive as defined by the requirements of International Electrotechnical Commission 601-2-13.

Sevoflurane, USP is a clear, colorless, liquid containing no additives. Sevoflurane, USP is not corrosive to stainless steel, brass, aluminum, nickel-plated brass, chrome-plated brass or copper beryllium. Sevoflurane, USP is nonpungent. It is miscible with ethanol, ether, chloroform, and benzene, and it is slightly soluble in water. Sevoflurane, USP is stable when stored under normal room lighting conditions according to instructions. No discernible degradation of sevoflurane, USP occurs in the presence of strong acids or heat. When in contact with alkaline CO_2 absorbents (e.g. Baralyme® and to a lesser extent soda lime) within the anesthesia machine, Sevoflurane, USP can undergo degradation under certain conditions. Degradation of sevoflurane, USP is minimal, and degradants are either undetectable or present in non-toxic amounts when used as directed with fresh absorbents. Sevoflurane, USP degradation and subsequent degradant formation are enhanced by increasing absorbent temperature, increased sevoflurane, USP concentration, decreased fresh gas flow and desiccated CO_2 absorbents (especially with potassium hydroxide containing absorbents e.g. Baralyme).

Sevoflurane, USP alkaline degradation occurs by two pathways. The first results from the loss of hydrogen fluoride with the formation of pentafluoroisopropenyl fluoromethyl ether, (PIFE, $C_4H_2F_6O$), also known as Compound A, and trace amounts of pentafluoromethoxy isopropyl fluoromethyl ether, (PMFE, $C_5H_6F_6O$), also known as Compound B. The second pathway for degradation of sevoflurane, USP, which occurs primarily in the presence of desiccated CO_2 absorbents, is discussed later.

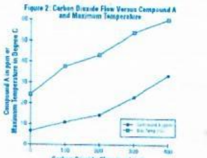
In the first pathway, the defluorination pathway, the production of degradants in the anesthesia circuit results from the extraction of the acidic proton in the presence of a strong base (KOH and/or NaOH) forming an alkene (Compound A) from sevoflurane, USP similar to formation of 2-bromo-2-chloro-1,1-difluoro ethylene (BCDFE) from halothane. Laboratory simulations have shown that the concentration of these degradants is inversely correlated with the fresh gas flow rate (See Figure 1).

Figure 1: Fresh Gas Flow Rate versus Compound A Levels in a Circle Absorber System



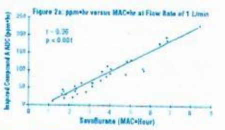
Since the reaction of carbon dioxide with absorbents is exothermic, the temperature increase will be determined by quantities of CO_2 absorbed, which in turn will depend on fresh gas flow in the anesthesia circle system, metabolic status of the patient, and ventilation. The relationship of temperature produced by varying levels of CO_2 and Compound A production is illustrated in the following in vitro simulation where CO_2 was added to a circle absorber system.

Figure 2: Carbon Dioxide Flow Versus Compound A and Maximum Temperature



Compound A concentration in a circle absorber system increases as a function of increasing CO_2 absorbent temperature and composition (Baralyme producing higher levels than soda lime), increased body temperature, and increased minute ventilation, and decreasing fresh gas flow rates. It has been reported that the concentration of Compound A increases significantly with prolonged dehydration of Baralyme. Compound A exposure in patients also has been shown to rise with increased sevoflurane, USP concentrations and duration of anesthesia. In a clinical study in which sevoflurane, USP was administered to patients under low flow conditions for ≥2 hours at flow rates of 1 Liter/minute, Compound A levels were measured in an effort to determine the relationship between MAC hours and Compound A levels produced. The relationship between Compound A levels and sevoflurane, USP exposure are shown in Figure 2a.

Figure 2a: ppm•hr versus MAC•hr at Flow Rate of 1 L/min



Compound A has been shown to be nephrotoxic in rats after exposures that have varied in duration from one to three hours. No histopathologic change was seen at a concentration of up to 270 ppm for one hour. Sporadic single cell necrosis of proximal tubule cells has been reported at a concentration of 114 ppm after a 3-hour exposure to Compound A in rats. The LC_{50} reported at 1 hour is 1050-1090 ppm (male-female) and, at 3 hours, 350-490 ppm (male-female).

An experiment was performed comparing sevoflurane, USP plus 75 or 100 ppm Compound A with an active control to evaluate the potential nephrotoxicity of Compound A in non-human primates. A single 8-hour exposure of Sevoflurane, USP in the presence of Compound A produced single-cell renal tubular degeneration and single-cell necrosis in cynomolgus monkeys. These changes are consistent with the increased urinary protein, glucose level and enzymic activity noted on days one and three on the clinical pathology evaluation. This nephrotoxicity produced by Compound A is dose and duration of exposure dependent.

At a fresh gas flow rate of 1 L/min, mean maximum concentrations of Compound A in the anesthesia circuit in clinical settings are approximately 20 ppm (0.002%) with soda lime and 30 ppm (0.003%) with Baralyme in adult patients; mean maximum concentrations in pediatric patients with soda lime are about half those found in adults. The highest concentration observed in a single patient with Baralyme was 61 ppm (0.0061%) and 32 ppm (0.0032%) with soda lime. The levels of Compound A at which toxicity occurs in humans is not known.

The second pathway for degradation of sevoflurane, USP occurs primarily in the presence of desiccated CO_2 absorbents and leads to the dissociation of sevoflurane, USP into hexafluoroisopropanol (HFIP) and formaldehyde. HFIP is inactive, non-genotoxic, rapidly glucuronidated and cleared by the liver. Formaldehyde is present during normal metabolic processes. Upon exposure to a highly desiccated absorbent, formaldehyde can further degrade into methanol and formate. Formate can contribute to the formation of carbon monoxide in the presence of high temperature that can be associated with desiccated Baralyme®. Methanol can react with Compound A to form the methoxy addition product Compound B. Compound B can undergo further HF elimination to form Compounds C, D and E.

Sevoflurane, USP degradants were observed in the respiratory circuit of an experimental anesthesia machine using desiccated CO_2 absorbents and maximum sevoflurane, USP concentrations (8%) for extended periods of time (≥2 hours). Concentrations of formaldehyde observed with desiccated soda lime in this experimental anesthesia respiratory circuit were consistent with levels that could potentially result in respiratory irritation. Although KOH containing CO_2 absorbents are no longer commercially available, in the laboratory experiments, exposure of sevoflurane, USP to the desiccated KOH containing CO_2 absorbent, Baralyme, resulted in the detection of substantially greater degradant levels.

CLINICAL PHARMACOLOGY

Sevoflurane, USP is an inhalational anesthetic agent for use in induction and maintenance of general anesthesia. Minimum alveolar concentration (MAC) of sevoflurane, USP in oxygen for a 40-year-old adult is 2.1%. The MAC of sevoflurane, USP decreases with age (see **DOSAGE AND ADMINISTRATION** for details).

Pharmacokinetics

UPTAKE AND DISTRIBUTION

Solubility

Because of the low solubility of sevoflurane, USP in blood (blood/gas partition coefficient @ 37°C = 0.63-0.69), a minimal amount of sevoflurane, USP is required to be dissolved in the blood before the alveolar partial pressure is in equilibrium with the arterial partial pressure. Therefore there is a rapid rate of increase in the alveolar (end-tidal) concentration (F_A) toward the inspired concentration (F_I) during induction.

Induction of Anesthesia

In a study in which seven healthy male volunteers were administered 70% N_2O/30% O_2 for 30 minutes followed by 1.0% sevoflurane, USP and 0.6% isoflurane for another 30 minutes the F_A/F_I ratio was greater for sevoflurane, USP than isoflurane at all time points. The time for the concentration in the alveoli to reach 50% of the inspired concentration was 4-8 minutes for isoflurane and approximately 1 minute for sevoflurane, USP.

F_A/F_I data from this study were compared with F_A/F_I data of other halogenated anesthetic agents from another study. When all data were normalized to isoflurane, the uptake and distribution of sevoflurane, USP was shown to be faster than isoflurane and halothane, but slower than desflurane. The results are depicted in Figure 3.

Recovery from Anesthesia

The low solubility of sevoflurane, USP facilitates rapid elimination via the lungs. The rate of elimination is quantified as the rate of change of the alveolar (end-tidal) concentration following termination of anesthesia (F_A), relative to the last alveolar concentration (F_{AO}) measured immediately before discontinuance of the anesthetic. In the healthy volunteer study described above, rate of elimination of sevoflurane, USP was similar compared with desflurane, but faster compared with either halothane or isoflurane. These results are depicted in Figure 4.

Figure 3: Ratio of Concentration of Anesthetic in Alveolar Gas to Inspired Gas

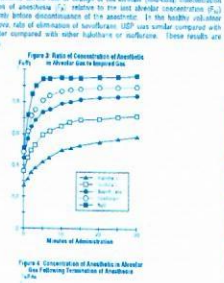
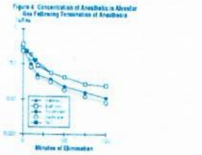


Figure 4: Concentration of Anesthetic in Alveolar Gas Following Termination of Anesthesia



Yasuda N, Lockhart S, Eger EI II, et al: Comparison of kinetics of sevoflurane and isoflurane in humans. Anesth Analg 72:316, 1991.

Protein Binding

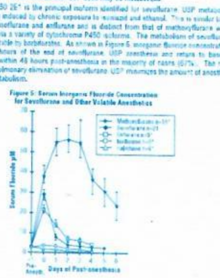
The effects of sevoflurane, USP on the displacement of drugs from serum and tissue proteins have not been investigated. Other fluorinated volatile anesthetics have been shown to displace drugs from serum and tissue proteins in vitro. The clinical significance of this is unknown. Clinical studies have shown no untoward effects when sevoflurane, USP is administered to patients taking drugs that are highly bound and have a small volume of distribution (e.g. phenytoin).

Metabolism

Sevoflurane, USP is metabolized by cytochrome P450 2E1, to hexafluoroisopropanol (HFIP) with release of inorganic fluoride and CO_2. Once formed HFIP is rapidly conjugated with glucuronic acid and eliminated as a urinary metabolite. No other metabolic pathways for sevoflurane, USP have been identified. In vivo metabolism studies suggest that approximately 5% of the sevoflurane, USP dose may be metabolized.

Cytochrome P450 2E1 is the principal isoform identified for sevoflurane, USP metabolism and this may be induced by chronic exposure to isoniazid and ethanol. This is similar to the metabolism of isoflurane and enflurane and is distinct from that of methoxyflurane which is metabolized via a variety of cytochrome P450 isoforms. The metabolism of sevoflurane, USP is not inducible by barbiturates. As shown in Figure 5, inorganic fluoride concentrations peak within 2 hours of the end of sevoflurane, USP anesthesia and return to baseline concentrations within 48 hours post-anesthesia in the majority of cases (67%). The rapid and extensive pulmonary elimination of sevoflurane, USP minimizes the amount of anesthetic available for metabolism.

Figure 5: Serum Inorganic Fluoride Concentration for Sevoflurane and Other Volatile Anesthetics



Cousins M.J., Greenstein L.R., Hitt B.A., et al: Metabolism and renal effects of enflurane in man. Anesthesiology 44:44; 1976* and Sevo-93-044

Legend
Pre-Anesth. = Pre-anesthesia

Fluoride Ion

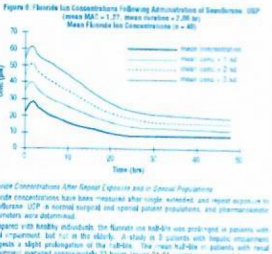
Up to 3.5% of the sevoflurane, USP dose appears in the urine as inorganic fluoride. Studies on fluoride indicate that up to 50% of fluoride clearance is nonrenal (via fluoride being taken up into bone).

PHARMACOKINETICS OF FLUORIDE ION

Fluoride ion concentrations are influenced by the duration of anesthesia, the concentration of sevoflurane, USP administered, and the composition of the anesthetic gas mixture. In studies where anesthesia was maintained purely with sevoflurane, USP for periods ranging from 1 to 6 hours, peak fluoride concentrations ranged between 12 µM and 90 µM. As shown in Figure 6, peak concentrations occur within 2 hours of the end of anesthesia and are less than 25 µM (475 ng/mL) for the majority of the population after 10 hours. The half-life is in the range of 15-23 hours.

It has been reported that following administration of methoxyflurane, serum inorganic fluoride concentrations >50 µM were correlated with the development of vasopressin-resistant, polyuric, renal failure. In clinical trials with sevoflurane, USP there were no reports of toxicity associated with elevated fluoride ion levels.

Figure 6: Fluoride Ion Concentrations Following Administration of Sevoflurane, USP (mean MAC = 1.27, mean duration = 2.06 hr) Mean Fluoride Ion Concentrations (n = 48)



Fluoride Concentrations After Repeat Exposure and in Special Populations

Fluoride concentrations have been measured after single, extended, and repeat exposure to sevoflurane, USP in normal surgical and special patient populations, and pharmacokinetic parameters were determined.

Compared with healthy individuals, the fluoride ion half-life was prolonged in patients with renal impairment, but not in the elderly. A study in 8 patients with hepatic impairment suggests a slight prolongation of the half-life. The mean half-life in patients with renal impairment averaged approximately 33 hours (range 21-61 hours) as compared to a mean of approximately 21 hours (range 10-48 hours) in normal healthy individuals. The mean half-life in the elderly (greater than 65 years) approximated 24 hours (range 18-72 hours). The mean half-life in individuals with hepatic impairment was 23 hours (range 16-47 hours). Mean maximal fluoride values (C_{max}) determined in individual studies of special populations are displayed below.

Table 1: Fluoride Ion Estimates in Special Populations Following Administration of Sevoflurane, USP

	n	Age (yr)	Duration (hr)	Dose (MAC•hr)	C_{max} (µM)
PEDIATRIC PATIENTS					
Anesthetic					
Sevoflurane-O_2	76	0 - 11	0.8	1.1	12.6
Sevoflurane-O_2	40	1 - 11	2.2	3.0	18.0
Sevoflurane/N_2O	25	5 - 13	1.9	2.4	21.3
Sevoflurane/N_2O	42	0 - 18	2.4	2.2	18.4
Sevoflurane/N_2O	40	1 - 11	2.0	2.6	15.3
ELDERLY	33	65 - 93	2.6	1.4	25.6
RENAL	21	29 - 83	2.5	1.0	26.1
HEPATIC	8	42 - 79	3.6	2.2	30.6
OBESE	35	24 - 73	3.0	1.7	38.0

n = number of patients studied.

Pharmacodynamics

Changes in the depth of sevoflurane, USP anesthesia rapidly follow changes in the inspired concentration.

In the sevoflurane, USP clinical program, the following recovery variables were evaluated:

1. **Time to events measured from the end of study drug:**
   - Time to removal of the endotracheal tube (extubation time)
   - Time required for the patient to open his/her eyes on verbal command (emergence time)
   - Time to respond to simple command (e.g., squeeze my hand) or demonstrates purposeful movement (response to command time, orientation time)
2. **Recovery of cognitive function and motor coordination was evaluated based on:**
   - psychomotor performance tests (Digit Symbol Substitution Test [DSST], Trieger Dot Test)
   - the results of subjective (Visual Analog Scale [VAS]) and objective (objective pain-discomfort scale [OPDS]) measurements
   - time to administration of the first post-anesthesia analgesic medication
   - assessments of post-anesthesia patient status
3. **Other recovery times were:**
   - time to achieve an Aldrete Score of ≥ 8
   - time required for the patient to be eligible for discharge from the recovery area, per standard criteria at site
   - time when the patient was eligible for discharge from the hospital
   - time when the patient was able to sit up or stand without dizziness

Some of these variables are summarized as follows:

Table 2: Induction and Recovery Variables for Evaluable Pediatric Patients in Two Comparative Studies: Sevoflurane versus Halothane

Time to End-Point (min)	Sevoflurane Mean ± SEM	Halothane Mean ± SEM
Induction	2.0 ± 0.2 (n=294)	2.7 ± 0.2 (n=252)
Emergence	11.3 ± 0.7 (n=293)	15.8 ± 0.8 (n=252)
Response to command	13.7 ± 1.0 (n=271)	19.3 ± 1.1 (n=230)
First analgesia	52.2 ± 8.5 (n=216)	67.6 ± 10.6 (n=150)
Eligible for recovery discharge	76.5 ± 2.0 (n=292)	81.1 ± 1.9 (n=246)

n = number of patients with recording of events.

Table 3: Recovery Variables for Evaluable Adult Patients in Two Comparative Studies: Sevoflurane versus Isoflurane

Time to Parameter (min)	Sevoflurane Mean ± SEM	Isoflurane Mean ± SEM
Emergence	7.7 ± 0.3 (n=395)	9.1 ± 0.3 (n=348)
Response to command	8.1 ± 0.3 (n=395)	9.7 ± 0.3 (n=345)
First analgesia	42.7 ± 3.0 (n=269)	52.9 ± 4.2 (n=228)
Eligible for recovery discharge	87.6 ± 5.3 (n=244)	79.1 ± 5.2 (n=252)

n = number of patients with recording of events.

Table 4: Meta-Analyses for Induction and Emergence Variables for Evaluable Adult Patients in Comparative Studies: Sevoflurane versus Propofol

Parameter	No. of Studies	Sevoflurane Mean ± SEM	Propofol Mean ± SEM
Mean maintenance anesthesia exposure	3	1.0 MAC•hr ± 0.5 (n=259)	7.2 mg/kg/hr ± 2.6 (n=258)
Time to induction (min)	1	3.1 ± 0.18* (n=93)	2.2 ± 0.18** (n=93)
Time to emergence (min)	3	8.6 ± 0.57 (n=255)	11.0 ± 0.57 (n=260)
Time to respond to command (min)	3	9.9 ± 0.60 (n=257)	12.1 ± 0.60 (n=260)
Time to first analgesia (min)	3	43.8 ± 2.79 (n=177)	57.9 ± 3.68 (n=179)
Time to eligibility for recovery discharge (min)	3	116.0 ± 4.15 (n=257)	115.6 ± 3.98 (n=261)

*Propofol induction of one sevoflurane group = mean of 178.6 mg ± 72.5 SD (n=165)
**Propofol induction of all propofol groups = mean of 170.2 mg ± 60.6 SD (n=245)
n = number of patients with recording of events.

CARDIOVASCULAR EFFECTS

Sevoflurane, USP was studied in 14 healthy volunteers (18-35 years old) comparing sevoflurane-O_2 (Sevo/O_2) to sevoflurane-N_2O/O_2 (Sevo/N_2O/O_2) during 7 hours of anesthesia. During controlled ventilation, hemodynamic parameters measured are shown in Figures 7-10:

Figure 7: Heart Rate

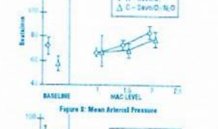


Figure 8: Mean Arterial Pressure

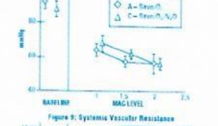


Figure 9: Systemic Vascular Resistance

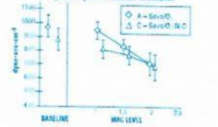
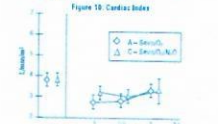


Figure 10: Cardiac Index



Sevoflurane, USP is a dose-related cardiac depressant. Sevoflurane, USP does not produce increases in heart rate at doses less than 2 MAC.

A study investigating the epinephrine induced arrhythmogenic effect of sevoflurane, USP versus isoflurane in adult patients undergoing transsphenoidal hypophysectomy demonstrated that the threshold dose of epinephrine (i.e., the dose at which the first sign of arrhythmia was observed) producing multiple ventricular arrhythmias was 5 mcg/kg with both sevoflurane, USP and isoflurane. Consequently, the interaction of sevoflurane, USP with epinephrine appears to be equal to that seen with isoflurane.

Clinical Trials

Sevoflurane, USP was administered to a total of 3185 patients prior to sevoflurane, USP NDA submission. The types of patients are summarized as follows:

Table 5: Patients Receiving Sevoflurane, USP in Clinical Trials

Type of Patients	Number Studied	
ADULT	2223	
Cesarean Delivery		29
Cardiovascular and patients at risk of myocardial ischemia		246
Neurosurgical		22
Hepatic impairment		8
Renal impairment		35
PEDIATRIC	962	

Clinical experience with these patients is described below.

ADULT ANESTHESIA

The efficacy of sevoflurane, USP in comparison to isoflurane, enflurane, and propofol was investigated in 3 outpatient and 25 inpatient studies involving 3591 adult patients. Sevoflurane, USP was found to be comparable to isoflurane, enflurane, and propofol for the maintenance of anesthesia in adult patients. Patients administered sevoflurane, USP showed shorter times (statistically significant) to some recovery events (extubation, response to command, and orientation) than patients who received isoflurane or propofol.

Mask Induction

Sevoflurane, USP has a nonpungent odor and does not cause respiratory irritability. Sevoflurane, USP is suitable for mask induction in adults. In 196 patients, mask induction was smooth and rapid, with complications occurring with the following frequencies: cough 6%, breathholding 6%, agitation 6%, laryngospasm 5%.

Ambulatory Surgery

Sevoflurane, USP was compared to isoflurane and propofol for maintenance of anesthesia supplemented with N_2O in two studies involving 786 adult (18-84 years of age) ASA Class I, II or III patients. Shorter times to emergence and response to commands (statistically significant) were observed with sevoflurane, USP compared to isoflurane and propofol.

Table 6: Recovery Parameters in Two Outpatient Surgery Studies: Least Squares Mean ± SEM

	Sevoflurane/ N_2O	Isoflurane/ N_2O	Sevoflurane/ N_2O	Propofol/ N_2O
Mean Maintenance Anesthesia Exposure ± SD	0.64 ± 0.03 MAC•hr (n=245)	0.66 ± 0.03 MAC•hr (n=249)	0.8 ± 0.5 MAC•hr (n=166)	7.3 ± 2.3 mg/kg/hr (n=166)
Time to Emergence (min)	8.2 ± 0.4 (n=246)	9.3 ± 0.3 (n=251)	8.3 ± 0.7 (n=137)	10.4 ± 0.7 (n=142)
Time to Respond to Commands (min)	8.5 ± 0.4 (n=246)	9.8 ± 0.4 (n=248)	9.1 ± 0.7 (n=139)	11.5 ± 0.7 (n=143)
Time to First Analgesia (min)	45.9 ± 4.7 (n=160)	59.1 ± 6.0 (n=252)	46.1 ± 5.4 (n=83)	60.0 ± 4.7 (n=85)
Time to Eligibility for Discharge from Recovery Area (min)	87.6 ± 5.3 (n=244)	79.1 ± 5.2 (n=252)	103.1 ± 3.8 (n=139)	105.1 ± 3.7 (n=143)

n = number of patients with recording of recovery events.

Inpatient Surgery

Sevoflurane, USP was compared to isoflurane and propofol for maintenance of anesthesia supplemented with N_2O in two multicenter studies involving 741 adult ASA Class I, II or III (18-92 years of age) patients. Shorter times to emergence, command response, and first post-anesthesia analgesia (statistically significant) were observed with sevoflurane, USP compared to isoflurane and propofol.

Table 7: Recovery Parameters in Two Inpatient Surgery Studies: Least Squares Mean ± SEM

	Sevoflurane/ N_2O	Isoflurane/ N_2O	Sevoflurane/ N_2O	Propofol/ N_2O
Mean Maintenance Anesthesia Exposure ± SD	1.27 MAC•hr ± 0.05 (n=271)	1.58 MAC•hr ± 0.06 (n=282)	1.43 MAC•hr ± 0.94 (n=93)	7.0 mg/kg/hr ± 2.9 (n=92)
Time to Emergence (min)	11.0 ± 0.6 (n=270)	16.4 ± 0.6 (n=281)	8.8 ± 1.2 (n=92)	13.2 ± 1.2 (n=92)
Time to Respond to Commands (min)	12.8 ± 0.7 (n=270)	18.4 ± 0.7 (n=281)	11.0 ± 1.20 (n=92)	14.4 ± 1.21 (n=91)
Time to First Analgesia (min)	46.1 ± 3.0 (n=233)	59.4 ± 3.2 (n=242)	37.8 ± 3.3 (n=82)	49.2 ± 3.3 (n=79)
Time to Eligibility for Discharge from Recovery Area (min)	139.2 ± 15.6 (n=268)	165.9 ± 16.3 (n=282)	148.4 ± 8.9 (n=92)	141.4 ± 8.9 (n=92)

n = number of patients with recording of recovery events.

PEDIATRIC ANESTHESIA

The concentration of sevoflurane, USP required for maintenance of general anesthesia is age-dependent (see **DOSAGE AND ADMINISTRATION**). Sevoflurane, USP or halothane was used to anesthetize 1620 pediatric patients aged 1 day to 18 years, and ASA physical status I or II (948 sevoflurane, USP 672 halothane). In one study involving 90 infants and children, there were no clinically significant decreases in heart rate compared to awake values at 1 MAC. Systolic blood pressure decreased 15-20% in comparison to awake values following administration of 1 MAC sevoflurane, USP; however, clinically significant hypotension requiring immediate intervention did not occur. Overall incidences of bradycardia [more than 20 beats/min lower than normal (80 beats/min)] in comparative studies was 3% for sevoflurane, USP and 7% for halothane. Patients who received sevoflurane, USP had slightly faster emergence times (12 vs. 19 minutes) and a higher incidence of post-anesthesia agitation (14% vs. 10%).

Sevoflurane, USP (n=91) was compared to halothane (n=89) in a single-center study for elective repair or palliation of congenital heart disease. The patients ranged in age from 9 days to 11.8 years with an ASA physical status of II, III, and IV (18%, 68%, and 13% respectively). No significant differences were demonstrated between treatment groups with respect to the primary outcome measures: cardiovascular decompensation and severe arterial desaturation. Adverse event data was limited to the study outcome variables collected during surgery and before institution of cardiopulmonary bypass.

Mask Induction

Sevoflurane, USP has a nonpungent odor and is suitable for mask induction in pediatric patients. In controlled pediatric studies in which mask induction was performed, the incidence of induction events is shown below (see **ADVERSE REACTIONS**).

Table 8: Incidence of Pediatric Induction Events

	Sevoflurane (n=836)	Halothane (n=660)
Agitation	14%	11%
Cough	6%	10%
Breathholding	5%	6%
Secretions	3%	3%
Laryngospasm	2%	2%
Bronchospasm	<1%	0%

n = number of patients.

Ambulatory Surgery

Sevoflurane, USP (n=518) was compared to halothane (n=382) for the maintenance of anesthesia in pediatric outpatients. All patients received N_2O and many received fentanyl, midazolam, bupivacaine, or lidocaine. The time to eligibility for discharge from post-anesthesia care units was similar between agents (see **CLINICAL PHARMACOLOGY** and **ADVERSE REACTIONS**).

CARDIOVASCULAR SURGERY

Coronary Artery Bypass Graft (CABG) Surgery

Sevoflurane, USP was compared to isoflurane as an adjunct with opioids in a multicenter study of 273 patients undergoing CABG surgery. Anesthesia was induced with midazolam (0.1-0.3 mg/kg), vecuronium (0.1-0.2 mg/kg), and fentanyl (5-15 mcg/kg). Both isoflurane and sevoflurane, USP were administered at loss of consciousness in doses of 1.0 MAC and titrated until the beginning of cardiopulmonary bypass to a maximum of 2.0 MAC. The total dose of fentanyl did not exceed 25 mcg/kg. The average MAC dose was 0.49 for sevoflurane, USP and 0.53 for isoflurane. There were no significant differences in hemodynamics, cardioactive drug use, or ischemia incidence between the two groups. Outcome was also equivalent. In this small multicenter study, sevoflurane, USP appears to be as effective and as safe as isoflurane for supplementation of opioid anesthesia for coronary bypass grafting.

Non-Cardiac Surgery Patients at Risk for Myocardial Ischemia

Sevoflurane-N_2O was compared to isoflurane-N_2O for maintenance of anesthesia in a multicenter study in 214 patients, age 40-87 years who were at mild-to-moderate risk for myocardial ischemia and were undergoing elective non-cardiac surgery. Forty-six percent (46%) of the operations were cardiovascular, with the remainder evenly divided between gastrointestinal and musculoskeletal and small numbers of other surgical procedures. The average duration of surgery was less than 2 hours. Anesthesia induction usually was performed with thiopental (2-5 mg/kg) and fentanyl (1-5 mcg/kg). Vecuronium (0.1-0.2 mg/kg) was also administered to facilitate intubation, muscle relaxation or immobility during surgery. The average MAC dose was 0.49 for both anesthetics. There was no significant difference between the anesthetic regimens for intraoperative hemodynamics, cardioactive drug use, or ischemic incidents, although only 83 patients in the sevoflurane, USP group and 85 patients in the isoflurane group were successfully monitored for ischemia. The outcome was also equivalent in terms of adverse events, death, and postoperative myocardial infarction. Within the limits of this small multicenter study in patients at mild-to-moderate risk for myocardial ischemia, sevoflurane, USP was a satisfactory equivalent to isoflurane in providing supplemental inhalation anesthesia to intravenous drugs.

CESAREAN SECTION

Sevoflurane, USP (n=29) was compared to isoflurane (n=27) in ASA Class I or II patients for the maintenance of anesthesia during cesarean section. Newborn evaluations and recovery events were recorded. With both anesthetics, Apgar scores averaged 8 and 9 at 1 and 5 minutes, respectively.

Use of sevoflurane, USP as part of general anesthesia for elective cesarean section produced no untoward effects in mother or neonate. Sevoflurane, USP and isoflurane demonstrated equivalent recovery characteristics. There was no difference between sevoflurane, USP and isoflurane with regard to the effect on the newborn, as assessed by Apgar Score and Neurological and Adaptive Capacity Score (average=29.5). The safety of sevoflurane, USP in labor and vaginal delivery has not been evaluated.

NEUROSURGERY

Three studies compared sevoflurane, USP to isoflurane for maintenance of anesthesia during neurosurgical procedures. In a study of 20 patients, there was no difference between sevoflurane, USP and isoflurane with regard to recovery from anesthesia. In 2 studies, a total of 22 patients with intracranial pressure (ICP) monitors received either sevoflurane, USP or isoflurane. There was no difference between sevoflurane, USP and isoflurane with regard to ICP response to inhalation of 0.5, 1.0, and 1.5 MAC inspired concentrations of volatile agent during N_2O-O_2-fentanyl anesthesia. During progressive hyperventilation from $PaCO_2$ = 40 to $PaCO_2$ = 30, ICP response to hypocarbia was preserved with sevoflurane, USP at both 0.5 and 1.0 MAC concentrations. In patients at risk for elevations of ICP, sevoflurane, USP should be administered cautiously in conjunction with ICP-reducing maneuvers such as hyperventilation.

HEPATIC IMPAIRMENT

A multicenter study (2 sites) compared the safety of sevoflurane, USP and isoflurane in 16 patients with mild-to-moderate hepatic impairment utilizing the lidocaine MEGX assay for assessment of hepatocellular function. All patients received intravenous propofol (1-3 mg/kg) or thiopental (2-7 mg/kg) for induction and succinylcholine, vecuronium, or atracurium for intubation. Sevoflurane, USP or isoflurane was administered in either 100% O_2 or up to 70% N_2O/O_2. Neither drug adversely affected hepatic function. No serum inorganic fluoride level exceeded 45 µM/L, but sevoflurane, USP patients had prolonged terminal disposition of fluoride, as evidenced by longer inorganic fluoride half-life than patients with normal hepatic function (23 hours vs. 10-48 hours).

RENAL IMPAIRMENT

Sevoflurane, USP was evaluated in renally impaired patients with baseline serum creatinine >1.5 mg/dL. Fourteen patients who received sevoflurane, USP were compared with 12 patients who received isoflurane. In another study, 21 patients who received sevoflurane, USP were compared with 20 patients who received enflurane. Creatinine levels increased in 7% of patients who received sevoflurane, USP, 8% of patients who received isoflurane, and 10% of patients who received enflurane. Because of the small number of patients with renal insufficiency (baseline serum creatinine greater than 1.5 mg/dL) studied, the safety of sevoflurane, USP administration in this group has not yet been fully established. Therefore, sevoflurane, USP should be used with caution in patients with renal insufficiency (see **WARNINGS**).

INDICATIONS AND USAGE

Sevoflurane, USP is indicated for induction and maintenance of general anesthesia in adult and pediatric patients for inpatient and outpatient surgery.

Sevoflurane, USP should be administered only by persons trained in the administration of general anesthesia. Facilities for maintenance of a patent airway, artificial ventilation, oxygen enrichment, and circulatory resuscitation must be immediately available. Since level of anesthesia may be altered rapidly, only vaporizers producing predictable concentrations of sevoflurane, USP should be used.

CONTRAINDICATIONS

Sevoflurane, USP can cause malignant hyperthermia. It should not be used in patients with known sensitivity to sevoflurane, USP or to other halogenated agents nor in patients with known or suspected susceptibility to malignant hyperthermia.

WARNINGS

Although data from controlled clinical studies at low flow rates are limited, findings taken from patient and animal studies suggest that there is a potential for renal injury which is presumed due to Compound A. Animal and human studies demonstrate that sevoflurane, USP administered for more than 2 MAC•hours and at fresh gas flow rates of <2 L/min may be associated with proteinuria and glycosuria.

While a level of Compound A exposure at which clinical nephrotoxicity might be expected to occur has not been established, it is prudent to consider all of the factors leading to Compound A exposure in humans, especially duration of exposure, fresh gas flow rate, and concentration of sevoflurane, USP. During sevoflurane, USP anesthesia the clinician should adjust inspired concentration and fresh gas flow rate to minimize exposure to Compound A. To minimize exposure to Compound A, sevoflurane, USP exposure should not exceed 2 MAC•hours at flow rates of 1 to <2 L/min. Fresh gas flow rates <1 L/min are not recommended.

Because clinical experience in administering sevoflurane, USP to patients with renal insufficiency (creatinine >1.5 mg/dL) is limited, its safety in these patients has not been established.

Sevoflurane, USP may be associated with glycosuria and proteinuria when used for long procedures at low flow rates. The safety of low flow sevoflurane, USP on renal function was evaluated in patients with normal preoperative renal function. One study compared sevoflurane, USP (N=98) to an active control (N=90) administered for ≥2 hours at a fresh gas flow rate of ≤1 Liter/minute. Per study defined criteria (Hou et al.) one patient in the sevoflurane, USP group developed elevations of creatinine, in addition to glycosuria and proteinuria. This patient received sevoflurane, USP at fresh gas flow rates of ≤800 mL/minute. Using these same criteria, there were no patients in the active control group who developed treatment emergent elevations in serum creatinine.

Sevoflurane, USP may present an increased risk in patients with known sensitivity to volatile halogenated anesthetic agents. KOH containing CO_2 absorbents are not recommended for use with sevoflurane, USP.

Reports of QT prolongation, associated with torsade de pointes (in exceptional cases, fatal), have been received. Caution should be exercised when administering sevoflurane to susceptible patients (e.g. patients with congenital Long QT Syndrome or patients taking drugs that can prolong the QT interval).

Malignant Hyperthermia

In susceptible individuals, potent inhalation anesthetic agents, including sevoflurane, USP, may trigger a skeletal muscle hypermetabolic state leading to high oxygen demand and the clinical syndrome known as malignant hyperthermia. Sevoflurane can induce malignant hyperthermia in genetically susceptible individuals, such as those with certain inherited ryanodine receptor mutations. The clinical syndrome is signaled by hypercapnia, and may include muscle rigidity, tachycardia, tachypnea, cyanosis, arrhythmias, and/or unstable blood pressure. Some of these nonspecific signs may also appear during light anesthesia, acute hypoxia, hypercapnia, and hypovolemia.

In clinical trials, one case of malignant hyperthermia was reported. In addition, there have been postmarketing reports of malignant hyperthermia. Some of these cases have been fatal. Treatment of malignant hyperthermia includes discontinuation of triggering agents (e.g., sevoflurane, USP), administration of intravenous dantrolene sodium (consult prescribing information for intravenous dantrolene sodium intravenous for additional information on patient management), and application of supportive therapy. Supportive therapy may include efforts to restore body temperature, respiratory and circulatory support as indicated, and management of electrolyte-fluid-acid-base abnormalities. Renal failure may appear later, and urine flow should be monitored and sustained if possible.

Perioperative Hyperkalemia

Use of inhaled anesthetic agents has been associated with rare increases in serum potass[...] levels that have resulted in cardiac arrhythmias and death in pediatric patients d[...] the postoperative period. Patients with latent as well as overt neuromuscular [...] particularly Duchenne muscular dystrophy, appear to be most vulnerable. C[...] use of succinylcholine has been associated with most, but not all, of these c[...] patients also experienced significant elevations in serum creatine kinase [...] some cases, changes in urine consistent with myoglobinuria. Despi[...] presentation to malignant hyperthermia, none of these patients exhibit[...] of muscle rigidity or hypermetabolic state. Early and aggressive [...] hyperkalemia and resistant arrhythmias is recommended, as is [...] latent neuromuscular disease.

PRECAUTIONS

During the maintenance of anesthesia, increasing [...] USP produces dose-dependent decreases in blood [...] insolubility in blood, these hemodynamic changes [...] volatile anesthetics. Excessive decreases in bloo[...] related to depth of anesthesia and may be corr[...] of sevoflurane, USP.

Rare cases of seizures have been repor[...] **PRECAUTIONS - Pediatric Use and A**[...]

The recovery from general anest[...] discharged from the post-anesth[...]

MPP-HK85

Drug Interactions

In clinical trials, no significant adverse reactions occurred with other drugs commonly used in the perioperative period, including central nervous system depressants, autonomic drugs, skeletal muscle relaxants, antimicrobial agents, hormones and synthetic substitutes, blood derivatives, and cardiovascular drugs.

INTRAVENOUS ANESTHETICS

Sevoflurane, USP administration is compatible with barbiturates, propofol, and other commonly used intravenous anesthetics.

BENZODIAZEPINES AND OPIOIDS

Benzodiazepines and opioids would be expected to decrease the MAC of sevoflurane, USP in the same manner as with other inhalational anesthetics. Sevoflurane, USP administration is compatible with benzodiazepines and opioids as commonly used in surgical practice.

NITROUS OXIDE

As with other halogenated volatile anesthetics, the anesthetic requirement for sevoflurane, USP is decreases when administered in combination with nitrous oxide. Using 50% N_2O the MAC equivalent dose requirement is reduced approximately 50% in adults, and approximately 25% in pediatric patients (see **DOSAGE AND ADMINISTRATION**).

NEUROMUSCULAR BLOCKING AGENTS

As is the case with other volatile anesthetics, sevoflurane USP increases both the intensity and duration of neuromuscular blockade induced by nondepolarizing muscle relaxants. When used to supplement dilantoin-N_2O anesthesia sevoflurane, USP and isoflurane equally potentiate neuromuscular block induced with pancuronium, vecuronium, or atracurium. Therefore, during sevoflurane, USP anesthesia, the dosage adjustments for these muscle relaxants are similar to those required with isoflurane.

Potentiation of neuromuscular blocking agents requires equilibration of muscle with delivered partial pressure of sevoflurane, USP. Reduced doses of neuromuscular blocking agents during induction of anesthesia may result in delayed onset of conditions suitable for endotracheal intubation or inadequate muscle relaxation.

Among available nondepolarizing agents only vecuronium, pancuronium and atracurium interactions have been studied during sevoflurane, USP anesthesia. In the absence of specific guidelines:

1. For endotracheal intubation, do not reduce the dose of nondepolarizing muscle relaxants.
2. During maintenance of anesthesia, the required dose of nondepolarizing muscle relaxants is likely to be reduced compared to that during N_2O-opioid anesthesia. Administration of supplemental doses of muscle relaxants should be guided by the response to nerve stimulation.

The effect of sevoflurane, USP on the duration of depolarizing neuromuscular blockade induced by succinylcholine has not been studied.

Hepatic Function

Results of evaluations of laboratory parameters (e.g., ALT, AST, alkaline phosphatase, and total bilirubin, etc.) as well as investigator-reported incidence of adverse events relating to liver function, demonstrate that sevoflurane, USP can be administered to patients with normal or mild-to-moderately impaired hepatic function. However, patients with severe hepatic dysfunction were not investigated.

Occasional cases of transient changes in postoperative hepatic function tests were reported with both sevoflurane, USP and reference agents. Sevoflurane, USP was found to be comparable to isoflurane with respect to these changes in hepatic function.

Very rare cases of mild, moderate and severe postoperative hepatic dysfunction or hepatitis with or without jaundice have been reported from postmarketing experiences. Clinical judgement should be exercised when sevoflurane, USP is used in patients with underlying hepatic conditions or under treatment with drugs known to cause hepatic dysfunction (see **ADVERSE REACTIONS**).

It has been reported that previous exposure to halogenated hydrocarbon anesthetics may increase the potential for hepatic injury.

Desiccated CO_2 Absorbents

An exothermic reaction occurs when sevoflurane, USP is exposed to CO_2 absorbents. This reaction is increased when the CO_2 absorbent becomes desiccated, such as after an extended period of dry gas flow through the CO_2 absorbent canisters. Rare cases of extreme heat, smoke, and/or spontaneous fire in the anesthesia breathing circuit have been reported during sevoflurane, USP use in connection with the use of desiccated CO_2 absorbent (specifically those containing potassium hydroxide (e.g., Baralyme®). KOH-containing CO_2 absorbents are not recommended for use with sevoflurane, USP. An unusually delayed rise in unexpected decline of inspired sevoflurane, USP concentration compared to the vaporizer setting may be associated with excessive heating of the CO_2 absorbent and abnormal breakdown of sevoflurane, USP.

As with other halogenated anesthetics, degradation and production of degradation products can occur when sevoflurane, USP is exposed to desiccated absorbents. When a clinician suspects that the CO_2 absorbent may be desiccated, it should be replaced. The color indicator of most CO_2 absorbents may not change upon desiccation. Therefore, the lack of significant color change should not be taken as an assurance of adequate hydration. CO_2 absorbents should be replaced routinely regardless of the state of the color indicator.

Carcinogenesis, Mutagenesis, Impairment of Fertility

Studies of carcinogenesis have not been performed for either sevoflurane, USP or Compound A. No mutagenic effect of sevoflurane, USP was noted in the Ames test, mouse micronucleus test, mouse lymphoma mutagenicity assay, human lymphocyte culture assay, micronucleus cell transformation assay, rat DNA adduct assay, and no chromosomal aberrations were induced in cultured mammalian cells.

Similarly, no mutagenic effect of Compound A was noted in the Ames test, the Chinese hamster chromosomal aberration assay and the in vivo mouse micronucleus assay. However, positive responses were observed in the human lymphocyte chromosome aberration assay. These responses were seen only at high concentrations and in the absence of metabolic activation (human S-9).

Pregnancy Category B. Reproduction studies have been performed in rats and rabbits at doses up to 1 MAC (minimum alveolar concentration) without CO_2 absorbent and have revealed no evidence of impaired fertility or harm to the fetus due to sevoflurane, USP at 0.2 MAC, the highest nontoxic dose. Developmental and reproductive toxicity studies of sevoflurane, USP in animals in the presence of strong alkalies (i.e., degradation of sevoflurane, USP and production of Compound A) have not been conducted. There are no adequate and well-controlled studies in pregnant women. Because animal reproduction studies are not always predictive of human response, sevoflurane, USP should be used during pregnancy only if clearly needed.

Labor and Delivery. Sevoflurane, USP has been used as part of general anesthesia for elective cesarean section in 20 women. There were no untoward effects in mother or neonate. (See **PHARMACODYNAMICS - Clinical Trials.**) The safety of sevoflurane, USP in labor and delivery has not been demonstrated.

Nursing Mothers. The concentrations of sevoflurane, USP in milk are probably of no clinical importance 24 hours after anesthesia. Because of rapid washout, sevoflurane, USP concentrations in milk are predicted to be below those found with many other volatile anesthetics.

Geriatric Use

MAC decreases with increasing age. The average concentration of sevoflurane, USP to achieve MAC in an 80 year old is approximately 50% of that required in a 20 year old.

Pediatric Use

Induction and maintenance of general anesthesia with sevoflurane, USP have been established in controlled clinical trials in pediatric patients aged 1 to 15 years (see **PHARMACODYNAMICS - Clinical Trials** and **ADVERSE REACTIONS**). Sevoflurane, USP has a non-pungent odor and is suitable for mask induction in pediatric patients.

The concentration of sevoflurane, USP required for maintenance of general anesthesia is age dependent. When used in combination with nitrous oxide, the MAC equivalent dose of sevoflurane, USP should be reduced in pediatric patients. MAC in premature infants has not been determined. (See **PRECAUTIONS - Drug Interactions** and **DOSAGE AND ADMINISTRATION** for recommendations in pediatric patients 1 day of age and older.)

The use of sevoflurane, USP has been associated with seizures (see **PRECAUTIONS** and **ADVERSE REACTIONS**). The majority of these have occurred in children and young adults starting from 2 months of age most of whom had no predisposing risk factors. Clinical judgement should be exercised when using sevoflurane, USP in patients who may be at risk for seizures.

ADVERSE REACTIONS

Adverse events are derived from controlled clinical trials conducted in the United States, Canada and Europe. The reference drugs were isoflurane, enflurane and propofol in adults and halothane in pediatric patients. The studies were conducted using a variety of premedications, other anesthetics, and surgical procedures of varying length. Most adverse events reported were mild and transient, and may reflect the surgical procedures, patient characteristics (including diseases) and/or medications administered.

Of the 5182 patients enrolled in the clinical trials, 2908 were exposed to sevoflurane, USP, including 110 adults and 207 pediatric patients who underwent mask induction. Each patient was counted once for each type of adverse event. Adverse events reported in patients in clinical trials and considered to be possibly or probably related to sevoflurane, USP are presented within each body system in order of decreasing frequency in the following listings. One case of malignant hyperthermia was reported in pre-registration clinical trials.

Adverse Events During the Induction Period (from Onset of Anesthesia by Mask Induction to Surgical Incision) Incidence >1%

	Adult Patients (N = 110)
Cardiovascular	Bradycardia 5%, Hypotension 4%, Tachycardia 2%
Nervous System	Agitation 7%
Respiratory System	Laryngospasm 8%, Apnea, obstruction 5%, Breathholding 5%, Cough increased 5%
	Pediatric Patients (N = 507)
Cardiovascular	Tachycardia 6%, Hypotension 4%
Nervous System	Agitation 13%
Respiratory System	Breathholding 3%, Cough increased 5%, Laryngospasm 5%, Apnea 2%
Digestive System	Increased salivation 2%

Adverse Events During Maintenance and Emergence Periods, Incidence >1% (N = 2906)

Body as a whole	Fever 1%, Shivering 6%, Hypothermia 1%, Movement 1%, Headache 1%
Cardiovascular	Hypotension 11%, Hypertension 2%, Bradycardia 5%, Tachycardia 2%
Nervous System	Somnolence 9%, Agitation 5%, Dizziness 4%, Increased salivation 4%
Digestive System	Nausea 25%, Vomiting 16%
Respiratory System	Cough increased 11%, Breathholding 2%, Laryngospasm 2%

Adverse Events, All Patients in Clinical Trials (N = 2906), All Anesthetic Periods, Incidence <1% (Reported in 3 or more patients)

Body as a whole	Asthenia, Pain
Cardiovascular	Arrhythmia, Ventricular Extrasystoles, Supraventricular Extrasystoles, Complete AV Block, Bigeminy, Hemorrhage, Inverted T Wave, Atrial Fibrillation, Atrial Arrhythmia, Second Degree AV Block, Syncope, S-T Depressed
Nervous System	Crying, Nervousness, Confusion, Hypertonia, Dry Mouth, Insomnia
Respiratory System	Sputum Increased, Apnea, Hypoxia, Wheezing, Bronchospasm, Hyperventilation, Pharyngitis, Hiccup, Hypoventilation, Dyspnea, Stridor
Metabolism and Nutrition	Increases in LDH, AST, ALT, BUN, Alkaline Phosphatase, Creatinine, Bilirubinemia, Glucosuria, Fibrinosis, Albuminuria, Hypophosphatemia, Acidosis, Hyperglycemia
Hemic and Lymphatic System	Leucocytosis, Thrombocytopenia
Skin and Special Senses	Amblyopia, Pruritus, Taste Perversion, Rash, Conjunctivitis
Urogenital	Urination Impaired, Urine Abnormality, Urinary Retention, Oliguria

See **WARNINGS** for information regarding malignant hyperthermia.

Post-Marketing Adverse Events

The following adverse events have been identified during post-approval use of Ultane (sevoflurane USP). Due to the spontaneous nature of these reports, the actual incidence and relationship of Ultane to these events cannot be established with certainty.

CNS

Seizures - Post-marketing reports indicate that sevoflurane, USP use has been associated with seizures. The majority of cases were in children and young adults, most of whom had no medical history of seizures. Several cases reported no concomitant medications and at least one case was confirmed by EEG. Although many cases were single seizures that resolved spontaneously or after treatment, cases of multiple seizures have also been reported. Seizures have occurred during, or soon after sevoflurane, USP induction, during emergence and during post-operative recovery up to a day following anesthesia.

Cardiac

Cardiac arrest

Hepatic

- Cases of mild, moderate and severe post-operative hepatic dysfunction or hepatitis with or without jaundice have been reported. Histological evidence was not provided for any of the reported hepatitis cases. In most of these cases, patients had underlying hepatic conditions or were under treatment with drugs known to cause hepatic dysfunction. Most of the reported events were transient and resolved spontaneously (see **PRECAUTIONS**).
- Hepatic necrosis
- Hepatic failure

Other

- Malignant hyperthermia (see **CONTRAINDICATIONS** and **WARNINGS**)
- Allergic reactions, such as rash, urticaria, pruritus, bronchospasm, anaphylactic or anaphylactoid reactions (see **CONTRAINDICATIONS**)
- Reports of hypersensitivity (including contact dermatitis, rash, dyskinesia, wheezing, chest discomfort, swelling face or anaphylactic reactions) have been received, particularly in association with long-term occupational exposure to inhaled anesthetic agents, including sevoflurane (see **OCCUPATIONAL CAUTION**).

Laboratory Findings

Transient elevations in glucose, low hormone tests, and white blood cell count may occur as with use of other anesthetic agents.

OVERDOSAGE

In the event of overdosage or what may appear to be overdosage the following action should be taken: discontinue administration of sevoflurane, USP; maintain a patent airway; institute assisted or controlled ventilation with oxygen; and maintain adequate cardiovascular function.

DOSAGE AND ADMINISTRATION

The concentration of sevoflurane, USP being delivered from a vaporizer during anesthesia should be known. This may be accomplished by using a vaporizer calibrated specifically for sevoflurane, USP. The administration of general anesthesia must be individualized based on the patient's response.

Replacement of Desiccated CO_2 Absorbents

When a clinician suspects that the CO_2 absorbent may be desiccated, it should be replaced. The exothermic reaction that occurs with sevoflurane, USP and CO_2 absorbents is increased when the CO_2 absorbent becomes desiccated, such as after an extended period of dry gas flow through the CO_2 absorbent canisters (see **PRECAUTIONS**).

Pre-anesthetic Medication

No specific premedication is either indicated or contraindicated with sevoflurane, USP. The decision as to whether or not to premedicate and the choice of premedication is left to the discretion of the anesthesiologist.

Induction:

Sevoflurane, USP has a non-pungent odor and does not cause respiratory irritability, making it suitable for mask induction in pediatrics and adults.

Maintenance:

Surgical levels of anesthesia can usually be achieved with concentrations of 0.5-3% sevoflurane, USP with or without the concomitant use of nitrous oxide. Sevoflurane, USP can be administered with any type of anesthesia circuit.

Table 9: MAC Values for Adults and Pediatric Patients According to Age

Age of Patient (years)	Sevoflurane in Oxygen	Sevoflurane in 65% N_2O/35% O_2
0 - 1 months *	3.3%	
1 - <6 months	3.0%	
6 months - <3 years	2.8%	2.0%@
3 - 12	2.5%	
25	2.6%	1.4%
40	2.1%	1.1%
60	1.7%	0.9%
80	1.4%	0.7%

* Neonates are full-term gestational age. MAC in premature infants has not been determined.

@ In 1 - <3 year old pediatric patients, 2.0% N_2O/40% O_2 was used.

Directions for Filling Vaporizers

UTILIZE AN ADAPTER, AS APPROPRIATE, WHEN FILLING THE VAPORIZER WITH SEVOFLURANE.

HOW SUPPLIED

Sevoflurane, USP, Volatile Liquid for Inhalation, is available as:

NDC 10019-651-64 - Aluminum bottle containing 250 mL sevoflurane, USP.

Safety and Handling

OCCUPATIONAL CAUTION:

There is an operator work exposure limit established for sevoflurane, USP. However, the National Institute for Occupational Safety and Health has recommended an 8 hour time-weighted average limit of 2 ppm for halogenated anesthetic agents in general. (2 ppm when coupled with exposure to N_2O) (see **ADVERSE REACTIONS**).

Storage

Store at controlled room temperature 15°-30°C (59°-86°F) (see USP).

The bottle cap should be replaced securely after each use of sevoflurane, USP.

Baxter

Manufactured for:
Baxter Healthcare Corporation
Deerfield, IL 60015, USA

For Product Inquiry 1 800 ANA DRUG (1-800-262-3784)

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