

## HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use OSMITROL safely and effectively. See full prescribing information for OSMITROL.

**OSMITROL (mannitol injection), for intravenous use**  
**Initial U.S. Approval: 1964**

### RECENT MAJOR CHANGES

Indications and Usage (removed, revised) (1)	11/2018
Contraindications (4)	11/2018
Warnings and Precautions (5.1, 5.2, 5.3, 5.4, 5.5, 5.6, 5.7)	11/2018

### INDICATIONS AND USAGE

OSMITROL is an osmotic diuretic, indicated for the reduction of:

- intracranial pressure and treatment of cerebral edema. (1)
- elevated intraocular pressure. (1)

### DOSAGE AND ADMINISTRATION

#### Administration Instructions (2.1):

- For intravenous infusion preferably into a large central vein.
- Prior to administration, correct fluid and electrolyte imbalances.

#### Recommended Dosage (2.2):

- The dosage, concentration and rate of administration depend on the age, weight and condition of the patient, including fluid requirement, urinary output and concomitant therapy.
- *Reduction of Intracranial Pressure:* 0.25 gram/kg administered every 6 to 8 hours as an intravenous infusion over 30 minutes.
- *Reduction of Intraocular Pressure:* 1.5 to 2 grams/kg of a 15% or 20% w/v solution as a single dose administered intravenously over at least 30 minutes.

### DOSAGE FORMS AND STRENGTHS

Injection (3):

- 5% (0.05 grams/mL): 5 grams of mannitol, USP per 100 mL in a single-dose 1000 mL flexible container
- 10% (0.1 grams/mL): 10 grams of mannitol, USP per 100 mL in a single-dose 500 mL flexible container
- 15% (0.15 grams/mL): 15 grams of mannitol, USP per 100 mL in a single-dose 500 mL flexible container
- 20% (0.2 grams/mL): 20 grams of mannitol, USP per 100 mL in single-dose 250 mL and 500 mL flexible containers

### CONTRAINDICATIONS

- Known hypersensitivity to mannitol. (4, 5.1)
- Anuria. (4, 5.2)
- Severe hypovolemia. (4, 5.4)
- Pre-existing severe pulmonary vascular congestion or pulmonary edema. (4, 5.5)
- Active intracranial bleeding except during craniotomy. (4)

### WARNINGS AND PRECAUTIONS

- **Hypersensitivity Reactions, including anaphylaxis:** Stop infusion immediately if hypersensitivity reactions develop. (5.1)
- **Renal Complications Including Renal Failure:** Risk factors include pre-existing renal failure, concomitant use of nephrotoxic drugs or other diuretics. Avoid use of nephrotoxic drugs. Discontinue OSMITROL if renal function worsens. (5.2, 8.6)
- **Central Nervous System (CNS) Toxicity:** Confusion, lethargy and coma may occur during or after infusion. Concomitant neurotoxic drugs may potentiate toxicity. Avoid use of neurotoxic drugs. Discontinue OSMITROL if CNS toxicity develops. (5.3)
- **Fluid and Electrolyte Imbalances, Hyperosmolarity:** Hypervolemia may exacerbate congestive heart failure, hyponatremia can lead to encephalopathy; hypo/hyperkalemia can result in cardiac adverse reactions in sensitive patients. Discontinue OSMITROL if fluid and/or electrolyte imbalances occur. (5.4)
- **Monitoring/Laboratory Tests:** Monitor fluid and electrolytes, serum osmolality and renal, cardiac and pulmonary function. Discontinue if toxicity develops. (5.5)
- **Infusion Site Reactions:** May include irritation and inflammation, as well as severe reactions (compartment syndrome) when associated with extravasation. (5.6)
- **Interference with Laboratory Tests:** High concentrations of mannitol may cause false low results of inorganic phosphorus blood concentrations. Mannitol may produce false positive results for blood ethylene glycol. (5.7, 7.6)

### ADVERSE REACTIONS

The most common adverse reactions are hypersensitivity reactions, renal failure, CNS toxicity, hypo/hypervolemia, hypo/hypermnatremia, hypo/hyperkalemia, and infusion site reactions. (6)

To report SUSPECTED ADVERSE REACTIONS, contact Baxter Healthcare at 1-866-888-2472 or FDA at 1-800-FDA-1088 or [www.fda.gov/medwatch](http://www.fda.gov/medwatch).

### DRUG INTERACTIONS

- **Nephrotoxic Drugs and Diuretics:** May increase the risk of renal failure; avoid concomitant use. (7.1, 7.2)
- **Neurotoxic Drugs:** May potentiate CNS toxicity of mannitol; avoid concomitant use. (7.3)
- **Drugs Affected by Electrolyte Imbalances:** May result in cardiac adverse reactions; monitor serum electrolytes and discontinue OSMITROL if cardiac status worsens. (7.4)
- **Renally Eliminated Agents:** Concomitant use may decrease the effectiveness of agents that undergo significant renal elimination. However, concomitant use of mannitol and lithium may increase risk of lithium toxicity. If concomitant use is necessary, frequently monitor lithium concentrations and for signs of toxicity. (7.5)

See 17 for PATIENT COUNSELING INFORMATION.

Revised: 11/2018

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## FULL PRESCRIBING INFORMATION

### 1 INDICATIONS AND USAGE

OSMITROL is indicated for:

- The reduction of intracranial pressure and treatment of cerebral edema;
- The reduction of elevated intraocular pressure.

### 2 DOSAGE AND ADMINISTRATION

#### 2.1 Important Preparation and Administration Instructions

- OSMITROL is for intravenous infusion preferably into a large central vein [see *Warnings and Precautions (5.5), Description (11)*].
- Prior to administration of OSMITROL, evaluate renal, cardiac and pulmonary status of the patient and correct fluid and electrolyte imbalances [see *Dosage and Administration (2.2)*].
- Do not administer OSMITROL simultaneously with blood products through the same administration set because of the possibility of pseudoagglutination or hemolysis.

#### Preparation

1. Tear overwrap down side at slit and remove solution container.
2. Visually inspect the container. If the outlet port protector is damaged, detached, or not present, discard container as solution path sterility may be impaired. Some opacity of the plastic due to moisture absorption during the sterilization process may be observed. This is normal and does not affect the solution quality or safety. The opacity will diminish gradually.
3. Check for minute leaks by squeezing inner bag firmly. If leaks are found, discard solution as sterility may be impaired.
4. Admixing OSMITROL with other medications is not recommended.
5. Do not administer unless solution is clear and seal is intact. Confirm the integrity of the bag. Use only if the bag is not damaged.
6. Inspect OSMITROL visually for particulate matter and discoloration prior to administration. If particulates or discoloration are present, discard the bag.
7. OSMITROL solutions may crystallize when exposed to low temperature. At higher concentrations, the solutions have a greater tendency to crystallize. Inspect OSMITROL for crystals prior to administration. If crystals are visible, re-dissolve by warming the solution up to 70°C, with agitation. Solutions should not be heated in water or in a microwave oven due to potential for product contamination or damage. Allow the solution to cool to room or body temperature before re-inspection for crystals and use.

## Administration

1. Suspend container from eyelet support.
2. Remove protector from outlet port at bottom of container.
3. Attach administration set. Refer to complete directions accompanying set.
4. Use administration sets with a final in-line filter because of the potential for OSMITROL crystals to form.
5. Set the vent to the closed position on a vented intravenous administration set to prevent air embolism.
6. Use a dedicated line without any connections to avoid air embolism.
7. Do not pressurize the OSMITROL bag to increase the flow rate, as air embolism can result if the residual air in the container is not fully evacuated prior to administration.
8. For single use only; discard unused portion.

## **2.2 Recommended Dosage**

Prior to administration of OSMITROL, evaluate renal, cardiac and pulmonary status of the patient and correct fluid and electrolyte imbalances.

The total dosage, concentration, and rate of administration depend on the age, weight, and condition of the patient being treated, including fluid requirement, electrolyte balance, serum osmolality, urinary output, and concomitant therapy.

The following outline of administration and dosage is only a general guide to therapy.

### Reduction of Intraocular Pressure

The recommended dosage is 1.5 to 2 grams/kg of a 20% w/v solution (7.5 to 10 mL/kg) or as a 15% w/v solution (10 to 13 mL/kg) as a single dose administered intravenously over at least 30 minutes. When used preoperatively, administer OSMITROL sixty to ninety minutes before surgery to achieve maximal reduction of intraocular pressure before operation.

### Reduction of Intracranial Pressure

Usually a maximum reduction in intracranial pressure can be achieved with a dose of 0.25 gram/kg given intravenously as an intravenous infusion over 30 minutes which may be repeated every six to eight hours.

During and following OSMITROL infusion, monitor fluid and electrolytes, serum osmolality, and renal, cardiac and pulmonary function. Discontinue OSMITROL if renal, cardiac, or pulmonary status worsens or CNS toxicity develops [*see Warnings and Precautions (5.2, 5.3, 5.4, 5.5)*].

## **3 DOSAGE FORMS AND STRENGTHS**

OSMITROL Injection:

- 5% (0.05 grams/mL): 5 grams of mannitol, USP per 100 mL in a single-dose 1000 mL flexible container
- 10% (0.1 grams/mL): 10 grams of mannitol, USP per 100 mL in a single-dose 500 mL flexible container
- 15% (0.15 grams/mL): 15 grams of mannitol, USP per 100 mL in a single-dose 500 mL flexible container
- 20% (0.2 grams/mL): 20 grams of mannitol, USP per 100 mL in single-dose 250 mL and 500 mL flexible containers

## 4 CONTRAINDICATIONS

OSMITROL is contraindicated in patients with:

- Known hypersensitivity to mannitol [*see Warnings and Precautions (5.1)*]
- Anuria [*see Warnings and Precautions (5.2)*]
- Severe hypovolemia [*see Warnings and Precautions (5.4)*]
- Pre-existing severe pulmonary vascular congestion or pulmonary edema [*see Warnings and Precautions (5.5)*]
- Active intracranial bleeding except during craniotomy

## 5 WARNINGS AND PRECAUTIONS

### 5.1 Hypersensitivity Reactions

Serious hypersensitivity reactions, including anaphylaxis, hypotension and dyspnea resulting in cardiac arrest and death have been reported with OSMITROL [*see Adverse Reactions (6)*].

Stop the infusion immediately if signs or symptoms of a suspected hypersensitivity reaction develop. Initiate appropriate therapeutic countermeasures as clinically indicated. During and following OSMITROL infusion, monitor the patient clinically and monitor laboratory tests for changes in fluid and electrolyte status. Discontinue OSMITROL if renal function worsens [*see Warnings and Precautions (5.5)*].

### 5.2 Renal Complications Including Renal Failure

Renal complications, including irreversible renal failure have been reported in patients receiving mannitol. In patients with severe renal impairment, a test dose should be utilized. A second test dose may be tried if there is an inadequate response, but no more than two test doses should be attempted [*see Dosage and Administration (2.2)*].

Reversible, oliguric acute kidney injury (AKI) has occurred in patients with normal pretreatment renal function who received large intravenous doses of OSMITROL.

Although the osmotic nephrosis associated with OSMITROL administration is in principle reversible, osmotic nephrosis in general is known to potentially proceed chronic or even end-stage renal failure. Monitor renal function closely during OSMITROL infusion. Patients with pre-existing renal disease, patients with conditions that put them at high risk for renal failure, or those receiving potentially nephrotoxic drugs or other diuretics, are at increased risk of renal failure following administration of OSMITROL. Avoid concomitant administration of nephrotoxic drugs (e.g., aminoglycosides) or, other diuretics with OSMITROL, if possible [see *Drug Interactions (7.1, 7.2)*].

Patients with oliguric AKI who subsequently develop anuria while receiving mannitol are at risk of congestive heart failure, pulmonary edema, hypertensive crisis, coma and death.

During and following OSMITROL infusion for the reduction in intracranial pressure, monitor the patient clinically and laboratory tests for changes in fluid and electrolyte status. Discontinue OSMITROL if CNS toxicity develops. [see *Warnings and Precautions (5.5)*].

OSMITROL should be administered with caution to patients with impaired renal function [see *Dosage and Administration (2.2)*].

If the urine output declines during OSMITROL infusion, the patient's clinical status should be closely monitored for developing renal impairment, and the OSMITROL infusion suspended, if necessary.

OSMITROL should not be administered in patients with renal dysfunction until volume and electrolytes have been restored [see *Warnings and Precautions (5.5)*].

### **5.3 Central Nervous System (CNS) Toxicity**

CNS toxicity manifested by, e.g., confusion, lethargy, coma has been reported in patients treated with mannitol, some resulting in death, in particular in the presence of impaired renal function CNS toxicity may result from high serum mannitol concentrations, serum hyperosmolarity resulting in intracellular dehydration within CNS, hyponatremia or other disturbances of electrolyte and acid/base balance secondary to mannitol administration [see *Warnings and Precautions (5.4)*].

At high concentrations, mannitol may cross the blood brain barrier and interfere with the ability of the brain to maintain the pH of the cerebrospinal fluid especially in the presence of acidosis.

In patients with preexisting compromise of the blood brain barrier, the risk of increasing cerebral edema (general and focal) associated with repeated or continued use of OSMITROL must be individually weighed against the expected benefits.

A rebound increase of intracranial pressure may occur several hours after the infusion. Patients with a compromised blood brain barrier are at increased risk.

Concomitant administration of neurotoxic drugs (e.g., aminoglycosides) with OSMITROL may potentiate neurotoxicity. Avoid concomitant use of neurotoxic drugs, if possible [see *Drug Interactions (7.3)*].

#### 5.4 Fluid and Electrolyte Imbalances, Hyperosmolarity

Depending on dosage and duration, administration of OSMITROL may result in hypervolemia leading to or exacerbating existing congestive heart failure. Accumulation of mannitol due to insufficient renal excretion increases risk of hypervolemia. Mannitol-induced osmotic diuresis may cause or worsen dehydration/hypovolemia and hemoconcentration. Administration of OSMITROL may also cause hyperosmolarity [see *Description (11)*].

Depending on dosage and duration of administration, electrolyte and acid/base imbalances may also result from transcellular shifts in water and electrolytes, osmotic diuresis and/or other mechanisms. Such imbalances may be severe and potentially fatal.

Imbalances that may result from OSMITROL administration include:

- Hyponatremia, dehydration and hemoconcentration
- Hyponatremia, which can lead to headache, nausea, seizures, lethargy, coma, cerebral edema, and death. Acute symptomatic hyponatremic encephalopathy is considered a medical emergency.
- Hypo/hyperkalemia. The development of electrolyte imbalances (e.g., hyperkalemia, hypokalemia) associated with mannitol administration may result in cardiac adverse reactions in patients receiving drugs that are sensitive to such imbalances (e.g., digoxin, agents that may cause QT prolongation, neuromuscular blocking agents) [see *Drug Interactions (7.4)*].
- Other electrolyte disturbances
- Metabolic acidosis/alkalosis

Pediatric patients less than two years of age, particularly preterm and term neonates, may be at higher risk for fluid and electrolyte abnormalities following OSMITROL administration due to decreased glomerular filtration rate and limited ability to concentrate urine [see *Use in Specific Populations (8.4)*]

During and following OSMITROL infusion for the reduction in intracranial pressure, monitor fluid and electrolyte status and discontinue OSMITROL if imbalances occur [see *Warnings and Precautions (5.5)*].

### **5.5 Monitoring/Laboratory Tests**

During and following OSMITROL infusion for the reduction in intracranial pressure, monitor:

- serum osmolarity, serum electrolytes (including sodium, potassium, calcium and phosphate) and acid base balance,
- the osmol gap
- signs of hypo- or hypervolemia, including urine output
- renal, cardiac and pulmonary function
- intracranial pressure

Discontinue OSMITROL if renal, cardiac, or pulmonary status worsens or CNS toxicity develops [see *Contraindications (4)*].

### **5.6 Infusion Site Reactions**

The infusion of hypertonic solutions through a peripheral vein, including OSMITROL at a concentration of 10% w/v or greater, may result in peripheral venous irritation, including phlebitis. Other severe infusion site reactions, such as compartment syndrome and swelling associated with extravasation, can occur with administration of OSMITROL [see *Adverse Reactions (6)*]. OSMITROL is preferably for administration into a large central vein [see *Dosage and Administration (2.1)*].

### **5.7 Interference with Laboratory Tests**

High concentrations of mannitol can cause false low results for inorganic phosphorus blood concentrations [see *Drug Interactions (7.6)*].

Mannitol may produce false positive results in tests for blood ethylene glycol concentrations [see *Drug Interactions (7.6)*].

## 6 ADVERSE REACTIONS

The following adverse reactions from voluntary reports or clinical studies have been reported with OSMITROL. Because many of these reactions were reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

- *Hypersensitivity reactions*: cardiac arrest, anaphylaxis, hypotension, dyspnea, hypertension, pyrexia, chills, sweating, cough, musculoskeletal stiffness, myalgia, urticarial/rash, pruritus, generalized pain, discomfort, nausea, vomiting, and headache. [see *Warnings and Precautions (5.1)*]
- *Renal Failure*: acute kidney injury, osmotic nephrosis, azotemia, anuria, hematuria, oliguria, polyuria [see *Warnings and Precautions (5.2)*]
- *CNS Toxicity*: coma, seizures, confusion, lethargy; rebound increase in intracranial pressure; dizziness [see *Warnings and Precautions (5.3)*]
- *Fluid and Electrolyte Imbalances*: hypovolemia, hypervolemia, peripheral edema, dehydration, hyponatremia, hypernatremia, hyperkalemia, hypokalemia; metabolic acidosis [see *Warnings and Precautions (5.4)*]
- *Infusion Site Reactions*: phlebitis, inflammation, pain, rash, erythema, pruritus; compartment syndrome and swelling associated with extravasation [see *Warnings and Precautions (5.6)*]
- *Cardiac and Respiratory Disorders*: congestive cardiac failure, pulmonary edema, palpitations
- *Gastrointestinal Disorders*: thirst, dry mouth
- *General Disorders*: asthenia, malaise

## 7 DRUG INTERACTIONS

### 7.1 Nephrotoxic Drugs

Concomitant administration of nephrotoxic drugs (e.g., cyclosporine, aminoglycosides) increases the risk of renal failure following administration of mannitol. Avoid use of nephrotoxic drugs with OSMITROL, if possible [see *Warnings and Precautions (5.2)*].

## 7.2 Diuretics

Concomitant administration of other diuretics may potentiate the renal toxicity of mannitol. Avoid use of other diuretics with OSMITROL, if possible [*see Warnings and Precautions (5.2)*].

## 7.3 Neurotoxic Drugs

Concomitant administration of systemic neurotoxic drugs (e.g., aminoglycosides) with OSMITROL may potentiate the CNS toxicity of mannitol. Avoid use of systemic neurotoxic drugs with OSMITROL, if possible [*see Warnings and Precautions (5.3)*].

## 7.4 Drugs Affected by Electrolyte Imbalances

The development of electrolyte imbalances (e.g., hyperkalemia, hypokalemia) associated with mannitol administration may result in cardiac adverse reactions in patients receiving drugs that are sensitive to such imbalances (e.g., digoxin, drugs that prolong the QT interval, neuromuscular blocking agents) [*see Warnings and Precautions (5.4)*]. During and following OSMITROL infusion, monitor serum electrolytes and discontinue OSMITROL if cardiac status worsens [*see Warnings and Precautions (5.5)*].

## 7.5 Renally Eliminated Drugs

Mannitol therapy may increase the elimination, and decrease the effectiveness of treatment with, drugs that undergo significant renal elimination. Concomitant administration of mannitol with lithium may initially increase the elimination of lithium but may also increase the risk of lithium toxicity if patients develop hypovolemia or renal impairment. In patients receiving lithium, consider holding lithium doses during treatment with OSMITROL. In patients requiring concomitant administration of lithium and OSMITROL, frequently monitor serum lithium concentrations and for signs of lithium toxicity.

## 7.6 Interference with Laboratory Tests

High concentrations of mannitol can cause false low results for inorganic phosphorus blood concentrations when an assay based on the conversion of phosphate (orthophosphate) to the phosphomolybdate complex is used.

Mannitol may produce false positive results in tests for blood ethylene glycol concentrations in which mannitol is initially oxidized to an aldehyde.

## **8 USE IN SPECIFIC POPULATIONS**

### **8.1 Pregnancy**

Animal reproduction studies have not been conducted with mannitol. It is also not known whether mannitol can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity. OSMITROL should be given to a pregnant woman only if clearly needed.

### **8.2 Labor and Delivery**

Studies have not been conducted to evaluate the effects of mannitol on labor and delivery. Caution should be exercised when administering OSMITROL during labor and delivery.

### **8.3 Nursing Mothers**

It is not known whether mannitol is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when OSMITROL is administered to a nursing woman.

### **8.4 Pediatric Use**

OSMITROL is approved for use in the pediatric population for the reduction of intracranial and intraocular pressure. Studies have not defined the optimal dose of OSMITROL in the pediatric population. The safety profile for mannitol use in pediatric patients is similar to adults at dosages described in labeling. However, pediatric patients less than two years of age, particularly preterm and term neonates, may be at higher risk for fluid and electrolyte abnormalities following OSMITROL administration due to decreased glomerular filtration rate and limited ability to concentrate urine [see *Warnings and Precautions (5.4)*]

### **8.5 Geriatric Use**

Mannitol is known to be substantially excreted by the kidney and the risk of adverse reactions to this drug may be greater in elderly patients with impaired renal function. Evaluate the renal, cardiac and pulmonary status of the patient and correct fluid and electrolyte imbalances prior to administration of OSMITROL [see *Warnings and Precautions (5.2, 5.3, 5.4, 5.5)*].

### **8.6 Renal Impairment**

Patients with pre-existing renal disease, patients with conditions that put them at high risk for renal failure, or those receiving potentially nephrotoxic drugs or other diuretics, are at increased risk of renal failure with administration of mannitol. Evaluate the renal, cardiac

and pulmonary status of the patient and correct fluid and electrolyte imbalances prior to administration of OSMITROL [see *Warnings and Precautions* (5.2, 5.3, 5.4, 5.5)].

## 10 OVERDOSAGE

Signs and symptoms of overdose with OSMITROL include renal failure and AKI, hypo/hypervolemia, hyperosmolarity and electrolyte imbalances, CNS toxicity (e.g., coma, seizures), some of which can be fatal [see *Warnings and Precautions* (5.2, 5.3, 5.4)].

Management of overdosage with OSMITROL is symptomatic and supportive. Discontinue the infusion and institute appropriate corrective measures with particular attention to renal, cardiac and pulmonary systems. Correct fluid and electrolyte imbalances.

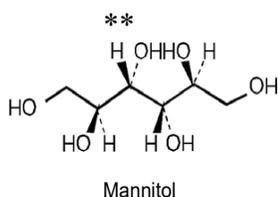
OSMITROL is dialyzable (hemodialysis and peritoneal dialysis), hemodialysis may increase OSMITROL elimination.

## 11 DESCRIPTION

OSMITROL is a sterile, nonpyrogenic solution of Mannitol, USP in a single-dose flexible container for intravenous administration. It contains no antimicrobial agents. Mannitol is a six carbon sugar alcohol prepared commercially by the reduction of dextrose. Although virtually inert metabolically in humans, it occurs naturally in fruits and vegetables. OSMITROL is an osmotic diuretic. The pH is adjusted with sodium hydroxide or hydrochloric acid. Composition, osmolarity, and pH are shown in Table 1.

Table 1	Size (mL)	Composition	*Osmolarity (mOsmol/L) (calc)	pH
		Mannitol, USP (g/L)		
5% OSMITROL	1000	50	274	5.0 (4.5 TO 7.0)
10% OSMITROL	500	100	549	5.0 (4.5 TO 7.0)
15% OSMITROL	500	150	823	5.0 (4.5 TO 7.0)
20% OSMITROL	250	200	1098	5.0 (4.5 TO 7.0)
	500			

\*Normal physiologic osmolarity range is approximately 280 to 310 mOsmol/L.



The plastic container is fabricated from a specially formulated polyvinyl chloride (PL 146 Plastic). The amount of water that can permeate from inside the container into the overwrap is insufficient to affect the solution significantly. Solutions in contact with the plastic container can leach out certain of its chemical components in very small amounts within the expiration period, e.g., di-2-ethylhexyl phthalate (DEHP), up to 5 parts per million. However, the safety of the plastic has been confirmed in tests in animals according to USP biological tests for plastic containers as well as by tissue culture toxicity studies.

## 12 CLINICAL PHARMACOLOGY

### 12.1 Mechanism of Action

Mannitol, when administered intravenously, exerts its osmotic diuretic effect as a solute of relatively small molecular size being largely confined to the extracellular space. Mannitol hinders tubular reabsorption of water and enhances excretion of sodium and chloride by elevating the osmolarity of the glomerular filtrate.

This increase in extracellular osmolarity affected by the intravenous administration of mannitol will induce the movement of intracellular water to the extracellular and vascular spaces. This action underlies the role of mannitol in reducing intracranial pressure, intracranial edema, and intraocular pressure.

### 12.3 Pharmacokinetics

#### Distribution

Mannitol distributes largely in the extracellular space in 20 to 40 minutes after intravenous administration. The volume of distribution of mannitol is approximately 17 L in adults.

#### Elimination

In subjects with normal renal function, the total clearance is 87 to 109 mL/min. The elimination half-life of mannitol is 0.5 to 2.5 hours.

#### *Metabolism*

Only relatively small amount of the dose administered is metabolized after intravenous administration of OSMITROL to healthy subjects.

#### *Excretion*

Mannitol is eliminated primarily via the kidneys in unchanged form. Mannitol is filtered by the glomeruli, exhibits less than 10% of tubular reabsorption, and is not secreted by tubular cells. Following intravenous administration, approximately 80% of an administered dose of mannitol is estimated to be excreted in the urine in three hours with lesser amounts thereafter.

### Specific Populations

#### *Patients with Renal Impairment*

In patients with renal impairment, the elimination half-life of mannitol is prolonged. In a published study, in patients with renal impairment including acute renal failure and end stage renal failure, the elimination half-life of mannitol was estimated at about 36 hours, based on serum osmolarity. In patients with renal impairment on dialysis, the elimination half-life of mannitol was reduced to 6 and 21 hours during hemodialysis and peritoneal dialysis, respectively. [see *Use in Specific Populations (8.6)*, *Overdosage (10)*].

## **16 HOW SUPPLIED/STORAGE AND HANDLING**

OSMITROL injection is supplied in single-dose, flexible VIAFLEX plastic containers and is available as follows:

<b>Code</b>	<b>Size (mL)</b>	<b>NDC</b>	<b>Product Name</b>
2D5604	1000	0338-0351-04	5% (0.05 g/mL mannitol, USP)
2D5613	500	0338-0353-03	10% (0.1 g/mL mannitol, USP)
2D5623	500	0338-0355-03	15% (0.15 g/mL mannitol, USP)
2D5632	250	0338-0357-02	20% (0.2 g/mL mannitol, USP)
2D5633	500	0338-0357-03	20% (0.2 g/mL mannitol, USP)

Do not remove container from overwrap until intended for use.

Exposure of pharmaceutical products to heat should be minimized. Avoid excessive heat. Store at room temperature (25°C); brief exposure up to 40°C does not adversely affect the product.

## **17 PATIENT COUNSELING INFORMATION**

Inform patients or caregivers of the following risks of OSMITROL:

- Hypersensitivity Reactions [see *Warnings and Precautions (5.1)*].
- Renal Failure [see *Warnings and Precautions (5.2)*]
- CNS Toxicity [see *Warnings and Precautions (5.3)*]

- Fluid and Electrolyte Imbalances, Hyperosmolarity [*see Warnings and Precautions (5.4)*]
- Infusion Site Reactions [*see Warnings and Precautions (5.6)*]

Manufactured by:

Baxter Healthcare Corporation

Deerfield, IL 60015 USA

Printed in USA

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