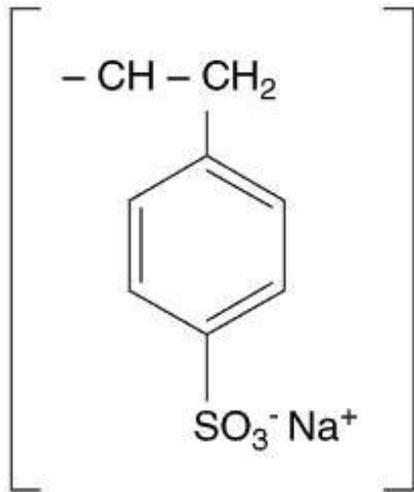


SODIUM POLYSTYRENE SULFONATE- sodium polystyrene sulfonate powder, for suspension

Belcher Pharmaceuticals, LLC

SODIUM POLYSTYRENE SULFONATE for suspension

Sodium Polystyrene Sulfonate, USP is a benzene, diethenyl-polymer, with ethenylbenzene, sulfonated, sodium salt and has the following structural formula:



The drug is a cream to light brown finely ground, powdered form of Sodium polystyrene sulfonate, USP a cation-exchange resin prepared in the sodium phase with an in vitro exchange capacity of approximately 3.1 mEq (in vivo approximately 1 mEq) of potassium per gram. The sodium content is approximately 100 mg (4.1 mEq) per gram of the drug. It can be administered orally or in an enema.

CLINICAL PHARMACOLOGY

As the resin passes along the intestine or is retained in the colon after administration by enema, the sodium ions are partially released and are replaced by potassium ions. For the most part, this action occurs in the large intestine, which excretes potassium ions to a greater degree than does the small intestine. The efficiency of this process is limited and unpredictably variable. It commonly approximates the order of 33 percent but the range is so large that definitive indices of electrolyte balance must be clearly monitored. Metabolic data are unavailable.

INDICATIONS AND USAGE

Sodium Polystyrene Sulfonate for suspension is indicated for the treatment of hyperkalemia.

CONTRAINDICATIONS

Sodium Polystyrene Sulfonate for suspension is contraindicated in the following

conditions: patients with hypokalemia, patients with a history of hypersensitivity to polystyrene sulfonate resins, obstructive bowel disease, neonates with reduced gut motility (postoperatively or drug induced) and oral administration in neonates (see PRECAUTIONS).

WARNINGS

Intestinal Necrosis

Cases of intestinal necrosis, which may be fatal, and other serious gastrointestinal adverse events (bleeding, ischemic colitis, perforation) have been reported in association with Sodium Polystyrene Sulfonate for suspension use. The majority of these cases reported the concomitant use of sorbitol. Risk factors for gastrointestinal adverse events were present in many of the cases including prematurity, history of intestinal disease or surgery, hypovolemia, and renal insufficiency and failure. Concomitant administration of sorbitol is not recommended (see PRECAUTIONS, Drug Interactions).

- Use only in patients who have normal bowel function. Avoid use in patients who have not had a bowel movement post-surgery.
- Avoid use in patients who are at risk for developing constipation or impaction (including those with history of impaction, chronic constipation, inflammatory bowel disease, ischemic colitis, vascular intestinal atherosclerosis, previous bowel resection, or bowel obstruction).
- Discontinue use in patients who develop constipation.

Alternative Therapy in Severe Hyperkalemia

Since effective lowering of serum potassium with Sodium Polystyrene Sulfonate for suspension may take hours to days, treatment with this drug alone may be insufficient to rapidly correct severe hyperkalemia associated with states of rapid tissue breakdown (e.g., burns and renal failure) or hyperkalemia so marked as to constitute a medical emergency. Therefore, other definitive measures, including dialysis, should always be considered and may be imperative.

Hypokalemia

Serious potassium deficiency can occur from therapy with Sodium Polystyrene Sulfonate for suspension. The effect must be carefully controlled by frequent serum potassium determinations within each 24 hour period. Since intracellular potassium deficiency is not always reflected by serum potassium levels, the level at which treatment with Sodium Polystyrene Sulfonate for suspension should be discontinued must be determined individually for each patient. Important aids in making this determination are the patient's clinical condition and electrocardiogram. Early clinical signs of severe hypokalemia include a pattern of irritable confusion and delayed thought processes.

Electrocardiographically, severe hypokalemia is often associated with a lengthened Q-T interval, widening, flattening, or inversion of the T wave, and prominent U waves. Also, cardiac arrhythmias may occur, such as premature atrial, nodal, and ventricular contractions, and supraventricular and ventricular tachycardias. The toxic effects of digitalis are likely to be exaggerated. Marked hypokalemia can also be manifested by severe muscle weakness, at times extending into frank paralysis.

Electrolyte Disturbances

Like all cation-exchange resins, Sodium Polystyrene Sulfonate for suspension is not totally selective (for potassium) in its actions, and small amounts of other cations such as magnesium and calcium can also be lost during treatment. Accordingly, patients receiving Sodium Polystyrene Sulfonate for suspension should be monitored for all applicable electrolyte disturbances.

Systemic Alkalosis

Systemic alkalosis has been reported after cation-exchange resins were administered orally in combination with nonabsorbable cation-donating antacids and laxatives such as magnesium hydroxide and aluminum carbonate. Magnesium hydroxide should not be administered with Sodium Polystyrene Sulfonate for suspension. One case of grand mal seizure has been reported in a patient with chronic hypocalcemia of renal failure who was given Sodium Polystyrene Sulfonate for suspension with magnesium hydroxide as laxative. (See PRECAUTIONS, Drug Interactions.)

PRECAUTIONS

Caution is advised when Sodium Polystyrene Sulfonate for suspension is administered to patients who cannot tolerate even a small increase in sodium loads (i.e., severe congestive heart failure, severe hypertension, or marked edema). In such instances compensatory restriction of sodium intake from other sources may be indicated.

In the event of clinically significant constipation, treatment with Sodium Polystyrene Sulfonate for suspension should be discontinued until normal bowel motion is resumed (see WARNINGS, Intestinal Necrosis).

DRUG INTERACTIONS

Antacids

The simultaneous oral administration of Sodium Polystyrene Sulfonate for suspension with nonabsorbable cation-donating antacids and laxatives may reduce the resin's potassium exchange capability.

Non-absorbable cation-donating antacids and laxatives

Systemic alkalosis has been reported after cation-exchange resins were administered orally in combination with nonabsorbable cation-donating antacids and laxatives such as magnesium hydroxide and aluminum carbonate. Magnesium hydroxide should not be administered with Sodium Polystyrene Sulfonate for suspension. One case of grand mal seizure has been reported in a patient with chronic hypocalcemia of renal failure who was given Sodium Polystyrene Sulfonate for suspension with magnesium hydroxide as a laxative.

Intestinal obstruction due to concretions of aluminum hydroxide when used in combination with Sodium Polystyrene Sulfonate for suspension has been reported.

Digitalis

The toxic effects of digitalis on the heart, especially various ventricular arrhythmias and A-V nodal dissociation, are likely to be exaggerated by hypokalemia, even in the face of serum digoxin concentrations in the "normal range". (See WARNINGS.)

Sorbitol

Concomitant use of Sorbitol with Sodium Polystyrene Sulfonate for suspension has been implicated in cases of intestinal necrosis, which may be fatal. Therefore, concomitant administration is not recommended. (See WARNINGS.)

Lithium

Sodium Polystyrene Sulfonate for suspension may decrease absorption of lithium.

Thyroxine

Sodium Polystyrene Sulfonate for suspension may decrease absorption of thyroxine.

Carcinogenesis, Mutagenesis, Impairment of Fertility

Studies have not been performed.

Pregnancy Category C

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

Nursing Mothers

It is not known whether this drug is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when Sodium Polystyrene Sulfonate for suspension is administered to a nursing woman.

Pediatric Use

The effectiveness of Sodium Polystyrene Sulfonate for suspension in pediatric patients has not been established. In neonates, Sodium Polystyrene Sulfonate for suspension should not be given by the oral route. In both children and neonates particular care should be observed with rectal administration, as excessive dosage or inadequate dilution could result in impaction of the resin.

Due to the risk of digestive hemorrhage or intestinal necrosis, particular care should be observed in premature infants or low birth weight infants.

ADVERSE REACTIONS

Sodium Polystyrene Sulfonate for suspension may cause some degree of gastric irritation. Anorexia, nausea, vomiting, and constipation may occur especially if high doses are given. Also, hypokalemia, hypocalcemia, hypomagnesemia and significant sodium retention, and their related clinical manifestations, may occur (see WARNINGS). Occasionally diarrhea develops. Large doses in elderly individuals may cause fecal impaction (see PRECAUTIONS). Rare instances of intestinal necrosis have been reported. Intestinal obstruction due to concretions of aluminum hydroxide, when used in combination with Sodium Polystyrene Sulfonate for suspension has been reported.

The following events have been reported from worldwide post marketing experience:

- Fecal impaction following rectal administration, particularly in children; Gastrointestinal concretions (bezoars) following oral administration;

- Ischemic colitis, gastrointestinal tract ulceration or necrosis which could lead to intestinal perforation; and,
- Rare cases of acute bronchitis and/or broncho-pneumonia associated with inhalation of particles of polystyrene sulfonate.

OVERDOSAGE

Overdosage may result in electrolyte disturbances including hypokalemia, hypocalcemia, and hypomagnesemia. Biochemical disturbances resulting from overdosage may give rise to clinical signs and symptoms of hypokalemia, including: irritability, confusion, delayed thought processes, muscle weakness, hyporeflexia, which may progress to frank paralysis and/or apnea. Tetany may occur. Electrocardiographic changes may be consistent with hypokalemia or hypocalcemia; cardiac arrhythmias may occur. Appropriate measures should be taken to correct serum electrolytes (potassium, calcium, magnesium), and the resin should be removed from the alimentary tract by appropriate use of laxatives or enemas.

DOSAGE AND ADMINISTRATION

Suspension of this drug should be freshly prepared and not stored beyond 24 hours.

The average daily adult dose of the resin is 15 g to 60 g. This is best provided by administering 15 g (approximately 4 level teaspoons) of Sodium Polystyrene Sulfonate for suspension one to four times daily. One gram of Sodium Polystyrene Sulfonate for suspension contains 4.1 mEq of sodium; one level teaspoon contains approximately 3.5 g of Sodium Polystyrene Sulfonate for suspension and 15 mEq of sodium. (A heaping teaspoon may contain as much as 10 g to 12 g of Sodium Polystyrene Sulfonate for suspension.) Since the in vivo efficiency of sodium-potassium exchange resins is approximately 33 percent, about one third of the resin's actual sodium content is being delivered to the body.

In smaller children and infants, lower doses should be employed by using as a guide a rate of 1 mEq of potassium per gram of resin as the basis for calculation.

Each dose should be given as a suspension in a small quantity of water or, for greater palatability, in syrup. The amount of fluid usually ranges from 20 mL to 100 mL, depending on the dose, or may be simply determined by allowing

3 mL to 4 mL per gram of resin. Healthcare professionals should follow full aspiration precautions when administering this product, such as placing and maintaining the patient in an upright position while the resin is being administered.

The resin may be introduced into the stomach through a plastic tube and, if desired, mixed with a diet appropriate for a patient in renal failure.

The resin may also be given, although with less effective results, in an enema consisting (for adults) of 30 g to 50 g every six hours. Each dose is administered as a warm emulsion (at body temperature) in 100 mL of aqueous vehicle. The emulsion should be agitated gently during administration. The enema should be retained as long as possible and followed by a cleansing enema.

After an initial cleansing enema, a soft, large size (French 28) rubber tube is inserted into

the rectum for a distance of about 20 cm, with the tip well into the sigmoid colon, and taped in place. The resin is then suspended in the appropriate amount of aqueous vehicle at body temperature and introduced by gravity, while the particles are kept in suspension by stirring. The suspension is flushed with 50 mL or 100 mL of fluid, following which the tube is clamped and left in place. If back leakage occurs, the hips are elevated on pillows or a knee-chest position is taken temporarily. A somewhat thicker suspension may be used, but care should be taken that no paste is formed, because the latter has a greatly reduced exchange surface and will be particularly ineffective if deposited in the rectal ampulla. The suspension is kept in the sigmoid colon for several hours, if possible. Then, the colon is irrigated with nonsodium containing solution at body temperature in order to remove the resin. Two quarts of flushing solution may be necessary. The returns are drained constantly through a Y tube connection. While the use of sorbitol is not recommended, particular attention should be paid to this cleansing enema if sorbitol has been used.

The intensity and duration of therapy depend upon the severity and resistance of hyperkalemia.

Sodium Polystyrene Sulfonate for suspension should not be heated for to do so may alter the exchange properties of the resin.

HOW SUPPLIED

Sodium Polystyrene Sulfonate for suspension is available as a cream to light brown, finely ground powder in bottles of 1 pound (454 g), NDC 62250-660-23.

Store at 20° to 25° C (68° to 77° F) with excursions permitted between 15° to 30° C (59° - 86° F) [See USP Controlled Room Temperature].

Call your doctor for medical advice about side effects. You may report side effects to Belcher Pharmaceuticals, LLC at 1-727-471-0850 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

Rx Only

Manufactured by

Belcher Pharmaceuticals, LLC

Largo, FL 33777

USA

Revised December 2015

L49I D-1512

PRINCIPAL DISPLAY PANEL - 454 g Bottle

Sodium Polystyrene Sulfonate for suspension

Read package insert.

Average adult dose: 15 g (approximately 4 level teaspoons) one to four time daily in water. See

complete prescribing information.

The effect must be carefully controlled by frequent serum potassium determinations within each 24 hour period. Sodium content approximately 60 mEq per 15 g.

Suspension should be freshly prepared and not stored beyond 24 hours. dispense in tight, light-resistant containers as defined in the official compendia.

Store at 20° to 25° C (68° to 77° F) with excursions permitted between 15° to 30° C (59° - 86° F) [See USP Controlled Room Temperature].

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**Mfg. by: Belcher Pharmaceuticals, LLC
Largo, Florida 33777
USA**

L660L-23 D-1512



N 3 62250 66023 7

LOT NO:
EXP DATE:

SODIUM POLYSTYRENE SULFONATE

sodium polystyrene sulfonate powder, for suspension

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:62250-660
Route of Administration	ORAL, RECTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM POLYSTYRENE SULFONATE (UNII: 1699G8679Z) (POLYSTYRENE SULFONIC ACID - UNII:70KO0R01RY)	SODIUM CATION	4.1 meq in 1 g

Product Characteristics

Color	brown (Cream to Light Brown Fine Powder)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62250-660-23	454 g in 1 BOTTLE; Type 0: Not a Combination Product	03/15/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA205727	03/15/2016	

Labeler - Belcher Pharmaceuticals, LLC (965082543)

Establishment

Name	Address	ID/FEI	Business Operations
Belcher Pharmaceuticals, LLC		965082543	manufacture(62250-660)

Revised: 10/2023

Belcher Pharmaceuticals, LLC