

# **SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE - sodium sulfate, potassium sulfate, magnesium sulfate solution, concentrate**

## **Affordable Pharmaceuticals, LLC**

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### **HIGHLIGHTS OF PRESCRIBING INFORMATION**

**These highlights do not include all the information needed to use SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE ORAL SOLUTION safely and effectively. See full prescribing information for SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE ORAL SOLUTION**

### **SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE ORAL SOLUTION**

**Initial U.S. Approval: 2010**

#### **RECENT MAJOR CHANGES**

Warnings and Precautions, Hypersensitivity Reactions (5.8) 11/2025

#### **INDICATIONS AND USAGE**

Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution is an osmotic laxative indicated for cleansing of the colon in preparation for colonoscopy in adult and pediatric patients 12 years of age and older. (1)

#### **DOSAGE AND ADMINISTRATION**

##### Preparation and Administration (2.2)

- Must dilute in water prior to ingestion.
- Administration of two bottles of SUPREP Bowel Prep Kit is required for a complete preparation for colonoscopy. One bottle is equivalent to one dose.
- Must consume additional water after each dose.
- Stop consumption of all fluids at least 2 hours before the colonoscopy.

##### Recommended Dosage and Administration

- Split-Dose (two-day) regimen consists of two doses of SUPREP Bowel Prep Kit: first dose during the evening prior to colonoscopy and second dose the next day, during the morning of colonoscopy. (2.1, 2.3, 2.4)
- Recommended SUPREP Bowel Prep Kit dosage is:
  - **Adults:** Two 6-ounce doses. (2.3)
  - **Pediatric Patients 12 Years of Age and Older:** Two 4.5-ounce doses. (2.4)
- For complete information on preparation before colonoscopy and administration of the dosage regimen, see full prescribing Information. (2.1, 2.2, 2.3, 2.4)

#### **DOSAGE FORMS AND STRENGTHS**

- Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (**for adults**): Two bottles each containing 6 ounces of an oral solution of 17.5 grams sodium sulfate, 3.13 grams potassium sulfate, and 1.6 grams magnesium sulfate. (3)
- Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (**for pediatric patients 12 years of age and older**): Two bottles each containing 4.5-ounces of an oral solution of 13.13 grams sodium sulfate, 2.35 grams potassium sulfate, and 1.2 grams magnesium sulfate. (3)

#### **CONTRAINDICATIONS**

- Gastrointestinal obstruction or ileus (4, 5.6)
- Bowel perforation (4, 5.6)
- Toxic colitis or toxic megacolon (4)
- Gastric retention (4)
- Hypersensitivity to any ingredient (4, 5.8)

#### **WARNINGS AND PRECAUTIONS**

- Risk of fluid and electrolyte abnormalities: Encourage adequate hydration, assess concurrent medications, and consider laboratory assessments prior to and after each use. (5.1, 7.1)
- Cardiac arrhythmias: Consider pre-dose and post-colonoscopy ECGs in patients at increased risk. (5.2)
- Seizures: Use caution in patients with a history of seizures and patients at increased risk of seizures,

including medications that lower the seizure threshold. (5.3, 7.1)

- Patients with renal impairment or taking concomitant medications that affect renal function: Use caution, ensure adequate hydration and consider laboratory testing. (5.4, 7.1)
- Suspected GI obstruction or perforation: Rule out the diagnosis before administration. (4, 5.6)
- Patients at risk for aspiration: Observe during administration. (5.7)
- Hypersensitivity reactions, including anaphylaxis: Inform patients to seek immediate medical care if symptoms occur. (5.8)

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#### ADVERSE REACTIONS

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Most common adverse reactions are:

- Adults (>2%): overall discomfort, abdominal distention, abdominal pain, nausea, and vomiting. (6.1)
- Pediatric Patients (>10%): nausea, abdominal pain, abdominal bloating and vomiting. (6.1)

**To report SUSPECTED ADVERSE REACTIONS, contact Affordable Pharmaceuticals, LLC at 1-800-514-5617 or FDA at 1-800-FDA-1088 or [www.fda.gov/medwatch](http://www.fda.gov/medwatch).**

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#### DRUG INTERACTIONS

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Drugs that increase risk of fluid and electrolyte imbalance. (7.1)

**See 17 for PATIENT COUNSELING INFORMATION and Medication Guide.**

**Revised: 11/2025**

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## **FULL PRESCRIBING INFORMATION**

### **1 INDICATIONS AND USAGE**

Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution is indicated for cleansing of the colon as a preparation for colonoscopy in adult and pediatric patients 12 years of age and older.

### **2 DOSAGE AND ADMINISTRATION**

#### **2.1 Dosage and Administration Overview**

Administration of two bottles of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution and additional water is required for a complete preparation for colonoscopy. One bottle of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution is equivalent to one dose. Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution is supplied in two dosage strengths [see *Dosage Forms and Strengths (3)*]. The recommended dosage is:

- Adults: Two 6-ounce doses [see *Dosage and Administration (2.3)*].
- Pediatric patients 12 years of age and older: Two 4.5-ounce doses [see *Dosage and Administration (2.4)*].

#### **2.2 Important Preparation and Administration Instructions**

- Correct fluid and electrolyte abnormalities before treatment with Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution [see *Warnings and Precautions (5.1)*]
- Must dilute Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution in

water before ingestion.

- Must consume additional water after each dose of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.
- On the day before colonoscopy, consume only a light breakfast or clear liquids (e.g., water, strained fruit juice without pulp, lemonade, plain coffee or tea, chicken broth gelatin dessert without fruit). On the day of the colonoscopy only consume clear liquids up to two hours prior to colonoscopy.
- Do not eat solid food or drink milk or eat or drink anything colored red or purple.
- Do not drink alcohol.
- Do not take other laxatives while taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.
- Do not take oral medications within one hour of starting each dose of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.
- If taking tetracycline or fluoroquinolone antibiotics, iron, digoxin, chlorpromazine, or penicillamine, take these medications at least 2 hours before and not less than 6 hours after administration of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution [see *Drug Interactions (7.2)*].
- Stop consumption of all fluids at least 2 hours prior to the colonoscopy.

### 2.3 Recommended Dosage and Administration for Adults

The recommended Split-Dose (two-day) regimen for **adults** consists of two 6-ounce doses of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution: the first dose during the evening prior to colonoscopy and the second dose the next day, during the morning of the colonoscopy.

Each dose consists of one bottle of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution with additional water. The total volume of liquid required for colon cleansing (using two bottles) is 3 quarts. The following are recommended dosage and administration instructions for adults:

Dose 1 - On the Day Prior to Colonoscopy:

- May consume a light breakfast, or only clear liquids (no solid food).
- In the evening before the procedure, pour the contents of one bottle of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution into the mixing container provided.
- Add cool drinking water to the 16-ounce fill line on the container, mix, and drink the entire amount.
- Drink two additional containers filled with water to the 16-ounce fill line over the next hour.

Dose 2 - Day of Colonoscopy:

- Continue to consume only clear liquids.
- In the morning (10 to 12 hours after the evening dose) on the day of the procedure,

pour the contents of the second bottle of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution into the mixing container provided.

- Add cool drinking water to the 16-ounce fill line on the container, mix, and drink the entire amount.
- Drink two additional containers filled with water to the 16-ounce fill line over the next hour.
- Complete all solution of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution and required water at least two hours prior to colonoscopy.

## **2.4 Recommended Dosage and Administration for Pediatric Patients 12 Years of Age and Older**

The recommended Split-Dose (two-day) regimen for pediatric patients 12 years of age and older consists of two 4.5-ounce doses of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution: the first dose during the evening prior to colonoscopy and the second dose the next day, during the morning of the colonoscopy.

Each dose consists of one bottle of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution with additional water. The total volume of liquid required for colon cleansing (using two bottles) is 2.25 quarts. The following are recommended dosage and administration instructions for pediatric patients 12 years of age and older and/or their caregivers:

Dose 1 – On the Day Prior to Colonoscopy:

- May consume a light breakfast, or only clear liquids (no solid food).
- In the evening before the procedure, pour the contents of one bottle of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution into the mixing container provided.
- Add cool drinking water to the 12-ounce fill line on the container, mix, and drink the entire amount.
- Drink two additional containers filled with water to the 12-ounce fill line over the next hour.

Dose 2 – Day of Colonoscopy:

- Continue to consume only clear liquids.
- In the morning (10 to 12 hours after the evening dose) on the day of the procedure, pour the contents of the second bottle of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution into the mixing container provided.
- Add cool drinking water to the 12-ounce fill line on the container, mix, and drink the entire amount.
- Drink two additional containers filled with water to the 12-ounce fill line over the next hour.

- Complete all solution of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution and required water at least two hours prior to colonoscopy.

### 3 DOSAGE FORMS AND STRENGTHS

- Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (**for adults**): Two bottles each containing 6 ounces of an oral solution of 17.5 grams sodium sulfate, 3.13 grams potassium sulfate, and 1.6 grams magnesium sulfate as a clear to slightly hazy liquid.
- Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (**for pediatric patients 12 years of age and older**): Two bottles each containing 4.5 ounces of an oral solution of 13.13 grams sodium sulfate, 2.35 grams potassium sulfate, and 1.2 grams magnesium sulfate as a clear to slightly hazy liquid.

When diluted as directed, the solution is clear and colorless.

### 4 CONTRAINDICATIONS

Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution is contraindicated in the following conditions:

- Gastrointestinal obstruction or ileus [*see Warnings and Precautions (5.6)*]
- Bowel perforation [*see Warnings and Precaution (5.6)*]
- Toxic colitis or toxic megacolon
- Gastric retention
- Hypersensitivity to any of the ingredients in Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution [*see Warnings and Precautions (5.8) and Description (11)*]

### 5 WARNINGS AND PRECAUTIONS

#### 5.1 Serious Fluid and Serum Chemistry Abnormalities

Advise all patients to hydrate adequately before, during, and after the use of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution. If a patient develops significant vomiting or signs of dehydration after taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution, consider performing post-colonoscopy lab tests (electrolytes, creatinine, and BUN).

Fluid and electrolyte disturbances can lead to serious adverse events including cardiac arrhythmias, seizures and renal impairment. Correct fluid and electrolyte abnormalities before treatment with Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution. Use Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution with caution in patients with conditions, or who are using medications, that increase the risk for fluid and electrolyte disturbances or may increase the risk of adverse events of seizure, arrhythmias, and renal impairment [*see Drug Interactions (7.1)*].

Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution can cause temporary elevations in uric acid [see *Adverse Reactions (6.1)*]. Uric acid fluctuations in patients with gout may precipitate an acute flare. The potential for uric acid elevation should be considered before administering Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution to patients with gout or other disorders of uric acid metabolism.

## **5.2 Cardiac Arrhythmias**

There have been rare reports of serious arrhythmias associated with the use of ionic osmotic laxative products for bowel preparation. Use caution when prescribing Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution for patients at increased risk of arrhythmias (e.g., patients with a history of prolonged QT, uncontrolled arrhythmias, recent myocardial infarction, unstable angina, congestive heart failure, or cardiomyopathy). Consider pre-dose and post-colonoscopy ECGs in patients at increased risk of serious cardiac arrhythmias.

## **5.3 Seizures**

There have been reports of generalized tonic-clonic seizures and/or loss of consciousness associated with use of bowel preparation products in patients with no prior history of seizures. The seizure cases were associated with electrolyte abnormalities (e.g., hyponatremia, hypokalemia, hypocalcemia, and hypomagnesemia) and low serum osmolality. The neurologic abnormalities resolved with correction of fluid and electrolyte abnormalities.

Use caution when prescribing Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution for patients with a history of seizures and in patients at increased risk of seizure, such as patients taking medications that lower the seizure threshold (e.g., tricyclic antidepressants), patients withdrawing from alcohol or benzodiazepines, or patients with known or suspected hyponatremia [see *Drug Interactions (7.1)*].

## **5.4 Use in Patients with Risk of Renal Injury**

Use Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution with caution in patients with impaired renal function or patients taking concomitant medications that may affect renal function (such as diuretics, angiotensin converting enzyme inhibitors, angiotensin receptor blockers, or non-steroidal anti-inflammatory drugs) [see *Drug Interactions (7.1)*]. These patients may be at risk for renal injury. Advise these patients of the importance of adequate hydration with Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution and consider performing baseline and post-colonoscopy laboratory tests (electrolytes, creatinine, and BUN) in these patients [see *Use in Specific Populations (8.6)*].

## **5.5 Colonic Mucosal Ulcerations and Ischemic Colitis**

Osmotic laxative products may produce colonic mucosal aphthous ulcerations, and there have been reports of more serious cases of ischemic colitis requiring hospitalization. Concurrent use of stimulant laxatives and Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution may increase these risks [see *Drug Interactions (7.3)*]. Consider the potential for mucosal ulcerations resulting from the bowel preparation when interpreting colonoscopy findings in patients with known or

suspect inflammatory bowel disease (IBD).

## 5.6 Use in Patients with Significant Gastrointestinal Disease

If gastrointestinal obstruction or perforation is suspected, perform appropriate diagnostic studies to rule out these conditions before administering Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution [see *Contraindications (4)*].

Use with caution in patients with severe active ulcerative colitis.

## 5.7 Aspiration

Patients with impaired gag reflex or other swallowing abnormalities are at risk for regurgitation or aspiration of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution. Observe these patients during administration of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution. Use with caution in these patients.

## 5.8 Hypersensitivity Reactions

Serious hypersensitivity reactions, including anaphylaxis, angioedema, dyspnea, rash, pruritus and urticaria have been reported with Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate [see *Adverse Reactions (6.2)*]. Inform patients of the signs and symptoms of anaphylaxis and instruct them to seek immediate medical care should signs and symptoms occur.

## 6 ADVERSE REACTIONS

The following important adverse reactions for bowel preparations are described elsewhere in the labeling:

- Serious Fluid and Serum Chemistry Abnormalities [see *Warnings and Precautions (5.1)*]
- Cardiac Arrhythmias [see *Warnings and Precautions (5.2)*]
- Seizures [see *Warnings and Precautions (5.3)*]
- Use in Patients with Risk of Renal Injury [see *Warnings and Precautions (5.4)*]
- Colonic Mucosal Ulceration and Ischemic Colitis [see *Warnings and Precautions (5.5)*]
- Patients with Significant Gastrointestinal Disease [see *Warnings and Precautions (5.6)*]
- Aspiration [see *Warnings and Precautions (5.7)*]
- Hypersensitivity Reactions [see *Warnings and Precautions (5.8)*]

### 6.1 Clinical Trials Experience

Because clinical studies are conducted under widely varying conditions, adverse reaction rates observed in the clinical studies of a drug cannot be directly compared to rates in clinical studies of another drug and may not reflect the rates observed in practice.

#### Adults

The safety of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution was evaluated in a multi-center, randomized, active controlled trial in 379 adult patients

undergoing colonoscopy [see *Clinical Studies (14)*].

### *Most Common Adverse Reactions*

Table 1 shows the most common adverse reactions reported in at least 2% of patients receiving Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution or the control (a bowel prep containing polyethylene glycol and electrolytes (PEG + E)) administered in split-dose (2-day) regimens.

**Table 1: Common Adverse Reactions\* in Adult Patients Undergoing Colonoscopy in a Randomized, Active Controlled Trial**

| Symptom              | Split-Dose (2-Day) Regimen  |                               |
|----------------------|---|-------------------------------|
|                      | Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution<br>%<br>N=190 | PEG + E product<br>%<br>N=189 |
| Overall Discomfort   | 54  | 67                            |
| Abdominal Distension | 40  | 52                            |
| Abdominal Pain       | 36  | 43                            |
| Nausea               | 36  | 33                            |
| Vomiting             | 8   | 4                             |

\* reported in at least 2% of patients

### *Laboratory Abnormalities*

Table 2 shows the most common laboratory abnormalities (at least 10% in either treatment group and more than 2% difference between groups) for patients who developed new abnormalities of important electrolytes and uric acid after completing the bowel preparation with either Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution or PEG+E administered as a split-dose (2-day) regimen.

**Table 2: Adult Patients with Normal Baseline Serum Chemistry with A Shift to an Abnormal Value While on the Split-Dose (2-Day) Regimen <sup>1</sup>**

|                         |   | Day of Colonoscopy<br>N (%) <sup>2</sup> | Day 30<br>N (%)<br><sup>2</sup> |
|-------------------------|---|--|---------------------------------|
| Bicarbonate (low)       | Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution | 20 (13)                                  | 7 (4)                           |
|                         | PEG + Electrolytes  | 24 (15)                                  | 4 (3)                           |
| Bilirubin, total (high) | Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution | 14 (9)                                   | 0 (0)                           |
|                         | PEG + Electrolytes  | 20 (12)                                  | 3 (2)                           |
| BUN (high)              | Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution | 2 (2)                                    | 14 (11)                         |
|                         | PEG + Electrolytes  | 4 (3)                                    | 19                              |

|                   |   |         |         |
|-------------------|---|---------|---------|
|                   |   |         | (15)    |
| Calcium (high)    | Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution | 16 (10) | 8 (5)   |
|                   | PEG + Electrolytes  | 6 (4)   | 6 (4)   |
| Chloride (high)   | Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution | 4 (2)   | 6 (4)   |
|                   | PEG + Electrolytes  | 20 (12) | 6 (4)   |
| Osmolality (high) | Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution | 8 (6)   | NA      |
|                   | PEG + Electrolytes  | 19 (13) | NA      |
| Uric acid (high)  | Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution | 27 (24) | 13 (12) |
|                   | PEG + Electrolytes  | 12 (10) | 20 (17) |

<sup>1</sup> The study was not designed to support comparative claims for the laboratory abnormalities reported in this table.

<sup>2</sup> Percent (n/N) of patients where N=number of patients with normal baseline who had abnormal values at the timepoint(s) of interest.

#### *Less Common Adverse Reactions*

AV Block (1 case) and CK increase.

#### *Adverse Reactions with Unapproved Use*

In another study of 408 adult patients, higher rates of the following adverse reactions and laboratory abnormalities were reported in patients treated with Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution as an evening-only (1-day) regimen compared to the split-dose (2-day) regimen.

- overall discomfort, abdominal distention, nausea, and vomiting
- total bilirubin (high), BUN (high), creatinine (high), osmolality (high), potassium (high) and uric acid (high)

Administration of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution in an evening-only (1-day) dosing regimen is not recommended.

#### Pediatrics 12 Years to 16 Years of Age

The safety of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution was evaluated in a single dose-ranging clinical trial of 89 pediatric patients aged 12 years to 16 years [see *Clinical Studies (14)*]. In 26 pediatric patients who received Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 4.5-ounce doses), the most common adverse reactions (> 10%) were nausea, abdominal pain, abdominal bloating, and vomiting.

## **6.2 Postmarketing Experience**

The following adverse reactions have been identified during post-approval use of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate. Because these reactions are 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: anaphylaxis, angioedema, dyspnea, rash, pruritus, urticaria [see Warning and Precautions (5.8)]*

## **7 DRUG INTERACTIONS**

### **7.1 Drugs That May Increase Risks of Fluid and Electrolyte Abnormalities**

Use caution when prescribing Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution to patients taking medications that increase the risk of fluid and electrolyte disturbances or may increase the risk of adverse events of seizure, arrhythmias, and prolonged QT in the setting of fluid and electrolyte abnormalities *[see Warnings and Precautions (5.1, 5.2, 5.3, 5.4)]*.

### **7.2 Potential for Reduced Drug Absorption**

Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution can reduce the absorption of other co-administered drugs *[see Dosage and Administration (2.1)]*.

- Administer oral medications at least one hour before starting each dose of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.
- Administer tetracycline and fluoroquinolone antibiotics, iron, digoxin, chlorpromazine, and penicillamine at least 2 hours before and not less than 6 hours after administration of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution to avoid chelation with magnesium.

### **7.3 Stimulant Laxatives**

Concurrent use of stimulant laxatives and Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution may increase the risk of mucosal ulceration or ischemic colitis. Avoid use of stimulant laxatives (e.g., bisacodyl, sodium picosulfate) while taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution *[see Warnings and Precautions (5.5)]*.

## **8 USE IN SPECIFIC POPULATIONS**

### **8.1 Pregnancy**

#### Risk Summary

There are no available data on Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution use in pregnant women to evaluate for a drug-associated risk of major birth defects, miscarriage, or adverse maternal or fetal outcomes. Animal reproductive studies have not been conducted with Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.

The estimated background risk of major birth defects and miscarriage for the indicated population is unknown. All pregnancies have a background risk of birth defect, loss, or other adverse outcomes. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2% to 4% and 15% to 20%, respectively.

## **8.2 Lactation**

### Risk Summary

There are no data available data on the presence of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution in human or animal milk, the effects on the breastfed child, or the effects on milk production. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution and any potential adverse effects on the breastfed child from Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution or from the underlying maternal condition.

## **8.4 Pediatric Use**

The safety and effectiveness of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 4.5-ounce doses) have been established for cleansing of the colon as a preparation for colonoscopy in pediatric patients 12 years of age and older. Use of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution in this age group is supported by evidence from an adequate and well-controlled trial of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution in adults and a single, dose-ranging, controlled trial in 89 pediatric patients 12 years to 16 years of age [see *Clinical Studies (14)*]. In the pediatric trial, Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 6-ounce doses) did not demonstrate additional treatment benefit and more patients reported gastrointestinal adverse reactions compared to Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 4.5-ounce doses). Therefore, Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 6-ounce doses) is not recommended for pediatric patients 12 years of age and older [see *Dosage and Administration (2.3)*]. The safety profile of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 4.5-ounce doses) in this pediatric population was similar to that seen in adults [see *Adverse Reactions (6.1)*].

The safety and effectiveness of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution in pediatric patients less than 12 years of age have not been established.

## **8.5 Geriatric Use**

Of the 375 patients who received Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution in clinical trials, 94 (25%) were 65 years of age or older, and 25 (7%) were 75 years of age or older. No overall differences in safety or effectiveness of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution, administered as the recommended split-dose (2-day) regimen, were observed between geriatric patients and younger patients. Geriatric patients reported more vomiting when Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution was given as a one-day preparation (not a recommended regimen). Elderly patients are more likely to have decreased hepatic, renal or cardiac function and may be more susceptible to adverse reactions resulting from fluid and electrolyte abnormalities [see *Warnings and Precautions (5.1)*].

## **8.6 Renal Impairment**

Use Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution with caution in patients with renal impairment or patients taking concomitant medications that may affect renal function. These patients may be at risk for renal injury. Advise these patients of the importance of adequate hydration before, during and after use of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution and consider performing baseline and post-colonoscopy laboratory tests (electrolytes, creatinine, and BUN) in these patients [see *Warnings and Precautions (5.4)*].

## 10 OVERDOSAGE

Overdosage of more than the recommended dose of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution may lead to severe electrolyte disturbances, as well as dehydration and hypovolemia, with signs and symptoms of these disturbances. [see *Warnings and Precautions (5.1, 5.2, 5.3)*]. Monitor for fluid and electrolyte disturbances and treat symptomatically.

## 11 DESCRIPTION

Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (**for adults**) is an osmotic laxative and is provided as two bottles each containing 6 ounces of solution. Each bottle contains: 17.5 grams sodium sulfate, 3.13 grams potassium sulfate, and 1.6 grams magnesium sulfate. Inactive ingredients include: citric acid USP, flavoring ingredients, malic acid FCC, sodium benzoate, NF, sucralose, purified water, USP.

Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (**for pediatric patients 12 years of age and older**) is an osmotic laxative and is provided as two bottles each containing 4.5 ounces of solution. Each bottle contains: 13.13 grams sodium sulfate, 2.35 grams potassium sulfate, and 1.2 grams magnesium sulfate. Inactive ingredients include: citric acid USP, flavoring ingredients, malic acid FCC, sodium benzoate, NF, sucralose, purified water, USP.

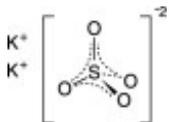
### Sodium Sulfate, USP

The chemical name is  $\text{Na}_2\text{SO}_4$ . The average Molecular Weight is 142.04. The structural formula is:



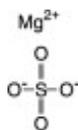
### Potassium Sulfate, FCC, purified

The chemical name is  $\text{K}_2\text{SO}_4$ . The average Molecular Weight is 174.26. The structural formula is:



### Magnesium Sulfate, USP

The chemical name is  $\text{MgSO}_4$ . The average Molecular Weight: 120.37. The structural formula is:



Each Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution also contains a polypropylene mixing container.

## 12 CLINICAL PHARMACOLOGY

### 12.1 Mechanism of Action

Sulfate salts provide sulfate anions, which are poorly absorbed. The osmotic effect of unabsorbed sulfate anions and the associated cations causes water to be retained within the gastrointestinal tract.

### 12.2 Pharmacodynamics

No formal pharmacodynamic studies have been conducted with Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.

### 12.3 Pharmacokinetics

#### Absorption and Elimination

After administration of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution in six healthy subjects, the time at which serum sulfate reached its highest point ( $T_{\text{max}}$ ) was approximately 17 hours after the first dose or approximately 5 hours after the second dose, and then declined with a half-life of 8.5 hours.

#### *Excretion*

Fecal excretion was the primary route of sulfate elimination.

#### Specific Populations

##### *Patients with Renal Impairment*

The disposition of sulfate after ingestion of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution was studied in patients (N=6) with moderate renal impairment (creatinine clearance of 30 to 49 mL/min). In patients with moderate renal impairment, mean AUC was 54% higher and mean  $C_{\text{max}}$  was 44% higher, than healthy subjects.

The mean sulfate concentrations in healthy subjects and in patients with moderate renal impairment returned to their respective baselines by Day 6 after dose initiation. Urinary excretion of sulfate over 30 hours after the first dose was approximately 16% lower in patients with moderate renal impairment than in healthy subjects. These differences are not considered clinically meaningful.

##### *Patients with Hepatic Impairment*

The disposition of sulfate after ingestion of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution was studied in patients (N=6) with mild to moderate hepatic impairment (Child-Pugh grades A and B). Systemic exposure of serum sulfate (AUC and  $C_{\text{max}}$ ) was similar between healthy subjects and patients with hepatic

impairment. The mean sulfate concentrations in healthy subjects and in patients with mild to moderate hepatic impairment returned to their respective baselines by Day 6 after dose initiation. Urinary excretion of sulfate over 30 hours after the first dose was similar between patients with hepatic impairment and healthy subjects.

## **13 NONCLINICAL TOXICOLOGY**

### **13.2 Animal Toxicology and/or Pharmacology**

The sulfate salts of sodium, potassium, and magnesium contained in Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution were administered orally (gavage) to rats and dogs up to 28 days up to a maximum daily dose of 5 grams/kg/day (approximately 0.9 and 3 times for rats and dogs, respectively, the recommended human dose of 44 grams/day or 0.89 grams/kg based on the body surface area). In rats, the sulfate salts caused diarrhea and electrolyte and metabolic changes, including hypochloremia, hypokalemia, hyponatremia, lower serum osmolality, and high serum bicarbonate. Significant renal changes included increased fractional sodium excretion, increased urinary sodium and potassium excretion, and alkaline urine in both males and females. In addition, creatinine clearance was significantly decreased in females at the highest dose. No microscopic renal changes were seen. In dogs, the sulfate salts caused emesis, excessive salivation, excessive drinking of water, and abnormal excreta (soft and/or mucoid feces and/or diarrhea) and increased urine pH and sodium excretion.

## **14 CLINICAL STUDIES**

### Adults

The colon cleansing efficacy of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution was evaluated in a randomized, single-blind, active-controlled, multicenter study in adult patients scheduled to have a colonoscopy. There were 363 adult patients included in the efficacy analysis. Patients ranged in age from 20 to 84 years (mean age 55 years) and 54% were female. Race distribution was 86% Caucasian, 9% African-American, and 5% other.

Patients were randomized to one of the following two colon preparation regimens: Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution or a marketed polyethylene glycol (PEG) plus electrolytes bowel preparation. In the Study Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution was administered as a split-dose (two-day) regimen. The PEG bowel prep was also given as a split-dose preparation according to its labeled instructions. Patients receiving Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution were limited to a light breakfast followed by clear liquids on the day prior to the day of colonoscopy; patients receiving the PEG bowel prep were allowed to have a normal breakfast and a light lunch, followed by clear liquids.

The primary efficacy endpoint was the proportion of patients with successful colon cleansing as assessed by the colonoscopists, who were not informed about the type of preparation received, as shown in Table 3. In the study, no clinically or statistically significant differences were seen between the group treated with Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution and the group treated with the

PEG bowel prep.

**Table 3: Proportion of Adult Patients with Successful Colon Cleansing Response Rates**

| <b>Treatment Group</b>   | <b>Regimen</b> | <b>N</b> | <b>Responders<sup>1</sup><br/>%<br/>(95% C. I.)</b> | <b>Sodium Sulfate,<br/>Potassium Sulfate<br/>and Magnesium<br/>Sulfate Oral<br/>Solution-PEG<br/>Difference<br/>(95% CI)</b> |
|--|----------------|----------|---|--|
| Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (with light breakfast) | Split-Dose     | 180      | <b>97%</b><br>(94%, 99%)                            | <b>2%</b> <sup>2</sup><br><br>(-2%, 5%)  |
| PEG bowel prep (with normal breakfast & light lunch)   | Split-Dose     | 183      | <b>96%</b><br>(92%, 98%)                            |  |

<sup>1</sup> Responders were patients whose colon preparations were graded excellent (no more than small bits of adherent feces/fluid) or good (small amounts of feces or fluid not interfering with the exam) by the colonoscopist.

<sup>2</sup> Does not equal difference in tabled responder rates due to rounding effects.

#### Pediatric Patients 12 Years to 16 Years of Age

Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution was evaluated for colon cleansing in a randomized, single-blind, multicenter, dose-ranging, active-controlled study in 89 pediatric patients 12 years to 16 years of age. The majority of patients were female (57%), white (78%), and of non-Hispanic or non-Latino ethnicity (91%). The mean age was 14 years. The median body weight was 60 kg (range 32 to 155 kg).

Patients were randomized to Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 6-ounce doses), Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 4.5-ounce doses) or oral PEG solution. Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 6-ounce doses) did not demonstrate additional treatment benefit and more patients reported gastrointestinal adverse reactions compared to Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 4.5-ounce doses); therefore, Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 6-ounce doses) is not recommended for pediatric patients 12 years of age and older [see *Dosage and Administration (2.4)*]. Patients in the Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 4.5-ounce doses) group took the preparation in a “split-dose” regimen, where the first dose was taken the evening before colonoscopy, with the second dose taken the morning of the exam. Patients in the control group took the preparation according to its approved labeling on the evening before colonoscopy.

Patients in the Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 4.5-ounce doses) group were allowed to have a light breakfast on the day before colonoscopy, followed by clear liquids until the colonoscopy is completed the following day. Patients in the control group subjects were permitted only clear liquids on the day prior to colonoscopy until completion of the colonoscopy the following day.

The primary efficacy endpoint was the proportion of patients with successful colon cleansing as assessed by the colonoscopists, who were not informed about the type of preparation received.

The percentage of responders and the associated 95% confidence intervals for the Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 4.5-ounce doses) and Oral PEG solution are shown in Table 4. Efficacy was similar between patients who weighed 65 kg or more (n=12) and those patients who weighed less than 65 kg (n=15) in the Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (two 4.5-ounce doses) arm.

**Table 4: Proportion of Pediatric Patients 12 Years to 16 Years of Age with Successful Colon Cleansing Response Rates**

| <b>Treatment Group</b>   | <b>Regimen</b>        | <b>N</b> | <b>Responders<sup>1</sup><br/>%<br/>(95% C. I.)</b> | <b>Sodium Sulfate,<br/>Potassium Sulfate and<br/>Magnesium Sulfate Oral<br/>Solution-PEG<br/>Difference<br/>(95% CI)</b> |
|--|-----------------------|----------|---|--|
| Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution<br><br>4.5 ounces per dose<br>(with light breakfast) | Split-Dose            | 26       | <b>85%</b><br>(71%, 99%)                            | <b>25%<sup>2</sup></b><br><br>(3%, 47%)  |
| Oral PEG solution<br>(with clear liquids only)   | Evening<br><br>Dosing | 32       | <b>59%</b><br>(42%, 76%)                            |  |

<sup>1</sup> Responders were patients whose colon preparations were graded excellent (no more than small bits of adherent feces/fluid) or good (small amounts of feces or fluid not interfering with the exam) by the colonoscopist.

<sup>2</sup> Does not equal difference in tabled responder rates due to rounding effects.

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

Each Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution **(for adults)** (NDC 10572-012-01) contains:

- Two bottles (NDC 10572-011-01) each containing 6-ounces of an oral solution of 17.5 grams sodium sulfate, 3.13 grams potassium sulfate, and 1.6 grams magnesium sulfate as a clear to slightly hazy liquid. When diluted as directed, the solution is clear and colorless.
- One (1) mixing container with a 16-ounce fill line.

Each Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution **(for pediatric patients 12 years of age and older)** (NDC 10572-112-01) contains:

- Two bottles (NDC 10572-111-01) each containing 4.5-ounces of an oral solution of 13.13 grams sodium sulfate, 2.35 grams potassium sulfate, and 1.2 grams

magnesium sulfate as a clear to slightly hazy liquid. When diluted as directed, the solution is clear and colorless.

- One (1) mixing container with a 12-ounce fill line.

Store at 20° to 25°C (68° to 77°F). Excursions permitted between 15° to 30°C (59° to 86°F). See USP controlled room temperature.

## **17 PATIENT COUNSELING INFORMATION**

Advise the patient and/or caregiver to read the FDA-approved patient labeling (Medication Guide).

Instruct patients or caregivers:

- Must dilute Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution before ingestion.
- Must consume additional water after each dose of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.
- On the day before colonoscopy, consume only a light breakfast or clear liquids (e.g., water, apple or orange juice without pulp, lemonade, coffee, tea, or chicken broth). On the day of the colonoscopy only consume clear liquids up to two hours prior to colonoscopy.
- Two doses of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution are required for a complete preparation for colonoscopy. One bottle of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution is equivalent to one dose.
- Do not take other laxatives while taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.
- Do not eat solid food or drink milk or eat or drink anything colored red or purple.
- Do not drink alcohol.
- Do not take oral medications within one hour of starting each dose of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.
- If taking tetracycline or fluoroquinolone antibiotics, iron, digoxin, chlorpromazine, or penicillamine, take these medications at least 2 hours before and not less than 6 hours after administration of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution [see *Drug Interactions (7.2)*].
- Stop consumption of all fluids at least 2 hours prior to colonoscopy.
- Contact their healthcare provider if they develop significant vomiting or signs of dehydration after taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution or if they experience cardiac arrhythmias or seizures [see *Warnings and Precautions (5.1, 5.2, 5.3)*].
- Seek immediate medical care should signs or symptoms of a hypersensitivity reaction occur [see *Warnings and Precautions (5.8)*].

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Braintree, MA 02185

U.S. Patent 6,946,149

## **MEDICATION GUIDE**

**Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution (so-DEE-um sul-fate, po-TASS ee-um sul-fate and mag-NEE-see-um sul-fate)**

Read and understand this Medication Guide **at least 2 days before** your colonoscopy and again before you start taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.

**What is the most important information I should know about Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution?**

**Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution and other bowel preparations can cause serious side effects, including:**

- **Serious loss of body fluid (dehydration) and changes in blood salts (electrolytes) in your blood. These changes can cause:**
  - o **abnormal heartbeats that can cause death.**
  - o **seizures.** This can happen even if you have never had a seizure.
  - o **kidney problems.**

**Your chance of having fluid loss and changes in body salts with Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution is higher if you:**

- have heart problems
- have kidney problems
- take water pills or non-steroidal anti-inflammatory drugs (NSAIDS)

**Tell your healthcare provider right away if you have any of these symptoms of a loss of too much body fluid (dehydration) while taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution:**

- o vomiting
- o dizziness
- o urinating less often than normal
- o headache

**See "What are the possible side effects of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution?" for more information about side effects.**

**What is Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution?**

Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution is a prescription medicine used by adults and children 12 years of age and older to clean the colon before a colonoscopy. Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution cleans your colon by causing you to have diarrhea. Cleaning your colon helps your healthcare provider see the inside of your colon more clearly during your colonoscopy.

**It is not known if Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution is safe and effective in children under 12 years of age.**

**Do not** take Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution if your healthcare provider has told you that you have:

- a blockage in your intestine (bowel obstruction).
- an opening in the wall of your stomach or intestine (bowel perforation).
- a very dilated intestine (toxic megacolon).

- problems with the emptying of food and fluid from your stomach (gastric retention).
- an allergy to any of the ingredients in Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution. See the end of this Medication Guide for a complete list of ingredients in Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.

**Before taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution, tell your healthcare provider about all of your medical conditions, including if you:**

- have problems with serious loss of body fluid (dehydration) and changes in blood salts (electrolytes).
- have gout.
- have heart problems including an irregular heartbeat, especially a condition called "QT prolongation".
- have a history of seizures or take medicines for seizures.
- are withdrawing from drinking alcohol or from taking benzodiazepines.
- have a low blood salt (sodium) level.
- have kidney problems or take medicines for kidney problems.
- have stomach or bowel problems including ulcerative colitis.
- have problems with swallowing or gastric reflux.
- are pregnant or plan to become pregnant. It is not known if Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution will harm your unborn baby. Talk to your healthcare provider if you are pregnant.
- are breastfeeding or plan to breastfeed. It is not known if Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution passes into your breast milk. You and your healthcare provider should decide if you will take Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution while breastfeeding.

**Tell your healthcare provider about all the medicines you take,** including prescription and over-the-counter medicines, vitamins, and herbal supplements. Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution may affect how other medicines work. Medicines taken by mouth may not be absorbed properly when taken within 1 hour before the start of each dose of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.

**Especially tell your healthcare provider if you take:**

- medicines for blood pressure or heart problems.
- medicines for kidney problems.
- medicines for seizures.
- water pills (diuretics).
- non-steroidal anti-inflammatory medicines (pain medicines).
- medicines for depression or mental health problems.
- laxatives. **Do not** take other laxatives while taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.

The following medicines should be taken at least 2 hours before starting Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution and not less than 6 hours after taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution:

- tetracycline
- fluoroquinolone antibiotics
- iron
- digoxin (Lanoxin)
- chlorpromazine

- penicillamine (Cuprimine, Depen)

Ask your healthcare provider or pharmacist for a list of these medicines if you are not sure if you are taking any of the medicines listed above.

Know the medicines you take. Keep a list of them to show your healthcare provider and pharmacist when you get a new medicine.

### **How should I take Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution?**

**See the Instructions for Use in the Patient Instructions for Use Booklet for dosing instructions. You must read, understand, and follow these instructions to take Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution the right way.**

- Take Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution exactly as your healthcare provider tells you to take it.
- Each bottle of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution must be mixed with water (diluted) before drinking.
- It is important for you to drink the additional prescribed amount of water listed in the Instructions for Use to prevent fluid loss (dehydration).
- One bottle of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution is equal to one dose.
- Two doses of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution are required for complete colonoscopy preparation.
- All people taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution should follow these general instructions starting 1 day **before** your colonoscopy:
  - eat only a light breakfast or clear liquids (for example: water, strained fruit juice without pulp, lemonade, plain coffee, or tea, chicken broth, gelatin dessert without fruit) on the day before your procedure.
  - only drink clear liquids the rest of the day and the next day until 2 hours before your colonoscopy. **Stop** drinking all fluids at least 2 hours before your colonoscopy.
  - after taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution if you have any bloating or feeling like your stomach is upset, wait to take your second dose until your stomach feels better.
- **While taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution, do not:**
  - take any other laxatives.
  - take any medicines by mouth (oral) within 1 hour of starting Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.
  - eat solid foods, drink dairy (such as milk), or drink alcohol while taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution and until after your colonoscopy.
  - eat or drink anything colored red or purple.

**Contact your healthcare provider right away** if after taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution you have severe vomiting, signs of dehydration, changes in consciousness such as feeling confused, delirious or fainting (loss of consciousness) or seizures.

## **What are the possible side effects of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution?**

Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution can cause serious side effects, including:

- See "**What is the most important information I should know about Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution?**"

- **Changes in certain blood tests.** Your healthcare provider may do blood tests after you take Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution to check your blood for changes. Tell your healthcare provider if you have any symptoms of too much fluid loss, including:

- o vomiting
- o nausea
- o bloating
- o stomach area (abdomen) cramping
- o dizziness
- o trouble drinking clear liquid
- o headache
- o urinate less than usual
- o trouble swallowing
- o seizures
- o heart problems
- o worsening gout

- **Ulcers of the bowel or bowel problems (ischemic colitis).** Tell your healthcare provider right away if you have severe stomach-area (abdomen) pain or rectal bleeding.

- **Serious allergic reactions.** Get medical help right away if you have any signs and symptoms of a serious allergic reaction while taking Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution, including:

- o difficulty breathing
- o itching
- o swelling of the face, lips, tongue or throat
- o raised red patches on your skin (hives)
- o skin rash

## **The most common side effects of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution in adults include:**

- o overall discomfort
- o stomach pain
- o vomiting
- o stomach bloating
- o nausea

## **The most common side effects of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution in children 12 to 16 years of age include:**

- o nausea
- o stomach bloating
- o stomach pain
- o vomiting

These are not all the possible side effects of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

## **How should I store Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution?**

- Store Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution at room temperature, between 68°F to 77°F (20°C to 25°C).

**Keep Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution and all medicines out of the reach of children.**

**General information about the safe and effective use of Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution.**

Medicines are sometimes prescribed for purposes other than those listed in a Medication Guide. Do not use Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution for a condition for which it was not prescribed. Do not give Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution to other people, even if they are going to have the same procedure you are. It may harm them. You can ask your pharmacist or healthcare provider for information about Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution that is written for health professionals.

**What are the ingredients in Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution?**

Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution is supplied in two dosage strengths. Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution comes in a carton containing two 6-ounce bottles, along with a 16-ounce polypropylene mixing container. Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Pediatric Oral Solution comes in a carton containing two bottles containing 4.5 ounces of oral solution, along with a 12-ounce polypropylene mixing container.

Each bottle contains:

**Active ingredients:** sodium sulfate, potassium sulfate and magnesium sulfate.

**Inactive ingredients:** citric acid, USP, flavoring ingredients, malic acid, FCC, sodium benzoate, NF, sucralose, purified water, USP.

Distributed by Affordable Pharmaceuticals, LLC  
Braintree, MA 02185, USA

For more information go to [www.affordablepharma.com](http://www.affordablepharma.com) or call 1-800-514-5617.

This Medication Guide has been approved by the U.S. Food and Drug Administration.  
Revised 11/2025

Principal Display Panel – Adult Carton Label

NDC 10572-012-01 U.S. Patent 6,946,149

Dispense the enclosed Medication Guide to each patient.

**Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution**

**For Adults**

(17.5g/3.13g/1.6g) per 6 ounces

**This carton contains:**

2 6-ounce (177 mL) bottles of liquid bowel prep

1 16-ounce mixing container

1 Patient booklet. Booklet includes:

1- Medication Guide

2- Patient Instructions

3- Full Prescribing Information

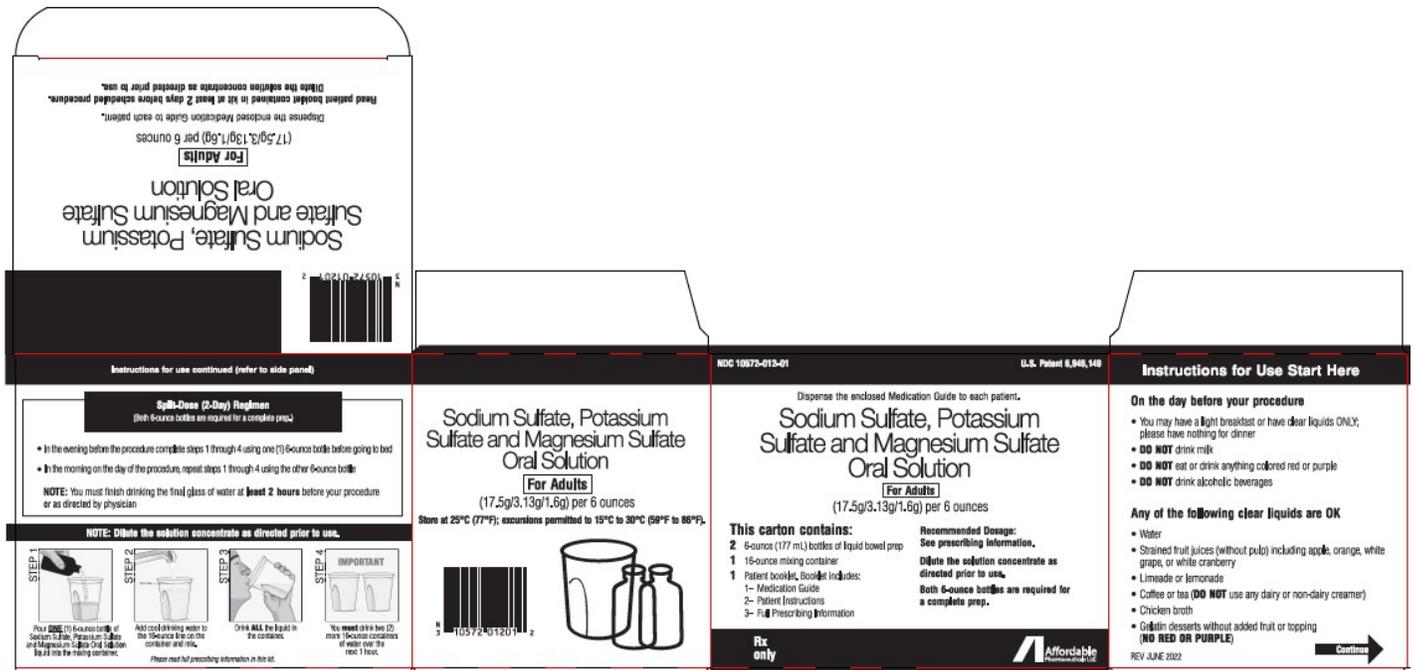
**Recommended Dosage:  
See prescribing information**

**Dilute the solution concentrate as directed prior to use.**

**Both 6-ounce bottles are required for a complete prep.**

Rx only

Affordable Pharmaceuticals, LLC



Principal Display Panel – Adult Bottle Label

**NDC 10572-011-01**

**Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution For Adults**

(17.5g/3.13g/1.6g) per 6 ounces

Dispense the enclosed Medication Guide to each patient.

This bottle contains 6 ounces (177 mL) of liquid bowel prep

**Directions:**

Dilute the solution concentrate prior to use. See enclosed booklet for complete dosage and administration instructions. Both 6-ounce bottles are required for a complete prep.

Keep this and other drugs out of reach of children.

Store at 25°C (77°F); excursions permitted to 15-30°C (59-86°F).

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**NDC 10572-011-01**

**Sodium Sulfate, Potassium  
Sulfate and Magnesium Sulfate  
Oral Solution**

**For Adults**

(17.5g/3.13g/1.6g) per 6 ounces

Dispense the enclosed Medication Guide to each patient.

This bottle contains 6 ounces (177 mL) of liquid bowel prep

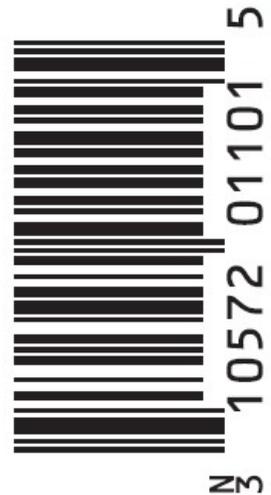
**Directions:**

Dilute the solution concentrate prior to use. See enclosed booklet for complete dosage and administration instructions. Both 6-ounce bottles are required for a complete prep.

Keep this and other drugs out of reach of children.

Store at 25°C (77°F); excursions permitted to 15-30°C (59-86°F).

Distributed by Affordable Pharmaceuticals, LLC, Braintree, MA **Rx only** Rev June '22



Unvarnished  
Area

Principal Display Panel – Pediatric Carton Label

**NDC 10572-112-01 U.S. Patent 6,946,149**

Dispense the enclosed Medication Guide to each patient.

**Sodium Sulfate, Potassium  
Sulfate and Magnesium Sulfate  
Oral Solution**

**For pediatric patients 12 years of age and older**

(13.13g/2.35g/1.2g) per 4.5 ounces

**This carton contains:**

**2** Bottles containing 4.5 ounces  
of liquid bowel prep

**1** 12-ounce mixing container

**1** Patient booklet. Booklet includes:

1- Medication Guide

2- Patient Instructions

3- Full Prescribing Information

**Recommended Dosage:**

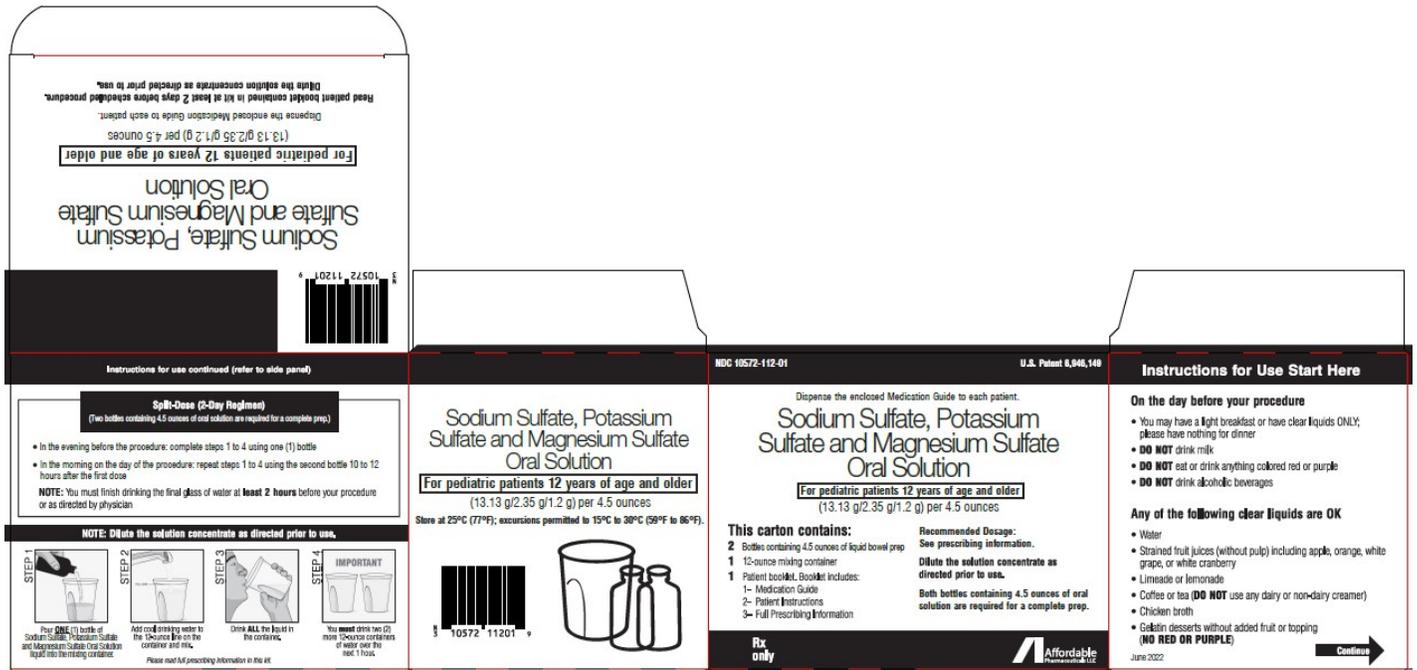
**See prescribing information**

**Dilute the solution concentrate as  
directed prior to use.**

**Both bottles containing 4.5 ounces of oral  
solutions are required for a complete prep.**

**Rx only**

Affordable Pharmaceuticals, LLC



Principal Display Panel – Pediatric Bottle Label

**NDC 10572-111-01**

**Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate) Oral Solution**

**For pediatric patients 12 years of age and older**  
(13.13g/2.35g/1.2g) per 4.5 ounces

Dispense the enclosed Medication Guide to each patient.

This bottle contains 4.5 ounces of liquid bowel prep

Directions:

Dilute the solution concentrate prior to use. See enclosed booklet for complete dosage and administration instructions. Both 6 bottles containing 4.5 ounces of oral solution are required for a complete prep.

Keep this and other drugs out of reach of children.

Store at 25°C (77°F); excursions permitted to 15-30°C (59-86°F).

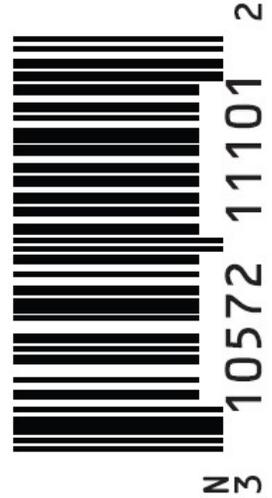
Distributed by Affordable Pharmaceuticals, LLC, Braintree, MA **Rx only** June 2022

NDC 10572-111-01

**Sodium Sulfate, Potassium Sulfate and Magnesium Sulfate Oral Solution**

**For pediatric patients 12 years of age and older**

(13.13 g/2.35 g/1.2 g) per 4.5 ounces



Dispense the enclosed Medication Guide to each patient.

This bottle contains 4.5 ounces of liquid bowel prep

**Directions:**

Dilute the solution concentrate prior to use. See enclosed booklet for complete dosage and administration instructions. Both bottles containing 4.5 ounces of oral solution are required for a complete prep.

Keep this and other drugs out of reach of children.

Store at 25°C (77°F); excursions permitted to 15°C to 30°C (59°F to 86°F).

Distributed by Affordable Pharmaceuticals, LLC, Braintree, MA **Rx only** June 2022

Unvarnished Area

**SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE**

sodium sulfate, potassium sulfate, magnesium sulfate solution, concentrate

**Product Information**

|                                |                         |                           |               |
|--------------------------------|-------------------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN PRESCRIPTION DRUG | <b>Item Code (Source)</b> | NDC:10572-012 |
| <b>Route of Administration</b> | ORAL                    |                           |               |

**Active Ingredient/Active Moiety**

| Ingredient Name  | Basis of Strength                   | Strength          |
|--|-------------------------------------|-------------------|
| <b>SODIUM SULFATE</b> (UNII: 0YPR65R21J) (SODIUM CATION - UNII:LYR4M0NH37)                         | SODIUM SULFATE                      | 17.5 g<br>in 1 mL |
| <b>POTASSIUM SULFATE</b> (UNII: 1K573LC5TV) (POTASSIUM CATION - UNII:295O53K152)                   | POTASSIUM SULFATE                   | 3.13 g<br>in 1 mL |
| <b>MAGNESIUM SULFATE, UNSPECIFIED FORM</b> (UNII: DE08037SAB) (MAGNESIUM CATION - UNII:T6V3LHY838) | MAGNESIUM SULFATE, UNSPECIFIED FORM | 1.6 g<br>in 1 mL  |

**Inactive Ingredients**

| Ingredient Name                           | Strength |
|---|----------|
| <b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU) |          |
| <b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)       |          |
| <b>MALIC ACID</b> (UNII: 817L1N4CKP)      |          |

CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)

### Product Characteristics

|          |               |              |  |
|----------|---------------|--------------|--|
| Color    |               | Score        |  |
| Shape    |               | Size         |  |
| Flavor   | BERRY (BERRY) | Imprint Code |  |
| Contains |               |              |  |

### Packaging

| # | Item Code        | Package Description  | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:10572-012-01 | 2 in 1 CARTON  | 08/05/2010           |                    |
| 1 |                  | 177.4 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product |                      |                    |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| NDA                | NDA022372                                | 08/05/2010           |                    |

## SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE

sodium sulfate, potassium sulfate, magnesium sulfate solution, concentrate

### Product Information

|                         |                         |                    |               |
|-------------------------|-------------------------|--------------------|---------------|
| Product Type            | HUMAN PRESCRIPTION DRUG | Item Code (Source) | NDC:10572-112 |
| Route of Administration | ORAL                    |                    |               |

### Active Ingredient/Active Moiety

| Ingredient Name  | Basis of Strength                   | Strength               |
|--|-------------------------------------|------------------------|
| <b>SODIUM SULFATE</b> (UNII: 0YPR65R21J) (SODIUM CATION - UNII:LYR4MONH37)                         | SODIUM SULFATE                      | 13.13 g<br>in 133.1 mL |
| <b>POTASSIUM SULFATE</b> (UNII: 1K573LC5TV) (POTASSIUM CATION - UNII:295053K152)                   | POTASSIUM SULFATE                   | 2.35 g<br>in 133.1 mL  |
| <b>MAGNESIUM SULFATE, UNSPECIFIED FORM</b> (UNII: DE08037SAB) (MAGNESIUM CATION - UNII:T6V3LHY838) | MAGNESIUM SULFATE, UNSPECIFIED FORM | 1.2 g<br>in 133.1 mL   |

### Inactive Ingredients

| Ingredient Name                           | Strength |
|---|----------|
| <b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU) |          |
| <b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)       |          |

|   |  |
|---|--|
| <b>MALIC ACID</b> (UNII: 817L1N4CKP)              |  |
| <b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP) |  |

### Product Characteristics

|                 |               |                     |  |
|-----------------|---------------|---------------------|--|
| <b>Color</b>    |               | <b>Score</b>        |  |
| <b>Shape</b>    |               | <b>Size</b>         |  |
| <b>Flavor</b>   | BERRY (BERRY) | <b>Imprint Code</b> |  |
| <b>Contains</b> |               |                     |  |

### Packaging

| # | Item Code        | Package Description  | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:10572-112-01 | 2 in 1 CARTON  | 01/05/2021           |                    |
| 1 |                  | 133.1 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product |                      |                    |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| NDA                | NDA022372                                | 08/05/2010           |                    |

**Labeler** - Affordable Pharmaceuticals, LLC (182815311)

Revised: 11/2025

Affordable Pharmaceuticals, LLC