

## Product Licence

**Natural Product Number:** 80027202

**Current Status:** Discontinued

**Brand Name:** Phosphate Novartis

**Licence holder:** Novartis Pharmaceuticals Canada Inc.

**Dosage form:** Tablet

**Recommended route of administration:** Oral

**Recommended Dose:**

Adults: Take 1 tablet 2 times daily. Take tablets only when dissolved in water. Use preferably 250 mL of diluent. Use only on the advice of a physician. Adjust dose according to patient response.

Children: (1-17 years): Take 0.5 tablets 2 times daily with meals.

**Recommended use or purpose:** Hypercalciuria, electrolyte replenisher.

**Risk Information:**

**Cautions and Warnings**

Use carefully in patients with cardiac disease treated with digitalis and in conditions where high potassium concentration may be encountered such as: Adrenal insufficiency, acute dehydration, severe renal insufficiency, in conditions such as in severe burns, where tissue breakdown is expected. Because of the sodium content of Phosphate-Novartis, use carefully in the presence of cardiac failure, liver cirrhosis or severe hepatic disease, edema, hypernatremia, hypertension and toxemia of pregnancy.

**Contra-Indications**

Do not use this medication if hyperphosphatemia is present or in the presence of severe impairment of renal function (less than 30% of normal).

**Known Adverse Reactions**

The following reactions have been reported: Nausea, vomiting, stomach pain, laxative effect or diarrhea, and less frequently: Fluid retention associated with swelling of feet and/or weight gain, hyperkalemia associated with confusion, tiredness or weakness; irregular or slow heart beat; numbness or tingling around lips, hands or feet; unexplained anxiety; breathing problems, hypernatremia associated with confusion; tiredness or weakness; convulsions; decrease in urine volume or in frequency of urine; fast heartbeat; headache or dizziness; increased thirst; hyperphosphatemia or hypocalcemic tetany associated with muscle cramps; numbness, tingling, pain, or weakness in hands or feet, shortness of breath or troubled breathing.

**List of Medicinal Ingredients:**

Medicinal Ingredients	Quantity (QTY)	Extract	Potency
Phosphorus	500 mg		
Potassium	123 mg		
Sodium	469 mg		

**List of Non-Medicinal Ingredients:**

- 1,2-Benzisothiazol-3(2H)-one,1,1-dioxide,sodium salt
- Citric Acid Anhydrous
- Orange flavour
- Polyethylene Glycol
- Potassium Bicarbonate
- Sodium Phosphate, Monobasic
- Sodium bicarbonate
- Sucrose

**Date of Licensing:** 2011-09-02