K PHOS ORIGINAL - potassium phosphate, monobasic tablet, soluble
Beach Products, Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

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K-PHOS® ORIGINAL
Potassium Acid Phosphate
Rx ONLY

DESCRIPTION
Each tablet contains potassium acid phosphate 500 mg. Each tablet yields approximately 114 mg of phosphorus and 144 mg of potassium or 3.7 mEq.

Inactive ingredients: Magnesium stearate, microcrystalline cellulose, silicon dioxide, starch, stearic acid.

ACTIONS
K-PHOS® ORIGINAL is a highly effective sodium-free urinary acidifier.

INDICATIONS AND USAGE
For use in patients with elevated urinary pH. K-PHOS® ORIGINAL helps keep calcium soluble and reduces odor and rash caused by ammoniacal urine. Also, by acidifying the urine, it increases the antibacterial activity of methenamine mandelate and methenamine hippurate.

CONTRAINDICATIONS
This product is contraindicated in patients with infected phosphate stones; in patients with severely impaired renal function (less than 30% of normal) and in the presence of hyperphosphatemia and hyperkalemia.

PRECAUTIONS
General
This product contains potassium and should be used with caution if regulation of this element is desired. Occasionally, some individuals may experience a mild laxative effect during the first few days of phosphate therapy. If laxation persists to an unpleasant degree, reduce the daily dosage until this effect subsides or, if necessary, discontinue the use of this product.

Caution should be exercised when prescribing this product in the following conditions: Cardiac disease (particularly in digitalized patients); severe adrenal insufficiency (Addison's disease); acute dehydration; severe renal insufficiency or chronic renal disease; extensive tissue breakdown (such as severe burns); myotonia congenita; hypoparathyroidism; and acute pancreatitis. Rickets may benefit from phosphate therapy, but caution should be exercised. High serum phosphate levels may increase the incidence of extraskeletal calcification.

Information for Patients
Patients with kidney stones may pass old stones when phosphate therapy is started and should be warned of this possibility. Patients should be advised to avoid the use of antacids containing aluminum, calcium, or magnesium, which may prevent the absorption of phosphate. To assure against gastrointestinal injury
associated with oral ingestion of concentrated potassium salt preparations, patients should be instructed to
dissolve tablets completely in an appropriate amount of water before taking.

**Laboratory Tests**
Careful monitoring of renal function and serum calcium, phosphorus and potassium may be required at
periodic intervals during potassium phosphate therapy. Other tests may be warranted in some patients,
depending on conditions.

**Drug Interactions**
The use of antacids containing magnesium, calcium, or aluminum in conjunction with phosphate
preparations may bind the phosphate and prevent its absorption. Potassium-containing medications or
potassium-sparing diuretics may cause hyperkalemia when used concurrently with potassium salts. Patients
should have serum potassium level determinations at periodic intervals. Concurrent use of salicylates may
lead to increased serum salicylate levels since excretion of salicylates is reduced in acidified urine. Serum
salicylate levels should be closely monitored to avoid toxicity.

**Carcinogenesis, Mutagenesis, Impairment of Fertility**
No long-term or reproduction studies in animals or humans have been performed with K-PHOS®
ORIGINAL to evaluate its carcinogenic, mutagenic, or impairment of fertility potential.

**Pregnancy**

**Teratogenic Effects**

**Pregnancy Category C**
Animal reproduction studies have not been conducted with K-PHOS® ORIGINAL. It is also not known
whether this product can cause fetal harm when administered to a pregnant woman or can affect
reproductive capacity. This product 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 this product is administered to a nursing woman.*

**ADVERSE REACTIONS**
Gastrointestinal upset (diarrhea, nausea, stomach pain, and vomiting) may occur with the use of potassium
phosphate. Also, bone and joint pain (possible phosphate-induced osteomalacia) could occur. The following
adverse effects may be observed with potassium administration: irregular heartbeat; dizziness; mental
confusion; weakness or heaviness of legs; unusual tiredness; muscle cramps; numbness, tingling, pain, or
weakness in hands or feet; numbness or tingling around lips; shortness of breath or troubled breathing.

**DOSAGE AND ADMINISTRATION**
Two tablets dissolved in 6-8 oz. of water 4 times daily with meals and at bedtime. For best results, let the
tablets soak in water for 2 to 5 minutes, or more if necessary, and stir. If any tablet particles remain
undissolved, they may be crushed and stirred vigorously to speed dissolution.

**HOW SUPPLIED**
White, scored tablet with the name BEACH and the number 1111 imprinted on each tablet. Bottles of 100
(NDC 0486-1111-01) and bottles of 500 (NDC 0486-1111-05) tablets.
STORAGE
Dispense in tight, light-resistant containers with child-resistant closures.
BEACH PHARMACEUTICALS, Div. of Beach Products, Inc., Tampa, FL 33611
Rev: 07/09B

PRINCIPAL DISPLAY PANEL - 100 Tablet Bottle Label
NDC 0486-1111-01
K-PHOS® Original
(Sodium-Free)
POTASSIUM ACID PHOSPHATE
URINARY ACIDIFIER
Rx ONLY
100 TABLETS

Beach

K-PHOS® Original (Sodium-Free)
POTASSIUM ACID PHOSPHATE
DESCRIPTION: Each tablet contains potassium acid phosphate 500 mg.
INACTIVE INGREDIENTS: Magnesium stearate, microcrystalline cellulose, silicon dioxide, starch, stearic acid
DOSAGE AND ADMINISTRATION: Two tablets dissolved in 6-8 oz. of water four times daily with meals and at bedtime. For best results, let the tablets soak in water for two to five minutes, or more if necessary, and stir. If any tablet particles remain undissolved, they may be crushed and stirred vigorously to speed dissolution. See package insert for complete prescribing information.

Beach

K PHOS ORIGINAL
potassium phosphate, monobasic tablet, soluble

Product Information

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<th>Item Code (Source)</th>
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Active Ingredient/Active Moiety

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<th>Basis of Strength</th>
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<td>Potassium Phosphate,</td>
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<td>(UNII:4J9FJ0HL51) (Phosphate Ion - UNII: NK08V8K8HR, Potassium Cation - UNII:295053K152)</td>
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## Inactive Ingredients

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<td>Silicon Dioxide</td>
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<td>Starch, Corn</td>
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<td>Stearic Acid</td>
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## Product Characteristics

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## Marketing Information

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## Labeler

- Beach Products, Inc. (032763633)

## Establishment

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Revised: 9/2014