

KALI CARBONICUM - potassium carbonate pellet
REMEDY MAKERS

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

KALI CARBONICUM 6C (Potassium Carbonate)

SHARP AND CUTTING PAIN, WEAKNESS, LABOR PAINS, OR OTHER INDICATIONS

POTASSIUM CARBONATE

WARNING:Keep this and all medications out of reach of children.

INDICATIONS: To be use according to standard homeopathic indications, for self-limiting conditions such as these indicated above or as directed by a physician.

WARNING: Use only if cap and seal are unbroken. If symptoms persist for more than 3 days or worsen, discontinue (STOP) use and consult your physician. As with any drug. If you are pregnant or nursing (breast-feeding) a baby, seek the advise of a health professional before using this product. Store tightly closed in a cool area.

Directions:(adult/children) Dissolve 3 or 4 pellets in mouth or under tongue 3 times a day or as directed by a physician. Children 2 years and older take 1/2 the adult dose.

Inactive Ingredient: Lactose and Sucrose. Free from yeast, wheat, corn and soy.

Questions or comments.(877)REM4YOU. Fax (909)594-4205 Pomona, CA. 91768. USA
www.remedymakers.com

Other information: Contain approx.145 - 150 pellets.



KALI CARBONICUM **6CH**

(Potassium Carbonate) Approx. 0.1pg per gram

**SHARP & CUTTING PAIN, WEAKNESS, LABOR PAINS,
OR OTHER INDICATIONS**

2 Drams (1/4 ounce) Lot # XXX-XXX-X Exp.XX/XXXX

Drug Facts: Active Ingredient Listed above. To be used according to standard homeopathic indications for self-limiting conditions such as those indicated above or as directed by a physician .

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NDC 10191-1665-2



**HOMEOPATHIC
MEDICINE**

KALI CARBONICUM

potassium carbonate pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10191-1665
Route of Administration	SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POTASSIUM CARBONATE (UNII: BQN1B9B9HA) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM CARBONATE	6 [hp_C]

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	
LACTOSE (UNII: J2B2A4N98G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10 19 1-16 65-2	145 in 1 VIAL, GLASS		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		12/08/2011	

Labeler - REMEDY MAKERS (018543582)

Revised: 12/2011

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