ALUMINA SILICATA- kaolin pellet Washington Homeopathic Products

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.

DRUG FACTS

ACTIVE INGREDIENTS

ALUMINA SILICATA 30C

USES

To relieve the symptoms of fainting.

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicines out of reach of children.

INDICATIONS

Indications: ALUMINA SILICATA Constriction

STOP USE AND ASK DOCTOR

If symptoms persist/worsen or if pregnant/nursing, stop use and consult your practitioner.

DIRECTIONS

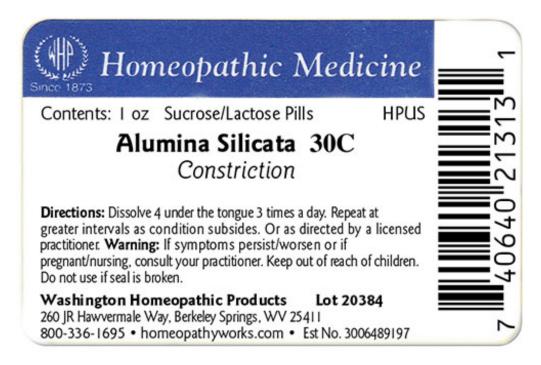
Adults: Dissolve 3 to 5 under the tongue three times a day or as directed by Lic. Practitioner. Take at greater intervals as condition subsides. Children: Dissolve 3 to 5 under the tongue three times a day or as directed by Lic. Practitioner. Take at greater intervals as condition subsides.

INACTIVE INGREDIENTS

Sucrose/Lactose

PRINCIPAL DISPLAY PANEL

The OTC potency range of ALUMINA SILICATA is 2x–30x, 1c–30c, 200c, 1m, 10m, 50m, and CM. Availability is subject to change.



All WHP single remedies are made to order; thus, the labels are printed on the same label stock as the orders are filled.

'Bottle Size' and 'Potency' vary on the label depending on customer choice.

Standard bottle sizes for pellet-form remedies are 2 dram, 4 dram, 1 ounce, 2 ounce, and 4 ounce.

| ALUMINA SILICATA kaolin pellet | | | | | | |
|-----------------------------------|--------------------------|--------------------|--------|---------------|-----|-----------|
| | | | | | | |
| Product Information | | | | | | |
| Product Type | HUMAN OTC DRUG | Item Code (Source) | | NDC:68428-874 | | |
| Route of Administration | ORAL | | | | | |
| | | | | | | |
| | | | | | | |
| Active Ingredient/Active I | Moiety | | | | | |
| Ingredient Name Basi | | | | s of Strength | | Strength |
| KAOLIN (UNII: 24H4NWX5CO) (K | AOLIN - UNII:24H4NWX5CO) | | KAOLIN | | | 30 [hp_C] |
| | | | | | | |
| Inactive Ingredients | | | | | | |
| Ingredient Name Strengt | | | | | gth | |
| SUCROSE (UNII: C151H8M554) | 0 | | | | | 0 |
| LACTOSE (UNII: J2B2A4N98G) | | | | | | |
| | | | | | | |
| | | | | | | |
| Product Characteristics | | | | | | |
| Color | white (white) | Score | | | | |
| Shape | | Size | | | | |
| Flavor | | Imprint Co | de | | | |
| Contains | | | | | | |

| Packaging | | | | | | | | |
|------------------------|----------------------|--|-------------------------|-----------------------|--|--|--|--|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date | | | | |
| 1 | NDC:68428-874- 03 | 75 in 1 VIAL, GLASS; Type 0: Not a Combination Product | 0 1/0 6/20 12 | | | | | |
| 2 | NDC:68428-874- 05 | 150 in 1 VIAL, GLASS; Type 0: Not a Combination Product | 0 1/0 6/20 12 | | | | | |
| 3 | NDC:68428-874-11 | 300 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product | 0 1/0 6/20 12 | | | | | |
| 4 | NDC:68428-874- 12 | 600 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product | 0 1/0 6/20 12 | | | | | |
| 5 | NDC:68428-874- 06 | 1200 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product | 0 1/0 6/20 12 | | | | | |
| | | | | | | | | |
| Marketing Information | | | | | | | | |
| Marketing Category | | | Marketing Start Date | Marketing End Date | | | | |
| unapproved homeopathic | | hic | 0 1/0 6 /20 12 | | | | | |
| | | | | | | | | |

Labeler - Washington Homeopathic Products (084929389)

Establishment

| Name | Address | ID/FEI | Business Operations |
|---------------------------------|---------|-----------|----------------------------|
| Washington Homeopathic Products | | 084929389 | manufacture(68428-874) |

Revised: 1/2012

Washington Homeopathic Products