

**KALI SULPHURICUM- potassium sulfate pellet**  
**Boiron**

*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 sulphuricum 6X**

Kali sulphuricum 6X

(\*\*contains 0.443 mg of the active ingredient per pellet)

Colds With Yellow Nasal Discharge\*

Stop use and ask a doctor if symptoms persist for more than 3 days or worsen

If pregnant or breast-feeding ask a health professional before use

Keep out of reach of children

Do not use if pellet dispenser seal is broken.

Contains approx 80 pellets.

How to dispense pellets? Turn tube upside down. Twist until 5 pellets are dispensed into cap. Carefully remove the cap and use it to pour pellets under the tongue.

\*CLAIMS BASED ON TRADITIONAL HOMEOPATHIC PRACTICE NOT ACCEPTED MEDICAL EVIDENCE. NOT FDA EVALUATED.

\*C,K,CK, and X are homeopathic dilutions: see BoironUSA.com/info for details.

lactose, sucrose

Adults and children: At the onset of symptoms, dissolve 5 pellets under the tongue 3 times a day until symptoms are relieved or as directed by a doctor.

1-800-BOIRON-1 (1-800-264-7661),

BoironUSA.com Info@boiron.com

Distributed by Boiron, Inc. Newtown Square, PA 19073

**Kali sulphuricum** **6<sup>x</sup>**

HPUS NDC 0220-2921-41

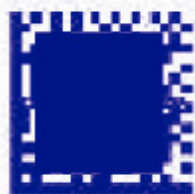
**BOIRON®**

HOMEOPATHIC MEDICINE Made in France

**Colds with yellow nasal discharge.\***

Kali sulphuricum

\*CLAIMS BASED ON TRADITIONAL HOMEOPATHIC PRACTICE NOT ACCEPTED MEDICAL EVIDENCE. NOT FDA EVALUATED.



Lot:

Exp:

6 x

3

06960

40803

1

Contains approx. 80 pellets.

US

Peel for Drugs Facts and instructions for use.

EOPATHIC PRACTICE,  
NOT FDA EVALUATED.

## Drug Facts

**Active ingredient\*\*:** See product name on front panel (contains 0.443 mg of the active ingredient per pellet).

**Uses:** See symptoms on front panel.

**Warnings:** Stop use and ask a doctor if symptoms persist for more than 3 days or worsen. If pregnant or breastfeeding, ask a health professional before use. Keep out of reach of children.

**Directions:** ■ Adults and children: At the onset of symptoms, dissolve 5 pellets under the tongue 3 times a day until symptoms are relieved or as directed by a doctor.

**Other information:** ■ Do not use if pellet dispenser seal is broken.

## Drug Facts (continued)

**Inactive ingredients:** lactose, sucrose

## Questions or comments?

BoironUSA.com Info@Boiron.com

1-800-BOIRON-1 (1-800-264-7661)

1-800-BOIRON-1 (1-800-204-7001)

\*\*C, K, CK, and X are homeopathic dilutions: see  
BoironUSA.com/info for details.

Distributed by Boiron Inc.  
Newtown Square, PA 19073

How to  
dispense pellets?



1 Turn tube upside down.



2 Twist until 5 pellets are dispensed into cap.



3 Carefully remove cap and use it to pour pellets under the tongue.

## KALI SULPHURICUM

potassium sulfate pellet

### Product Information

|                         |                |                    |               |
|-------------------------|----------------|--------------------|---------------|
| Product Type            | HUMAN OTC DRUG | Item Code (Source) | NDC:0220-2921 |
| Route of Administration | ORAL           |                    |               |

### Active Ingredient/Active Moiety

| Ingredient Name   | Basis of Strength | Strength |
|---|-------------------|----------|
| POTASSIUM SULFATE (UNII: 1K573LC5TV) (POTASSIUM CATION - UNII:295O53K152) | POTASSIUM SULFATE | 6 [hp_X] |

### Inactive Ingredients

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

### Product Characteristics

|          |       |              |     |
|----------|-------|--------------|-----|
| Color    | white | Score        |     |
| Shape    | ROUND | Size         | 4mm |
| Flavor   |       | Imprint Code |     |
| Contains |       |              |     |

### Packaging

| # | Item Code        | Package Description                             | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:0220-2921-41 | 80 in 1 TUBE; Type 0: Not a Combination Product | 03/03/1983           |                    |

| Marketing Information  |  |                      |                    |
|------------------------|--|----------------------|--------------------|
| Marketing Category     | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| unapproved homeopathic |  | 03/03/1983           |                    |

**Labeler** - Boiron (282560473)

**Registrant** - Boiron Inc. (014892269)

| Establishment |         |           |                        |
|---------------|---------|-----------|------------------------|
| Name          | Address | ID/FEI    | Business Operations    |
| Boiron        |         | 282560473 | manufacture(0220-2921) |

Revised: 4/2019

Boiron