**SILICEA BELLADONNA SPECIAL ORDER- silicea belladonna special order liquid**
**Uriel Pharmacy Inc.**

*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.*

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**Silicea Belladonna Special Order**

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 10 drops. Ages 2-11: 5 drops. Under age 2: Consult a doctor.

Active Ingredients: Argentum nitricum (Silver nitrate) 20X, Atropa belladonna (Nightshade) 20X, Silicea (Rock crystal) 20X

Inactive Ingredient: Distilled water

Use: Temporary relief of flu symptoms.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use. Do not use if safety seal is broken or missing.

REFRIGERATE AFTER OPENING.
BEST WHEN USED WITHIN 90 DAYS OF OPENING.

Questions? Call 866.642.2858 Uriel, East Troy, WI 53120 www.urielpharmacy.com


## SILICEA BELLADONNA SPECIAL ORDER

silicea belladonna special order liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:48951-8285 |
| **Route of Administration** | ORAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |

| | | |
|---|---|---|
| **SILVER** (UNII: 3M4G523W1G) (SILVER - UNII:3M4G523W1G) | SILVER | 20 [hp_X] in 1 mL |
| **ATROPA BELLADONNA** (UNII: WQZ3G9PF0H) (ATROPA BELLADONNA - UNII:WQZ3G9PF0H) | ATROPA BELLADONNA | 20 [hp_X] in 1 mL |
| **SILICON DIOXIDE** (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4) | SILICON DIOXIDE | 20 [hp_X] in 1 mL |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **WATER** (UNII: 059QF0KO0R) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:48951-8285-3 | 60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product | 09/01/2009 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved homeopathic | | 09/01/2009 | |

## Labeler - Uriel Pharmacy Inc. (043471163)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Uriel Pharmacy Inc. | | 043471163 | manufacture(48951-8285) |

Revised: 5/2018 Uriel Pharmacy Inc.