SANTONINUM- santonin liquid
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

SANTONINUM

USES

To relieve the symptoms of urinary problems.

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicines out of reach of children.

INDICATIONS

Indications:

SANTONINUM Urinary problems

STOP USE AND ASK DOCTOR

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

DIRECTIONS

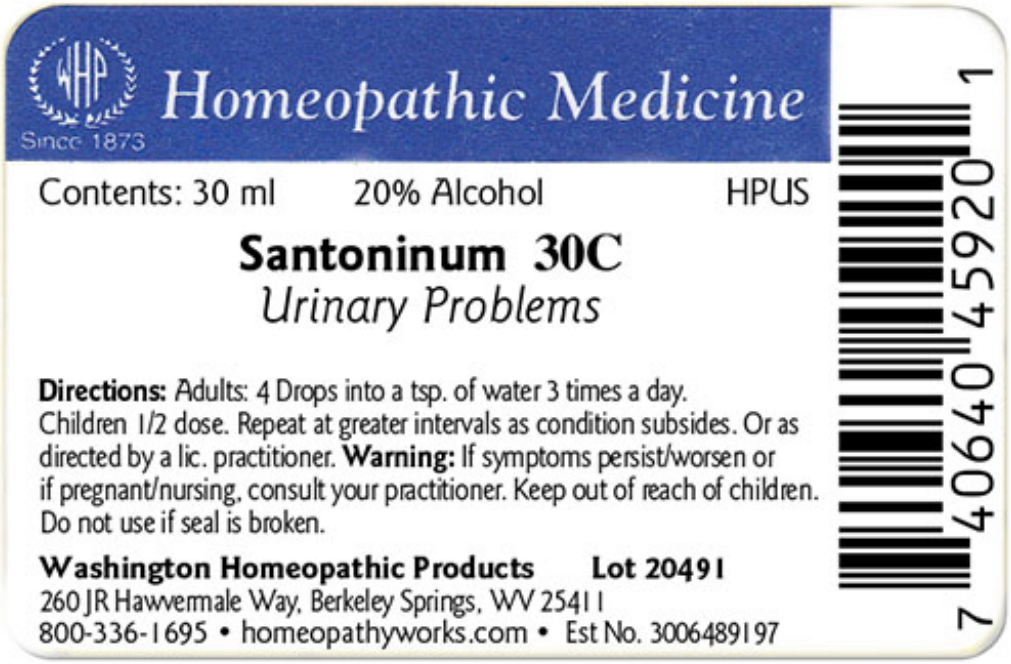
Adults: 4 drops into a tsp. of water 3 times a day. Children: 1/2 dose. Repeat at greater intervals as condition subsides. Or as directed bya lic. practitioner.

INACTIVE INGREDIENTS

Sucrose/Lactose

PRINCIPAL DISPLAY PANEL

The OTC potency range of SANTONINUM is 6x–30x, 3c–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,' 'Potency,' and 'Alcohol Percentage' vary on the label depending on customer choice.

Standard bottle sizes for dilution-form remedies are 15ml, 30ml, 50ml, and 100ml.

SANTONINUM

santonin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71919-609
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SANTONIN (UNII: 1VL8J38ERO) (SANTONIN - UNII:1VL8J38ERO)	SANTONIN	30 [hp_C] in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics			
Color	white (white)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71919-609-07	15 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	03/16/2010	
2	NDC:71919-609-08	30 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	03/16/2010	
3	NDC:71919-609-09	50 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	03/16/2010	
4	NDC:71919-609-10	100 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	03/16/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		03/16/2010	

Labeler - Washington Homeopathic Products (084929389)

Establishment

Name	Address	ID/FEI	Business Operations
Washington Homeopathic Products		084929389	manufacture(71919-609)

Revised: 3/2010 Washington Homeopathic Products