SPONGIA TOSTA- silicon dioxide 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

SPONGIA

USES

To relieve the symptoms of dry cough.

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicines out of reach of children.

INDICATIONS

Indications:

SPONGIA Dry Cough

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 SPONGIA 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.

SPONGIA TOSTA silicon dioxide pellet					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (S	ource)	NDC:6842	8-644
Route of Administration	ORAL				
Active Ingredient/Active	Moiety				
Ingre	dient Name		Basis of St	rength	Strength
SPONGIA OFFICINALIS SKELETON, ROASTED (UNII: 1PIP394IID) (SPONGIA OFFICINALIS SKELETON, ROASTED - UNII:1PIP394IID)			SPONGIA OFFICINALIS SKELETON, ROASTED		30 [hp_C]

Inactive Ingredients					
	Ingredient Name		Strength		
SUCROSE (UNII: C151H8M554)					
LACTOSE (UNII: J2B2A4N98G)					
Product Characteristics					
Color	white (white)	Score			
COIOI					
		Size			
Shape Flavor		Size Imprint Code			

Packaging					
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:68428- 644-03	75 in 1 VIAL, GLASS; Type 0: Not a Combination Product	11/20/2009		
2	NDC:68428- 644-05	150 in 1 VIAL, GLASS; Type 0: Not a Combination Product 11/20/2009			
3	NDC:68428- 644-11	28-300 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product11/20/2009			
4	NDC:68428- 644-12	600 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	11/20/2009		
5	NDC:68428- 644-06	1200 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	11/20/2009		
Marketing Information					
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
	approved meopathic		11/20/2009		

Labeler - Washington Homeopathic Products (084929389)

Establishment					
Name	Address	ID/FEI	Business Operations		
Washington Homeopathic Products		084929389	manufacture(68428-644)		

Revised: 10/2023

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