STELLARIA MEDIA- stellaria media 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

STELLARIA MED

USES

To relieve the symptoms of sluggish in morning.

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicines out of reach of children.

INDICATIONS

Indications:

STELLARIA MED Sluggish in morning

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 STELLARIA MED 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,' '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.

STELLARIA MEDIA											
stellaria media liquid											
Product Information											
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:71919-643							
Route of Administration	ORAL										
Active Ingredient/Active	Moioty										
	· · · · · · · · · · · · · · · · · · ·	D	asis of Strer	l.	Cauca zah						
				•	Strength						
STELLARIA MEDIA (UNII: 2H03479QVR) (STELLARIA MEDIA - UNII:2H03479QVR) STELLARIA MEDIA 30 [hp_C] in 1											
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:71919-643- 07	15 mL in 1 VIAL, GLASS; Type 0: Not a Combination Produ	ct 09/08/2010					
2	NDC:71919-643- 08	30 mL in 1 VIAL, GLASS; Type 0: Not a Combination Produ	act 09/08/2010					
3	NDC:71919-643- 09	50 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	09/08/2010					
4	NDC:71919-643- 10	100 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	09/08/2010					
Marketing Information								
Marketing Category		ory Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
unapproved homeopathi		thic	09/08/2010					

Labeler - Washington Homeopathic Products (084929389)

Establishment

Name	Address	ID/FEI	Business Operations
Washington Homeopathic Products		084929389	manufacture(71919-643)

Revised: 9/2010

Washington Homeopathic Products