

PLANTAGO MAJOR- plantago major 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

PLANTAGO

USES

To relieve the symptoms of restless sleep.

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicines out of reach of children.

INDICATIONS

Indications:

PLANTAGO Restless sleep

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 by a lic. practitioner.

INACTIVE INGREDIENTS

Sucrose/Lactose

PRINCIPAL DISPLAY PANEL

The OTC potency range of PLANTAGO 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.

PLANTAGO MAJOR

plantago major liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:719 19-543
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PLANTAGO MAJOR (UNII: W2469WNO6U) (PLANTAGO MAJOR - UNII:W2469WNO6U)	PLANTAGO MAJOR	30 [hp_C] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0K00R)	

Product Characteristics

Color	white (white)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start	Marketing End
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#	Item Code	Package Description	Date	Date
1	NDC:719 19-543-07	15 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	12/10/2009	
2	NDC:719 19-543-08	30 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	12/10/2009	
3	NDC:719 19-543-09	50 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	12/10/2009	
4	NDC:719 19-543-10	100 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	12/10/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		12/10/2009	

Labeler - Washington Homeopathic Products (084929389)

Establishment

Name	Address	ID/FEI	Business Operations
Washington Homeopathic Products		084929389	manufacture(719 19-543)

Revised: 12/2018

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