PASSIFLORA INCARNATA- passiflora incarnata flowering top 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

PASSIFLORA

USES

To relieve the symptoms of insomnia.

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicines out of reach of children.

INDICATIONS

Indications:

PASSIFLORA Insomnia

STOP USE AND ASK DOCTOR

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

DIRECTIONS

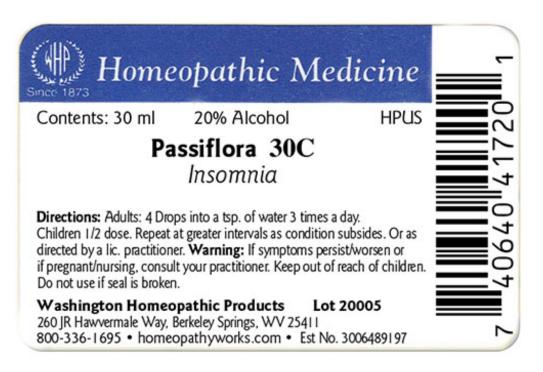
Adults: 4 drops into a tsp. of water 3 times a day. Children: 1/2 dose. Repeat at greater intervals as condition subsides. Or as directed by a lic. practitioner.

INACTIVE INGREDIENTS

Sucrose/Lactose

PRINCIPAL DISPLAY PANEL

The OTC potency range of PASSIFLORA 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.

PASSIFLORA INCARNATA

passiflora incarnata flowering top liquid

Prod	luct.	Info	ита	tion
Pron				

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71919-521	
Route of Administration	ORAL			

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PASSIFLORA INCARNATA FLO WERING TOP (UNII: CLF5YFS110) (PASSIFLORA INCARNATA FLOWERING TOP - UNII: CLF5YFS110)	PASSIFLORA INCARNATA FLOWERING TOP	30 [hp_C] in 1 mL

Inactive Ingredients

Ingredien	t Name	Strength
ALCOHOL (UNII: 3K9958V90M)		
WATER (UNII: 059QF0KO0R)		

Product Characteristics

1 Todact Characteristics				
Color	white (white)	Score		
Shape		Size		
Flavor		Imprint Code		

Contains

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:71919-521- 07	15 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	02/03/2010		
2	NDC:71919-521- 08	30 mL in 1 VIAL, GLASS; Type 0: Not a Combination Product	02/03/2010		
3	NDC:71919-521- 09	50 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	02/03/2010		
4	NDC:71919-521- 10	100 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	02/03/2010		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		02/03/2010	

Labeler - Washington Homeopathic Products (084929389)

Establishment				
Name	Address	ID/FEI	Business Operations	
Washington Homeopathic Products		084929389	manufacture(71919-521)	

Revised: 2/2010 Washington Homeopathic Products