

DIGITALINUM- digitalin 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

DIGITALINUM

USES

To relieve the symptoms of numbness.

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicines out of reach of children.

INDICATIONS

Indications:

DIGITALINUM Numbness

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 DIGITALINUM is 8x–30x, 4c–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.

DIGITALINUM

digitalin pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68428-933
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIGITALIN (UNII: Q15ECE254B) (DIGITALIN - UNII:Q15ECE254B)	DIGITALIN	30 [hp_C]

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	
LACTOSE (UNII: J2B2A4N98G)	

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:68428-933-03	75 in 1 VIAL, GLASS; Type 0: Not a Combination Product	05/26/2011	
2	NDC:68428-933-05	150 in 1 VIAL, GLASS; Type 0: Not a Combination Product	05/26/2011	
3	NDC:68428-933-11	300 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	05/26/2011	
4	NDC:68428-933-12	600 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	05/26/2011	
5	NDC:68428-933-06	1200 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	05/26/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		05/26/2011	

Labeler - Washington Homeopathic Products (084929389)

Establishment

Name	Address	ID/FEI	Business Operations
Washington Homeopathic Products		084929389	manufacture(68428-933)

Revised: 5/2011

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