DIGITALIS PUPUREA - digitalis 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

DIGITALIS

USES

To relieve the symptoms of worry.

KEEP OUT OF REACH OF CHILDREN

Keep this and all medicines out of reach of children.

INDICATIONS

Indications:

DIGITALIS Worry

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 DIGITALIS is 4x–30x, 2c–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.

DIGITALIS PUPUREA

digitalis pellet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68428-357
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIGITALIS (UNII: F1T8QT9U8B) (DIGITALIS - UNII:F1T8QT9U8B)	DIGITALIS	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:68428-357-03	75 in 1 VIAL, GLASS; Type 0: Not a Combination Product	02/03/2010	
2	NDC:68428-357-05	150 in 1 VIAL, GLASS; Type 0: Not a Combination Product	02/03/2010	
3	NDC:68428-357-11	300 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	02/03/2010	
4	NDC:68428-357-12	600 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	02/03/2010	
5	NDC:68428-357-06	1200 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)

Registrant - Washington Homeopathic Products (084929389)

Establishment

Name	Address	ID/FEI	Business Operations
Washington Homeopathic Products		084929389	manufacture(68428-357)

Revised: 12/2022 Washington Homeopathic Products