DIGITALIS PURPUREA- digitalis purpurea tablet
Rxhomeo Private Limited d.b.a. Rxhomeo, Inc

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.

ACTIVE INGREDIENT

DIGITALIS PURPUREA HPUS 6X and Higher

USES

Swollen Legs

INDICATIONS

Condition listed above or as directed by the physician

DOSAGE

Adults- Take 4 or 6 Tablets by mouth, three times daily or as suggested by physician. Children 2 years and older- take 1/2 the adult dose.

WARNINGS

This product is to be used for self-limiting conditions

If symptoms do not improve in 4 days, or worsen, discontinue use and seek assistance of health professional

As with any drug, if you are pregnant, or nursing a baby, seek professional advice before taking this product

Keep this and all medication out of reach of children

Do not use if capseal is broken or missing.

Close the cap tightly after use.

INACTIVE INGREDIENTS

Lactose

STORAGE

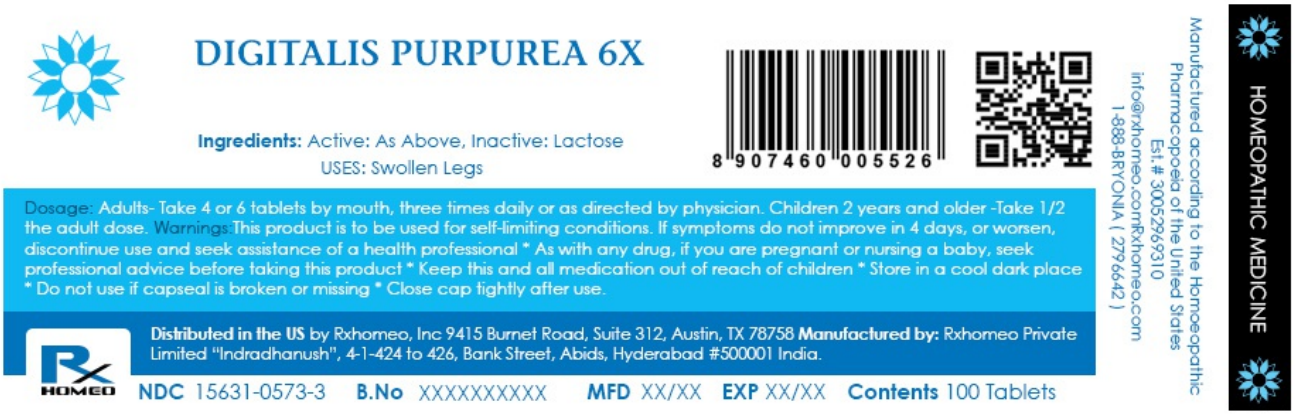
Store in a cool dark place

QUESTIONS OR COMMENTS

www.Rxhomeo.com | 1.888.2796642 | info@rxhomeo.com

Rxhomeo, Inc 9415 Burnet Road, Suite 312, Austin, TX 78758

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



DIGITALIS PURPUREA

digitalis purpurea tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:15631-0573
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIGITALIS (UNII: F1T8QT9U8B) (DIGITALIS - UNII:F1T8QT9U8B)	DIGITALIS	6 [hp_X]

Inactive Ingredients	
Ingredient Name	**Strength**
LACTOSE (UNII: J2B2A4N98G)	

Product Characteristics			
Color	white	**Score**	no score
Shape	ROUND	**Size**	3mm
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:15631-0573-0	1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:15631-0573-1	4 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:15631-0573-2	50 in 1 CONTAINER; Type 0: Not a Combination Product		
4	NDC:15631-0573-3	100 in 1 CONTAINER; Type 0: Not a Combination Product		

5	NDC:15631-0573-4	250 in 1 CONTAINER; Type 0: Not a Combination Product		
6	NDC:15631-0573-5	500 in 1 CONTAINER; Type 0: Not a Combination Product		
7	NDC:15631-0573-6	1000 in 1 CONTAINER; Type 0: Not a Combination Product		
8	NDC:15631-0573-7	10000 in 1 CONTAINER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		12/27/2015	

Labeler - Rxhomeo Private Limited d.b.a. Rxhomeo, Inc (650833994)

Establishment

Name	Address	ID/FEI	Business Operations
Rxhomeo, Inc		832534981	wholesale drug distributor(15631-0573)

Establishment

Name	Address	ID/FEI	Business Operations
Rxhomeo Private Limited d.b.a. Rxhomeo, Inc		650833994	manufacture(15631-0573) , label(15631-0573)

Revised: 1/2016 Rxhomeo Private Limited d.b.a. Rxhomeo, Inc