

PHYTOLACCA DECANDRA 200C- phytolacca decandra pellet

Paramesh Banerji Life Sciences LLC

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 Ingredient

Phytolacca decandra 200C HPUS

Inactive Ingredients

Sucrose, Lactose

Purpose

Throat pain, breast pain

Uses

Throat pain, breast pain

Warnings

If pregnant or breast feeding ask a health professional before use.

Keep out of reach of children

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Direction

Adult or child: Take three pills daily. Leaving a gap of 30 minutes after any food or as advised by your physician.

Manufactured by

Paramesh Banerji Life Sciences, LLC.

North Brunswick, NJ 08902, USA.

Tel: +1-732-743-5936,

Email: info@pblifesciences.com

www.pblifesciences.com/permapotent

Principal Display Panel

PermaPotent™
homeopathics NDC: 69152-1203-1

Lot No.:

**PHYTOLACCA
DECANDRA 200C**

HOMEOPATHIC MEDICINE

THROAT PAIN,
BREAST PAIN

96 PILLS (Approx.)

PRODUCT
OF USA

www.pblifesciences.com

PBL
PARAMESH
BANERJI
LIFE
SCIENCES

Mfg. Dt.:

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Phytolacca decandra 200C

Homeopathic Medicine

Throat pain, breast pain

96 Pills (Approx.)

Product of USA

PHYTOLACCA DECANDRA 200C

phytolacca decandra pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69 152-1203
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PHYTOLACCA AMERICANA ROOT (UNII: 11E6 V18 VEG) (PHYTOLACCA AMERICANA ROOT - UNII:11E6 V18 VEG)	PHYTOLACCA AMERICANA ROOT	200 [hp_C]

Inactive Ingredients

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

Product Characteristics

Color	white	Score	
Shape	ROUND	Size	4mm
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69152-1203-1	96 in 1 BOTTLE; Type 0: Not a Combination Product	08/26/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		08/26/2015	

Labeler - Paramesh Banerji Life Sciences LLC (079393726)

Establishment

Name	Address	ID/FEI	Business Operations
Paramesh Banerji Life Sciences LLC		079393726	manufacture(69152-1203) , pack(69152-1203) , label(69152-1203)

Revised: 4/2016

Paramesh Banerji Life Sciences LLC