

TRILIF 40 (NUMBER 80)- antimonium crudum, graphites 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 Ingredients

Antimonium crudum 30C HPUS, Graphites 30C HPUS

Inactive Ingredients

Sucrose, Lactose

Purpose

Clears and prevents warts and growths on skin

Uses

Clears and prevents warts and growths on skin

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

Principal Display Panel

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NDC: 69152-0080-1

TRILIF 40

NUMBER 80

HOMEOPATHIC MEDICINE

CLEARS & PREVENTS WARTS
AND GROWTHS ON SKINPRODUCT
OF USA

96 PILLS (Approx.)

Lot No.:

Mfg. Dt.:

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Trilif 40

Number 80

Homeopathic Medicine

Clears and prevents warts and growths on skin

96 Pills (Approx.)

Product of USA

TRILIF 40 (NUMBER 80)

antimonium crudum, graphites pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69152-0080
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ANTIMONY TRISULFIDE (UNII: F79059A38U) (ANTIMONY TRISULFIDE - UNII:F79059A38U)	ANTIMONY TRISULFIDE	30 [hp_C]
GRAPHITE (UNII: 4QQN74LH4O) (GRAPHITE - UNII:4QQN74LH4O)	GRAPHITE	30 [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:69152-0080-1	96 in 1 BOTTLE; Type 0: Not a Combination Product	07/16/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/31/2014	

Labeler - Paramesh Banerji Life Sciences LLC (079393726)

Establishment

Name	Address	ID/FEI	Business Operations
Paramesh Banerji Life Sciences LLC		079393726	manufacture(69152-0080) , pack(69152-0080) , label(69152-0080)

Revised: 11/2016

Paramesh Banerji Life Sciences LLC