

IBEALEE WART REMOVER- wart remover patch
Guangzhou Hanhai Trading Co., Ltd

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Active Ingredient

Salicylic Acid 40%

Purpose

WART REMOVER

Use

How to use:

Directions

- wash affected area and dry thoroughly
- apply tag patch
- after 24 hours, remove tag patch
- repeat procedure every 24 hours for up to 14 days for tags /warts removal and up to 12 weeks for warts, until the problem has cleared.
- may hot compress wart in warm water for 5 minutes to assist in removal

Warnings

Caution:

1. Not recommended to be used simultaneously with other medication.
2. If skin becomes red and swollen, or if burning, aching and other forms of irritation occur, cease using and seek medical advice at once.
3. It is normal to find bubbles in the colloid.
4. Keep out of reach of children.

Keep Out Of Reach Of Children

Do not place this product within the reach of children

Do not use

If allergy occurs, please stop using it

Inactive ingredients

Tea Tree Oil, Calendula, Cica Oil, Castor Oil, Argan Oil

Directions

Apply gently to skin

PRINCIPAL DISPLAY PANEL







IBEALEE WART REMOVER

wart remover patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83675-014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	40 g in 100

Inactive Ingredients

Ingredient Name	Strength
OITICICA OIL (UNII: Q4F4N7T4JN)	
TEA TREE OIL (UNII: VIF565UC2G)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
CASTOR OIL (UNII: D5340Y2I9G)	
ARGAN OIL (UNII: 4V59G5UW9X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83675-014-01	144 in 1 PATCH; Type 0: Not a Combination Product	09/26/2023	
2	NDC:83675-014-02	180 in 1 PATCH; Type 0: Not a Combination Product	09/26/2023	
3	NDC:83675-014-03	540 in 1 PATCH; Type 0: Not a Combination Product	09/26/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358B	09/26/2023	

Labeler - Guangzhou Hanhai Trading Co., Ltd (419707381)

Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Hanhai Trading Co., Ltd		419707381	label(83675-014) , manufacture(83675-014)

Revised: 9/2023

Guangzhou Hanhai Trading Co., Ltd