WART REMOVER- wart remover liquid
Shenzhen Mengyue Technology Co., Ltd.

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003 WART REMOVER LIQUID

Salicylic Acid 17%

Wart remover

USES
•To remove warts such as plantar wart, flat wart, common wart and corns, calluses.
•The plantar wart is recognized by its location only on the bottom of the foot it's tenderness, and the interruption of the footprint pattern.

WARNINGS

· The liquid can get easily volatilized & crystallized, tighten the cap after use.

· Avoid long-term contact with air while using.

· For external use only, don't contact with eyes or swallow.Children should use it under the supervision of adults.

DO NOT USE

• On damaged skin(cuts, abrasions, eczema, sunburn.

• If you are allergic to any of the ingredients in this product.

• If you are pregnant or breastfeeding.

N/A

ASK DOCTOR

See a doctor as you may have a more serious skin condition.

KEEP OUT OF REACHOF CHILDREN

•If swallowed, get medical help or contact a Poison Control Center right away.

DIRECTIONS

• Wash the affected area.

• May soak the wart in warm water for 5 minutes.

• Dry area thoroughly.

• Using the applicator(cotton swab),apply a layer of ointment to sufficiently cover each wart

• Allow it to fully absorb and cover it with a bandage as needed.

• Repeat this procedure once or twice daily as needed(until the wart is removed) for up to 12 weeks.

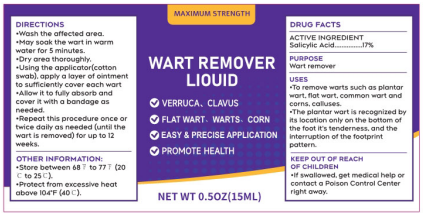
OTHER INFORMATION:

• Store between 68°F to 77°F(20°C to 25°C).

• Protect from excessive heat above 104°F(40°C).

INACTIVE INGREDIENTS
Shard paraffin, lanolin, phenol.

60x30mm



30*30*70mm

WART REMOVER

wart remover liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87255-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	17 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
PHENOL (UNII: 339NCG44TV)	
LANOLIN (UNII: 7EV65EAW6H)	
PARAFFIN (UNII: I9O0E3H2ZE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87255-003-01	15 mL in 1 BOX; Type 0: Not a Combination Product	11/13/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M028	11/13/2025	

Labeler - Shenzhen Mengyue Technology Co., Ltd. (511200055)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Mengyue Technology Co., Ltd.		511200055	manufacture(87255-003)

Revised: 11/2025 Shenzhen Mengyue Technology Co., Ltd.