

VKAVKA WART REMOVER- salicylic acid liquid
Shenzhen Lanxi Trading Co., Ltd.

87386-001 vkavka Wart Remover

Salicylic acid 12%

wart remover

Effectively removes both common warts and plantar warts

Common warts are identified by their raised, rough texture resembling a cauliflower's surface

Plantar warts specifically occur on weight-bearing areas of the foot, typically causing pressure pain when walking and noticeable disruption of the natural foot skin ridge patterns

Use only on the skin surface.

Keep away from fire and flame

Precise application: Cover only the wart, not surrounding skin

on irritated skin or on any area that is infected or reddened.

on moles, birthmarks, warts with hair growing from them, warts on the face or mucous membranes.

If you have a known hypersensitivity disorder or diabetes mellitus or poor blood circulation.

Avoid contact with eyes. If the product gets into the eye, flush with water for 15 minutes.

Avoid inhaling vapors.

Cap tightly and store at room temperature, away from heat.

Non-edible.

If discomfort persists.

Keep out of reach of children. If swallowed get medical help or contact the Poison Control Center (1-800-222-1222) right away.

Wash the affected area.

May soak wart in warm water for 5 minutes

Dry area thoroughly.

Use a brush to apply a sufficient amount to cover each wart.

Allow it to fully absorb and let dry.

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

Water

Sophora flavescens root

Cnidium monnieri whole

Hawthorn leaf with flower

Podophyllum
Stellera chamaejasm whole
Borneol

Drug Facts
Active ingredients **Purpose**
Salicylic acid 12%w/w wart remover

Indications & Uses

- Effectively removes both common warts and plantar warts
- Common warts are identified by their raised, rough texture resembling a cauliflower's surface
- Plantar warts specifically occur on weight-bearing areas of the foot, typically causing pressure pain when walking and noticeable disruption of the natural foot skin ridge patterns

Do not use

- on irritated skin or on any area that is infected or reddened.
- on moles, birthmarks, warts with hair growing from them, warts on the face or mucous membranes.
- If you have a known hypersensitivity disorder or diabetes mellitus or poor blood circulation.

Warnings

- Use only on the skin surface.
- Keep away from fire and flame.
- Precise application: Cover only the wart, not surrounding skin.

When using this product

- Avoid contact with eyes. If the product gets into the eye, flush with water for 15 minutes.
- Avoid inhaling vapors.
- Cap tightly and store at room temperature, away from heat.
- Non-edible.

Stop use and ask a doctor

- If discomfort persists.
- Keep out of reach of children. If swallowed get medical help or contact the Poison Control Center (1-800-222-1222) right away.

Directions

- Wash the affected area.
- May soak wart in warm water for 5 minutes.
- Dry area thoroughly.
- Use a brush to apply a sufficient amount to cover each wart.
- Allow it to fully absorb and let dry.
- Repeat this procedure once or twice daily as needed (until wart is removed) for up to 12 weeks.

Inactive ingredients

Water, Sophora flavescens, Cnidium monnieri, Hawthorn, Podophyllum, Stellera chamaejasm, Borneol.

Other information

Skin discoloration may occur during or after use.

vkavka®

salicylic acid liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:87386-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	0.12 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PODOPHYLLUM (UNII: 2S713A4VP3)	
SOPHORA FLAVESCENS ROOT (UNII: IYR6K8KQ5K)	
STELLERA CHAMAEJASME WHOLE (UNII: 80N0P0DD9J)	
CNIDIUM MONNIERI WHOLE (UNII: GYR30735RE)	
BORNEOL (UNII: M89NIB437X)	
HAWTHORN LEAF WITH FLOWER (UNII: 60M09R PY36)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87386-001-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/29/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M028	01/29/2026	

Labeler - Shenzhen Lanxi Trading Co., Ltd. (512086574)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lanxi Trading Co., Ltd.		512086574	manufacture(87386-001)

Revised: 1/2026

Shenzhen Lanxi Trading Co., Ltd.