EELHOE SKIN ADVANCED MOLE REMOVE SERUM- glycerin liquid
Shantou Eelhoe Daily Chemical Technology Co., Ltd.

MELALEUCA ALTERNIFOLIA FLOWERING TOP□HAMAMELIS VIRGINIANA (WITCH HAZEL) LEAF WATER

1: Soak the area with warts in warm water to soften the skin

2: If necessary, scrape off the hard skin. Apply the verruca solution to the affected area for 2-3 times to obtain the best effect.

3: The skin of the affected area will change during use and return to normal after a period of time.Restore your pink, smooth skin

Just a few drops on the blemish and the serum penetrates to the roct of the wart or skin tag. This triggers a massive influx of white blood cells to the blemish and quickly begins the removal process.

Please keep out of reach of children. Do not swallow.Please clean your hands before use to ensure the best results from the product. Discontinue use if signs of irritation or rash occur. Store in a cool and dry place.

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GLYCERIN□PROPYLENE GLYCOL□BUTYLENE GLYCOL□PEG-40 HYDROGENATED CASTOR OIL□SALICYLIC ACID

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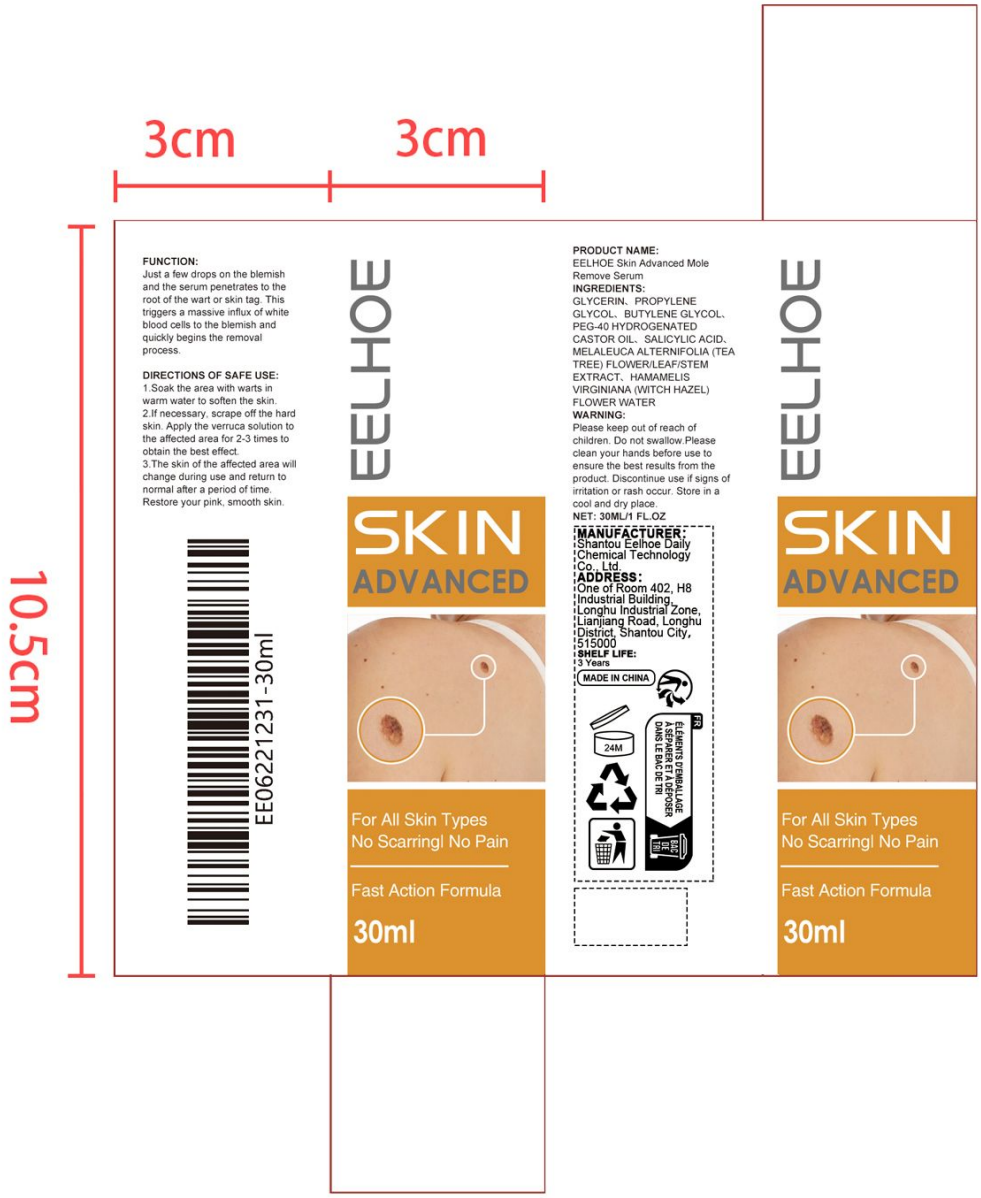
纸盒

规格尺寸：长3X宽3X高10.5CM



EELHOE SKIN ADVANCED MOLE REMOVE SERUM

glycerin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:85064-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MELALEUCA ALTERNIFOLIA FLOWERING TOP (UNII: 5AZ4S6N66F)	MELALEUCA ALTERNIFOLIA	0.006 mg

(MELALEUCA ALTERNIFOLIA FLOWERING TOP - UNII:5AZ4S6N66F)	FLOWERING TOP	in 30 mg
HAMAMELIS VIRGINIANA (WITCH HAZEL) LEAF WATER (UNII: 8FP93ED6H2) (HAMAMELIS VIRGINIANA (WITCH HAZEL) LEAF WATER - UNII:8FP93ED6H2)	HAMAMELIS VIRGINIANA (WITCH HAZEL) LEAF WATER	0.006 mg in 30 mg

Inactive Ingredients

Ingredient Name	Strength
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	9 mg in 30 mg
SALICYLIC ACID (UNII: O414PZ4LPZ)	0.006 mg in 30 mg
PEG-40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	0.03 mg in 30 mg
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	9 mg in 30 mg
GLYCERIN (UNII: PDC6A3C0OX)	11.952 mg in 30 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85064-012-01	30 mg in 1 BOTTLE; Type 0: Not a Combination Product	09/17/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	09/17/2025	

Labeler - Shantou Eelhoe Daily Chemical Technology Co., Ltd. (617290813)

Registrant - Shantou Eelhoe Daily Chemical Technology Co., Ltd. (617290813)

Establishment

Name	Address	ID/FEI	Business Operations
Shantou Eelhoe Daily Chemical Technology Co., Ltd.		617290813	manufacture(85064-012)

Revised: 11/2025 Shantou Eelhoe Daily Chemical Technology Co., Ltd.