

ANTI-SCAR DROPS 2084- anti-scar drops liquid

Professional Complementary Health Formulas

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.

C84

ACTIVE INGREDIENTS

Thuja occidentalis 3X, 6X, 12X, 24X, 30X
Aconitum napellus 6X
Alumen 6X
Silicea 6X
Calcarea phosphorica 6X, 12X
Thiosinaminum 6X, 12X
Nitricum acidum 8X
Graphites 12X
X-ray 60X
Arnica montana 100X

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of painful, sensitive, or itchy skin resulting from injury.*

*Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

WARNINGS

In case of overdose, get medical help or contact a poison control center right away.

Keep out of the reach of children.

If pregnant or breastfeeding, ask a healthcare professional before use.

DIRECTIONS

Place drops under tongue 30 minutes before/after meals. Adults and children 12 years and over: Take 10 drops up to 3 times per day. Consult a physician for use in children under 12 years of age.

OTHER INFORMATION

Tamper resistant. If seal is broken, do not use. After opening, close container tightly and store at room temperature away from heat.

INACTIVE INGREDIENTS

20% ethanol, purified water.

LABEL

Est 1985

Professional Formulas

Complementary Health

Anti-Scar Drops

Homeopathic Remedy

2 FL. OZ. (59 mL)



ANTI-SCAR DROPS 2084

anti-scar drops liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-2084
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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THUJA OCCIDENTALIS LEAF (UNII: 0T0DQN8786) (THUJA OCCIDENTALIS LEAF - UNII:0T0DQN8786)	THUJA OCCIDENTALIS LEAF	3 [hp_X] in 59 mL
ACONITUM NAPELLUS WHOLE (UNII: U0NQ8555JD) (ACONITUM NAPELLUS WHOLE - UNII:U0NQ8555JD)	ACONITUM NAPELLUS WHOLE	6 [hp_X] in 59 mL
POTASSIUM ALUM (UNII: 1L24V9R23S) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	POTASSIUM ALUM	6 [hp_X] in 59 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	6 [hp_X] in 59 mL
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J) (ANHYDROUS DIBASIC CALCIUM PHOSPHATE - UNII:L11K75P92J)	ANHYDROUS DIBASIC CALCIUM PHOSPHATE	6 [hp_X] in 59 mL
ALLYLTHIOUREA (UNII: 706IDJ14B7) (ALLYLTHIOUREA - UNII:706IDJ14B7)	ALLYLTHIOUREA	6 [hp_X] in 59 mL
NITRIC ACID (UNII: 411VRN1TV4) (NITRIC ACID - UNII:411VRN1TV4)	NITRIC ACID	8 [hp_X] in 59 mL
GRAPHITE (UNII: 4QQN74LH4O) (GRAPHITE - UNII:4QQN74LH4O)	GRAPHITE	12 [hp_X] in 59 mL
LACTOSE, X-RAY EXPOSED (1000 RAD) (UNII: LNT739I158) (LACTOSE, X-RAY EXPOSED (1000 RAD) - UNII:LNT739I158)	LACTOSE, X-RAY EXPOSED (1000 RAD)	60 [hp_X] in 59 mL
ARNICA MONTANA WHOLE (UNII: O80TY208ZW) (ARNICA MONTANA WHOLE - UNII:O80TY208ZW)	ARNICA MONTANA WHOLE	100 [hp_X] in 59 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63083-2084-2	59 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	08/15/1985	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		08/15/1984	

Labeler - Professional Complementary Health Formulas (167339027)

Registrant - Natural Pharmaceutical Manufacturing LLC (015624923)

Establishment

Name	Address	ID/FEI	Business Operations
Natural Pharmaceutical Manufacturing LLC		015624923	manufacture(63083-2084)