UMCKA COUGH ACONITUM NAPELLUS, BRYONIA ALBA ROOT, CALCIUM SULFIDE, SPONGIA OFFICINALIS SKELETON, ROASTED , TIN, PELARGONIUM SIDOIDES ROOT- umcka cough syrup
Schwabe North America, Inc.

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.

Umcka Cough Max Relief

Active Ingredients

Pelargonium sidoides 1X

Aconitum napellus 3X

Bryonia 3X

Hepar sulphuris calcareum 6C

Stannum metallicum 6C

Spongia tosta 3X

Inactive Ingredients

Citric Acid

English Ivy Leaf Extract

Ethyl Alcohol

Glycerin

Lactose Monohydrate

Maltodextrin

Potassium Sorbate

Purified Water

Sorbitol

Purpose

Shortens duration and reduces severity of symptoms associated with the common cold and throat/nasal/bronchial irritations: congestion, cough, hoarseness, sore throat, sneezing, stuffy/runny nose.

Temporarily relieves cough due to minor bronchial and throat irritations as may occur with a cold.

Indications & Usage

Shortens duration and reduces severity of symptoms associated with the common cold and throat/nasal/bronchial irritations: congestion, cough, hoarseness, sore throat, sneezing, stuffy/runny nose.

Temporarily relieves cough due to minor bronchial and throat irritations as may occur with a cold.

Doage & Administration

Directions

For best results, use at first sign of symptoms.

Continue to use for an additional 48 hours after symptoms cease.

Shake well before each use.

Use only with enclosed dosage cup.

Adults and Children 12 years of age and older: Take 1 ½ teaspoons (tsp) (7.5 mL) three times daily.

Children 6 to 11 years of age: Take 1 teaspoon (5 mL) three times daily.

Children under 6 years of age: Consult a physician.

Warnings

Sore throat warning: severe or persistent sore throat for more than 2 days or if accompanied by high fever, headache, nausea, vomiting, or rash may be serious.

Consult a physician promptly.

Ask Doctor

Ask a doctor before use if you have a persistent or chronic cough that lasts, is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema, or is accompanied by excessive phlegm (mucus).

Stop Use

Stop use and ask a doctor if new symptoms occur, symptoms worsen or do not get better within 7 days, fever worsens or lasts more than 3 days, cough lasts more than 7 days or occurs with rash or persistent headache.

These could be a sign of a serious condition.

Overdose

In case of overdose, seek medical help or contact a Poison Control Center immediately.

Pregnancy or Breast Feeding

If pregnant or breast-feeding, ask a healthcare professional before use.

Keep out of reach of children

Keep out of reach of children.

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umcka cough syrup

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53499-5864
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PELARGONIUM SIDOIDES ROOT (UNII: H6J53HEX8E) (PELARGONIUM SIDOIDES ROOT - UNII:H6J53HEX8E)	PELARGONIUM SIDOIDES ROOT	1 [hp_X] in 120 mL
ACONITUM NAPELLUS (UNII: U0NQ8555JD) (ACONITUM NAPELLUS - UNII:U0NQ8555JD)	ACONITUM NAPELLUS	3 [hp_X] in 120 mL
BRYONIA ALBA ROOT (UNII: T7J046YI2B) (BRYONIA ALBA ROOT - UNII:T7J046YI2B)	BRYONIA ALBA ROOT	3 [hp_X] in 120 mL
CALCIUM SULFIDE (UNII: 1MBW07J51Q) (CALCIUM SULFIDE - UNII:1MBW07J51Q)	CALCIUM SULFIDE	6 [hp_C] in 120 mL
TIN (UNII: 387GMG9FH5) (TIN - UNII:387GMG9FH5)	TIN	6 [hp_C] in 120 mL
SPONGIA OFFICINALIS SKELETON, ROASTED (UNII: 1PIP394IID) (SPONGIA OFFICINALIS SKELETON, ROASTED - UNII:1PIP394IID)	SPONGIA OFFICINALIS SKELETON, ROASTED	3 [hp_X] in 120 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
HEDERA HELIX LEAF (UNII: ZP9XFG71A7)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53499-5864-4	1 in 1 BOX	10/01/2012	11/30/2025
1		120 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/01/2012	11/30/2025

Labeler - Schwabe North America, Inc. (831153908)

Establishment

Name	Address	ID/FEI	Business Operations
Schwabe North America		831153908	manufacture(53499-5864)

Revised: 4/2024 Schwabe North America, Inc.