TINNITUS CONTROL - arnica montana, chininum sulphuricum, ferrum metallicum, kali phosphoricum, natrum sulphuricum, pulsatilla, silicea, thiosinaminum, spray
Liddell Laboratories, 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.

Tinnitus Control

ACTIVE INGREDIENTS: Arnica montana 30X, Chininum sulphuricum 30X, Ferrum Metallicum 30X, Kali phosphoricum 30X, Natrum sulphuricum 30X, Pulsatilla 30X, Silicea 30X, Thiosinaminum 30X.

INDICATIONS: Helps relieve symptoms of Tinnitus.

WARNINGS: If pregnant or breast feeding, seek the advice of a doctor before use.

Do not use if you have ever had an allergic reaction to this product or any of its ingredients.

Stop use and ask a doctor if symptoms persist, worsen or if new symptoms occur.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Do not use if tamper evident seal around neck of bottle is missing or broken.

DIRECTIONS: Adults and Children over 12: Spray twice under the tongue 3 times per day. Children under 12: Consult a doctor prior to use.

INACTIVE INGREDIENTS: Alcohol 20% v/v, Purified water.

INDICATIONS: Helps relieve symptoms of Tinnitus.

Distributed by:

Nutralogy

2049 North Lincoln Street

Burbank, CA 91504

Made in the USA

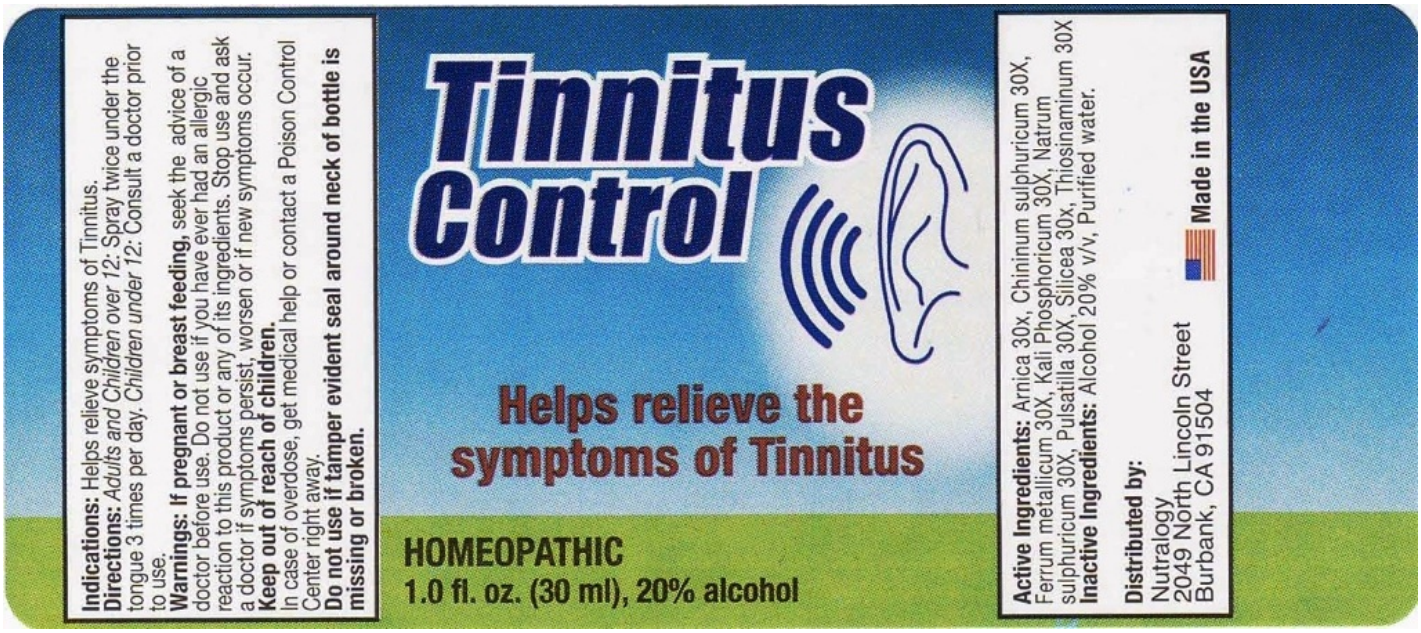
KEEP OUT OF REACH OF CHILDREN. In case of overdose, get medical help or contact a Poison Control Center right away.

Tinnitus Control

Helps relieve the symptoms of Tinnitus

HOMEOPATHIC

1.0 fl. oz. (30 ml)



TINNITUS CONTROL

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50845-0130
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	30 [hp_X] in 1 mL
QUININE SULFATE (UNII: KF7Z0E0Q2B) (QUININE - UNII:A7V27PHC7A)	QUININE SULFATE	30 [hp_X] in 1 mL
IRON (UNII: E1UOL152H7) (IRON - UNII:E1UOL152H7)	IRON	30 [hp_X] in 1 mL
POTASSIUM PHOSPHATE, DIBASIC (UNII: CI71S98N1Z) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM PHOSPHATE, DIBASIC	30 [hp_X] in 1 mL
SODIUM SULFATE (UNII: 0YPR65R21J) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM SULFATE	30 [hp_X] in 1 mL
PULSATILLA VULGARIS (UNII: I76KB35JEV) (PULSATILLA VULGARIS - UNII:I76KB35JEV)	PULSATILLA VULGARIS	30 [hp_X] in 1 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	30 [hp_X] in 1 mL
ALLYLTHIOUREA (UNII: 706IDJ14B7) (ALLYLTHIOUREA - UNII:706IDJ14B7)	ALLYLTHIOUREA	30 [hp_X] in 1 mL

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50845-0130-1	30 mL in 1 BOTTLE, SPRAY		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		04/26/2011	

Labeler - Liddell Laboratories, Inc. (832264241)

Establishment

Name	Address	ID/FEI	Business Operations
Liddell Laboratories, Inc.		832264241	manufacture

Revised: 4/2011 Liddell Laboratories, Inc.