

SECALE NICOTIANA- secale nicotiana liquid
Uriel Pharmacy 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.

Secale Nicotiana

Directions: FOR ORAL USE.

Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow.

Active Ingredients: Betula (Silver birch leaves) 3X, Betula (Silver birch bark) 4X, Nicotiana (Tobacco) 6X, Secale corn. e grano (Ergot) 6X, Arteria poplitea (Bovine popliteal artery) 8X, Galena (Lead glance) 8X

Inactive Ingredients: Water, Salt

Use: Temporary relief of headache.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use.

Questions? Call 866.642.2858 Uriel, East Troy, WI 53120 shopuriel.com

SECALE NICOTIANA

secale nicotiana liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-8176
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
BETULA PUBESCENS LEAF (UNII: 845OH00300) (BETULA PUBESCENS LEAF - UNII:845OH00300)	BETULA PUBESCENS LEAF	3 [hp_X] in 1 mL
BETULA PUBESCENS BARK (UNII: 3R504894L9) (BETULA PUBESCENS BARK - UNII:3R504894L9)	BETULA PUBESCENS BARK	4 [hp_X] in 1 mL
TOBACCO LEAF (UNII: 6YR2608RSU) (TOBACCO LEAF - UNII:6YR2608RSU)	TOBACCO LEAF	6 [hp_X] in 1 mL
CLAVICEPS PURPUREA SCLEROTIUM (UNII: 01G9XEA93N) (CLAVICEPS PURPUREA SCLEROTIUM - UNII:01G9XEA93N)	CLAVICEPS PURPUREA SCLEROTIUM	6 [hp_X] in 1 mL
BOS TAURUS ARTERY (UNII: R2M88A4HSI) (BOS TAURUS ARTERY - UNII:R2M88A4HSI)	BOS TAURUS ARTERY	8 [hp_X] in 1 mL
LEAD SULFIDE (UNII: 2425D15SYM) (LEAD - UNII:2P299V784P)	LEAD SULFIDE	8 [hp_X] in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-8176-1	10 in 1 BOX	09/01/2009	
1		1 mL in 1 AMPULE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2009	

Labeler - Uriel Pharmacy Inc. (043471163)

Establishment

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-8176)

Revised: 10/2023

Uriel Pharmacy Inc.