

SECALE QUARTZ- secale quartz pellet
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 Quartz

Directions: FOR ORAL USE ONLY.

Dissolve pellets under the tongue 3-4 times daily. Ages 12 and older: 10 pellets. Ages 2-11: 5 pellets. Under age 2: Consult a doctor.

Active Ingredients: Sanguinaria (Bloodwort) 6X, Secale corn. e grano (Ergot) 6X, Quartz (Rock crystal) 30X

Inactive Ingredient: Organic sucrose

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. Contains sugar. Diabetics and persons intolerant of sucrose (sugar): Consult a doctor before use. 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. Do not use if safety seal is broken or missing.

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

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Secale Quartz
Homeopathic Pellets
net wt. 1 oz (30g)

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Uriel, East Troy, WI 53120
www.urielpharmacy.com Lot:

SECALE QUARTZ

secale quartz pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-8180
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SANGUINARIA CANADENSIS ROOT (UNII: N9288CD508) (SANGUINARIA CANADENSIS ROOT - UNII:N9288CD508)	SANGUINARIA CANADENSIS ROOT	6 [hp_X]
CLAVICEPS PURPUREA SCLEROTIUM (UNII: 01G9XEA93N) (CLAVICEPS PURPUREA SCLEROTIUM - UNII:01G9XEA93N)	CLAVICEPS PURPUREA SCLEROTIUM	6 [hp_X]
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	30 [hp_X]

Inactive Ingredients	
Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	

Product Characteristics			
Color	white	Score	no score
Shape	ROUND	Size	3mm
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-8180-2	1350 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	09/01/2009	

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(48951-8180)