

SCORODITE SACCHARUM SPECIAL ORDER- scorodite saccharum special order 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.

Scorodite Saccharum Special Order

Directions: FOR ORAL USE.

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

Active Ingredients: Oxalis (Wood sorrel) 6X, Scorodite (Nat. ferric arsenate) 6X, Glandulae Parathyreoidea (Bovine parathyroid glands) 10X, Sacch. Sacchari (Cane sugar) 10X

Inactive Ingredients: Water, Salt, Lactose

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. Contains traces of lactose. 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 Made by Uriel, East Troy, WI 53120 www.urielpharmacy.com

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



**Scorodite
Saccharum
Special Order**

Homeopathic Ampules
net vol. 0.3 fl. oz (10 x 1 ml)

Scorodite Saccharum s.o.

SCORODITE SACCHARUM SPECIAL ORDER

scorodite saccharum special order liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-8170
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BOS TAURUS PARATHYROID GLAND (UNII: SMV71DDI4T) (BOS TAURUS PARATHYROID GLAND - UNII:SMV71DDI4T)	BOS TAURUS PARATHYROID GLAND	6 [hp_X] in 1 mL

OXALIS ACETOSELLA LEAF (UNII: U1W3U02EW0) (OXALIS ACETOSELLA LEAF - UNII:U1W3U02EW0)	OXALIS ACETOSELLA LEAF	6 [hp_X] in 1 mL
FERROUS ARSENATE (UNII: 129CO35H12) (FERROUS ARSENATE - UNII:129CO35H12)	FERROUS ARSENATE	6 [hp_X] in 1 mL
SUGARCANE (UNII: 81H2R5AOH3) (SUGARCANE - UNII:81H2R5AOH3)	SUGARCANE	10 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
LACTOSE (UNII: J2B2A4N98G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-8170-1	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(48951-8170)