EVERTOX SPECIAL BATH PREPARATIONS - silicon dioxide powder KMTR Co., Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Drug Facts

monzonite

silica compound

relieve atopic symptom (itching, inflammation and pain), relieve ring worm, relieve stress

keep out of reach of the children

apply on evertox per 100 liter approximately application part: whole body apply for 5~30 minutes preferably during bath keep cool and dry after use

do not open the case, in case the case opened by user fault, please keep the ingredients away immediately

for external use only

EVERTOX EVERTOX SPECIAL BATH PREPARATION

Drug Facts

Active Ingredients

Monzonite™ 80 wt%....... Purifier, Filter (FDA Medical device Class 1_NSF ANSI42_ DWTU)

- Effects
 - Relieve atopic symptom (itching, inflammation and pain), relieve ring worm,

Warning

Do not open the case, in case the case opened by users' fault, please keep the

Direction

- Apply One Evertox per 100 liter approximately application part Whole body apply for 5~30minute preferably during bath keep cool and dry after use

Other Information

- store between 1-45degree C (33.8 -113 degree F) avoid freezing and excessive heat above 60 degree C (140 degree F)

Inactive Ingredient

EVERTOX SPECIAL BATH PREPARATIONS

silicon dioxide powder

Product Information									
Product T ype	HUMAN OTC DRUG	Item Code (Sour	ce) NDC:694	NDC:69474-1001					
Route of Administration	TOPICAL								
Active Ingredient/Active Moiety									
Active Ingredient/Active N	Aoiety								
Active Ingredient/Active N	Aoiety Ingredient Name		Basis of Strength	Strength					
Active Ingredient/Active N SILICON DIO XIDE (UNII: ETJ7Z62	Ingredient Name	II:ETJ7Z6XBU4)	Basis of Strength SILICON DIOXIDE	Strength 20 g in 100 g					

		Ingredient Nam						
	Strength							
SILV	ER (UNII: 3M4G523V	V1G)						
SOD	IUM ALUMINIUM S	ILICATE (UNII: 058TS43PSM)						
Packaging								
#	Item Code	Package Description	Marketing Start Date Mark		rketing End Date			
1 NC	C:69474-1001-1	65 g in 1 BOTTLE						
Ma	rketing Infor	rmation						
Mar	keting Category	Application Number or Monograph Citation		Marketing Start Date		Marketing End Date		
unapr	proved drug other			0 1/11/20 15				
11	0							

Labeler - KMTR Co., Ltd. (688263356)

Registrant - KMTRCo., Ltd. (688263356)

Establishment

Name	Address	ID/FEI	Business Operations
KMTR Co., Ltd.		688263356	manufacture(69474-1001)

Revised: 1/2015

KMTR Co., Ltd.