CHAMOMILLA E RAD. 12 SPECIAL ORDER- chamomilla e rad. 12 special order 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.

Chamomilla e rad. 12 Special Order

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 Ingredient: Chamomilla e rad. 12X

Inactive Ingredient: Organic sucrose

Use: Temporary relief of cramps.

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



CHAMOMILLA E RAD. 12 SPECIAL ORDER chamomilla e rad. 12 special order pellet						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:48951-31	97	
Route of Administration	ORAL					
Active Ingredient/Active Moie	etv					
In	Basis of Strength		Strength			
MATRICARIA RECUTITA (UNII: G0 R4	MATRICA	RIA RECUTITA	12 [hp_X]			
Inactive Ingredients						
I	ngredient Name			Strength		

SUCROSE (UNII: C151H8 M554)								
Product Characteristics								
С	olor		white	Score		no score		
S	hape		ROUND	Size		3mm		
Fl	lavor			Imprint Code				
С	ontains							
_								
Packaging								
#	Item Code		Package Description		Marketing Star	t Marketing End		
	Item Code		- actinge 2 coort		Date	Date		
1	NDC:48951-3197-	1350 in 1 l Product	BOTTLE, GLASS; Type 0:	-	Date	Date		
1	NDC:48951-3197-		-	-		Date		
1	NDC:48951-3197-		-	-		Date		
1	NDC:48951-3197-	Product	BOTTLE, GLASS; Type 0:	•		Date		
N	NDC:48951-3197- 2	Product format	BOTTLE, GLASS; Type 0:	Not a Combination				
N	NDC:48951-3197- 2	Product formatory Approx	BOTTLE, GLASS; Type 0:	Not a Combination	09/01/2009			

Labeler - Uriel Pharmacy Inc. (043471163)

Establishment						
Name	Address	ID/FEI	Business Operations			
Uriel Pharmacy Inc.		043471163	manufacture(48951-3197)			
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Revised: 6/2018

Uriel Pharmacy Inc.