SUNSCREEN- zinc oxide sunscreen ointment
Private Label Select Ltd CO

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

SUNSCREEN

zinc oxide sunscreen ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62932-151
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	20 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
OLIVE OIL (UNII: 6UYK2W1W1E)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
COCONUT OIL (UNII: Q9L0O73W7L)	
CASTOR OIL (UNII: D5340Y2I9G)	
JOJOBA OIL (UNII: 724GKU717M)	
TOCOPHEROL (UNII: R0ZB2556P8)	
SHEA BUTTER (UNII: K49155WL9Y)	

Product Characteristics			
Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:62932-151-20	84 g in 1 TUBE; Type 0: Not a Combination Product	01/31/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	01/31/2016	

Labeler - Private Label Select Ltd CO (005415463)

Revised: 5/2017 Private Label Select Ltd CO