

**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.*

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**RCP Mineral Sunscreen SPF30**

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RECOVERY CARE PRODUCTS



*Mineral Sunscreen*

**Scar Protection**

**Broad Spectrum SPF 30**

net wt 0.5 oz / 14 g

**Drug Facts:**

Active Ingredients: Zinc Oxide 20%  
 Purpose: Sunscreen  
 Uses: Helps prevent sunburn

**Directions:**

Please adhere to your physician's protocol for this product\* · Wound must be fully closed and no longer scabbed · Gently glide over scar 15 minutes before sun exposure · Reapply: every two hours, after sweating and swimming · Children under 6 months: consult your physician · For optimal results, use in conjunction with RCP Scar Management\*

\*For more information on protocols visit:

[www.rcpcare.com/recoveryresources](http://www.rcpcare.com/recoveryresources)

USA Manufactured for and Distributed by:  
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[www.rcpcare.com](http://www.rcpcare.com)

**SUNSCREEN**

zinc oxide sunscreen ointment

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:62932-188
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	25 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
CARNAUBA WAX (UNII: R12CBM0EIZ)	
HELIANTHUS ANNUUS SEED WAX (UNII: 42DG15CHXV)	
TOCOPHEROL (UNII: R0ZB2556P8)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
CASTOR OIL (UNII: D5340Y2I9G)	
SHEA BUTTER (UNII: K49155WL9Y)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
COCONUT OIL (UNII: Q9L0O73W7L)	
CANDELILLA WAX (UNII: WL0328HX19)	
COCOA BUTTER (UNII: 512OYT1CRR)	

**Product Characteristics**

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62932-188-29	14 g in 1 TUBE; Type 0: Not a Combination Product	11/30/2018	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	11/30/2018	

**Labeler** - Private Label Select Ltd CO (005415463)**Registrant** - Private Label Select Ltd CO (005415463)