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.

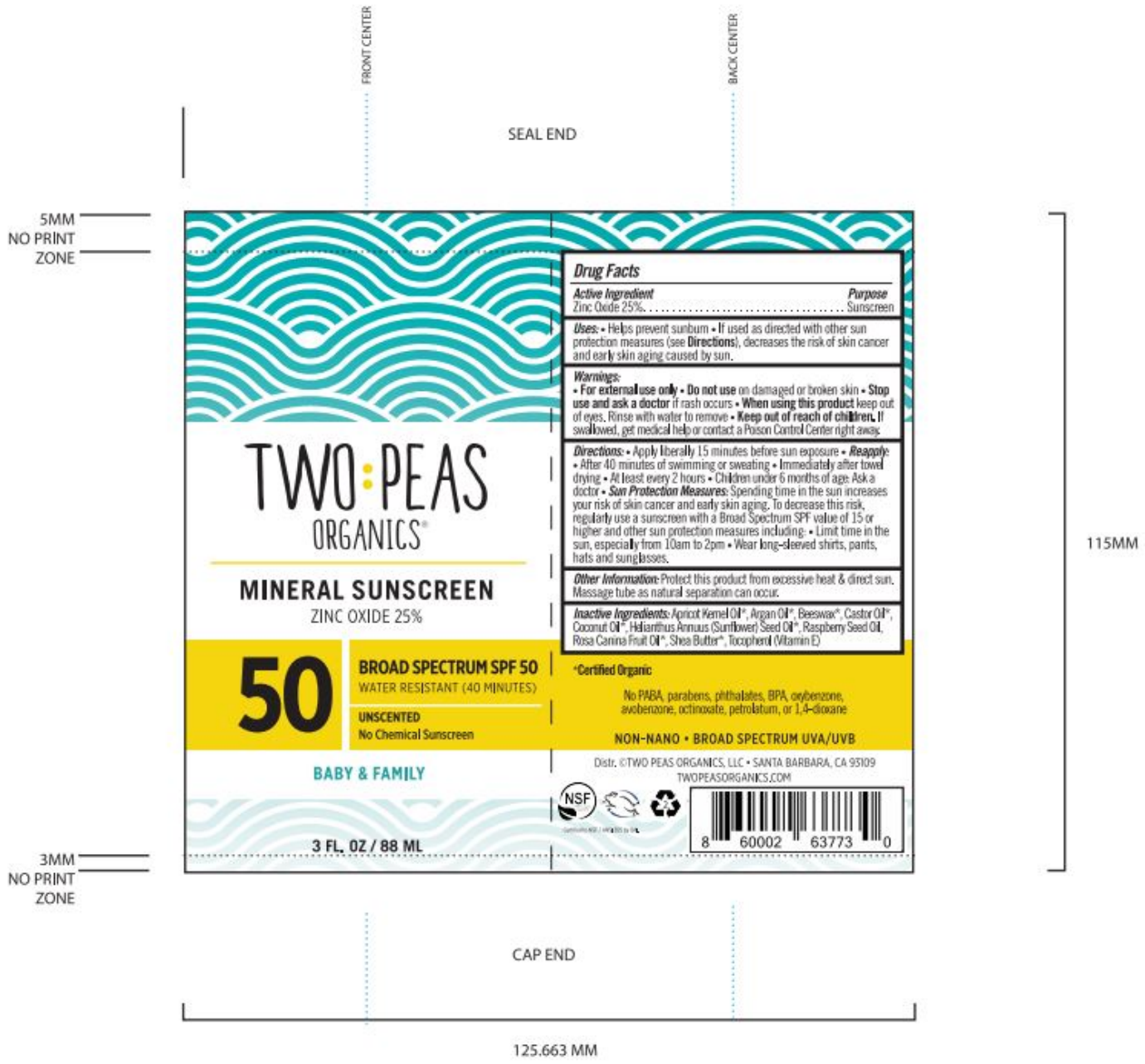
Two Peas Mineral Sunscreen, SPF 50

OTC - ACTIVE INGREDIENTS SECTION

OTC - PURPOSE SECTION

OTC - KEEP OUT OF REACH OF CHILDREN SECTION

ARTWORK



SUNSCREEN

zinc oxide sunscreen ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62932-253
Route of Administration	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
WHITE WAX (UNII: 7G1J5DA97F)	
.BETA.-TOCOPHEROL (UNII: 9U6A490501)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
CASTOR OIL (UNII: D5340Y2I9G)	
RASPBERRY SEED OIL (UNII: 9S8867952A)	
APRICOT KERNEL OIL (UNII: 54JB35T06A)	
.GAMMA.-TOCOPHEROL (UNII: 8EF1Z1238F)	
COCONUT OIL (UNII: Q9L0O73W7L)	
SHEA BUTTER (UNII: K49155WL9Y)	
ARGAN OIL (UNII: 4V59G5UW9X)	
ROSA CANINA SEED (UNII: 4503R1M9UT)	
.ALPHA.-TOCOPHEROL, D- (UNII: N9PR3490H9)	
.DELTA.-TOCOPHEROL (UNII: JU84X1II0N)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62932-253-47	85 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	01/01/2020	

Labeler - Private Label Select Ltd CO (005415463)

Revised: 1/2022

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