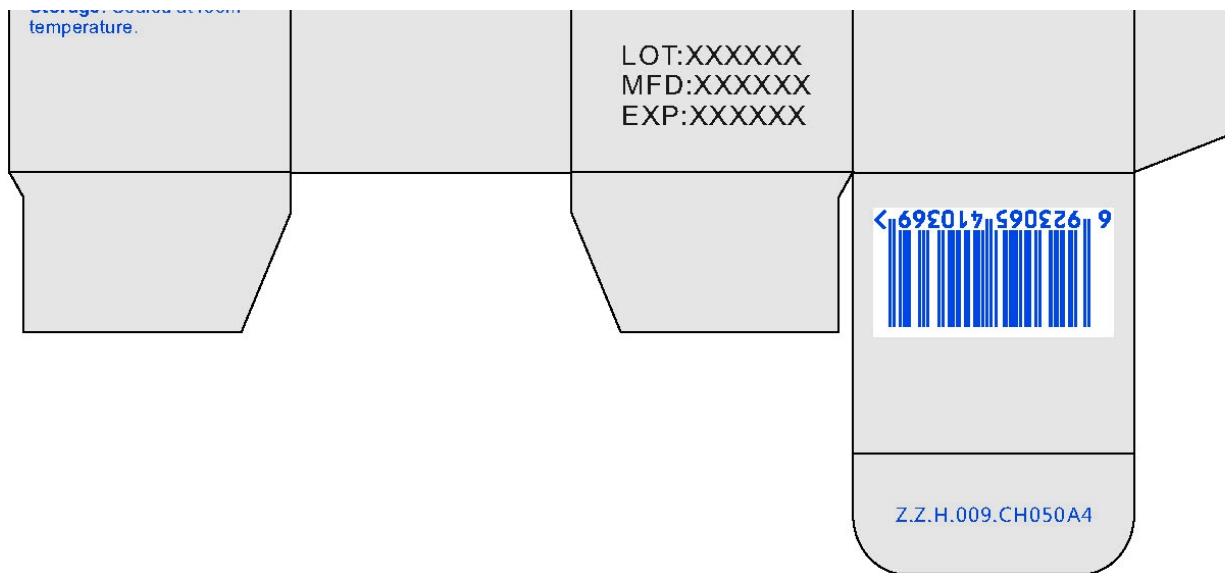


SUNBLOCK CREAM- zinc oxide, octisalate, titanium dioxide cream
Zhuhai Yasha Medical Instrument Co., Ltd.

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.

50g in a bottle NDC: 80376-001-01





SUNBLOCK CREAM

zinc oxide, octisalate, titanium dioxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80376-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTISALATE (UNII: 4X49 Y0596 W) (OCTISALATE - UNII:4X49 Y0596 W)	OCTISALATE	9 g in 100 g
TITANIUM DIOXIDE (UNII: 15FIX9 V2JP) (TITANIUM DIOXIDE - UNII:15FIX9 V2JP)	TITANIUM DIOXIDE	6 g in 100 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	3 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)	3 g in 100 g
WATER (UNII: 059QF0K00R)	45.8 g in 100 g
PHENOXYETHANOL (UNII: HIE492ZZ3T)	0.2 g in 100 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80376-001-01	50 g in 1 BOTTLE; Type 0: Not a Combination Product	08/31/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	08/31/2020	

