ECO 10 MINERAL SUN PRO PUFF TYPE - zinc oxide powder
MIZON 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.

Drug Facts

zinc oxide

Mica,Silica ,Triethoxycaprylylsilane,Iron Oxides,Dimethicone,Aluminum Hydroxide,Glycerine, Iron Oxides,Methicone,Bismuth Oxychloride,Iron Oxides,Hydrolyzed Algin,Laminaria Digitata Extract, Phormidium Persicinum Extract,Undaria Pinnatifida Extract ,Plankton extract,Phenethyl Alcohol , Chlorella vulgaris extract,Pancratium maritimum extract,Crithmum Maritimum Extract, Helichrysum italicum extract, Ulex europaeus leaf/root/stem extract

UVA/UVB protection

keep out of reach of the children

Apply it to your face after all of your basic skincare. If you need better UV protect effect, apply it more whenever and wherever needed

1. If following symptoms occur, stop use and consult a doctor : red spots, swelling, itcihng, irritation, or symtoms on applied skin under direct sunlight.
2. Do not use on scarred skin, or if you have dermatitis or eczema
3. Keep the cap close on its product
4. Keep away from direct sunlight or heat.
5. This product blocks UVA rays and ultraviolet protection effectiveness have been measured by the Food and Drug Administration methods and standards

for topical use only

ECO 10 MINERAL SUN PRO PUFF TYPE

zinc oxide powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:57718-040
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Zinc oxide (UNII: SOI2LOH54Z) (Zinc oxide - UNII:SOI2LOH54Z)	Zinc oxide	15.004 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
DIMETHICONE (UNII: 92RU3N3Y1O)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
GLYCERIN (UNII: PDC6A3C0OX)	
BISMUTH OXYCHLORIDE (UNII: 4ZR792I587)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57718-040-01	12 g in 1 CONTAINER		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	07/03/2014	

Labeler - MIZON CO., LTD. (557815570)

Registrant - MIZON CO., LTD. (557815570)

Establishment

Name	Address	ID/FEI	Business Operations
MIZON CO., LTD.		557815570	manufacture(57718-040)

Revised: 7/2014 MIZON CO., LTD.