#### AVEENO POSITIVELY MINERAL SENSITIVE SKIN SUNSCREEN BROAD SPECTRUM SPF 50- zinc oxide lotion Johnson & Johnson Consumer Inc.

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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## Aveeno Positively Mineral Sensitive Skin Sunscreen

#### Broad spectrum SPF 50

**Drug Facts** 

## **Active ingredients**

Zinc Oxide (21.6%)

#### Purpose

Sunscreen

#### Uses

- helps prevent sunburn
- if used as directed with other sun protection measures (see **Directions**), decreases the risk of skin cancer and early skin aging caused by the sun

#### Warnings

#### For external use only

Do not use on damaged or broken skin

#### When using this product

• keep out of eyes. Rinse with water to remove.

#### Stop use and ask a doctor if rash occurs

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

#### Directions

- apply generously and evenly 15 minutes before sun exposure
- reapply:
  - after 80 minutes of swimming or sweating
  - immediately after towel drying
  - at least every 2 hours

- **Sun Protection Measures.** Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a Broad Spectrum SPF value of 15 or higher and other sun protection measures including:
  - limit time in the sun, especially from 10 a.m. 2 p.m.
  - wear long-sleeved shirts, pants, hats, and sunglasses
- children under 6 months of age: Ask a doctor

### Other information

- protect this product from excessive heat and direct sun
- may stain some fabrics

## Inactive ingredients

Water, C12-15 Alkyl Benzoate, Dimethicone, Glycerin, Phenoxyethanol, Phenyl Trimethicone, Styrene/Acrylates copolymer, Octyl dodecyl Citrate Crosspolymer, Cetyl PEG/PPG-10/1 Dimethicone, Polyhydroxystearic acid, Silica, Ethyl methicone,Cetyl Dimethicone, Triethoxycaprylylsilane, Glyceryl Behenate, Sodium Chloride,Acrylates/Dimethicone copolymer, Chlorphenesin, Phenethyl alcohol, Avena Sativa (Oat) Kernel flour, Caprylyl Glycol,Cetyl Dimethicone/Bis-Vinyldimethicone Crosspolymer, Chrysanthemum Parthenium (Feverfew) Flower/Leaf/Stem juice

## Questions?

886-428-3366; Outside US, dial collect 215-273-8755 www.aveeno.com

Distributed by: JOHNSON & JOHNSON CONSUMER INC. Skillman, NJ 08558

## PRINCIPAL DISPLAY PANEL - 88 mL Tube Label

DERMATOLOGIST RECOMMENDED

Aveeno ® POSITIVELY MINERAL™

sensitive skin sunscreen

**BROAD SPECTRUM SPF 50** 

naturally-sourced 100% zinc oxide active ingredient sweat + water resistant (80 min) 3.0 fl. oz (88 mL)



## AVEENO POSITIVELY MINERAL SENSITIVE SKIN SUNSCREEN BROAD SPECTRUM SPF 50

zinc oxide lotion

Product Information					
Product Type	HUMAN OTC DRUG	Item Cod	e (Source)	NDC:69968	-0395
Route of Administration	TOPICAL				
Active Ingredient/Active	e Moiety				
Ingred	lient Name		Basis of Streng	gth St	rength
ZINC OXIDE (UNII: SOI2LOH54Z)	(ZINC OXIDE - UNII:SOI2LOF	154Z)	ZINC OXIDE	216 m	g in 1 mL
Inactive Ingredients					
	Ingredient Nar	ne			Strength
WATER (UNII: 059QF0KO0R)					
ALKYL (C12-15) BENZOATE (UI	NII: A9EJ3J61HQ)				
DIMETHICONE (UNII: 92RU3N3Y1	.0)				
GLYCERIN (UNII: PDC6A3C0OX)					
PHENOXYETHANOL (UNII: HIE49	2ZZ3T)				
PHENYL TRIMETHICONE (UNII: I	DR0K5NOJ4R)				
BUTYL METHACRYLATE/METHY (UNII: V5RS026Q0H)	L METHACRYLATE/METH	ACRYLIC AC	ID/STYRENE CROSS	POLYMER	
OCTYLDODECYL CITRATE CRO	SSPOLYMER (UNII: X323T6	QO4M)			
CETYL PEG/PPG-10/1 DIMETHIC	CONE (HLB 5) (UNII: 035JKJ	76MT)			
POLYHYDROXYSTEARIC ACID (	2300 MW) (UNII: YXH47AO	JOF)			
SILICON DIOXIDE (UNII: ETJ7Z6)					
ETHYL METHICONE (8 MPA.S)					
CETYL DIMETHICONE 25 (UNII:					
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GLYCERYL DIBEHENATE (UNII: F	•				
SODIUM CHLORIDE (UNII: 451W	. ,				
CHLORPHENESIN (UNII: 1670DAL					
<b>PHENYLETHYL ALCOHOL</b> (UNII: <b>OATMEAL</b> (UNII: 8PI54V663Y)	ML9LGA/408)				
CAPRYLYL GLYCOL (UNII: 00YIU	543811)				
CETYL DIMETHICONE/BIS-VINY					
FEVERFEW (UNII: Z64FK7P217)					
Deckening					
Packaging					

# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:69968- 0395-3	88 mL in 1 TUBE; Type 0: Not a Combination Product	10/01/2018	11/27/2024
<b>2</b> NDC:69968- 0395-1	14 mL in 1 TUBE; Type 0: Not a Combination Product	10/01/2018	01/12/2023

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part352	10/01/2018	11/27/2024			

# Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 5/2023

Johnson & Johnson Consumer Inc.