

## **ZINC OXIDE - zinc oxide ointment**

**RiteAid**

*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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### **drug facts**

<b>Active ingredient</b>	<b>Purpose</b>
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Zinc Oxide.....	Skin protectant
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### **Uses**

- dries the oozing and weeping od poison - ivy - oak - sumac
- helps treat and prevent diaper rash
- protects chafed skin associated with diaper rash and helps protect from wetness

### **Keep out of the reach of children.**

If swallowed, get medical help or contact a Poison Control Center right away.

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### **Warnings**

**For external use only.**

**When using this product** do not get into eyes

### **Stop use and ask a doctor if:**

- **condition worsens - symptoms last more than 7 days or clear up and occur again within a few days.**

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

### **Directions**

**- apply as needed**

For diaper rash

- change wet and soiled diapers promptly
- cleanse the diaper area, and allow to dry
- apply liberally as often as necessary, with each diaper change, especially at bedtime or anytime when exposure to wet diapers may be prolonged

**- children under 12 years**

ask a doctor

**Other information**

Store at 20 degrees to 25 degrees C (68 degrees to 77 degrees F)

**Inactive Ingredients**

beeswax, mineral oil, petrolatum



## ZINC OXIDE

zinc oxide ointment

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-0081
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	200 mg in 1 g

## Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	
WHITE WAX (UNII: 7G1J5DA97F)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-0081-3	57 g in 1 TUBE		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	05/19/2009	

**Labeler** - RiteAid (014578892)

**Registrant** - Pharma Pac, LLC (140807475)

## Establishment

Name	Address	ID/FEI	Business Operations
Pharma Pac, LLC		140807475	manufacture

Revised: 2/2011

Rite Aid