# ZINC OXIDE 20%- zinc oxide ointment Trifecta Pharmaceuticals USA

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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#### Zinc Oxide Ointment 20%

#### **Drug Facts**

#### **Active Ingredient**

Zinc Oxide 20%

#### Purpose

Skin Protectant

#### Uses

- Protects skin
- Protects and releives chafed skin due to diaper rash helps seal out wetness
- Dries the oozing and weeping of poison ivy, oak and sumac

# Keep out of the reach of children

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

# Warnings

# For External Use Only

- When using this product avoid contact with eyes
- Stop use and ask a doctor if condition worsens or does not improve within 7 days. This may be a sign of a serious condition.

# Directions

- Apply ointment liberally as often as necessary
- For the treatment of diaper rash, change wet and soiled diapers promptly
- Cleanse the diaper area and allow to dry
- Apply with each diaper change, especially at bedtime or anytime when exposure to wet diaper may be prolonged

# **Other Information**

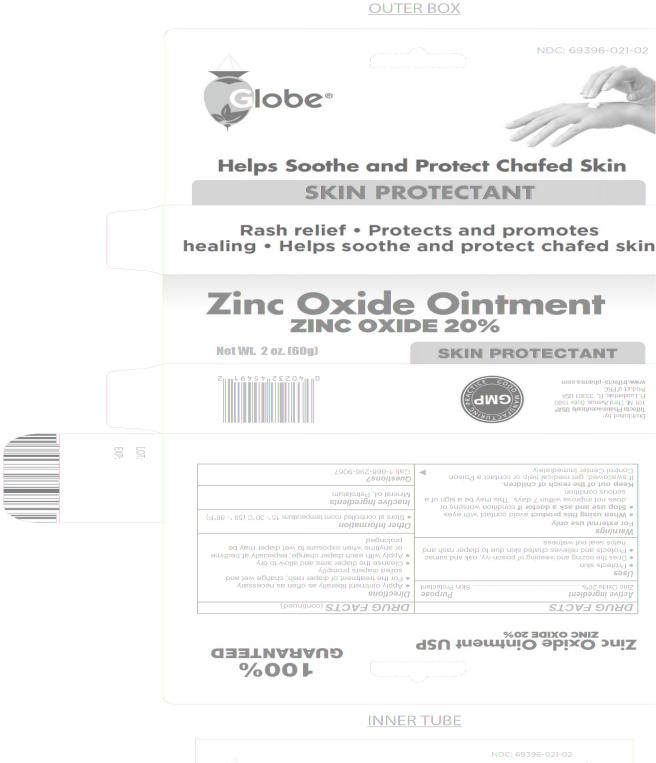
• Store at controlled room temperature 15° - 30°C (59° - 86°F)

# **Inactive Ingredients**

Mineral Oil, Petrolatum

# **Distributed By:**

Trifecta Pharmaceuticals USA, LLC. 101 NE Third Avenue, Ste. 1500 Ft. Lauderdale, FL. 33301, USA. Product of PRC www.trifecta-pharma.com





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Zinc Oxide 20% Skin Protectant	Center immediately.
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condition.	Questions? C coll 1-886-206-0067

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Pı	oduct Informati	ion					
Product Type			HUMAN OTC DRUG	Item Code (Source)		NDC:69396-021	
Ro	ute of Administrat	ion	TOPICAL				
Ac	tive Ingredient/	Active Moi	ety				
Ingredient Name				<b>Basis of Strength</b>		Strength	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH5			C OXIDE - UNII:SOI2LOH54Z)		ZINC OXIDE		0.2 g in 1 g
In	active Ingredier	nts					
Ingredient Name						Strength	
			Ingredient Name			Str	ength
MI	NERAL OIL (UNII: T	5L8T28FGP)	Ingredient Name			Stro	ength
	NERAL OIL (UNII: T Irolatum (UNII: 4		Ingredient Name			Stro	ength
PE'	<b>FROLATUM</b> (UNII: 4		Ingredient Name			Stro	eng th
РЕ <sup>г</sup> Ра	rrolatum (unii: 4 ckaging	IT6 H12 B N9 U)		Marke	ting Start Date		
PE' Pa #	rrolatum (UNII: 4 ckaging Item Code	IT6 H12 B N9 U)	Ingredient Name Package Description	<b>Marke</b> 01/26/20	ting Start Date		
РЕ Ра #	rrolatum (UNII: 4 ckaging Item Code	1 in 1 BOX		01/26/20	•		
РЕ Ра #	rrolatum (UNII: 4 ckaging Item Code	1 in 1 BOX	Package Description	01/26/20	•		
PE' Pa # 1 1	rrolatum (UNII: 4 ckaging Item Code	1 in 1 BOX 60 g in 1 TUBE	Package Description	01/26/20	•		
PE Pa 1 1 1	rrolatum (UNII: 4 ckaging Item Code NDC:69396-021-02	1 in 1 BOX 60 g in 1 TUBE <b>rmation</b>	Package Description	0 1/26 /20 o duct	•	Market	ength ting End Date ting End Date

Labeler - Trifecta Pharmaceuticals USA (079424163)

Revised: 1/2018