CRITIC AID SKIN- zinc oxide paste
Coloplast Manufacturing US, LLC

Critic-Aid® Skin Paste

Drug Facts

Active ingredients

Zinc Oxide: 20%

Purpose

Skin Protectant

Uses

- Helps protect minor skin irritation due to diaper rash
- Protects chafed skin due to diaper rash
- Helps seal out wetness

Warnings

For external use only.

When using this product:

- avoid contact with eyes

Consult a doctor:

- if condition worsens or does not improve within 7 days.

Do not use on:

- deep or puncture wounds
- animal bites
- serious burns

Keep out of reach of children.

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

Directions

- Clean the affected area and dry thoroughly
- Apply a thin layer of product over affected area twice daily (morning and night), or as directed by a doctor.
- Change wet and soiled absorbent brief/diapers promptly.

Inactive ingredients

Cellulose Gum (CMC), Dimethicone, Petrolatum

Questions or comments?

Call toll free 1-800-533-0464

Manufactured by: Coloplast A/S
Holtedam 1, DK-3050 Humlebaek, Denmark

Distributed by: Coloplast Corp
1601 W River Rd. N, Minneapolis, MN 55411 USA

PRINCIPAL DISPLAY PANEL - 170 g Tube Label

NDC 11701-050-32

Critic-Aid® Skin Paste

Skin Protectant

Thick Moisture
Barrier Paste

For Minor to
Severe Skin Irritation
Due to Incontinence

Product #1947

Coloplast

NET WT. 6 OZ. (170 g)

NDC 11701-050-32

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SCP SAMPLE
12345 67890 5

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See crimp for lot no. and expiration date
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Distributed by: Coloplast Corp
1601 W River Rd. N, Minneapolis, MN 55411 USA
www.coloplast.us
Product #1947 ©2014-10, Coloplast Corp
Made in the USA

K14-667

CRITIC AID SKIN

zinc oxide paste

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11701-050
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)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11701-050-33	71 g in 1 TUBE; Type 0: Not a Combination Product	06/15/2009	
2	NDC:11701-050-32	170 g in 1 TUBE; Type 0: Not a Combination Product	06/15/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M015	06/15/2009	

Labeler - Coloplast Manufacturing US, LLC (110326675)

Registrant - Coloplast Corp (847436391)

Establishment

Name	Address	ID/FEI	Business Operations
Coloplast Manufacturing US, LLC		110326675	MANUFACTURE(11701-050)

Revised: 12/2023 Coloplast Manufacturing US, LLC