

BAZA CLEAR- petrolatum ointment
Coloplast Manufacturing US, LLC

Baza® Clear
Skin Protectant
Moisture Barrier Ointment

For Skin Irritation Due to Incontinence

Drug Facts

Active ingredient

Petrolatum, 96%

Purpose

Skin Protectant

Uses Helps treat and prevent minor skin irritation due to diaper rash. Helps prevent and temporarily protects chafed, chapped or cracked skin. Helps seal out wetness.

Warnings

When using this product

- avoid contact with eyes
- not to be applied over deep or puncture wounds or lacerations.

Consult a doctor

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

For external use only.

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

Directions

- apply liberally as often as necessary.

For Diaper Rash: cleanse the diaper area and allow to dry

- apply liberally as often as necessary, with each diaper change especially at bedtime or any time when exposure to wet diapers may be prolonged.

Inactive ingredients benzethonium chloride, BHT, cod liver oil, fragrance, green 6, sodium caseinate (CONTAINS MILK)

See crimp for lot no. and expiration date

Manufactured by: Coloplast A/S

DK-3050 Humlebaek, Denmark

Distributed by: Coloplast Corp.

Minneapolis, MN 55411 U.S.A.

1-800-533-0464 www.us.coloplast.com

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Made in the U.S.A.

G8-303

PRINCIPAL DISPLAY PANEL - NET WT. 5 OZ. (142g)

NDC 11701-048-014

Baza[®] Clear

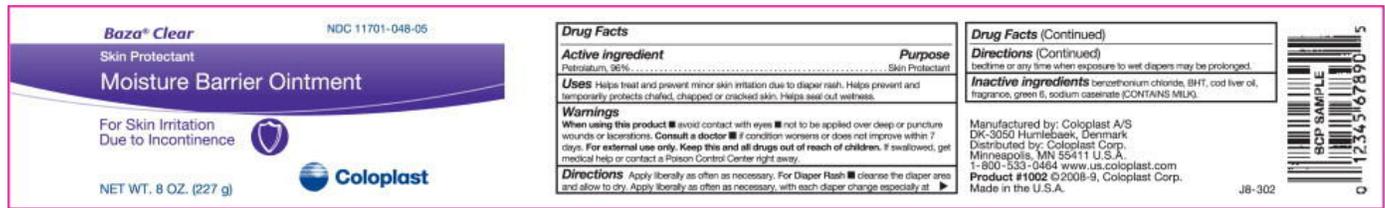
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3-11701-04805-6

BAZA CLEAR				
petrolatum ointment				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11701-048	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	960 mg in 1 g	
Inactive Ingredients				
	Ingredient Name	Strength		
	COD LIVER OIL (UNII: BBL281NWFG)			
	SODIUM CASEINATE (UNII: 7473P66J9E)			
	BENZETHONIUM CHLORIDE (UNII: PH41D05744)			
	BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)			
	D&C GREEN NO. 6 (UNII: 4QP5U84YF7)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:11701-048-22	4 g in 1 PACKET; Type 0: Not a Combination Product	06/15/2009	
2	NDC:11701-048-24	50 g in 1 TUBE; Type 0: Not a Combination Product	06/15/2009	
3	NDC:11701-048-14	142 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	M016	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(11701-048)

Revised: 12/2023

Coloplast Manufacturing US, LLC