DG BABY DIAPER RASH- zinc oxide ointment
DOLGENCORP 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.

DRUG FACTS

ACTIVE INGREDIENT

ZINC OXIDE 40%

PURPOSE

SKIN PROTECTANT

USES

- HELPS TREAT AND PREVENT 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. IF CONTACT OCCURS, RINSE THOROUGHLY WITH WATER

STOP USING THIS PRODUCT AND ASK A DOCTOR IF

- CONDITION WORSENS
- SYMPTOMS LAST MORE THAN 7 DAYS OR CLEAR UP AND OCCUR AGAIN WITHIN A FEW DAYS

DO NOT USE ON

- DEEP PUNCTURE WOUNDS
- ANIMAL BITES
- SERIOUS BURNS

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY

DIRECTIONS

- CHANGE WET AND SOILED DIAPERS PROMPTLY
- CLEAN THE DIAPER AREA AND ALLOW TO DRY
- APPLY OINTMENT LIBERALLY AS OFTEN AS REQUIRED WITH EACH DIAPER CHANGE, ESPECIALLY AT BEDTIME OR ANYTIME WHEN EXPOSURE TO WET DIAPERS MAY BE PROLONGED.

OTHER INFORMATION

STORE BETWEEN 20°-25°C (68°-77°F)

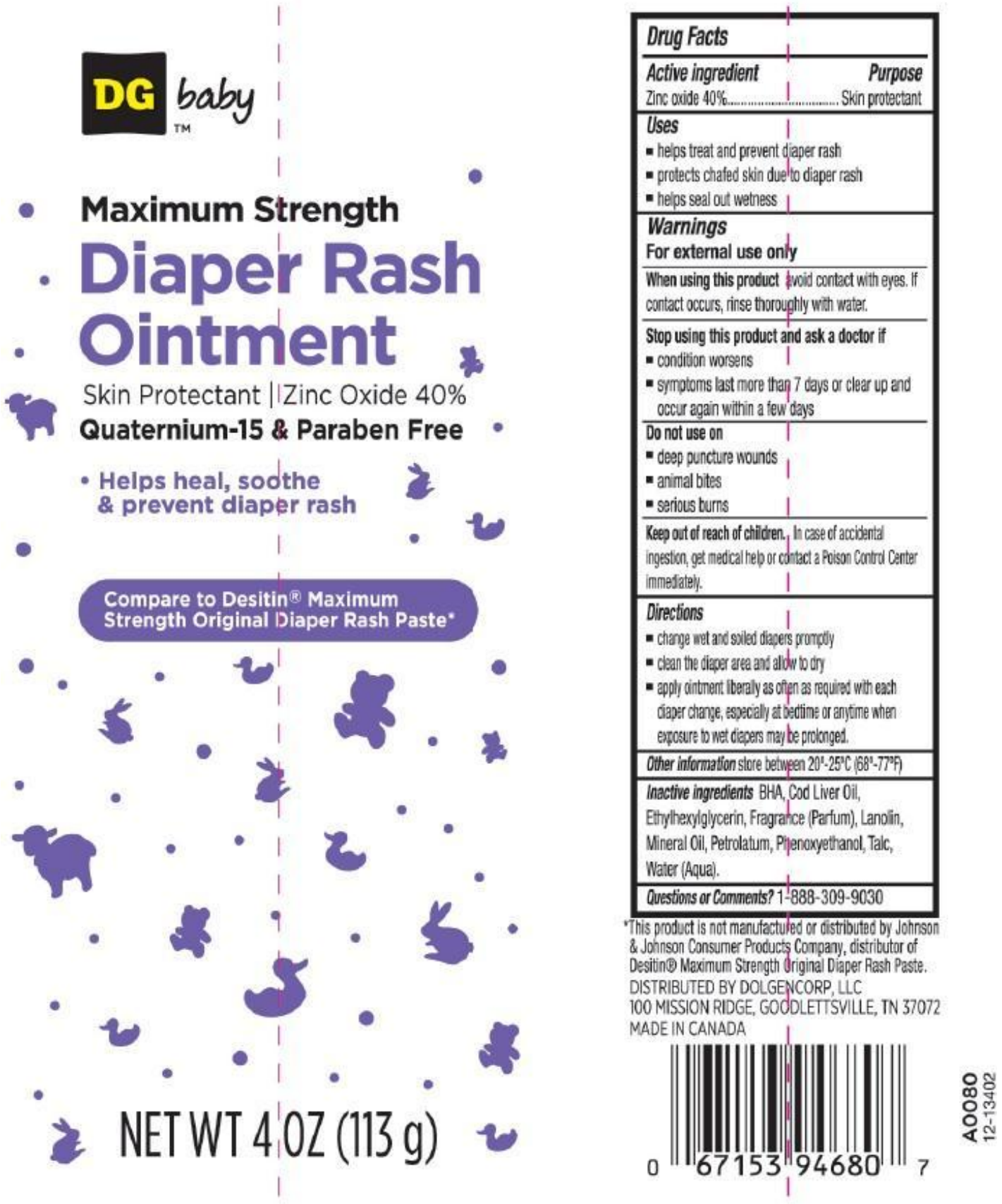
INACTIVE INGREDIENTS

BHA, COD LIVER OIL, ETHYLHEXYLGLYCERIN, FRAGRANCE (PARFUM), LANOLIN, MINERAL OIL, PETROLATUM, PHENOXYETHANOL, TALC, WATER (AQUA)

QUESTIONS OR COMMENTS?

1-888-309-9030

LABEL COPY



DG BABY DIAPER RASH

zinc oxide ointment

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-334
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	40 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)	
COD LIVER OIL (UNII: BBL281NWFG)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
LANOLIN OIL (UNII: OVV5IIJ58F)	
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TALC (UNII: 7SEV7J4R1U)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-334-04	113 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	07/01/2015	

Labeler - DOLGENCORP INC (068331990)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(55910-334)

Revised: 7/2015 DOLGENCORP INC