BABYSONS- dexpanthenol ointment Laboratorios Quimica Son's, S.A. de C.V

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

BABYSONS

Drug Facts

Active ingredient

Dexpanthenol 5%

Purpose

Skin Protectant

Uses

- Helps treat and prevent diaper rash
- Protects chafed skin due to diaper rash and helps seal out wetness

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 reach of children.

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

Directions

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

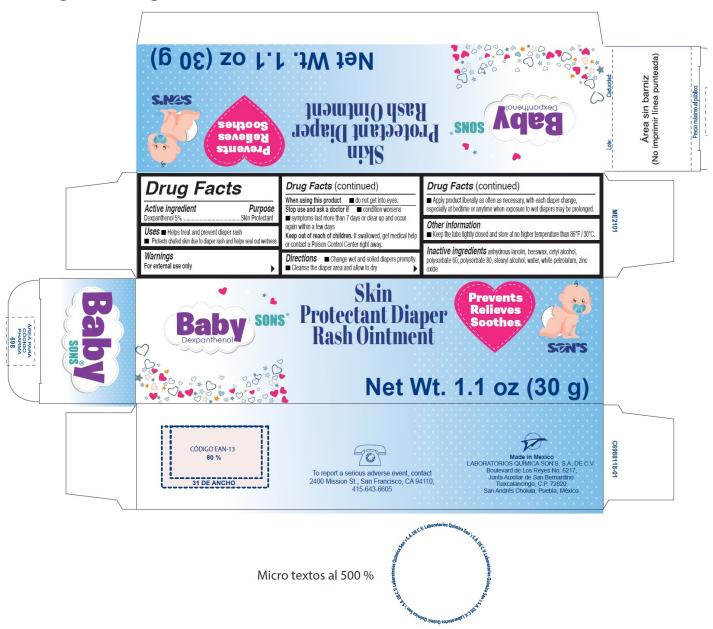
Other information

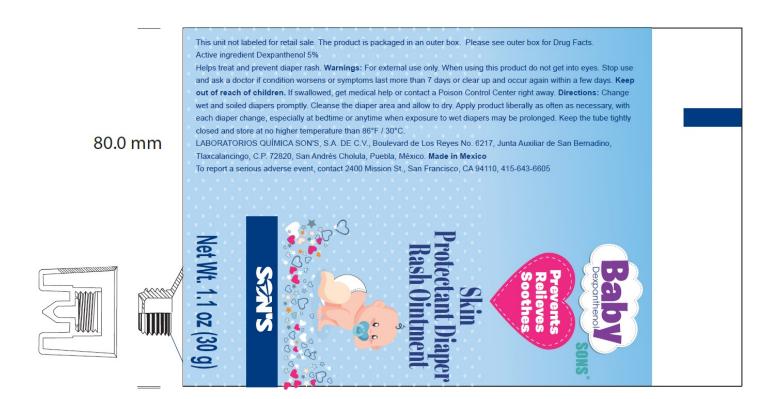
Keep the tube tightly closed and store at no higher temperature than 86°F / 30°C.

Inactive ingredients

anhydrous lanolin, beeswax, cetyl alcohol, polysorbate 60, polysorbate 80, stearyl alcohol, water, white petrolatum, zinc oxide

Package Labeling:





BABYSONS

dexpanthenol ointment

Product Information	n
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:73519-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

DEXPANTHENOL (UNII: 106C93RI7Z) (DEXPANTHENOL - UNII:106C93RI7Z) DEXPANTHENOL 50 mg in 1 g

Inactive Ingredients Ingredient Name Strength LANOLIN (UNII: 7EV65EAW6H) YELLOW WAX (UNII: 2ZA36H0S2V) CETYL ALCOHOL (UNII: 936JST6JCN) POLYSORBATE 60 (UNII: CAL22UVI4M) POLYSORBATE 80 (UNII: 6OZP39ZG8H) STEARYL ALCOHOL (UNII: 2KR89I4H1Y) WATER (UNII: 059QF0KO0R) PETROLATUM (UNII: 4T6H12BN9U) ZINC OXIDE (UNII: SOI2LOH54Z)

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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73519-001- 30	1 in 1 BOX	06/15/2020	
1		30 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	
unapproved drug other		06/15/2020		

Labeler - Laboratorios Quimica Son's, S.A. de C.V (815603378)

Establishment				
Name	Address	ID/FEI	Business Operations	
Laboratorios Quimica Son's, S.A. de C.V		815603378	manufacture(73519-001)	

Revised: 10/2024 Laboratorios Quimica Son's, S.A. de C.V