BABYO2- colloidal silver and glycerin solution
Oxigenesis, Inc.

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.

BabyO2™

Active ingredients	**Purpose**
Colloidal Silver 2 mg	Skin Protectant
Glycerin (Plant) 2 mg.....	Skin Protectant

Uses

Relieves and prevents the discomfort and symptoms from: diaper rash, chaffed skin, minor burns and abrasions.

Warnings

For external use only.

Do not use over puncture wounds, infections or lacerations.

Avoid contact with the eyes.

Stop use and ask a doctor if condition worsens or does not improve in 7 days.

Keep out of reach of children.

Directions

Change wet and solid diapers promptly. Cleanse diaper area and dry. Apply ointment liberally as often necessary with every diaper change, especially at bedtime.

Other information

Sstore at 20°-25°C (68° -70° F)

Inactive ingredients

Distilled water, Sodium Chloride, Hydroxypropyl Starch Phosphate.

Marketed & Distributed by:

Oxigenesis, Inc.
2917 Union Road, Suite B
Paso Robles, CA 93446 USA
805-549-0275

BABYO2

colloidal silver and glycerin solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72136-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SILVER (UNII: 3M4G523W1G) (SILVER - UNII:3M4G523W1G)	SILVER	2 mg in 240 mL

GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	2 mg in 240 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE NA-22 (UNII: VMP9781061)	
HYDROXYPROPYL CORN STARCH (5% SUBSTITUTION BY WEIGHT) (UNII: 9M44R3409A)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72136-005-01	240 mL in 1 TUBE; Type 0: Not a Combination Product	02/26/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		02/26/2019	

Labeler - Oxigenesis, Inc. (006774725)

Registrant - Oxigenesis, Inc. (006774725)

Establishment

Name	Address	ID/FEI	Business Operations
Oxigenesis, Inc.		006774725	manufacture(72136-005)

Revised: 2/2019 Oxigenesis, Inc.