CRC SUNSCREEN TOWEL CRC- sunscreen towel cloth
CRC Industries, 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.

Active Ingredient	*Purpose*
Homosalate (2%)	Sunscreen
Octinoxate (7.5%)	Sunscreen
Octisalate (4%)	Sunscreen
Oxybenzone (5%)	Sunscreen

Uses

- Provides high protection against sunburn.
- Retains SPF after 80 minutes of sweating or activity in the water.

Warnings

For external use only.

Flammable. Keep away from heat and flame when applying this product.

When using this product

- Keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor

- If rash or irritation develops and lasts.

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

Directions

- Unfold towel.
- Apply evenly before sun exposure.
- For children under 6 months, consult a doctor.
- Reapply after towel drying, swimming or sweating.

Inactive Ingredients

Acrylates/Octylacrylamide Copolymer, Ethanol, Fragrance

Towel

Water/Sweat Resistant
Greaseless
UVA/UVB Protection
Aminobenzoic Acid (PABA)-free

SPF 30+

Contains 1:
8" x 10" (20.3 cm x 25.4 cm) Premoistened Towel

See precautionary statements on reverse side.

NO. 04105

Sun Alert: Limiting sun exposure, wearing protective clothing and using sunscreens may reduce the risks of skin aging, skin cancer, and other harmful effects of the sun.

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For Additional Information, Read the Material Safety Data Sheet for this Product.

CRC

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Warminster, PA 18974
www.crcindustries.com
Made in U.S.A. 10A

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CRC SUNSCREEN TOWEL CRC

sunscreen towel cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66348-4105
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HOMOSALATE (UNII: V06SV4M95S) (SALICYLIC ACID - UNII:O414PZ4LPZ)	HOMOSALATE	20 g in 856 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (2-ETHYLHEXYL 4-PHENYLBENZOPHENONE-2'-CARBOXYLATE - UNII:93NOD9WBCS)	OCTINOXATE	75 g in 856 mL
OCTISALATE (UNII: 4X49Y0596W) (SALICYLIC ACID - UNII:O414PZ4LPZ)	OCTISALATE	40 g in 856 mL

OXYBENZONE (UNII: 95OOS7VE0Y) (BENZOPHENONE - UNII:701M4TTV9O)	OXYBENZONE	50 g in 856 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	80.1 g in 856 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66348-4105-1	9 mL in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	03/01/2010	

Labeler - CRC Industries, Inc. (069880029)

Registrant - ITW Dymon (103307604)

Establishment

Name	Address	ID/FEI	Business Operations
ITW Dymon		103307604	manufacture

Revised: 3/2010 CRC Industries, Inc.