TOTALLY FUN AFTER SUN- lidocaine hcl gel
MJ Products Association, LLC

Totally Fun After Sun

Drug Facts

Active Ingredients

Lidocaine HCL 1.50%

PURPOSE

Topical Anesthetic

Indications

For the Temporary relief of pain and itching due to sunburn, minor skin irritations, minor cuts, minor burns, scrapes and insect bites.

WARNINGS

FOR EXTERNAL USE ONLY

Do not use

- in large quantites, particularly over raw or blistered areas.
- Avoid contact with eyes.

Stop use and ask a doctor

- if condition gets worse, symptoms last for more than 7 days, or symptoms clear up and occur again with a few days.
- If sumptoms persist for more than seven days, discontinue use and consult physician.

Keep out of reach of children.

If swallowed, consult physician.

Directions

- Adults and children two-years of age or older: Apply to affected area not more than three to four times daily.
- Children under two-years of age: consult a physician.

Additional information

Store at room temperature.

Other Ingredients

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Carbomer, Ethylhexylglycerin, : Polyethylene Terephthalate, Polymethyl Methacrylate, Polyurethan-33, FD&C Yellow #5, D&C Red #7; Parfum (Orange Fragrance), Phenoxyethanol, Terephthalate, Triethanolamine. *Glitter*

Questions or comments?

Contact us directly at 1-888-601-0441

Distr. by/par: Sunshine & Glitter

MJ Products Association LLC

9750 NW 17th Street, Miami, FL 33172

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Package Labeling

TOTALLY FUN AFTER SUN

lidocaine hcl gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72085-145
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	15 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
POLY(METHYL METHACRYLATE; 450000 MW) (UNII: Z47NNT4J11)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
D&C RED NO. 7 (UNII: ECW0LZ41X8)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72085-145-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/01/2018	

Labeler - MJ Products Association, LLC (081022202)

Revised: 11/2023 MJ Products Association, LLC