

SUNBURNT ADVANCED AFTER-SUN- calendula officinalis, echinacea angustifolia, cantharis vesicatoria gel

Welmedix LLC

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

SunBurnt® Advanced After-Sun Gel

Drug Facts

<i>Active ingredients</i>	<i>Purposes</i>
Each dose contains:	
8 parts Cantharis vesicatoria 3X HPUS	Sunburns, rawness
1 part Calendula officinalis TINC HPUS	Promotes skin healing
1 part Echinacea angustifolia TINC HPUS	Itching, burning, redness

The letters 'HPUS' indicate that the active ingredients are officially monographed in the Homeopathic Pharmacopoeia of the United States. TINC and 3X are homeopathic dilutions: see www.sunburnt.com for details.

Uses

Temporarily relieves symptoms of mild and moderate sunburn such as:

- overheated skin
- redness
- rawness
- itching
- dryness

Product Uses are based on the Homeopathic Materia Medica. These Uses have not been evaluated by the Food and Drug Administration.

Warnings

For external use only

Do not use

- on severe sunburn
- on broken skin
- if you have a rash
- if you have an allergy or hypersensitivity to any ingredients

When using this product

- avoid contact with eyes or mucous membranes

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

If pregnant or breast feeding, ask a health professional before use.

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

Directions

- for best results, clean skin and apply immediately to affected area
- adults and children 2 years of age and older: apply 2-3 times a day, as needed, or as directed by your doctor
- children under 2 years of age: consult a doctor

Other information

- store at 15-30°C (59-86°F)
- do not use if tamper-evident seal under cap is compromised or missing

Inactive ingredients

aloe barbadensis leaf juice, carbomer, ethyl alcohol, panthenol, phenoxyethanol, sodium hyaluronate, sodium hydroxide, water

Questions or Comments?

1-888-565-2876 Monday through Friday, 9am-5pm EST

Dist. by: **Welmedix Consumer Healthcare**
Princeton, New Jersey 08540

PRINCIPAL DISPLAY PANEL - 170 g Tube Carton

SUN

BURNT®

ADVANCED SUN RECOVERY

AFTER-SUN GEL

MUCH MORE THAN ALOE®

Immediately cools and rehydrates skin

Soothing relief for sunburned skin

*Helps reduce the appearance
of redness & peeling*

INSTANTLY

COOLING

ULTRA

HYDRATING

NON-STICKY

FORMULA

HOMEOPATHIC

Net Wt 6 oz (170 g)



**SUN
BURNT.**

ADVANCED SUN RECOVERY



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AFTER-SUN GEL

MUCH MORE THAN ALOE®

Immediately cools and rehydrates skin

Soothing relief for sunburned skin

Helps reduce the appearance

**INGREDIENTS
WE AVOID USING
IN SUNBURNT GEL**

NO
SULFATES

NO
PARABENS

NO
GLYCOLS

NO
PHTHALATES

NO
DYES

NO
FRAGRANCE



of redness & peeling

INSTANTLY COOLING



ULTRA HYDRATING

NON-STICKY FORMULA

HOMEOPATHIC

Net Wt 6 oz (170 g)



SUN BURNT

ADVANCED SUN RECOVERY

MUCH MORE THAN ALOE®

SunBurnt's non-sticky formula blends natural actives + aloe for soothing relief

MUCH MORE THAN ALOE®

SunBurnt® After-Sun Gel combines a unique, non-sticky blend of actives sourced from Mother Nature + aloe to provide soothing relief for mild & moderate sunburn.



CALENDULA



ECHINACEA



CANTHARIS

+



ALOE



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U.S. Patent Nos. 7,959,955 and 8,221,802

Money Back Guarantee
 For more details, visit sunburnt.com

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Dist. by: **Welmedix Consumer Healthcare**
 Princeton, New Jersey 08540
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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:24330-210
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD) (CALENDULA OFFICINALIS FLOWER - UNII:P0M7O4Y7YD)	CALENDULA OFFICINALIS FLOWER	1 [hp_Q] in 0.1 g
ECHINACEA ANGUSTIFOLIA (UNII: VB06AV5US8) (ECHINACEA ANGUSTIFOLIA - UNII:VB06AV5US8)	ECHINACEA ANGUSTIFOLIA	1 [hp_Q] in 0.1 g
LYTTA VESICATORIA (UNII: 3Q034RO3BT) (LYTTA VESICATORIA - UNII:3Q034RO3BT)	LYTTA VESICATORIA	3 [hp_X] in 0.1 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	
ALCOHOL (UNII: 3K9958V90M)	
PANTHENOL (UNII: WV9CM0O67Z)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24330-210-06	1 in 1 CARTON	09/01/2016	
1		170 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:24330-210-03	1 in 1 CARTON	09/01/2016	
2		71 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:24330-210-01	1 in 1 POUCH	08/01/2015	09/01/2016
3		7.1 g in 1 TUBE; Type 0: Not a Combination Product		
4	NDC:24330-210-21	1 in 1 CARTON	08/01/2015	08/02/2015
4		21.3 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 homeopathic		01/01/2007	

Labeler - Welmedix LLC (830387812)

Revised: 3/2018

Welmedix LLC