

ARC SKIN PROTECTANT- allantoin ointment
Anjon Biologics, Inc

NATURAL CREAM ARC

Protection Against Radiation Dermatitis

Indication for use •

ARC[®] is a pre-radiation topical cream treatment for the protection against and management of Radiation Dermatitis.

Directions •

For topical use only. Apply ARC[®] twice daily to all skin areas that will be exposed to radiation, or as directed by a health care professional. Can be applied 1-2 hours prior to treatment. Massage gently into the skin until absorbed. Continue use until treatment cycle is complete. Washing of the treated area will not reduce skin protection. ARC[®] does not interfere with the effectiveness of radiation treatment.

Ingredients •

Each gram contains 5 mg (0.5%) Allantoin in a cream base of

Aloe Vera Leaf Gel, Bees Wax White, BHT, Cetyl Alcohol, DMDM H, Glycerin, Glyceryl Stearate SE, Isopropyl Myristate, Mineral Oil, Nipagin, Nipasol, Polydimethylsiloxane (PDMS), Polysorbate 60, Polysorbate 20, Sesame Oil, Shea Butter, Sorbitan Tristearate, Water

Store at room temperature. Avoid excessive heat. Do not freeze.

Warnings

For external use only. Should you have an adverse reaction, consult your physician.

Keep out of reach of children.

NATURAL CREAM

THERGY

Cancer Care Solutions

Las Vegas, NV 89117

thergy.com

Made in USA

Packaging

NATURAL CREAM

ARC

Protection Against
Radiation Dermatitis



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Net Wt 100gr (3.5 oz)


Las Vegas, NV 89117
thergy.com
Made In USA
NDC 69148-001-02

ARC SKIN PROTECTANT

allantoin ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69148-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WHITE WAX (UNII: 7G1J5DA97F)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
MINERAL OIL (UNII: T5L8T28FGP)	
METHYLPARABEN SODIUM (UNII: CR6K9C2NHK)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SESAME OIL (UNII: QX10HYY4QV)	
SHEA BUTTER (UNII: K49155WL9Y)	
SORBITAN TRISTEARATE (UNII: 6LUM696811)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69148-001-00	118.6 g in 1 TUBE; Type 0: Not a Combination Product	08/06/2014	04/22/2024
2	NDC:69148-001-01	60 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2018	
3	NDC:69148-001-02	100 g in 1 TUBE; Type 0: Not a Combination Product	04/22/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	08/06/2014	

Labeler - Anjon Biologics, Inc (054126951)**Registrant** - Anjon Biologics, Inc (054126951)

