MAXIMUM STRENGTH MOISTURE BARRIER- petrolatum and zinc oxide ointment
Derma Sciences Canada Inc.

Maximum Strength Moisture Barrier Ointment

Petrolatum 50%

Zinc Oxide 16%

Discontinue use and contact a physician if condition worsens.

Uses

Temporarily protects and helps relieve minor skin irritation and itching due to rashes

Directions

- Carefully cleanse the area and pat dry thoroughly. Apply liberally to desired areas as often as necessary.

Warnings

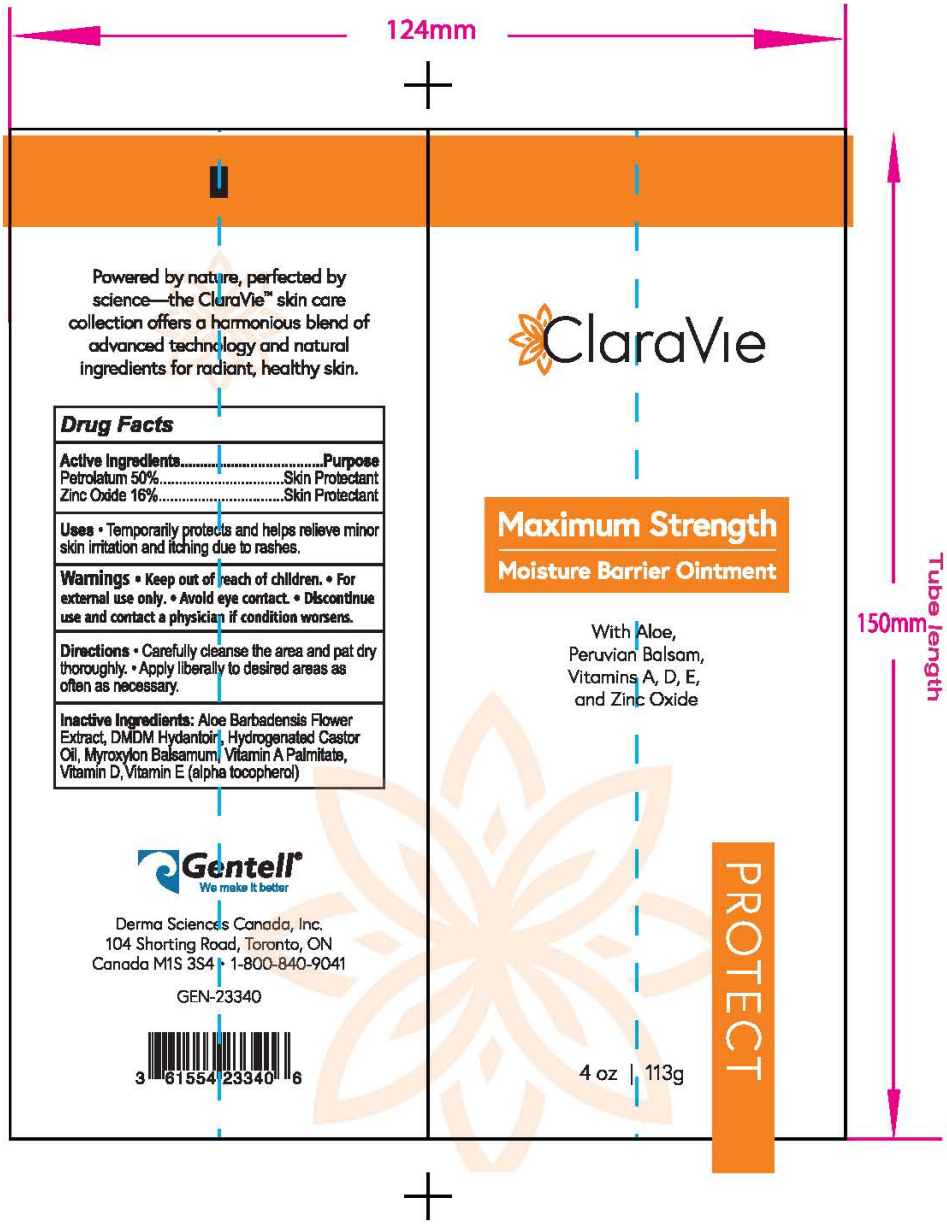
Keep out of reach of children. For external use only. Avoid Eye contact. Discontinue use and contact a physician if condition worsens.

Aloe Barbadensis Flower Extract, DMDM Hydantoin; Hydrogenated Castor Oil; Myroxylon Balsamum, Vitamin A Palminate, Vitamin D, Vitamin E (Alpha Tocopherol)

Skin Protectant

Keep out of reach of children

1-800-840-9041



Date	: 10,02,2025
Revise No	: 1
Name Product	ClaraVie Moisture Barrier Ointment 4oz_OL _V3_020725
DIAMETER	: 40 x 150
Code	:
Approved by	:
	OFFSET 307 151 Black 15% 151

Signed _____ *Date* _____

(Unsigned approval via return email, indicating such approval, will be accepted)

MAXIMUM STRENGTH MOISTURE BARRIER

petrolatum and zinc oxide ointment

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:64772-203
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	0.5 g in 1 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.16 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	
VITAMIN D (UNII: 9VU1KI44GP)	
MYROXYLON BALSAMUM WHOLE (UNII: 5OU0811D9J)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64772-203-01	113.4 g in 1 TUBE; Type 0: Not a Combination Product	11/13/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	11/13/2014	

Labeler - Derma Sciences Canada Inc. (200564891)

Establishment

Name	Address	ID/FEI	Business Operations
Derma Sciences Canada Inc.		200564891	manufacture(64772-203) , label(64772-203)