

LEVOCETIRIZINE DIHYDROCHLORIDE 2% / MUPIROCIN 2% / TRIAMCINOLONE ACETONIDE 0.025% - levocetirizine dihydrochloride 2% / mupirocin 2% / triamcinolone acetonide 0.025% cream
Sincerus Florida, LLC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, [click here](#).

LEVOCETIRIZINE DIHYDROCHLORIDE 2% / MUPIROCIN 2% / TRIAMCINOLONE ACETONIDE 0.025%

Directions for use



Directions for use

As directed by Physician.

Apply topically. For external use only. Wash hands after use.

Store at controlled room temperature (20-25C).

Sincerus Florida, LLC (800) 604-5032

3265 W McNab Rd, Pompano Beach, FL 33069

To report suspected adverse reactions, contact

Sincerus Florida, LLC at (800) 604-5032, or FDA at www.FDA.gov/MedWatch or (800) FDA-1088.

Office use only. Not for resale.



Sincerus Florida, LLC. Adverse reactions



Directions for use

As directed by Physician.

Apply topically. For external use only.

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3265 W McNab Rd, Pompano Beach, FL 33069

To report suspected adverse reactions, contact

Sincerus Florida, LLC at (800) 604-4444

at www.FDA.gov/MedWatch or (800) 604-4444

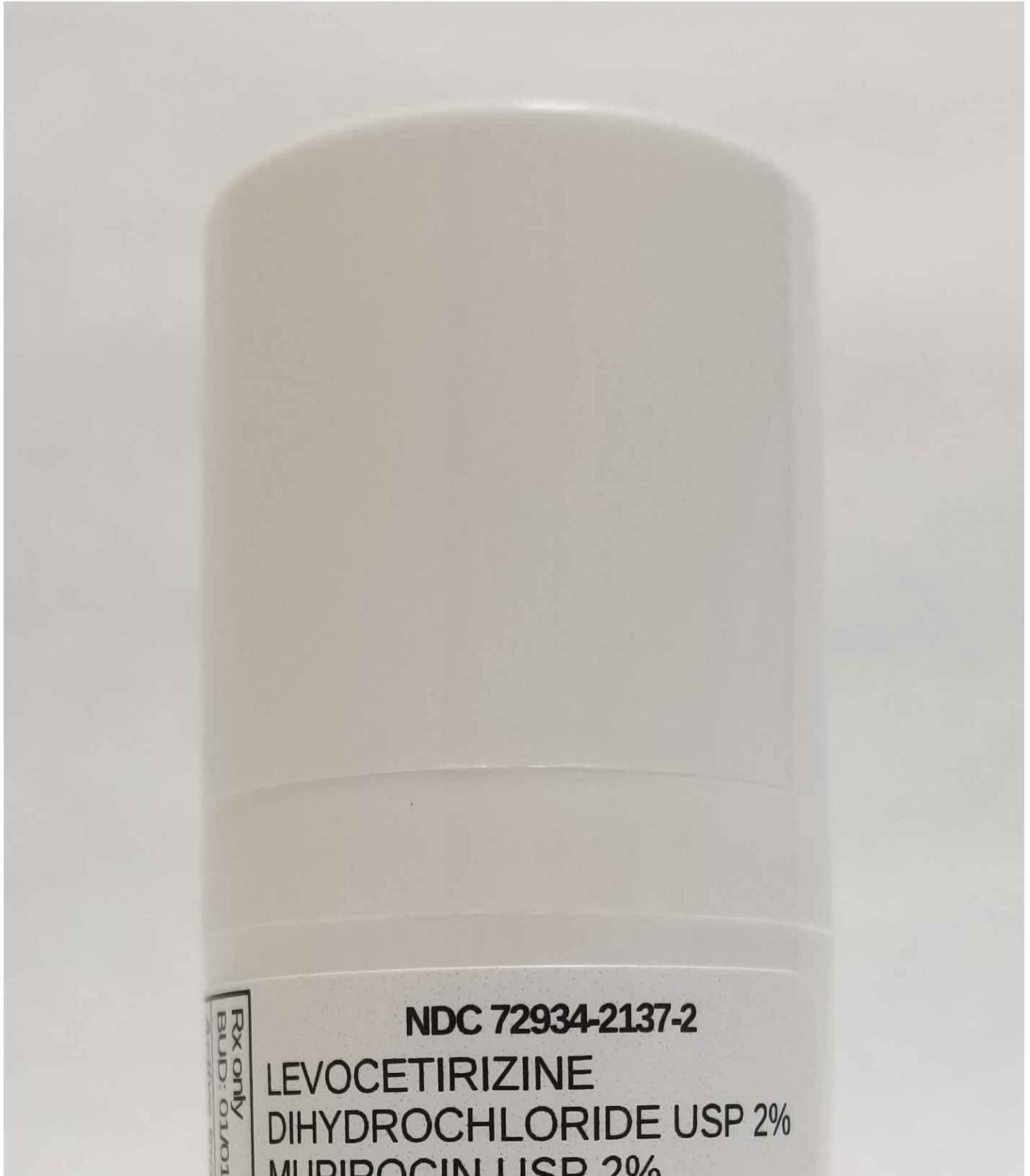
Office use only. Not for patient use.



Active, inactive



NDC 72934- 2137-2 LEVOCETIRIZINE DIHYDROCHLORIDE USP 2% / MUPIROCIN USP 2% / TRIAMCINOLONE ACETONIDE USP 0.025%. Cream 30gm





LEVOCETIRIZINE DIHYDROCHLORIDE 2% / MUPIROCIN 2% / TRIAMCINOLONE ACETONIDE 0.025%

levocetirizine dihydrochloride 2% / mupirocin 2% / triamcinolone acetoneide 0.025% cream

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:72934-2137
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LEVOCETIRIZINE DIHYDROCHLORIDE (UNII: SOD6A38AGA) (LEVOCETIRIZINE - UNII:6U5EA9RT2O)	LEVOCETIRIZINE DIHYDROCHLORIDE	2 g in 100 g
MUPIROCIN (UNII: D0GX8630A5) (MUPIROCIN - UNII:D0GX8630A5)	MUPIROCIN	2 g in 100 g
TRIAMCINOLONE ACETONIDE (UNII: F446C597KA) (TRIAMCINOLONE ACETONIDE - UNII:F446C597KA)	TRIAMCINOLONE ACETONIDE	0.025 g in 100 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72934-2137-2	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/15/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/15/2019	

Labeler - Sincerus Florida, LLC (080105003)**Establishment**

Name	Address	ID/FEI	Business Operations
Sincerus Florida, LLC		080105003	manufacture(72934-2137)

Revised: 5/2019

Sincerus Florida, LLC