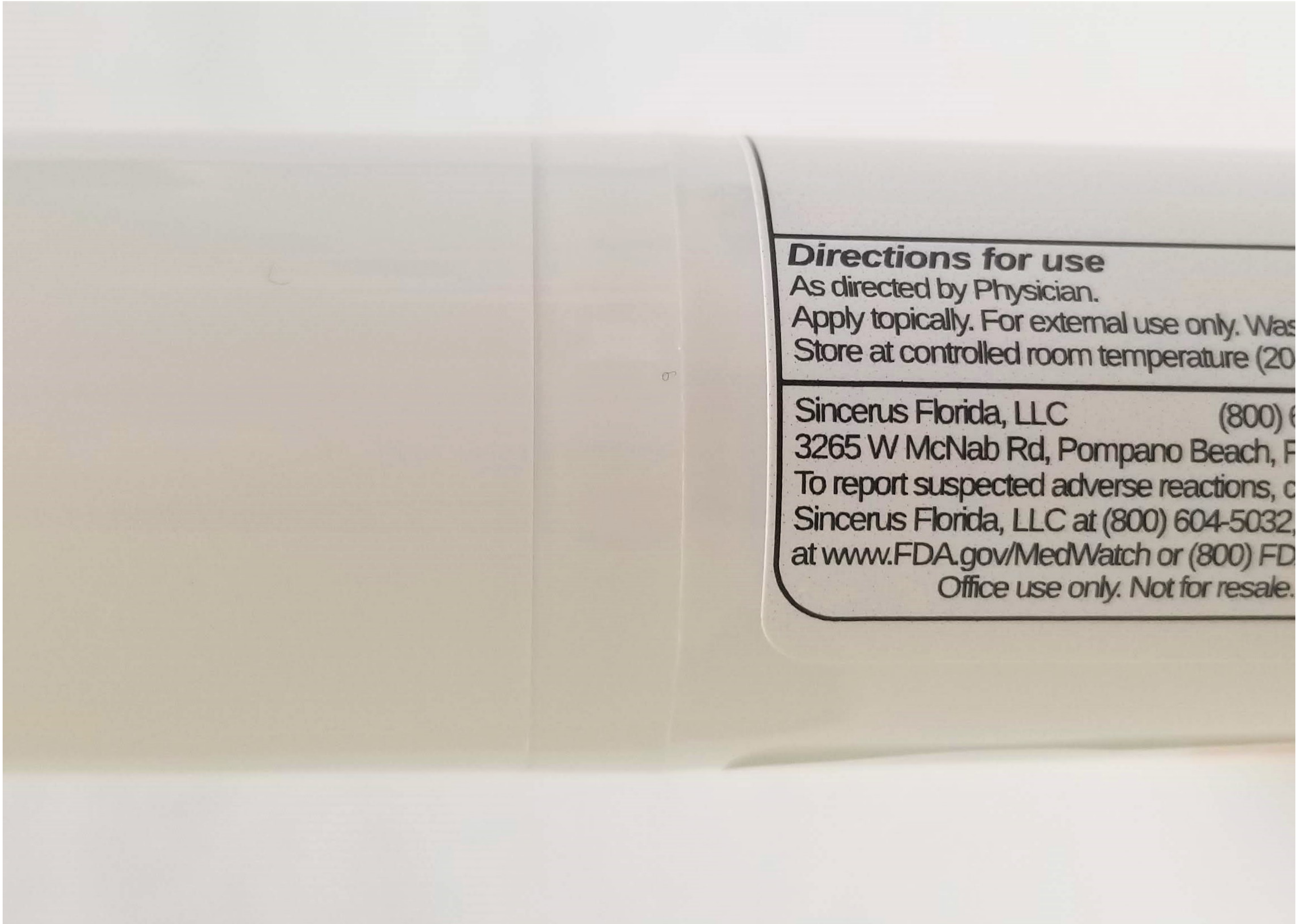
METRONIDAZOLE 1% / MUPIROCIN 2%- metronidazole 1% / mupirocin 2% ointment
Sincerus Florida, LLC

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.

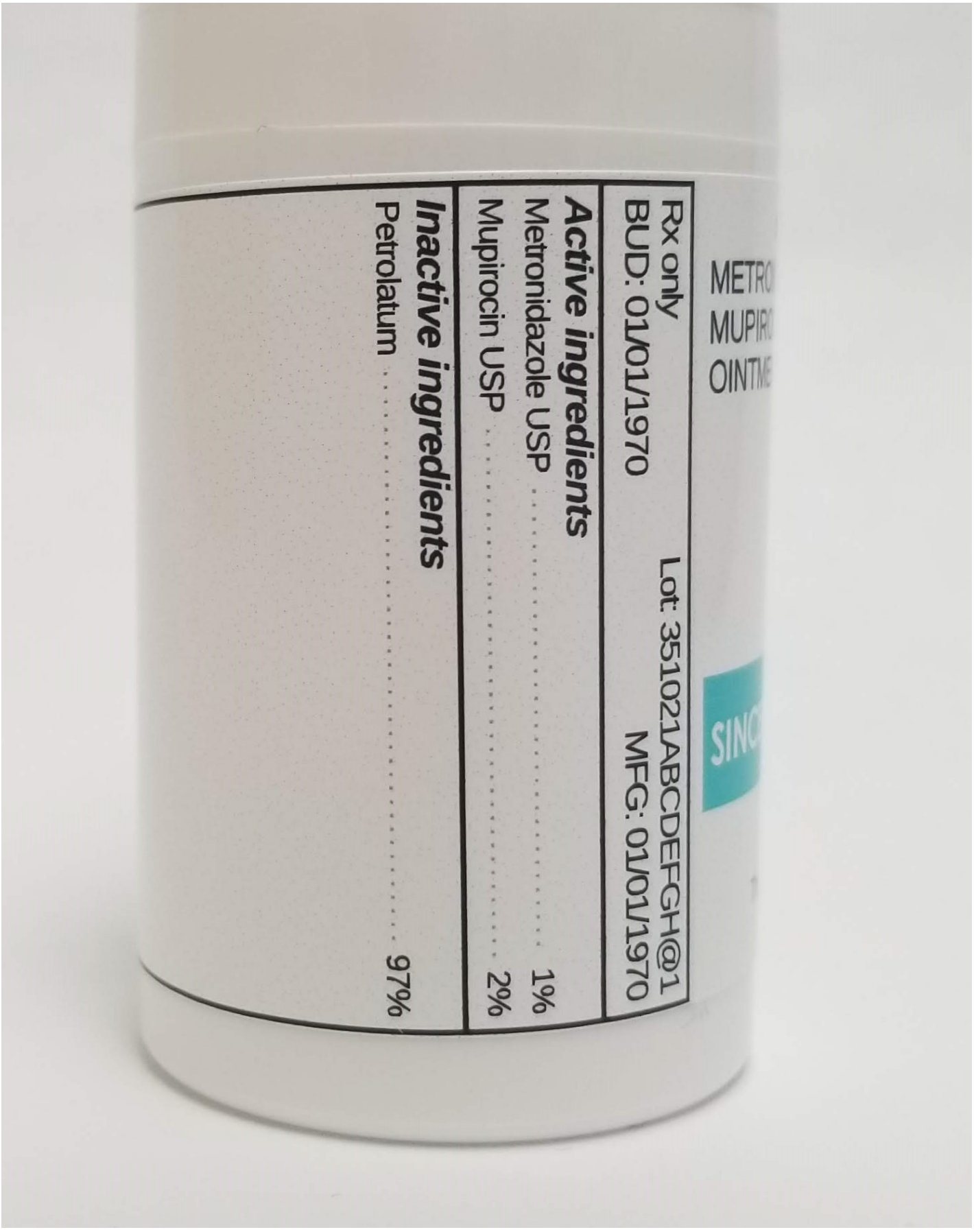
METRONIDAZOLE 1% / MUPIROCIN 2%

Directions for use

Sincerus Florida, LLC. Adverse reactions


Directions for use
As directed by Physician.
Apply topically. For external use only. Was
Store at controlled room temperature (20
Sincerus Florida, LLC (800)
3265 W McNab Rd, Pompano Beach, F
To report suspected adverse reactions, c
Sincerus Florida, LLC at (800) 604-5032,
at www.FDA.gov/MedWatch or (800) FD
Office use only. Not for resale.

Active, inactive



NDC 72934- 5144-2 METRONIDAZOLE USP 1% / MUPIROCIN USP 2%, Ointment 30gm


NDC 72934-5144-2
METRONIDAZOLE USP 1%
MUPIROCIN USP 2%
OINTMENT 30gm
Rx only

METRONIDAZOLE 1% / MUPIROCIN 2%

metronidazole 1% / mupirocin 2% ointment

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:72934-5144
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
METRONIDAZOLE (UNII: 140QMO216E) (METRONIDAZOLE - UNII:140QMO216E)	METRONIDAZOLE	1 g in 100 g
MUPIROCIN (UNII: D0GX863OA5) (MUPIROCIN - UNII:D0GX863OA5)	MUPIROCIN	2 g in 100 g

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date

#	Item Code	Package Description	Date	Marketing End Date
1	NDC:72934-5144-2	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(72934-5144)

Revised: 5/2019 Sincerus Florida, LLC