141055 HYDROQUINONE 6% - 141055 hydroquinone 6% emulsion 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.

141055 HYDROQUINONE 6%

Directions for use





Sincerus Florida, LLC adverse reactions

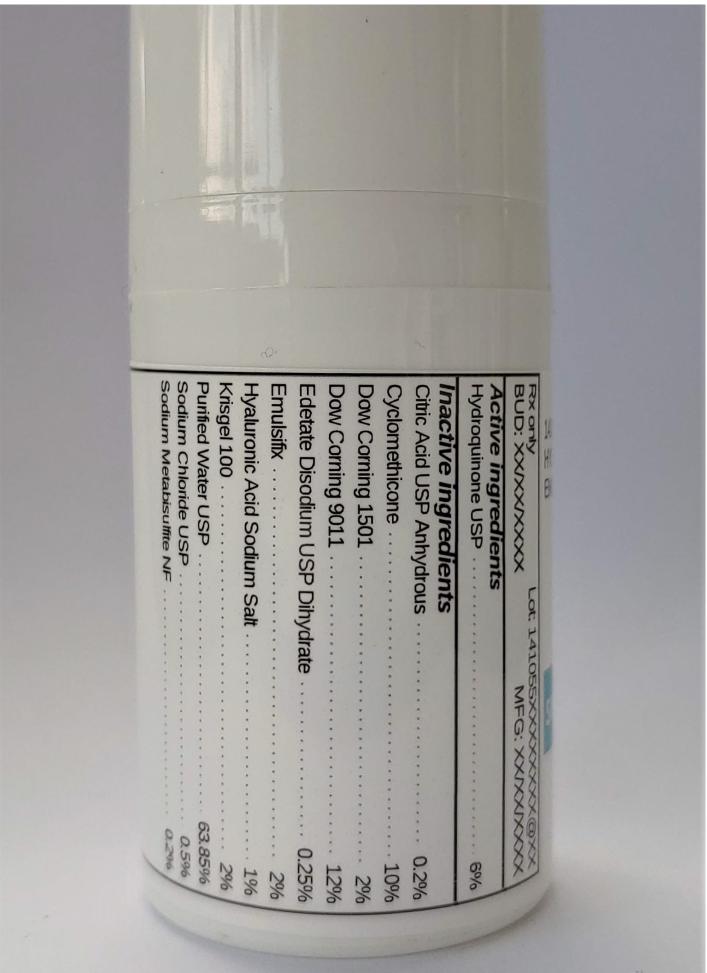
Sincerus Florida, LLC (8 3265 W McNab Rd, Pompano Bea To report suspected adverse reactic Sincerus Florida, LLC at (800) 604at www.FDA.gov/MedWatch or (80 Office use only. Not for re

Directions for use As directed by Physician. Apply topically. For external use on Store at controlled room temperat.



Active Inactive







NDC 72934-6230-2 141055 HYDROQUINONE 6% Emulsion 30 gm





141055 HYDROQUINONE 6%

141055 hydroquinone 6% emulsion

Product Information

Product Type

HUMAN PRESCRIPTION DRUG

Item Code (Source)

NDC:72934-6230

Route of Administration			TOPICAL									
Active Ingredient/Active Moiety												
			Ingredient Name		Basis of Strength S			rength				
н	YDRO Q UINO NE (U	NII: XV74	1AE)	HYDROQUINONE 6 g in 100 g			n 100 g				
Product Characteristics												
Color			yellow (Beige)	Sc	ore							
Shape				Siz	ize							
Flavor				Im	nprint Code							
Contains												
Packaging												
					Marke	ting Start Marke		eting	ting End			
#	Item Code		Package Description		Date		Date					
1	NDC:72934-6230- 2	30 g in 1 Product	30 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		07/02/2020							
N	Marketing Information											
	Marketing Catego		olication Number or Monograph Citatio	n	Marketing Start Date		Marketing End Date		nd Date			
ur	napproved drug other	r			07/02/2020							

Labeler - Sincerus Florida, LLC (080105003)

Establishment											
Name	Address	ID/FEI	Business Operations								
Sincerus Florida, LLC		080105003	manufacture(72934-6230)								

Revised: 7/2020

Sincerus Florida, LLC