MEOLY MILIA REMOVER SERUM- milia remover serum liquid
Consilii LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Active Ingredient

Didecyl dimethyl 0.025%

Purpose

Disinfectant

Use

temporarily relieves these symptoms dueto hay fever or other upper respiratory allergies

Warnings

For external use only. Stop using and ask a doctorif rash or irritation happens.

Do not use

on irritated skin on any area that is infected or reddened - Do not use it if pregnant or if there is any allergic reaction to this product - Avoid contact with eyes.

When Using

For external use only Stop using and ask a doctorif rash or irritation happens.

Stop Use

Stop using and ask a doctorif rash or irritation happens. If contact with eyesoccurs, rinse thoroughly with water, Store in a coolshady place.

Keep Oot Of Reach Of Children

If product gets into eyes, flush with water for 15 minutes

If swallowed, get medical help or contact a Poison Control Center right away

Ask Doctor

Stop using and ask a doctorif rash or irritation happens.

Directions

Apply daily after cleansing and toning. Twice a day for at least 4 weeks.
Dispense a pea-size amount of the essence onto the problematic area and use fingertips to massage until fully absorbed.
Keep using it for a period of time until the milia doesn’t appear again.

Inactive ingredients

Glycerin, salicyle acid, dextrin,xanthan gum, amylopectin, tocopheryl acetate, dodecyl dimethyl benzyl ammonium chloride 0.025% ± 10%

PACKAGE LABEL

Size / Package size: 40mm x 40mm x 100mm
Bottle label size: 113mm x 40mm

MEOLY MILIA REMOVER SERUM

milia remover serum liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83299-013
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIDECYLDIMONIUM (UNII: Z7F472XQPA) (DIDECYLDIMONIUM - UNII:Z7F472XQPA)	DIDECYLDIMONIUM	0.025 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
AMYLOPECTIN, CORN (UNII: TQI9LJM246)	
GLYCERIN (UNII: PDC6A3C0OX)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ICODEXTRIN (UNII: 2NX48Z0A9G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83299-013-01	30 g in 1 BOTTLE; Type 0: Not a Combination Product	08/14/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M005	08/14/2023	

Labeler - Consilii LLC (118891890)

Establishment

Name	Address	ID/FEI	Business Operations
Consilii LLC		118891890	manufacture(83299-013) , label(83299-013)

Revised: 8/2023 Consilii LLC