

ANIOSGEL 85 NPC- ethanol gel
Laboratoires Anios

Ethanol 70.0% (w/w) (equivalent to 76.91% v/v)

Antiseptic Hand Rub

Uses

A hand rub to decrease bacteria on the skin.

Warnings

- **For external use only**
- **FLAMMABLE. Keep away from fire or flame, heat sparks and sources of static discharge.**

Do not use

In eyes

When using this product

- If in eyes, rinse promptly and thoroughly with water.
- Discontinue use if irritation and redness develop.

Stop use and ask a doctor if skin irritatin or redness occurs for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a poison Control Center right away.

Apply product onto hands, spread thoroughly and rub until dry.

Other information

- Store at 41 - 77 °F (5 - 25 °C)
- See Safety Data Sheet (SDS)
- For emergency medical information in USA, call (800)726-2308

Inactive ingredients Water (aqua), Isopropyl Alcohol, Glycerin, Acrylates/C10-30 Alkyl Acrylate Cross-polymer, PEG-5 Cocoate, Bisabolol, Aminomethyl Propanol, Methylpropanediol

Questions or comments? Call (800)726-2308 Monday/Friday 9:00am to 5:00pm (UTC-4)

Principal display panel and representative label

ANIOSGEL

85 NPC

HAND SANITIZING

GEL HAND-RUB

Active ingredient: ethanol 70.0% w/w (equivalent to 76.91% v/v)

17 fl oz (500 mL)

Manufactured by:

Laboratoires Anios and Ecolab Company

1, rue de l'Espoir, Lezennes, France

ANIOSGEL

85 NPC

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GEL HAND-RUB**



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CF16440500MADR-200811

ANIOSGEL 85 NPC

Drug Facts

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- Call Angelini Pharma Inc.: (800)726-2308
- Monday/Friday 9:00am to 5:00pm (UTC-4)



Manufactured by :
Laboratoires ANIOS
1, rue de l'Espoir, Lezennes, France

LOT



EXP



Distributor/Importer :
Angelini Pharma Inc.
9200 Corporate Blvd., Suite 100
Rockville, MD 20820
(800)726-2308
www.angelini-us.com

CF1644855500MADV-200811

ANIOSGEL 85 NPC

ethanol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62169-203
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	377.5 mL in 500 mL

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
GLYCERIN (UNII: PDC6A3C0OX)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
DIETHYLAMINOETHYL PEG-5 COCOATE (UNII: UUG92PR9NR)	
LEVOMENOL (UNII: 24WE03BX2T)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
METHYLPROPANEDIOL (UNII: N8F53B3R4R)	

Product Characteristics

Color		Score	
Shape	FREEFORM	Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62169-203-12	12 in 1 PACKAGE	03/07/2018	08/31/2027
1	NDC:62169-203-17	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
OTC Monograph Drug	505G(a)(3)	04/01/2014	08/31/2027

Labeler - Laboratoires Anios (268309216)

Revised: 2/2025

Laboratoires Anios