SANINTA ANTISEPTIC WIPE- alcohol liquid LL Pharma

Drug Facts - Saninta Antiseptic Wipe - Individual Wipe & Box

Active Ingredient

Isopropyl alcohol 70% v/v

Purpose

Antiseptic

Use

Antiseptic wipe to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame.

Do not use

Do not use

- In children less than 2 months of age
- On open skin wounds

When using this product

When using this product Keep out of eyes, ears and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away at 1-800-222-1222.

Directions

- Wipe hands thoroughly and carefully for at least 30 seconds. Allow to dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information

- Store between 15°C 30°C (59°-86°F)
- Avoid freezing and excessive heat above 40°C (104°F).

Inactive ingredients

Purified Water USP

Questions or Comments?

Questions or Comments? Tel.: 1-833-336-6159

Package Label - Principal Display Panel

NDC: 81351-001-01

Product: 001-01

Effective for hand sanitizing

For personal hand hygiene to help prevent the spread of bacteria

Saninta

Antiseptic Wipe

Antiseptic Alcohol Wipe

Isopropyl Alcohol 70% USP

51 Unit large wipes

8" x 4 3/4 " wipes

Manufactured by:

LL Pharma Inc.

5789 rue Cypihot, Saint-Laurent (Montreal), Quebec H4S 1R3

Made in Canada

www.llpharmainc.com

LOT #:

EXP.:

F001

VER. 01

RE: 01-2021



SANINTA ANTISEPTIC WIPE

Inactive Ingredients

alcohol liquid					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:81351-001		1-001	
Route of Administration	TOPICAL				
Active Ingredient/Active Moiety					
Ingredient Name			Basi: Stren		Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)			ISOPROPYL	ALCOHOL	0.7 L in 1 L

Strength

Ingredient Name

WATER (UNII: 059QF0KO0R)

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81351- 001-01	51 in 1 BOX	03/09/2021	
1		0.0027 L in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	03/09/2021	
3 1 3			

Labeler - LL Pharma (206925146)

Registrant - LL Pharma (206925146)

Establishment			
Name	Address	ID/FEI	Business Operations
LL Pharma Inc.		206925146	manufacture(81351-001) , analysis(81351-001) , label(81351-001) , pack(81351-001)

Revised: 10/2023 LL Pharma