NEW LIFE ADVANCED FORMULA TOPICAL HAND SANITIZER- alcohol spray
New Life Products, Inc.

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.

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new life ™ ADVANCED FORMULA Topical Spray Hand Sanitizer

Drug Facts

Active Ingredient

Alcohol 80% v/v

Purpose

Antiseptic

Uses • Hand sanitizer 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 • In children less than 2 months of age

• On open skin wounds.

Stop use and ask doctor if irritation and redness develops. These may be signs of a serious condition.

KEEP OUT OF REACH OF CHILDREN. If swallowed, get medical help or contact a Poison Control Center right away.

Directions • Spray enough product on hands to cover completely. Rub hands together until dry.
• Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information • Store between 15-30 C(59-86 F)
• Avoid freezing and excessive heat above 40 C(104 F)

Inactive ingredients: Purified Water USP, Glycerin, Hydrogen Peroxide.

Kills Germs +

Helps Reduce The Risk Of Infections

NON-STERILE SOLUTION

MADE IN USA

AFAM CONCEPT, Inc.

Chicago, IL 60629

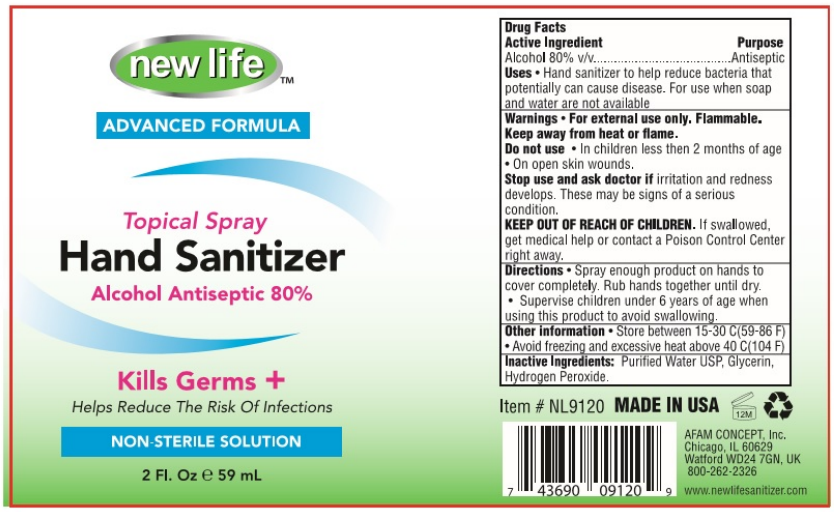
Watford WD24 7GN, UK

800-262-2326

www.newlifesanitizer.com

Packaging

59 ml Pack



236 ml Pack

NEW LIFE ADVANCED FORMULA TOPICAL HAND SANITIZER

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49681-200
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49681-200-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/21/2020	
2	NDC:49681-200-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/21/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/21/2020	

Labeler - New Life Products, Inc. (117815140)

Establishment

Name	Address	ID/FEI	Business Operations
New Life Products, Inc.		117815140	manufacture(49681-200)

Revised: 2/2023 New Life Products, Inc.