ALCOHOL- alcohol liquid ALCOHOL- alcohol gel Hello Bello

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.

DRUG FACTS

ACTIVE INGREDIENT

Ethyl Alcohol - 62.0%

PURPOSE

Antiseptic

USES:

For hand washing to decrease bacteria on skin

• Recommended for repeated use.

WARNINGS:

For external use only

- Flammable
- Keep away from fire/flame
- Do not use this product in or near eyes. If contact occurs rinse thoroughly with water
- Keep out of reach of children
- If swallowed, get medical help or contact a Poison Control Center right away
- Children under six years of age should be supervised.

DIRECTIONS:

Dispense an adequate amount in your palm to cover all surfaces of hands completely

- Rub hands together until dry
- Does not require rinsing.

OTHER INFORMATION:

Do not store above 104°F/40°C.

• May discolor some fabrics.

INACTIVE INGREDIENTS:

Aloe Barbadensis Leaf Juice. Calendula Officinalis Flower Extract Chamomilla Recutita (Matricaria) Flower Extract, Glycerin, Water

Principal Display Panel – 59 mL Bottle Label

 $\begin{array}{c} \text{hello} \\ \text{bello}^{\text{TM}} \end{array}$

Fragrance-Free

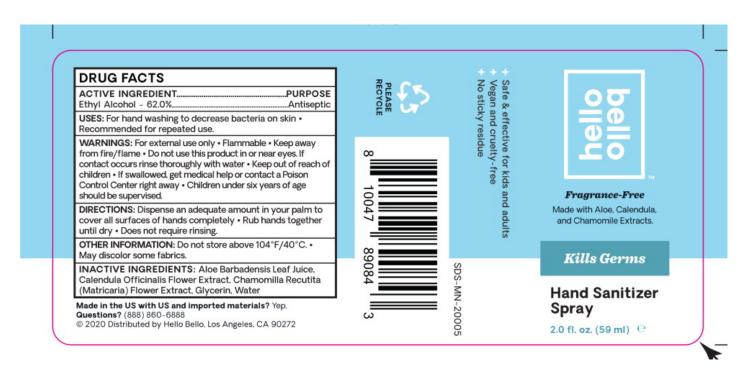
Made with Aloe, Calendula, and Chamomile Extracts.

- + Safe & effective for kids and adults
- + Vegan and cruelty-free
- + No sticky residue

Kills Germs

Hand Sanitizer Spray

2.0 fl. oz. (59 ml) e



Principal Display Panel – 118 mL Bottle Label

Non-Sterile Solution

hello belloTM

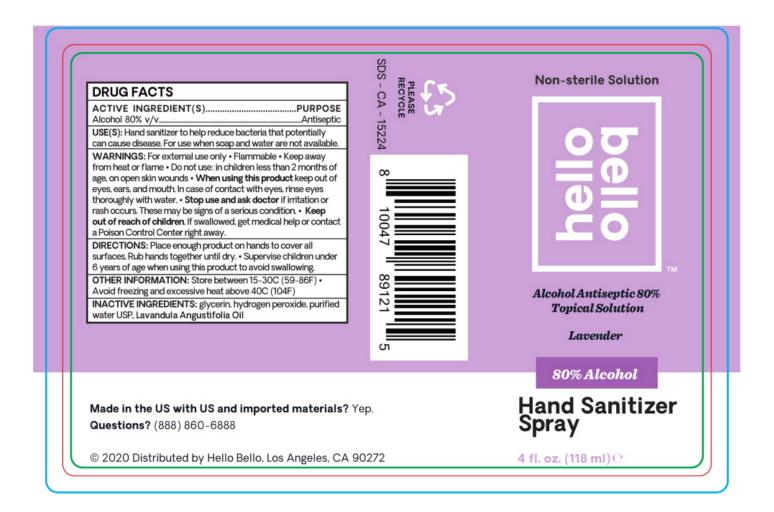
Alcohol Antiseptic 80% Topical Solution

Lavender

80% Alcohol

Hand Sanitizer Spray

4 fl. oz. (118 ml) e



Principal Display Panel – 118 mL Bottle Label

Non-Sterile Solution

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Alcohol Antiseptic 80% Topical Solution

80% Alcohol

Hand Sanitizer Spray

4 fl. oz. (118 ml) e



Principal Display Panel – 236 mL Bottle Label

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Fragrance-Free

70% Alcohol

Hand Sanitizer Gel

8 fl. oz. (236 ml) e



ALCOHOL

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73418-030
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	62 mL in 100 mL	

Inactive Ingredients		
Ingredient Name	Strength	
Aloe Vera Leaf (UNII: ZY81Z83H0X)		
Calendula Officinalis Flower (UNII: P0M7O4Y7YD)		
Chamomile (UNII: FGL3685T2X)		
Glycerin (UNII: PDC6A3C0OX)		
Water (UNII: 059QF0KO0R)		

	Pac	ckaging			
ı	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 N	DC:73418-030-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	04/01/2020		

ALCOHOL

alcohol liquid

Pro	duct	Information	
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:73418-031

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient NameBasis of StrengthStrengthAlcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)Alcohol80 mL in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
GLYCERIN (UNII: PDC6A3C0OX)		
Hydrogen Peroxide (UNII: BBX060AN9V)		
Water (UNII: 059QF0KO0R)		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:73418-031-01	118 mL in 1 BOTTLE: Type 0: Not a Combination Product	04/01/2020		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/01/2020	

ALCOHOL

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73418-032
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		
GLYCERIN (UNII: PDC6A3C0OX)			
Hydrogen Peroxide (UNII: BBX060AN9V)			
Water (UNII: 059QF0KO0R)			
Lavender Oil (UNII: ZBP1YXW0H8)			

ı	Packaging				
ı	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:73418-032-01	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/01/2020	

ALCOHOL

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73418-033
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	70 mL in 100 mL

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)			
GLYCERIN (UNII: PDC6 A3C0 OX)			
CARBO MER HO MO PO LYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)			
TROLAMINE (UNII: 9O3K93S3TK)			

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:73418-033-01	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2020		
Marketing Information				
with Kethig iii	or mucron			
Marketing Catego		Marketing Start Date	Marketing End Date	
	ry Application Number or Monograph Citation	Marketing Start Date 04/01/2020	Marketing End Date	

Labeler - Hello Bello (081135707)

Establishment			
Name	Address	ID/FEI	Business Operations
Unconditional Love		081135707	MANUFACTURE(73418-030, 73418-031, 73418-032, 73418-033)

Revised: 2/2021 Hello Bello