

W/CLIP PDQ INSTANT HAND SANITIZER- ethyl alcohol
Universal Distribution Center LLC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

W/CLIP PDQ INSTANT HAND SANITIZER

INSTANT HAND SANITIZER Citrus Zest

Drug Facts

Active ingredient

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Uses

hand sanitizer to help reduce bacteria on the skin.

Warnings

For external use only.

Flammable. Keep away from heat and flame.

When using this product avoid contact with face, eyes, and broken skin. In case of eye contact, flush with plenty of water and seek medical advice.

Stop use and ask a doctor if irritation or redness develops.

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

Directions

• wet hands thoroughly with product and rub into skin until dry. • Children under 6 years of age should be supervised by an adult when using.

Inactive ingredients

aqua(water), acrylates/c10-30 alkyl acrylate crosspolymer, aminomethyl propanol, aloe barbadensis leaf juice, CI 14700, CI 19140, denatonium benzoate, fragrance, glycerine,

propylene glycol, tocopheryl acetate

INSTANT HAND SANITIZER Lavender

Drug Facts

Active ingredient

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Uses

hand sanitizer to help reduce bacteria on the skin.

Warnings

For external use only.

Flammable. Keep away from heat and flame.

When using this product avoid contact with face, eyes, and broken skin. In case of eye contact, flush with plenty of water and seek medical advice.

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Directions

• wet hands thoroughly with product and rub into skin until dry. • Children under 6 years of age should be supervised by an adult when using.

Inactive ingredients

aqua(water), acrylates/c10-30 alkyl acrylate crosspolymer, aminomethyl propanol, aloe barbadensis leaf juice, CI 17200, CI 42090, denatonium benzoate, fragrance, glycerine, propylene glycol, tocopheryl acetate

INSTANT HAND SANITIZER Aloe Vera

Drug Facts

Active ingredient

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Uses

hand sanitizer to help reduce bacteria on the skin.

Warnings

For external use only.

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Directions

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Inactive ingredients

aqua(water), acrylates/c10-30 alkyl acrylate crosspolymer, aminomethyl propanol, aloe barbadensis leaf juice, CI 19140, CI 42090, denatonium benzoate, fragrance, glycerine, propylene glycol, tocopheryl acetate

INSTANT HAND SANITIZER Classic

Drug Facts

Active ingredient

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Uses

hand sanitizer to help reduce bacteria on the skin.

Warnings

For external use only.

Flammable. Keep away from heat and flame.

When using this product avoid contact with face, eyes, and broken skin. In case of eye contact, flush with plenty of water and seek medical advice.

Stop use and ask a doctor if irritation or redness develops.

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

Directions

- wet hands thoroughly with product and rub into skin until dry.
- Children under 6 years of age should be supervised by an adult when using.

Inactive ingredients

aqua(water), acrylates/c10-30 alkyl acrylate crosspolymer, aminomethyl propanol, aloe barbadensis leaf juice, denatonium benzoate, fragrance, glycerine, propylene glycol, tocopheryl acetate

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Packaging

Kit Label



Kit Components Label



Drug Facts Label - Citrus Zest

Drug Facts

Active ingredient Purpose
Ethyl Alcohol 70% v/v.....Antimicrobial

Uses hand sanitizer to help reduce bacteria on the skin.

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Inactive ingredients aqua(water), acrylates/c10-30 alkyl acrylate crosspolymer, aminomethyl propanol, aloe barbadensis leaf juice, CI 14700, CI 19140, denatonium benzoate, fragrance, glycerine, propylene glycol, tocopheryl acetate

Drug Facts Label - Lavender

Drug Facts

Active ingredient Purpose
Ethyl Alcohol 70% v/v.....Antimicrobial

Uses hand sanitizer to help reduce bacteria on the skin.

Warnings For external use only.

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Directions ■ wet hands thoroughly with product and rub into skin until dry. ■ Children under 6 years of age should be supervised by an adult when using.

Inactive ingredients aqua(water), acrylates/c10-30 alkyl acrylate crosspolymer, aminomethyl propanol, aloe barbadensis leaf juice, CI 17200, CI 42090, denatonium benzoate, fragrance, glycerine, propylene glycol, tocopheryl acetate

Drug Facts Label - Aloe Vera

Drug Facts

Active ingredient Purpose
Ethyl Alcohol 70% v/v.....Antimicrobial

Uses hand sanitizer to help reduce bacteria on the skin.

Warnings For external use only.

Flammable. Keep away from heat and flame.

When using this product avoid contact with face, eyes, and broken skin. In case of eye contact, flush with plenty of water and seek medical advice.

Stop use and ask a doctor if irritation or redness develops.

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

Directions ■ wet hands thoroughly with product and rub into skin until dry. ■ Children under 6 years of age should be supervised by an adult when using.

Inactive ingredients aqua(water), acrylates/c10-30 alkyl acrylate crosspolymer, aminomethyl propanol, aloe barbadensis leaf juice, CI 19140, CI 42090, denatonium benzoate, fragrance, glycerine, propylene glycol, tocopheryl acetate

Drug Facts Label - Classic

Drug Facts

Active ingredient Purpose
Ethyl Alcohol 70% v/v.....Antimicrobial

Uses hand sanitizer to help reduce bacteria on the skin.

Warnings For external use only.

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Inactive ingredients aqua(water), acrylates/c10-30 alkyl acrylate crosspolymer, aminomethyl propanol, aloe barbadensis leaf juice, denatonium benzoate, fragrance, glycerine, propylene glycol, tocopheryl acetate

W/CLIP PDQ INSTANT HAND SANITIZER

ethyl alcohol kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52000-423
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-423-24	1 in 1 BOX; Type 0: Not a Combination Product	04/09/2025	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	6 BOTTLE	328.2 mL
Part 2	6 BOTTLE	328.2 mL
Part 3	6 BOTTLE	328.2 mL
Part 4	6 BOTTLE	328.2 mL

Part 1 of 4

INSTANT HAND SANITIZER CITRUS ZEST

ethyl alcohol gel

Product Information

Item Code (Source)	NDC:52000-424
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Route of Administration	TOPICAL
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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)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	

GLYCERIN (UNII: PDC6A3C0OX)					
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)					
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)					
Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:52000-424-24	54.7 mL in 1 BOTTLE; 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)	04/09/2025		
Part 2 of 4					
INSTANT HAND SANITIZER LAVENDER					
ethyl alcohol gel					
Product Information					
Item Code (Source)		NDC:52000-425			
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)					
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)					
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)					
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)					
D&C RED NO. 33 (UNII: 9DBA0SBB0L)					
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)					
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)					
GLYCERIN (UNII: PDC6A3C0OX)					
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)					
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)					

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-425-24	54.7 mL in 1 BOTTLE; 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)	04/09/2025	

Part 3 of 4	
INSTANT HAND SANITIZER ALOE VERA	
ethyl alcohol gel	

Product Information	
Item Code (Source)	NDC:52000-426
Route of Administration	TOPICAL

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

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging			
#	Item Code	Package Description	Marketing End Date

1	NDC:52000-426-24	54.7 mL in 1 BOTTLE; Type 0: Not a Combination Product		
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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/09/2025	

Part 4 of 4

INSTANT HAND SANITIZER CLASSIC

ethyl alcohol gel

Product Information

Item Code (Source)	NDC:52000-427
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)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-427-24	54.7 mL in 1 BOTTLE; 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)	04/09/2025	
Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/09/2025	

Labeler - Universal Distribution Center LLC (019180459)