

**UNITED AIRLINES POLARIS KIT- sodium monofluorophosphate, alcohol
Buzz Export Services Pty Ltd (Buzz Products)**

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.

United Airlines Polaris Kit

Drug Facts

Active ingredient

Sodium monofluorophosphate 0.76% (0.15% w/v fluoride ion)

Purpose

Anticavity

Use

helps protect against cavities

Warnings

Keep out of the reach of children under 6 years of age.

If more than used for brushing is accidentally swallowed, get medical help or contact a Poison Control Center right away.

Directions

adults and children 2 years of age and older	brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or physician
children 2 to 6 years	use only a pea sized amount and supervise child's brushing and rinsing (to minimize swallowing)
children under 2 years	ask a dentist or physician

Inactive ingredients

dicalcium phosphate dihydrate, water, sorbitol, sodium lauryl sulfate, flavor, cellulose gum, tetrasodium pyrophosphate, sodium saccharin

Questions or comments?

Call to-free 1-800-468-6502

Drug Facts

Active Ingredient

Ethyl Alcohol 75% (v/v)

Purpose

Antiseptic

Uses

- To help reduce bacteria on the skin.

Warnings

- For external use only.
- Do not dispose of wipes in flush toilets.

Flammable

- Keep away from fire or flame.

When using this product

- avoid contact with eyes.
- If contact occurs, rinse thoroughly with water.

Stop using and ask a doctor if

- irritation or redness develops and lasts.

Keep out of reach of children

- In case of accidental ingestion, get medical help or contact a poison control center immediately.

Directions

- Apply to hands or surface.
- No rinsing required.
- Discard after use.

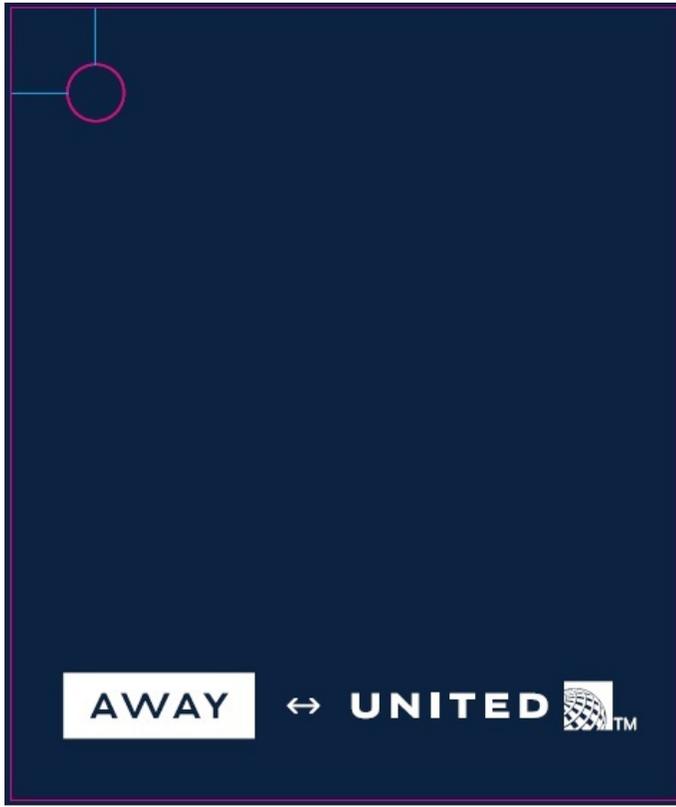
Other Information

- Store below 110°F (43°C).
- May discolor certain fabrics or surfaces.

Inactive ingredients

Aqua (Water), Glycerin, Polyquaternium-37, Melaleuca Alternifolia Leaf Oil.

Package Labeling:69491-036-01



TOOTHBRUSH MADE IN CAMBODIA. COLGATE TOOTHPASTE MADE IN THAILAND. DISTRIBUTED BY BUZZ PRODUCTS.
Consciously designed using less plastic.



DENTAL KIT

Contains:
TOOTHBRUSH
COLGATE TOOTHPASTE 5G

READ THIS FULL INFORMATION BEFORE USE.

TOOTHPASTE DIRECTIONS FOR USE:
Adults and children 2 years and older: Brush teeth thoroughly after meals or at least twice a day, or as directed by a dentist.
Children 2-6 years: Use a pea-sized amount and supervise child to minimize swallowing.
Children under 2 years: Ask a dentist.

WARNINGS:
Keep out of reach of children under 6 years of age. If more than used for brushing is accidentally swallowed, get medical help or contact a Poison Control Center right away.

INGREDIENTS:
Contains: Sodium Monofluorophosphate 0.76% (0.15% W/v Fluoride Ion) Anticavity.
Ingredients: Dicalcium Phosphate Dihydrate, Water, Sorbitol, Sodium Lauryl Sulfate, Flavor, Cellulose Gum, Tetrasodium Pyrophosphate, Sodium Saccharin.

Package Labeling:42555-060-94



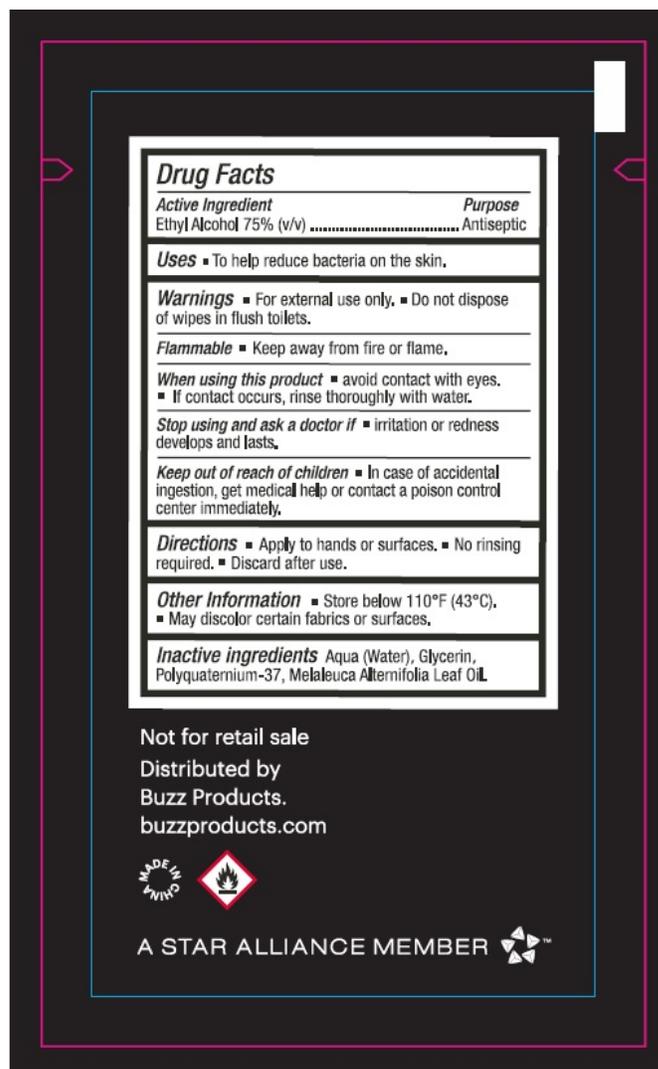
Drug Facts		Drug Facts (continued)	
Active ingredient Sodium monofluorophosphate 0.76% (0.15% w/v fluoride ion) ...Anticavity	Purpose Anticavity	Directions	
Use helps protect against cavities		adults and children 2 years of age and older	brush teeth thoroughly, preferably after each meal or a least twice a day, or as directed by a dentist or physician
Warnings Keep out of the reach of children under 6 years of age. If more than used for brushing is accidentally swallowed, get medical help or contact a Poison Control Center right away.		children 2 to 6 years	use only a pea sized amount and supervise child brushing and rinsing (to minimize swallowing)
		children under 2 years	ask a dentist or physician
		Inactive ingredients dicalcium phosphate dihydrate, water, sorbitol, sodium lauryl sulfate, flavor, cellulose gum, tetrasodium pyrophosphate, sodium saccharin	
		Questions or comments? Call toll-free 1-800-468-6502	

Read this full drug facts information before use.

Package Labeling:69491-035-01



FRONT



BACK

UNITED AIRLINES POLARIS KIT

sodium monofluorophosphate, alcohol kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69491-036
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69491-036-01	1 in 1 KIT	02/15/2022	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 TUBE	5 g
Part 2	1 PATCH	5 mL

Part 1 of 2

COLGATE ANTICAVITY

sodium monofluorophosphate paste, dentifrice

Product Information

Item Code (Source) NDC:42555-060

Route of Administration DENTAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	7.6 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
SODIUM PYROPHOSPHATE (UNII: O352864B8Z)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor	MINT	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42555-060-94	5 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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Part 2 of 2

SANITIZER WIPE IN SACHET

alcohol cloth

Product Information

Item Code (Source)	NDC:69491-035
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
TEA TREE OIL (UNII: VIF565UC2G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69491-035-01	1 in 1 PACKAGE		
1		5 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	02/15/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part355	02/15/2022	

Labeler - Buzz Export Services Pty Ltd (Buzz Products) (747317902)

Revised: 1/2023

Buzz Export Services Pty Ltd (Buzz Products)