WEEKENDER CLEAN TRAVEL KIT- benzalkonium chloride
Talus Corporation

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.

Weekender Clean Travel Kit

Drug Facts

Active Ingredients

Benzalkonium Chloride 0.133% w/v

Purpose

First Aid Antiseptic

Use

Antiseptic Cleansing of face, hands and body without soap and water. Air dries in seconds.

Do Not Use

In the eyes or apply over alrge areas of the body.

Stop Use

If irritation, redness or other symptoms develop. Consult a doctor if the condition persists or gets worse.

Caution Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

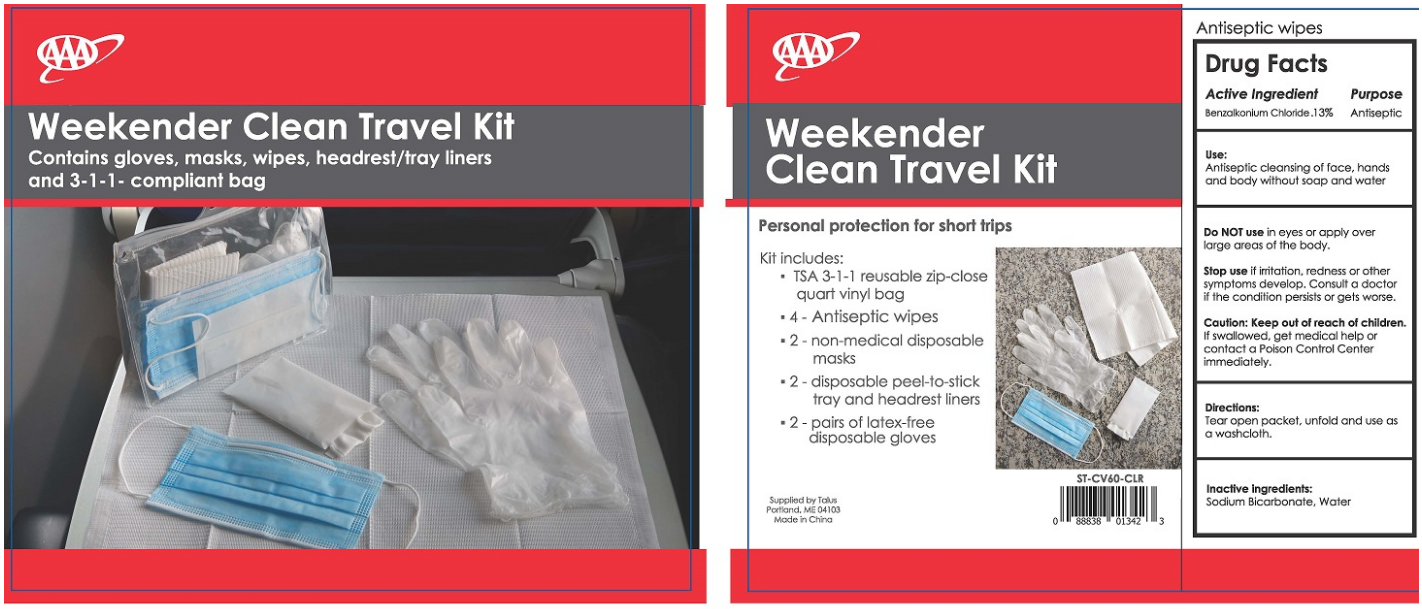
Directions

Tear open packet, unfold and use as a washcloth.

Inactive Ingredients

Sodium Bicarbonate, Water

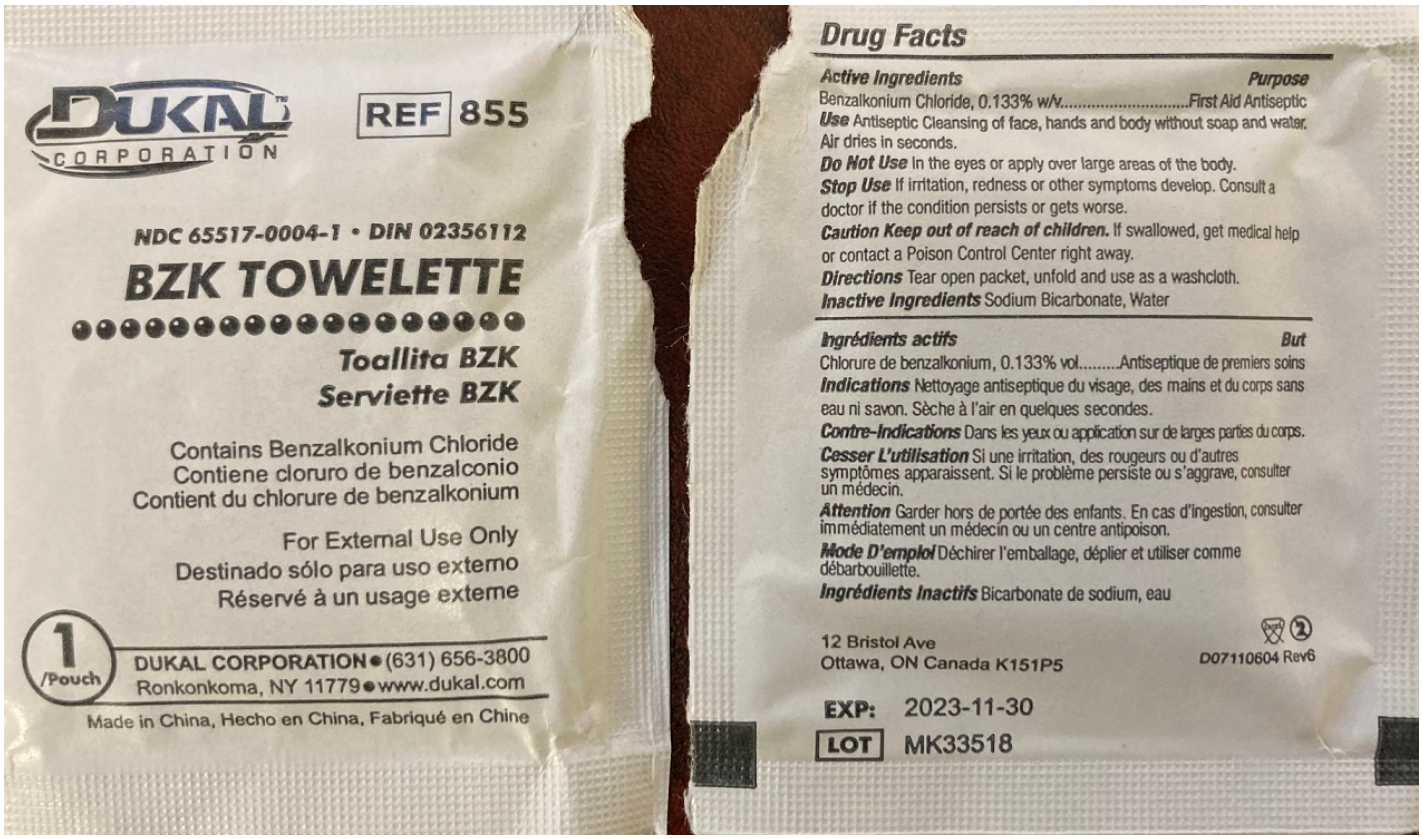
Package Labeling:Kit



Package ID: ZT-ST-CV60-CLR0X
Product ID: ST-CV60-CLR
Package type: Insert (2 pieces - a front and a back)
Colors: CMYK + Pantone 485CV
Size:17.78 cm x 15.24 cm each
Paper stock: 127 g paper board

Acknowledge receipt of file
Make hand sample to be sure of fit in bag
Suggest changes if needed
If no changes are needed - proceed to production
These are to be printed separately and NOT inserted into the final product

Package Labeling:81330-000-00



WEEKENDER CLEAN TRAVEL KIT

benzalkonium chloride kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81330-002

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81330-002-01	1 in 1 KIT	12/23/2020	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	4 PATCH	5.6 mL

Part 1 of 1

DUKAL BZK TOWELETTE

benzalkonium chloride cloth

Product Information

Item Code (Source)	NDC:81330-000
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.33 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81330-000-01	4 in 1 PACKET		
1		1.4 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	12/23/2020	

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Labeler - Talus Corporation (147961023)

Revised: 1/2021

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