

**CRAYOLA WILD BLUE YONDER ROLL ON HAND SANITIZER - benzalkonium chloride gel
Health-Tech, 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.

Drug Facts

Active Ingredients

Active Ingredients: Benzalkonium Chloride - 0.13%

Purpose

Antimicrobial

Uses

For hand sanitizing to decrease bacteria on skin

For external use only. Keep out of reach of children, except under adult supervision. If swallowed, get medical help or contact a Poison Control center right away. Do not use in the eyes. Discontinue use if irritation or redness develops. If condition persists, consult a doctor.

Directions

Remove cap. roll over palms and fingers to wet hands thoroughly. Allow to dry without wiping.

Aloe, Vera, Cetrimonium Chloride, Citrus Fragrance, Disodium EDTA, D and C red 33, FD and C yellow 5, Luvigel, Water

wild blue yonder label



MM15

wild blue yonder blister card

MM16



hand sanitizing roll-on

with Convenient Carabiner Clip!

Drug Facts

Active Ingredient

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Directions

Remove cap. Roll over palms and fingers to wet hands thoroughly. Allow to dry without wiping.

Inactive Ingredients

Aloe Vera, Cetrimonium chloride, DI Water, Disodium EDTA, Fragrance, Luvigel

* See Inner Label for Color Information

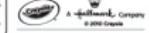
Dist. by Health-Tech, Inc. Boynton Beach, FL 33426



Health-Tech, Inc. guarantees the quality of this Crayola Product. If this product does not perform properly, contact us at: 800-600-3961
M - T 8:00 to 4:30, F 8:00 to 3:30 ET

OFFICIAL LICENSED PRODUCT

ABOUT US



075595-05025-9

www.crayola.com

Crayola roll on shipper label

HEALTH TECH, INC
3622 QUANTUM BLVD
BOYNTON BEACH, FL
33426 800-600-2861

CRAYOLA HAND SANITIZER
15 ML PORTABLE ROLL ON
ITEM: 5025

CASE PACK
12/6
CASE QTY 72 PCS

NDC 48871-006-01 24 PCS
NDC 48871-007-01 24 PCS
NDC 48871-009-01 24 PCS



00075595300033

MM17

CRAYOLA WILD BLUE YONDER ROLL ON HAND SANITIZER

benzalkonium chloride gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48871-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Benzalkonium Chloride (UNII: F5UM2KM3W7) (benzalkonium - UNII:7N6JUD5X6Y)	Benzalkonium Chloride	.0013 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Aloe vera leaf (UNII: ZY81Z83H0X)	
Cetrimonium Chloride (UNII: UC9PE95IBP)	
Edetate Disodium (UNII: 7FLD91C86K)	
water (UNII: 059QF0K00R)	

Product Characteristics

Color	red (D and C Red 33) , green (D and C green 5)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48871-006-01	4 in 1 CASE		
1		6 in 1 CARTON		
1		1 in 1 BLISTER PACK		
1		20 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333	09/01/2010	

Labeler - Health-Tech, Inc. (084007889)

Establishment

Name	Address	ID/FEI	Business Operations
Health-Tech, Inc.		084007889	manufacture

Revised: 5/2010

Health-Tech, Inc.