CRAYOLA SUNGLOW YELLOW 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.

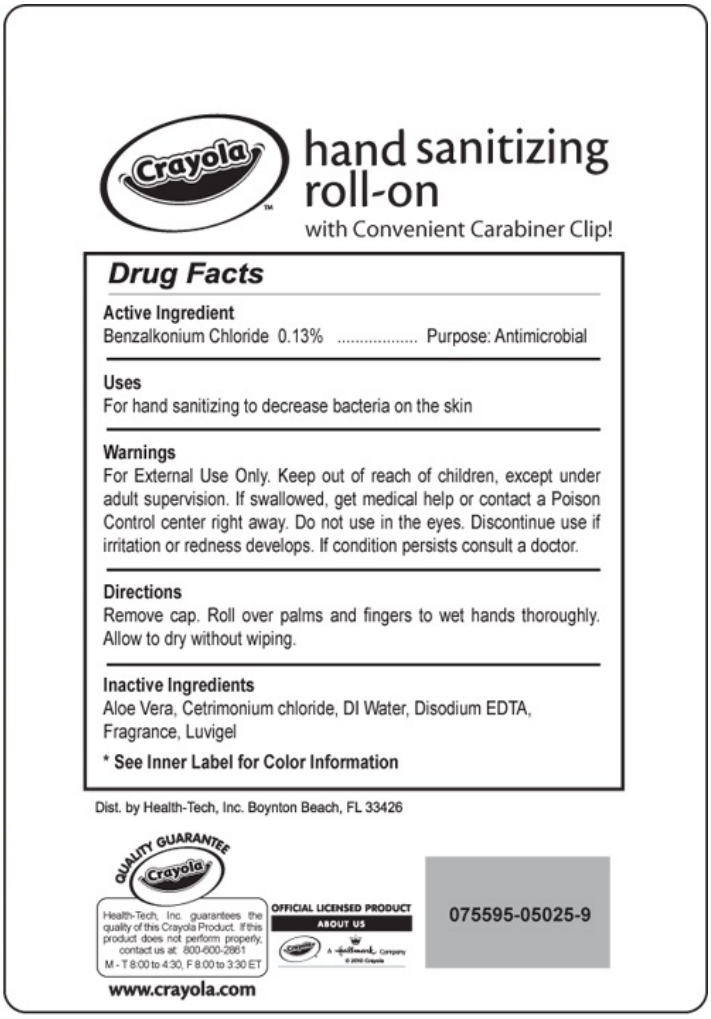
Aloe, Vera, Cetrimonium Chloride, Citrus Fragrance, Disodium EDTA, D and C red 33, FD and C yellow 5, Luvigel, Water

Crayola Sun Glow Yellow roll on label

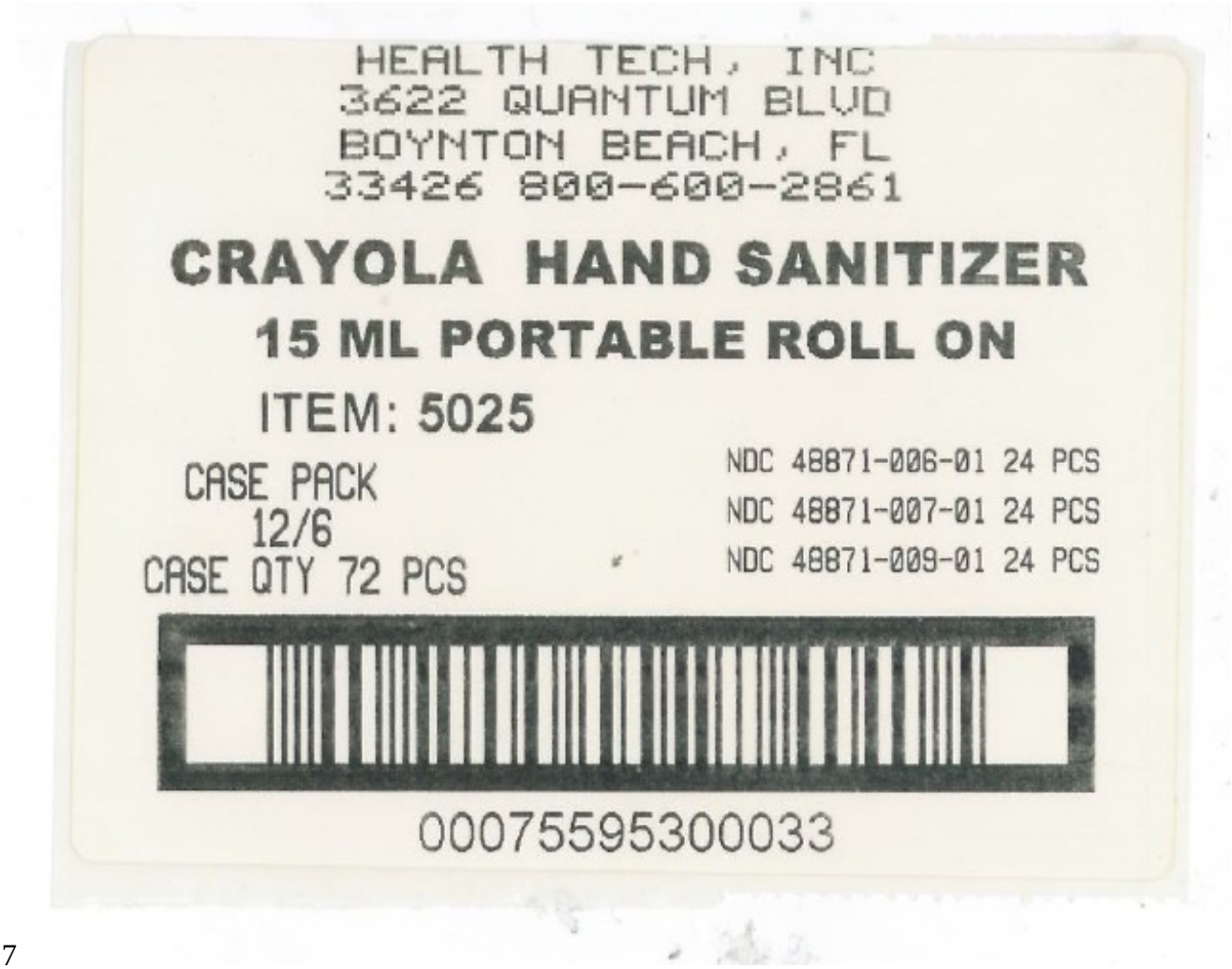
MM20

Crayola blister card

MM21



Crayola roll on shipper label



MM17

CRAYOLA SUNGLOW YELLOW ROLL ON HAND SANITIZER

benzalkonium chloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48871-009
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: 059QF0KO0R)	

Product Characteristics			
Color	yellow (FD and C Yellow 5)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48871-009-01	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.