ADVANCED HAND SANITIZER- alcohol gel Blue Cross Laboratories, 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.

Advanced Hand Sanitizer

Active Ingredient(s)

Ethyl Alcohol 70% v/v. Purpose: Antiseptic

Use

- To decrease bacteria on the skin that could cause disease.
- recommended for repeated use

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

Stop use and ask a doctor if

• Skin irritation develops

Warnings

For external use only. Flammable. Keep away from heat or flame

When using this product

- Keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- Avoid contact with broken skin
- Do not inhale or ingest

Directions

- Wet hands thoroughly with product and allow to dry without wiping
- For children under 6, use only under adult supervision
- not recommend for infants

Other information

- do not store above 105°F
- may discolor some fabrics
- harmful to wood finishes and plasics

Inactive ingredients water, glycerin, propylene glycol, carbomer

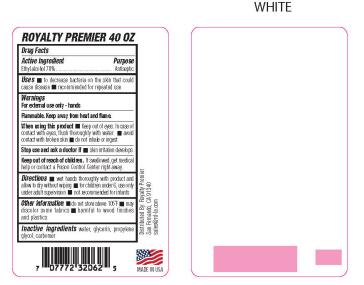
Royalty Premier

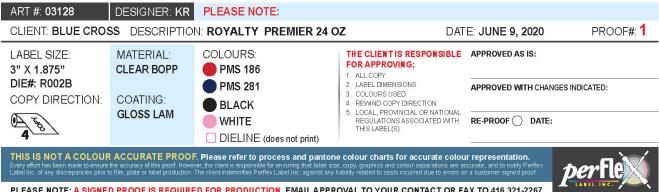
Advanced

Hand Sanitizer

Fights Germs and Bacteria

40 FL OZ (1.18 L)





PLEASE NOTE: A SIGNED PROOF IS REQUIRED FOR PRODUCTION. EMAIL APPROVAL TO YOUR CONTACT OR FAX TO 416 321-2267.



ADVANCED HAND SANITIZER

alcohol gel								
Product Informa	ition							
Product Type		HUMAN OTC DRUG	Item Code	(Source)	NDC:2	2431-254		
Route of Administr	ation	TOPICAL						
Active Ingredier	nt/Active Moi	etv						
Ingredient Name Basis of Strength								
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL					Strength mL in 100 L			
Inactive Ingredi	ents							
Ingredient Name								
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)								
GLYCERIN (UNII: PDC6A3C0OX)								
WATER (UNII: 059QF0KO0R)								
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)								
Packaging								
# Item Code		Package Description	Ma	rketing Start Date	Marke	eting End Date		
1 NDC:22431-254-01	1.18 L in 1 BOTT	LE; Type 0: Not a Combination	Product 06/	17/2020				
Marketing Information								
Marketing Catego		ion Number or Monograph (Citation M	arketing Start Date	Mark	eting End Date		
OTC monograph not f				/17/2020				

Labeler - Blue Cross Laboratories, Inc (008298879)

Establishment									
Name	Address	ID/FEI	Business Operations						
Blue Cross Laboratories, Inc		008298879	manufacture(22431-254)						

Revised: 6/2020

Blue Cross Laboratories, Inc