PAW PATROL HAND SANITIZER- ethyl alcohol liquid
Ashtel Studios, 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.

Paw Patrol Hand Sanitizer

Drug Facts

Active Ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Use To help reduce bacteria and germs on the skin.

WARNING Flammable. Keep away from fire or flame. For external use only • Stop use and ask a doctor if irritation or redness develops and persists.

• **Keep out of reach of children**. • In case of accidental digestion, seek professional assistance or contact a Poison Control Center immediately.

Directions • Place enough product in palm to cover hands and rub hands together briskly until dry. • Children under 6, use only under adult supervision. • Not recommended for infants.

Other Information • Do not store above 100°F (38°C). • May discolor some fabrics. • Harmful to wood finishes and plastics.

Inactive Ingredients • Deionized Water, Glycerine, Carbomer, Vitamin E, Fragrance, Aloe Barbadensis Gel, Triethanolamine, Propylene Glycol.

KILLS 99% OF GERMS

Smart Care®

QUESTIONS OR COMMENTS?

1-877-274-8358 Toll Free in USA • 1-909-434-0911 International COPYRIGHTS AND TRADEMARKS GRANTED OR PENDING WORLDWIDE SMARTCAREUS.COM • ONTARIO, CA **ITEM#: 70229-24**

Packaging

Drug Facts

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PAW PATROL HAND SANITIZER

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70108-018
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TROLAMINE (UNII: 9O3K93S3TK)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:70108-018-01	53 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/26/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/26/2019	

Labeler - Ashtel Studios, Inc. (148689180)