TONKA HAND SANITIZER TNK 200- alcohol gel
Gold Orient International Limited

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.

Tonka Hand Sanitizer TNK 200

Active Ingredient(s)

Alcohol 62% v/v.

Purpose

Antiseptic

Use

For hand-washing to decrease bacteria on the skin, only when water is not available

Warnings

For external use only. Flammable. Keep away from fire or flames

Do not use

- in children less than 2 months of age
- on open skin wounds

Do not get into eyes.

if contact occurs, rinse eyes thoroughly with water

irritation and redness develop

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

Directions

wet hands thoroughly with product and allow to dry without wiping

Other information

- Store between 66 to 77F (20-25C)
- Do not store above 110F (43C)
- You may report a serious adverse reaction to this product to Report Reaction, LLC, PO Box 22 Plainsboro, NJ 08536

Inactive ingredients

Water, Glycerin, Acrylates/ C10-30 Alkyl Acrylates Crosspolymer, Fragrance, Polysorbate 20, Sodium Hydroxide, Red 33, Blue1

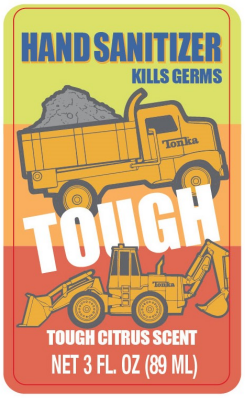
Package Label - Principal Display Panel

HAND SANITIZER

SCENT:
CITRUS

HAND SANITIZER
PANTONE
7683 @ 50%

FRONT



BACK

TONKA HAND SANITIZER TNK 200

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51522-018
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**
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51522-018-01	1 in 1 POUCH	03/30/2020	
1		89 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - Gold Orient International Limited (679905914)

Establishment

Name	Address	ID/FEI	Business Operations
Gold Orient International Limited		679905914	manufacture(51522-018)

Revised: 7/2020 Gold Orient International Limited