

**QS PLUS PLUS INSTANT HAND SANITIZER- benzealkonium chloride liquid
ABC Compounding Co., 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.

QS Plus Plus Instant Hand Sanitizer 6475 Drug Facts and Label

Drug Facts Box OTC-Active Ingredient Section

benzalkonium chloride USP 0.13%

Drug Facts Box OTC-Indications & Usage Section

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

Drug Facts Box OTC-Warnings Section

For external use only

Drug Facts Box OTC-Purpose Section

Antiseptic

Drug Facts Box-OTC When Using Section

do not get into eyes

if contact occurs, rinse eyes thoroughly with water

Drug Facts Box-OTC Stop Use Section

irritation and redness develop

Drug Facts Box-OTC Keep Out Of Reach Of Children Section

If swallowed, get medical help or contact a Poison Control Center right away

Drug Facts Box-OTC Dosage & Administration Section

press pump twice to deliver two squirts (about a quarter size) of foaming product onto the palm of your hand

rub hands together until dry

wash hands with soap and water at earliest opportunity

Drug Facts Box-OTC Inactive Ingredient Section

water, glycerine, dimethicone, DMDM hydantoin, iodopropynyl butylcarbamate, methylchloroisothiazolinone, methylisothiazolinone, fragrance

QS Plus Plus Instant Hand Sanitizer

UP TO 150 APPLICATIONS!

QS PLUS PLUS INSTANT HAND SANITIZER

KILLS 99.99% OF MOST COMMON GERMS THAT CAUSE ILLNESS IN AS LITTLE AS 15 SECONDS!

- Non-Alcoholic • Non-Flammable
- Pleasant Fragrance

Manufactured by: ABC Compounding Co., Inc.
P. O. Box 14247, Atlanta, GA 30321-0247

NET CONTENTS: 50 ml (1.7 oz)

44571407519 Batch No.: XXXX

Drug Facts

Active Ingredient

benzalkonium chloride 0.13%.....

Use for hand-washing to decrease b water is not available

Warnings

For external use only

When using this product

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III if contact occurs, rinse eyes thro

Stop use and ask a doctor if

III irritation and redness develop

Keep out of reach of children. If sv contact a Poison Control Center righ

Directions: III press pump twice a quarter size) of foaming product or III rub hands together until dry. III water at earliest opportunity

Inactive Ingredients

wa. DMDM hydantoin, iodopropynyl buty thiazolinone, methylisothiazolinone, f

QS Plus Plus Instant Hand Sanitizer

QS PLUS PLUS INSTANT HAND SANITIZER

benzealkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62257-457
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
IODOPROPYNYL BUTYL CARBAMATE (UNII: 603P14DHEB)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62257-457-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	02/20/2019	
2	NDC:62257-457-18	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/20/2019	
3	NDC:62257-457-14	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/20/2019	
4	NDC:62257-457-12	1000 mL in 1 BAG; Type 0: Not a Combination Product	02/20/2019	
5	NDC:62257-457-17	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/20/2019	
6	NDC:62257-457-16	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/20/2019	
7	NDC:62257-457-24	115 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/20/2019	
8	NDC:62257-457-15	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/20/2019	
9	NDC:62257-457-55	208200 mL in 1 DRUM; Type 0: Not a Combination Product	02/20/2019	
10	NDC:62257-457-19	18900 mL in 1 PAIL; Type 0: Not a Combination Product	05/12/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	02/20/2019	

Labeler - ABC Compounding Co., Inc. (003284353)

Registrant - ABC Compounding Co., Inc. (003284353)

Establishment

Name	Address	ID/FEI	Business Operations
ABC Compounding Co., Inc.		003284353	manufacture(62257-457)

Revised: 5/2020

ABC Compounding Co., Inc.