WINTER IN VENICE HOSPITAL GRADE HAND SANITIZER- alcohol gel WIV Global 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.

Winter in Venice HAND SANITISER

Drug Facts

Active ingredient

Alcohol Denat. 72%

Purpose

Antimicrobial

Use • Hand Sanitiser to help reduce bacteria on the skin.

Warnings

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

When using this product

• Keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor

• if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children

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

Directions • Dispense a small amount onto dry hands and massage well into skin. • Do not rinse.

Other information • Store under 110°F (43°C)

Inactive ingredients: Aqua, Glycerin, Carbomer, Rosa Multiflora Extract, Benzalkonium Chloride, Phenoxyethanol, Parfum, Triethanolamine, Ethylhexylglycerin, Citronellol, Geraniol.

Questions? +44 845 652 1223

KILLS 99.9% OF GERMS

CERTIFIED AND PASSED HOSPITAL GRADE AS PER EU STANDARDS

KILLS GERMS WITHOUT WATER

KILLS 99.9% OF BACTERIA AND CORONAVIRUS

WINTER IN VENICE HOSPITAL GRADE ANTIBACTERIAL HAND SANITISER PROVIDES LONG-LASTING ANTIMICROBIAL PROTECTION FOR THE SKIN WITHOUT SOAP AND WATER.

- EN 14476
- EN 1276
- EN 1500
- EN 1650
- EN 13624
- EN 13727
- Antibacterial according to EN 1276, EN 1500 and EN 13727
- Virucidal according to EN 14476
- Fungicidal and Yeasticidal according to EN 1650 and EN 13624

Manufactured for

Winter in Venice

Dalton House, 60 Windsor Avenue,

London, SW19 2RR. UNITED KINGDOM

www.winterinvenice.co.uk

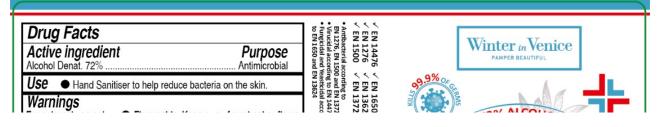
Designed and formulated in the UK

Assembled in Turkey

Packaging

Pack size - 100 mL







Pack size - 500 mL



WINTER IN VENICE HOSPITAL GRADE HAND SANITIZER

alcohol gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78609-124	
Route of Administration	TOPICAL			

ength
100 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
GLYCERIN (UNII: PDC6A3C0OX)		
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)		
ROSA MULTIFLORA WHOLE (UNII: XZ57EQ287T)		
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
TROLAMINE (UNII: 9O3K93S3TK)		
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)		
.BETACITRONELLOL, (R)- (UNII: P01OUT964K)		
GERANIOL (UNII: L837108USY)		

	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:78609-124-10	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2020		
ľ	2 NDC:78609-124-50	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/02/2020		
	3 NDC:78609-124-55	5000 mL in 1 CAN; Type 0: Not a Combination Product	05/26/2020		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	05/26/2020			

Labeler - WIV Global Limited (211563238)

Revised: 10/2020 WIV Global Limited