

BIOWASH HAND ANTISEPTIC- hi iq water spray Hand Sanitizer LLC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Active Ingredient

Hydrogen Cation .77%

Purpose

Antiseptic

Uses

For spraying on hands or hard surfaces in order to help prevent the spreading of bacteria & harmful germs.

Warnings

For external use only. Do not use if you are allergic to any ingredients. Stop use if rash develops and ask a doctor right away.

Keep Out of Reach of Children

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Directions

Spray solution on hard surfaces or hands. Allow hands to dry. For maximum efficacy on surfaces, we recommend drying with a BioFoam™ Mitt, a BioFoam™ Sponge, or a BioFoam™ Wipe.

Other Safety Information

Store under 110° F (43° C)

Inactive Ingredients

Water, Urea

BIOWASH HAND ANTISEPTIC

WATER BASED - ALCOHOL & BLEACH FREE - CHILD & PET SAFE

bioWASH



The Smart Choice!

HAND ANTISEPTIC

ERASE GERMS
HYPOALLERGENIC
NON-IRRITATING
CHILD SAFE



Made in the USA
NDC # 76701-315-02

GetBioWash.com

Powered by
Hy-IQ



1.7 FL OZ
(50.28ml)



Manufactured by Hand Sanitizer LLC, 850 Kaliste Saloom Rd, Suite 212 Lafayette, LA 7058, ©US BioSolutions LLC

Drug Facts

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BIOWASH HAND ANTISEPTIC

hi iq water spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76701-315
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROGEN CATION (UNII: 5046UKT60S) (HYDROGEN CATION - UNII:5046UKT60S)	HYDROGEN CATION	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	940 mg in 1 mL
UREA (UNII: 8W8T17847W)	50 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:76701-315-02	50.275 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/11/2021	
2	NDC:76701-315-08	237 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/11/2021	
3	NDC:76701-315-01	3785.41 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/11/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/11/2021	

Labeler - Hand Sanitizer LLC (117473019)

Establishment

Name	Address	ID/FEI	Business Operations
Hand Sanitizer LLC		117473019	manufacture(76701-315) , label(76701-315) , pack(76701-315)

Revised: 8/2021

Hand Sanitizer LLC