HYDRA PEARL HAND SANITIZER- antiseptic gel gel NOVAGEL HAND SANITIZER- antiseptic gel gel IVOGEL HAND SANITIZER- antiseptic gel gel HYDRA PEARL LIQUID- sanitizing hand spray liquid PlasmaDent 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.

Nova Gel





Advanced Hand Sanitizer

Original Scent

Leaves Hands Feeling Soft

8 FL OZ (236.6 mL)

Active Ingredient

Active Ingredient Purpose

Ethyl Alcohol 70% v/v ------Antiseptic

Purpose

Use- To decrease bacteria on the skin

Use

Use-To decrease bacteria on the skin

Keep out of the reach of children

Keep out of the reach of children

Warnings

Warnings

Flammable. Keep away from fire or flame.

For external use only

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stope use and ask a doctor if irritation or rash appears and lasts.

Keep out of reach of children. If swallowed get medical help or contact a poisen control center right away.

Directions

Directions-wet hands thoroughly with product and allow to dry without wiping. For children under 6 use only under adult supervision. Not recommended for infants.

Other information

Other information- Do not store above 110 °F (43 °C)

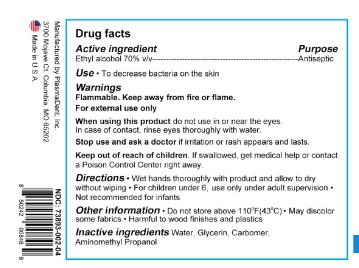
May discolor some fabrics.

Harmful to wood finishes and plastics

Inactive Ingredient Section

Inactive Ingredient- Water, Glycerin, Carbomer, Aminomethyl propanol, fragrance

Hydra Pearl







Leaves Hands Feeling Soft and Clean

10 FL OZ (296 mL)

Advanced Hand Sanitizer
Original Scent
Leaves Hands Feeling Soft
8 FL OZ (236.6 mL)

Hydra Pearl Sanitizing Spray







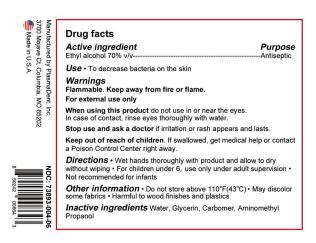


Alcohol Antiseptic • Non-Sterile • Topical Solution

1 Gallon (3785 mL)

Ivogel Hand Sanitizer

Ivogel Hand Sanitizer
70% Alcohol Unscented
Moisterizing Gel
Leaves hands feeling soft





HYDRA PEARL HAND SANITIZER

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:73893-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	0.1 g in 100 mL	
GLYCERIN (UNII: PDC6A3C0OX)	2 g in 100 mL	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	0.3 g in 100 mL	
WATER (UNII: 059QF0KO0R)	32.8 g in 100 mL	

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73893- 002-02	236.6 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	08/12/2020	
2	NDC:73893- 002-03	473.2 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	08/12/2020	
3	NDC:73893- 002-04	295.7 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	08/12/2020	
4	NDC:73893- 002-05	946.4 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	08/12/2020	
5	NDC:73893- 002-06	3800 mL in 1 CONTAINER; Type 0: Not a Combination Product	08/12/2020	
6	NDC:73893- 002-11	59.1 mL in 1 TUBE; Type 0: Not a Combination Product	08/12/2020	
7	NDC:73893- 002-08	18900 mL in 1 CONTAINER; Type 0: Not a Combination Product	10/02/2020	
8	NDC:73893- 002-12	189000 mL in 1 DRUM; Type 0: Not a Combination Product	02/28/2022	

Marketing Information				
Marketing Application Number or Monograph Marketing Start Marketing End Category Citation Date Date				
OTC monograph not final	part333A	08/12/2020		

NOVAGEL HAND SANITIZER

antiseptic gel gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:73893-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M)	ALCOHOL	70 ml in 100 ml

Inactive Ingredients	
Ingredient Name	Strength
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	0.1 g in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)	2 g in 100 mL
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	0.3 g in 100 mL
WATER (UNII: 059QF0KO0R)	32.8 g in 100 mL

Pa	ckaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73893- 001-01	59.1 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/30/2020	
2	NDC:73893- 001-02	236.6 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/30/2020	
3	NDC:73893- 001-03	473.2 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	04/06/2020	
4	NDC:73893- 001-04	295.7 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	04/15/2020	
5	NDC:73893- 001-05	946.4 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	04/15/2020	
6	NDC:73893- 001-06	3800 mL in 1 CONTAINER; Type 0: Not a Combination Product	04/15/2020	
7	NDC:73893- 001-07	9500 mL in 1 CONTAINER; Type 0: Not a Combination Product	04/15/2020	
8	NDC:73893- 001-08	18900 mL in 1 CONTAINER; Type 0: Not a Combination Product	04/15/2020	
9	NDC:73893- 001-09	113600 mL in 1 DRUM; Type 0: Not a Combination Product	04/15/2020	
10	NDC:73893- 001-10	189300 mL in 1 DRUM; Type 0: Not a Combination Product	04/15/2020	
11	NDC:73893- 001-11	59.1 mL in 1 TUBE; Type 0: Not a Combination Product	07/20/2020	
12	NDC:73893- 001-12	354.9 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	08/26/2020	

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

IVOGEL HAND SANITIZER

antiseptic gel gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:73893-004

Route of Administration TOPICAL

Active Ingredient/Active Moiety

- 10 11 g. Canoni, 10 11 C. J.		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	0.1 g in 100 mL	
GLYCERIN (UNII: PDC6A3C0OX)	2 g in 100 mL	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	0.3 g in 100 mL	
WATER (UNII: 059QF0KO0R)	32.8 g in 100 mL	

Pa	ckaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73893- 004-01	59.1 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	10/23/2020	
2	NDC:73893- 004-02	236.6 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	10/23/2020	
3	NDC:73893- 004-03	473.2 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	10/23/2020	
4	NDC:73893- 004-04	295.7 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	10/23/2020	
5	NDC:73893- 004-05	946.4 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	10/23/2020	
6	NDC:73893- 004-06	3800 mL in 1 CONTAINER; Type 0: Not a Combination Product	10/23/2020	
7	NDC:73893- 004-07	9500 mL in 1 CONTAINER; Type 0: Not a Combination Product	10/23/2020	
8	NDC:73893- 004-08	18900 mL in 1 CONTAINER; Type 0: Not a Combination Product	10/23/2020	
9	NDC:73893- 004-09	113600 mL in 1 DRUM; Type 0: Not a Combination Product	10/23/2020	
10	NDC:73893- 004-10	189300 mL in 1 DRUM; Type 0: Not a Combination Product	10/23/2020	
11	NDC:73893- 004-11	59.1 mL in 1 TUBE; Type 0: Not a Combination Product	10/23/2020	
12	NDC:73893- 004-12	354.9 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	10/23/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/23/2020	

HYDRA PEARL LIQUID

sanitizing hand spray liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:73893-003

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

80 mL in 100 mL

Inactive Ingredients

Ingredient Name

GLYCERIN (UNII: PDC6A3C0OX)

Strength

1.45 mL in 100 mL

WATER (UNII: 059QF0KOOR)

18.55 mL in 100 mL

Packaging Marketing Start Marketing End Package Description Item Code **Date Date** 1 NDC:73893-3785 mL in 1 BOTTLE; Type 0: Not a Combination 10/09/2020 003-01 Product 2 NDC:73893-946 mL in 1 BOTTLE; Type 0: Not a Combination 11/09/2020 003-02 **Product** 3 NDC:73893-473 mL in 1 BOTTLE; Type 0: Not a Combination 11/09/2020 003-03 4 NDC:73893-227304 mL in 1 DRUM; Type 0: Not a Combination 02/28/2022 003-04 Product

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/09/2020	

Revised: 2/2022 PlasmaDent Inc