CLINDACARE- clindamycin, ibuprofen, chlorhexidine gluconate
Brisk Pharmaceuticals

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.

ClindaCare

Rx Only

NDC 73614-922-03

ClindaCare™

Each Pack Contains:

Clindamycin Hydrochloride Capsules, USP 150 mg - One 30 ct Bottle

Ibuprofen Tablets, USP 800 mg - One 30 ct Bottle

Chlorhexidine Gluconate Oral Rinse, USP 0.12% - Two 118 ml Bottles

Brisk Pharmaceuticals

Packaged by:

Unit Dose Solutions Inc.,

Morrisville, NC 27560

Distributed by:

Brisk pharmaceuticals,

Richardson, TX 75081

Keep out of reach of children.

Storage: Store at 20°C to 25°C

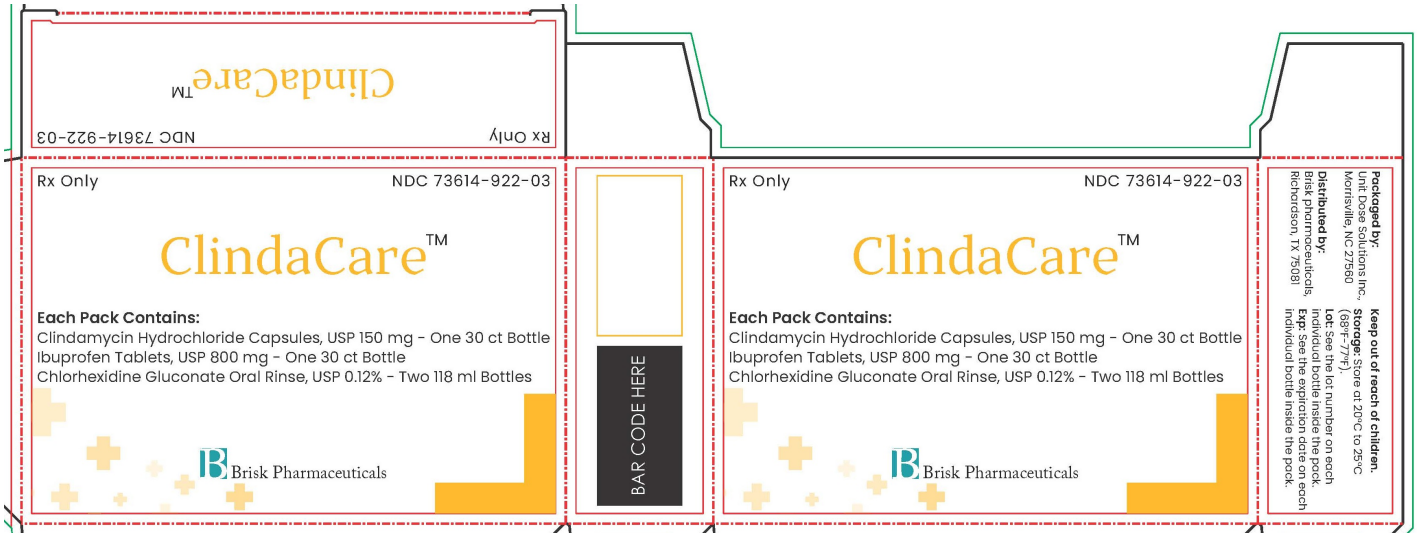
(68°F-77°F).

Lot: See the lot number on each

individual bottle inside the pack.

Exp: See the expiration date on each

individual bottle inside the pack.



Each capsule contains:
Clindamycin Hydrochloride equivalent to 150 mg Clindamycin

Keep out of reach of children
Store at 20°C - 25°C (68°F - 77°F)

This medication bottle is intended for distribution only as part of ClindaCare™ pack.

73614-204-03

CLINDAMYCIN HYDROCHLORIDE Capsules, USP

150 mg

Rx Only 30 Capsules

Brisk Pharmaceuticals

Repackaged
Unit Dose S
Morrisville,
Distributed
Brisk Pharm
Richardson

Each film coated tablet contains:
Ibuprofen, USP 800 mg

Keep out of reach of children
Store at 20°C - 25°C (68°F - 77°F)

This medication bottle is intended for distribution only as part of CRAPAZ™ pack.

73614-201-03

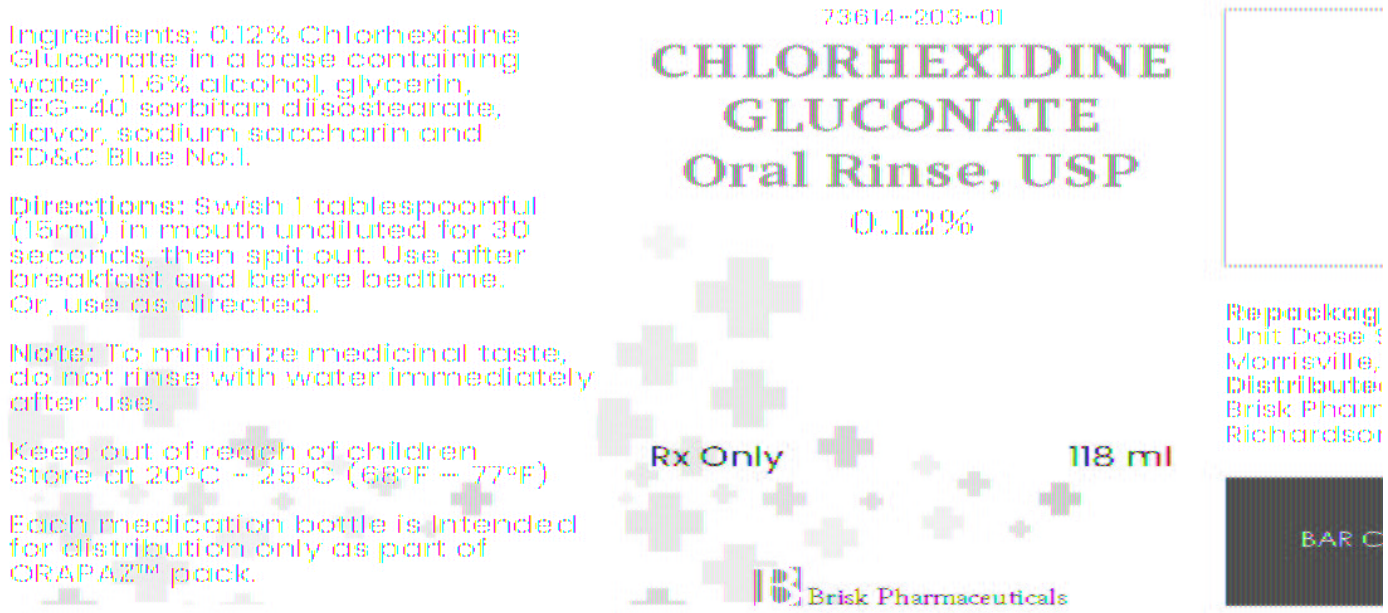
IBUPROFEN Tablets, USP

800 mg

Rx Only 30 Tablets

Brisk Pharmaceuticals

Repackaged
Unit Dose S
Morrisville,
Distributed
Brisk Pharm
Richardson



CLINDACARE

clindamycin, ibuprofen, chlorhexidine gluconate kit

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:73614-922

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73614-922-03	1 in 1 CARTON; Type 1: Convenience Kit of Co-Package	02/10/2025	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	30
Part 2	1 BOTTLE	118 mL
Part 3	1 BOTTLE	118 mL
Part 4	1 BOTTLE	30

Part 1 of 4

IBUPROFEN

ibuprofen tablet, film coated

Product Information

Item Code (Source)	NDC:73614-201
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	800 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
TALC (UNII: 7SEV7J4R1U)	
STARCH, CORN (UNII: O8232NY3SJ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white	**Score**	no score
Shape	OVAL (CAPLET)	**Size**	19mm
Flavor		**Imprint Code**	BI8
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73614-201-03	30 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA202413	05/23/2024	

Part 2 of 4

CHLORHEXIDINE GLUCONATE ORAL RINSE

chlorhexidine gluconate oral rinse solution

Product Information	
Item Code (Source)	NDC:73614-203
Route of Administration	DENTAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E) (CHLORHEXIDINE - UNII:R4KO0DY52L)	CHLORHEXIDINE GLUCONATE	1.2 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
PEG-40 SORBITAN DIISOSTEARATE (UNII: JL4CCU7I1G)	

Product Characteristics			
Color	blue	**Score**	
Shape		**Size**	
Flavor	MINT	**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73614-203-01	118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075561	07/27/2023	

Part 3 of 4

CHLORHEXIDINE GLUCONATE ORAL RINSE

chlorhexidine gluconate oral rinse solution

Product Information	
Item Code (Source)	NDC:73614-203
Route of Administration	DENTAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E) (CHLORHEXIDINE - UNII:R4KO0DY52L)	CHLORHEXIDINE GLUCONATE	1.2 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
PEG-40 SORBITAN DIISOSTEARATE (UNII: JL4CCU7I1G)	

Product Characteristics			
Color	blue	**Score**	
Shape		**Size**	
Flavor	MINT	**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73614-203-01	118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075561	07/27/2023	

Part 4 of 4

CLINDAMYCIN HYDROCHLORIDE

clindamycin hydrochloride capsule

Product Information

Item Code (Source)	NDC:73614-204
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CLINDAMYCIN HYDROCHLORIDE (UNII: T20OQ1YN1W) (CLINDAMYCIN - UNII:3U02EL437C)	CLINDAMYCIN	150 mg

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B71O)	
TALC (UNII: 7SEV7J4R1U)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	

Product Characteristics

Color	green (LIGHT GREEN) , blue (LIGHT BLUE)	**Score**	no score
Shape	CAPSULE	**Size**	19mm
Flavor		**Imprint Code**	G;143
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73614-204-03	30 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA216957	03/10/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		02/10/2025	

Labeler - Brisk Pharmaceuticals (117250794)

Revised: 2/2025 Brisk Pharmaceuticals