BSS - balanced salt solution solution Alcon Laboratories, Inc.

BSS[®]
Sterile Irrigating Solution
(balanced salt solution)

DESCRIPTION:

BSS[®] Sterile Irrigating Solution is a sterile balanced salt solution, each mL containing sodium chloride (NaCl) 0.64%, potassium chloride (KCl) 0.075%, calcium chloride dihydrate (CaCl₂•2H₂O) 0.048%, magnesium chloride hexahydrate (MgCl₂•6H₂O) 0.03%, sodium acetate trihydrate (C₂H₃NaO₂•3H₂O) 0.39%, sodium citrate dihydrate (C₆H₅Na₃O₇•2H₂O) 0.17%, sodium hydroxide and/or hydrochloric acid (to adjust pH), and water for injection. The pH is approximately 7.5. The osmolality is approximately 300 mOsm/Kg.

CLINICAL PHARMACOLOGY:

BSS Sterile Irrigating Solution is an isotonic solution for use in irrigating tissues of the eyes.

INDICATIONS AND USAGE:

For use as an extraocular and intraocular irrigating solution during ocular surgical procedure involving perfusion of the eye with an expected maximum duration of less than 60 minutes.

WARNINGS:

- NOT FOR INJECTION OR INTRAVENOUS INFUSION.
- Do not use unless outer overwrap is intact, product is clear, seal is intact and bag is undamaged.
- Do not use if product is discolored or contains a precipitate.
- SINGLE patient use only. The contents of this bag should not be used in more than one patient.
- This solution contains no preservative, unused contents should be discarded.

PRECAUTIONS:

Open under aseptic conditions only.

Studies suggest that intraocular irrigating solutions which are iso-osmotic with normal aqueous fluids should be used with caution in diabetic patients undergoing vitrectomy since intraoperative lens changes have been observed.

There have been reports of corneal clouding and edema following ocular surgery in which BSS Sterile Irrigating Solution was used as an irrigating solution.

ADVERSE REACTIONS

Irrigation or any other trauma may result in corneal swelling or bullous keratopathy.

Post-operative inflammatory reactions as well as incidents of corneal edema and corneal decompensation have been reported.

DOSAGE AND ADMINISTRATION

This irrigating solution should be used according to standard format for each surgical procedure. Follow directions of the particular administration set to be used. Remove the outer overwrap. Clean and disinfect the rubber stopper by using a sterile alcohol wipe. Insert the spike aseptically into the bag through the target area of the rubber stopper. Allow the fluid to flow and remove air from the tubing before irrigation begins.

HOW SUPPLIED

BSS[®] Sterile Irrigating Solution is supplied in a clear polypropylene bag using a grey butyl stopper and aluminum seal packaged in a clear, multilayered plastic outer overwrap.

500 mL in a bag: **NDC** 0065-1795-04.

STORAGE: Store at 36° - 77°F (2° - 25°C).

9012628-1115

Alcon®

A Novartis company

Distributed by:

Alcon Laboratories, Inc. Fort Worth, Texas 76134 USA © 2012, 2015 Novartis

PRINCIPAL DISPLAY PANEL SINGLE PATIENT USE ONLY

Rx Only

NDC 0065-1795-04

BSS®

Sterile Irrigating Solution

(balanced salt solution)

Each mL contains: sodium chloride 0.64%, potassium chloride 0.075%, calcium chloride ${}^{\bullet}2H_2O$ 0.048%, magnesium chloride ${}^{\bullet}6H_2O$ 0.03%, sodium acetate ${}^{\bullet}3H_2O$ 0.39%, sodium citrate ${}^{\bullet}2H_2O$ 0.17%, sodium hydroxide and/or hydrochloric acid (to adjust pH), and water for injection. The pH is approximately 7.5. The osmolality is approximately 300 mOsm/Kg.

NOT FOR I.V. USE

Read accompanying insert.

WARNINGS: NOT FOR INJECTION OR INTRAVENOUS INFUSION. Do not use unless outer overwrap is intact, product is clear, seal is intact and bag is undamaged. Do not use if product is discolored or contains a precipitate. Discard unused contents. Do not use this bag for more than one patient.

STORAGE: Store at 36° - 77° F (2° - 25°C).

STERILE 500 mL

Alcon®

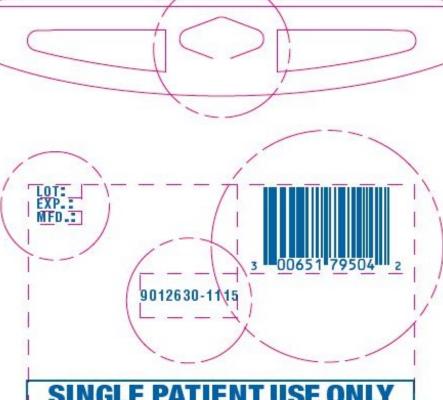
Alcon Laboratories, Inc. Fort Worth, Texas 76134 USA

© 2012, 2015 Novartis

LOT: EXP.:

MFD.:

9012630-1115



SINGLE PATIENT USE ONLY

Rx Only

NDC 0065-1795-04

BSS® Sterile Irrigating Solution

(balanced salt solution)

Each mL contains: sodium chloride 0.64%, potassium chloride 0.075%, calcium chloride •2H2 0 0.048%, magnesium chloride •6H2O 0.03%, sodium acetate • 3H2O 0.39%, sodium citrate • 2H2O 0.17%, sodium hydroxide and/or hydrochloric acid (to adjust pH), and water for injection. The pH is approximately 7.5. The osmolality is approximately 300 m0sm/Kg.

NOT FOR I.V. USE

Read accompanying insert.

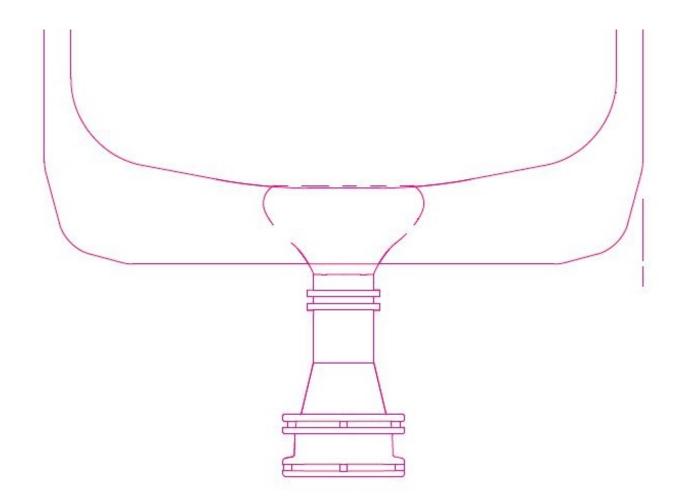
WARNINGS: NOT FOR INJECTION OR INTRAVENOUS INFUSION. Do not use unless outer overwrap is intact, product is clear, seal is intact and bag is undamaged. Do not use if product is discolored or contains a precipitate. Discard unused contents. Do not use this bag for more than one patient. STORAGE: Store at 36° - 77° F (2° - 25°C).

STERILE

500 m L

Alcon Laboratories, Inc. Fort Worth, Texas 76134 USA © 2012, 2015 Novartis





Storage: Store at 36° - 77°F (2° - 25°C).

Do not use unless outer overwrap is intact.

9012629-1115

Storage: Store at 36° - 77°F (2° - 25°C).

Do not use unless outer overwrap is intact.

9012629-1115

BSS

balanced salt solution solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0065-1795
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37) SODIUM CHLORIDE 6.4 mg in 1 mL POTASSIUM CHLORIDE (UNII: 660 YQ98 I10) (POTASSIUM CATION -POTASSIUM CHLORIDE | 0.75 mg in 1 mL UNII:295O53K152) 0.48 mg CALCIUM CHLORIDE (UNII: M4I0 D6 VV5M) (CALCIUM CATION - UNII:2M8 3C4R6 ZB) CALCIUM CHLORIDE in 1 mL MAGNESIUM CHLORIDE (UNII: 02F3473H9O) (MAGNESIUM CATION -MAGNESIUM 0.3 mg in 1 mL UNII:T6 V3LHY838) **CHLORIDE** SODIUM ACETATE (UNII: 4550 K0 SC9B) (Sodium Cation - UNII:LYR4M0 NH37) **SODIUM ACETATE** $3.9 \ mg \ in \ 1 \ mL$ SODIUM CITRATE (UNII: 1Q73Q2JULR) (SODIUM CATION - UNII:LYR4M0 NH37) SODIUM CITRATE 1.7 mg in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
SO DIUM HYDRO XIDE (UNII: 55X0 4QC32I)		
HYDRO CHLO RIC ACID (UNII: QTT17582CB)		
WATER (UNII: 059QF0KO0R)		

ı	Packaging			
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:0065-1795-04	500 mL in 1 BAG; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA020742	03/28/1969	

Labeler - Alcon Laboratories, Inc. (008018525)

Registrant - Alcon Laboratories, Inc. (008018525)

Establishment				
Name	Address	ID/FEI	Business Operations	
Alcon Research Ltd		007672236	MANUFACTURE(0065-1795)	

Revised: 1/2016 Alcon Laboratories, Inc.