

AquaLase™ Solution
(balanced salt solution)
Sterile

DESCRIPTION: AquaLase™ 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: AquaLase™ 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 package is clear, seal is intact, container is undamaged.
- Do not use if product is discolored or contains a precipitate.
- SINGLE patient use only. The contents of this bottle should not be used in more than one patient.
- The use of additives with this solution may cause corneal decompensation.
- 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 or edema following ocular surgery in which AquaLase™ Solution was used as an irrigating solution.

ADVERSE REACTIONS: Irrigation or any trauma to the corneal endothelium may result in corneal swelling or bullous keratopathy.

Postoperative inflammatory reactions as well as incidents of corneal edema and corneal decompensation have been reported.

DOSAGE AND ADMINISTRATION: The irrigating solution should be used according to standard format for each surgical procedure. **Note:** Pull the tab to remove the outer aluminum ring and dust cover. Insert the bottle into the AquaLase™ Liquefaction Device. See the equipment directions for specific instructions.

HOW SUPPLIED: 90 mL of solution in a 100 mL polypropylene single dose bottle which contains a collapsible bag with a rubber stopper and aluminum seal.

NDC 0065-0795-90

STORAGE: Store at 36° to 77°F (2° to 25°C). DO NOT FREEZE.

Rx Only

Alcon logo

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Lot:	Each mL containing sodium chloride 0.64%, potassium chloride 0.075%, calcium chloride ·2H ₂ O 0.048%, magnesium chloride ·6H ₂ O 0.03%, sodium acetate ·3H ₂ O 0.39%, sodium citrate ·2H ₂ O 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. STERILE 90 mL SINGLE USE ONLY [NO LATEX SYMBOL]	Rx Only NOT FOR IV USE SINGLE DOSE UNIT Read accompanying insert. WARNINGS: NOT FOR INJECTION OR INTRAVENOUS INFUSION. Do not use unless product is clear, seal is intact and container is undamaged. Do not use if product is discolored or contains a precipitate. Discard unused contents. Do not use this container on more than one patient. Do not use additives with this product. Tissue damage could result if other drugs are added to product.
Exp:		STORAGE: Store at 36° - 77°F (2° - 25°C). DO NOT FREEZE.
Barcode		ALCON LOGO Alcon Laboratories, Inc. Fort Worth Texas 76134 USA Printed in USA ©2002 Alcon, Inc. Par Number-Revision Date