VERION® IMAGE GUIDED SYSTEM IMPORTANT PRODUCT INFORMATION/VERION® REFERENCE UNIT AND VERION® DIGITAL MARKER. CAUTION: Federal (USA) law restricts this device to sale by, or on the order of, a physician. **INTENDED USES:** The Verion® Reference Unit is a preoperative measurement device that captures and utilizes a high-resolution reference image of a patient's eye. In addition, the Verion® Reference Unit provides preoperative surgical planning functions to assist the surgeon with planning cataract surgical procedures. The Verion® Reference Unit also supports the export of the reference image, preoperative measurement data, and surgical plans for use with the Verion® Digital Marker and other compatible devices through the use of a USB memory stick. The Verion® Digital Marker links to compatible surgical microscopes to display concurrently the reference and microscope images, allowing the surgeon to account for lateral and rotational eye movements. In addition, details from the Verion® Reference Unit surgical plan can be overlaid on a computer screen or the physician's microscope view. **CONTRAINDICATIONS:** The following conditions may affect the accuracy of surgical plans prepared with the Verion® Reference Unit: a pseudophakic eye, eye fixation problems, a non-intact cornea, or an irregular cornea. In addition, patients should refrain from wearing contact lenses during the reference measurement as this may interfere with the accuracy of the measurements. The following conditions may affect the proper functioning of the Verion® Digital Marker: changes in a patient's eye between preoperative measurement and surgery, an irregular elliptic limbus (e.g., due to eye fixation during surgery, and bleeding or bloated conjunctiva due to anesthesia). In addition, the use of eye drops that constrict sclera vessels before or during surgery should be avoided. **WARNINGS:** Only properly trained personnel should operate the Verion® Reference Unit and Verion® Digital Marker. Use only the provided medical power supplies and data communication cable. Power supplies for the Verion® Reference Unit and the Verion® Digital Marker must be uninterruptible. Do not use these devices in combination with an extension cord. Do not cover any of the component devices while turned on. The Verion® Reference Unit uses infrared light. Unless necessary, medical personnel and patients should avoid direct eye exposure to the emitted or reflected beam. **PRECAUTIONS:** To ensure the accuracy of Verion® Reference Unit measurements, device calibration and the reference measurement should be conducted in dimmed ambient light conditions. Only use the Verion® Digital Marker in conjunction with compatible surgical microscopes. **ATTENTION:** Refer to the user manuals for the Verion® Reference Unit and the Verion® Digital Marker for a complete description of proper use and maintenance of these devices, as well as a complete list of contraindications, warnings and precautions.

LENSX® LASER IMPORTANT PRODUCT INFORMATION FOR CATARACT TREATMENT. CAUTION: United States Federal Law restricts this device to sale and use by or on the order of a physician or licensed eye care practitioner. **INDICATION:** The LenSx® Laser is indicated for use in patients undergoing cataract surgery for removal of the crystalline lens. Intended uses in cataract surgery include anterior capsulotomy, phacofragmentation, and the creation of single plane and multi-plane arc cuts/incisions in the cornea, each of which may be performed either individually or consecutively during the same procedure. **RESTRICTIONS:** Patients must be able to lie flat and motionless in a supine position. Patient must be able to understand and give an informed consent. Patients must be able to tolerate local or topical anesthesia. Patients with elevated IOP should use topical steroids only under close medical supervision. **CONTRAINDICATIONS:** Corneal disease that precludes applanation of the cornea or transmission of laser light at 1030 nm wavelength. Descemetocele with impending corneal rupture. Presence of blood or other material in the anterior chamber. Poorly dilating pupil, such that the iris is not peripheral to the intended diameter for the capsulotomy. Conditions which would cause inadequate clearance between the intended capsulotomy depth and the endothelium (applicable to capsulotomy only). Previous corneal incisions that might provide a potential space into which the gas produced by the procedure can escape. Corneal thickness requirements that are beyond the range of the system. Corneal opacity that would interfere with the laser beam. Hypotony or the presence of a corneal implant. Residual, recurrent, active ocular or eyelid disease, including any corneal abnormality (for example, recurrent corneal erosion, severe basement membrane disease). History of lens or zonular instability. Any contraindication to cataract or keratoplasty. This device is not intended for use in pediatric surgery. **WARNINGS:** The LenSx® Laser System should only be operated by a physician trained in its use. The LenSx® Laser delivery system employs one sterile disposable Patient Interface consisting of an applanation lens and suction ring. The Patient Interface is intended for single use only. The disposables used in conjunction with ALCON® instrument products constitute a complete surgical system. Use of disposables other than those manufactured by Alcon may affect system performance and create potential hazards. The physician should base patient selection criteria on professional experience, published literature, and educational courses. Adult patients should be scheduled to undergo cataract extraction. **PRECAUTIONS:** Do not use cell phones or pagers of any kind in the same room as the LenSx® Laser. Discard used Patient Interfaces as medical waste. **COMPLICATIONS:** Capsulotomy, phacofragmentation, or cut or incision decentration. Incomplete or interrupted capsulotomy, fragmentation, or corneal incision procedure. Capsular tear. Corneal abrasion or defect. Pain. Infection. Bleeding. Damage to intraocular structures. Anterior chamber fluid leakage, anterior chamber collapse. Elevated pressure to the eye. **ATTENTION:** Refer to the LenSx® Laser Operator's Manual for a complete listing of indications, warnings and precautions.

CENTURION® VISION SYSTEM IMPORTANT PRODUCT INFORMATION. CAUTION: Federal (USA) law restricts this device to sale by, or on the order of, a physician. As part of a properly maintained surgical environment, it is recommended that a backup IOL Injector be made available in the event the AutoSert® IOL Injector Handpiece does not perform as expected. **INDICATION:** The Centurion® Vision System is indicated for emulsification, separation, irrigation, and aspiration of cataracts, residual cortical material and lens epithelial cells, vitreous aspiration and cutting associated with anterior vitrectomy, bipolar coagulation, and intraocular lens injection. The AutoSert® IOL Injector Handpiece is intended to deliver qualified AcrySof® intraocular lenses into the eye following cataract removal. The AutoSert® IOL Injector Handpiece achieves the functionality of injection of intraocular lenses. The AutoSert® IOL Injector Handpiece is indicated for use with the AcrySof® lenses SN60WF, SN6AD1, SN6AT3 through SN6AT9, as well as approved AcrySof® lenses that are specifically indicated for use with this inserter, as indicated in the approved labeling of those lenses. **WARNINGS:** Appropriate use of Centurion® Vision System parameters and accessories is important for successful procedures. Use of low vacuum limits, low flow rates, low bottle heights, high power settings, extended power usage, power usage during occlusion conditions (beeping tones), failure to sufficiently aspirate viscoelastic prior to using power, excessively tight incisions, and combinations of the above actions may result in significant temperature increases at incision site and inside the eye, and lead to severe thermal eye tissue damage. Good clinical practice dictates the testing for adequate irrigation and aspiration flow prior to entering the eye. Ensure that tubings are not occluded or pinched during any phase of operation. The consumables used in conjunction with ALCON® instrument products constitute a complete surgical system. Use of consumables and handpieces other than those manufactured by Alcon may affect system performance and create potential hazards. **AEs/COMPLICATIONS:** Inadvertent actuation of Prime or Tune while a handpiece is in the eye can create a hazardous condition that may result in patient injury. During any ultrasonic procedure, metal particles may result from inadvertent touching of the ultrasonic tip with a second instrument. Another potential source of metal particles resulting from any ultrasonic handpiece may be the result of ultrasonic energy causing micro abrasion of the ultrasonic tip. **ATTENTION:** Refer to the Directions for Use and Operator's Manual for a complete listing of indications, warnings, cautions and notes.

DUOVISC® OVD BRIEF STATEMENT. DESCRIPTION: DUOVISC® Viscoelastic System is designed to give two Viscoelastic materials with different physico-chemical properties that can be used differently and/or sequentially to perform specific tasks during a cataract procedure. DUOVISC® Viscoelastic System consists of VISCOAT® Ophthalmic Viscosurgical Device and PROVISC® Ophthalmic Viscosurgical Device. **CAUTION:** Federal (USA) law restricts this device to sale by, or on the order of, a physician. **DESCRIPTION:** VISCOAT® (Sodium Chondroitin Sulfate – Sodium Hyaluronate) Ophthalmic Viscosurgical Device **INDICATIONS:** VISCOAT® OVD is indicated for use as an ophthalmic surgical aid in anterior segment procedures including cataract extraction and intraocular lens (IOL) implantation. VISCOAT® OVD maintains a deep anterior chamber during anterior segment surgeries, enhances visualization during the surgical procedure, and protects the corneal endothelium and other ocular tissues. The viscoelasticity of the solution maintains the normal position of the vitreous face and prevents formation of a flat chamber during surgery. **WARNINGS:** Failure to follow assembly instructions or use of an alternate cannula may result in cannula detachment and potential patient injury. **PRECAUTIONS:** Precautions are limited to those normally associated with the surgical procedure being performed. Although sodium hyaluronate and sodium chondroitin sulfate are highly purified biological polymers, the physician should be aware of the potential allergic risks inherent in the use of any biological material. **ADVERSE REACTIONS:** VISCOAT® OVD has been extremely well tolerated in human and animal studies. A transient rise in intraocular pressure in the early postoperative period may be expected due to the presence of sodium hyaluronate, which has been shown to effect such a rise. It is therefore recommended that VISCOAT® OVD be removed from the anterior chamber by thorough irrigation and/or aspiration at the end of surgery to minimize postoperative IOP increases. Do not overfill anterior chamber. **ATTENTION:** Please refer to the directions for use for a complete listing of indications, warnings and precautions. **DESCRIPTION:** PROVISC® (Sodium Hyaluronate) Ophthalmic Viscosurgical Device **INDICATIONS:** PROVISC® OVD is indicated for use as an ophthalmic surgical aid in the anterior segment during cataract extraction and intraocular lens (IOL) implantation. Ophthalmic viscoelastics serve to maintain a deep anterior chamber during anterior segment surgery allowing reduced trauma to the corneal endothelium and surrounding ocular tissues. They help push back the vitreous face and prevent formation of a flat chamber during surgery. **PRECAUTIONS:** Postoperative increases in intraocular pressure have been reported with sodium hyaluronate products. The IOP should be carefully monitored and appropriate therapy instituted if significant increases should occur. It is recommended that PROVISC® OVD be removed by irrigation and/or aspiration at the close of surgery. Do not overfill anterior chamber. Although sodium hyaluronate is a highly purified biological polymer, the physician should be aware of the potential allergic risks inherent in the use of any biological material; care should be used in patients with hypersensitivity to any components in this material. Cannula assembly instructions should be followed to prevent patient injury. **ADVERSE REACTIONS:** Postoperative inflammatory reactions such as hypopyon and iritis have been reported with the use of ophthalmic viscoelastics, as well as incidents of corneal edema, corneal decompensation, and a transient rise in intraocular pressure. It is therefore recommended that PROVISC® OVD be removed from the anterior chamber by thorough irrigation and/or aspiration at the end of surgery to minimize postoperative IOP increases. Do not overfill anterior chamber. **ATTENTION:** Please refer to the directions for use for a complete listing of indications, warnings and precautions.

ACRYSOF® IQ INTRAOCULAR LENSES IMPORTANT PRODUCT INFORMATION. CAUTION: Federal (USA) law restricts this device to the sale by or on the order of a physician. **INDICATIONS:** The AcrySof® IQ posterior chamber intraocular lens is intended for the replacement of the human lens to achieve visual correction of aphakia in adult patients following cataract surgery. This lens is intended for placement in the capsular bag. **WARNINGS/PRECAUTIONS:** Careful preoperative evaluation and sound clinical judgment should be used by the surgeon to decide the risk/benefit ratio before implanting a lens in a patient with any of the conditions described in the Directions for Use labeling. Caution should be used prior to lens encapsulation to avoid lens decentrations or dislocations. Studies have shown that color vision discrimination is not adversely affected in individuals with the AcrySof® Natural IOL and normal color vision. The effect on vision of the AcrySof® Natural IOL in subjects with hereditary color vision defects and acquired color vision defects secondary to ocular disease (e.g., glaucoma, diabetic retinopathy, chronic uveitis, and other retinal or optic nerve diseases) has not been studied. Do not resterilize; do not store over 45° C; use only sterile irrigating solutions such as BSS® or BSS PLUS® Sterile Intraocular Irrigating Solutions. **ATTENTION:** Reference the Directions for Use labeling for a complete listing of indications, warnings and precautions.

ORA™ SYSTEM IMPORTANT PRODUCT INFORMATION. CAUTION: Federal (USA) law restricts this device to the sale by or on the order of a physician. **INTENDED USE:** The ORA™ System uses wavefront aberrometry data in the measurement and analysis of the refractive power of the eye (i.e., sphere, cylinder and axis measurements) to support cataract surgical procedures. **CONTRAINDICATIONS:** The ORA™ System is contraindicated for patients: who have progressive retinal pathology such as diabetic retinopathy, macular degeneration or any other pathology that the physician deems would interfere with patient fixation; who have corneal pathology such as Fuchs' EBMD, keratoconus, advanced pterygium impairing the cornea or any other pathology that the physician deems would interfere with the measurement process; whose preoperative regimen includes residual viscous substances left on the corneal surface such as lidocaine gel or viscoelastics; with visually significant media opacity (such as prominent floaters or asteroid hyalosis) what will either limit or prohibit the measurement process; or who have received retro or peribulbar block or any other treatment that impairs their ability to visualize the fixation light. In addition, utilization of iris hooks during an ORA™ System image capture is contraindicated, because the use of iris hooks will yield inaccurate measurements. **WARNINGS AND PRECAUTIONS:** Significant central corneal irregularities resulting in higher order aberrations might yield inaccurate refractive measurements. Post refractive keratectomy eyes might yield inaccurate refractive measurement. The safety and effectiveness of using the data from the ORA™ System have not been established for determining treatments involving higher order aberrations of the eye such as coma and spherical aberrations. The ORA™ System is intended for use by qualified health personnel only. Improper use of this device may result in exposure to dangerous voltage or hazardous laser-like radiation exposure. Do not operate the ORA™ System in the presence of flammable anesthetics or volatile solvents such as alcohol or benzene, or in locations that present an explosion hazard. **ATTENTION:** Refer to the ORA™ System Operator's Manual for a complete description of proper use and maintenance of the ORA™ System, as well as a complete list of contraindications, warnings and precautions.

1. Bala C, Meades K. Electron Microscopy of femtosecond laser capsulotomy edge: An inter-platform comparison. *J Cat Refract Surg* 2014; 40:1382-1389.
2. Roberts TV, Lawless M, Bali SJ, et al. Surgical outcomes and safety of femtosecond laser cataract surgery: a prospective study of 1500 consecutive cases, *Ophthalmology*, 2013;120:227-233.
3. Mayer W, Klaproth O, Hengerer F, Kohnen T. Impact of crystalline lens opacification on effective phacoemulsification time in femtosecond laser-assisted cataract surgery. *Am J Ophthalmol*. 2014;157(2):426-432.
4. Prickett A, Hou J, Jain S, Cortina S, de la Cruz J. Initial resident experience performing cataract surgery with and without femtosecond laser. Presented at UIC Alumni Day; June 21, 2013.
5. Alcon data on file. (LuxOR®)
6. Zacharias J. Thermal characterization of phacoemulsification probes operated in axial and torsional modes. *Journal of Cataract & Refractive Surgery*. 2015;41(1):208-216.
7. Han W, Miller KM. Heat production: Longitudinal versus torsional phacoemulsification. *J Cataract Refract Surg*. 2009;35(10):1799-1805.
8. Christakis PG, Braga-Mele RM. Intraoperative performance and postoperative outcome comparison of longitudinal, torsional, and transversal phacoemulsification machines. *J Cataract Refract Surg*. 2012;38(2):234-241.
9. Sharif-Kashani, et al. Comparison of occlusion break responses and vacuum rise times of phacoemulsification systems. *BMC Ophthalmology*. 2014;14:96.
10. Nicoli M, Miller K, Dimalanta R, Loke D. Jules Stein Eye Institute, UCLA. IOP Stability Measurement and Comparison Between Gravity-Fed and Actively Controlled Phacoemulsification Systems. 2014.
11. Solomon K, Lorente R, Cionni R, Fanney D. Prospective, randomized clinical study using a new phaco system with intraocular system target pressure control. Presented at ASCRS-ASOA Symposium and Congress; April 25-29, 2014; Boston, USA.
12. Vasavada A, Ong M, Cordova D, Hartzer M. Protective effect ophthalmic viscosurgical devices (OVDs) against hydrogen peroxide-induced oxidative damage to corneal endothelial cells: an in-vitro model. Presented at American Society of Cataract and Refractive Surgeons; April 3-8, 2009; San Francisco, CA.
13. Petroll WM, Jafari M, Lane SS, Jester JV, Cavanagh HD. Quantitative assessment of ophthalmic viscosurgical device retention using in vivo confocal microscopy. *J Cataract Refract Surg*. 2005;31(12):2363-2368.
14. Lindstrom RL, Ong M. Protective effect of OVDs against hydrogen peroxide-induced oxidative damage to corneal endothelial cells: in vitro model. Presented at ASCRS; March 26, 2011; San Diego, CA.
15. DisCoVisc® Package Insert.
16. Alcon data on file. VerifEye™ Technology incorporates the VerifEye® Technology validation software, but VerifEye™ Technology was not available at the time of the study.
17. Ianchulev T, Hoffer K, Yoo S, et al. Intraoperative refractive biometry for predicting intraocular lens power calculation after prior myopic refractive surgery. *Ophthalmology*. 2014;121(1):57-60.2.
18. Wentzech MG, et al. Effect of haptic design on change in axial lens position after cataract surgery. *J Cataract Refract Surg*. 2004;30(1):45-51.
19. Koshy JJ, Nishi Y, Hirnschall N, et al. Rotational stability of a single-piece toric acrylic intraocular lens. *J Cataract Refract Surg*. 2010;36(10):1665-1670.
20. Nejima R, et al. Prospective intrapatient comparison of 6.0-millimeter optic single-piece and 3-piece hydrophobic acrylic foldable intraocular lenses. *Ophthalmology*. 2006;113(4):585-590.
21. Linnola RJ, Sund M, Wönen R,Pihlajaniemi T. Adhesion of soluble fibronectin, laminin, and collagen type IV to intraocular lens materials. *J Cataract Refract Surg*. 1999;25:1486-1491.
22. AcrySof® IQ ReSTOR® IOL Clinical trial data on file (models SN6AD1 and SN6AD3). Fort Worth, TX; Alcon Laboratories, Inc.
23. AcrySof® IQ IOL Directions for Use.

Alcon
a Novartis company

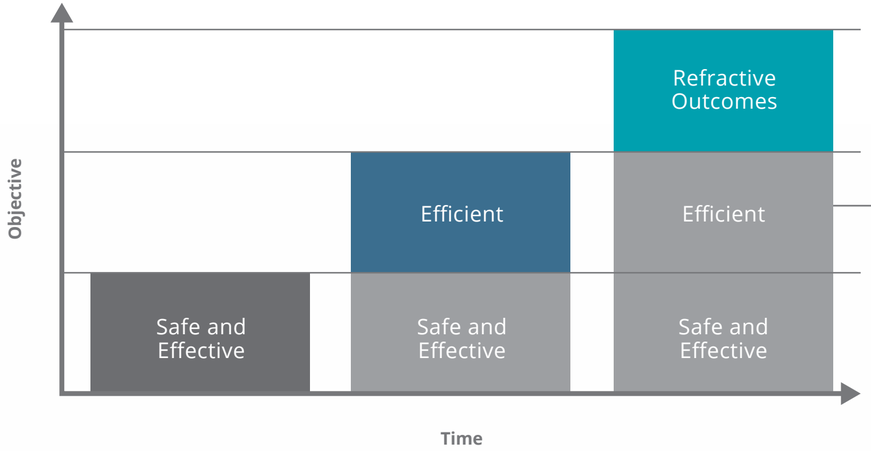
The Cataract Refractive Suite BY ALCON | Advancing CATARACT SURGERY

LEADING TECHNOLOGY FOR THE OUTCOMES ERA

Alcon
a Novartis company

The Cataract Refractive Suite BY ALCON | Advancing CATARACT SURGERY

Evolution of Cataract Technology

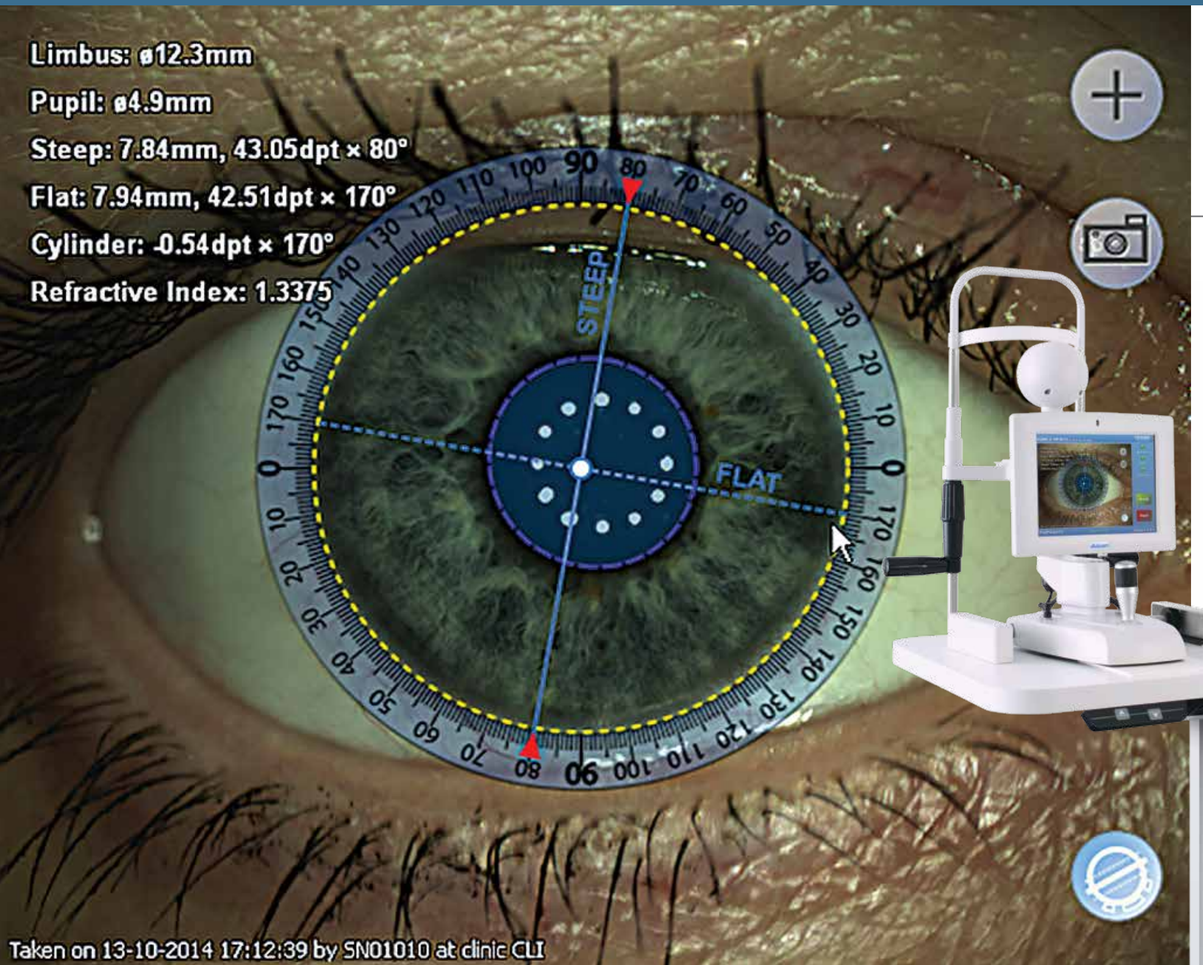


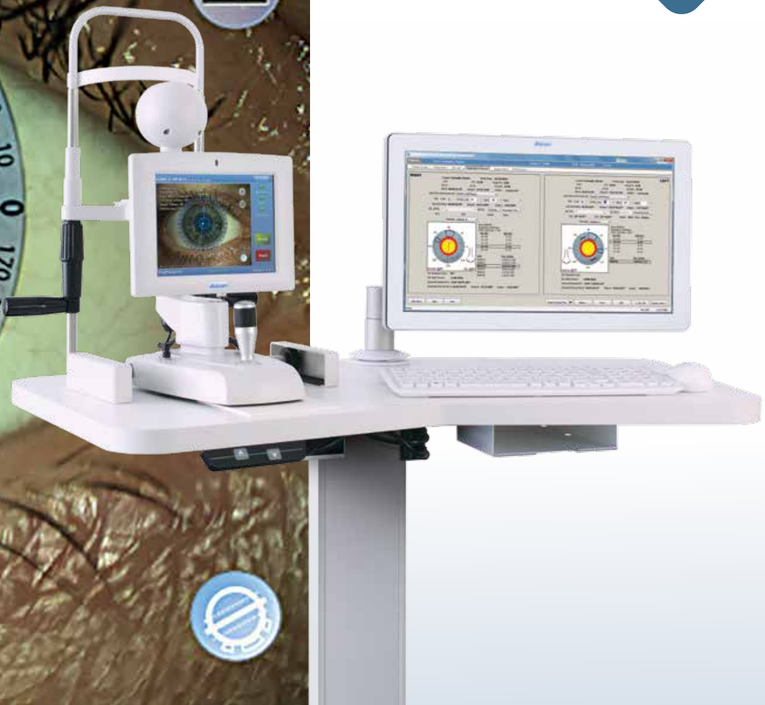
"For more than two thousand years, cataract surgery has been viewed as a rehabilitative procedure. Only within the last 15 years has it come to be seen as also a refractive procedure."
—Dr. Warren Hill

The Vision and Technology for the New Era of Cataract Surgery

Refractive outcomes are the new pursuit in modern cataract surgery. This is what patients expect and how physicians will be measured. Delivering targeted refractive outcomes is possible with leading-edge technologies, working together throughout each step of the procedure. The Cataract Refractive Suite by Alcon is designed to provide a seamless experience, from pre- to post-op, that empowers surgeon decision-making at each step to deliver advanced technology outcomes.

Individually, each technology is engineered for best-in-class performance. Together, they work as an integrated ensemble designed to provide the best possible refractive outcome. Efficient workflow. Greater confidence. Optimized outcomes. The era of refractive outcomes is here—and the Cataract Refractive Suite by Alcon is leading the way.





Verion® Image Guided System

Enhances the functionality of key platform technologies throughout the cataract refractive procedure.

- Create a complete surgical plan that moves with you from the clinic to the LenSx® Laser
- Generates digital guidance overlays for precise placement of incisions, IOLs and capsulotomies
- Accounts for cyclorotation with patient eye registration and tracking

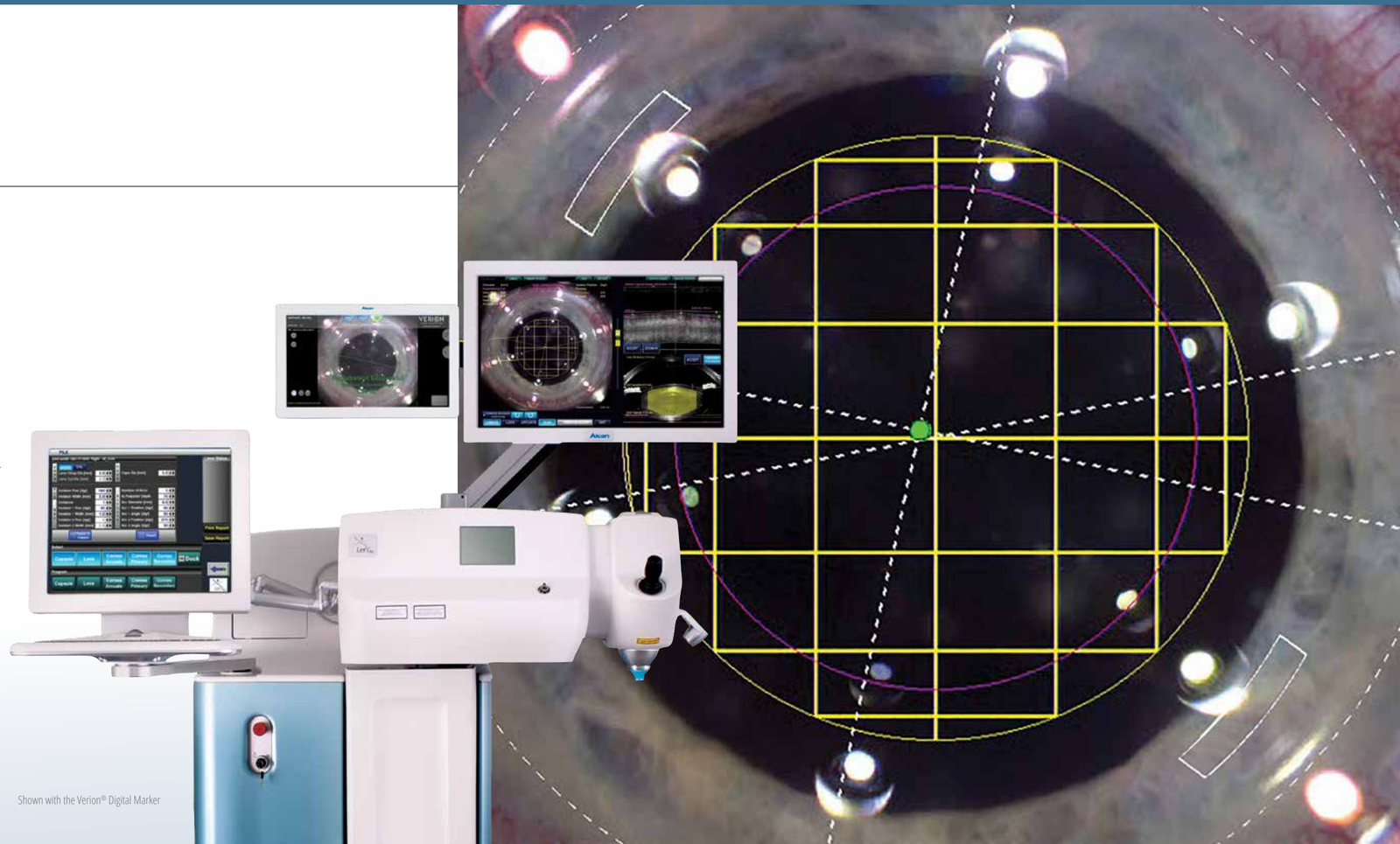
Seamless Integration
The Verion® Image Guided System allows you to create a complete surgical plan that moves with you from the clinic to the OR.

LenSx® Laser

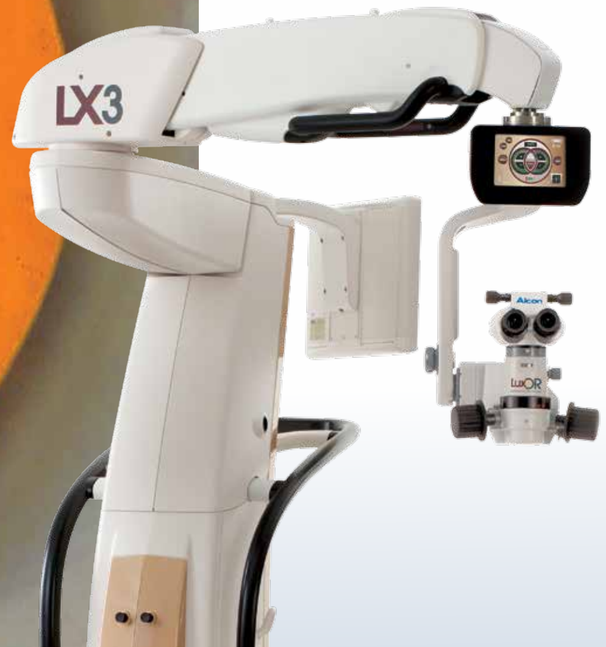
Proven precision and consistency[1,2] from the global leader in laser cataract refractive surgery.

- Advanced automation and rapid treatment times to help maximize surgical outcomes
- Automatically accounts and adjusts for cyclorotation when paired with the Verion® Digital Marker
- Lens fragmentation patterns for more efficient phacoemulsification[3,4]
- Simple SoftFit™ Patient interface that fixates the eye, meaning no need for strapping down the patients head.

The Verion® Image Guided System and the LenSx® Laser work together to provide enhanced levels of precision.



Shown with the Verion® Digital Marker



LuxOR® LX3 Ophthalmic Microscope

Superior visualization that turns the ordinary into extraordinary:

- Expanded illumination field with a 6x larger, highly stable red reflex zone[5]
- Greater red reflex stability during patient eye movement and other intraoperative maneuvers[5]
- Increased depth of focus for crisp visualization of multiple planes of the eye at once

When combined with Alcon's Cataract Refractive Diagnostic technologies, you can access digital markers and image-guided overlays right in your ocular.

Experience the Centurion® Effect

Superior phaco performance[6-8] through superior engineering.

The Centurion® Vision System's innovative phacoemulsification performance helps to enhance the OR experience for you and your patients.

- **Active Fluidics™:** Dynamic IOP management designed to enhance chamber stability[9,10]
- **Centurion® Energy Delivery:** Harmonizing OZil® Torsional Phaco delivery with increased fluidic control to maximize performance[11]
- **INTREPID® Integration:** Micro-incisional portfolio designed to enhance surgeon control

Monitor your Centurion® Vision System metrics in real time, directly in your ocular, when paired with the Verion® Image Guided System.

The Family of OVDs by Alcon

The Alcon portfolio contains OVD brands that can deliver exceptional performance, with a powerful combination of:

- Chondroitin sulfate for superior endothelial protection and retention[12-14]
- Mechanical protection, space maintenance and clarity[15]

The ORA™ System with VerifEye+™ Technology

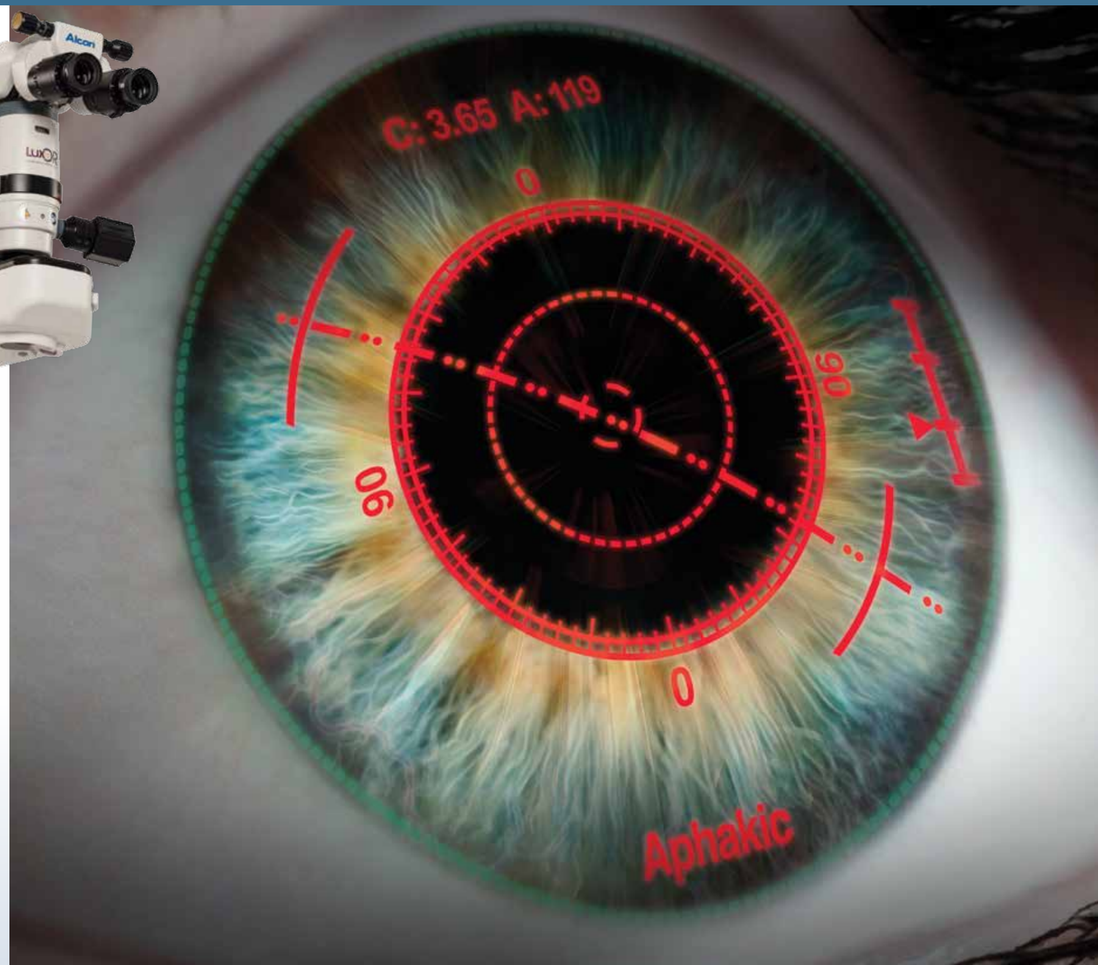
Empower surgical decision-making, in real-time, for your advanced technology outcomes.[16]

- Real-time, streaming information and guidance in your ocular
- Assess refractive impact of critical IOL decisions (power and cylinder) before implantation during the aphakic state
- Convenient, real-time guidance to help accurately place toric IOLs or LRIs[16]

Cataract procedures utilizing the ORA™ System have been clinically proven to:

- Significantly improve astigmatic outcomes by accounting for anterior and posterior corneal astigmatism[16]
- Significantly increase accuracy of lens power selection and decrease prediction error in prior myopic LASIK patients[17]

With surgical decision-making, having the complete picture matters. Real-time information while you operate means improved astigmatic outcomes.[16,17]

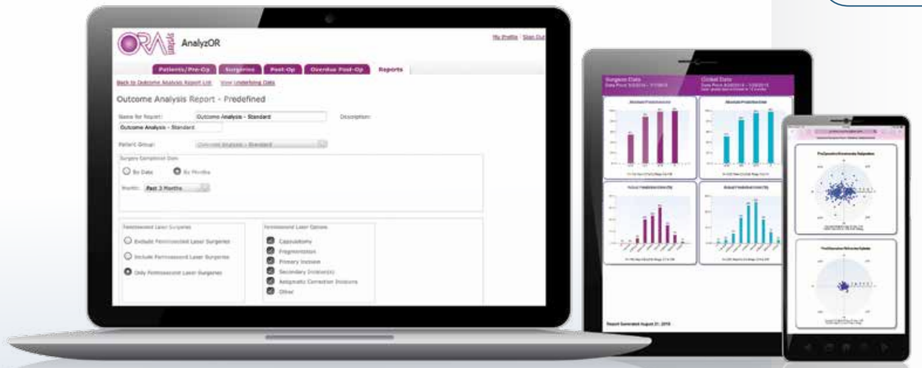


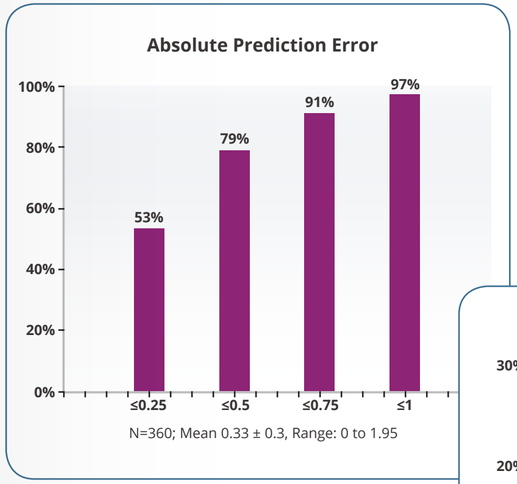
The ORA™ System with AnalyzOR™

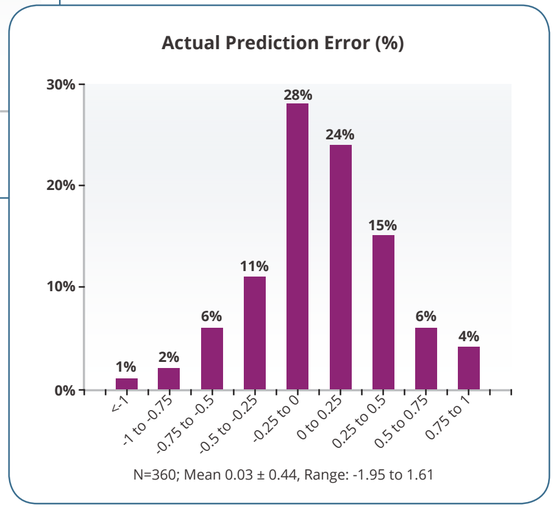
Dynamic variable optimization and robust reporting.

- Regular updates optimize surgical variables based on trending real-world outcomes
- Your personal coefficients are customized further based on your individual post-op data
- Collects and analyzes post-op data to help you enhance accuracy over time

More accurate calculations can help improve cataract refractive outcomes.[16,17]







The AcrySof® Family of IOLs

A full portfolio of advanced technology lenses designed to maximize cataract refractive outcomes.

- Treat astigmatism at the time of surgery with AcrySof® IQ Toric IOL
- Designed to reduce spectacle dependence over a range of distances with the AcrySof® IQ ReSTOR® Family of IOLs

Plus confidence with the AcrySof® Advantage.

High-performance BioMechanics

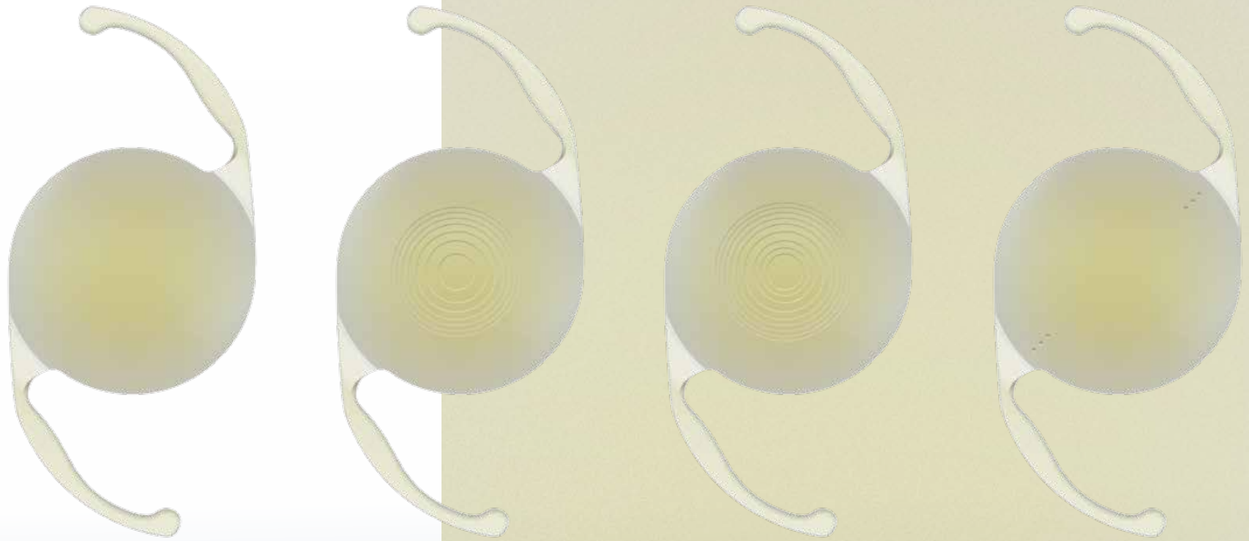
- STABLEFORCE® haptics allow for capsular bag stability and centration[18,19]
- Single-piece design delivers excellent refractive predictability[20]

Unique BioMaterial

- Bioadhesion to the capsular bag[21]

Advanced BioOptics

- UV and blue light filtration[22]
- Aspheric design for image quality[23]



AcrySof® IQ Monofocal IOL

AcrySof® IQ ReSTOR® +2.5 D IOL

AcrySof® IQ ReSTOR® +3.0 D IOL

AcrySof® IQ Toric IOL

A Platform for Performance Today and Tomorrow

Are you building your practice with the future in mind? When you invest in a piece of technology, you want to ensure that it will grow along with your practice. Every piece of the Cataract Refractive Suite by Alcon is engineered to work together for a truly integrated solution.

Whether you purchase one piece or the entire suite, you'll get the building blocks of a complete platform designed to enhance efficiency, improve workflow and provide the best possible refractive outcome. And when you add to your suite later, you'll have the confidence that everything will work together.