

EVO
Visian ICL™

PHAKIC INTRAOCULAR LENS

Evolution in Visual Freedom™

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EVO Visian™ Implantable Collamer™ Lens (EVO ICL™)

DIRECTIONS FOR USE

PRODUCT INFORMATION

Please review this product information completely before performing your initial clinical procedure. All physicians must complete the STAAR Surgical EVO ICL Physician Certification Program; special attention is placed on sizing methodologies for determination of EVO ICL overall diameter. Improper EVO ICL size may lead to adverse events ranging from mild to severe.

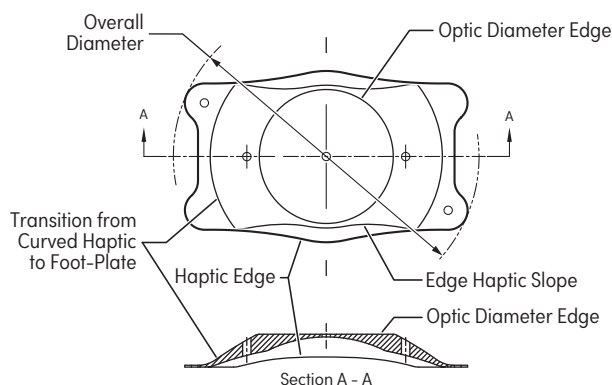
DEVICE DESCRIPTION

EVO ICL features a single piece lens design with a concave/convex optic zone of 4.9 to 5.8 mm diameter (according to model and diopter) and a 0.36 mm diameter central hole in the optic known as the KS-AquaPORT™. The lens is manufactured in four overall diameters: 12.1, 12.6, 13.2, 13.7 mm to accommodate different eye sizes. The lenses are capable of being folded and implanted through an incision of 3.5 mm or less. The lenses are manufactured from a proprietary ultraviolet (UV) radiation absorbing polymer containing hydroxyethylmethacrylate (HEMA) and porcine collagen. The 10% UV cut-offs for STAAR's phakic IOL lens family are:

- 377 nm for the thinnest central thickness lens, -5.5 D, and
- 388 nm for the thickest central thickness lens, +10.0 D.

Table 1: VICMO Models

Brand Name	Model Name	Dioptic Power (D)	Overall Diameter (mm)	Optic Diameter (mm)	Haptic Design
EVO Visian ICL	VICMO 12.1	-0.5 to -18.0	12.1	4.9 to 5.8	Flat, plate
EVO Visian ICL	VICMO 12.6	-0.5 to -18.0	12.6	4.9 to 5.8	Flat, plate
EVO Visian ICL	VICMO 13.2	-0.5 to -18.0	13.2	4.9 to 5.8	Flat, plate
EVO Visian ICL	VICMO 13.7	-0.5 to -18.0	13.7	4.9 to 5.8	Flat, plate



VICMO Diagram

INDICATIONS

EVO ICL is indicated for use in phakic eye treatment in patients 21–60 years of age for:

- The correction/reduction of myopia in patients ranging from -0.5 D to -20.0 D at the spectacle plane.
- With an anterior chamber depth (ACD) equal to or greater than 2.8 mm, as measured from the corneal endothelium to the anterior lens capsule.

MODE OF ACTION

EVO ICL is intended to be placed entirely within the posterior chamber directly behind the iris and in front of the anterior surface of the natural lens of a phakic patient. When correctly positioned, the lens functions as a refractive element to optically correct/reduce myopia.

CONTRAINDICATIONS

EVO ICL is contraindicated in the presence of any of the following circumstances and/or conditions:

1. Patients with low/abnormal corneal endothelial cell density, Fuchs' dystrophy or other corneal pathology.
2. Ocular hypertension in either eye.
3. Any cataract in the operative eye or non-traumatic cataract in the fellow eye.
4. Persons under the age of 21 years.
5. Primary Open Angle or Narrow Angle Glaucoma.
6. Narrow anterior chamber angles (i.e. less than Grade III as determined by gonioscopic exam).
7. Pregnant or nursing.
8. Previous or pre-existing ocular disease that would preclude post-operative visual acuity of 0.477 logMAR (20/60 Snellen) or better.
9. Patients who are amblyopic or blind in the fellow eye.
10. Implantation of a lens in an eye with an anterior chamber depth (ACD), as measured from the corneal endothelium to the anterior lens capsule, less than 2.8 mm.

COMPLICATIONS AND ADVERSE REACTIONS

Adverse reactions and complications due to, or following surgery and implantation of any EVO ICL may include, but are not limited to: Hyphema, Non-reactive Pupil, Pupillary Block, Additional YAG Iridotomy, Secondary Glaucoma, Cataract, Intraocular Infection, Uveitis/Iritis, Retinal Detachment, Vitritis, Corneal Edema, Macular Edema, Corneal Decompensation, Over/Under Correction, Significant Glare and/or Halos (under night driving conditions), Hypopyon, Increased Astigmatism, Loss of BSCVA, Decentration/Subluxation, IOP Elevation from Baseline, Corneal Endothelial Cell Loss, Iris Pigment Dispersion, Secondary Surgical Intervention to Remove/Replace/Reposition the Lens, Peripheral Anterior Synechia (PAS), Iris Synechia to Implant, Conjunctival Irritation, Vitreous Loss.

PRECAUTIONS

1. Do not attempt to sterilize.
2. Do not autoclave.
3. The lens should not be exposed to any solutions other than the normally used intraocular irrigating solutions (e.g. isotonic saline, Balanced Salt Solution (BSS), viscoelastic, etc.).
4. The lens should be handled carefully. No attempts should be made to reshape or cut any portion of the lens or to apply undue pressure to the lens optical portion with a sharp object.
5. Do not allow the lens to dry in air. The lens should be stored in sterile BSS solution during surgery.
6. The long-term effect of the lens has not been determined. Therefore, physicians should continue to monitor implant patients postoperatively on a regular basis.
7. Safety and effectiveness of the lens has not been established in patients with: unstable refractive error in either eye, keratoconus, history of clinical signs of iritis/uveitis, synechia, pigment dispersion syndrome, pseudoexfoliation, insulin-dependent diabetes or diabetic retinopathy, history of previous ocular surgery including refractive corneal surgery.
8. Implantation of a lens may result in a decrease in corneal endothelial cell density.

CALCULATION OF LENS POWER AND SIZING

The lens power and size calculation should be performed by the surgeon using the STAAR OCOS Calculation Software. Using the software potentially prevents calculation errors that may result in secondary surgery due to refractive surprise, excessive vaulting, lens rotation, IOP elevation from baseline, etc. During the U.S. FDA trial for the ICM/TICM, the white to white and ACD (from the corneal endothelium to the anterior lens capsule) were used to determine the ICL overall diameter. There are some reports suggesting that white to white corneal measurements do not correlate with sulcus to sulcus measurements. Recent publications indicate that new imaging technologies may provide optimal visualization and measurement of the intraocular dimensions involved in phakic intraocular lens implantation.

LENS PREPARATION

Verify that the level of the liquid fills at least 2/3 of the vial. The thermoformed tray and vial should be opened in a sterile field. Record serial number on operative report to retain traceability of the lens. Remove the aluminum cap and stopper from the vial. Remove the lens from the vial. The lens should not be exposed to a dry environment (air) for more than one minute.
CAUTION: Do not use if package has been opened or damaged.
CAUTION: Do not allow the lens to dry after removal from the glass vial.

ADMINISTRATION AND INSTRUCTION FOR USE

Implantation of EVO ICL should only be attempted by a surgeon who is highly skilled in the required surgical technique. The following procedure is recommended for implantation of EVO ICL.
The patient should be prepared for surgery according to the surgeon’s standard operating procedure. A clear scleral or corneal tunnel wound incision of 3.5 mm or less should be used, followed by filling of the anterior and posterior chamber with an appropriate viscoelastic. The lens is then folded using a MICROSTAAR™ injector MSI-PF or MSI-TF with SFC-45 cartridge and injected into the anterior chamber of the eye. Please refer to the product insert or loading guide provided with the injector for instructions regarding proper loading and injection of the lens using the MICROSTAAR injection system. Verify correct orientation of the lens in the eye and that the lens is not inverted. If the pupil remains sufficiently dilated, the lens should be well centered and positioned under the iris in front of the natural lens of a phakic patient, so that the footplates are placed in the sulcus. Complete removal of the viscoelastic material from the eye must be performed after completion of the surgical procedure and before the eye is closed (without sutures). From this point the operation can proceed according to the surgeon’s standard procedure. Dispose of any single use accessories that may have become contaminated with bodily fluids during the procedure as biohazardous waste according to standard surgical biohazard waste disposal procedure. Postoperative medical care of the patient should also follow the surgeon’s standard procedure.

WARNINGS

- 1. Check the label of the lens package for proper lens model and power.
- 2. Open the package to verify the dioptric power of the lens.
- 3. Handle the lens by the haptic portion. Do not grasp the optic with forceps and never touch the center of the optic once the lens is placed inside the eye.
- 4. Complete removal of viscoelastic from the eye after completion of the surgical procedure is essential. STAAR Surgical recommends a low molecular weight 2% hydroxypropyl methylcellulose (HPMC) or dispersive, low viscosity ophthalmic viscosurgical device.
- 5. STAAR Surgical recommends using the MICROSTAAR MSI-PF or MSI-TF with SFC-45 cartridge injection systems to insert the lens in the folded state.

NOTE: The primary viscoelastic used during the US FDA clinical trial was a low molecular weight 2% hydroxypropyl methylcellulose preparation.

SUMMARY OF SAFETY AND CLINICAL PERFORMANCE

A Summary of Safety and Clinical Performance (SSCP) for the ICL family of lenses can be found in the European database on medical devices (Eudamed) at <https://ec.europa.eu/tools/eudamed>. The Basic UDI-DI (BUDI-DI) used to search for the ICL family of lenses on the website is 764013516ICLV.

CLINICAL TRIAL ICM

Summary Findings of the Clinical Studies:

The Model ICM Implantable Collamer Lenses were found to be safe and effective as refractive elements to optically reduce moderate to high myopia.

Table 2: Patient Demographics
696 Eyes Treated of 404 Patients

Sex		
Male	158	(39.1%)
Female	246	(60.9%)
Ethnic Origin		
Caucasian	348	(86.1%)
Black	6	(1.5%)
Hispanic	26	(6.4%)
Other	24	(5.9%)

Mean Age 37.1 years

Table 3: Adverse Events
A summary of adverse events reported in the 696 eyes enrolled during the clinical trial (at any postoperative exam) is presented below:

Adverse Event	N	%
ICL Removal Due to Elevated IOP	2	0.3
ICL Replacement (Due to Incorrect Sizing)	6	0.9
ICL Repositioning	4	0.6
ICL Removal Due to Cataract	5	0.7
Other Secondary Surgical Interventions	6	0.8

Table 4: Best Spectacle Corrected Visual Acuity with Time for Patients with PREOP BSCVA 20/20 or better
The Implantable Collamer Lens for Myopia

	Preop n%	1 Week n%	1 Month n%	3 Months n%	6 Months n%	12 Months n%	24 Months n%
≤20/20	439/439 (100.0%)	376/417 (90.2%)	403/419 (96.2%)	391/404 (96.8%)	368/386 (95.3%)	270/280 (96.4%)	76/82 (92.7%)
≤20/25	439/439 (100.0%)	408/417 (97.8%)	416/419 (99.3%)	403/404 (99.8%)	384/386 (99.5%)	279/280 (99.6%)	82/82 (100.0%)
≤20/32	439/439 (100.0%)	414/417 (99.3%)	419/419 (100.0%)	404/404 (100.0%)	386/386 (100.0%)	279/280 (99.6%)	82/82 (100.0%)
≤20/40	439/439 (100.0%)	416/417 (99.8%)	419/419 (100.0%)	404/404 (100.0%)	386/386 (100.0%)	279/280 (99.6%)	82/82 (100.0%)
≤20/80	439/439 (100.0%)	417/417 (100.0%)	419/419 (100.0%)	404/404 (100.0%)	386/386 (100.0%)	280/280 (100.0%)	82/82 (100.0%)
≤20/200	439/439 (100.0%)	417/417 (100.0%)	419/419 (100.0%)	404/404 (100.0%)	386/386 (100.0%)	280/280 (100.0%)	82/82 (100.0%)
>20/200	0/439 (0.0%)	0/417 (0.0%)	0/419 (0.0%)	0/404 (0.0%)	0/386 (0.0%)	0/280 (0.0%)	0/82 (0.0%)
Not Reported	0	11	4	3	5	0	1
Total	439	422	421	405	391	280	83

**Table 5: Uncorrected Visual Acuity over Time for Patients with PREOP BSCVA 20/20 or better
The Implantable Collamer Lens for Myopia**

	Preop n%	1 Week n%	1 Month n%	3 Months n%	6 Months n%	12 Months n%	24 Months n%
≤20/20	0/439 (0.0%)	162/421 (38.5%)	197/421 (46.8%)	210/404 (52.0%)	200/391 (51.2%)	158/278 (56.8%)	33/83 (39.8%)
≤20/25	0/439 (0.0%)	248/421 (58.9%)	278/421 (66.0%)	281/404 (69.6%)	274/391 (70.1%)	206/278 (74.1%)	44/83 (53.0%)
≤20/32	0/439 (0.0%)	316/421 (75.1%)	351/421 (83.4%)	338/404 (83.7%)	323/391 (82.6%)	235/278 (84.5%)	63/83 (75.9%)
≤20/40	0/439 (0.0%)	364/421 (86.5%)	377/421 (89.5%)	369/404 (91.3%)	346/391 (88.5%)	253/278 (91.0%)	71/83 (85.5%)
≤20/80	0/439 (0.0%)	408/421 (96.9%)	409/421 (97.1%)	397/404 (98.3%)	382/391 (97.7%)	271/278 (97.5%)	80/83 (96.4%)
≤20/200	0/439 (0.0%)	421/421 (100.0%)	421/421 (100.0%)	404/404 (100.0%)	391/391 (100.0%)	278/278 (100.0%)	83/83 (100.0%)
>20/200	439/439 (100.0%)	0/421 (0.0%)	0/421 (0.0%)	0/404 (0.0%)	0/391 (0.0%)	0/278 (0.0%)	0/83 (0.0%)
Not Reported	0	3	1	3	1	3	0
Total	439	422	421	405	391	280	83

**Table 6: Manifest Refraction Spherical Equivalent with Time
The Implantable Collamer Lens for Myopia**

Spherical Equivalent (D)	Preop n%	1 Week n%	1 Month n%	3 Months n%	6 Months n%	12 Months n%	24 Months n%
≥1.01	0 (0.0%)	4 (0.6%)	7 (1.1%)	6 (1.0%)	5 (0.8%)	4 (0.9%)	0 (0.0%)
+1.00 to +0.01	0 (0.0%)	97 (15.0%)	119 (18.1%)	121 (19.6%)	104 (17.5%)	68 (15.2%)	14 (10.3%)
0.00 to -1.00	0 (0.0%)	399 (61.8%)	405 (61.7%)	374 (60.6%)	356 (59.8%)	283 (63.3%)	75 (55.1%)
-1.01 to -2.00	0 (0.0%)	103 (15.9%)	81 (12.3%)	81 (13.1%)	91 (15.3%)	58 (13.0%)	36 (26.5%)
-2.01 to -6.00	0 (0.0%)	40 (6.2%)	41 (6.3%)	31 (5.0%)	36 (6.1%)	32 (7.2%)	11 (8.1%)
-6.01 to -10.00	284 (40.8%)	3 (0.5%)	3 (0.5%)	4 (0.6%)	3 (0.5%)	2 (0.4%)	0 (0.0%)
-10.01 to -15.00	318 (45.7%)	0 (0.0%)	0 (0.0%)	0 (0.0%)	0 (0.0%)	0 (0.0%)	0 (0.0%)
-15.01 to -20.00	88 (12.6%)	0 (0.0%)	0 (0.0%)	0 (0.0%)	0 (0.0%)	0 (0.0%)	0 (0.0%)
<-20.00	6 (0.9%)	0 (0.0%)	0 (0.0%)	0 (0.0%)	0 (0.0%)	0 (0.0%)	0 (0.0%)
Total	696 (100.0%)	646 (100.0%)	656 (100.0%)	617 (100.0%)	595 (100.0%)	447 (100.0%)	136 (100.0%)
Mean	-11.408	-0.658	-0.566	-0.526	-0.580	-0.623	-0.857

ADVERSE EVENT REPORTING

Adverse Reactions and/or potentially sight-threatening complications that may reasonably be regarded as lens related must be reported immediately to STAAR Surgical and the competent authority of the EU Member State where the patient is established. This information is being requested from surgeons in order to document potential long-term effects of EVO ICL implantation.

HOW SUPPLIED

EVO ICL is supplied sterile and non-pyrogenic in a sealed vial containing BSS. The vial is sealed within a thermoformed tray placed in a box with labels and product information. Sterility is assured until the expiration date indicated on package label, if the tray and vial seal are not punctured or damaged. EVO ICL is steam sterilized. A Patient Implant Card, implant card instructions and labels are supplied in the unit package. This card, which includes a link to important safety information regarding the implanted lens should be completed by the healthcare provider and given to the patient to keep as a permanent record of the implant, and a resource to show to any eye care practitioner seen in the future.

EXPIRATION DATE

The expiration date on the device package is the sterility expiration date. This device must not be used past the indicated sterility expiration date.

RETURN POLICY FOR STAAR EVO ICL

Contact STAAR Surgical. EVO ICL must be returned dry. Do not attempt to re-hydrate the lens.

WARRANTY AND LIMITATION OF LIABILITY

STAAR Surgical warrants that reasonable care was taken in making this product. STAAR Surgical shall not be responsible for any incidental or consequential loss, damage, or expense which arises directly or indirectly from the use of this product. To the extent permitted by law, STAAR Surgical's sole liability from any and all causes pursuant to EVO ICL shall be limited to the replacement of EVO ICL which is returned to and found to be defective by STAAR Surgical. This warranty is in lieu of and excludes all other warranties not expressly set forth herein, whether expressed or implied by operation of law or otherwise, including but not limited to, or any implied merchantability or fitness for use.

STORAGE

Store the lens at room/ambient temperature.

CAUTION

- Do not autoclave the lens. Do not store the lens at temperatures greater than 40°C. Do not freeze. If temperature requirements are not met, return the lens to STAAR Surgical.
- STAAR Surgical EVO ICL and disposable accessories are packaged and sterilized for single use only. Cleaning, reuse and/or resterilization are not applicable to these devices. If one of these devices were reused after cleaning and/or resterilization, it is highly probable that it would be contaminated and the contamination could result in infection and/or inflammation.

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SYMBOL GLOSSARY

	Medical device		Authorized representative in the European Community
	Do not re-use		CE conformity marking per European Council Directive 93/42/EEC or European Council Regulation (EU) 2017/745
	Do not re-sterilize		Manufacturer
	Do not use if the product sterile barrier system or its packaging is compromised		Date of manufacture
	Body diameter (Optic diameter)		Country of manufacture – United States
	Overall diameter		Country of manufacture – Switzerland
	Single sterile barrier system with protective packaging outside		Unique Device Identifier
	Use-by date		Catalogue number
	Diopter		Right eye
	Date		Left eye
	Caution		Serial number
	Contains biological material of animal origin		Consult electronic instructions for use
	U.S. (Federal) law restricts this device to sale by or on the order of a physician		Health care center or Doctor
	Store at room/ambient temperature. Do not freeze. Do not expose to temperature greater than 40°C		
	Sterilized using steam		