


IOL Product Guide






AcrySof[®]
ACRYLIC FOLDABLE IOL

Alcon[®]




ACRYSOF® ReSTOR® Aspheric IOL

	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS*	A-CONSTANT
	SN6AD3	6.0	Apodized Diffractive Aspheric	13.0	0°		+10.0 to +30.0 +31.0 to +34.0 (1.0 diopter increments)	118.9

ACRYSOF® ReSTOR® IOL

	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS*	A-CONSTANT
	SN60D3	6.0	Apodized Diffractive	13.0	0°		+6.0 to +30.0	118.1
	SA60D3	6.0	Apodized Diffractive	13.0	0°		+10.0 to +30.0	118.1
	MN60D3	6.0	Apodized Diffractive	13.0	10°		+6.0 to +30.0	118.3

ACRYSOF® Toric IOL


	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS*	A-CONSTANT
	SN60T3	6.0	Biconvex Toric	13.0	0°		+6.0 to +30.0 Spherical Equivalent 1.50 Cylinder	118.4
	SN60T4	6.0	Biconvex Toric	13.0	0°		+6.0 to +30.0 Spherical Equivalent 2.25 Cylinder	118.4
	SN60T5	6.0	Biconvex Toric	13.0	0°		+6.0 to +30.0 Spherical Equivalent 3.00 Cylinder	118.4

Visit www.acrysoftoriccalculator.com
Unless otherwise noted, IOLs are available in 0.5 increments.
Not all lenses available in all countries.


FOR U.S. ORDERS CALL 1-800-TO-ALCON (1-800-862-5266)

OUTSIDE U.S. CONTACT YOUR LOCAL REPRESENTATIVE


ACRYSOF® IQ IOL

	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS [#]	A-CONSTANT
	SN60WF	6.0	Aspheric	13.0	0°		+6.0 to +30.0	118.7


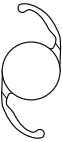
ACRYSOF® IQ IOL with ACRYCERT® Delivery System

	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS [#]	A-CONSTANT
	SN60WS	6.0	Aspheric	13.0	0°		+6.0 to +30.0	118.7

ACRYSOF® Single-Piece - Natural IOL

	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS [#]	A-CONSTANT
	SN60AT	6.0	Anterior Asymmetric Biconvex	13.0	0°		+6.0 to +30.0 +31.0 to +40.0 (1.0 diopter increments)	118.4

ACRYSOF® Single-Piece IOL

	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS [#]	A-CONSTANT
	SA30AT	5.5	Anterior Asymmetric Biconvex	13.0	0°		+10.0 to +30.0	118.4
	SA60AT	6.0	Anterior Asymmetric Biconvex	13.0	0°		+6.0 to +30.0 +31.0 to +40.0 (1.0 diopter increments)	118.4


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ACRYSOF® Multipiece - Natural

	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS [#]	A-CONSTANT
	MN60AC	6.0	Anterior Asymmetric Biconvex	13.0	10°		+6.0 to +30.0	118.4

ACRYSOF® Multipiece

	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS [#]	A-CONSTANT
	MA30AC	5.5	Anterior Asymmetric Biconvex	12.5	5°		+10.0 to +30.0	118.4
	MA60AC	6.0	Anterior Asymmetric Biconvex	13.0	10°		+6.0 to +30.0	118.4
	MA30BA[‡]	5.5	Biconvex	12.5	5°		+10.0 to +30.0	118.9

ACRYSOF® Multipiece - Posterior Convex

	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS [#]	A-CONSTANT
	MA60BM[‡]	6.0	Biconvex	13.0	10°		+6.0 to +30.0	118.9
	MA50BM	6.5	Biconvex	13.0	10°		+6.0 to +30.0	118.9

Visit www.acrysoftoriccalculator.com

[‡] Not available in U.S.



[#] Unless otherwise noted, IOLs are available in 0.5 increments.

Not all lenses available in all countries.

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ACRYSOF® Multipiece - EXPAND® Series









	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS [#]	A-CONSTANT
	MA60MA	6.0	Meniscus	13.0	5°		minus 5.0 to plus 5.0 (1.0 diopter increments)	118.9
	MN60MA	6.0	Meniscus	13.0	5°		minus 5.0 to plus 5.0 (1.0 diopter increments)	118.9

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[#] Unless otherwise noted, IOLs are available in 0.5 increments.
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Single-Piece PMMA


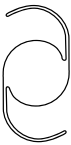

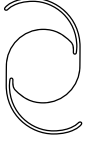

	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS [#]	A-CONSTANT
	MC40BD	5.0	Biconvex	12.5	5° SLANT™		+10.0 to +30.0	118.7
	MZ40BD	5.0	Biconvex	11.5	5° SLANT		+10.0 to +30.0	118.7
	MX40BD[‡]	5.0	Biconvex	12.0	5° SLANT		+10.0 to +30.0	118.7
	LX10BD	5.25	Biconvex	12.0	5° SLANT	SLIMPLANT™ DESIGN	+10.0 to +30.0	118.7
	MZ30BD	5.5	Biconvex	12.0	5° SLANT		+10.0 to +30.0	118.7
	LX90BD	5.75	Biconvex	12.0	5° SLANT	SLIMPLANT™ DESIGN	+10.0 to +30.0	118.7
	MZ60BD	6.0	Biconvex	12.5	5° SLANT		+4.0 to +30.0 +31.0 to +34.0 (1.0 diopter increments)	118.7
	LC80BD	6.25	Biconvex	13.0	5° SLANT	SLIMPLANT™ DESIGN	+10.0 to +30.0	118.7

‡ Not available in U.S.
Unless otherwise noted, IOLs are available in 0.5 increments.
† SLIMPLANT Reg. U.S. Patent & TM Office
Not all lenses available in all countries.



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Single-Piece PMMA




	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS [#]	A-CONSTANT
	MC60BD	6.0	Biconvex	13.5	5° SLANT™		+10.0 to +30.0	118.7
	MC50BD	6.5	Biconvex	13.5	5° SLANT		+10.0 to +30.0	118.7
	CZ70BD	7.0	Biconvex	12.5	5° SLANT	Eyelet	+10.0 to +30.0	118.8
	CR5BUO	7.0	Biconvex	13.5	10°		+10.0 to +30.0	119.0
	CR70BU	7.0	Biconvex	13.5	10°	Eyelet	+4.0 to +30.0	119.0

Single-Piece PMMA EXPAND® Series

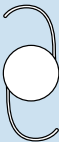




	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS [#]	A-CONSTANT
	MZ60MD	6.0	Meniscus	12.5	5° SLANT		minus 3.0 to plus 3.0 (1.0 diopter increments)	118.7
	MZ60PD	6.0	Plano-concave	12.5	5° SLANT		minus 10.0 to minus 4.0 (1.0 diopter increments)	118.7

[#] Unless otherwise noted, IOs are available in 0.5 increments.
Not all lenses available in all countries.

ReFORM® Capsular Tension Rings

	MODEL NUMBER	EXPANDED	COMPRESSIBLE TO
	ACTR10	12.3 mm	10 mm
	ACTR11	13.0 mm	11 mm
	ACTR12	14.5 mm	12 mm

MONOFLEX™† PMMA


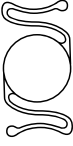




	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC ANGULATION	OTHER FEATURES	DIOPTIC RANGE/ INCREMENTS#	A-CONSTANT
	MC30BA	5.5	Biconvex	13.5	5°		+10.0 to +30.0	118.7
	MC60BM	6.0	Biconvex	13.5	10°		+4.0 to +6.0 (1.0 diopter increments) +7.0 to +30.0 +31.0 to +34.0 (1.0 diopter increments)	118.7
	MC60CM	6.0	Convexoplano	13.5	10°		+2.0 to +3.0 (1.0 diopter increments†) +5.0 to +30.0 +31.0 to +34.0 (1.0 diopter increments) +4.0 (1.0 diopter increments)	116.6
	MC50BM	6.5	Biconvex	14.0	10°		+10.0 to +30.0	118.7
	MC70CM	7.0	Convexoplano	14.0	10°		+2.0 to +3.0 (1.0 diopter increments†) +4.0 (1.0 diopter increments) +5.0 to +30.0 +31.0 to +34.0 (1.0 diopter increments)	116.4

† MONOFLEX Reg. U.S. Patent & TM Office
Unless otherwise noted, IOLs are available in 0.5 increments.
‡ Not available in U.S.
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FOR U.S. ORDERS CALL 1-800-TO-ALCON (1-800-862-5266)

OUTSIDE U.S. CONTACT YOUR LOCAL REPRESENTATIVE

Anterior Chamber

	MODEL NUMBER	OPTIC SIZE (mm)	OPTIC TYPE	LENGTH (mm)	HAPTIC VAULT	DIOPTIC RANGE/ INCREMENTS [#]	ACD ¹	A-CONSTANT
	MTA2U0	5.5	Convexoplano	12.0	0.5 mm	+10.0 to +28.0	3.39	115.3
	MTA3U0	5.5	Convexoplano	12.5	0.5 mm	+5.0 to +9.0 (1.0 diopter increments) +10.0 to +28.0 +29.0 to +30.0 (1.0 diopter increments)	3.39	115.3
	MTA4U0	5.5	Convexoplano	13.0	0.5 mm	+5.0 to +9.0 (1.0 diopter increments) +10.0 to +28.0 +29.0 to +30.0 (1.0 diopter increments)	3.39	115.3
	MTA5U0	5.5	Convexoplano	13.5	0.5 mm	+5.0 to +9.0 (1.0 diopter increments) +10.0 to +28.0 +29.0 to +30.0 (1.0 diopter increments)	3.39	115.3
	MTA6U0	5.5	Convexoplano	14.0	0.5 mm	+10.0 to +28.0	3.39	115.3
	MTA7U0	5.5	Convexoplano	14.5	0.5 mm	+10.0 to +28.0	3.39	115.3

¹ ACD is based on the Binkhorst relationship as presented in Holladay, J.T., Musgrove, K.H., Prager, T.C., Lewis, J.W., Chandler, T.Y., and Ruiz, R.S., "A three-part system for refining intraocular lens power calculations," *Journal of Cataract and Refractive Surgery*, Vol. 14, pp. 17-24, 1988.

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IOL Delivery Systems

MONARCH® II IOL Delivery System

The MONARCH® II IOL Delivery System combines a reusable titanium handpiece and a sterile single use cartridge for enhanced implantation of the ACRYSoF® IOL. Advanced design enables the surgeon to view and verify lens orientation throughout the implantation process. Simplified loading, controlled consistent delivery, and ease of implantation are the benefits that MONARCH® II can provide.

Part No.: 8065977771
• MONARCH® II IOL Injector

• Ten (10) MONARCH® II Cartridges per box

MONARCH® Cartridges	ACRYSoF® Lens Model	
8065-977757 A Cartridge	MA50BM	MN60D3
	MA60AC	MN60AC
	MA60BM	MN60MA
	MA60MA	
8065-977758 B Cartridge	SN60AT	MA30AC
	SN60D3	MA30BA
	SN60WF	SA30AL
	SN60T3	SA30AT
	SN60T4	SA60AT
	SN60T5	SA60D3
	SN6AD3	
8065-977759 C Cartridge	SN60AT	+6.0 to +27.0 D
	SN60D3	+10.0 to +27.0 D
	SN60WF	+6.0 to +30.0 D
	SN60T3	+16.0 to +25.0 D
	SN60T4	+16.0 to +25.0 D
	SN60T5	+16.0 to +25.0 D
	SA30AT	+16.0 to +25.0 D
	SA60AT	+6.0 to +27.0 D
	SA60D3	+10.0 to +27.0 D
	SN6AD3	+10.0 to +27.0 D

Part No.: 560.01
The Alcon®/Grieshaber® MONARCH® II Loading Forceps is for fully controlled handling of the ACRYSoF® Single-Piece and ACRYSoF® Natural IOLs from its packaging into the MONARCH® II Delivery System.

MONARCH® III IOL Delivery System

The MONARCH® III IOL Delivery System combines a reusable titanium handpiece and a sterile single use cartridge for enhanced implantation of the ACRYSoF® IOL. Advanced design enables the surgeon to view and verify lens orientation throughout the implantation process. Simplified loading, controlled consistent delivery, and ease of implantation are the benefits that MONARCH® III can provide.

Part No.: 8065977773
• MONARCH® III IOL Injector

• Ten (10) MONARCH® Cartridges per box

MONARCH® Cartridges	ACRYSoF® Lens Model	
8065-977758 B Cartridge	SN60AT	MA30AC
	SN60D3	MA30BA
	SN60WF	SA30AL
	SN60T3	SA30AT
	SN60T4	SA60AT
	SN60T5	SA60D3
	SN6AD3	
8065-977759 C Cartridge	SN60AT	+6.0 to +27.0 D
	SN60D3	+10.0 to +27.0 D
	SN60WF	+6.0 to +30.0 D
	SN60T3	+16.0 to +25.0 D
	SN60T4	+16.0 to +25.0 D
	SN60T5	+16.0 to +25.0 D
	SA30AT	+16.0 to +25.0 D
8065-977763 D Cartridge	SA60AT	+6.0 to +27.0 D
	SA60D3	+10.0 to +27.0 D
	SN6AD3	+10.0 to +30.0 D
	SN60WF	+6.0 to +27.0 D

Part No.: 560.01
The Alcon®/Grieshaber® MONARCH® Loading Forceps is for fully controlled handling of the ACRYSoF® Single-Piece and ACRYSoF® Natural IOLs from its packaging into the MONARCH® Delivery System.

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LENS GLIDES

Part No.: 8065183907

- Available in boxes of ten (10) packages
- Two glides per package: one 17.0 mm x 6.65 mm, one 24.6 mm x 6.65 mm
- For use with single- or double-glide insertion techniques



Part No.: 8065183921

- Available in boxes of thirty-six (36) packages
- One glide per package: 5.0 mm x 35.0 mm



ACRYPAK® FOLDER

Part No.: 8065977725

- Designed specifically for multi-axis folding of the ACRYSOF® IOL
- Ten (10) ACRYPAK® Folders per box

PREMARKET APPROVAL

All lenses and capsular tension rings listed in the guide are premarket approved by the FDA unless otherwise noted.

A-CONSTANTS & ACDs

There are two basic classes of formulae (theoretical and regression analysis) used to calculate dioptric power for an intraocular lens implant. The theoretical formulae (i.e., Binkhorst, Colenbrander, Holladay, and others) require an estimation of anterior chamber depth, while regression analysis formulae (SRK, SRKII and SRK/T) by Sanders, Retzlaff, and Kraff require an A-constant. Both of these numbers will vary depending on the model utilized by the surgeon. All of the numbers listed within the IOL catalog are presented as guidelines only and are good starting points for the implant power calculations. We recommend that you develop your own A-constant or estimated anterior chamber depth measurements based on your experience with a particular implant model, surgical technique, measuring equipment and postoperative results. A-constant for all convexoplano models assumes optic orientation with plano side placed posteriorly.

DIOPTRIC POWERS

Alcon offers a wide range of dioptric powers. Numerous models are offered in a range from +4.0 to +34.0 diopters. The EXPAND® Series extends our diopter range to -10.0 diopters.

EXPAND® Series Lenses

Powers for the EXPAND® Series were determined using a modern theoretical formula, SRK/T, and should not be considered interchangeable with powers derived for any other lens styles or formulae. For a reference chart on IOL power using the SRK/T formula for EXPAND® Series lenses, contact your Alcon representative.

PATENTS

SLANT® -	U.S. Patent Nos. 5,282,855 and 5,160,345
SLIMPLANT® -	U.S. Patent No. 5,197,981
ACRYSOF® IOL Material -	U.S. Patent No. 5,290,892, 5,603,774, 5,403,901, 5,433,746, 5,674,960 and 5,861,031
ACRYSOF® Haptic Attachment Method -	U.S. Patent No. 5,523,029
MONARCH® -	U.S. Patent Nos. 6,010,510, 5,947,976 and 6,083,231
ACRYSOF® Single-Piece Lens -	U.S. Patent No. 5,716,403
Blue light filtering chromophore	U.S. Patent No. 5,470,932, 5,543,504
ACRYPAK®	U.S. Patent No. D349342
ReSTOR®	U.S. Patent No. 5,699,142
TORIC	U.S. Patent No. 5,716,403

TRADEMARKS

† Note:

SLANT Reg. U.S. Patent & TM Office
 SLIMPLANT Reg. U.S. Patent & TM Office
 MONOFLEX Reg. U.S. Patent & TM Office

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DESCRIPTION: Alcon® UV-absorbing Single Piece PMMA and MONOFLEX™ PMMA posterior chamber lenses and Single Piece PMMA anterior chamber lenses are optical implants for the replacement of the human crystalline lens in the visual correction of aphakia in adult patients following cataract removal. These lenses have biconvex, convexoplano, convexoplano with Hoffer™* ridge modification, or meniscus optics with supporting haptics.

ACRYSOF® Posterior Chamber Intraocular Lenses are indicated for the replacement of the human lens to achieve visual correction of aphakia in adult patients following cataract removal. These lenses are intended for placement in the capsular bag.

CAUTION: Federal law restricts this device to sale by or on the order of a physician.

WARNINGS: 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. Some adverse reactions that have been associated with the implantation of intraocular lenses are: hypopyon, intraocular infection, acute corneal decompensation, and secondary surgical intervention.

PRECAUTIONS: Do not resterilize; do not store over 45°C; use only sterile irrigating solutions such as BSS® or BSS PLUS®.

ATTENTION: Reference the Product Insert for a complete listing of warnings & precautions as they may vary by lens.

ACRYSOF® RESTOR® SN6AD3/SN60D3/MN60D3/SA60D3/MA60D3

CAUTION: Federal law restricts this device to sale by or on the order of a physician.

INDICATIONS: The **ACRYSOF®** ReSTOR® Apodized Diffractive Optic Posterior Chamber Intraocular Lens (IOL) is intended for primary implantation for the visual correction of aphakia secondary to removal of a cataractous lens in adult patients with or without presbyopia, who desire near, intermediate and distance vision with increased spectacle independence. The lens is intended to be placed in the capsular bag.

WARNINGS: 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. Some adverse reactions that have been associated with the implantation of intraocular lenses are: hypopyon, intraocular infection, acute corneal decompensation, macular edema, pupillary block, retinal detachment, and secondary surgical intervention (including but not limited to lens repositioning, biometry error, visual disturbances or patient dissatisfaction). As a result of the multifocality, some visual effects (halos or radial lines around point sources of light at night) may also be expected due to the superposition of focused and unfocused multiple images. A reduction in contrast sensitivity may also be experienced by some patients, especially in low lighting conditions such as driving at night. In order to achieve optimal visual performance with this lens, emmetropia must be targeted. Patients with significant preoperative or expected postoperative astigmatism >1.0D may not achieve optimal visual outcomes. Care should be taken to achieve IOL centration, as lens decentration may result in a patient experiencing visual disturbances under certain lighting conditions.

PRECAUTIONS: Do not resterilize. Do not store over 45° C. Use only sterile irrigating solutions such as BSS® or BSS PLUS® Sterile Intraocular Irrigating Solution. Clinical studies with the **ACRYSOF®** ReSTOR® IOL indicated that posterior capsule opacification (PCO), when present, developed earlier into clinically significant PCO. 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. The long-term effects of filtering blue light and the clinical efficacy of that filtering on the retina have not been conclusively established.

ATTENTION: Reference the Physician Labeling/Directions for Use for a complete listing of indications, warnings and precautions.

ACRYSOF® Toric IOL Models SN60T3, SN60T4, and SN60T5

CAUTION: Federal law restricts this device to sale by or on the order of a physician.

INDICATIONS: **ACRYSOF®** Toric IOL Models SN60T3, SN60T4, and SN60T5 Posterior Chamber Intraocular lenses are intended for primary implantation in the capsular bag of the eye for the visual correction of aphakia and pre-existing corneal astigmatism secondary to the removal of a cataractous lens in adult patients with or without presbyopia, who desire improved uncorrected distance vision, reduction of residual refractive cylinder and increased spectacle independence for distance vision.

WARNINGS: 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. Toric IOLs should not be implanted if the posterior capsule is ruptured, if the zonules are damaged, or if a primary posterior capsulotomy is planned. Rotation can reduce astigmatic correction, if necessary lens repositioning should occur as early as possible prior to lens encapsulation. All viscoelastics should be removed from both the anterior and posterior sides of the lens; residual viscoelastics may allow the lens to rotate.

PRECAUTIONS: 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. The long-term effects of filtering blue light and the clinical efficacy of that filtering on the retina have not been conclusively established. 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.

The ACRYSOF® IQ Aspheric Natural & ACRYSOF® Natural IOLs

CAUTION: Federal law restricts this device to sale by or on the order of a physician.

INDICATIONS: **ACRYSOF®** IQ Aspheric Natural Intraocular lenses and the **ACRYSOF®** Natural Posterior Chamber Intraocular Lenses (IOLs) are indicated for the replacement of the human lens to achieve visual correction of aphakia in adult patients following cataract surgery. These lenses are intended for placement in the capsular bag.

WARNINGS: 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. Some adverse reactions that have been associated with the implantation of intraocular lenses are: hypopyon, intraocular infection, acute corneal decompensation and secondary surgical intervention. Caution should be used prior to lens encapsulation to avoid lens decentrations or dislocations.

PRECAUTIONS: 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. The long-term effects of filtering blue light and the clinical efficacy of that filtering on the retina have not been conclusively established. 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 Physician Labeling/Directions for Use for a complete listing of indications, warnings and precautions. The long-term effects of filtering blue light and the clinical efficacy of that filtering on the retina have not been conclusively established.

Holladay IOL Surgeon
Factor Conversion Table

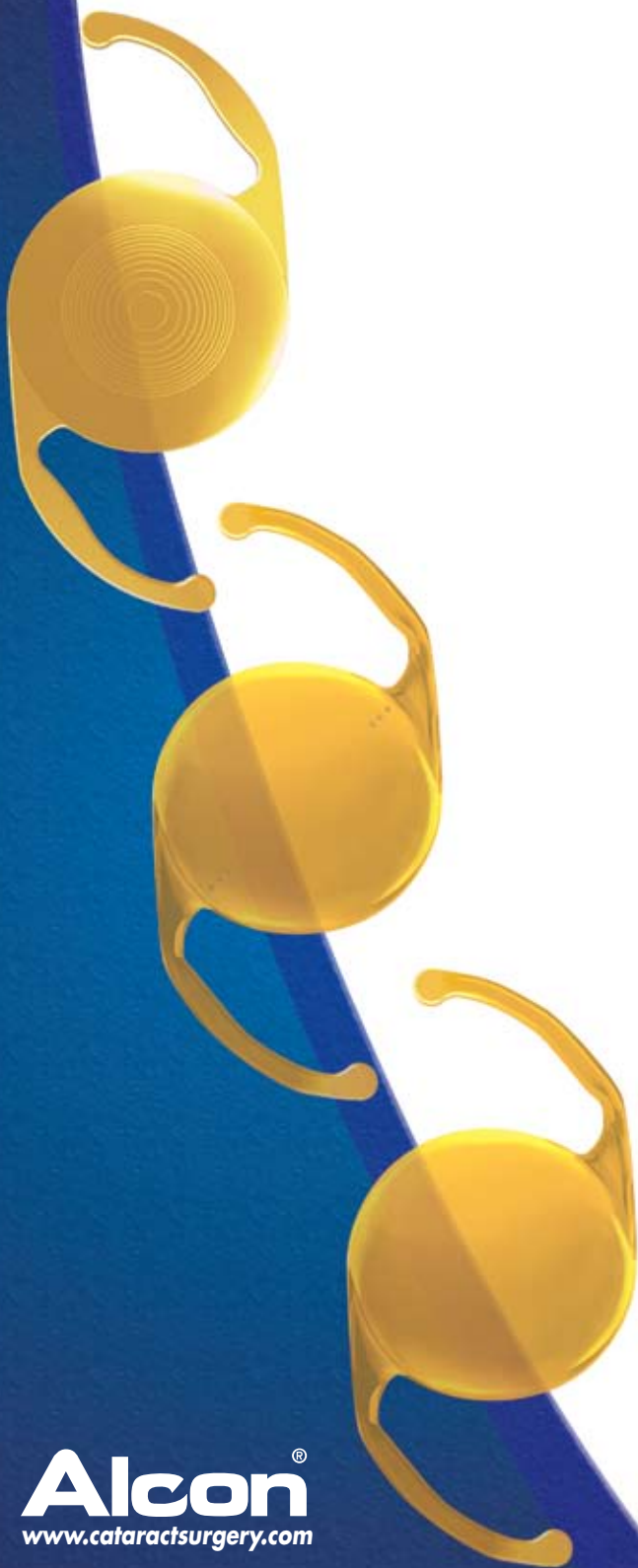
A-CONSTANT	S FACTOR	ACD
110.0	-3.31	0.30
110.1	-3.25	0.36
110.2	-3.19	0.41
110.3	-3.14	0.47
110.4	-3.08	0.53
110.5	-3.02	0.59
110.6	-2.97	0.65
110.7	-2.91	0.70
110.8	-2.85	0.76
110.9	-2.80	0.82
111.0	-2.74	0.88
111.1	-2.68	0.94
111.2	-2.63	1.00
111.3	-2.57	1.06
111.4	-2.51	1.11
111.5	-2.46	1.17
111.6	-2.40	1.23
111.7	-2.34	1.29
111.8	-2.29	1.35
111.9	-2.23	1.40
112.0	-2.17	1.46
112.1	-2.12	1.52
112.2	-2.06	1.58
112.3	-2.00	1.64
112.4	-1.95	1.70
112.5	-1.89	1.76
112.6	-1.84	1.81
112.7	-1.78	1.87
112.8	-1.72	1.93
112.9	-1.66	1.99
113.0	-1.61	2.05
113.1	-1.55	2.11
113.2	-1.50	2.16
113.3	-1.44	2.22
113.4	-1.38	2.28
113.5	-1.32	2.34
113.6	-1.27	2.40
113.7	-1.21	2.46
113.8	-1.16	2.51
113.9	-1.10	2.57
114.0	-1.04	2.63
114.1	-0.98	2.69
114.2	-0.93	2.75
114.3	-0.87	2.81
114.4	-0.82	2.86
114.5	-0.76	2.92
114.6	-0.70	2.98
114.7	-0.64	3.04
114.8	-0.59	3.10
114.9	-0.53	3.16

Holladay IOL Surgeon
Factor Conversion Table

A-CONSTANT	S FACTOR	ACD
115.0	-0.48	3.21
115.1	-0.42	3.27
115.2	-0.36	3.33
115.3	-0.31	3.39
115.4	-0.25	3.45
115.5	-0.19	3.51
115.6	-0.14	3.56
115.7	-0.08	3.62
115.8	-0.02	3.68
115.9	0.03	3.74
116.0	0.09	3.80
116.1	0.15	3.86
116.2	0.20	3.91
116.3	0.26	3.97
116.4	0.32	4.03
116.5	0.37	4.09
116.6	0.43	4.15
116.7	0.49	4.21
116.8	0.54	4.26
116.9	0.60	4.32
117.0	0.66	4.38
117.1	0.71	4.44
117.2	0.77	4.50
117.3	0.83	4.56
117.4	0.88	4.62
117.5	0.94	4.67
117.6	1.00	4.73
117.7	1.05	4.79
117.8	1.11	4.85
117.9	1.17	4.91
118.0	1.22	4.96
118.1	1.28	5.02
118.2	1.34	5.08
118.3	1.39	5.14
118.4	1.45	5.20
118.5	1.51	5.26
118.6	1.56	5.32
118.7	1.62	5.37
118.8	1.68	5.43
118.9	1.73	5.49
119.0	1.79	5.55
119.1	1.85	5.61
119.2	1.90	5.66
119.3	1.96	5.72
119.4	2.02	5.78
119.5	2.07	5.84
119.6	2.1	5.90
119.7	2.19	5.96
119.8	2.24	6.02
119.9	2.30	6.07
120.0	2.36	6.13

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