TECNIS® TORIC II 1-PIECE IOL
Hydrophobic Acrylic

HAPTICS OFFSET FOR 3-POINTS OF FIXATION

Anterior Side

TECNIS® IOL WAVEFRONT-DESIGN TORIC ASPHERIC SURFACE

Posterior Side

HAPTICS HAVE A SQUARED AND FROSTED DESIGN

13.0 mm OVERALL LENGTH

ANTERIOR CYLINDER AXIS MARKS DENOTE IOL MERIDIAN WITH LOWEST POWER

ProTEC FROSTED, CONTINUOUS 360° POSTERIOR SQUARE EDGE

6.0 mm OPTIC DIAMETER
INDICATIONS AND IMPORTANT SAFETY INFORMATION

Rx Only

ATTENTION

Reference the Directions for Use labeling for a complete listing of Indications and Safety Information.

INDICATIONS

The TECNIS® Toric II 1-Piece IOL posterior chamber lens is indicated for the visual correction of aphakia and pre-existing corneal astigmatism of one diopter or greater in adult patients with or without presbyopia in whom a cataractous lens has been removed by phacoemulsification and who desire improved uncorrected distance vision, reduction in residual refractive cylinder, and increased spectacle independence for distance vision. The device is intended to be placed in the capsular bag.

WARNINGS

Physicians considering lens implantation should weigh the potential risk/benefit ratio for any circumstances described in the TECNIS® Toric II 1-Piece IOL Directions for Use that could increase complications or impact patient outcomes. The clinical study did not show evidence of effectiveness for the treatment of preoperative corneal astigmatism of less than one diopter. The TECNIS® Toric II 1-Piece IOL should not be placed in the ciliary sulcus.

Rotation of the TECNIS® Toric II 1-Piece IOL away from its intended axis can reduce its astigmatic correction. Misalignment greater than 30° may increase postoperative refractive cylinder.

PRECAUTIONS

Accurate keratometry and biometry in addition to the use of the TECNIS® Toric Calculator (www.TecnisToricCalc.com) are recommended to achieve optimal visual outcomes. The safety and effectiveness of the toric intraocular lens have not been substantiated in patients with certain preexisting ocular conditions and intraoperative complications. Refer to the TECNIS® Toric II 1-Piece IOL Directions for Use for a complete description of the preexisting conditions and intraoperative complications. All preoperative surgical parameters are important when choosing a toric lens for implantation. Variability in any of the preoperative measurements can influence patient outcomes. All corneal incisions were placed temporally in the clinical study. When the insertion system is used improperly, the haptics of the TECNIS® Toric II 1-Piece IOL may become broken. Please refer to the specific instructions for use provided with the insertion instrument or system. Do not reuse, resterilize, or autoclave.

ADVERSE EVENTS

The most frequently reported cumulative adverse event that occurred during the TECNIS® Toric 1-Piece IOL clinical trial was surgical re-intervention which occurred at a rate of 3.4% (lens repositioning procedures and retinal repair procedures).

To learn more and to view important safety information, please review the TECNIS Toric II IOL Directions For Use (DFU).

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