

Break free from tradition. Unleash the power of the PanOptix[®] IOL.

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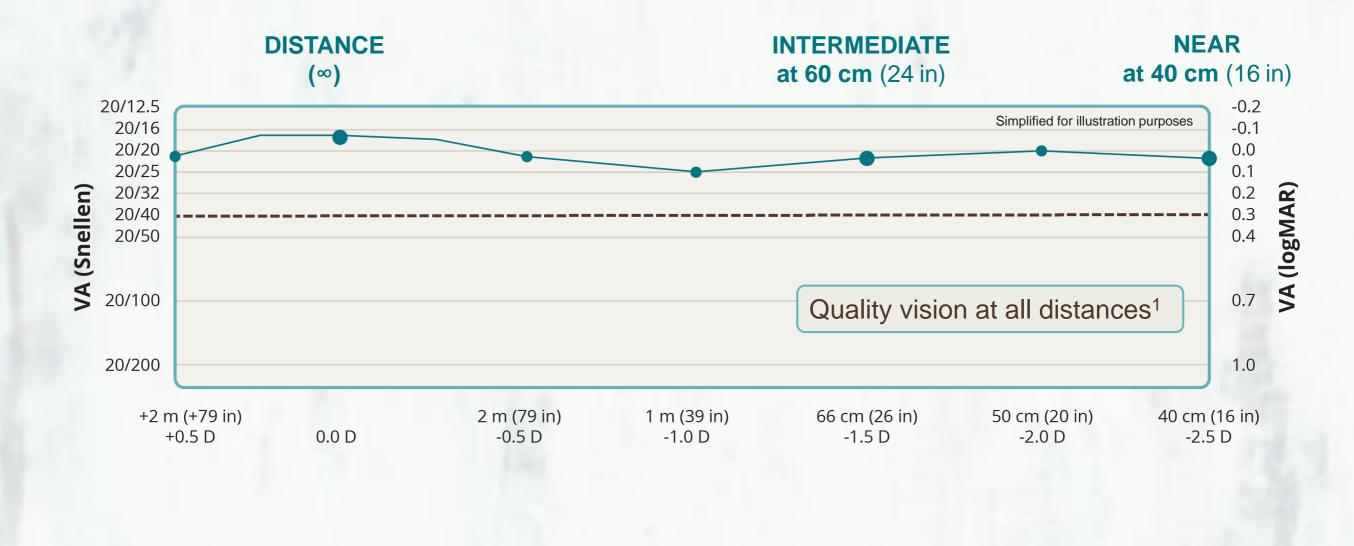
THRILL







20/20 NEAR, INTERMEDIATE AND DISTANCE VISION IS NOW POSSIBLE^{*, †,1}



*Based on mean value of binocular defocus curve at near, intermediate and distance at 6 months (n=127).

[†]Snellen VA was converted from logMAR VA. A Snellen notation of 20/20⁻² or better indicates a logMAR VA of 0.04 or better, which means 3 or more of the 5 ETDRS chart letters in the line were identified correctly.



ENLIGHTEN® OPTICAL TECHNOLOGY

The PanOptix[®] lens is an advanced trifocal IOL equipped with **ENLIGHTEN**[®] Optical Technology — a proprietary design that optimizes intermediate without compromising exceptional near and distance vision.

88% total light utilization at a 3 mm pupil size²

(Light allocation: 50% distance, 25% intermediate, 25% near)

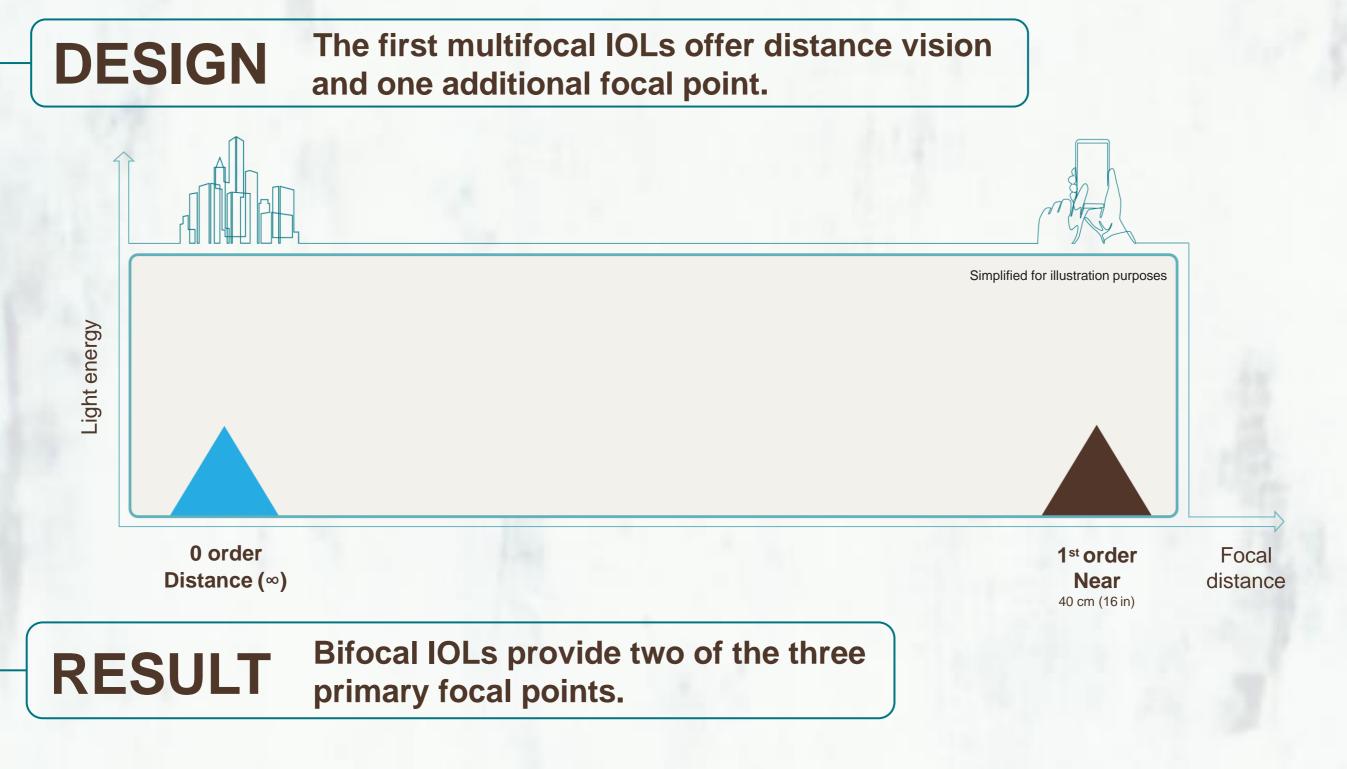
Reduces dependence on pupil size^{2,3} with a 4.5 mm diffractive zone

4.5 mm





THE ENLIGHTEN® OPTICAL DESIGN STORY: STEP 1 BIFOCAL IOLS



REFERENCES

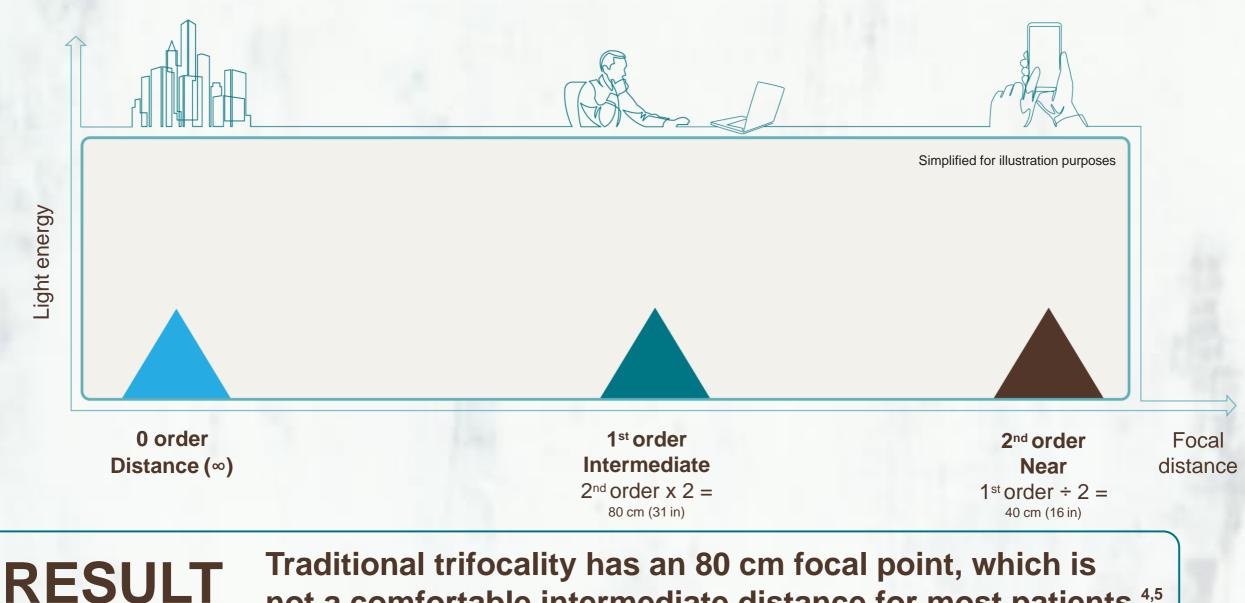


DESIGN



THE ENLIGHTEN® OPTICAL DESIGN STORY: STEP 2 TRADITIONAL TRIFOCALITY

Based on the rules of diffractive optics, trifocality requires sequential, equally spaced focal points.



not a comfortable intermediate distance for most patients.^{4,5}

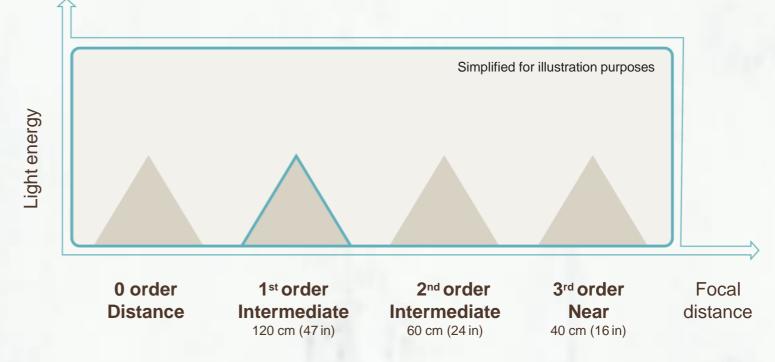
REFERENCES





THE ENLIGHTEN® OPTICAL DESIGN STORY: STEP 3 PANOPTIX® ADVANCED TRIFOCAL IOL

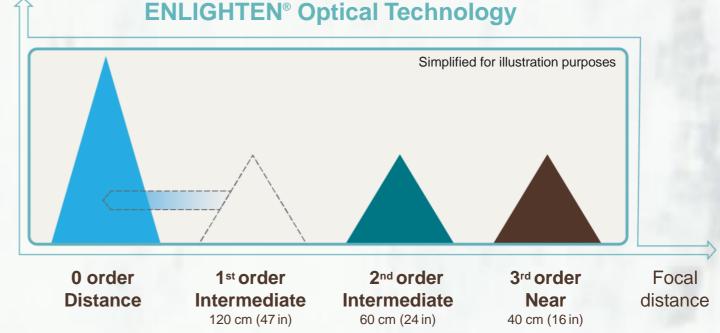
DESIGN To create **ENLIGHTEN®** Optical Technology, we first created an additional focal point at 120 cm.



Then, we redirected the new 1st order intermediate focal point's light energy to distance.

MENU

Light energy



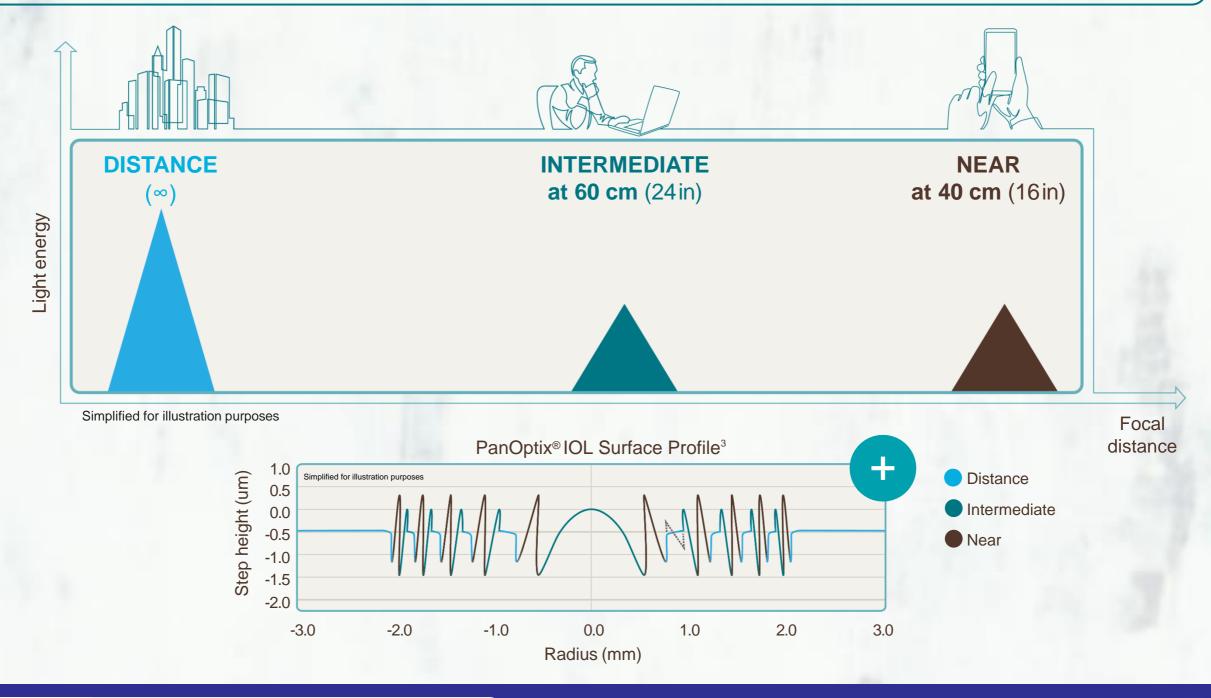
REFERENCES





THE ENLIGHTEN® OPTICAL DESIGN STORY: STEP 4 REIMAGINE MULTIFOCAL PERFORMANCE

RESULT ENLIGHTEN® Optical Technology optimizes intermediate without compromising near and distance vision.



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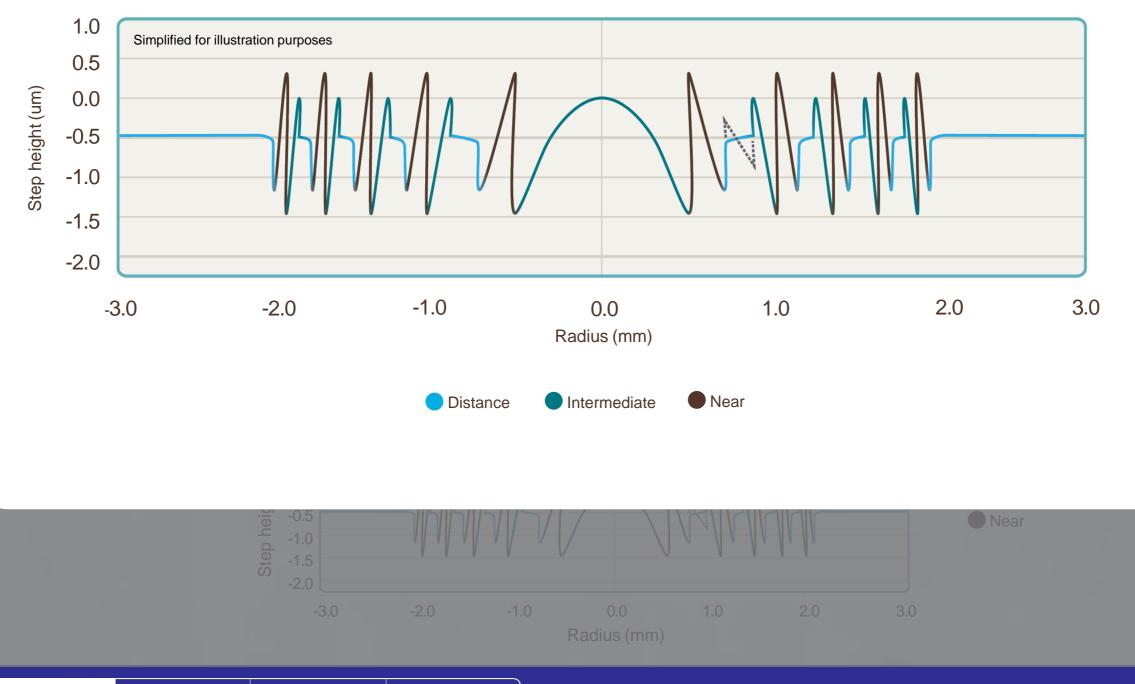


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THE ENLIGHTEN® OPTICAL DESIGN STORY: STEP 4

REIMAGINE MULTIFOCAL PERFORMANCE

PanOptix® IOL Surface Profile³



⇒





UNDENIABLE PATIENT SATISFACTION⁶

Data collected 6 months post-op:

99.2%

of patients would have had the same lens implanted again.**,‡,6

**n=127

[‡]Response to the following question on IOLSAT questionnaire (Ver. 1.0, Dec. 20, 2018) at 6 months post-op: "Given your vision today, if you had to do it all over, would you have the same lenses implanted again?"

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REFERENCES





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Break free from tradition and unleash the power of the PanOptix® IOL.

20/20 near, intermediate and distance vision is now possible*,t,1

Proprietary ENLIGHTEN® Optical Technology

99.2% of patients would have had the same lens implanted again^{‡,6}

^{*}Based on mean value of binocular defocus curve at near, intermediate and distance at 6 months (n=127). [†]Snellen VA was converted from logMAR VA. A Snellen notation of 20/20⁻² or better indicates a logMAR VA of 0.04 or better, which means 3 or more of the 5 ETDRS chart letters in the line were identified correctly. [‡]Response to the following question on IOLSAT questionnaire (Ver. 1.0, Dec. 20, 2018) at 6 months post-op: "Given your vision today, if you had to do it all over, would you have the same lenses implanted again?"

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Available in toric on the platform you can rely on

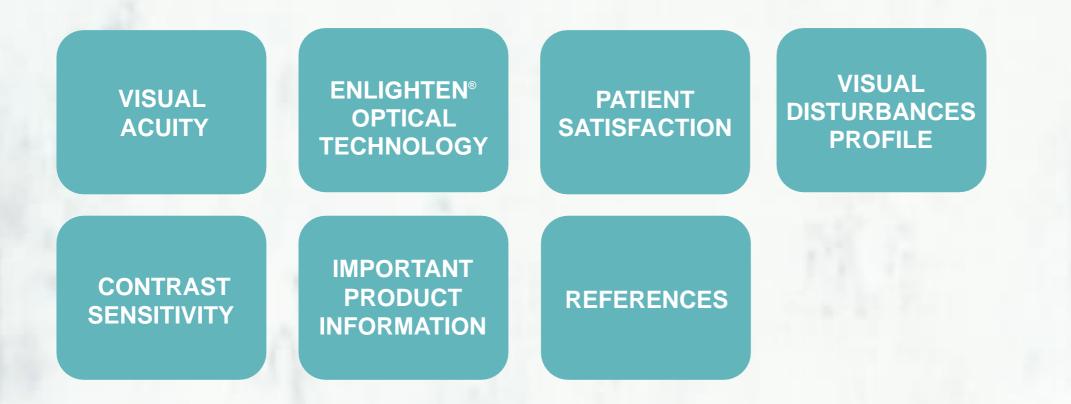
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Select a chapter to learn more about the **PanOptix® IOL**.



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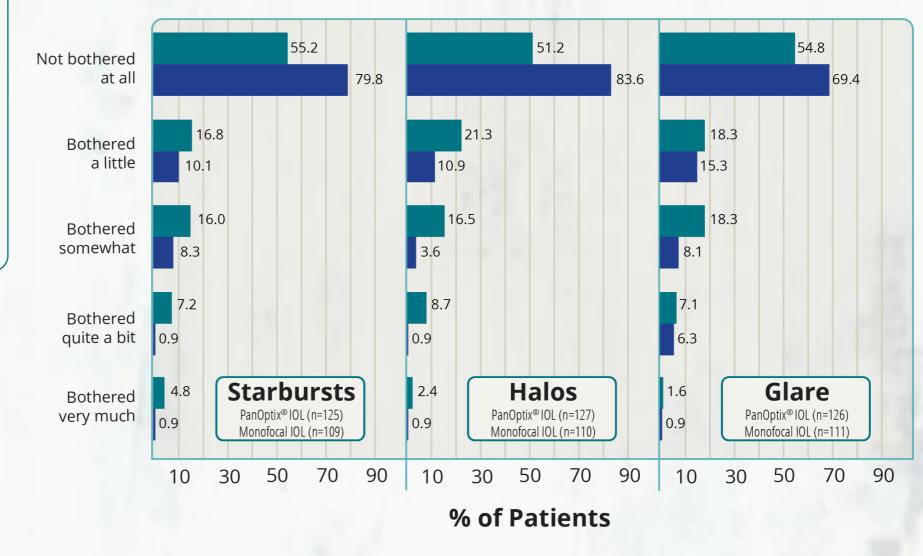
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PATIENT-REPORTED VISUAL DISTURBANCES⁶

Results from a **patient-prompted** and **validated** QUVID questionnaire at 6 months when asked: "In the past 7 days, how much were you bothered with starbursts, halos and glare?"



Ratings of How Much Patients Were Bothered by Visual Symptoms at 6 Months

PanOptix[®] IOL (n=129 patients) Monofocal IOL (n=114 patients)

4.8% by Starbursts (n=125)
2.4% by Halos (n=127)

much by:

Most bothersome visual

disturbances. Percent of

patients bothered very

• 1.6% by Glare (n=126)

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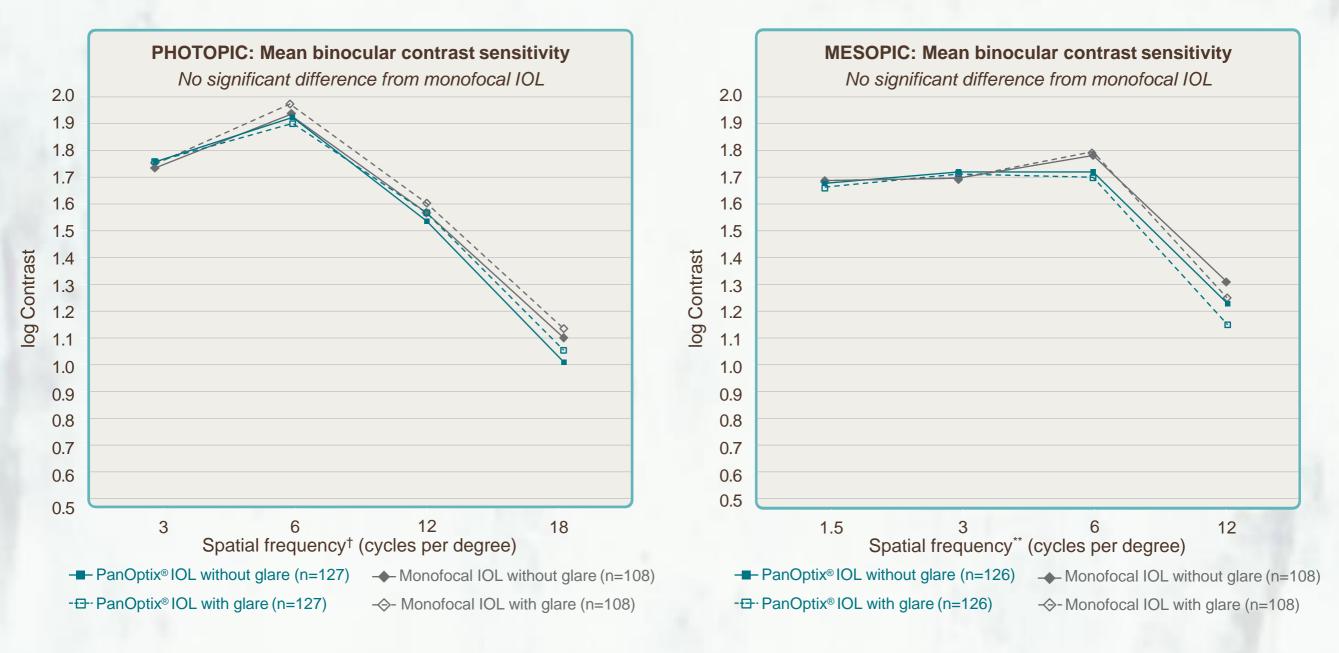
REFERENCES





CONTRAST SENSITIVITY

The **PanOptix**[®] **IOL** delivers excellent contrast sensitivity compared to a monofocal IOL at 6 months — in both photopic and mesopic conditions.^{*,1}



*Contrast above 0.3 is clinically meaningful. †At 85 cd/m2 in photopic conditions. **At 3 cd/m2 in mesopic conditions.

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REFERENCES

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IMPORTANT PRODUCT INFORMATION

AcrySof® IQ PanOptix® Family of Trifocal IOLs 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 PanOptix[®] Trifocal IOLs include AcrySof[®] IQ PanOptix[®] and AcrySof[®] IQ PanOptix[®] Toric and are indicated for primary implantation in the capsular bag in the posterior chamber of the eye for the visual correction of aphakia in adult patients, with less than 1 diopter of pre-existing corneal astigmatism, in whom a cataractous lens has been removed. The lens mitigates the effects of presbyopia by providing improved intermediate and near visual acuity, while maintaining comparable distance visual acuity with a reduced need for eyeglasses, compared to a monofocal IOL. In addition, the AcrySof[®] IQ PanOptix[®] Toric Trifocal IOL is indicated for the reduction of residual refractive astigmatism.

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. Physicians should target emmetropia and ensure that IOL centration is achieved. For the AcrySof® IQ PanOptix® Toric Trifocal IOL, the lens 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. Some visual effects may be expected due to the superposition of focused and unfocused multiple images. These may include some perceptions of halos or starbursts, as well as other visual symptoms. As with other multifocal IOLs, there is a possibility that visual symptoms may be significant enough that the patient will request explant of the multifocal IOL. A reduction in contrast sensitivity as compared to a monofocal IOL may be experienced by some patients and may be more prevalent in low lighting conditions. Therefore, patients implanted with multifocal IOLs should exercise caution when driving at night or in poor visibility conditions. Patients should be advised that unexpected outcomes could lead to continued spectacle dependence or the need for secondary surgical intervention (e.g., intraocular lens replacement or repositioning). As with other multifocal IOLs, patients may need glasses when reading small print or looking at small objects. Posterior capsule opacification (PCO) may significantly affect the vision of patients with a copy of the Patient Information Brochure, available from Alcon, informing them of possible risks and benefits associated with the AcrySof® IQ PanOptix® Trifocal IOLs.

ATTENTION: Reference the Directions for Use labeling for each IOL for a complete listing of indications, warnings and precautions.







REFERENCES

- 1. AcrySof[®] IQ PanOptix[®] Directions for Use.
- 2. Alcon Data on File.
- 3. Alcon Data on File.
- 4. Carson D, et al. Optical bench performance of 3 trifocal intraocular lenses. J Cataract Refract Surg. 2016;42:1361-1367.
- 5. Kohnen T, et al. Visual performance of a quadrifocal (trifocal) intraocular lens following removal of the crystalline lens. Am J Ophthalmol. 2017;184:52-62.
- 6. Alcon Data on File.

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