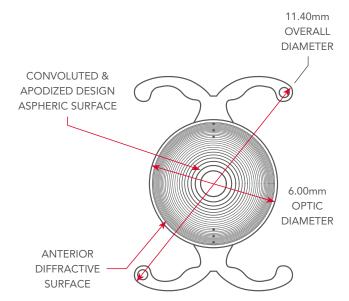
ONLINE TORIC CALCULATOR WITH **ABULAFIA-KOCH** FORMULA:

WWW.PHYSIOLTORIC.EU



FINEVISION TORIC

Trifocal Toric Hydrophilic



Description

					505	~				
Model	POD FT									
Material	26% Hydrophilic Acrylic									
Overall diameter	11.40mm									
Optic diameter	6.00mm									
Optic	Biconvex Aspheric Toric Trifocal									
Haptic design	Double C-loop & Posterior Angulated Haptic									
Filtration	UV & Blue Light									
Refractive index	1.46									
Abbe number	58									
Additional power (IOL plane)	+1.75D & +3.50D									
Injection system	Medicel Accuject 2.0 up to 24.5D and Medicel Accuject 2.1/2.2 up to 35D									
Spherical power	+6D to +35D (0.5D steps)									
Cylinder power (IOL plane)	1.00 - 1.50 - 2.25 - 3.00 - 3.75 - 4.50 - 5.25 - 6.00D									
Suggested A constant ¹					Interferometry					
	Hoffer Q: pACD			5.59						
	Holladay 1: Sf			1.83						
	Barrett: LF			1.86						
	SRK/T: A Haigis²: a0; a1; a2				118.95					
					1.36; 0.4; 0.1					
	POD FT 1.0	POD FT 1.5	POD FT 2	2.25	POD FT 3.0	POD FT 3.75	POD FT 4.5	POD FT 5.25	POD FT 6.0	
Cylinder power at IOL plane	1.00D	1.50D	2.250)	3.00D	3.75D	4.50D	5.25D	6.00D	
Cylinder power at corneal plane ³	0.68D	1.03D	1.55D		2.06D	2.57D	3.08D	3.60D	4.11D	

¹ Values estimated only: surgeons are recommended to personalize their A-constant based on their surgical techniques and equipment, experience with the lens model and postoperative results.

² Not optimized.

³ Savini G., J Cataract Refract Surg 2013; 39:1900-1903.

Note: The FINEVISION TORIC intraocular lens is not FDA approved.

bvimedical.com

Product Information

Manufacturer	PhysIOL s.a Liège Science Park Allée des Noisetiers 4 B-4031 Belgium +32 4 361 05 49 physiol@bvimedical.com			
Certificate information	CE: Certificate N° CE658516 ISO 13485:2016: Certificate N° MD658518 MDSAP: Certificate N° MDSAP 691544			
Shelf life	Five (5) years from manufacturing date			
Intended Use	Intended use (for all IOLs): The posterior chamber intraocular lens which is intended to be placed into the capsular bag for the replacement of the human lens to achieve the visual correction of aphakia in adult patients in whom the cataractous lens has been removed by extracapsular cataract extraction.			
Indication for use	The lens should be used as intended in adult patients, with pre-existing astigmatism, surgically treated for cataract, with possibly associated presbyopia, who desire improved uncorrected far vision, useful near and intermediate visual functions and reduced spectacle dependence.			
Product Composition	No products of animal or human origin are present in the implant. The implant is made of the HELIOFLEX material, composed of an acrylate copolymer Hydroxyethyl methacrylate (HEMA) and Methyl methacrylate (MMA), including a UV and blue light filter			
For sterile product	All IOLs from PhysIOL are steam sterilized			
Packaging Material	Holder (Polypropylene) Container (Polypropylene) Storage liquid (0.9% NaCl solution) Aluminium lid (Aluminium Gold) Container label (paper) Blister PP (Polypropylene) Tyvek lid			
Product Class	MDD Class IIb Sterile, According to European Medical Device Directive 93/42/EEC Not available in the United States			

