

ALCON IOL POWER CALCULATION CONSULT FORM

This form is being sent in response to your unsolicited request for support with IOL calculations. Should you have any questions or need assistance in completing this form, please contact Alcon Medical Information Services at **800.757.9785**.

Alcon IOL Power Calculation Assistance Policy:

Alcon performs intraocular lens (IOL) power calculations in response to unsolicited requests for assistance from surgeons using Alcon IOLs. This service is provided as a courtesy to Alcon customers and requests for IOL calculations will only be accepted for Alcon products. In order to help determine an accurate calculation, it may be necessary for Alcon to contact the requesting surgeon for confirmation of submitted information and/or additional data to support the request. Alcon may decline calculation services for lack of sufficient reliable data or for misuse of these services.

Calculation services require a minimum of two business days after all required data have been received for non-post-refractive ReSTOR and Toric IOL model requests; "rush" or immediate services are not available.

Calculation services for all other models and for post-refractive IOL requests are not available at this time.

Please circle one:

Accept Terms

Decline Terms

Surgeon Name _____ Today's Date ____/____/____

Phone# _____ Fax# _____

Email Address: _____ Account # _____

Surgeon Attestation:

I, as the operating surgeon, attest that the data and information provided herein are true and accurate. I request that Alcon provide me with calculation confirmation using the Holladay 2 formula to assist me in making a decision regarding intraocular lens power selection. I acknowledge that Alcon's service is dependent on the accuracy of my data. I understand that Alcon does not intend to provide medical or surgical advice, does not warrant or guarantee the accuracy or completeness of this information and disclaims any liability for unanticipated refractive or surgical outcomes. I release Alcon from any liability for this service. I further acknowledge that the selection of the specific intraocular lens model, A-constant and power used for this patient is solely my responsibility as the operating surgeon.

Surgeon Initials:

Patient Details

Patient Age _____ Patient ID _____ Gender: M F

Calculation Assistance Requested for: OD OS OU

Date of Surgery: OD ____/____/____ OS ____/____/____

FOR VALIDATION PURPOSES, PLEASE PROVIDE RELEVANT DATA FOR BOTH EYES EVEN IF REQUEST IS ONLY FOR ONE EYE.

FAX COMPLETED FORM TO ALCON AT 800.757.9786 or EMAIL to medinfo@alconlabs.com

ALCON IOL POWER CALCULATION CONSULT FORM

Surgeon Name _____ Fax# _____ Patient ID _____
FOR VALIDATION PURPOSES, PLEASE PROVIDE DATA FOR BOTH EYES, EVEN IF REQUEST IS FOR ONE EYE. Should you have any questions or need assistance in completing this form, please contact Alcon Medical Information Services at 800.757.9785.

OD	
Preoperative Data	
Pre-Operative Refraction: _____ x _____ <input type="checkbox"/> Pre-Cataract "Most Plus" (Preferred) <input type="checkbox"/> Current	
Horiz White to White _____ mm	Phakic Lens Thickness (rec. for AL < 22.0mm) _____ mm
K1 _____ D @ _____	K2 _____ D @ _____ Method for Ks: _____
Axial Length _____ mm	Phakic ACD _____ mm
Method used for Axial Length: <input type="checkbox"/> Contact <input type="checkbox"/> Immersion <input type="checkbox"/> IOLMaster / LENSTAR	
Does the patient have a history of any of the following:	
Keratoconus: <input type="checkbox"/> Yes <input type="checkbox"/> No	Scleral Buckle: <input type="checkbox"/> Yes <input type="checkbox"/> No Silicone Oil: <input type="checkbox"/> Yes <input type="checkbox"/> No
Other Ocular Pathology: <input type="checkbox"/> Yes <input type="checkbox"/> No	If yes, please explain: _____
Keratorefractive Surgery: <input type="checkbox"/> Yes <input type="checkbox"/> No	If yes, please explain: _____
Surgical Plan Alcon Lens Model _____ Lens Constant _____ Optimized ? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Target Ref _____ D	IOL Power _____ D Formula Used _____

OS	
Preoperative Data	
Pre-Operative Refraction: _____ x _____ <input type="checkbox"/> Pre-Cataract "Most Plus" (Preferred) <input type="checkbox"/> Current	
Horiz White to White _____ mm	Phakic Lens Thickness (rec. for AL < 22.0mm) _____ mm
K1 _____ D @ _____	K2 _____ D @ _____ Method for Ks: _____
Axial Length _____ mm	Phakic ACD _____ mm
Method used for Axial Length: <input type="checkbox"/> Contact <input type="checkbox"/> Immersion <input type="checkbox"/> IOLMaster / LENSTAR	
Does the patient have a history of any of the following:	
Keratoconus: <input type="checkbox"/> Yes <input type="checkbox"/> No	Scleral Buckle: <input type="checkbox"/> Yes <input type="checkbox"/> No Silicone Oil: <input type="checkbox"/> Yes <input type="checkbox"/> No
Other Ocular Pathology: <input type="checkbox"/> Yes <input type="checkbox"/> No	If yes, please explain: _____
Keratorefractive Surgery: <input type="checkbox"/> Yes <input type="checkbox"/> No	If yes, please explain: _____
Surgical Plan Alcon Lens Model _____ Lens Constant _____ Optimized ? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Target Ref _____ D	IOL Power _____ D Formula Used _____

CALCULATION WILL INCLUDE HOLLADAY 2 FORMULA ONLY

All data points may be required to process calculation

Comments: _____

****REQUIRED****

SURGEON SIGNATURE:

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