





LENSTEC, INC.  
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## CUSTOMER RETURN AUTHORIZATION FORM

### EVENT SPECIFICS: (Please check all boxes that apply to this event)

Serial Number: \_\_\_\_\_

Date of Surgery: \_\_\_\_\_

Did the product have any Patient Contact?

Yes

No

Was the lens itself:

Destroyed

Discarded

Lost

N/A

If yes, please state: \_\_\_\_\_

Was this an issue due to:

Handling/User Error? (NO product complaint)

Defective Product?

Other\*

Specifically (check any/all that apply):

Loading Issues

Debris on Lens

Folding / Unfolding Issues

Cartridge Defective

Stuck in Delivery System

Broken Haptic

Other (note below)

Cracked / Torn Lens

Cartridge Lot #: \_\_\_\_\_

Was the IOL explanted/removed from the eye?

Yes

No

If yes, Date of implant \_\_\_\_\_

Date of Explant \_\_\_\_\_

Was the incision enlarged to remove IOL?

Yes

No

Was there any patient injury? (if yes, please explain)

Yes

No

Was another lens used? \_\_\_\_\_

\*If yes, same model? \_\_\_\_\_

\*Notes / Other:

### LENSTEC STAFF USE ONLY

Awareness Date: \_\_\_\_\_

Return Authorization Number Assigned: \_\_\_\_\_

By/Date: \_\_\_\_\_

Send completed Return Authorization form along with device(s) to Lenstec Customer Service at the address above.