

COMPLAINT HANDLING FORM

CUSTOMER INFORMATION	
FACILITY NAME:	SURGEON NAME:
ADDRESS LINE 1:	ADDRESS LINE 2:
CITY: STATE: ZIP:	CONTACT PERSON:
ACCOUNT NUMBER:	PHONE:
DATE OF PROCEDURE:	EMAIL:
INCIDENT DESCRIPTION	
Describe the incident in detail, including when it occurred (before, during or after surgery)? List any other devices, instruments and/or accessories that were used with the product during the case.	
PATIENT IMPACT YES NO If yes, please explain how the patient was impacted.	
CLINICAL CONSEQUENCES YES NO	If yes, please explain how the patient was treated and the date of treatment.



PRODUCT DETAIL	
ITEM NUMBER:	
ITEM DESCRIPTION:	
LOT/SERIAL NUMBER:	
EXPIRATION DATE:	
Is the product available for return to FCI? YES NO	
If the product is not available for return, is a photo of the product available to submit for review?	
IMPORTANT	
WHEN RETURNING A PRODUCT THAT HAS HAD PATIENT CONTACT, CUSTOMER MUST PROVIDE PROOF OF DECONTAMINATION.	
PLEASE HAVE PROOF CLEARLY MARKED ON BOX.	
CUSTOMER SIGNATURE DATE	
CONFIRMS THAT THE INFORMATION PROVIDED HEREIN IS COMPLETE, ACCURATE AND TRUE.	
To be completed by FCI Ophthalmics:	
IMMEDIATE ACTION	
Complaint Reported to Quality YES NO	

Date Reported: _____

RMA Number Provided to Customer:

Completed By:

Complaints must be submitted within 48 hours of incident