McLaren Flint Implanted / Extended Device Record

☐ Implanted ☐ Explanted Device Record	
Date: Surgeon:	
Procedure/Locations:	
Place manufacturer product label in the space below, and/or write in the following information:	
Device description:	
Name:	
S.N./Lot #:	
Catalog #:	
Manufacturer:	
Model #:	
Quantity:	
Exp. Date:	
<u>Tissues Package Integrity</u>	Reconstitution:
Supplier Package intact: ☐ Yes ☐ No ☐ N/A	Reconstitution: ☐ Yes ☐ No
Product Package intact: ☐ Yes ☐ No	Reconstitution by:
Temperature Maintained: ☐ Yes ☐ No	Reconstitution Date/time:
Product Label Legible: ☐ Yes ☐ No	Reconstitution instructions available: ☐ Yes ☐ No ☐ N/A
	Type of Fluid:
	Lot # of Fluid:
	Exp. Date of Fluid:
Implants sterilized in house: Date of sterilization:	
CPD: Sterilization Load Number:	
OR: Sterilizer Used:	Cycle Count #:
In the event of an Explant, Provide the following information:	
Reason for removal:	
Final Disposition:	
In the event of a cardiology Implant/Explant, provide the following information:	
Length of warranty:	
Parameter settings:	

IMPLANTED/EXTENDED DEVICE RECORD

M-17459 REV. 12/2016

PT.

MR.#/P.M.