UNIVERSITY OF TOLEDO HEALTH SCIENCE CAMPUS DEVICE TRACKING REPORT

(Use when manufacturer's registration document is not available)

Section 1:	Device Status	Status change date:	_
	Receipt	Date Device Received:	
	Returned to Manufacturer Implantation or Distribution	Date device put in service:	
	Returned to Inventory		
	Removed from Service Removed from Patient		
	Death of Patient		
0 11 0	D : 1 C ::		
Section 2: Device Information			
Manufacturer I	Name:	ECRI Manufacturer Code:	
Address:	7	Rep name:	
City:	State/Provin	nce: Zip/Postal Code:	<u>-</u>
Device Type:		ECRI Device Code:	AFFIX DEVICE LABEL IF AVAILABLE
Model Name:	Model #: Single Patient	ECRI Device Code: S/N:	
interided 03c.	Jangie Fatient Multiple Fatient		
Device Type:		ECRI Davica Coda:	AFFIX DEVICE LABEL IF
Model Name:	Model #:	ECRI Device Code: S/N:	AVAILABLE
Intended Use:	Single Patient Multiple Patient		
			AFFIX DEVICE LABEL IF
Device Type:	Model #	ECRI Device Code: S/N:	AVAILABLE
Intended Use:	Single Patient	S/N	
Section 3: Patient Information			
Patient Name:			
Medical Record No.: Y N Hospital Consent Form Signed			
Social Security	/ No.:	N Releas	e of Social Security Number
Section 4: Physician Information			
	•	Tolophono. (
Address	ne printed:	Telephone: () City: State:	7in·
Physician invo	Ivement: Prescribing Implanting	Explanting Primary Care/Following	ZIP
Physician Nam	ne printed:	Telephone: () City: State:	7!
Address:	Ivement: Prescribing Implanting	State:State:State:	Zip:
Employee:	Dep	partment:	Date://
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