(place patient label here)

Patient Name:



Version 1 Approved 8/22/19

<u>Please complete this form for all patients that have a cardiac implanted electronic device</u> (pacemaker/ICD) and it is anticipated that ELECTROCAUTERY or RADIOFREQUENCY may be required during their procedure.

PATIF				
17(112)	NT'S PRIMARY CARDIOLOGIST: Dr			
TREAT	TING SURGEON: Dr			
(Mr. /	Ms.)	[DOB: (procedure).	is scheduled for
	ry Date: y Site:			
<i>-Card -Bene</i>	section is completed by: liologist on-call for afterhours pro efis Cardiac Rhythm Management		er occurrence	
	Type of device -	DEFIBRIL	ATOR	
2.	Brand and model – 🗆 Boston Scien	ntific :	🗆 Medtron	nic:
	St. Jude Medical:	_ 🗆 Biotronik:_		Sorin/ELA:
3.	Current mode and lower rate limit	t		_
4.	Is the patient pacemaker depende	ent? - YES	S NO	
5.	Would you suggest applying a mag	gnet during caute	ery and/or radiofr	equency? - YES NO
6.	How will the device respond to a magnet? – Please mark as applicable.			
	a ICD THERAPIES AF	RE DISABLED		
	b PACE ASYNCHROM	NOUS AT:	BPM	
	c OTHER:			
	Would you suggest manually repro	ogramming the d	levice prior to pro	cedure? - YES NO
7.				her:
7.	If yes, how? - 🗆 VOO: b	opm 🗌 DOO:		

Date:_____Time:____