



CRH Preoperative Pacemaker/Defibrillator Information Form

Dr (Cardiologist) Date of Request:
You or your practice has been identified as managing this patient's implanted cardiac rhythm management
device. Please complete the appropriate section of this form to assist in preparation of the proposed surgery.
Preoperative cardiac evaluation may have been requested from the primary cardiologist.
TO BE COMPLETED BY SURGEON'S OFFICE STAFF:
Patient: DOB:
Procedure:
Date of Procedure: Surgeon:
Diagnosis:
Location of Procedure : CRMC VB ASC
Planned Postoperative Disposition: Outpatient Extended Recovery
□ Inpatient: Expected LOS? Clinical Reasons
TO BE COMPLETED BY CARDIOLOGIST/STAFF:
☐ Pacemaker ☐ Defibrillator (ICD/AICD)
Manufacturer:
Indications for Implantation:
Date of Last Interrogation:
Device Location: □ right chest □ left chest □ other
Is patient pacemaker dependent?
FOR PACEMAKERS:
Will magnet application <u>temporarily</u> convert device to an asynchronous pacing mode?
If magnet is used, does device need interrogation and/or does patient need to follow up with cardiologist as an
outpatient? Yes No
FOR DEFIBRILLATOR (ICD/AICD):
Will magnet application temporarily disable anti-tachycardia therapies? Yes No
Will magnet application permanently change any device settings? Yes No Unknown *
* If unknown, device needs interrogation prior to procedure to determine if magnet use is safe.
If magnet is used, does device need interrogation and/or does patient need to follow up with cardiologist as an
outpatient? Yes No
ADDITIONAL INFORMATION OR RECOMMENDATIONS:
ADDITIONAL INI ONNIATION ON RECOMMENDATIONS.
Cardiologist/PA/NP Signature: Phone/Pager:
Cardiologist/PA/NP Signature: Phone/Pager:
Date: Time:

667-POPMD-001(08/19) Patient Label

Fax this form and any associated documentation to: CRMC PSAT Fax: 757-312-6297 OR VBASC Fax: 757-312-6877