

## **PORT HURON**

## PACEMAKER INDICATION FORM

## (For Initial Implants or Generator Replacements)

 $\underline{\textbf{INSTRUCTIONS:}} \ \textbf{Review and complete all section(s)}. \ \underline{\textbf{Must}} \ \textbf{be completed for initial implants} \\ \textbf{and generator replacements} \ \textbf{prior to the initiation of the procedure}.$ 

Patient Name:		<del> </del>				
A. Indication(s) for cardiac pacemaker Imp	ant or Replacement:					
(Check <u>all</u> that apply)						
<ul> <li>Patient qualifies for cardiac resynchronization therapy</li> <li>Generator replacement is indicated</li> <li>Third degree (complete) atrioventricular (AV) block</li> <li>Second degree atrioventricular (AV) block, Mobitz type I or II</li> <li>Sinoatrial node dysfunction</li> </ul>						
			[ ] Sick sinus syndrome/ Tachycardia-bradycard	[ ] Sick sinus syndrome/ Tachycardia-bradycardia syndrome		
			[ ] Congenital heart block			
			[ ] Congenital heart disease			
			[ ] Obstructive hypertrophic cardiomyopathy			
· · · · · · · · · · · · · · · · · · ·	Sustained pause-dependent ventricular tachycardia, with or without QT prolongation					
[ ] Pacemaker insertion in advance of AV node a	·					
[ ] Other cause of non-reversible symptomatic by	oradycardia, specify:					
(Check <u>all</u> that apply) [ ] Dizziness [ ] Lightheadedness [ ] Syncope/n [ ] Exercise intolerance caused by poor heart rat [ ] Other	e response to exertion [ ] S					
C. Check one of the following:						
[ ] Patient's bradycardia, tachycardia-bradycard	a syndrome, or sustained pa	ause-dependent ventricular				
tachycardia does not have a temporary or revers	sible cause;					
OR						
[ ] Patient receives necessary medical therapy the	nat contributes to bradycard	ia that cannot be safely				
reduced or withdrawn due to the underlying cor	dition of:					
Provider Signature:	Date:	Time:				
Cath Lab Reviewer Signature:	Date:	Time:				
Scheduled Procedure Date:						

