Système de gestion de l'accès aux services



Requisition Form

ELECTROPHYSIOLOGY

Sections for Referring Physicians

decitions for receiving rangements					
Date of Request Year Month Day					
User (Additional Information)	, Area code				
Chart number of referring hospital Chart number (if known) of consulted hospital	Telephone number in case of emergency				
Referring Institution					
Referring Hospital Name	ite				
Referring Physician Specialty	Permit number				
Information					
User's Place of Origin: Home Transfer (Referring Hospital) Ward: Hospitalized – Internal Emergency – Internal Referring Hospital Contact Person Area code Fax number					
E-mail	Denominalized Code (if faxed)				
User's Year Month Day Year Month Day non-availability: From To	y Reason				
Infection	Substance Abuse				
☐ MRSA¹+ ☐ VRE²+ ☐ Other:	Yes \(\sum \) No				
Previous Electrophysiology Study , Date , Hospital					
Yes No					
Consent to release information Signed Not	signed				
Reason for Request					
□ Primo procedure □ Redo □ Diagnostic study □ Tilt table □ Internal cardioversion □ External cardioversion □ Defibrillator test □ Implantable monitor					
Pacemaker Ab	lation				
Simple New Implant Double Replacement of electrode	Complete AV Node Transeptal				
Biventricular Header replacement Upgrade	Pulmonary veins				
Repositioning of electrode Header repositioning	Adult (≥ 18 years) ☐ 3D Mapping				
Double Biventricular Electrode extraction Pediatric (< 18 years or < 30 kg) Removal					
Dependant:	General anesthesia				
Other: Referral:	Service Dr				

 $^{^{\}rm 1}$ MRSA: Methicillin-Resistant Staphylococcus aureus – $^{\rm 2}$ VRE: Vancomycin-Resistant Enterococci

	Name and Surname
User Identification	

Clinical Information					
Cardiac					
Insufficiency Classification (NYHA): 1 2 Myocardial Infarction: Acute < 1 week	\square 3 \square 4 \square < 3 months \square > 3 months				
Myocardiai imarction:	☐ < 3 months ☐ > 3 months				
Potentially Malignant Arrhythmia Yes No	Hemodynamically Unstable	☐ Yes ☐ No			
Recurrent Arrhythmia Yes No	Sudden Death	☐ Yes ☐ No			
Malignant Ventricular Arrhythmia Yes No	Nocturnal Pause	☐ Yes ☐ No			
Right Bundle Branch Block Yes No	Prophylaxis	☐ Yes ☐ No			
Left Bundle Branch Block	Node Re-entry	☐ Yes ☐ No			
1st Degrée A-V Block Yes No	Temporary Transvenous Pacemak	xer Yes No			
2 nd Degrée A-V Block Yes No	Syncope	☐ Yes ☐ No			
3 rd Degrée A-V Block ☐ Yes ☐ No	Atrial Tachycardia	☐ Yes ☐ No			
Congenital Cardiopathy Yes No	S.V.T.	☐ Yes ☐ No			
Rapid Atrial Fibrillation Yes No	V.T.	☐ Yes ☐ No			
Atrial Flutter Yes No	W.P.W.	☐ Yes ☐ No			
Valvular Insufficiency: Yes No If yes:	Aortic Mitral Pulmo	nary Tricuspid			
Valvular Stenosis: Yes No If yes:	Aortic Mitral Pulmon	nary Tricuspid			
Prosthetic Valve:	☐ Aortic ☐ Mitral ☐ Pulmor	nary Tricuspid			
Ejection fraction: Unknown % Te	est: Echocardiography Angiogra	phy Nuclear Medecine			
Myocardial Scintigraphy – Anterior Ischemia or Multiple Iscl	nemic Zones: Yes No				
Rythm Strip of Arrhythmia Available: Yes (If sent)	☐ No				
Medication					
To be Days stopped before	To be Days stopped before	To be Days stopped before			
ASA (Aspirin) Clopidogre		none (Rythmol)			
Amiodarone (Cardarone) Digoxin	Quinidin				
	ide (Rythmodan)				
☐ Calcium Channel Blocker ☐ ☐ ☐ Flecainide	(Tambocor)	e (Coumadin)			
☐ Other: ☐					
Metabolic Disease					
Creatinine: µmol/L					
Diabetes: ☐ Yes ☐ No If yes: ☐ Trea	ted by diet				
Vascular Disease Allergies					
Previous CVA: Yes No lodine Latex Penicilin Other:					
Remarks Medical Summary					
	Inc	luded L To follow L			
Referring Name (please print)	Signature	Year Month Day			

	Access to Electrophysiology – Priority Classification (CMQ ⁽¹⁾ – RQCT ⁽²⁾)					
User's Origin	Procedure		Clinical Cardiac – Information	Priority	Delays	
User is			Hemodynamically Unstable	1	< 24 hours	
hospitalized, in			Temporary transvenous Pacemaker	1	< 24 hours	
the emergency or transferred from another hospital		User hospitalized for one of the diagnostics or other severe symptoms shown under section (Clinical Cardiac – Information)	2	< = 48 hours		
	Pacemaker/Defibrillator					
	New implant	Without dependance		3	<= 2 weeks	
	Replacement of electrode or pacemaker	With dependance		3	<= 2 weeks	
	Repositioning of electrode or pacemaker	With dependance		3	<= 2 weeks	
	Electrode extraction			3	<= 2 weeks	
User coming from home	Replacement of electrode or pacemaker	Without dependance		4	< = 4 weeks	
	Repositioning of electrode or pacemaker	Without dependance		4	< = 4 weeks	
	Upgrade			4	< = 4 weeks	
	Removal			4	< = 4 weeks	
	Ablation		Rapid atrial fibrillation	3	<= 2 weeks	
	Ablation		Potentially malignant	3	<= 2 weeks	
	Ablation		Arrhythmia	4	< = 4 weeks	
	Ablation		Atrial flutter	4	< = 4 weeks	
	Ablation		Wolf-Parkinson-White	4	< = 4 weeks	
	Ablation	Redo	Syncope	4	< = 4 weeks	
	Defibrillator test			3	<= 2 weeks	
	Implantable monitor			4	< = 4 weeks	
	Diagnostic study			4	< = 4 weeks	
	Internal cardioversion			5	< = 3 months	
	External cardioversion			5	< = 3 months	
	Ablation		(Without any other specifications)	5	< = 3 months	
	Tilting table			5	< = 3 months	

Insufficiency Classification (NYHA)(3)		
Class	Class Description	
Class 1	Users with no limitation of activities; they suffer no symptoms from ordinary activities.	
Class 2	Users with slight, mild limitation of activity; they are comfortable with rest or with mild exertion.	
Class 3	Users with marked limitation of activity; they are comfortable only at rest.	
Class 4	Users who should be at complete rest, confined to bed or chair; any physical activity brings on discomfort and symptoms occur at rest.	

⁽¹⁾ CMQ: Collège des médecins du Québec

⁽²⁾ RQCT: Réseau québécois de cardiologie tertiaire

⁽³⁾ NYHA: New York Heart Association