1.	I here	by authorize		ar	d/or such associates (or assistants as may be selected by	
	said provider to perform the following procedure(s) which has (have) been explained to me: Insertion of a Cardiac Reshynchronization Biventricular pacemaker (CRT-P) an electrical device which is inserted under the skin of the upper chest to assist the electrical and pumping function of the heart through small wire(s) placed in the heart.						
2.			l and pumping function of the heart d for my condition(s) has (have)	-	•		
		· · · · · · · · · · · · · · · · · · ·	·	<u> </u>			
3. I recognize that, during the course of the operation, post operative care, medi conditions may necessitate additional or different procedures than set forth a his or her assistants or designees, to perform such surgical or other procipudgment necessary and desirable.					bove. I therefore authorize my above named provider, and		
4.	I have been informed that there are significant risks such as severe loss of blood, infection and cardiac arrest that can lead to death or permanent or partial disability, which may occur from the performance of any procedure. Other risks include the potential hazard of prolonged or frequent radiation exposure to include but not limited to the following, short term and rare side effect: skin irritation, skin ulcers a small increase in the lifetime risk of cancer, Female (childbearing age) a small potential hazard to fetus. These risks can be serious and possibly fatal. I acknowledge that no warranty or guarantee has been made to me as to result or cure.						
6.	the dir possib arrest Any tis	rection of a provid ble damage to vital and/or brain deatl	er as may be deemed necessary.	I understand , liver and kid causes.	that all anesthetics in Iney and that in some	ist, CRNA or other qualified party undo volve risks of complications and seriou cases may result in paralysis, cardiac re with accustomed practice.	
7.	l recog a) c) (check	gnize that I have the the nature and chathe alternative form side effects, and an proposed treatment treatment, including kone) My provider has in	nformed me of the above points to at I do not want to be told of the ak	b) the ad) the r	nticipated results of the p ecognized serious possibl	oroposed treatment; e risks,	
Ο.		I consent to the ti	ransfusion of Blood and Blood Prod at to a blood transfusion during this			Non Blood Medical Management).	
l ce	rertify that this form has been fully explained to me, that I have read it and or				have had it read to me, and that I understand its contents.		
Pati	ent's Nam	ne (printed)			_		
Pati	ent / (Pare	ent if patient is a minor) /	Authorized Representative		Date	Time	
Rela	ntionship i	if Authorized Representat	ive		_		
Wit	ness to Pa	atient / Legal Guardian Sig	nature		Date	Time	
l co tre	onfirm t atment	as well as the risks a		g with the tre	atment. I have offered to	al risks and alternatives to the proposed answer any questions and have fully derstands what I have explained.	
PR	OVIDER	SIGNATURE:			Date:	Time:	
	Page 1 of 1	1				PATIENT INFORMATION	
		507120	CHI Franciscan CONSENT FOR PROCEDU	RE/TREAT	MENT		

ELECTROPHYSIOLOGY STUDY WITH

RADIOFREQUENCY

(05/10/2019)

597130