atient Name:		GRAND VIEW HEALTH 700 Lawn Avenue Sellersville, PA 18960	
Patient Birthdate:		OUTPATIENT AICD / PPM MANAGEMENT	
<u> NR</u> Affix Patient Label			
		AICD - Automated Internal Cardiac Defibrillator PPM - Permanent Pacemaker	
Patient Name		Date of Birth	
Device Type	Model #	Serial #	
Device Company			
Medtronic - 1.800.345.4943	Guidant - 1.800.CARDIAC	St. Jude's - 1.800.722.3423 Biotronic 1.800.547.0394	
A representative from Device Convolution Notes:	ompany has been confirmed by	our office and will be present for OR procedure device support.	
inactivate therapies. When prod defibrillator WILL NOT affect a	cedure is finished, remove magr any bradycardia pacing functi uring time that a magnet is over	hagnet over device and tape in place during procedure to het and therapies will be active. Donut magnets over a ion. the device, immediately remove magnet to activate device defibrillation therapies as per ACLS guidelines.	
Consult Cardiology post proced			
_			
Patient has Guidant defibriliator Patient should remain monitore		gnet application. Requires interrogation to deactivate detection. post procedure.	
Pacemaker Instructions			
Patient is Pacemaker depender applied. (Actual paced rate will		uring cautery. Device will pace at a fixed rate when magnet is et rate of device)	
over-sensing. (When a mag	net is placed over pacemake	net. Use short bursts of cautery to minimize or and patient is NOT pacemaker dependent, it derlying, intrinsic rhythm and possibly promote	
Consult Cardiology post proced	ure if device malfunction is susp	pected.	
Other Instructions			
Interrogate post-procedure			
 Does not require interrogation p 	ost-procedure		
Fax Form to Surgeon's Office:		device management and that they do not replace any necessary	
communication with the physician it	r needed to ensure safe care for		
communication with the physician it	r needed to ensure safe care for	· 	