



GWTG-HF Limited Case Record Form (CRF)

August 2026

Patient ID:			
DEMOGRAPHIC DATA			
Patient First Name		Patient Last Name	
Date of Birth		Age	
Sex	☐ Male ☐ Female ☐ Unknown		
Patient Gender Identify	☐ Male ☐ Female ☐ Female-to-Male (FTM)/Transgender Male/Trans Man ☐ Male-to-Female (MTF)/Transgender Female/Trans Woman ☐ Genderqueer, neither exclusively male nor female ☐ Additional gender category or other. _____ ☐ Did not disclose.		
Patient-Identified Sexual Orientation	☐ Straight or heterosexual ☐ Lesbian or gay ☐ Queer, pansexual, and/or questioning ☐ Something else; please specify. _____ ☐ Don't know ☐ Declined to answer		
☐ Homeless		☐ Foreign Address	
Street Address	City		
	State		
	Patient Zip Code	_____ - _____	
External Tracking ID	_____		
Race and Ethnicity			
Race and/or Ethnicity	☐ American Indian or Alaska Native ☐ Asian ☐ Asian Indian ☐ Chinese ☐ Filipino ☐ Japanese ☐ Korean ☐ Vietnamese ☐ Other Asian ☐ Black or African American ☐ Hispanic or Latino ☐ Middle Eastern or North African ☐ Native Hawaiian or Pacific Islander ☐ Native Hawaiian ☐ Guamanian or Chamorro ☐ Samoan ☐ Other Pacific Islander ☐ White ☐ UTD		
Select Hispanic Origin Group(s):	☐ Mexican, Mexican American, Chicano/a ☐ Cuban ☐ Puerto Rican ☐ Another Hispanic, Latino, or Spanish Origin		
ARRIVAL AND ADMISSION INFORMATION			
Internal Tracking ID:	_____	Physician/Provider NPI:	_____
+ Arrival Date/Time:	___/___/___ __:___	☐ Unknown Date/UTD	
Admission Date:	___/___/___		

Point of Origin for Admission or Visit:	☐ Non-Healthcare Facility Point of Origin ☐ Clinic ☐ Transfer From a Hospital (Different Facility) ☐ Transfer From a Skilled Nursing Facility (SNF) or Intermediate Care Facility (ICF) ☐ Transfer From Another Health Care Facility ☐ Emergency Room ☐ Information Not Available ☐ Transfer From a Hospice and is Under a Hospice Plan of Care or is Enrolled in a Hospice Program		
Discharge Date/Time	___/___/___ __:___		
Medical History			
Medical History (Select all that apply):			
☐ Anemia ☐ Atrial Fib (chronic or recurrent) ☐ Atrial Flutter (chronic or recurrent) ☐ ATTR-CM ☐ Hereditary ☐ Wild-type ☐ Barostim Device ☐ CAD ☐ Cancer ☐ Cardiac Ablation ☐ CardioMEMs (implantable hemodynamic monitor) ☐ Chronic Kidney Disease (CKD) ☐ COPD or Asthma ☐ CRT-D (cardiac resynchronization therapy with ICD) ☐ CRT-P (cardiac resynchronization therapy-pacing only) ☐ Current Pregnancy (up to 6 weeks postpartum) ☐ CVA/TIA ☐ Depression ☐ Diabetes ☐ Dialysis (chronic) ☐ Emerging Infectious Disease ☐ SARS-COV-2 (COVID-19) ☐ Other infectious respiratory pathogen ☐ Familial hypercholesterolemia	☐ Heart failure ☐ Heart Transplant ☐ Hyperlipidemia ☐ Hypertension ☐ Hypertriglyceridemia ☐ Hypertrophic Cardiomyopathy ☐ ICD only ☐ Long COVID ☐ Mechanical Heart Valve ☐ Mitral Stenosis ☐ Obesity/Overweight ☐ Optimizer System ☐ Pacemaker ☐ Peripheral Vascular Disease ☐ Postpartum (6 weeks to 12 months postpartum) ☐ Prior CABG ☐ Prior MI ☐ Pulmonary Hypertension ☐ Pulmonary arterial hypertension ☐ Pulmonary hypertension due to left heart disease ☐ Pulmonary hypertension due to lung disease and/or hypoxia ☐ Chronic thromboembolic pulmonary hypertension (CTEPH) ☐ Pulmonary hypertension with unclear multifactorial mechanisms ☐ Unknown/Not Documented ☐ Prior PCI ☐ Sleep-Disordered Breathing ☐ TAVR ☐ TMVR ☐ Tricuspid Valve procedure ☐ Valvular Heart Disease ☐ Ventricular assist device ☐ Watchman Device		
☐ No Medical History			
Currently Pregnant?	☐ Yes, currently pregnant ☐ No, postpartum up to 6 weeks		
History of cigarette smoking? (In the past 12 months)	☐ Yes	☐ No	
History of vaping or e-cigarette use in the past 12 months?	☐ Yes	☐ No/ND	

Heart Failure History			
Known history of HF prior to this admission?	☐ Yes	☐ No	
Diagnosis			
Heart Failure Diagnosis	☐ Heart Failure with CAD	☐ Heart Failure, no CAD	☐ Heart Failure, Secondary Diagnosis
Atrial Fibrillation (At presentation or during hospitalization)	☐ Yes	☐ No	
Atrial Flutter (At presentation or during hospitalization)	☐ Yes	☐ No	
New Diagnosis of Diabetes	☐ Yes	☐ No	☐ Not Documented
Active bacterial or viral infection at admission or during hospitalization	☐ None ☐ Bacterial infection ☐ Emerging Infectious Disease ☐ SARS-COV-2 (COVID-19) ☐ Influenza ☐ Seasonal Cold ☐ Other viral infection		
Medications at Admission			
Medications Used Prior to Admission: <i>[Select all that apply]</i>			
☐ Patient on no meds prior to admission ☐ ACE Inhibitor - ACE Medication Name - ACE Medication Dosage - ACE Medication Frequency ☐ Angiotensin receptor blocker (ARB) - ARB Medication Name - ARB Medication Dosage - ARB Medication Frequency ☐ Angiotensin Receptor Neprilysin Inhibitor (ARNI) - ARNI Medication Name - ARNI Medication Dosage - ARNI Medication Frequency ☐ Antiarrhythmic ☐ Anticoagulation Therapy ☐ Warfarin ☐ Direct Thrombin Inhibitor ☐ Factor Xa Inhibitor ☐ Other ☐ Antiplatelet agent (excluding aspirin) ☐ Aspirin ☐ Beta-Blocker - Beta Blocker Class - Beta Blocker Medication Name - Beta Blocker Medication Dosage - Beta Blocker Medication Frequency ☐ Ca Channel Blocker ☐ Digoxin ☐ Diuretic ☐ Thiazide/Thiazide-like ☐ Loop		☐ GLP-1 Receptor Agonists - GLP-1 Medication Name - GLP-1 Dosage - GLP-1 Frequency ☐ Hydralazine ☐ Ivabradine ☐ Lipid lowering agent (Any) ☐ Statin ☐ Other Lipid lowering agent ☐ Mavacamten at Admission ☐ Mineralocorticoid Receptor Antagonist (MRA) - MRA Medication Name - MRA Medication Dosage - MRA Medication Frequency ☐ Nitrate ☐ Omega-3 fatty acid supplement ☐ Renin Inhibitor ☐ SGLT2 Inhibitor - SGLT2 Medication Name - SGLT2 Medication Dosage - SGLT2 Medication Frequency ☐ Vericiguat ☐ Other Anti-Hyperglycemic Medications: ☐ DPP-4 Inhibitors ☐ Insulin ☐ Metformin ☐ Sulfonylurea ☐ Thiazolidinedione ☐ Other Oral Agents ☐ Other injectable/subcutaneous agents ☐ Other Medications Prior to Admission	
Exams/Labs at Admission			
Height	_____ inches, cm		
Weight	_____ lbs, Kg		

Labs (Closest to Admission)	Serum Creatinine (Admission)	_____	☐ mg/dL	☐ μmol/L	☐ Not Available
	Potassium (K+) (Admission)	_____	☐ mEq/L	☐ mmol/L	☐ Not Available
	EKG QRS Duration (ms)	_____	☐ Not Available		
	+EKG QRS Morphology	☐ Normal ☐ LBBB	☐ RBBB ☐ NS-IVCD	☐ Paced ☐ Not available	
CLINICAL CODES					
ICD-10-CM Principal Diagnosis Code		_____			
IN-HOSPITAL CARE					
Procedures					
☐ No Procedures ☐ Cardiac Cath/Coronary Angiography ☐ CardioMEMs (implantable hemodynamic monitor) ☐ Coronary Artery Bypass Graft ☐ CRT-P (cardiac resynchronization therapy-pacing only) ☐ Dialysis or Ultrafiltration unspecified ☐ ICD only ☐ Mechanical Ventilation ☐ PCI ☐ Right Cardiac Catheterization ☐ TMVR ☐ Tricuspid Valve Procedure			☐ Atrial Fibrillation Ablation or Surgery ☐ Cardiac Valve Surgery ☐ Cardioversion ☐ CRT-D (cardiac resynchronization therapy with ICD) ☐ Dialysis ☐ ECMO ☐ Intra-aortic Balloon Pump ☐ Left Ventricular Assist Device ☐ Mechanical Heart Valve ☐ Pacemaker ☐ PCI with stent ☐ Stress Testing ☐ TAVR ☐ Transplant (Heart) ☐ Ultrafiltration		
EF – Quantitative	_____ %	Obtained:	☐ This Admission ☐ Within the last year ☐ > 1 year ago		
EF – Qualitative	☐ Not Applicable ☐ Normal or mild dysfunction ☐ Qualitative moderate/severe dysfunction m Performed/results not available ☐ Planned after discharge ☐ Not performed	Obtained:	☐ This Admission ☐ Within the last year ☐ > 1 year ago		
Documented LVSD?	☐ Yes	☐ No			
LVF Assessment?	☐ Yes	☐ No	☐ Not done, Reason Documented		
Was the patient ambulating by the end of hospital day 2?	☐ Yes	☐ No	☐ Not Documented		
Was DVT prophylaxis initiated by the end of hospital day 2?	☐ Yes	☐ No	☐ Contraindicated		
Influenza Vaccination	☐ Influenza vaccine was given during this hospitalization during the current flu season ☐ Influenza vaccine was received prior to admission during the current flu season, not during this hospitalization ☐ Documentation of patient's refusal of influenza vaccine ☐ Allergy/Sensitivity to influenza or if medically contraindicated ☐ Vaccine not available ☐ None of the above/Not Documented/UTD				

COVID-19 Vaccination	☐ COVID-19 vaccine was given during this hospitalization ☐ COVID-19 vaccine was received prior to admission, not during this hospitalization ☐ Documentation of patient's refusal of COVID-19 vaccine ☐ Allergy/Sensitivity to COVID-19 or if medically contraindicated ☐ Vaccine not available ☐ None of the above/Not Documented/UTD			
COVID-19 Vaccination Date	____/____/_____ ☐ Unknown			
Is there documentation that this patient was included in a COVID-19 vaccine trial?	☐ Yes ☐ No/ND			
Pneumococcal Vaccination	☐ Pneumococcal vaccine was given during this hospitalization ☐ Pneumococcal vaccine was received in the past, not during this hospitalization ☐ Documentation of patient's refusal of pneumococcal vaccine ☐ Allergy/sensitivity or if medically contraindicated to pneumococcal vaccine ☐ None of the above/Not Documented/UTD			
DISCHARGE INFORMATION				
What was the patient's discharge disposition on the day of discharge?	1 – Home 2 – Hospice – Home 3 – Hospice – Health Care Facility 4 – Acute Care Facility 5 – Other Health Care Facility		6 – Expired 7 – Left Against Medical Advice/AMA 8 – Not documented or Unable to Determine (UTD)	
If Other Health Care Facility:	☐ Skilled Nursing Facility (SNF) ☐ Inpatient Rehabilitation Facility (IRF) ☐ Long Term Care Hospital (LTCH)		☐ Intermediate Care Facility (ICF) ☐ Other	
Skilled Nursing Facility	_____			
When is the earliest physician/APN/PA documentation of comfort measures only?	☐ Day 0 or 1 ☐ Day 2 or after		☐ Timing unclear ☐ Not Documented	
Labs (Closest to Discharge)	Serum Creatinine (Discharge)	_____	☐ mg/dL	☐ µmol/L
	Potassium (K+) (Discharge)	_____	☐ mEq/L	☐ mmol/L
Discharge Medications				
ACE Prescribed?	☐ Yes ☐ No ☐ NC (None-Contraindicated)			
ACE Medication/ Dosage/ Frequency	Medication:	Dosage:	Frequency:	
Contraindications or Other Documented Reason(s) for Not Providing ACEI:	☐ Contraindicated ☐ Angioedema ☐ Hypotensive patient who was at immediate risk of cardiogenic shock ☐ Pregnancy ☐ Renal Artery Stenosis ☐ Other Contraindications ☐ Not Eligible ☐ Not Tolerant ☐ Patient Enrolled in Clinical Trial ☐ Patient Reason			

	☐ System Reason ☐ Other Reason		
ARB Prescribed?	☐ Yes ☐ No ☐ NC (None-Contraindicated)		
ARB Medication / Dosage / Frequency	Medication:	Dosage:	Frequency:
Contraindications or Other Documented Reason(s) For Not Providing ARB:	☐ Contraindicated ☐ Hypotensive patient who was at immediate risk of cardiogenic shock ☐ Pregnancy ☐ Renal Artery Stenosis ☐ Other Contraindications ☐ Not Eligible ☐ Not Tolerant ☐ Patient Enrolled in Clinical Trial ☐ Patient Reason ☐ System Reason ☐ Other Reasons		
ARNI Prescribed?	☐ Yes ☐ No ☐ NC (None-Contraindicated)		
ARNI Medication/Dosage/Frequency	Medication:	Dosage:	Frequency:
Contraindications or Other Documented Reason(s) for Not Providing ARNI at Discharge:	☐ Contraindicated ☐ Allergy ☐ Angioedema ☐ Hyperkalemia ☐ Hypotension ☐ Pregnancy ☐ Renal Artery Stenosis ☐ Renal dysfunction defined as creatinine > 2.5 mg/dL in men or > 2.0 mg/dL in women ☐ Other Contraindications ☐ Not Eligible ☐ Not Tolerant ☐ Patient Enrolled in Clinical Trial ☐ Patient Reason ☐ System Reason ☐ Other Reasons		
Reasons for not switching to ARNI at discharge:	☐ Yes ☐ No	☐ ARNI was prescribed at discharge	
If Yes,	☐ NYHA Class I ☐ NYHA Class IV		
Beta Blocker Prescribed?	☐ Yes ☐ No ☐ NC (None-Contraindicated)		
Beta Blocker Class	☐ Evidence-Based Beta Blocker ☐ Non-Evidence-Based Beta Blocker ☐ Unknown Class		
Contraindications or Other Documented Reason(s) For Not Providing Beta Blockers:	☐ Contraindicated ☐ Asthma ☐ Fluid Overload ☐ Low Blood Pressure ☐ Patient recently treated with an intravenous positive inotropic agent ☐ Other Contraindications ☐ Not Eligible ☐ Not Tolerant		

	☐ Patient Enrolled in Clinical Trial ☐ Patient Reason ☐ System Reason		
Beta Blocker Medication/Dosage/Frequency	Medication:	Dosage:	Frequency:
SGLT2 Inhibitor Prescribed?	☐ Yes ☐ No ☐ NC Medication: Dosage: Frequency:		
Contraindications or Other Documented Reason(s) For Not Providing SGLT2 Inhibitor:	☐ Contraindicated ☐ Patient currently on dialysis ☐ Ketoacidosis ☐ Known hypersensitivity to the medication ☐ Type I diabetes (not approved for use in patients with Type I diabetes due to increased risk of ketoacidosis) ☐ Other Contraindications ☐ Not Eligible ☐ Not Tolerant ☐ Patient Enrolled in Clinical Trial ☐ Patient Reason ☐ System Reason ☐ Other Reason		
GLP-1 Receptor Agonist Prescribed at Discharge?	☐ Yes ☐ No ☐ NC (None-Contraindicated)		
GLP-1 Receptor Agonist Medication/Dosage/Frequency	Medication:	Dosage:	Frequency:
Contraindications or Other Documented Reason(s) for Not Providing GLP-1 Receptor Agonist:	☐ Contraindicated ☐ Allergy ☐ Pancreatitis ☐ Medullary Thyroid Cancer ☐ Multiple Endocrine Neoplasia – Type 2 ☐ Other Contraindications ☐ Not Eligible ☐ Not Tolerant ☐ Patient Enrolled in Clinical Trial ☐ Patient Reason ☐ System Reason ☐ Other Reason		
Mineralocorticoid Receptor Antagonist (MRA) Prescribed?	☐ Yes ☐ No ☐ NC (None-Contraindicated)		
MRA Medication/Dosage/Frequency	Medication:	Dosage:	Frequency:
Was there a dose increase since prior to admission?	☐ Yes ☐ No/ND		
Potassium ordered or planned after discharge?	☐ Yes ☐ No/ND		
Renal function test scheduled	☐ Yes ☐ No/ND		
Contraindications or Other Documented Reason(s) for Not Providing Mineralocorticoid Receptor Antagonist (MRA) at Discharge	☐ Contraindicated ☐ Allergy due to MRA ☐ Hyperkalemia ☐ Renal dysfunction defined as creatinine >2.5 mg/dL in men or >2.0 mg/dL in women. ☐ Other contraindications ☐ Not Eligible ☐ Not Tolerant ☐ Patient Enrolled in Clinical Trial ☐ Patient Reason		

	☐ System Reason ☐ Other Reason				
Anticoagulation Therapy Prescribed?	☐ Yes ☐ No ☐ NC (None-Contraindicated)				
Anticoagulation Therapy Class	☐ Warfarin ☐ Direct Thrombin Inhibitor		☐ Factor Xa Inhibitor ☐ Factor Xia Inhibitor ☐ Other		
	Medication:		Dosage:	Frequency:	
Hydralazine Nitrate Prescribed?	☐ Yes ☐ No ☐ NC (None-Contraindicated)				
Other Anti-hyperglycemic Prescribed? (regardless of indication)	☐ Yes ☐ No ☐ NC				
Other Antihyperglycemic Class/Medication (regardless of indication):	Class:		Medication:		
	Class:		Medication:		
	Class:		Medication:		
Diuretics Prescribed?	☐ Yes ☐ No ☐ NC				
Diuretic Medication Class 1:	☐ Loop Diuretics ☐ Thiazide Diuretics ☐ Potassium-Sparing Diuretics Carbonic Anhydrase Inhibitors				
Diuretic Medication 1:	Medication:		Dosage:	Frequency:	
Diuretic Medication Class 2:	☐ Loop Diuretics ☐ Thiazide Diuretics ☐ Potassium-Sparing Diuretics Carbonic Anhydrase Inhibitors				
Diuretic Medication 2:	Medication:		Dosage:	Frequency:	
Clopidogrel Prescribed?	☐ Yes ☐ No ☐ NC				
Clopidogrel Dosage/Frequency	Dosage:		Frequency:		
Ivabradine Prescribed?	☐ Yes ☐ No ☐ NC				
Contraindications or Other Documented Reason(s) For Not Providing Ivabradine:	☐ Contraindicated ☐ Allergy to Ivabradine ☐ Patient 100% atrial or ventricular paced ☐ Other Contraindications ☐ Not Eligible ☐ NYHA class I or IV ☐ Not in sinus rhythm ☐ New Onset of HF ☐ Not treated with maximally tolerated dose beta blockers or beta blockers contraindicated ☐ Not Tolerant ☐ Patient Enrolled in Clinical Trial ☐ Patient Reasons ☐ System Reasons ☐ Other Medical Reasons				
Lipid Lowering Medication Prescribed?	☐ Yes ☐ No ☐ NC				
Lipid Lowering Class/Medication/Dosage/Frequency	Class:		Medication:	Dosage:	Frequency:
	Class:		Medication:	Dosage:	Frequency:
	Class:		Medication:	Dosage:	Frequency:

Omega-3 Prescribed?		☐ Yes ☐ No ☐ NC			
Other Medications					
☐ Antiarrhythmic (Discharge) ☐ Amiodarone ☐ Dofetilide ☐ Sotalol ☐ Other antiarrhythmics		☐ Ca Channel Blocker (Discharge) ☐ Digoxin (Discharge) ☐ Aficamten ☐ Mavacamten at Discharge		☐ Nitrate (Discharge) ☐ Omecamtiv ☐ Ranolazine ☐ Renin Inhibitor (Discharge) ☐ Vericiguat ☐ Other Anti-Hypertensive ☐ Other medications at discharge	
DISCHARGE INSTRUCTIONS					
Activity Level	☐ Yes	☐ No	Diet (Salt restricted)	☐ Yes	☐ No
Follow-up	☐ Yes	☐ No	Medications	☐ Yes	☐ No
Symptoms Worsening	☐ Yes	☐ No	Weight Monitoring	☐ Yes	☐ No
Follow-up Visit Scheduled	☐ Yes	☐ No	Date/Time of first follow-up visit:	__/__/__ __:__	
Location of first follow-up visit:			☐ Office Visit m Home Health Visit	☐ Telehealth m Not Documented	
Medical or Patient Reason for no follow-up appointment being scheduled?			☐ Yes	☐ No	
Follow-up Phone Call Scheduled	☐ Yes	☐ No	Date/Time of first follow-up phone call:	__/__/__ ☐ Unknown	
Follow-up appointment scheduled for diabetes management?	☐ Yes	☐ No/ND ☐ NC	Date of diabetes management follow-up visit:	__/__/__(MM/DD/YYYY) ☐ Unknown	
Other Risk Interventions					
TLC (Therapeutic Lifestyle Change) Diet	☐ Yes	☐ No	☐ Not Documented	☐ Not Applicable	
Referred to Outpatient Cardiac Rehab Program	☐ Yes	☐ No	☐ Not Documented	☐ Not Applicable	
Referral to Outpatient HF Management Program	☐ Yes	☐ No	☐ Not Documented	☐ Not Applicable	
Referral My HF Guide/AHA Heart Failure Interactive Workbook	☐ Yes	☐ No	☐ Not Documented	☐ Not Applicable	
Provision of at least 60 minutes of Heart Failure Education by a qualified educator	☐ Yes	☐ No	☐ Not Documented	☐ Not Applicable	
Is there documentation that a Palliative Care Consultation took place during this episode of care?	☐ Yes	No/ND			
Advanced Care Plan Documented Or Discussed?	☐ Yes	No/ND			
Surrogate Decision Maker Documented Or Discussed?	☐ Yes	No/ND			
Advance Directive Executed	☐ Yes		☐ No		
Post Discharge Transition					
Care Transition Record Transmitted	☐ By the seventh post-discharge day ☐ Exists, but not transmitted by the seventh post-discharge day ☐ No Care Transition Record/UTD				
Care Transition Record Includes	☐ All were included (<i>Check all yes</i>)				
	Discharge Medications			☐ Yes	☐ No
	Follow-up Treatment(s) and Service(s) Needed			☐ Yes	☐ No
	Procedures Performed During Hospitalization			☐ Yes	☐ No

	Reason for Hospitalization	☐ Yes	☐ No
	Treatment(s)/Service(s) Provided	☐ Yes	☐ No
Health Related Social Needs Assessment			
During this admission, was a standardized health related social needs form or assessment completed?	☐ Yes	☐ No/ND	
If yes, identify the areas of unmet social need. (select all that apply):	☐ None ☐ Education ☐ Employment ☐ Financial Strain ☐ Food ☐ Living Situation/Housing	☐ Mental Health ☐ Personal Safety ☐ Substance Use Disorder ☐ Transportation Barriers ☐ Utilities	