American Heart Association.
Get With The Guidelines.
Heart Failure

GWTG-HF Case Record Full Form (CRF) **June 2023**

<table>
<tr><td colspan="4">Patient ID:</td></tr>
<tr><td colspan="4">DEMOGRAPHICS TAB</td></tr>
<tr><td colspan="4">Demographics</td></tr>
<tr><td>Sex</td><td colspan="3">○ Male ○ Female ○ Unknown</td></tr>
<tr><td>Patient Gender Identity</td><td colspan="3">○ 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.</td></tr>
<tr><td>Patient-Identified Sexual Orientation</td><td colspan="3">○ Straight or heterosexual
○ Lesbian or gay
○ Queer, pansexual, and/or questioning
○ Something else; please specify. ______________.
○ Don't know
○ Declined to answer</td></tr>
<tr><td>Date of Birth</td><td>__/__/____ (MM/DD/YYYY)</td><td>Patient Postal Code</td><td>________ - _______</td></tr>
<tr><td>Payment Source</td><td colspan="3">❑ Medicare Title 18
❑ Medicaid Title 19
❑ Medicare – Private/HMO/PPO/Other
❑ Medicaid – Private/HMO/PPO/Other
❑ Private/HMO/PPO/Other
❑ VA/CHAMPVA/Tricare
❑ Self-pay/No Insurance
❑ Other/Not Documented/UTD</td></tr>
<tr><td>External Tracking ID</td><td colspan="3">________________</td></tr>
<tr><td colspan="4">Race and Ethnicity</td></tr>
<tr><td>Race</td><td colspan="3">❑ American Indian or Alaska Native
❑ Asian
  ❑ Asian Indian
  ❑ Chinese
  ❑ Filipino
  ❑ Japanese
  ❑ Korean
  ❑ Vietnamese
  ❑ Other Asian
❑ Black or African American
❑ Native Hawaiian or Pacific Islander
  ❑ Native Hawaiian
  ❑ Guamanian or Chamorro
  ❑ Samoan
  ❑ Other Pacific Islander
❑ White
❑ UTD</td></tr>
<tr><td>Hispanic Ethnicity</td><td colspan="3">○ Yes ○ No/UTD</td></tr>
<tr><td>Select Hispanic Origin Group(s):</td><td colspan="3">❑ Mexican, Mexican American, Chicano/a
❑ Cuban
❑ Puerto Rican
❑ Another Hispanic, Latino, or Spanish Origin</td></tr>
<tr><td colspan="4">ADMISSIONS TAB</td></tr>
<tr><td colspan="4">Arrival and Admission</td></tr>
<tr><td>Internal Tracking ID</td><td>________________</td><td>Physician/Provider NPI</td><td>________________</td></tr>
</table>

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Arrival Date/Time	___/___/_____ ___: ____	Admission Date	___/___/_____
Transferred in (from another ED?)	❍ Yes	❍ No	
Point of Origin for Admission or Visit	❍ 1. Non-Healthcare Facility Point of Origin ❍ 2. Clinic ❍ 4. Transfer from a Hospital (Different Facility) ❍ 5. Transfer from a Skilled Nursing Facility (SNF) or Intermediate Care Facility (ICF)	❍ 6. Transfer from another Health Care Facility ❍ 7. Emergency Room ❍ 9. Information not available ❍ F. Transfer from Hospice and is Under a Hospice Plan of Care or 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
- ❑ CAD
- ❑ CardioMEMs (implantable hemodynamic monitor)
- ❑ COPD or Asthma
- ❑ CRT-D (cardiac resynchronization therapy with ICD)
- ❑ CRT-P (cardiac resynchronization therapy-pacing only)
- ❑ CVA/TIA
- ❑ Depression
- ❑ Diabetes
- ❑ Dialysis (chronic)
- ❑ Emerging Infectious Disease
  - ❍ MERS
  - ❍ SARS-COV-1
  - ❍ SARS-COV-2 (COVID-19)
  - ❍ Other infectious respiratory pathogen
- ❑ Familial hypercholesterolemia
- ❑ No Medical History
- ❑ Heart failure
- ❑ Heart Transplant
- ❑ Hyperlipidemia
- ❑ Hypertension
- ❑ ICD only
- ❑ Kidney Transplant
- ❑ Left Ventricular Assist Device
- ❑ Pacemaker
- ❑ Peripheral Vascular Disease
- ❑ Prior CABG
- ❑ Prior MI
- ❑ Prior PCI
- ❑ Renal insufficiency - chronic (SCr>2.0)
- ❑ Sleep-Disordered Breathing
- ❑ TAVR
- ❑ TMVR
- ❑ Tricuspid Valve procedure
- ❑ Valvular Heart Disease
- ❑ Ventricular assist device

Diabetes Type:	❍ Type 1 ❍ Type 2 ❍ ND
Diabetes Duration:	❍ <5 years ❍ 5 - <10 years ❍ 10 - <20 years ❍ >=20 years ❍ Unknown
Sleep-Disordered Breathing Type:	❑ Obstructive ❑ Central ❑ Mixed ❑ Unknown/Not Documented

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Equipment used at home:	❑ O2 ❑ CPAP ❑ BiPAP ❑ Adaptive Servo-Ventilation ❑ None ❑ Unknown/Not Documented
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 Etiology: Check if history of:	❑ Ischemic/CAD ❑ Non-Ischemic ❑ Alcohol/Other Drug ❑ Chemotherapy ❑ Familial ❑ Hypertensive ❑ Postpartum ❑ Viral ❑ Other Etiology ❑ Unknown Etiology
Known history of HF prior to this admission?	❍ Yes ❍ No
# of hospital admissions in past 6 mo. for HF:	❍ 0 ❍ 1 ❍ 2 ❍ >2 ❍ Unknown
❑ Patient Listed for Transplant	
DIAGNOSIS	
Heart Failure Diagnosis	❍ Heart Failure, primary diagnosis, with CAD ❍ Heart Failure, primary diagnosis, no CAD ❍ Heart Failure, secondary diagnosis
Atrial Fibrillation (At presentation or during hospitalization)	❍ Yes ❍ No ❑ Documented New Onset?
Atrial Flutter (At presentation or during hospitalization)	❍ Yes ❍ No ❑ Documented New Onset?
New Diagnosis of Diabetes	❍ Yes ❍ No ❍ Not Documented
Basis for Diagnosis	❑ HbA1c ❑ Fasting Blood Sugar ❑ Oral Glucose Tolerance ❑ Test Other
Characterization of HF at admission or when first recognized	❍ Acute Pulmonary Edema ❍ Dizziness/Syncope ❍ Dyspnea ❍ ICD Shock/Sustained Ventricular Arrhythmia ❍ Pulmonary Congestion ❍ Volume overload/Weight Gain ❍ Worsening fatigue ❍ Other
Other Conditions Contributing to HF Exacerbation *Select all that apply*	❑ Arrhythmia ❑ Pneumonia/respiratory process ❑ Noncompliance - medication ❑ Worsening Renal Failure ❑ Ischemia/ACS ❑ Uncontrolled HTN ❑ Noncompliance – dietary ❑ Other
Active bacterial or viral infection at admission or during hospitalization	❍ None/ND ❍ Bacterial infection ❍ Emerging Infectious Disease ❑ SARS-COV-1 ❑ SARS-COV-2 (COVID-19) ❑ MERS ❑ Other infectious respiratory pathogen ❍ Influenza ❍ Seasonal Cold ❍ Other Viral Infection

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New Diagnosis of ATTR-CM	❍ Yes ❍ Hereditary ❍ Wild-Type ❍ Unknown/Not Documented ❍ No ❍ Not Documented

MEDICATIONS AT ADMISSION

Medications Used Prior to Admission: *[Select all that apply]*

- ❑ Patient on no meds prior to admission
- ❑ ACE Inhibitor
- ❑ Angiotensin receptor blocker (ARB)
- ❑ Angiotensin Receptor Neprilysin Inhibitor (ARNI)
- ❑ Antiarrhythmic
- ❑ Anticoagulation Therapy
  - ❍ Warfarin
  - ❍ Direct Thrombin Inhibitor
  - ❍ Factor Xa Inhibitor
  - ❍ Other
- ❑ Anti-hyperglycemic medications:
  - ❑ DPP-4 Inhibitors
  - ❑ GLP-1 receptor agonist
  - ❑ Insulin
  - ❑ Mavacamten
  - ❑ Metformin
  - ❑ Mineralocorticoid Receptor Antagonist (MRA)
  - ❑ Sulfonylurea
  - ❑ Thiazolidinedione
  - ❑ Other Oral Agents
  - ❑ Other injectable/subcutaneous agents
- ❑ Antiplatelet agent (excluding aspirin)
- ❑ Aspirin
- ❑ Beta-Blocker
- ❑ Ca channel blocker
  - ❑ Other injectable/subcutaneous agents
- ❑ Digoxin
- ❑ Diuretic
  - ❍ Thiazide/Thiazide-like
  - ❍ Loop
- ❑ Hydralazine
- ❑ Ivabradine
- ❑ Finerenone
- ❑ Lipid lowering agent (Any)
  - ❍ Statin
  - ❍ Other Lipid lowering agent
- ❑ Nitrate
- ❑ Omega-3 fatty acid supplement
- ❑ Renin Inhibitor
- ❑ SGLT2 Inhibitor
- ❑ Vericiguat
- ❑ Other Medications Prior to Admission

Symptoms (Closest to Admission) *Select all that apply*	❍ Chest Pain ❍ Orthopnea ❍ Palpitations	❍ Dyspnea at rest ❍ Fatigue ❍ PND	❍ Dyspnea on Exertion ❍ Decreased appetite/early satiety ❍ Dizziness/lightheadedness/syncope

EXAMS/LABS AT ADMISSION

Height	____________ ❍ inches ❍ cm			❍ Height ND
Weight	____________ ❍Lbs. ❍Kgs.			❍ Weight ND
Waist Circumference	____________ ❍ inches ❍ cm			❍ Waist Circumference ND
BMI	____________ (Automatically Calculated)			
Systolic	________			
Diastolic	________			
	❍ BP ND			
Respiratory Rate (breaths per minute)	________			
JVP (cm):	❍ Yes ❍ No	❍ Unknown	JVP Value ________	
Rales:	❍ Yes ❍ No	❍ Unknown	Rales Value ________	❍ <1/3 ❍ ≥1/3 ❍ N/A

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Lower Extremity Edema	❍ Yes ❍ No	❍ Unknown	Lower Extremity Value	❍ Trace ❍ 1+ ❍ 2+ ❍ 3+ ❍ 4+ ❍ N/A	
Lipids	TC: _____ mg/dL	HDL: _____ mg/dL	LDL: _____ mg/dL	TG: _____ mg/dL	❑ Lipids Not Available

Labs (Closet to Admission)					
Sodium (Na+)	_____	❍ mEq/L	❍ mmol/L	❍ mg/dL	❑ Not Available
Hgb	_____	❍ g/dL	❍ g/L		❑ Not Available
Albumin	_____	❍ g/dL	❍ g/L		❑ Not Available
BNP	_____	❍ pg/mL	❍ pmol/L	❍ ng/L	❑ Not Available
NT-proBNP	_____	❍ pg/mL	❍ ng/L		❑ Not Available
Serum Creatinine	_____	❍ mg/dL	❍ µmol/L		❑ Not Available
BUN	_____	❍ mg/dL	❍ µmol/L		❑ Not Available
Troponin (Peak)	__________ ❍ng/mL ❍ug/L	❍ T ❍ I ❍ hs-I ❍ hs-T	❍ Normal ❍ Abnormal		❑ Not Available
Potassium (K+)	_____	❍ mEq/L	❍ mmol/L	❑ Not Available	
Ferritin (ng/mL)	__________				
HbA1C	__________ %	❑ Not Available			
Fasting Blood Glucose (mg/dL)	______________	❑ 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	_________		
ICD-10-CM Other Diagnoses Codes	1.	2.	3.
	4.	5.	6.
	7.	8.	9
	10.	11.	12.
ICD-10-PCS Principal Procedure Code	________	Date: __/__/____	❍ Date UTD
ICD-10-PCS Other Principal Procedure Codes	1.	Date: __/__/____	❍ Date UTD
	2.	Date: __/__/____	❍ Date UTD
	3.	Date: __/__/____	❍ Date UTD
	4.	Date: __/__/____	❍ Date UTD
	5.	Date: __/__/____	❍ Date UTD

IN-HOSPITAL

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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 ❑ 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 ❍ Performed/results not available ❍ Planned after discharge ❍ Not performed	Obtained:	❍ This Admission ❍ Within the last year ❍ > 1 year ago
Mitral Valve Regurgitation (MR) on echocardiogram	❍ Not applicable ❍ None ❍ Trace/trivial ❍ 1+ or Mild ❍ 2+ or Moderate ❍ 3+ or Moderate to Severe ❍ 4+ or Severe		
Documented LVSD?	❍ Yes	❍ No	
LVF Assessment?	❍ Yes	❍ No	❍ Not done, Reason Documented
Oral Medications during hospitalization *Select all that apply*	❑ None ❑ ARNI ❑ ARB	❑ Hydralazine Nitrate ❑ Mineralocorticoid Receptor Antagonist (MRA)	❑ ACE Inhibitor ❑ Beta Blocker ❑ SGLT2 Inhibitor
IV Iron	❑ Yes	❑ No	❑ Not documented

Parenteral Therapies during hospitalization
Select all that apply
	❑ None
	❑ Dopamine
	❑ Dobutamine
	❑ Iron
		❑ Loop Diuretics
		  ❑ Continuous Infusion
		  ❑ Intermittent bolus
		❑ Milrinone
		❑ Nesiritide Nitroglycerine
		❑ Other IV Vasodilator
		❑ Vasopressin antagonist	

Was the patient ambulating at the end of hospital day 2?	❍ Yes	❍ No	❍ Not Documented
Was DVT prophylaxis initiated by the end of hospital day 2?	❍ Yes	❍ No/Not Documented	❍ Contraindicated
DVT prophylaxis type	❑ Low dose unfractionated heparin (LDUH) ❑ Low molecular weight heparin (LMWH) ❑ Warfarin	❑ Factor Xa Inhibitor ❑ Direct thrombin inhibitor ❑ Venous foot pumps (VFP) ❑ Intermittent pneumatic compression devices (IPC)	

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	❑ Other
Was DVT or PE (pulmonary embolus) documented?	❍ Yes ❍ No/Not Documented
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 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	________________ ❑ ND		
If Home, special discharge circumstances:	❍ Home Health Care ❍ Homeless	❍ International ❍ Prison/Incarcerated	❍ None/UTD
Primary Cause of Death	❍ Cardiovascular	❍ Non-Cardiovascular	❍ Unknown
If Cardiovascular:	❍ Acute Coronary Syndrome	❍ Worsening Heart Failure	❍ Sudden Death ❍ Other
When is the earliest physician/APN/PA documentation of comfort measures only?		❍ Day 0 or 1 ❍ Day 2 or after	❍ Timing unclear ❍ Not Documented
Symptoms (closest to discharge)	❍ Worse ❍ Unchanged	❍ Better, Symptomatic ❍ Better, Asymptomatic	❍ Unable to determine

Vital Signs (closest to Discharge)	Weight	________ ❍ Lbs. ❍ Kgs.	❍ Not Documented
	Heart Rate (bpm)	________	❍ Not Documented
	Systolic		❍ Not Documented

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	Diastolic					
Exam (Closest to Discharge)	JVP:	❍ Yes	❍ No	❍ Unknown	If Yes, ______ cm	
	Rales:	❍ Yes ❍ No	❍ Unknown	If Yes,	❍ <1/3	❍ ≥1/3
					❍ N/A	
	Lower Extremity Edema	❍ Yes ❍ No	❍ Unknown	If Yes,	❍ Trace ❍ 1+	❍ 2+ ❍ 3+
					❍ 4+ ❍ N/A	
Labs (Closest to Discharge)	Sodium (Na+)	___________	❍ mEq/L	❍ mmol/L	❍ mg/dL	❑ Unavailable
	BNP	___________	❍ pg/mL	❍ pmol/L	❍ ng/L	❑ Unavailable
	Serum Creatinine	___________________	❍ mg/dL	❍ µmol/L	❑ Unavailable	
	BUN	___________________	❍ mg/dL	❍ µmol/L	❑ Unavailable	
	eGFR (mL/min)					
	NT-proBNP (pg/mL)	___________________________	❑ Not Documented			
	Potassium (K+)	________	❍ mEq/L	❍ mmol/L	❍ mg/dL	❑ Unavailable
	Urinary Albumin (mg/dL)					
	Urinary Creatinine (mg/dL)					
	Urinary Albumin-to-Creatinine Ratio (UACR) (mg/g)					
	Ferritin (mg/mL)	_______	❑ Unavailable			

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 ❑ Hypotensive patient who was at immediate risk of cardiogenic shock ❑ Hospitalized patient who experienced marked azotemia ❑ 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		

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<table>
<tr><td></td><td colspan="3">❑ Hospitalized patient who experienced marked azotemia
❑ Other Contraindications
❑ Not Eligible
❑ Not Tolerant
❑ Patient Enrolled in Clinical Trial
❑ Patient Reason
❑ System Reason
❑ Other Reasons</td></tr>
<tr><td>ARNI Prescribed?</td><td colspan="3">❍ Yes ❍ No ❍ NC (None-Contraindicated)</td></tr>
<tr><td>ARNI Medication/Dosage/Frequency</td><td>Medication:</td><td>Dosage:</td><td>Frequency:</td></tr>
<tr><td>Contraindications or Other Documented Reason(s) for Not Providing ARNI at Discharge:</td><td colspan="3">❑ Contraindicated
❑ ACE inhibitor use within the prior 36 hours
❑ Allergy
❑ Hyperkalemia
❑ Hypotension
❑ 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</td></tr>
<tr><td>Reasons for not switching to ARNI at discharge:</td><td>❍ Yes
❍ No</td><td colspan="2">❍ ARNI was prescribed at discharge</td></tr>
<tr><td>If Yes,</td><td colspan="3">❍ NYHA Class I
❍ NYHA Class IV</td></tr>
<tr><td>Beta Blocker Prescribed?</td><td colspan="3">❍ Yes ❍ No ❍ NC (None-Contraindicated)</td></tr>
<tr><td>Beta Blocker Class</td><td colspan="3">❍ Evidence-Based Beta Blocker
❍ Non-Evidence-Based Beta Blocker
❍ Unknown Class</td></tr>
<tr><td>Contraindications or Other Documented Reason(s) For Not Providing Beta Blockers:</td><td colspan="3">❑ 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</td></tr>
<tr><td>Beta Blocker Medication/Dosage/Frequency</td><td>Medication:</td><td>Dosage:</td><td>Frequency:</td></tr>
<tr><td rowspan="2">SGLT2 Inhibitor Prescribed?</td><td colspan="3">❍ Yes ❍ No ❍ NC</td></tr>
<tr><td>Medication:</td><td>Dosage:</td><td>Frequency:</td></tr>
<tr><td>Contraindications or Other Documented Reason(s) For Not Providing SGLT2 Inhibitor:</td><td colspan="3">❑ 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</td></tr>
</table>

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	❑ 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 ❑ Other	
	Medication:	Dosage:	Frequency:
Anticoagulation Contraindication(s):	❑ Contraindicated ❑ Allergy to or complication r/t anticoagulation therapy (hx or current) ❑ Risk for bleeding or discontinued due to bleeding ❑ Serious side effect to medication ❑ Terminal illness/Comfort Measures Only ❑ Other Contraindications ❑ Not Eligible ❑ Not Tolerant ❑ Patient Enrolled in Clinical Trial ❑ Patient Reason ❑ System Reason ❑ Other		
Hydralazine Nitrate Prescribed?	❍ Yes ❍ No ❍ NC (None-Contraindicated)		
Contraindications or Other Documented Reason(s) For Not Providing Hydralazine Nitrate:	❑ Contraindicated ❑ Not Eligible ❑ Not Tolerant ❑ Patient Enrolled in Clinical Trial ❑ Patient Reason ❑ System Reason ❑ Other Reasons		
Anti-hyperglycemic Prescribed?	❍ Yes ❍ No ❍ NC		

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Antihyperglycemic Class/Medication	Class:		Medication:	
	Class:		Medication:	
	Class:		Medication:	
ASA Prescribed?	❍ Yes ❍ No ❍ NC (None-Contraindicated)			
ASA Medication/Dosage/Frequency	Medication:		Dosage:	Frequency:
Other Antiplatelets Prescribed?	❍ Yes ❍ No ❍ NC (None-Contraindicated)			
Other Antiplatelets Medication/Dosage/Frequency	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) ❑ Diuretic (Discharge) ❑ Loop Diuretic ❑ Thiazide Diuretic ❑ Mavacamten		❑ Nitrate (Discharge) ❑ Omecamtiv ❑ Ranolazine ❑ Renin Inhibitor (Discharge) ❑ Vericiguat ❑ Other Anti-Hypertensive ❑ Other medications at discharge	

Other Therapies		
ICD Counseling?	❍ Yes	❍ No
Reason for not counseling	❍ Yes	❍ No
Documented Medical Reason(s) for Not Counseling?	❑ ICD or CRT-D device in patient ❑ Multiple or significant comorbidities	❑ Limited Life Expectancy ❑ other reasons not eligible for ICD (e.g. EF>35%, new onset HF) ❑ Other reasons for not counseling
ICD Placed or Prescribed?	❍ Yes	❍ No

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Reason(s) for Not Placing or Prescribing?	❍ Yes	❍ No
Documented Reason(s) for Not Placing or Prescribing ICD Therapy?	❑ Contraindications ❑ Not receiving optimal medical therapy ❑ Patient Reason ❑ System Reason	❑ Any other physician documented reason including AMI in prior 40 days, recent revascularization, recent onset HF

CRT-D Placed or Prescribed?	❍ Yes	❍ No
CRT-P Placed or Prescribed?	❍ Yes	❍ No
Reason for not Placing or Prescribing?	❍ Yes	❍ No

Documented Reason(s) for Not Placing or Prescribing CRT Therapy?	❑ Contraindications ❑ Not receiving optimal medical therapy ❑ Not NYHA functional Class III or ambulatory Class IV ❑ Patient Reason	❑ Any other physician documented reason including AMI in prior 40 days, recent revascularization, recent onset of HF ❑ System Reason
Risk Interventions		
Smoking Cessation Counseling Given	❍ Yes	❍ No
Smoking Cessation Therapies Prescribed (select all that apply)	❑ Treatment Not Specified ❑ Counseling Only ❑ Over the Counter Nicotine Replacement Therapy	❑ Prescription Medications ❑ Other

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 ❍ Home Health Visit	❍ Telehealth ❍ 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:	__/__/____	
Follow-up appointment scheduled for diabetes management?	❍ Yes	❍ No	Date of diabetes management follow-up visit:	__/__/____	

Other Risk Interventions				
TLC (Therapeutic Lifestyle Change) Diet	❍ Yes	❍ No	❍ Not Documented	❍ Not Applicable
Obesity Weight Management	❍ Yes	❍ No	❍ Not Documented	❍ Not Applicable
Activity Level/Recommendation	❍ Yes	❍ No	❍ Not Documented	❍ Not Applicable
Referred to Outpatient Cardiac Rehab Program	❍ Yes	❍ No	❍ Not Documented	❍ Not Applicable
Anticoagulation Therapy Education	❍ Yes	❍ No	❍ Not Documented	❍ Not Applicable
Was Diabetes Teaching provided?	❍ Yes	❍ No	❍ Not Documented	❍ Not Applicable
PT/INR Planned Follow-Up	❍ Yes	❍ No	❍ Not Documented	❍ Not Applicable
Referral to Sleep Study	❍ Yes	❍ No	❍ Not Documented	❍ Not Applicable
Referral to Outpatient HF Management Program	❍ Yes	❍ No	❍ Not Documented	❍ Not Applicable
Outpatient HF Management Program Type(s):	❑Telemanagement	❑ Home Visit	❑ Clinic-based	

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Referral to AHA My HF Guide/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
Advanced Care Plan/Surrogate Decision Maker Documented Or Discussed?	❍ Yes ❍ No ❍ Not Documented ❍ Not Applicable
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 Transmitted Includes	❑ All were included *(Check all yes)*		
	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
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 of the areas of unmet social need listed ❑ Education ❑ Employment ❑ Financial Strain ❑ Food ❑ Living Situation/Housing ❑ Mental Health ❑ Personal Safety ❑ Substance Abuse ❑ Transportation Barriers ❑ Utilities		

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