State of GWTG-Heart Failure 2018 Tuesday February 13, 2018



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Heart.org/QualityHI

Our Presenters





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Northwestern Medicine

American Heart Association Get With The Guidelines- HF

"Guideline Directed Care Algorithms"

> Clyde W. Yancy, MD, MSc Professor of Medicine, Professor, Medical Social Science Chief, Cardiology Associate Director, Bluhm CV Institute & Vice-Dean, Diversity & Inclusion Northwestern University, FSM & Deputy Editor, JAMA Cardiology

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No relevant disclosures

Heart Failure Awareness Week 2018





New ACC/AHA/HFSA Guidelines

Yancy et al 2017 ACC/AHA/HFSA Heart Failure Focused Update

2017 ACC/AHA/HFSA Focused Update of the 2013 ACCF/AHA Guideline for the Management of Heart Failure

A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines and the Heart Failure Society of America

Developed in Collaboration With the American Academy of Family Physicians, American College of Chest Physicians, and International Society for Heart and Lung Transplantation

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Michael R. Bristow et al. JCHF 2017;5:772-781

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Stages, Phenotypes and Treatment of HF

At Risk for Heart Failure Heart Failure STAGE A STAGE B STAGE C STAGE D At high risk for HF but Structural heart disease Structural heart disease without structural heart but without signs or with prior or current Refractory HF disease or symptoms of HF symptoms of HF symptoms of HF e.g., Patients with: HTN Atherosclerotic disease e.g., Patients with: • DM e.g., Patients with: Refractory Previous MI e.g., Patients with: · Marked HF symptoms at Obesity Development of symptoms of HF • LV remodeling including Structural heart Known structural heart disease and Metabolic syndrome symptoms of HF at rest, despite rest disease LVH and low EF HF signs and symptoms GDMT or Recurrent hospitalizations Asymptomatic valvular Patients despite GDMT disease Using cardiotoxins · With family history of cardiomyopathy HF*p*EF HE/EF THERAPY THERAPY THERAPY THERAPY THERAPY Goals Goals Goals Goals Goals Control symptoms Control symptoms Improve HRQOL · Heart healthy lifestyle · Prevent HF symptoms Control symptoms Patient education Prevent vascular, Prevent further cardiac Improve HRQOL Prevent hospitalization Reduce hospital coronary disease remodeling Prevent hospitalization Prevent mortality readmissions Establish patient's end- Prevent LV structural Prevent mortality Drugs for routine use • Diuretics for fluid retention Drugs of-life goals abnormalities ACEI or ARB as Strategies ACEL or ARB. Options appropriate Druas Identification of comorbidities. · Beta blockers Advanced care Beta blockers as ACEL or ARB in Aldosterone antagonists measures appropriate Heart transplant appropriate patients for Treatment Drugs for use in selected patients Chronic inotropes vascular disease or DM Diuresis to relieve symptoms In selected patients Hydralazine/isosorbide dinitrate Temporary or permanent Statins as appropriate of congestion MCS ICD ACEL and ARB. Follow guideline driven Digoxin Experimental surgery or Revascularization or indications for comorbidities, drugs Palliative care and valvular surgerv as In selected patients e.g., HTN, AF, CAD, DM appropriate CRT hospice Revascularization or valvular • ICD ICD deactivation

surgery as appropriate



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Yancy C, et al. JACC, 2013

 Revascularization or valvular surgery as appropriate











Biomarkers Indications for Use



*Other biomarkers of injury or fibrosis include soluble ST2 receptor, galectin-3, and high-sensitivity troponin. ACC indicates American College of Cardiology; AHA, American Heart Association; ADHF, acute decompensated heart failure; BNP, B-type natriuretic peptide; COR, Class of Recommendation; ED, emergency department; HF, heart failure; NT-proBNP, Nterminal pro-B-type natriuretic peptide; NYHA, New York Heart Association; and pts, patients.







Blood Pressure (BP) Thresholds and Recommendations for Treatment and Follow-Up (continued on next slide)







Hypertension

Treating Hypertension to Reduce the Incidence of HF

COR	LOE	Recommendations	Comment/ Rationale
I	B-R	In patients at increased risk, stage A HF, the optimal blood pressure in those with hypertension should be less than 130/80 mm Hg.	NEW: Recommendation reflects new RCT data.





Simplified Schematic of the Renin– Angiotensin–Aldosterone System



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von Lueder T G et al. Circ Heart Fail. 2013;6:594-605

Northwestern Medicine*

Simplified Schematic of the Natriuretic Peptide System (NPS)



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Medicine' von Lueder T G et al. Circ Heart Fail. 2013;6:594-605

Northwestern

Pharmacological Treatment for Stage C HF With Reduced EF

Renin-Angiotensin System Inhibition With ACE-Inhibitor or ARB or ARNI

COR	LOE	Recommendations	Comment/ Rationale
I	ARNI: B-R	In patients with chronic symptomatic HF <i>r</i> EF NYHA class II or III who tolerate an ACE inhibitor or ARB, replacement by an ARNI is recommended to further reduce morbidity and mortality.	NEW: New clinical trial data necessitated this recommendation.





Pharmacological Treatment for Stage C HF With Reduced EF

Renin-Angiotensin System Inhibition With ACE-Inhibitor or ARB or ARNI

COR	LOE	Recommendations	Comment/ Rationale
III: Harm	B-R	ARNI should not be administered concomitantly with ACE inhibitors or within 36 hours of the last dose of an ACE inhibitor.	NEW: Available evidence demonstrates a potential signal of harm for a concomitant use of ACE inhibitors and ARNI.
III: Harm	C-EO	ARNI should not be administered to patients with a history of angioedema.	NEW: New clinical trial data.





Pharmacological Treatment for Stage C HF With Reduced EF

Ivabradine

COR	LOE	Recommendations	Comment/ Rationale
lla	B-R	Ivabradine can be beneficial to reduce HF hospitalization for patients with symptomatic (NYHA class II-III) stable chronic HF <i>r</i> EF (LVEF ≤35%) who are receiving GDEM*, including a beta blocker at maximum tolerated dose, and who are in sinus rhythm with a heart rate of 70 bpm or greater at rest.	NEW: New clinical trial data.

*In other parts of the document, the term "GDMT" has been used to denote guideline-directed management and therapy. In this recommendation, however, the term "GDEM" has been used to denote this same concept in order to reflect the original wording of the recommendation that initially appeared in the "2016 ACC/AHA/HFSA Focused Update on New Pharmacological Therapy for Heart Failure: An Update of the 2013 ACCF/AHA Guideline for the Management of Heart Failure". *Helping Cardiovascular Professionals*







JAMA

From: Table. Demonstrated Benefits of Evidence-Based Therapies for Patients Inhibitor Thera With Heart Failure and Reduced Ejection Fraction

Evidence-Based Therapy	Relative Risk Reduction in All-Cause Mortality in Pivotal Randomized Clinical Trial(s), %	NNT to Prevent All-Cause Mortality Over Time	NNT for All-Cause Mortality ^a
ACEI/ARB	17	22 over 42 mo	77
ARNI ^b	16	36 over 27 mo	80
β-Blocker	34	28 over 12 mo	28
Aldosterone antagonist	30	9 over 24 mo	18
Hydralazine/ nitrate	43	25 over 10 mo	21
CRT	36	12 over 24 mo	24
ICD	23	14 over 60 mo	70

Abbreviations: ACEI, angiotensin-converting enzyme inhibitor;

ARB, angiotensin receptor blocker; ARNI, angiotensin receptor neprilysin inhibitor; CRT cardiac resynchronization therapy; ICD, implantable cardioverter defibrillator, NNT, number needed to treat.

^a Standardized to 12 months.

^{Demo b} Benefit of ARNI therapy incremental to that achieved with ACEI therapy. For

- the other medications shown, the benefits are based on comparisons to placebo control.
- place







Treatment of HFrEF Stage C and D



Continue GDMT with serial reassessment & optimized dosing/adherence

†Hydral-Nitrates green box: The combination of ISDN/HYD with ARNI has not been robustly tested. BP response should be carefully monitored. ‡See 2013 HF guideline.

§Participation in investigational studies is also appropriate for stage C, NYHA class II and III HF.

ACEI indicates angiotensin-converting enzyme inhibitor; ARB, angiotensin receptor-blocker; ARNI, angiotensin receptor-neprilysin inhibitor; BP, blood pressure; bpm, beats per minute; C/I, contraindication; COR, Class of Recommendation; CrCI, creatinine clearance; CRT-D, cardiac resynchronization therapy-device; Dx, diagnosis; GDMT, guideline-directed management and therapy; HF, heart failure; HFrEF, heart failure with reduced ejection fraction; ICD, implantable cardioverter-defibrillator; ISDN/HYD, isosorbide dinitrate hydral-nitrates; K+, potassium; LBBB, left bundle-branch block; LVAD, left ventricular assist device; LVEF, left ventricular ejection fraction; MI, myocardial infarction; NSR, normal sinus rhythm; and NYHA, New York Heart Association.







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🥕 PDF Article

2017 ACC Expert Consensus Decision Pathway for Optimization of Heart Failure Treatment: Answers to 10 Pivotal Issues About Heart Failure With Reduced Ejection Fraction A Report of the American College of Cardiology Task Force on Expert Consensus Decision Pathways

Clyde W. Yancy, James L. Januzzi Jr., Larry A. Allen, Javed Butler, Leslie L. Davis, Gregg C. Fonarow, Nasrien E. Ibrahim, Mariell Jessup, JoAnn Lindenfeld, Thomas M. Maddox, Frederick A. Masoudi, Shweta R. Motiwala, J. Herbert Patterson, Mary Norine Walsh and Alan Wasserman



10 Principles for Successful Treatment of Heart Failure

How to implement How to address GDMT... challenges with... I. Initiate & Switch III. Referral Treatment algorithm for Triggers for referral to guideline-directed medical HF specialist (Table 6) therapy including novel (Table 15) IV. Care Coordination therapies (Figure 2 and 3) Essential skills for a II. Titration HF team (Table 7) Target doses of select Infrastructure for team-based guideline-directed heart failure HF care (Table 8) therapy (Tables 1, 2, 3, 4, 5) V. Adherence Considerations for monitoring Causes of non-adherence (Table 9) Interventions for adherence (Table 10, 11) VI. Specific Patient Cohorts Evidence based recommendations and assessment of risk for special cohorts: African Americans: older adults: frail (Table 12)

VII. Cost of Care Strategies to reduce cost (Table 13)

Helpful information for completion of prior authorization forms (Table 14)

How to manage...

VIII. Increasing Complexity Ten pathophysiologic targets in HFrEF and treatments

Ten principles and actions to guide optimal therapy

IX. Comorbidities

Common cardiac and non-cardiac comorbidities with suggested actions (Table 16)

X. Palliative/Hospice Care Seven principles and actions to consider regarding palliative care



Clvde W. Yancv et al. JACC 2018:71:201-230 Helping Cardiovascular Professionals Learn. Advance. Heal. ©2018 by American College of Cardiology





Guidelines Failure 2017 ACCF/AHA Heart



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New Guideline Takeaway messages:

- New effective medical therapies have now been fully incorporated in evidence based guideline directed treatment algorithms
- There is an increasing complexity in the treatment of HFrEF; this will require careful assessment of the clinical context/scenario
- Powerful new data should drive the PREVENTION of heart failure
- Avoiding entry into the "HF Club" is the best therapeutic approach







GWTG-HF Update and Reducing Readmissions Safely

Gregg C. Fonarow, MD FACC, FAHA, FHFSA Eliot Corday Chair of Cardiovascular Medicine and Science Co-Chief UCLA Division of Cardiology Director, Ahmanson-UCLA Cardiomyopathy Center, Los Angeles, CA

GWTG-HF Hospital Participation

GWTG-Heart Failure Enrolled Hospitals Data through Dec. 2017



Column1

GWTG-HF: Hospitalization Episodes Entered



Number of records

ACEI/ARB or ARNI at Discharge* Percent of heart failure patients with left ventricular systolic dysfunction (LVSD) and without angiotensin converting enzyme inhibitor (ACEI) and angiotensin receptor blocker (ARB) or angiotensin-receptor/neprilysin inhibitor (ARNI) contraindications who are prescribed an ACEI, ARB, or ARNI at hospital discharge.

Time Period: 01/2010 - 12/2018



Benchmark Group	Time Period	Numerator	Denominator	% of Patients
All Hospitals	2010	35872	37898	94.7%
All Hospitals	2011	36964	38801	95.3%
All Hospitals	2012	35837	37352	95.9%
All Hospitals	2013	35671	37075	96.2%
All Hospitals	2014	35848	37180	96.4%
All Hospitals	2015	36780	391.28	94.0%
All Hospitals	2016	38290	40900	93.6%
All Hospitals	2017	35533	38466	92.4%
All Hospitals	2018	588	648	90.7%

Post Discharge Appointment for Heart Failure Patients Percent of eligible heart failure patients for whom a follow-up appointment was scheduled and documented including location, date, and time for follow up visits, or home health visit.

Time Period: 01/2010 - 12/2018



Data For: Post Discharge Appointment for Heart Failure Patients					
Benchmark Group	Time Period	Numerator	Denominator	% of Pati ents	
All Hospitals	2010	323	96637	0.3%	
All Hospitals	2011	14502	103989	13.9%	
All Hospitals	2012	45670	98353	46.4%	
All Hospitals	2013	61635	96032	64.2%	
All Hospitals	2014	69578	99138	70.2%	
All Hospitals	2015	77999	104070	74.9%	
All Hospitals	2016	89868	114796	78.3%	
All Hospitals	2017	90172	113236	79.6%	
All Hospitals	2018	1501	1974	76.0%	

Measure LV Function* HF patients with documentation in the hospital record that left ventri cular function (LVF) was assessed before arrival, during hospitalization, or is planned for after discharge. Time Period: 01/2010 - 12/2018



Benchmark Group	Time Period	Numerator	Denominator	% of Patients
All Hospitals	2010	113920	115315	98.8%
All Hospitals	2011	121799	126173	96.5%
All Hospitals	2012	117686	122109	96.4%
All Hospitals	2013	119209	120417	99.0%
All Hospitals	2014	123850	125221	98.9%
All Hospitals	2015	129739	131656	98.5%
All Hospitals	2016	143365	145452	98.6%
All Hospitals	2017	141378	143303	98.7%
All Hospitals	2018	2411	2496	96.6%

Evidence-Based Specific Beta Blockers* Percent of HF patients who were prescribed evidence-based specific beta blockers (Bisoprolol, Carvedilol, Metoprolol succinate CR/XL) at discharge Time Period: 01/2010 - 12/2018



Benchmark Group	Time Period	Numerator	Denominator	% of Patients			
All Hospitals	2010	24745	46684	53.0%			
All Hospitals	2011	29183	48921	59.7%			
All Hospitals	2012	39599	47349	83.6%			
All Hospitals	2013	42096	47411	88.8%			
All Hospitals	2014	43588	48270	90.3%			
All Hospitals	2015	46667	51330	90.9%			
All Hospitals	2016	49529	54363	91.1%			
All Hospitals	2017	47965	52233	91.8%			
All Hospitals	2018	822	224	93.0%			

Beta Blocker at Discharge Percent of patients on Beta blockers at discharge Time Period: 01/2010 - 12/2018



Benchmark Group	Time Period	Numerator	Denominator	% of Patients
All Hospitals	2010	44441	46684	95.2%
All Hospitals	2011	47119	48921	96.3%
All Hospitals	2012	46176	47349	97.5%
All Hospitals	2013	46484	47411	98.0%
All Hospitals	2014	47487	48270	98.4%
All Hospitals	2015	50357	51330	98.1%
All Hospitals	2016	53220	54363	97.9%
All Hospitals	2017	51291	52292	98.1%
All Hospitals	2018	895	910	98.4%

Angiotensin Receptor-Neprilysin Inhibitor (ARNI) at Discharge Percentage of eligible patients with heart failure who are prescribed an ARNI at hospital discharge.

Time Period: 01/2010 - 12/2018



Aldosterone Antagonist at discharge* Percent of heart failure patients with left ventricular ejection fraction <=35% or a qualitative assessment of moderate/severe dysfunction with no contraindications or documented intolerance who were prescribed Aldosterone Antagonist at discharge.

Time Period: 01/2010 - 12/2018



Hydralazine Nitrate at Discharge* Black Heart failure patients with left ventri cular systolic dysfunction (LVSD) with no contraindications or documented intolerance who were prescribed a combination of hydralazine and isosorbide dimitrate at discharge. Note this treatment is recommended in addition to ACEI or ARB and beta blocker therapy at discharge.



Influenza Vaccination During Flu Season Percent of patients that received an influenza vaccination prior to discharge during flu season Time Period: 01/2010 - 12/2018



Data For: Influenza Vaccination During Flu Season						
Benchmark Group	Time Period	Numerator	Denominator	% of Pahents	I	
All Hospitals	2010	16900	58193	29.0%		
All Hospitals	2011	23095	59573	38.8%		
All Hospitals	2012	30710	55181	\$5.7%		
All Hospitals	2013	34258	52781	64.9%		
All Hospitals	2014	36688	55519	66.1%		
All Hospitals	2015	40711	55645	73.2%		
All Hospitals	2016	45562	59355	76.8%		
All Hospitals	2017	42497	56131	75.7%		
All Hospitals	2018	1390	2319	59.9%		
Anticoagulation for Atrial Fibrillation or Atrial Flutter Percent of patients with chronic or recurrent atrial fibrillation or atrial flutter at high risk for thromboembolism, according to CHADS2 risk stratification prescribed Anticoagulation at discharge.

Time Period: 01/2010 - 12/2018



Data For: Anticoagulation for Atrial Fibrillation or Atrial Flutter				
Benchmark Group	Time Period	Numerator	Denominator	% of Patients
All Hospitals	2010	12076	16806	71.9%
All Hospitals	2011	15021	19367	77.6%
All Hospitals	2012	21317	26697	79.8%
All Hospitals	2013	24686	30147	81.9%
All Hospitals	2014	25241	30383	83.1%
All Hospitals	2015	30247	35821	84.4%
All Hospitals	2016	36405	42450	85.8%
All Hospitals	2017	38409	44201	86.9%
All Hospitals	2018	687	757	90.8%

CRT-D or CRT-P Placed or Prescribed at Discharge

Percent of heart failure patients with left ventricular ejection fraction less than or equal to 35% with a QRS duration of 120 ms or above and Left Bundle Branch Block or QRS 150ms or above regardless of QRS morphology, with no contraindications, documented intolerance, or any other reason against who have CRT-D or CRT-P, had CRT-D or CRT-P placed, or were prescribed CRT-D or CRT-P at discharge.



Time Period: 01/2010 - 12/2018



Follow-up Visit Within 7 Days or Less Percent of eligible patients with a follow-up visit scheduled within 7 days or less from time of hospital discharge Time Period: 01/2010 - 12/2018



🔳 All	Hospitals
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	Dat	a For: Follow-up Visit Within 7	Days or Less		
Benchmark Group	Time Period	Numerator	Denominator	% of Patients	ī
All Hospitals	2010	12201	64547	18.9%	
All Hospitals	2011	18063	81895	22.1%	
All Hospitals	2012	34691	92622	37.5%	
All Hospitals	2013	46600	92964	50.1%	
All Hospitals	2014	53839	94779	56.8%	
All Hospitals	2015	60614	100154	60.5%	
All Hospitals	2016	70170	111873	62.7%	
All Hospitals	2017	72548	110652	65.6%	
All Hospitals	2018	1247	1966	63 494	_

Outpatient Cardiac Rehab Program Referral Percent of heart failure patients referred to outpatient cardiac rehab program.

Time Period: 01/2010 - 12/2018



Evidence-Based HFrEF Therapies

Guideline Recommended Therapy	Relative Risk Reduction in Mortality	Number Needed to Treat for Mortality	NNT for Mortality (standardized to 36 months)	Relative Risk Reduction in HF Hospitalizations
ACEI/ARB	17%	22 over 42 months	26	31%
ARNI	16%	36 over 27 months	27	21%
Beta-blocker	34%	28 over 12 months	9	41%
Aldosterone Antagonist	30%	9 over 24 months	6	35%
Hydralazine/Nitrate	43%	25 over 10 months	7	33%
CRT	36%	12 over 24 months	8	52%
ICD	23%	14 over 60 months	23	NA
Ivabradine	NA	NA	NA	26%

Updated from Fonarow GC, et al. Am Heart J. 2011;161:1024-1030.

Influence of Sacubitril/Valsartan on Readmission Rates After HF Hospitalization: PARADIGM HF



30 Day All Cause Readmission Odds Ratio: 0.74; 95% CI 0.56-0.97

30 Day HF Readmission Odds Ratio: 0.62; 95% CI 0.45-0.87

2,383 investigator-reported HF hospitalizations, of which 1,076 (45.2%) occurred in subjects assigned to sacubitril/valsartan and 1,307 (54.8%) occurred in subjects assigned to enalapril.

Desai, A.S. et al. J Am Coll Cardiol. 2016;68(3):241-8.

Hospital Readmission Reduction Program

- Up to 3% cut to all DRGs for readmissions over the expected %
- Up to 1% in fiscal year 2013, 2% in fiscal year 2014, and 3% in fiscal year 2015 and beyond
- Initially AMI, heart failure, and pneumonia
- Expand to COPD, CABG, PCI, and other vascular conditions in 2015
- 10 year decrease in reimbursement to hospitals \$7.1 billion
- Public reporting began in 2010 and the hospital financial penalties began October 2012 (beginning of fiscal year 2013)

Medicare Penalizing 2,211 Hospitals For Excess Readmissions

HRRP Impact: Decreasing 30-Day HF Readmissions Accompanied by Increasing 30 Day Risk-Adjusted Mortality



5,200 additional deaths in 2014 may be related to the HRRP

10,400 additional deaths a year if previous declines in mortality had continued

	Year							
Outcomes	2008	2009	2010	2011	2012	2013	2014	Delta
30-Day Risk Adjusted Readmission with HRRP	23.5%	23.5%	23.4%	23.0%	22.5%	21.6%	21.4%	-2.1%
30-Day Mortality after discharge with HRRP	7.9%	8.1%	8.4%	8.7%	8.8%	9.1%	9.2%	+1.3%
30-Day Mortality after discharge without HRRP (projected)	7.9%	7.8%	7.5%	7.2%	7.0%	6.7%	6.6%	-1.3%

Fonarow GC et al JACC 2017 Oct 10;70(15):1931-1934 Data from Dharmarajan K et al. J Am Med Assoc. 2017;318:270-278.

Has HRRP Reporting of Hospital Readmission Rates and Penalties Affected Patient Outcomes?

JAMA Cardiology | Original Investigation

Association of the Hospital Readmissions Reduction Program Implementation With Readmission and Mortality Outcomes in Heart Failure

Ankar Gupta, MD, PhD; Larry A, Alker, MD, MHS; Deepak L, Bhatt, MD, MPH; Margueritte Cox, MS, MGIST; Adam D, DeVore, MD, MHS; Paul A, Heidenreich, MD, MS; Adrian F, Hernandez, MD, MHS; Eric D, Peterson, MD, MH+R Jinand A, Matsouaka, PhD; Olyde W, Yanoy, MD, MS; Gregg C, Forarow, MD

+ Supplemental content

IMPORTANCE Public reporting of hospitals' 30-day risk-standardized readmission rates following neart failure hospitalization and the financial penalization of hospitals with higher rates have been associated with a reduction in 30-day readmissions but have raised concerns regarding the potential for unintended consequences.

ouscrive: To examine the association of the Hospital Readmissions Reduction Program (HRRP) with readmission and mortality outcomes among patients hospitalized with heart failure within a prospective clinical registry that allows for detailed risk adjustment.

DESIGN_SETTING. AND PARTICIPANTS Interrupted time-series and survival analyses of index heart failure hospitalizations were conducted from January 1. 2006, to Desember 31. 2014. This study included 115 245 fee-for-service Medicare beneficiaries across 416 US hospital sites participating in the American Heart Association Get With The Guidelines-Heart Failure registry. Data analysis took place from January 1. 2017, to June 8, 2017.

EXPOSURES Time Intervals related to the HRRP were before the HRRP implementation (January 1, 2006, to March 31, 2010), during the HRRP implementation (April 1, 2010, to September 30, 2012), and after the HRRP penalties went into effect (October 1, 2012, to December 31, 2014).

MAIN OUTCOMES AND MEASURES Risk-adjusted 30-day and 1-year all-cause readmission and mortality rates.

RESULTS The mean (SD) age of the study population (n = 115.245) was 80.5 (8.4) years, 65.292 (54.6%) were womer, and 91.996 (BI.39%) were writher and 10.27 (9.7%) were black. The 30-day risk-adjust der readmission rate declined from 20.0% before the HRRP implementation to 81.4% in the HRRP penalties phase thraard rate (HR) after vs before the HRRP implementation, 0.91, 95% (C. 0.87-0.95; P < .001). In contrast, the 30-day risk-adjusted mortality rate increased tom 27.5% before the HRRP implementation to 8.0% in the HRRP penalties phase (HR after vs before the HRRP implementation, 18, 95% C, 10.1-21, P < .001). The I-year risk-adjusted readmission and mortality rates followed a similar pattern as the 30-day outcomes. The I-year risk-adjusted mortality rates followed a similar pattern as the 30-day outcomes. The I-year risk-adjusted mortality rates followed a similar pattern as the 30-day outcomes. The I-year risk-adjusted mortality rates followed a similar bortor the HRRP implementation.

CONCLUSIONS AND RELEVANCE Among fee-for-service Medicare beneficiaries discharged after heart failure hospitalizations, implementation of the HRRP was temporally associated with a reduction in 30-day and 1-year readmissions but an increase in 30-day and 1-year mortality. If confirmed, this failing may require reconsideration of the HRRP in heart failure.

JAMA Cardiol. 2018;3(1):44-53. doi:101001/jamacardio.2017.4265 Published online November 12, 2017. Author Affiliations: Author affiliations are listed at the end of this article.

Corresponding Aethors: Grogg C. Fonarow, MD, Division of Cardiology, Ahmarson-ULLA Cardiomyopathy Center, Ronals Reagen-UCLA Medical Centers, (Rosals LeConte Ave, Room 47-123 CHS, Los Angeles, CA900095-1679 (cfroarrow/endent.ucla.edu). The 30-day risk-adjusted readmission rate declined from 20.0% before the HRRP implementation to 18.4% in the HRRP penalties phase (hazard ratio (HR) after vs before the HRRP implementation, 0.91; 95%CI, 0.87-0.95; *P* < .001).

In contrast, the 30-day risk-adjusted mortality rate increased from 7.2% before the HRRP implementation to 8.6% in the HRRP penalties phase (HR after vs before the HRRP implementation, 1.18; 95%CI, 1.10-1.27; *P* < .001).

The 1-year risk-adjusted mortality rate increased from 31.3% to 36.3% (HR, 1.10; 95%Cl, 1.06-1.14; P < .001) after vs before the HRRP implementation.

Increase in Risk-Adjusted Mortality after the HRRP Implementation among FFS Medicare Beneficiaries Hospitalized for HF

Study	GWTG-HF Registry linked to FFS Medicare Data ¹	100% Sample of FFS Medicare Data ²	5% Random Sample of FFS Medicare Data ³
Risk Adjustment	Clinical	Administrative	Administrative
Time Period	Pre-HRRP (2006-2010) vs Post-HRRP (2012-2014)	2008 to 2014	2010 to 2012
30-Day Mortality	1.4% ↑	1.3% ↑	-
90-Day Mortality	-	2.2%↑	-
1-Year Mortality	5.0% ↑	-	3.3%↑

1. Gupta et al. JAMA Cardiol 2017; doi:10.1001/jamacardio.2017.4265.

2. Dharmarajan et al. JAMA 2017;318:270-278.

3. Khera et al. Circ Heart Fail 2017; 10:e004402.

Conclusions

- GWTG-HF is focused on improving on meaningful processes of care and patient-centered outcomes
- The CMS 30 day readmission metric is fundamentally flawed in measuring quality and driving patient benefit
- The CMS HRRP has created a perfect storm for suboptimal care, both by side-stepping the best interests of the patient and by thwarting assessment of risk for both clinicians, in their care, and for CMS in its attempt at adjudication and penalty assignment to hospitals
- It is critical to move entirely away from artificial metrics and penalties and toward greater direct responsibility of health care systems for quality, safety, and value, with any potential rewards linked to long-term patientcentered benefit, through innovative approaches to care



Heart Failure Treatments in Special Populations

Adam DeVore, MD, MHS Assistant Professor of Medicine Duke University School of Medicine

Heart.org/QualityHF



PARADIGM-HF Baseline Characteristics

Table 1. Characteristics of the Patients at Baseline.*				
Characteristic	LCZ696 (N=4187)	Enalapril (N = 4212)		
Age — yr	63.8±11.5	63.8±11.3		
Female sex — no. (%)	879 (21.0)	953 (22.6)		
Race or ethnic group — no. (%)†				
White	2763 (66.0)	2781 (66.0)		
Black	213 (5.1)	215 (5.1)		
Asian	759 (18.1)	750 (17.8)		
Other	452 (10.8)	466 (11.1)		
Region — no. (%)				
North America	310 (7.4)	292 (6.9)		
Latin America	713 (17.0)	720 (17.1)		
Western Europe and other‡	1026 (24.5)	1025 (24.3)		
Central Europe	1393 (33.3)	1433 (34.0)		

McMurray JJ et al. N Engl J Med 2014;371:993-1004.



Populations of Interest

- Elderly
- Females
- Racial and ethnic minorities
- Specific cardiomyopathies
- Comorbid conditions



Heart Failure Care in the Elderly



Mozaffarian D. et al. Circulation. 2015 Jan 27;131(4):e29-322



Heart Failure Care in the Elderly

- High prevalence and poor outcomes
- Different presentations (e.g., different causes of peripheral edema)
- More likely to have non-CV causes of symptoms and more likely to have comorbid conditions (e.g., hypertension, atrial fibrillation)
- Low lean body mass and impaired renal function may increase adverse effects from medical therapy (e.g., hyperkalemia with MRAs or increased risk of digoxin toxicity)
- Increased risk of polypharmacy
- May require more frequent visits and laboratory monitoring

2/13/2018



Heart Failure Care in the Elderly in 2018





Race/Ethnic Differences in Outcomes Among Hospitalized Medicare Patients With Heart Failure and Preserved Ejection Fraction



Boback Ziaeian, MD, PHD,^{a,b} Paul A. Heidenreich, MD, MS,^c Haolin Xu, MS,^d Adam D. DeVore, MD, MHS,^{d,e} Roland A. Matsouaka, PHD,^{d,f} Adrian F. Hernandez, MD, MHS,^{d,e} Deepak L. Bhatt, MD, MPH,^g Clyde W. Yancy, MD,^h Gregg C. Fonarow, MDⁱ

ABSTRACT

OBJECTIVES This study analyzed HFpEF patient characteristics and clinical outcomes according to race/ethnicity and adjusted for patient and hospital characteristics along with socioeconomic status (SES).

BACKGROUND The proportion of hospitalizations for heart failure with preserved ejection fraction (HFpEF) has increased over the last decade. Whether the short- and long-term outcomes differ between racial/ethnic groups is not well described.

METHODS The Get With The Guidelines-Heart Failure registry was linked to Medicare administrative data to identify



Precision Medicine in Heart Failure?





AHA SCIENTIFIC STATEMENT

Current Diagnostic and Treatment Strategies for Specific Dilated Cardiomyopathies

A Scientific Statement From the American Heart Association

The intent of this American Heart Association (AHA) scientific statement is to summarize our current understanding of dilated cardiomyopathies. There is special emphasis on recent developments in diagnostic approaches and therapies for specific cardiomyopathies. Recommendations in this document are based on published studies, published practice guidelines from the American College of Cardiology (ACC)/AHA¹ and other organizations,^{2,3} and the multidisciplinary expertise of the writing group. Existing evidence in epidemiology, classification, diagnosis, and management of specific cardiomyopathies is usually Biykem Bozkurt, MD, PhD, FAHA, Chair Monica Colvin, MD, FAHA Jennifer Cook, MD, FAHA Leslie T. Cooper, MD, FAHA Anita Deswal, MD, MPH, FAHA Gregg C. Fonarow, MD,



Important Comorbidites in Heart Failure





EMPA-REG: Hospitalizations for Heart Failure



Zinman B et al. *N Engl J Med* 2015 Fitchett D et al. *Eur Heart J* 2016



Conclusions

- HF care in the elderly deserves special consideration to improve outcomes and decrease risk of adverse effects
- Opportunities for precision medicine exist in HF through the study of differences in biology by race/ethnicity and specific cardiomyopathies
- Comorbid conditions in HF are common and may offer opportunities to improve care
- Pam Peterson will speak next on women with heart failure



Women with Heart Failure

Pamela N Peterson, MD MSPH

Associate Professor of Medicine University of Colorado Anschutz Medical Center Denver Health Medical Center

Lifetime Risk of Heart Failure



Loyd-Jones DM et al. Circulation 2002; 106:3068

Incidence of HF with Preserved vs. Reduced EF in Men and Women



Ho JE et al. Circ Heart Fail 2013

No Differences in In-Hospital Mortality by Gender or LVEF



Hsich EM et al. Am Heart J 2012

Characteristics by Sex Among those with LVEF <40%



	Female	Male
Age	74	69
Hypertension	74	71
Diabetes	42	40
CAD	48	55
Anemia	17	13
Valvular Disease	12	10
Atrial Fibrillation	26	30
Depression	11	7

Hsich EM et al. Am Heart J 2012



	Female	Male
Age	79	74
Hypertension	81	78
Diabetes	45	48
CAD	41	50
Anemia	24	20
Valve Disease	14	11
Atrial Fibrillation	34	35
Depression	13	9

Hsich EM et al. Am Heart J 2012

No Sex Differences in Treatment of HF

- Women are under-represented in RCTs
- However, available data:
 - Stratified analyses of RCTs
 - Pooled data/ meta-analyses
 - Observational data
- Guidelines do not differ based on sex
- All quality metrics apply equally to men and women

Quality Metrics in Women vs. Men

	Unadjusted		Multivariable	
Characteristic	OR	95% CI	Adjusted OR*	95% CI
Complete set of written instructions at time of discharge	0.95	0.92-0.97	0.97	0.94-1.01
Documentation of evaluation of LV function	0.91	0.88-0.94	0.81	0.76-0.86
ACEI/ARB prescription for LVSD	1.01	0.94-1.07	1.03	0.96-1.11
Adult smoking cessation counseling	1.01	0.94-1.09	1.06	0.95-1.19
β-blocker prescription for LVSD	0.89	0.84-0.95	0.94	0.87-1.03
Defect-free measure (100% compliance with all 5 primary measures)	1.13	1.1–1.16	0.98	0.95-1.01
Composite quality measure	0.97	0.95-0.99	0.96	0.94-0.99
Warfarin at discharge for patients with atrial fibrillation	0.85	0.81-0.89	0.91	0.86-0.96
Evidence based β -blockers prescription for LVSD	0.93	0.89-0.98	1.02	0.97-1.08
Aldosterone antagonists prescription for LVSD	0.95	0.89-1.02	1.06	0.99-1.13
Black patients with LVSD prescribed hydralazine/isosorbide dinitrate	0.82	0.67-1.01	0.80	0.66-0.96
ICD in patients with LVEF ≤35% (before admission or placed during admission)	0.61	0.56-0.67	0.70	0.65-0.75

Sex differences in ICD Counseling 2011-2014 ICD Counseling



Among those counseled, women and men were similarly likely to receive an ICD (OR 1.13; 0.99-1.29)

Hess PL, et al. Circulation 2016

Improvement in care and reduction in sex differences with GWTG participation

100 80 % Treated 60 40 20 0 Baseline 1 year 2 years 3 years 4+years Men Years in Program

Complete Set of Discharge Instructions

Women

Klein L, et al. Circ Heart Fail 2011

Improvement in care and reduction in sex differences with GWTG participation



Evaluation of Left Ventricular Function

Klein L, et al. Circ Heart Fail 2011









Advanced Heart Failure: Marking a Difference



Larry A. Allen, MD, MHS GWTG-HF Webinar






Clinical course of heart failure



Timing of transplant, LVAD, hospice



Difficult to know where a patient is ...



I-NEED-HELP

I: IV inotropes N: NYHA IIIB/IV Natriuretic peptides persistently elevated E: End-organ dysfunction **E**: Ejection fraction <25% **D**: Defibrillator shocks H: Hospitalizations >1 E: Edema, escalating diuretics L: Low blood pressure, high heart rate **P**: Prognostic medication – progressive intolerance or down-titration of GDMT

- Right heart cath? Palliative care?
- Referral to Advanced HF Center?



Not for Everyone: Complex Trade-Offs









McIlvennan, et al. Circ Heart Fail. 2014

P=0.008



McIlvennan, et al. Circ Heart Fail. 2014





- MagLev: no bearings, less friction/heat
- Large rotor gaps: less shear, hemolysis
- Artificial pulse: flush clot, angiodysplasia
- Smaller size: easier implant

MOMENTUM HM3 Endpoints

Table 2. Noninferiority and Superiority Analyses in the Intention-to-Treat Population.*

Variable	Centrifugal-Flow Pump Group (N=152)		Axial-Flow Pump Group (N=142)	
	no. of patients	% (95% CI)	no. of patients	% (95% CI)
Noninferiority analysis				
Primary end point	131	86.2 (79.7–91.2)	109	76.8 (68.9–83.4
Superiority analyses				
Primary end point	131	86.2 (79.7–91.2)	109	76.8 (68.9–83.4)
First event that resulted in failure to reach the primary end point				
Did not receive the assigned implant	1	0.7 (0-3.6)	4	2.8 (0.8-7.1)
Had disabling stroke	6	3.9 (1.5–8.4)	4	2.8 (0.8–7.1)
Underwent reoperation to replace or remove pumpt	1	0.7 (0–3.6)	11	7.7 (3.9–13.4)
Died within 6 months after implantation	13	8.6 (4.6–14.2)	14	9.9 (5.5–16.0)

Mehra M, et al. NEJM. 2016

Transplant remains the Gold Standard



Heart transplant outcomes



- Average Age of Recipient: 54 years old
- Median Survival 10.7 years 1992-2001 cohort
 - Better in post 2002 cohorts
- 93% 1 year survival

ISHLT 2013 Report. J Heart Lung Txplt. 2013; 32: 952.



Which option?





Factor	LVAD	Transplant
Survival, median	4-5 yr	10-13 yr
Quality of life (and swimming)	++	+++
RV failure and ventricular tachycardia	Maybe	Yes
Complications	!!!!	!!!
Stroke, infection, bleeding, HF		
Rejection, infection, cancer, CKD, DM		
Availability of therapy	Unlimited	Limited
Cost	\$\$\$\$	\$\$\$

Option^B

- 1) Advanced age (median HF hosp 78 years)
- 2) Comborbidity (50% have 5+ diagnoses)



Final Perspective

- 6,000,000 with HF
- 2,400,000 (40%) HFrEF
- 240,000 (10%) with stage D
- 60,000 (25%) may benefit from advanced Rx (LVAD/Tx)
- 2,800 transplants
- 4,000 LVADs

... but large benefit in carefully selected patients



GWTG-HF - STATE OF THE ART

Quality of Life in Heart Failure -A Goal Not to be Missed

Nancy M. Albert PhD, CCNS, CHFN, NE-BC, FAHA, FHFSA, FAAN

February 2018



life is why-

Quality of Life in Heart Failure-A Goal Not to be Missed

Objective:

 Discuss the value of understanding quality of life data in patients with heart failure



Quality of Life in HF

Perspectives

Efficacy of Treatments from *Health Care Providers*

- Based on parameters
 - Clinical status
 - Hemodynamics
 - Neurohormonal status
 - Echo/MRI indices

Efficacy of Treatments from *Patients*

- Based on:
 - Functional capacity
 - Exercise performance
 - Psychological status
 - Frequency of

lization

- **1) Under represented in clinical trials**
- 2) No universal definition of quality of life endpoints
- 3) Difficult to standardize data collection

American Heart Stroke Association

life is why≃

Quality of Life Tools in HF 25 tools discussed in the literature

Instrument Name Descri	ption
Minnesota Living w HF Q	21 items; lifestyle limitations; ↓ score = û QoL
Kansas City Cardiomyopathy Q	12/23 items; physical, symptoms, QoL, social impact and self- efficacy; î score = î QoL
Euro HF QoL Q	40 items; functional status, etc.; 企 score = 企 QoL
EuroQ-5D (generic; assesses problems)	VAS; mobility, self-care, usual activities, pain & anxiety/depression domains; & score = 1 QoL
Chronic HF Q	20 items; dyspnea, fatigue, emotional function domains; î score = î QoL
Qual of Life in Severe HF	26 items; physical activity + VAS of life satisfaction- social/emotional; & score = & QoL (less impairment)
Medical Outcomes Study 36- item Short Form	36 items; 8 subscales; assesses negative health aspects; 1 score = 1 QoL
Nottingham Health Profile	38 items based on WHO classification of disabilities;
Sickness Impact Profile	136 Y/N items; 12 areas of pts. life; ♣ score = î QoL

Quality of Life in HF Correlates of *QoL*

- 1037 older ambulatory adults, (KCCQ & EQ-5D)¹
 - Tools were highly related; rho, 0.815
 - Factors associated with worse QoL:
 - ^u Older age, female
 - **Worse functional class**
 - Higher Charlson comorbidity index
 - Recent hospitalization for HF
- 1136 (MLHFQ)² & 52 (KCCQ)³ hospitalized adults
 - QoL improved during hospitalization³ and after discharge in all patients;^{2,3} despite intervention vs. control group²
 - 1. Comi´n-Colet J et al. Rev Esp Cardiol. 2016;69(3):256-271.
 - 2. Riegel B et al. Nurs Res. 2002;51(4):209-18.
- 3. Sauser K, et al. J Card Fail. 2014;20(5):378.e11-5.



Quality of Life in HF Event-Free Survival; by MLHFQ

425 pts. from ESCAPE study; 3 Month Event*



life is why~

Moser DK, et al. *J Cardiac Fail*. 2009;15(9):763-

Quality of Life in HF Event-Free Survival by *Change in* MLHFQ

425 pts. from ESCAPE study; 6 Month Event



Quality of Life in HF

1x Self-Rated Health (SF-12)* Score Predicts Healthcare Utilization





Chamberlain AM, et al. J Am. Heart Assoc. 2014;28(3):e000931.

life is why≃

Quality of Life in HF Mortality Meta-Analysis (x17 papers w >100 pts ea)



Quality of Life in HF-A Goal NOT to Be Missed

When it comes to HF, ~ 44% of patients do not recognize early HF symptoms,¹ & most patients do not recognize HF exacerbation²



Assessment of physical functioning / symptoms via a HR-QoL tool may optimize assessment & treatment ⇒ optimize QoL



- 1. Riegel B, et al. *Heart Lung* 2018; ePub Jan 3.
- 2. Lee S, Reigel B. J Cardiovasc Nurs 2017; ePub Aug 30

Value of Assessing QoL

- If physical health impairments lead to hospitalization or mortality, and change in QoL score 1 month post hospitalization can predict early (60 day to 6 month) event free survival
 - QoL score should be assessed at hospitalization and 1 month after discharge
 - To provide *future* hospitalization/survival risk
 - To help patients understand rationale for implementing interventions known to improve QoL



QoL Goals

- If we help patients understand QOL goals as part of usual care education (based on score improvements known to be associated with improved health status)
- We might enhance patient engagement and empowerment in HF self care



life is why~

Quality of Life in HF Predictors of *Future Health Status*

1458 pts. from EVEREST study; 6 Month Outcomes





Allen LA, et al. Circ. Cardiovasc Qual Outcomes. 2011;4(4):389-398.

life is why~

QoL Goals

- More research is needed to determine if:
 - A standard HF-related QoL tool should be systematically used
 - Tool administration should be standardized in the OPD (every ? months) and hospital at admission/post-discharge (? 30 days)

u To determine CHANGE in scores

 Tool administration and FU *burden* is feasible - time to administer (~ 7 minutes), resources needed to administer/score/share results, communication with patient







life is why-



Contact Us to Learn More

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Thank you for your active participation and contributions to GWTG-Heart Failure!