

# Profile of sacubitril/valsartan in the treatment of heart failure: patient selection and perspectives

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**Abstract:** With an estimated prevalence of 5.8 million in the USA and over 23 million people worldwide, heart failure (HF) is growing in epidemic proportions. Despite the use of guideline-directed medical therapies such as angiotensin-converting enzyme inhibitors, beta-adrenergic blockers, angiotensin receptor blockers, and mineralocorticoid receptor antagonists for chronic systolic HF for almost two decades, HF remains a leading cause of morbidity, mortality, and health care expenditures. The Prospective Comparison of Angiotensin Receptor-Nephrilysin Inhibitor with Angiotensin-Converting Enzyme Inhibitor to Determine Impact on Global Mortality and Morbidity in Heart Failure (PARADIGM-HF) trial provided compelling evidence for the cardiovascular and mortality benefit of sacubitril/valsartan when compared to enalapril in patients with heart failure and reduced ejection fraction (HFrEF). Sacubitril/valsartan performed better than enalapril across various HFrEF patient characteristics and showed substantial benefit in patients with other common comorbidities. Following the trial, the US Food and Drug Administration approved this drug for the treatment of HF. Various international HF consensus guidelines endorse sacubitril/valsartan as a class I recommendation for the management of symptomatic HFrEF. Although this high-quality clinical study is the largest and the most globally represented trial in HFrEF patients, concerns have been raised regarding the generalizability of the trial results in real-world HF population. The gaps in US Food and Drug Administration labeling and guideline recommendations might lead to this medication being used in a larger population than it was studied in. In this review, we will discuss the current role of sacubitril/valsartan in the management of HF, concerns related to PARADIGM-HF and answers, shortcomings of this novel drug, effects on patient characteristics, real-world eligibility, and the role of ongoing and further investigations to clarify the profile of sacubitril/valsartan in the management of HF.

**Keywords:** sacubitril/valsartan, Entresto, HFrEF, systolic heart failure, LCZ696, angiotensin receptor neprilysin inhibitor

## Introduction

Heart failure (HF) is associated with significant morbidity, mortality, and health care expenditure. HF is classified based on left ventricular ejection fraction (LVEF) into HF with reduced EF (HFrEF) with an LVEF <40% and HF with preserved EF (HFpEF) with an LVEF ≥50%.<sup>1</sup> An EF between 40% and 49% is considered an intermediate zone and is termed as HF with borderline EF or HF with mid-range EF. Epidemiologic data indicate that HFpEF and HFrEF contribute equally to the total HF population.<sup>1</sup> HFpEF patients have a similar post-discharge mortality risk and equally high rates of rehospitalization, compared to patients with HFrEF.<sup>2</sup> With an estimated prevalence of 5.8 million in the USA and over 23 million people worldwide, HF is growing in

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epidemic proportions.<sup>3</sup> The cost of HF in the USA was around \$30 billion in 2012, a number that is projected to increase to around \$70 billion by the year 2030.<sup>4</sup>

Acute decompensated HF (ADHF) is the clinical syndrome of new onset or worsening HF symptoms and signs requiring urgent treatment.<sup>5</sup> In the USA, ADHF exacerbations result in around one million hospitalizations yearly and contribute largely to the overall HF health care expenditure.<sup>4</sup> Hospitalization for ADHF serves as a poor prognostic indicator with ~30% and 50% readmission rates at 1 and 6 months, respectively, and a 1-year all-cause mortality as high as 30%.<sup>6,7</sup> The estimated survival rate after the diagnosis of HF is 50% at 5 years and 10% at 10 years.<sup>8</sup> Despite the use of guideline-directed medical therapies such as angiotensin-converting enzyme inhibitors (ACEIs), beta-adrenergic blockers, angiotensin receptor blockers (ARBs), and mineralocorticoid receptor antagonists (MRAs) as cornerstone medical therapies for chronic systolic HF for almost two decades, HF remains a leading cause of morbidity, mortality, and health care expenditures in the USA and worldwide.

Advances in our understanding of the renin–angiotensin–aldosterone (RAAS) pathway and natriuretic peptide system, lessons learned from randomized trials of natriuretic peptide system augmentation, and pharmaco-innovation led to the creation and validation of combination sacubitril/valsartan (Entresto™ [LCZ696]; Novartis) for the treatment of HFrEF. The Prospective Comparison of Angiotensin Receptor-Nepilysin Inhibitor with Angiotensin-Converting Enzyme Inhibitor to Determine Impact on Global Mortality and Morbidity in Heart Failure (PARADIGM-HF) trial provided compelling evidence for the cardiovascular (CV) and mortality benefit of sacubitril/valsartan when compared to enalapril (an ACEI) in patients with HFrEF.<sup>9</sup> Numerous post hoc analyses of the original trial extended the benefits of this innovative medication across a multitude of clinical characteristics.<sup>10</sup> Following the trial, the US Food and Drug Administration (FDA) approved this drug for the treatment of HF. International HF consensus guidelines now endorse sacubitril/valsartan as a class I recommendation for the management of HFrEF.<sup>11–13</sup> In this review, we will discuss the current role of sacubitril/valsartan in the management of HF, shortcomings of this novel drug, effects on patient characteristics, real-world eligibility, and the role of ongoing and further investigations to clarify the profile of sacubitril/valsartan in the management of HF.

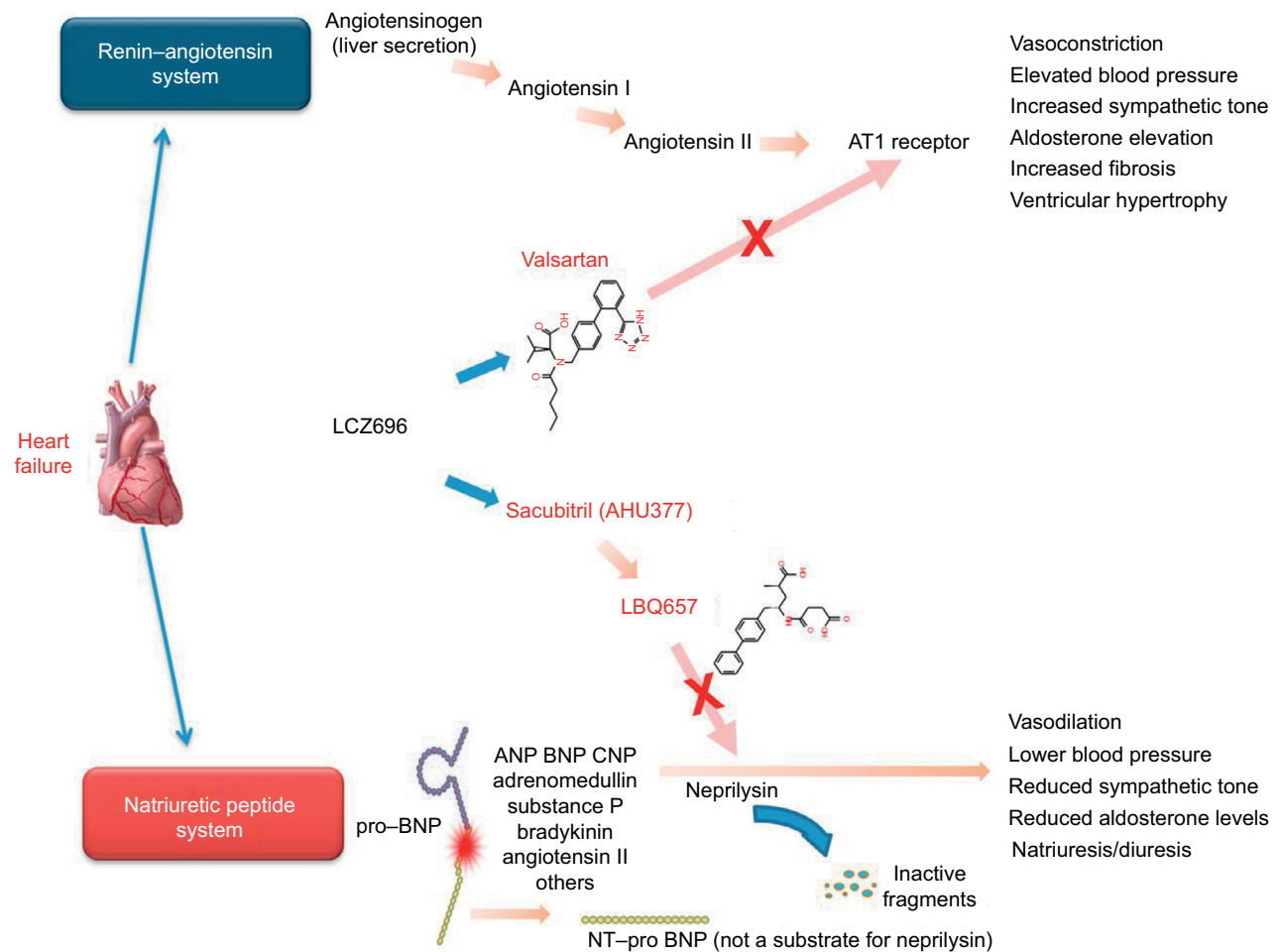
## The PARADIGM-HF trial

LCZ696 is a novel, orally active, first-in-class angiotensin receptor-nepilysin inhibitor (ARNI), which is a 1:1

combination of valsartan (an ARB) and the neprilysin inhibitor (NEPI) sacubitril.<sup>14</sup> Sacubitril (AHU377) is a prodrug, which upon ingestion is rapidly metabolized to an active NEPI moiety LBQ657.<sup>14</sup> The mechanism of action and the effects of LCZ696 on the CV system in HF are explained in Figure 1.<sup>15</sup> Preclinical trials of this combination provided evidence for simultaneous neprilysin inhibition and angiotensin receptor blockade.<sup>10</sup> Favorable safety and efficacy data of this combination from a Phase III randomized controlled trial in hypertensive HFpEF patients paved the way for the larger PARADIGM-HF trial.<sup>9</sup>

The PARADIGM-HF was a Phase III double-blind controlled trial that randomized 8,442 patients with New York Heart Association (NYHA) class II–IV HF and an LVEF of  $\leq 40\%$  (later amended to  $\leq 35\%$ ) to receive twice daily dosing of either 200 mg of LCZ696 or 10 mg of enalapril in addition to standard medical therapy for HF (Table 1).<sup>9</sup> Patients were eligible for the study if they had elevated plasma B-type natriuretic peptide (BNP) level  $\geq 150$  pg/mL (or  $\geq 100$  pg/mL if hospitalized for HF in previous 12 months) or elevated N-terminal pro-BNP (NT-proBNP) level  $\geq 600$  pg/mL (or  $\geq 400$  pg/mL if hospitalized for HF in previous 12 months). At 27-month follow-up, LCZ696 was associated with a 20% decrease in the composite primary endpoint of death from CV causes or hospitalization for HF, compared to enalapril. Additionally, LCZ696 reduced death from any CV cause by 20%, first hospitalization for worsening HF by 21%, all-cause mortality by 16%, and improved the symptoms and quality of life as measured on the Kansas City Cardiomyopathy Questionnaire (Table 1).<sup>9,16</sup> Numerous post hoc analyses have demonstrated the superiority of sacubitril/valsartan over enalapril, irrespective of age, systolic blood pressure (SBP), LVEF, baseline HF risk scores, and glycemic status.<sup>10</sup>

LCZ696 exhibited additional clinical benefit over enalapril in HF patients by attenuating progression of the HF disease process. Fewer LCZ696-treated patients required intensification of medical treatment for HF or an emergency department visit for worsening HF.<sup>17</sup> Patients in the LCZ696 group had 23% fewer hospitalizations for worsening HF evident within the first 30 days of randomization and were less likely to have implantation of an HF device or cardiac transplantation.<sup>17</sup> Patients taking sacubitril/valsartan had early and sustained reduction in biomarkers of myocardial wall stress and injury (NT-proBNP and troponin) and increased plasma BNP levels versus enalapril.<sup>17</sup> Putative placebo analysis of LCZ696 using results from previous landmark placebo-controlled trials of RAAS blockade in HF showed significantly greater reductions in CV death, HF hospitalization, and all-cause mortality with LCZ696.<sup>18</sup> Findings



**Figure 1** Mechanism of action of sacubitril/valsartan in heart failure.

**Notes:** Heart failure stimulates both the renin-angiotensin system and the natriuretic peptide system. LCZ696 is composed of two molecular moieties, the angiotensin receptor blocker valsartan and the neprilysin inhibitor prodrug sacubitril (AHU377). Valsartan blocks the AT1 receptor. Sacubitril is converted enzymatically to the active neprilysin inhibitor LBQ657, which inhibits neprilysin, an enzyme that breaks down ANP, BNP, and CNP, as well as other vasoactive substances. NT-proBNP is not a substrate for neprilysin. Reprinted from *JACC: Heart Failure*, Volume 2/Edition 6, Vardeny O, Miller R, Solomon SD, Combined neprilysin and renin-angiotensin system inhibition for the treatment of heart failure, Pages 663–670, Copyright (2014), with permission from Elsevier.<sup>15</sup>

**Abbreviations:** ANP, atrial natriuretic peptide; AT1, angiotensin type I; BNP, brain (or B-type) natriuretic peptide; CNP, C-type natriuretic peptide; NT-proBNP, N-terminal pro-BNP.

from the PARADIGM-HF trial and subsequent secondary analyses incited a lot of excitement in the HF community, as the discovery of sacubitril/valsartan proves momentous in the relatively stagnant field of HF pharmacotherapy.

## International consensus recommendation for sacubitril/valsartan in HF

Following the trial, the USA FDA reviewed LCZ696 under its priority review program with a fast-track designation and approved this drug for the treatment of all NYHA II–IV HFrEF patients in July 2015.<sup>19</sup> The USA FDA labeling is overarching, recommending the medication for all NYHA function class II–IV HFrEF patients with serum potassium  $\leq 5.2$  mmol/L, and no contraindication or intolerance to ACEI/

ARB. The same year, the European Medical Agency approved the use of sacubitril/valsartan in adult patients for treatment of symptomatic chronic HFrEF.<sup>20</sup>

In their 2016 focused update on HF guidelines, the American College of Cardiology, the American Heart Association, and the Heart Failure Society of America recommended replacing an ACEI or ARB with an ARNI in patients with chronic symptomatic HFrEF, NYHA class II or III, currently tolerating an ACEI or ARB, to further reduce morbidity and mortality (class I recommendation).<sup>11</sup> The guideline also recommended against concomitant use of ARNI with ACEI or within 36 hours of last ACEI dose, as it can precipitate angioedema (class III recommendation).<sup>11</sup>

The Canadian Cardiovascular Society was the first to incorporate the use of sacubitril/valsartan in HF patients.<sup>12</sup> In

**Table 1** The PARADIGM-HF trial<sup>9,16</sup>

Study (first author, year); Aim	Study population	Design	Outcomes
PARADIGM-HF (McMurray et al, 2014); <sup>9</sup> comparison of efficacy of LCZ696 versus enalapril in patients with HFrEF (LVEF ≤35%)	n=8,442; mean age 64 years; mean LVEF 29±6%; 70% NYHA class II; 78% male; 66% white; 5% black; 60% ICM; 100% on either an ACEI or an ARB; 93% on beta-blockers	Multicenter, randomized, double-blind study of LCZ696 200 mg twice daily versus enalapril 10 mg twice daily for a median of 27 months 2-week enalapril run-in period and a 4–6-week LCZ696 run-in period prior to randomization to ensure target dose tolerability	<ul style="list-style-type: none"> <li>• ARR 4.7%, HR 0.8, NNT 21 in LCZ696 arm for the composite of CV death or HF hospitalization</li> <li>• ARR 3.2%, HR 0.8, NNT 32 in LCZ696 arm for any CV death</li> <li>• 21% reduction (HR=0.79) in the first worsening HF hospitalization in LCZ696 arm</li> <li>• 16% reduction (HR=0.84) in all-cause mortality in LCZ696 arm</li> <li>• Improved KCCQ scores in LCZ696 arm at 8 months</li> <li>• Drug discontinuation due to adverse events (10.7% versus 12.3% in LCZ696 versus enalapril arms, <math>p=0.03</math>)</li> <li>• 14% of patients receiving LCZ696 experienced hypotension, compared to 9% in the enalapril group (<math>p&lt;0.001</math>), no significant difference in drug discontinuation due to hypotension</li> <li>• 19 cases of angioedema in LCZ696 arm versus 10 cases in enalapril group (<math>p=0.13</math>)</li> <li>• Cough (11.3% in the LCZ696 group versus 14.3% in the enalapril group, <math>p&lt;0.001</math>)</li> <li>• Elevated serum potassium &gt;6.0 mmol/L (4.3% in the LCZ696 group versus 5.6% in the enalapril group, <math>p=0.007</math>)</li> <li>• Elevated serum creatinine ≥2.5 mg/dL (3.3% in the LCZ696 group versus 4.5% in the enalapril group, <math>p=0.007</math>)</li> </ul>

**Notes:** Yandrapalli S, Aronow WS, Mondal P, et al. Limitations of sacubitril/valsartan in the management of heart failure. *American Journal of Therapeutics*. Volume 24, Issue 2: Pages 234–239.<sup>16</sup>

**Abbreviations:** ACEI, angiotensin-converting enzyme inhibitor; ARB, angiotensin receptor blocker; ARR, absolute risk reduction; CV, cardiovascular; HF, heart failure; HFrEF, heart failure with reduced ejection fraction; HR, hazard ratio; ICM, ischemic cardiomyopathy; KCCQ, Kansas City Cardiomyopathy Questionnaire; LVEF, left ventricular ejection fraction; NNT, number needed to treat; NYHA, New York Heart Association; PARADIGM-HF, Prospective Comparison of Angiotensin Receptor-Nephrilysin Inhibitor with Angiotensin-Converting Enzyme Inhibitor to Determine Impact on Global Mortality and Morbidity in Heart failure.

their 2014 update on Heart Failure Management Guidelines, the Canadian guidelines recommend the use of LCZ696 in place of an ACEI or an ARB in patients with mild to moderate HF, LVEF <40%, an elevated natriuretic peptide level or hospitalization for HF in the past 12 months, a serum potassium <5.2 mmol/L, an estimated glomerular filtration rate (eGFR) ≥30 mL/min, and treated with appropriate doses of guideline-directed medical therapy, with close surveillance of serum potassium and creatinine (Conditional Recommendation; High-Quality Evidence).<sup>12</sup> The Canadian HF guidelines limit the use of ARNI in patients who met the PARADIGM trial entry criteria.

The 2016 European Society of Cardiology (ESC) HF guidelines recommend the use of sacubitril/valsartan as an ACEI replacement to further reduce the risk of death and HF hospitalization in ambulatory patients with HFrEF (LVEF <35%) who remain symptomatic despite optimal

treatment with ACEI, a beta-blocker, and an MRA (class IB recommendation).<sup>13</sup> In contrast to the American College of Cardiology, the American Heart Association, and the Heart Failure Society of America HF guidelines, the ESC guidelines specify having an LVEF cutoff of 35% prior to initiation of sacubitril/valsartan. The ESC guidelines also require the patient to be tried on an MRA before initiating an ARNI.

## Concerns with PARADIGM-HF and answers

Current recommendations for the use of sacubitril/valsartan in HFrEF patients are based on the results of one trial, the PARADIGM-HF. Although this study is the largest and a globally represented high-quality clinical trial in HFrEF patients that evaluated key outcomes, it is not exempt from criticism. Since the publication of the original trial results in 2014, several physicians and researchers have raised

apprehensions regarding the generalizability of the trial results with regards to the study design, the effect of target medication doses achieved on clinical outcomes, study population representativeness of the real-world HFrEF patients, and the effect of ethnic and geographic variations on outcomes.<sup>16</sup>

PARADIGM-HF was initiated without a Phase II safety trial, and hence, the investigators employed steps to test the tolerability of LCZ696 before randomization with a run-in phase for enalapril and then for LCZ696.<sup>10</sup> More than 10% of the original study population dropped out during the LCZ696 run-in phase.<sup>9</sup> Concern existed regarding randomizing only the patients who tolerated LCZ696 during the run-in phase of the trial, as such a design can exclude patients who are intolerant to the drug and can affect study outcomes and external validity. However, a post hoc analysis of the trial data showed that run-in noncompletion did not affect the magnitude of the treatment benefit of LCZ696 over enalapril.<sup>21,22</sup> Low blood pressure, low eGFR, and more advanced HF (higher NTproBNP level) were associated with higher risk for run-in noncompletion, suggesting that these patients should undergo closer monitoring during the up-titration of LCZ696 or the conversion of patients from enalapril to LCZ696.<sup>21,22</sup>

The mean target dose achieved for enalapril (18.9 mg/day) was lower than the maximum recommended dose in HF, whereas the maximum recommended target dose was achieved for sacubitril/valsartan.<sup>16</sup> This raises the concern as to whether the maximal dose of valsartan produced more RAAS blockade leading to better clinical outcomes when compared to a moderate dose of enalapril, as no trials showed head-to-head comparison of the study doses of valsartan and enalapril as used in the PARADIGM-HF.<sup>16</sup> The mean target dose for enalapril achieved in most placebo-controlled landmark HF trials providing significant CV benefit was also <20 mg/day (16.6 mg/day in the Studies of Left Ventricular Dysfunction and 18.4 mg/day in the Cooperative North Scandinavian Enalapril Survival Study).<sup>21</sup> It is important to note that in routine clinical practice, <30% HF patients achieved the target dose of enalapril due to complications including hypotension and hyperkalemia.<sup>21</sup> Although LCZ696-treated patients had greater risk of developing symptomatic hypotension (14% versus 9%,  $p < 0.001$ ; Table 1), it did not result in significantly more medication discontinuation than enalapril patients, suggesting better tolerance with LCZ696.<sup>9,21</sup>

Black patients have been less represented in HF trials, although they have poorer outcomes. The lower proportion of black patients in the PARADIGM-HF trial (5%) may have limited the ability of this trial to detect a significant

difference in the frequency of angioedema with LCZ696.<sup>16</sup> This is particularly concerning, as black patients had significantly more instances of angioedema compared to enalapril in the Omapatrilat Cardiovascular Treatment versus Enalapril trial.<sup>16</sup> Future studies should shed light on this association.

## Profile of sacubitril/valsartan in HF and patient selection perspectives

In this section, we will discuss the profile of sacubitril/valsartan in HF with regards to patient characteristics and future perspectives for its use, with a mention of currently undergoing clinical trials. Registry data on HFrEF patients from Acute Decompensated Heart Failure National Registry (ADHERE) and Organized Program to Initiate Lifesaving Treatment in Hospitalized Patients With Heart Failure (OPTIMIZE-HF) were used to provide comparisons with PARADIGM-HF patient population in available baseline demographic characters (Table 2).<sup>9,23,24</sup>

OPTIMIZE-HF is a large US national registry and performance improvement program for patients hospitalized for HF.<sup>23</sup> ADHERE is a large multicenter registry designed to compile a large clinical database on the clinical characteristics, management, and outcomes of patients hospitalized for HF across the USA.<sup>24</sup> When compared to the HFrEF population in these registries, the PARADIGM-HF study population is relatively younger and male dominant. Additionally, the population has less black subjects, higher mean LVEF, lower SBP at randomization, excluded chronic kidney disease (CKD) stage 4 and 5 (eGFR <30 mL/min/1.73 m<sup>2</sup>) patients, and most patients in NYHA functional class II, with almost no NYHA class IV patients. ADHERE and OPTIMIZE-HF represent real-world US HF patients; hence, these differences in patient characteristics have the potential to affect clinical outcomes in real-world HFrEF patients.

## Age

Similar to the real-world HF population, mortality and HF hospitalization rate increased with age in PARADIGM-HF.<sup>9</sup> The median patient age in the trial was 63.8 years, which is lower than the real-world HFrEF population and those in ADHERE (mean 70 years) and OPTIMIZE-HF (mean 70 years) registry (Table 2).<sup>9,23,24</sup> This difference could have important clinical considerations with regards to better outcomes noticed in this relatively younger ambulatory HFrEF cohort. It is interesting to note that the clinical benefit observed in the sacubitril/valsartan group was consistent across prespecified age groups including patients <55 and >75 years, when compared to enalapril.<sup>25</sup> HF is a progressive



**Table 2** Baseline patient characteristics in various reduced ejection fraction heart failure patient study groups

Study population characteristics	OPTIMIZE- HF <sup>23</sup>	ADHERE <sup>24</sup>	PARADIGM-HF <sup>9</sup>
Patients with HFrEF (among those with data on LVEF assessment)	20,118	25,865	8,442
Mean age (SD), years	70 (14)	70 (14)	63.8 (11.5)
Sex	62% male	60% male	78% male
Race			66% Caucasian
	21% black	22% black	5% black
			18% Asian
Etiology of HF	54% ICM	59% population	60% ICM
	17% hypertensive HF	had CAD	11.5% Hypertensive HF
Hypertension	66%	69%	70%
Diabetes mellitus	39%	40%	34.6%
CKD		26%	–
Atrial fibrillation	28%		37%
LVEF±SD (%)	24±8		29.5±6.2
Mean SBP (mmHg)	135±31		121 (15)
NYHA functional class	44% class IV		5% class I
			70% class II
			24% class III
			0.7% class IV

**Abbreviations:** ADHERE, Acute Decompensated Heart Failure National Registry; CAD, coronary artery disease; CKD, chronic kidney disease; HF, heart failure; HFrEF, heart failure with reduced ejection fraction; ICM, ischemic cardiomyopathy; LVEF, left ventricular ejection fraction; NYHA, New York Heart Association; OPTIMIZE-HF, Organized Program to Initiate Lifesaving Treatment in Hospitalized Patients With Heart Failure; PARADIGM-HF, Prospective Comparison of Angiotensin Receptor-Nephrilysin Inhibitor with Angiotensin-Converting Enzyme Inhibitor to Determine Impact on Global Mortality and Morbidity in Heart failure; SBP, systolic blood pressure.

disease and advances in severity with age, so initiation of sacubitril/valsartan at a younger age in HFrEF patients who have a less-advanced disease stage will provide more clinical and survival benefit.

## Sex

The study population in PARADIGM-HF is predominantly male (78%) with a rate higher than that observed for HFrEF patients in the ADHERE (60%) and OPTIMIZE-HF registries (62%) (Table 2).<sup>9</sup> This proportion is higher when compared to real-world HF patients in whom HFrEF is almost equally prevalent sex-wise. However, in prespecified subgroup analysis of the PARADIGM-HF, the clinical benefit of sacubitril/valsartan was not affected by sex.<sup>9</sup> ARNI can be used without reservation in HFrEF patients of either sex.

## Race/ethnicity

As mentioned earlier, caution should be exercised when prescribing ARNI to black HFrEF patients, as available safety data about angioedema is limited in this underrepresented population in the PARADIGM-HF.<sup>9,16</sup> The prospective comparison of ARNI with ACE inhibitor to determine the novel beneficial treatment value in Japanese HF patients (PARALLEL-HF) trial is now being conducted to assess the safety and efficacy of sacubitril–valsartan in Japanese HFrEF population. This is a multicenter, randomized, double-blind study with similar design, phases, inclusion and exclusion criteria as in PARADIGM-HF trial (Table 3).<sup>26</sup>

## Regional demographic variations

Significant regional differences in baseline demographics such as age, symptoms, comorbidities, background therapies, and event rates between patients enrolled from different geographic locations exist in the PARADIGM-HF cohort.<sup>27</sup> A post hoc analysis of the original trial demonstrated a consistent clinical benefit of sacubitril/valsartan across geographic regions, despite having significant baseline regional demographic variations.<sup>27</sup> These findings are encouraging for the global use of this novel pharmacotherapy in HFrEF patients to provide reduction in HF morbidity and mortality across the world.

## NYHA functional class

Majority of the study population in the PARADIGM-HF consisted of NYHA class II patients (70%).<sup>9</sup> A nominally significant interaction between NYHA class at randomization and the effect of treatment was seen for the primary endpoint (better outcomes with NYHA class I–II;  $p=0.03$ ; without adjustment for multiple comparisons), but not for the interaction between NYHA class and the effect on death from CV causes ( $p=0.76$ ).<sup>9</sup> Although the interaction is nominal, it confers a greater CV benefit to sacubitril/valsartan when initiated early in the course of HFrEF. With almost 75% patients in the trial classified as either NYHA class I or II, the power of this group to identify an interaction if present may be increased. Nevertheless, initiation of sacubitril/valsartan early in the disease course of HFrEF is encouraged.

**Table 3** Currently undergoing trails of sacubitril/valsartan which will answer important clinical questions

Trial name (identifier)	Type, phase	Study start–end	n	Study population	Intervention	Control arm	Primary endpoint	Secondary endpoint	Duration of study
PIONEER-HF (NCT02554890) <sup>38</sup>	RCT, Phase 4	April 2016–April 2018	736	HFpEF patients hospitalized for ADHF	Sacubitril–valsartan	Enalapril	Percentage change from baseline in NT-proBNP	Incidence of symptomatic hypotension, hyperkalemia, angioedema; change in high-sensitivity troponin, urinary cGMP, BNP to NT-proBNP ratio	8 weeks
TRANSITION (NCT02661217) <sup>39</sup>	RCT, Phase 4	February 2016–March 2018	1,000	HFpEF patients hospitalized for ADHF	Initiation of sacubitril–valsartan pre-discharge	Initiation of sacubitril–valsartan post-discharge (up to 14 days)	Percentage of patients taking target dose of sacubitril–valsartan 200 mg BID at 10 weeks post-randomization	Number of patients who achieved and maintained either the dose of sacubitril–valsartan 100 and/or 200 mg BID for at least 2 weeks; number of patients who, regardless of previous dose changes, achieved any dose of sacubitril–valsartan BID for at least 2 weeks; percentage of patients who permanently discontinued from treatment due to adverse events	26 weeks
PARADISE-AMI (NCT02924727) <sup>40</sup>	RCT, Phase 3	April 2016–July 2019	4,650	Post-MI with LVEF ≤40%	Sacubitril–valsartan	Ramipril	Composite endpoint of CV death, HF hospitalization, or outpatient HF	Time to first occurrence of composite of CV death or HF hospitalization, composite of HF hospitalization or outpatient HF, composite of CV death, nonfatal spontaneous myocardial infarction, or nonfatal stroke, total number of recurrent confirmed composite endpoints, time to all-cause mortality	32 months
PARAGON-HF (NCT01920711) <sup>28</sup>	RCT, Phase 3	July 2014–March 2019	4,829	HFpEF, elevated NT-proBNP, structural heart disease	Sacubitril–valsartan	Valsartan	Composite of CV death and total (first and recurrent) HF hospitalizations	Change in KCCQ score, change in NYHA functional class, time to first composite renal endpoint (renal death, ESRD, or ≥50% decline in eGFR), time to all-cause mortality	57 months
UK HARP-III (ISRCTN11958993) <sup>42</sup>	RCT	January 2014–January 2017	400	Proteinuric CKD in the last 3 months with eGFR 20–60 mL/min/1.73 m <sup>2</sup>	Sacubitril–valsartan	Irbesartan	Change in eGFR	Change in urine albumin:creatinine ratio, pharmacokinetics at 3 months	12 months

(Continued)

Table 3 (Continued)

Trial name (identifier)	Type, phase	Study start–end	n	Study population	Intervention	Control arm	Primary endpoint	Secondary endpoint	Duration of study
PARALLEL-HF (NCT02468232) <sup>26</sup>	RCT, Phase 3	June 2015–November 2018	225	Same as PARADIGM-HF trial	Sacubitril–valsartan	Enalapril	Composite of CV death or HF hospitalization	Change in NT-proBNP; time to the first occurrence of CV death, HF hospitalization, or intensification of treatments due to documented episode(s) of worsening HF; change in NYHA functional class; number of patients with reported total adverse events, serious adverse events and death; composite of CV death and total number of HF hospitalizations; clinical composite score; time to all-cause mortality; number of hospitalized patients; number of hospital admissions; number of days in ICU; number of rehospitalizations; number of emergency visits for HF; changes in NT-proBNP; changes in blood amino-terminal propeptide of procollagen type III; changes in urine cGMP	40 months
PARALLAX (NCT03066804) <sup>29</sup>	RCT, Phase 3	June 2017–December 2019	2,200	HFpEF, NYHA class II–IV, KCCQ <75, structural heart disease	Sacubitril–valsartan	Enalapril, valsartan, or placebo	Change in NT-proBNP	Change in KCCQ clinical summary score; percentage of patients with $\geq 5$ -point deterioration or improvement in KCCQ CSS; change in the 6-minute walk test; change in NYHA functional class; change in SF-36 physical component summary score	24 weeks

**Abbreviations:** ADHF, acute decompensated heart failure; BID, twice daily; c-GMP, cyclic-guanosine monophosphate; CKD, chronic kidney disease; CSS, clinical summary score; CV, cardiovascular; eGFR, estimated glomerular filtration rate; ESRD, end-stage renal disease; HF, heart failure; HFpEF, heart failure with preserved ejection fraction; HFrEF, heart failure with reduced ejection fraction; ICU, intensive care unit; KCCQ, Kansas City Cardiomyopathy Questionnaire; LVEF, left ventricular ejection fraction; MI, myocardial infarction; NT-proBNP, N-terminal pro-brain natriuretic peptide; NYHA, New York Heart Association; PARADISE-MI, Prospective ARNI versus ACE Inhibitor Trial to Determine Superiority in Reducing Heart Failure Events After MI; PARAGON-HF, Prospective comparison of Angiotensin Receptor-neprilysin inhibitor with ARB Global Outcomes in HF; PARALLAX, A Randomized, Double-blind Controlled Study Comparing LCZ696 to Medical Therapy for Comorbidities in HFpEF Patients; PARALLEL-HF, Study of Efficacy and Safety of LCZ696 in Japanese Patients With Chronic Heart Failure and Reduced Ejection Fraction; PIONEER-HF, Comparison of Sacubitril/Valsartan versus Enalapril on Effect on NT-proBNP in Patients Stabilized from an Acute Heart Failure Episode; RCT, randomized controlled trial; SF-36, 36-Item Short Form Survey TRANSITION, Therapy in HFrEF Patients After an Acute Decompensation Event; UK HARP-III, UK Heart and Renal Protection III.



## Type of HF

Sacubitril/valsartan is currently approved for use in chronic HFrEF patients. Having demonstrated safety in HFpEF population and efficacy in hypertensive patients in earlier trials, sacubitril/valsartan has the potential to show substantial clinical benefit in the HFpEF population, since hypertension is an important risk factor for HFpEF. The Phase III, randomized controlled, Prospective comparison of Angiotensin Receptor-neprilysin inhibitor with ARB Global Outcomes in HF with preserved ejection fraction (PARAGON-HF) trial is evaluating the effects of sacubitril/valsartan versus valsartan on the primary composite outcome of CV death and HF hospitalization in HFpEF patients and is expected to be completed in 2019 (Table 3).<sup>28</sup> The PARAGON-HF will also compare treatment benefit of sacubitril/valsartan versus valsartan on functional class, change in Kansas City Cardiomyopathy Questionnaire score, and time to deterioration in renal function.<sup>28</sup> The randomized, double-blind controlled study comparing LCZ696 to medical therapy for comorbidities in HFpEF patients (PARALLAX) is a 24-week, multicenter, parallel-group, active controlled study which will evaluate the effect of sacubitril/valsartan on NT-proBNP levels, symptoms, exercise function, and safety, compared to individualized medical management of comorbidities (with enalapril, valsartan, or placebo) in HFpEF patients (Table 3).<sup>29</sup>

## Etiology of HF

HFrEF has different etiologies depending on age, sex, geography, and race. Clinical outcomes and response to medical therapies in HFrEF patients will vary depending on etiology. In the PARADIGM-HF trial, 60% of the study population had ischemic cardiomyopathy (ICM), which is comparable to that observed in HFrEF patients in the ADHERE and the OPTIMIZE-HF registries (Table 2).<sup>9,23,24</sup> This raises concern of the benefit of sacubitril/valsartan in patients with non-ICM (NICM). A post hoc analysis of the PARADIGM-HF classified the study population based on etiology of HF into patients with ICM (60%) and NICM (40%).<sup>30</sup> NICM consisted of idiopathic HF (19%), hypertensive HF (11.5%), and other causes of HF (9.5%).<sup>30</sup> The investigators noted that patients with NICM were younger, more likely female, and had higher NT-proBNP levels than those with ICM.<sup>30</sup> With regards to primary composite endpoint and CV death, the benefit of sacubitril/valsartan over enalapril was consistent across the different etiology subgroups.<sup>30</sup> The authors concluded that sacubitril/valsartan is effective in both ICM and NICM including idiopathic and hypertensive causes of NICM. These

findings have important considerations when using this ARNI in real-world HFrEF patients and suggest that the complex pathophysiologic interactions of ARNI with the RAAS are independent of the type of insult resulting in HF.

## Left ventricular ejection fraction

In the PARADIGM-HF trial, the risk of all clinical outcomes increased with decreasing LVEF (mean 29.5%), and LVEF was noted to be a strong predictor of outcomes.<sup>9</sup> In a post hoc analysis, sacubitril/valsartan was effective across the LVEF spectrum and there was no evidence of heterogeneity for the primary endpoint, CV death, HF hospitalization, or all-cause mortality.<sup>31</sup> These findings suggest that the benefit of sacubitril/valsartan is not LVEF dependent and may have implications in its testing in HFpEF patients.

## Intracardiac devices

Patients with cardiac devices were underrepresented in the PARADIGM-HF trial. Around 15% had implantable cardioverter-defibrillator and around 7% patients had cardiac resynchronization therapy.<sup>9</sup> Despite this limitation, the magnitude of clinical benefit of sacubitril/valsartan on sudden death did not differ in patients with or without an implantable cardioverter-defibrillator.<sup>32</sup> ARNI use will provide additional CV and survival benefit in symptomatic HFrEF patients, irrespective of them having an intracardiac device.

## ADHF and post-acute myocardial infarction patients

PARADIGM-HF excluded patients who had an acute coronary event within the last 3 months of randomization.<sup>9</sup> The benefit of initiating or uptitrating neurohormonal antagonist during an episode of ADHF is not clear.<sup>33</sup> However, it is well known that initiation of beta-blocker, ACEI, and an MRA after acute myocardial infarction (MI) with associated left ventricle systolic dysfunction reduces the rate of hospitalization for HF and mortality.<sup>34-36</sup> In a post hoc analysis of the PARADIGM-HF trial, the benefit of sacubitril-valsartan in reducing CV mortality or hospitalization for HF was consistent in all patient subgroups, with no significant difference in patients with no prior HF hospitalization compared to those with recent HF hospitalization (<3 months from randomization).<sup>37</sup>

Although available data do not support the use of this ARNI in ADHF patients and in post-acute MI patients, ongoing trials will shed more light. Comparison of Sacubitril/Valsartan versus Enalapril on Effect on NT-proBNP in Patients Stabilized from an Acute Heart Failure Episode (PIONEER-HF) is a multicenter, randomized, double-blind, 8-week study,

with the aim being to evaluate the effect of sacubitril–valsartan versus enalapril on changes in NT-proBNP in HFrEF patients who have been stabilized following hospitalization for ADHF. The primary outcome is the percentage change in NT-proBNP from baseline to weeks 4 and 8 (Table 3).<sup>38</sup>

The Comparison of Pre- and Post-discharge Initiation of LCZ696 Therapy in HFrEF Patients After an Acute Decompensation Event (TRANSITION) trial is a multicenter, randomized, open-label study, with an estimated enrollment of 1,000 patients, aiming to compare the initiation of sacubitril/valsartan after stabilization of patients with ADHF at any point prior to discharge with post-discharge initiation (up to 14 days). The primary outcome is the percentage of patients who are receiving sacubitril/valsartan 200 mg twice daily at 10 weeks after randomization (Table 3).<sup>39</sup> This trial will also evaluate the safety of initiation of sacubitril/valsartan in ADHF and the percentage of patients who permanently discontinued from treatment due to adverse events.<sup>39</sup>

The Prospective ARNI versus ACE Inhibitor Trial to Determine Superiority in Reducing Heart Failure Events After MI (PARADISE-MI) trial is a Phase III, randomized controlled study conducted in post-acute MI patients without prior chronic HF, with LVEF<40%, and/or pulmonary congestion at the time of randomization. The PARADISE-MI trial will evaluate the benefit of sacubitril/valsartan versus ramipril in reducing the occurrence of the primary composite endpoint of CV death, HF hospitalization, and outpatient HF (time-to-first event analysis), as shown in Table 3.<sup>40</sup>

## CKD and hyperkalemia

HFrEF patients frequently have CKD with varying degrees of eGFRs. Worsening renal function and hyperkalemia can limit the successful use or up-titration of RAAS blockade agents in HF patients.<sup>41</sup> The PARADIGM-HF trial excluded patients with eGFR<30 mL/min/1.73 m<sup>2</sup>, thereby limiting safety data on its use in CKD stage 4 and 5 patients. Interestingly, it was observed that LCZ696-treated patients had significantly less instances of elevated serum creatinine  $\geq 2.5$  mg/dL (3.3% in the LCZ696 group versus 4.5% in the enalapril group,  $p=0.007$ ) compared to enalapril-treated patients. The UK Heart and Renal Protection III (UK HARP-III) trial will compare LCZ696 against irbesartan (an ARB) in patients with proteinuric CKD (eGFR 20–60 mL/min/1.73 m<sup>2</sup>) to assess the short-term safety and efficacy of LCZ696 in this population with respect to changes in eGFR (Table 3).<sup>42</sup> If the results from the UK HARP-III trial favor LCZ696, longer duration studies to evaluate the effects of LCZ696 in delaying the progression of CKD to end-stage renal disease should be planned.

With regards to severe hyperkalemia, LCZ696-treated patients had lower instances of elevated serum potassium  $>6$  mmol/L (4.3% in the LCZ696 group versus 5.6% in the enalapril group,  $p=0.007$ ; Table 1).<sup>9</sup> Moreover, among patients treated with an MRA, severe hyperkalemia was more likely with enalapril treatment than with LCZ696 (3.1 versus 2.2 per 100 patient-years,  $p=0.02$ ), suggesting that LCZ696 might decrease the risk of hyperkalemia compared to enalapril, when added to MRA therapy in patients with HFrEF.<sup>43</sup> Although further safety data will be available from currently ongoing trials, sacubitril/valsartan therapy might facilitate the up-titration of other RAAS blocking agents with proven mortality benefit in HFrEF patients without increasing the incidence of dose-limiting adverse events.

## Diabetes mellitus

In the PARADIGM-HF trial, it was observed that patients with diabetes and pre-diabetes had a higher risk of the primary composite outcome.<sup>9</sup> Sacubitril/valsartan reduced the risk of CV death and HF hospitalizations compared with enalapril, irrespective of the glycemic status.<sup>44</sup> A recently published post hoc analysis of the PARADIGM-HF trial found that sacubitril/valsartan decreased hemoglobin A1C levels by 0.26% during the first year of follow-up, compared to a 0.16% reduction with enalapril ( $p=0.0023$ ).<sup>45</sup> Over 3 years, HbA1c levels remained persistently lower in patients treated with sacubitril/valsartan compared to enalapril, with an overall reduction of 0.14% (95% CI: 0.06–0.23,  $p=0.0055$ ).<sup>45</sup> In addition, 29% fewer sacubitril/valsartan-treated patients initiated insulin therapy to achieve glycemic control (7% versus 10% patients, hazard ratio 0.71, 95% CI: 0.56–0.90;  $p=0.0052$ ).<sup>45</sup> These results are interesting as they hypothesize an additional metabolic benefit and emphasize the extracardiac pleiotropic effects of sacubitril/valsartan in HFrEF patients in addition to its compelling CV efficacy.

## Eligibility of the real-world HF patients for sacubitril/valsartan use

The US FDA labeling for sacubitril/valsartan is very broad, recommending this medication for all NYHA function class II–IV HFrEF patients with serum potassium  $\leq 5.2$  mmol/L and no contraindication or intolerance to ACEI/ARB.<sup>19</sup> Differences exist between international consensus guidelines for ARNI indications in HFrEF, which might affect real-world patient groups who will be prescribed this drug. Also, concerns exist regarding the generalizability of the results of PARADIGM-HF to the entire real-world HFrEF patients, as the PARADIGM-HF study population represents

only a minority of the real-world HFREF population. Several researchers investigated the differences in eligibility of sacubitril/valsartan initiation in the real-world HFREF population based on FDA labeling versus PARADIGM-HF enrollment criteria.

Parikh et al evaluated patient population from the Get With The Guidelines-Heart Failure (GWTG-HF) registry to characterize patients' eligibility and potential barriers for sacubitril/valsartan initiation according to the criteria set forth in FDA labeling and PARADIGM-HF.<sup>46</sup> The GWTG-HF registry is composed of patients who were admitted for worsening HF or developed significant HF symptoms during a hospitalization.<sup>46</sup> Among 28,932 hospitalizations of HFREF patients, 20,083 (69%) patients were eligible for sacubitril/valsartan initiation based on FDA labeling and 11,018 (38%) based on PARADIGM-HF criteria (including natriuretic peptide cutoffs), as shown in Table 4.<sup>46</sup> Of the 20,083 who were eligible based on FDA criteria, only 55% would have been eligible for sacubitril/valsartan initiation based on

PARADIGM-HF criteria. Based on ESC guidelines, this number would go down further, as only 41% patients in the PARADIGM-HF eligible group were on an MRA. The most common reasons for ineligibility (using PARADIGM-HF criteria) were: LVEF >35%, discharge SBP <100 mmHg, and not being on ACEI/ARB therapy.<sup>21</sup> Patients eligible for sacubitril/valsartan based on PARADIGM-HF criteria had significantly lower 30-day and 1-year mortality rates, compared to those who were excluded based on FDA-labeling or PARADIGM-HF entry criteria.<sup>46</sup>

Perez et al performed a single-center chart review of 210 NYHA class II–IV patients admitted to their HF service and had post-discharge follow-up, to determine the eligibility of sacubitril/valsartan initiation using FDA approval criteria and PARADIGM-HF trial enrollment criteria.<sup>47</sup> Of the 149 (71% total) patients who were eligible based on FDA labeling, only 54 (26% total) were eligible based on PARADIGM-HF enrollment criteria (Table 4).<sup>47</sup> The authors found that the patients who did not meet PARADIGM-HF enrollment

**Table 4** Studies evaluating the real-world eligibility of sacubitril/valsartan in HFREF patients

Population	HFREF patients who were admitted for worsening HF or developed significant HF symptoms during a hospitalization (from The Get With The Guidelines-Heart Failure Registry)	NYHA class II–IV HF patients who were discharged from the HF services of the Cleveland Clinic and had post-discharge follow-up	Ambulatory NYHA class II–IV chronic HFREF patients in the Swedish Heart Failure Registry (SwedeHF)	Chronic HFREF patients with available NT-proBNP measurements and on target doses ACEI/ARB referred to a community HF clinic in the UK
Total no. of patients	28,932	210	12,914	1,396
Number of patients meeting FDA labeling (%) for sacubitril/valsartan initiation	20,083 (69%)	149 (71%)	6,452 (50%)	–
Number of patients meeting PARADIGM-HF criteria (%)	11,018 (38%)	54 (26%)	5,015 (39%)	172 (21%)
Ratio of PARADIGM-HF like patients to FDA labeling eligible patients	55%	37%	78%	–
Reason for exclusion based on FDA labeling	Existing contraindication to ACEI/ARB; low discharge SBP	N/A	Not on at least enalapril 10 mg daily or equivalent	–
Reason for exclusion from PARADIGM-HF	LVEF 35%–40%, low discharge SBP <100 mmHg	SBP ≤100 mmHg, eGFR ≤30 mL/min/1.73 m <sup>2</sup> , not on ACEI or ARB, not on beta-blocker, serum K ≥5.2 mmol/L	Low natriuretic peptide level, SBP <100 mmHg, eGFR ≤30 mL/min/1.73 m <sup>2</sup> , serum K >5.2 mmol/L	Not on target dosing of ACEI/ARB, lack of limiting symptoms, plasma NT-proBNP <600 ng/mL

**Abbreviations:** ACEI, angiotensin-converting enzyme inhibitor; ARB, angiotensin receptor blocker; eGFR, estimated glomerular filtration rate; FDA, US Food and Drug Administration; HF, heart failure; HFREF, heart failure with reduced ejection fraction; LVEF, left ventricular ejection fraction; NT-proBNP, N-terminal pro-brain natriuretic peptide; NYHA, New York Heart Association; PARADIGM-HF, Prospective Comparison of Angiotensin Receptor-Nephrilysin Inhibitor with Angiotensin-Converting Enzyme Inhibitor to Determine Impact on Global Mortality and Morbidity in Heart failure; SBP, systolic blood pressure.

criteria were sicker with higher NYHA functional class, lower SBP, greater NT-proBNP level, lower eGFR, and they were less likely to be on ACEI, beta-blocker, and an MRA.<sup>47</sup> These findings reiterate the significant evidence gap with regards to FDA labeling of sacubitril/valsartan and proves concerning for the safety of sacubitril/valsartan as it can be used now in real-world HF population which was not represented in the PARADIGM-HF trial.

The above-mentioned analyses are primarily in hospitalized patients, whereas PARADIGM-HF enrolled ambulatory patients. European investigators analyzed registry data to establish the eligibility of ambulatory patients with HFrEF for treatment with sacubitril/valsartan. Simpson et al analyzed the Swedish Heart Failure Registry (SwedeHF) to determine the eligibility of patients with HFrEF for treatment with sacubitril/valsartan, according to the criteria used in PARADIGM-HF (Table 4).<sup>48</sup> They identified 12,914 ambulatory patients with symptomatic (NYHA class II–IV) HFrEF (LVEF <40%), of which 6,452 were treated with at least enalapril 10 mg daily or equivalent.<sup>48</sup> Of these 6,452 patients treated with enalapril 10 mg daily or equivalent, 5,015 (77.7%) were eligible for treatment with sacubitril–valsartan using PARADIGM-HF inclusion criteria and the most common reason for ineligibility was low natriuretic peptide level (17.6%).<sup>48</sup> Very few ambulatory symptomatic HFrEF patients on enalapril were ineligible due to a low eGFR of <30 mL/min/1.73 m<sup>2</sup> (87/6,452; 1.3%) or serum potassium >5.2 mmol/L (51/6,452; 0.7%).<sup>48</sup> Based on these findings, 38.8% of the ambulatory SwedeHF patients with symptomatic HFrEF were eligible for sacubitril/valsartan initiation. The findings from this study make sense in a broader perspective, as ambulatory HF patients are less sick than hospitalized patients and have lower natriuretic peptide levels, higher eGFR, and lower serum potassium levels.

These findings are in contrast with those reported by Pellicori et al who studied patient population in a community HF referral clinic in the UK between 2001 and 2014.<sup>49</sup> They identified 1,396 chronic HFrEF patients with LVEF ≤40% and available NT-proBNP measurements (Table 4).<sup>49</sup> Of the 379 patients (27% of total) who were on target dose of ACEI/ARB, only 172 patients (45% of 379) were eligible for sacubitril–valsartan initiation based on PARADIGM-HF enrollment criteria.<sup>49</sup> Lack of symptoms and low NT-proBNP <600 ng/L levels were the most common reasons for ineligibility.<sup>49</sup> A further 122 patients became eligible during follow-up. Based on this study, in a community-based chronic HFrEF population, only 21% (294/1,396) were eligible for sacubitril/valsartan initiation based on PARADIGM-HF enrollment

criteria, a number that increased to 60% (701/1,396) when background medications were ignored.<sup>49</sup> However, it should be remembered that being on an ACEI/ARB is a requirement prior to ARNI initiation. It is also important to note that this study is based on a single referral HF clinic, whereas the SwedeHF included a broader HFrEF population, which can explain the difference in the proportions of ambulatory patients who were eligible for sacubitril/valsartan initiation.

Zeymer et al compared PARADIGM-HF enrollment criteria with retrospective data from three German chronic HFrEF (LVEF <40%) registries/studies and reported that patients enrolled in PARADIGM-HF trial represented about 50% of the general population of patients with HFrEF in these registries.<sup>50</sup> It is interesting to note that only 5.3%–20.4% patients in the registries were treated with enalapril. Overall, it is evident that the patient population studied in the PARADIGM-HF does not represent the entire real-world HFrEF population largely because of the lack of symptoms (NYHA class I), not being on an ACEI/ARB, or low levels of natriuretic peptide. It is obvious from these findings that sacubitril/valsartan cannot be prescribed for the entire HFrEF population, given that a few major exclusions in the trial represent a significant number of real-world HFrEF population. It is also apparent that the differences in FDA labeling and consensus recommendations may result in this ARNI being used in a larger HFrEF population than it is evaluated in.

## Conclusion

Although the PARADIGM-HF trial provided compelling evidence for the superiority of angiotensin receptor-NEP inhibition over enalapril in HFrEF patients, it is important to note that certain subgroups of patients who represent a significant proportion of the real-world HF population were underrepresented or excluded from the original trial. Real-world eligibility data suggest that only 20%–40% of the HFrEF patients will be eligible for sacubitril/valsartan initiation based on current guidelines. The differences in FDA labeling and international consensus recommendations may result in this ARNI being used in a larger HFrEF population than it is studied in or recommended for, raising important safety concerns. It is crucial to clarify this evidence gap to prevent unwarranted complications in HFrEF patients in whom sacubitril/valsartan will be prescribed. Variations in inter-consensus guidelines must be also clarified, as these differences might affect patient selection. Despite these reservations, sacubitril/valsartan is an important breakthrough for the management of HFrEF and should be used in appropriately selected patients to provide morbidity and mortality



reduction. Available evidence suggests that sacubitril/valsartan might have a metabolic and nephroprotective benefit in HFrEF patients and proposes a significant CV benefit of this ARNI, irrespective of the type or etiology of HF. Ongoing trials testing this combination across various comorbidity profiles will expand the current indications of this game-changer medication in the management of HF.

## Disclosure

The authors report no conflicts of interest in this work.

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