

# Clinical outcomes after onset of heart failure with left ventricular ejection fraction ≥40% in the US

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## Introduction

- Heart failure (HF) with preserved ejection fraction is acknowledged as the area of CV medicine that has the largest unmet need.
- In the recent FINEARTS–HF trial, finerenone reduced the risk of worsening HF events/cardiovascular (CV) death in HF patients with LVEF ≥40%.
- We aimed to describe the profile of patients with HF who might be eligible for future treatment with finerenone, their healthcare resource use (HCRU), mortality, and CV outcomes.

## Methods

- We conducted a retrospective cohort study using the US Optum® de-identified electronic health record (EHR) dataset.
- We included patients aged ≥18 years with a first inpatient or outpatient HF diagnosis (ICD-10 codes) from Jan 2020 to Dec 2023. Patients were required to have ≥1 LVEF value of 5–95% primarily derived using natural language processing of unstructured physician notes within ±90 days of the first HF code. The index date was the date of first HF diagnosis or index LVEF value, whichever came second.
- We excluded patients with <365 days continuous data before the index date and those with a previous record of heart transplant or semi-permanent ventricular assist device.
- Patients were followed from the index date until 31 Dec 2023, 1 year after follow-up, use of a ventricular assist device or heart transplant, or death, whichever came first.
- We evaluated patient characteristics in the year before the index date, and HCRU, all-cause mortality, CV outcomes in the first year of follow-up. This was undertaken for the base cohort (LVEF ≥40%) and the following four subcohorts:

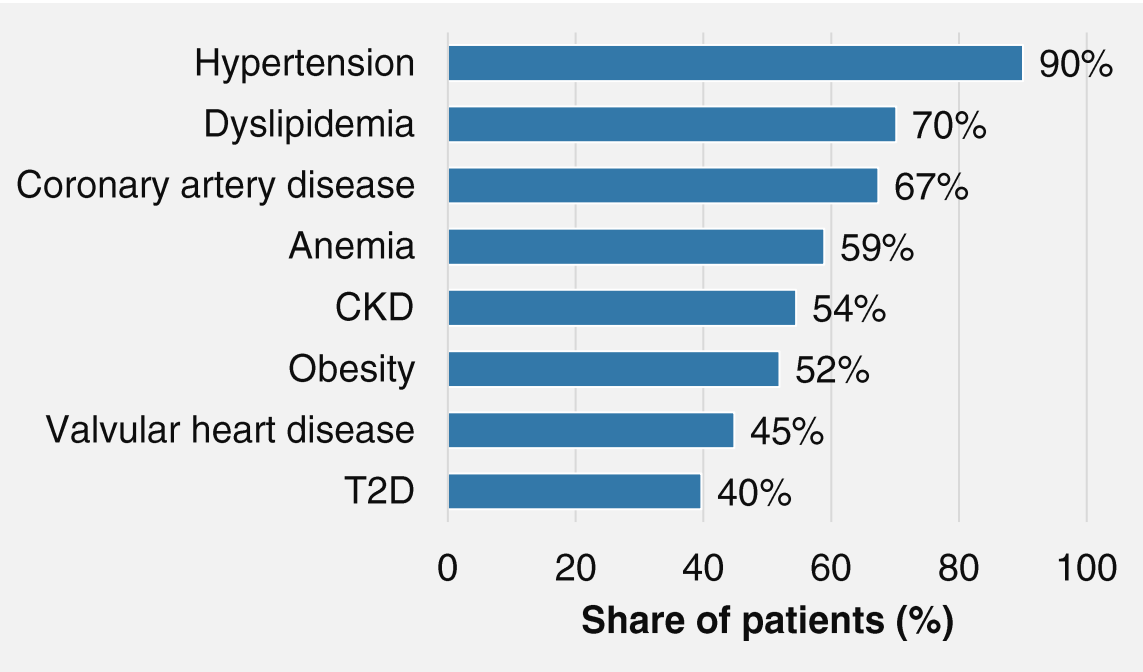
	Eligibility criteria
Patients with T2D	Application of diabetes phenotyping algorithm <sup>a</sup> , and classification of T1D/T2D using a combination of ICD-10 codes + abnormal lab result or prescription for a diabetes-targeting medication, in the year before the index date.
Patients with CKD	ICD-10-CM code for CKD or two CKD qualifying events <sup>b</sup> ≥90 days apart in the year before the index date.
Patients with T2D + CKD	Patients meeting the T2D and CKD criteria above.
Patients meeting the FINEARTS–HF trial criteria	Aged ≥40 years at the index date, ≥1 eGFR value in the year before the index date, index eGFR value ≥25ml/min/1.73m <sup>2</sup> .

<sup>a</sup> Developed by Sun & Hernandez-Boussard for extraction of diabetes patients from EHRs.  
<sup>b</sup> Clinical lab values of either 25 ≤ eGFR < 90 ml/min/1.73m<sup>2</sup> or 30 mg/g ≤ UACR ≤ 5000 mg/g.

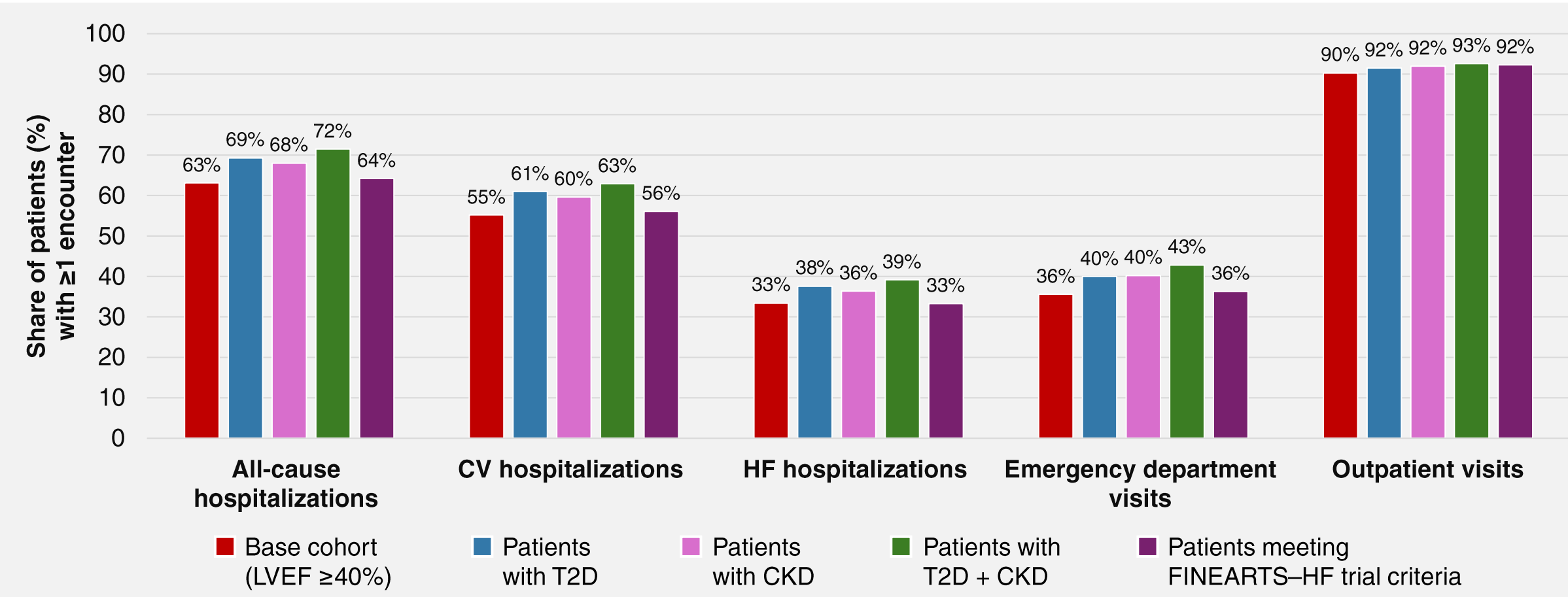
## Results

### Patient characteristics

- The base cohort included a total 201,343 patients with incident HF and LVEF ≥40%; median (Q1, Q3) age at the index date was 73 (63, 81) years; 50% were male. Race distribution was 79% White, 14% African–American, 2% Asian, 5% missing.
- Median (Q1, Q3) follow-up was 11.1 (3.2, 23.5) months.
- Baseline treatments (in the year before the index date) included steroidal MRAs (5%) and SGLT2is (3%).
- Prevalence of baseline comorbidities was high (**Fig 1**).



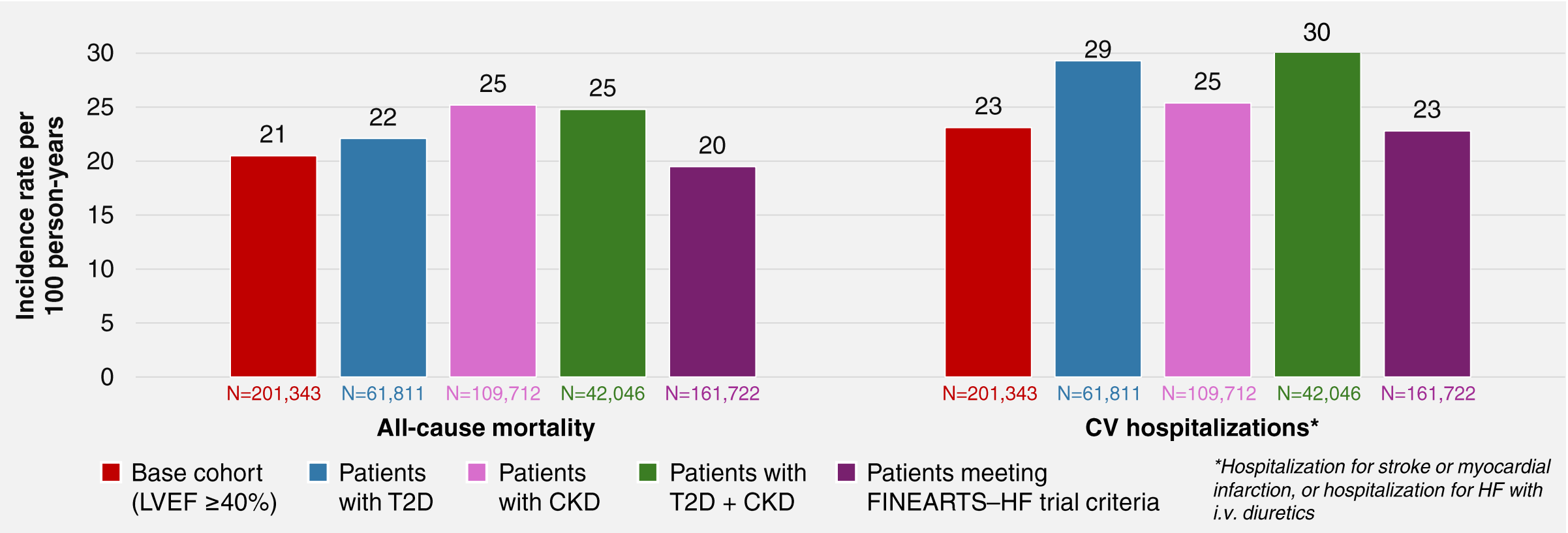
**Fig 1.** Comorbidities (in the year before the index date) among the base cohort of HF patients (LVEF ≥40%).



**Fig 2.** Healthcare resource use in the first year after HF diagnosis.

### Healthcare resource use

- During the first year after HF diagnosis, 63% of patients had an all-cause hospitalization, 55% had a CV hospitalization, and 33% had a HF hospitalization – higher rates were observed in patients with CKD and T2D at baseline (**Fig 2**).
- The mean (SD) number of days hospitalized was 7 (±13); 5.7 (±11) for CV hospitalizations, and 3 (±8) for HF hospitalizations.



**Fig 3.** All-cause mortality rates and incidence rates of CV hospitalizations in the first year after HF diagnosis.

### All-cause mortality and CV outcomes

- A total of 14% of patients died within the first year of follow-up, corresponding to a mortality rate of 21 per 100 person-years (**Fig 3**).
- Mortality rates and incidence rates of CV events were highest among the CKD and CKD + T2D subcohorts (**Fig 3**).

## Conclusion

- These findings from real-world clinical practice in the US suggest that patients with newly-diagnosed HF and LVEF ≥40% have high HCRU and a high risk of CV events, particularly those with comorbid CKD.
- HCRU and incidence rates of CV outcomes were higher in newly-diagnosed HF patients with pre-existing CKD and T2D.
- The care and outcomes of these patients could potentially be improved with new available treatments.

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