

Clinical outcomes after onset of heart failure with left ventricular ejection fraction $\geq 40\%$ 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 $\geq 40\%$.
- 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 $\geq 40\%$) and the following four subcohorts:

Eligibility criteria	
Patients with T2D	Application of diabetes phenotyping algorithm ^a , 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 ^b ≥ 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 $\geq 25 \text{ ml/min}/1.73\text{m}^2$.

^aDeveloped by Sun & Hernandez-Boussard for extraction of diabetes patients from EHRs.

^bClinical lab values of either $25 \leq \text{eGFR} < 90 \text{ ml/min}/1.73\text{m}^2$ or $30 \text{ mg/g} \leq \text{UACR} \leq 5000 \text{ mg/g}$.

Results

Patient characteristics

- The base cohort included a total 201,343 patients with incident HF and LVEF $\geq 40\%$; 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).

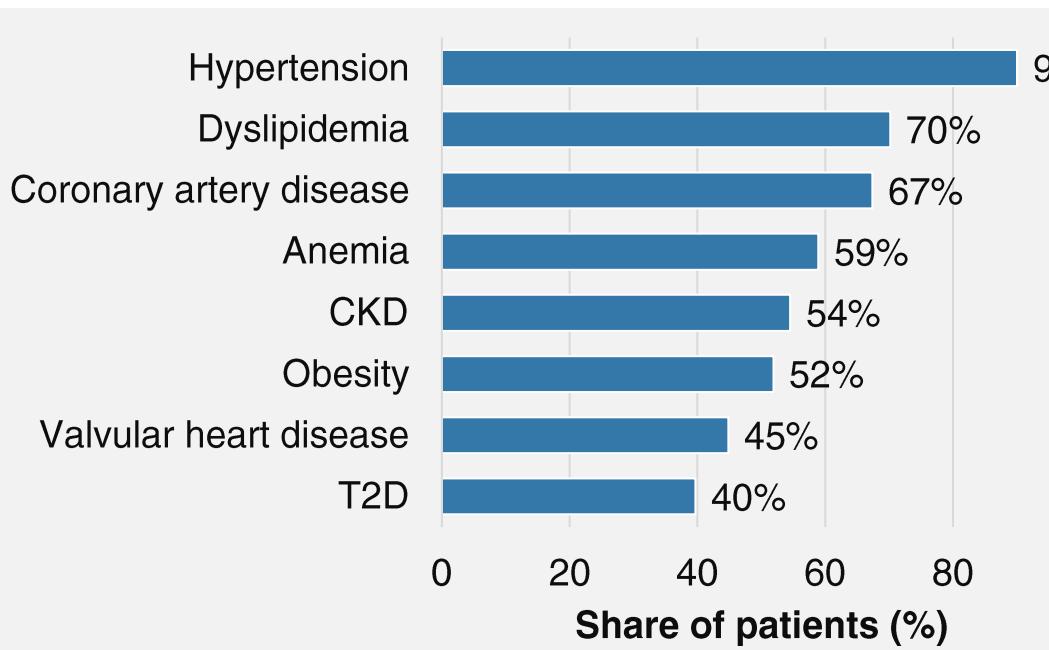


Fig 1. Comorbidities (in the year before the index date) among the base cohort of HF patients (LVEF $\geq 40\%$).

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.

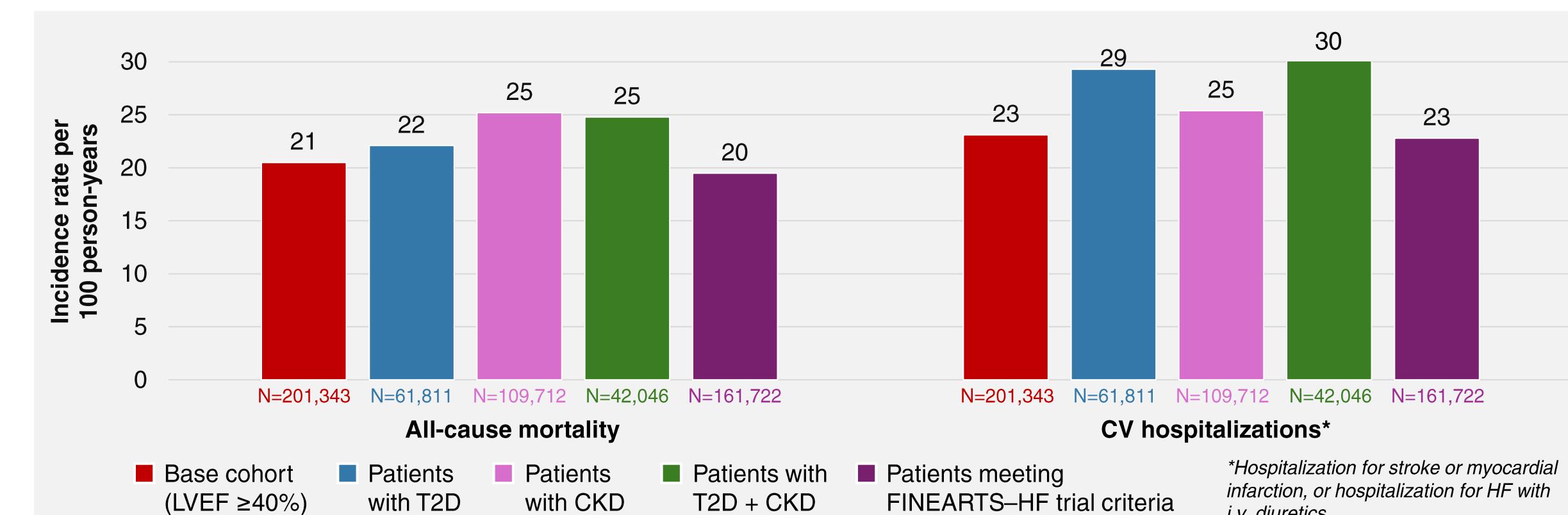


Fig 3. All-cause mortality rates and incidence rates of CV hospitalizations in the first year after HF diagnosis.

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).

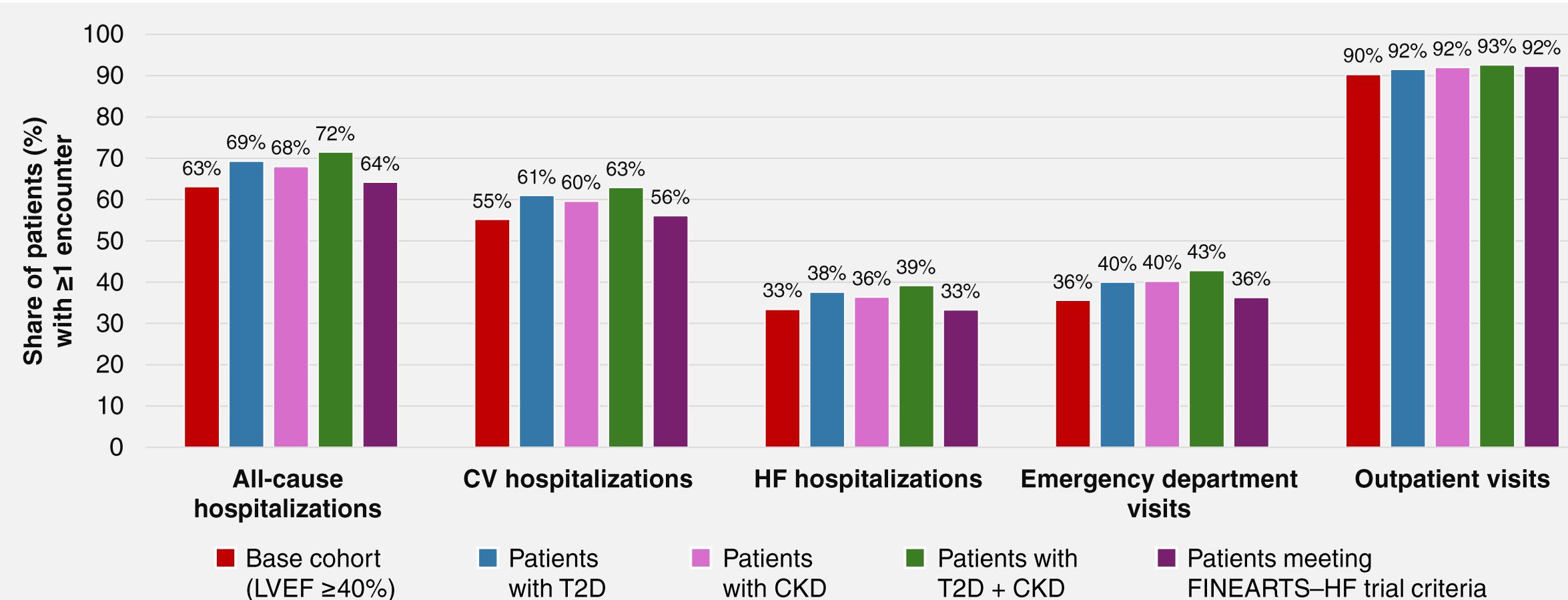


Fig 2. Healthcare resource use in the first year after HF diagnosis.

Conclusion

- These findings from real-world clinical practice in the US suggest that patients with newly-diagnosed HF and LVEF $\geq 40\%$ 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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