Association between heart failure medication at discharge and heart failure readmission

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Purpose

In selected groups of heart failure (HF) patients, angiotensin-converting-enzyme inhibitors (ACEI), angiotensin-receptor blockers (ARB), beta-blockers (BB) and mineralocorticoid-receptor antagonists (MRA) reduce the risk of readmission for HF. Diuretics, are indispensable for most HF patients.

Aim of this large Dutch population-based cohort study is to compare hospital readmission rates of patients with HF, prescribed core HF medications versus nonuse in a real-world scenario.

Methods and results

Medication at hospital discharge based on dispensing data from the **Dutch PHARMO Database Network including 22,476 patients with a diagnosis of HF between 2001 and 2015**.

Median follow-up was 29.3 months.

One third of patients were readmitted for HF. Propensity scores were calculated as a proxy for comorbidities and hazard ratios were adjusted (HRadj) accordingly.

ACEI and ARB were not associated with readmission. Only β 1-selective BB (sBBHF: bisoprolol, metoprolol and nebivolol) decreased risk of readmission (HRadj 0.94; 95%CI 0.90-0.99). Carvedilol, a β - and partly α 1-blocking agent (HRadj 1.25; 95%CI 1.14-1.38), MRA (HRadj 1.09; 95%CI 1.03-1.15) and diuretics (HRadj 1.14; 95%CI 1.06-1.23) were associated with an increased risk of readmission (Fig.).

Fig. Hazard ratios* of heart failure readmission and heart failure medication



*HR versus non-use; adjusted for age, gender, number of other drugs, year of admission, propensity score of particular medication (based on baseline covariates and co-medication); sBBHF: selective beta1-receptor blocker with a market autorisation for HF

Baseline characteristics

Total number of patients	22,476	
Patients readmitted %	29.9%	
Age: mean (SD)	76.8 (10.9)	
Gender: female %	50.9%	
ength of stay in hospital (days)	6 (3-11)	
Medication profile on discharge		

ACEI/ARB	62.7%
• ACEI	47.1%
• ARB	17.3%
Beta-blocker	59.6%
• sBBHF	46.0%
Carvedilol	5.4%
MRA	37.0%
Diuretics	81.8%

Conclusions

Selective beta-blockers for HF might be preferred to the nonselective beta-blocker carvedilol whereas these are considered equivalent in current guidelines. Further investigations are necessary to confirm our results in other real-world HF cohorts.

