



# Sodium-glucose cotransporter-2 inhibitors after heart transplantation

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# **BACKGROUND**

Sodium-glucose cotransporter-2 inhibitors (**SGTL2i**): dapagliflozin and empagliflozin, are widely used to manage adult patients with type 2 diabetes mellitus (DM), heart failure (HF), in all ejection fraction subgroups, and slowing the progression or reducing the risk failure of kidney disease.

In recent years, safety and efficacy have been published on their use in kidney transplant patients, however, there is no evidence in heart transplant (HT) recipients.

# **PURPOSE**

To evaluate safety, tolerability and effectiveness of SGTL2i in HT recipients.

#### MATERIAL AND METHODS

- Retrospective descriptive cohort study
- Patients included: All adults undergoing HT from January 2016 to July 2023 treated with SGLT2i in a tertiary hospital
- Demographic, clinical and pharmacological data were recorded.
- Outcome measures: Body Mass Index (BMI), glycated hemoglobin (HbA1c) and Creatinine at baseline and 6 months, adverse events and number of hospitalizations in patients with HF after SGLT2i initiation
- <u>Statystical analysis</u>: Wilcoxon test to compare the evolution of efficacy variables (BMI, HbA1c, creatinine) at baseline and 6 months after starting glyflozines was applied

#### **RESULTS**

# Demographic, clinical and pharmacological results

154 HT recipients were on SGLT2i



**Age**: 62.1 [50.9 – 63.4] years old

Dapagliflozin: 9 (32.1%)
Empagliflozin: 19 (67.9%)



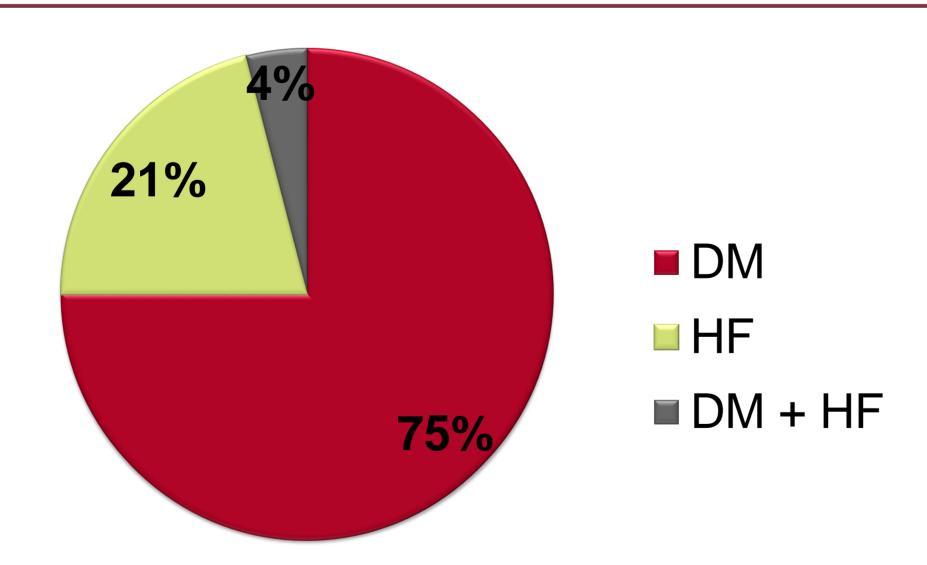


Figure 1. Indications of SGLT2i

# Effectiveness

Variable	Median (IQR)	p-value
HbA1c	-0.85 (-6.10; 0.00)	< 0.001
BMI	1.36 (-2.26; 0.80)	0.064
Creatinine	3.00 (-11.00; 33.00)	0.402

 Table 1: Wilcoxon Test comparing variables at baseline and at six months.

The non-parametric test for paired data showed that only HbA1c was significantly reduced after SGLT2i initiation.

## Safety and tolerability

Adverse Events: 3 (10.7%) patients:

- ✓ 2 patients suffered urinary tract infections.
- ✓ 1 patient suffered head instability.

2 patients discontinued SGTL2i, one after 4 months due to intolerance and the other after 11 months because of HbA1c normalization.

22 (78.6%) were DM (DM and DM+HF)

> 18 (81.8%) patients were on a combined hypoglycemic therapy.

> 7 (25.0%) patients developed it after HT.

## CONCLUSIONS

Our results show that SGTL2i are well-tolerated in HT recipients. Although these data are consistent with findings in renal recipients<sup>1</sup>, further investigation is needed.



