



SGLT2 INHIBITORS: CARDIORENAL BENEFIT SUMMARY



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	CARDIOVASCULAR	HEART FAILURE	RENAL
CANAGLIFLOZIN	<p>FDA-APPROVED BENEFIT ★</p> <p>Evidence: Lower rate of MACE composite</p> <ul style="list-style-type: none"> Individual components NSD <p>Population: T2DM with ASCVD or mult. RFs</p> <p>Trial: CANVAS Program (2017)</p>	<p>EVIDENCE OF BENEFIT ★</p> <p>Evidence: Lower rate of HF hospitalization</p> <ul style="list-style-type: none"> Not primary outcome of trial <p>Population: T2DM with ASCVD or mult. RFs</p> <p>Trial: CANVAS Program (2017)</p>	<p>FDA-APPROVED OF BENEFIT ★</p> <p>Evidence: Lower rate of renal composite</p> <ul style="list-style-type: none"> Primary benefit slowing CKD progression Mortality benefit not demonstrated <p>Population: T2DM with CKD plus albuminuria receiving ACEi or ARB therapy</p> <p>Trial: CREDENCE (2019)</p>
DAPAGLIFLOZIN	<p>NEUTRAL EFFECT ★</p> <p>Evidence: Similar rate of MACE composite</p> <ul style="list-style-type: none"> Individual components NSD <p>Population: T2DM with ASCVD or mult. RFs</p> <p>Trial: DECLARE-TIMI 58 (2019)</p>	<p>FDA-APPROVED BENEFIT ★</p> <p>Evidence: Lower rate of HF hospitalization and cardiovascular death composite</p> <ul style="list-style-type: none"> Morbidity and mortality benefit w/ HFrEF Only morbidity benefit w/ HFpEF <p>Population: Heart failure with/without T2DM</p> <p>Trials: DAPA-HF (2019) and DELIVER (2022)</p>	<p>FDA-APPROVED OF BENEFIT ★</p> <p>Evidence: Lower rate of renal composite</p> <ul style="list-style-type: none"> Primary benefit slowing CKD progression Mortality benefit not demonstrated <p>Population: CKD (with/without T2DM) plus albuminuria receiving ACEi or ARB therapy</p> <p>Trial: DAPA-CKD (2020)</p>
EMPAGLIFLOZIN	<p>FDA-APPROVED BENEFIT ★</p> <p>Evidence: Lower rate of MACE composite</p> <ul style="list-style-type: none"> Mortality benefit demonstrated <p>Population: T2DM with ASCVD</p> <p>Trial: EMPA-REG OUTCOME (2016)</p>	<p>FDA-APPROVED BENEFIT ★</p> <p>Evidence: Lower rate of HF hospitalization and cardiovascular death composite</p> <ul style="list-style-type: none"> Only morbidity benefit demonstrated (HFrEF or HFpEF) <p>Population: Heart failure with/without T2DM</p> <p>Trials: EMPEROR-Reduced (2020) and EMPEROR-Preserved (2021)</p>	<p>FDA-APPROVED OF BENEFIT ★</p> <p>Evidence: Lower rate of renal composite</p> <ul style="list-style-type: none"> Primary benefit slowing CKD progression Mortality benefit not demonstrated <p>Population: CKD (with/without T2DM) plus albuminuria receiving ACEi or ARB therapy</p> <p>Trial: EMPA-KIDNEY (2023)</p>

Adapted from Table 9.2 of the 2025 American Diabetes Association Standards of Care

All mentioned evidence is sources from randomized-controlled trials comparing SGLT2i to placebo. Within each benefit group (e.g., cardiovascular, heart failure, renal) the included SGLT2is are given a ranking of bronze, silver, or gold based on the level of benefit demonstrated (e.g., neutral, morbidity only, morbidity and mortality). These rankings are subjective and are not a substitute for medical advice.

CKD: chronic kidney disease **HFpEF:** heart failure preserved ejection fraction **HF:** heart failure **HFrEF:** heart failure reduced ejection fraction **MACE:** major adverse cardiac events (e.g., cardiovascular death, myocardial infarction, and stroke)

NSD: not significantly different **Renal composite:** end-stage renal disease, sustained renal function decline from baseline or cardiovascular/renal death **RFs:** risk factors