

EMPA-KIDNEY

Empagliflozin in Patients With Chronic Kidney Disease

International, Randomized, Parallel-Group, Double-Blind, Placebo-Controlled Trial

OBJECTIVE: To evaluate the effect of empagliflozin treatment on the progression of kidney disease and CV disease and to examine the safety profile among participants with chronic kidney disease.

6,609
PATIENTS

INCLUSION CRITERIA:

- Race-adjusted eGFR of at least 20 but less than 45mL/minute/1.73m²
- Urinary albumin-to-creatinine ratio of at least 200 at screening visit
- Clinically appropriate dose of single agent RAS inhibitor with or without diabetes







PRIMARY ENDPOINT

The primary outcome, progression of kidney disease or death from CV cause, occurred in 13.1% in the empagliflozin group and 16.9% in the placebo group, p<0.001

SECONDARY ENDPOINT

For Empagliflozin vs. Placebo:

Hospitalization for heart failure or death from CV cause: 4.0% vs. 4.6%, p=0.15

Hospitalization for any cause:

24.8 events/100 patient-years vs. 29.2 events/100 person-years, p=0.003

Death from any cause: 4.5% vs. 5.1%, p=0.21

Progression of kidney disease: 11.6% vs. 15.2%, HR 0.71; 95% CI 0.62-0.81

CONCLUSION

Among patients with chronic kidney disease with or without diabetes, empagliflozin led to a lower risk of progression of kidney disease or death from CV cause than placebo.