Budget impact analysis of empagliflozin for adults with Chronic Kidney Disease (CKD) in Greece

Koulentaki M,¹ Vlahakos D,² Tsimihodimos V,³ Smyrnaios C,⁴ Delli E,⁴ Karpouzos G,⁴ Karathanou F,¹ Kourlaba G,⁵

1. Econcare Lp, Athens, Greece; 2. Scientific Advisor and President of the Scientific Council at Bioiatriki Group; 3. Department of Internal Medicine, School of Medicine, University of Ioannina, Ioannina, Greece; 4. Boehringer Ingelheim, Athens, Greece; 5. Faculty of Health Sciences, University of Peloponnese, Tripoli, Greece.

Objective

- \odot
- To investigate the budgetary impact from the introduction of empagliflozin as add-on treatment to Standard of Care (SoC) for adults with Chronic Kidney Disease (CKD) in Greece.



EE623

Scan the QR code to view the e-poster

Introduction

 CKD is characterized by kidney abnormalities lasting over 3 months, classified by cause, glomerular filtration rate (GFR), and albuminuria (CGA staging)¹.

Results

 The total number of the eligible population was 409,013 patients per year and the number of patients treated with empagliflozin+SoC in the new market scenario was estimated to increase from 4,090 in 2025 to 26,586 patients in 2029.



- Research shows low CKD awareness worldwide, leading to late-stage diagnoses and reduced treatment options²⁻⁴.
- Due to limited screening practices, approximately 8,236 per 100,000 individuals with CKD remain undiagnosed in Greece, with an increase of 3.7% over a five-year time horizon⁵.
- Global CKD prevalence is estimated at 9.1%, with stages three to five ranging from 4.1% to 10.6% ^{6,7}.
- Traditional CKD treatments include Angiotensin-converting enzyme (ACE) inhibitors, angiotensin-receptor blockers (ARBs), statins, and antiplatelets, which slow progression but do not halt it, with limited options until recent advancements.
- Sodium-glucose transport protein 2 (SGLT-2) inhibitors, including dapagliflozin, canagliflozin, and empagliflozin, now offer new treatment options, showing benefits in CKD management^{8,9}.
- The EMPA-KIDNEY trial¹⁰⁻¹² found empagliflozin reduced CKD progression and cardiovascular death to 13.1% compared to 16.9% with placebo, proving effective and safe.

Methods

- The evaluation of economic consequences is derived from the comparison between the economic burden of CKD in a world without empagliflozin, with a burden in a world with empagliflozin, over a five-year period (2025-2029).
- The total costs for both scenarios are calculated by multiplying the per-patient costs with the global number of patients, allowing estimation of the budget impact of introducing the new intervention to the market.
- Epidemiological data, as retrieved from the literature and local clinical experts estimates, were used to estimate the number of people with CKD eligible for treatment with empagliflozin (**Table 1**).

- Due to faster CKD progression in the SoC arm, patients deteriorate more quickly and incur higher costs, leading to increased annual total costs from year three onward compared to the dapagliflozin and empagliflozin arms (**Table 4**).
- Over the five-year horizon, the inclusion of empagliflozin led to a decrease in public expenditures, with an average annual total cost savings of €17,620,805 (Figure 1).
- Results of sensitivity analysis were found fairly insensitive.

Table 4. Budget impact results of all CKD patients (€)

	2025	2026	2027	2028	2029
Current Scenario	1,178,531,969	1,268,078,778	1,537,182,746	1,605,908,663	1,735,840,916
New Scenario	1,178,564,831	1,269,851,984	1,533,277,580	1,599,297,253	1,726,930,619
Total budget impact	32,862	1,773,206	-3,905,166	-6,611,409	-8,910,297
Cumulative budget impact	32,862	1,806,067	-2,099,099	-8,710,508	-17,620,805

Figure 1. Net and cumulative budget impact



- Market share of the analysis for both scenarios were calculated based on a local clinical experts' estimation (**Table 2**).
- The drug acquisition costs, costs of events and complications, and adverse event costs were derived from the literature and are detailed as total costs per category per treatment over a 5-year period (**Table 3**).
- A sensitivity analysis was performed, individually varying several model parameters and assumptions to assess the key drivers and robustness of the base case findings.

Table 1. Estimated target patient population

Parameter	Number of patients	Source
Prevalence of CKD (12.86%)	1,115,991	Local clinical experts estimation; Stafylas et al. ¹³
Diagnosed patients with CKD (61,86%)	690,317	Local clinical experts
Patients who are diagnosed and treated with SoC treatment (59%)	409,013	Local clinical experts
People who have type 2 diabetes (12.5%)	51,127	Local clinical experts; Kibria et al. ¹⁴ ;

Hounkpatin et al.¹⁵

Note: According to clinical experts estimates, 7.7% of CKD patients are Urine albumin-creatinine ratio (uACR ≥22.6 mg/mmol) (and without T2D)

Table 2. Market shares with and without empagliflozin in all CKD patients

Thoronoutio options	Current market (scenario without Empagliflozin)					
Therapeutic options	2025	2026	2027	2028	2029	
Empagliflozin+SoC	0.0%	0.0%	0.0%	0.0%	0.0%	
SoC	85.0%	80.0%	75.0%	70.0%	65.0%	
Dapagliflozin+SoC	15.0%	20.0%	25.0%	30.0%	35.0%	
Total	100%	100%	100%	100%	100%	
Thoropoutio options	Projected market (scenario with Empagliflozin)					
Therapeutic options	2025	2026	2027	2028	2029	
Empagliflozin+SoC	5.0%	10.0%	21.3%	25.0%	32.5%	
SoC	85.0%	75.0%	57.5%	50.0%	40.0%	
Dapagliflozin+SoC	10.0%	15.0%	21.3%	25.0%	27.5%	
Total	100%	100%	100%	100%	100%	

Conclusion

This budget impact analysis indicates that introducing empagliflozin for CKD treatment in Greece provides significant clinical benefits with manageable budget impacts in the first two years and appears cost-saving for the public payer over five years.

Abbreviations

ACEi, angiotensin-converting enzyme. inhibitors; ARBs, angiotensin-receptor blockers; CGA, Cause, GFR category and albuminuria category; CKD, Chronic Kidney Disease; EGFR, estimated GFR; ESKD, end-stage kidney disease; GFR, glomerular filtration rate; SGLT-2, sodium-glucose transport protein 2; SoC, Standard of care; T2D, Type 2 diabetes.

References

Table 3. Five year costs per cost category for CKD patients (€)

	Empagliflozin + SoC	SoC alone	Dapagliflozin
Monitoring	14,495	15,098	14,481
Treatment	5,401	1,270	5,343
Adverse events	1	0	1
Total kidney replacement therapy	2,381	6,184	2,343
Total End-stage kidney disease	708	740	716
Cardiovascular complications	9,221	9,054	9,200
Anemia	520	551	521
Other chronic kidney disease complications	1,708	1,765	1,703
Bone and mineral disorder	1,413	1,439	1,412
Acute kidney injury	300	348	297
Infections	650	633	650
Cancers	13	13	13

1. KDIGO.Org, KDIGO 2012 Clinical Practice Guideline for the Evaluation and Management of Chronic Kidney Disease. 2013. 3(1); 2. Hsiao, L.-L. (2018), Raising awareness, screening and prevention of chronic kidney disease: It takes more than a village. Nephrology, 23: 107-111; 3. Stolpe, S. et al. High Unawareness of Chronic Kidney Disease in Germany. Int J Environ Res Public Health. 2021 Nov 9;18(22):11752.; 4. Chu CD, et al. Patient Awareness of CKD: A Systematic Review and Meta-analysis of Patient-Oriented Questions and Study Setting. Kidney Med. 2021 Jun 1;3(4):576-585; 5. Chertow GM, et al. Projecting the clinical burden of chronic kidney disease at the patient level (Inside CKD): a microsimulation modelling study. EClinicalMedicine. 2024 May 2;72:102614.; 6. Bikbov, B. et al. Global, regional, and national burden of chronic kidney disease, 1990–2017: a systematic analysis for the Global Burden of Disease Study 2017. The Lancet 395, 709–733 (2020); 7. Reichel, H. et al. Chronic kidney disease progression and mortality risk profiles in Germany: results from the Chronic Kidney Disease Outcomes and Practice Patterns Study. Nephrology Dialysis Transplantation 35, 803–810 (2020); 8. Tuttle, K. R. et al. SGLT2 Inhibition for CKD and Cardiovascular Disease in Type 2 Diabetes: Report of a Scientific Workshop Sponsored by the National Kidney Foundation. Diabetes 70, 1–16 (2021); 9. CT.gov. EMPA-KIDNEY (The Study of Heart and Kidney Protection With Empagliflozin). Available from: <u>https://clinicaltrials.gov/ct2/show/study/NCT03594110</u>; 10. Herrington, W. G. et al. The potential for improving cardio-renal outcomes by sodium-glucose co-transporter-2 inhibition in people with chronic kidney disease: a rationale for the EMPA-KIDNEY study. Clin Kidney J 11, 749–761 (2018); 11. Herrington, W. G. et al. Design, recruitment, and baseline characteristics of the EMPA-KIDNEY trial. Nephrology Dialysis Transplantation 37, 1317–1329 (2022); 12. Empagliflozin in Patients with Chronic Kidney Disease. New England Journal of Medicine 388, 117–127 (2023); 13. Stafylas, M. et al. #5632 The clinical and economic burden of Chronic Kidney Disease in Greece. (2023).); 14. Hounkpatin, H. O. et al. Prevalence of chronic kidney disease in adults in England: comparison of nationally representative cross-sectional surveys from 2003 to 2016. BMJ Open 10, e038423 (2020).

Disclosure statement

- The authors meet criteria for authorship as recommended by the International Committee of Medical Journal Editors (ICMJE).
- The authors did not receive payment related to the development of the abstract/poster.
- Koulentaki M., MSc of Econcare Lp provided writing, editorial support, and formatting assistance, which was contracted and funded by Boehringer Ingelheim .
- Boehringer Ingelheim was given the opportunity to review the manuscript for medical and scientific accuracy as well as intellectual property considerations.
- The study was supported and funded by Boehringer Ingelheim

