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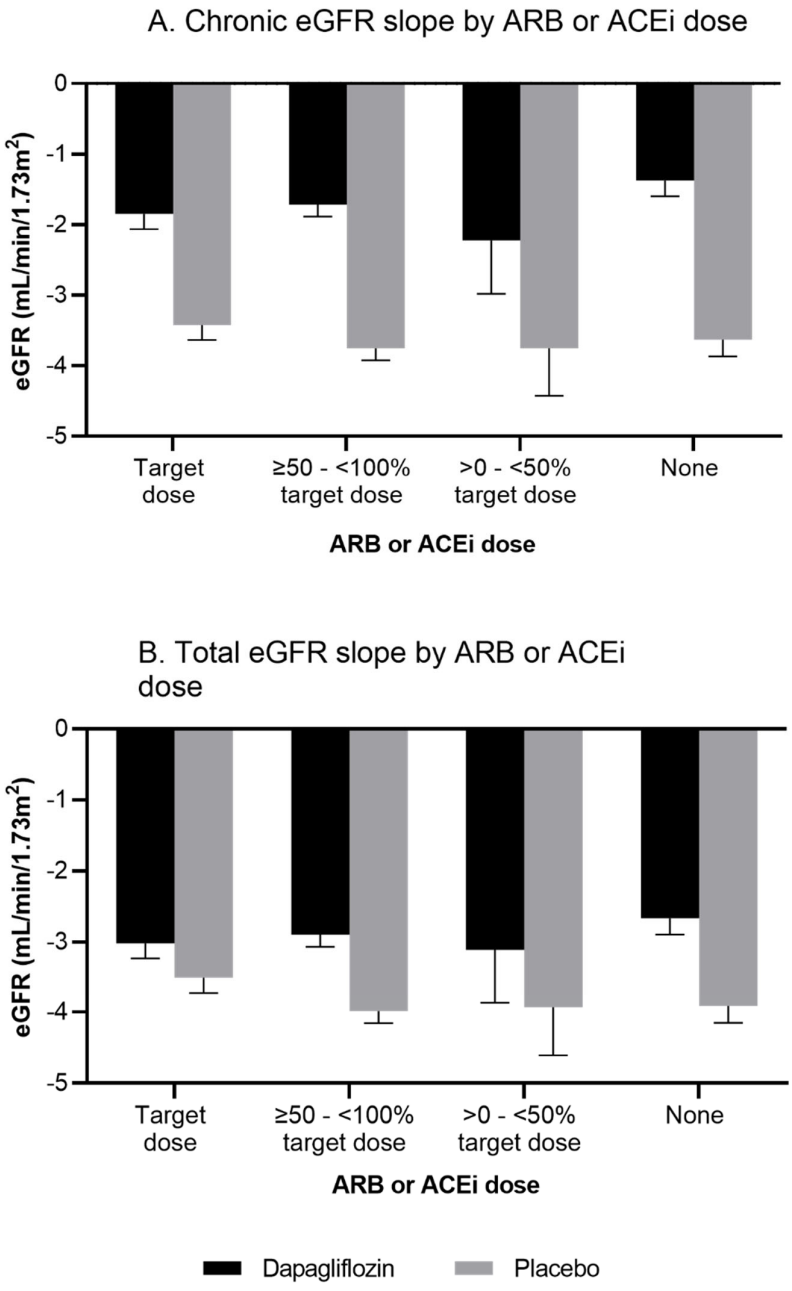
Table S1: Serious adverse events (SAEs) and adverse events (AEs) leading to discontinuation in patients stratified by ACEi/ARB dose

AE, n/N (%)	Dapagliflozin	Placebo
Any AE leading to discontinuation of the study drug		
None	3/64 (4.7)	8/74 (10.8)
<50%	24/547 (4.4)	22/517 (4.3)
50 to <100%	54/934 (5.8)	51/932 (5.5)
100% or more	37/604 (6.1)	42/626 (6.7)
Any SAE*		
None	22/64 (34.4)	29/74 (39.2)
<50%	161/547 (29.4)	161/517 (31.1)
50 to <100%	278/934 (29.8)	325/932 (34.9)
100% or more	172/604 (28.5)	214/626 (34.2)

*Includes death. Categories refer to the percent of maximum labelled antihypertensive dose.
ACEi, angiotensin-converting enzyme inhibitor; ARB, angiotensin receptor blockers.

Figure S1: eGFR decline between dapagliflozin 10 mg and placebo based in ARB or ACEi dose (A) Chronic slope, (B) Total slope



Chronic eGFR slope was determined from Week 2 to the end of the study; Total eGFR slope was determined over the entire study period

ACEi, angiotensin-converting enzyme inhibitor; ARB, angiotensin receptor blocker; eGFR, estimated glomerular filtration rate.