Supplementary material – Gritter et al.

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Figure S7: Correlations between the change in office systolic blood pressure (BP) with baseline blood pressure, urinary sodium (Na⁺) and potassium (K⁺) excretion, and estimated glomerular filtration rate (eGFR).

Table S1: Baseline characteristics associated with a smaller or lager increase in plasma

 potassium after KCl supplementation.

Variable	Univariable regression		Multivariable regression	
	β (95% CI)	P	β (95% CI)	P
Female sex	0.112 (-0.022, 0.246)	0.100	0.091 (-0.042, 0.224)	0.182
Type 2 diabetes mellitus	0.118 (-0.003, 0.239)	0.056	0.060 (-0.058, 0.177)	0.320
Renin-angiotensin system	0.079 (-0.077, 0.235)	0.322	0.175 (0.027, 0.323)	0.021
inhibitor use				
Beta blocker use	0.147 (0.028, 0.266)	0.015	0.110 (-0.005, 0.225)	0.062
Diuretic use	-0.110 (-0.229, 0.008)	0.068	-0.152 (-0.270, -0.035)	0.011
Age, per 10 years increase	0.069 (0.015, 0.123)	0.013	0.068 (0.010, 0.126)	0.021
Baseline plasma potassium, per	-0.036 (-0.098, 0.027)	0.264	-0.114 (-0.183, -0.045)	0.001
0.5 mmol/L increase				
Baseline plasma bicarbonate,	-0.022 (-0.039, -0.005)	0.011	-0.021 (-0.040, -0.002)	0.033
mmol/L				
Baseline eGFR, per 10	-0.090 (-0.157, -0.023)	0.008	-0.069 (-0.138, 0.001)	0.053
mL/min/1.73 m ² increase				
Baseline urine potassium, per	-0.041 (-0.064, -0.017)	0.001	-0.021 (-0.046, 0.003)	0.089
10 mmol/day increase				

Table S2: Treatment of participants with plasma potassium > 6.0 mmol/L after KCl supplementation.

Participant	Plasma potassium	Treatment	Plasma
	after 2 weeks KCl		potassium at
	supplementation		follow-up
5035	6.2 mmol/L	Sodium bicarbonate 3 x 1 g/day	4.9 mmol/L
		for 2 days	
7038	6.4 mmol/L	Sodium polystyrene sulfonate 2 x	3.9 mmol/L
		30 g/day for 3 days; temporary	
		discontinuation of losartan	
7041	6.9 mmol/L	Sodium polystyrene sulfonate 2 x	4.3 mmol/L
		30 g/day for 3 days; temporary	
		discontinuation of irbesartan	
8069	6.3 mmol/L	Sodium polystyrene sulfonate 1 x	4.9 mmol/L
		15 g/day for 3 days; temporary	
		discontinuation of lisinopril	
8103	6.9 mmol/L	Sodium polystyrene sulfonate 2 x	3.9 mmol/L
		30 g/day; temporary	
		discontinuation of lisinopril	

Figure S1: Flowchart of screened and included patients.

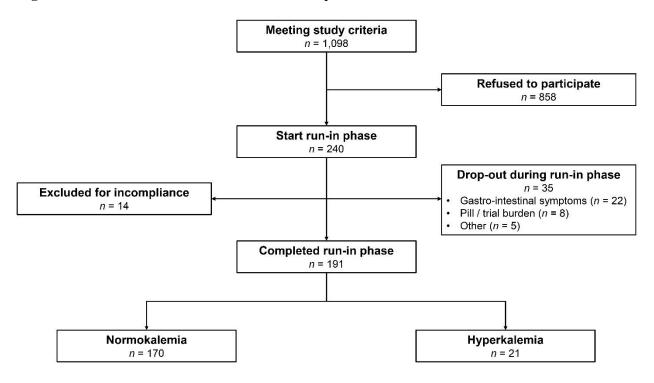


Figure S2: Change in urine potassium (K^+) excretion in participants with or without an increase in plasma K^+ concentration after KCl supplementation.

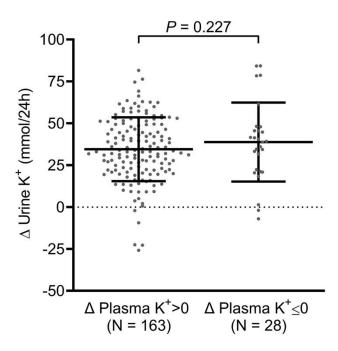


Figure S3: Change in plasma potassium (K^+) after KCl supplementation classified by sex, presence of diabetes mellitus, and the use of renin-angiotensin inhibitors, beta blockers, or diuretics.

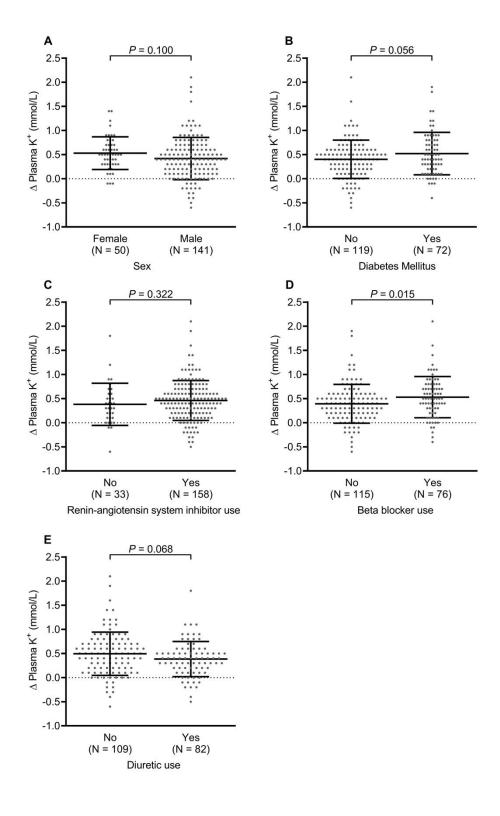


Figure S4: Correlations between the change in plasma potassium (K⁺) after KCl supplementation with age and selected baseline laboratory measurements.

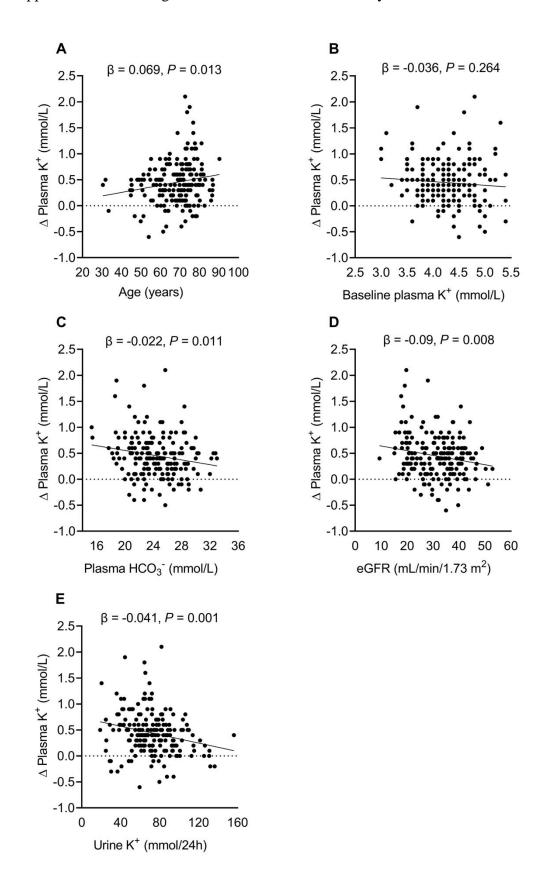


Figure S5: Exploratory analysis of baseline characteristics that were associated with a smaller or lager increase in plasma potassium after KCl supplementation for two weeks with the addition of ethnicity.

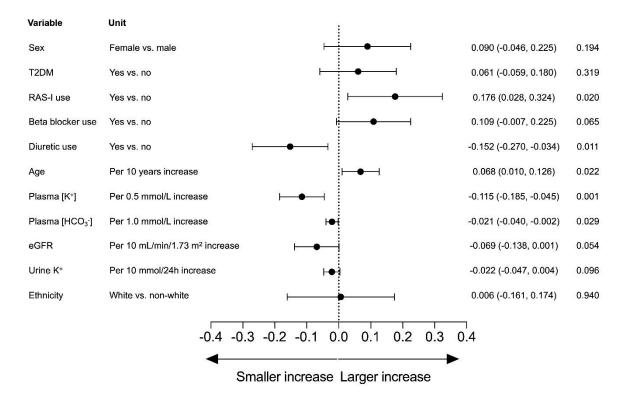


Figure S6: Change in urine potassium (K⁺) excretion in patients with or without hyperkalemia after KCl supplementation.

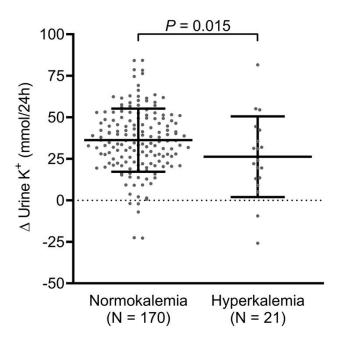
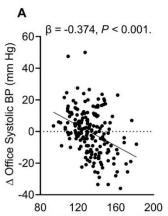
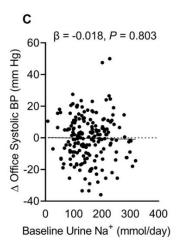
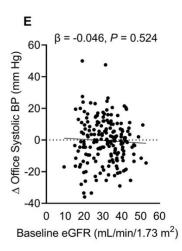


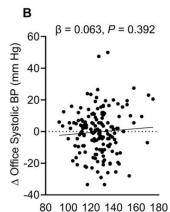
Figure S7: Correlations between the change in office systolic blood pressure (BP) after potassium chloride supplementation with baseline office and 24-hour systolic blood pressure, urinary sodium (Na⁺) and potassium (K⁺) excretion, and estimated glomerular filtration rate (eGFR).



Baseline Office Systolic BP (mm Hg)







Baseline 24h Systolic BP (mm Hg)

