

Healthcare Costs and All-cause Mortality Following a Hyperkalaemia Event and Reduction of RAASi Therapy in Sweden

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Maria K. Svensson,¹ Kun Kim,^{2,3} Thomas Cars,⁴ Naveen Rao,⁵ Eva Lesén,⁶ Krister Järbrink⁶

¹Department of Medical Sciences, Renal Medicine, Uppsala University, Uppsala, Sweden; ²Health Economics, AstraZeneca, Stockholm, Sweden; ³Department of Neurobiology, Care Sciences and Society, Karolinska Institute, Sweden; ⁴Sence Research AB, Uppsala, Sweden; ⁵Health Economics and Payer Evidence CVRM, AstraZeneca, Cambridge, UK; ⁶CVRM Evidence, AstraZeneca, Gothenburg, Sweden

Introduction

- Guidelines recommend use of renin-angiotensin-aldosterone system inhibitor (RAASi) therapy at the maximum tolerated dose to achieve optimal cardiorenal benefit in patients with chronic kidney disease (CKD) and/or heart failure (HF).¹⁻⁴
- Risk of hyperkalemia (HK) presents a barrier to achieving guideline-directed target dosing with RAASi; with RAASi often being discontinued or down-titrated.⁵
- Current international guidelines recommend the use of novel potassium binders to manage HK and facilitate RAASi therapy optimization in individuals with CKD and/or HF.¹⁻⁴

Aims

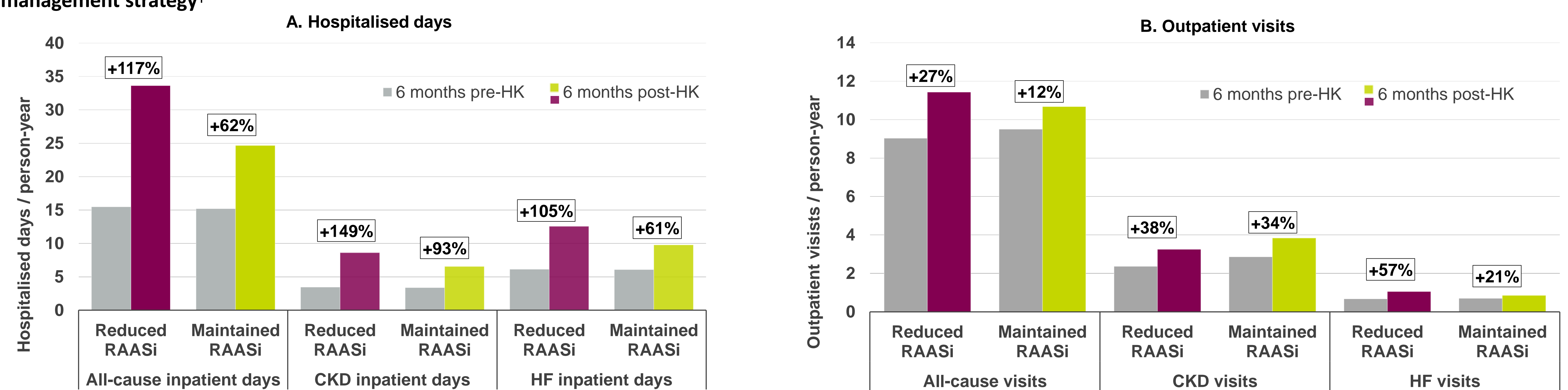
To assess healthcare costs and all-cause mortality associated with reduced RAASi treatment following a HK event in Swedish patients with CKD and/or HF.

Methods

- This observational study used data from national registries linked with health records from two large Swedish regions (Stockholm and Skåne).
- Inclusion criteria:
 - Patients aged ≥ 18 years with HF and/or CKD, and RAASi use at baseline.
 - Index HK event (ICD-10 code E87.5) or serum potassium >5.0 mmol/L during Mar 2018–Jul 2020.
- Propensity score matching (1:1) was used to balance the cohorts who reduced versus maintained their pre-HK RAASi treatment on demographics, comorbidities, baseline medications (incl. RAASi), and hospitalised days prior to index.
- Change in healthcare utilisation was measured as outpatient visits and hospitalised days (all-cause and related to CKD and HF) per person-year within 6 months before vs after the index HK event.
- Costs were based on national DRG-weights and 2021 years national cost in Swedish krona (SEK).
- Days Alive Outside Hospital (DAOH) was calculated for each individual patient, as total follow-up time (6 months) – (days in hospital + days beyond the date of death until 6 months after index). Patients who were lost to follow-up within these 6 months were excluded from these analyses.
- All-cause mortality within 6 months after index was analysed using the Kaplan-Meier method.

Results

Figure 1A and 1B. Hospitalised days (A) and outpatient visits (B) per person-year during 6 months before and 6 months after the index hyperkalaemia event, by RAASi management strategy[†]



The propensity score matched cohorts included 6998 patients each. [†]Patients were defined as having reduced or maintained their pre-HK RAASi therapy based on filled prescriptions, or lack thereof within 120 days pre- versus post HK index event. CKD, chronic kidney disease; RAASi, renin-angiotensin-aldosterone system inhibitor.

Figure 2. Healthcare costs 6 months before and after a hyperkalaemia event, per person-year

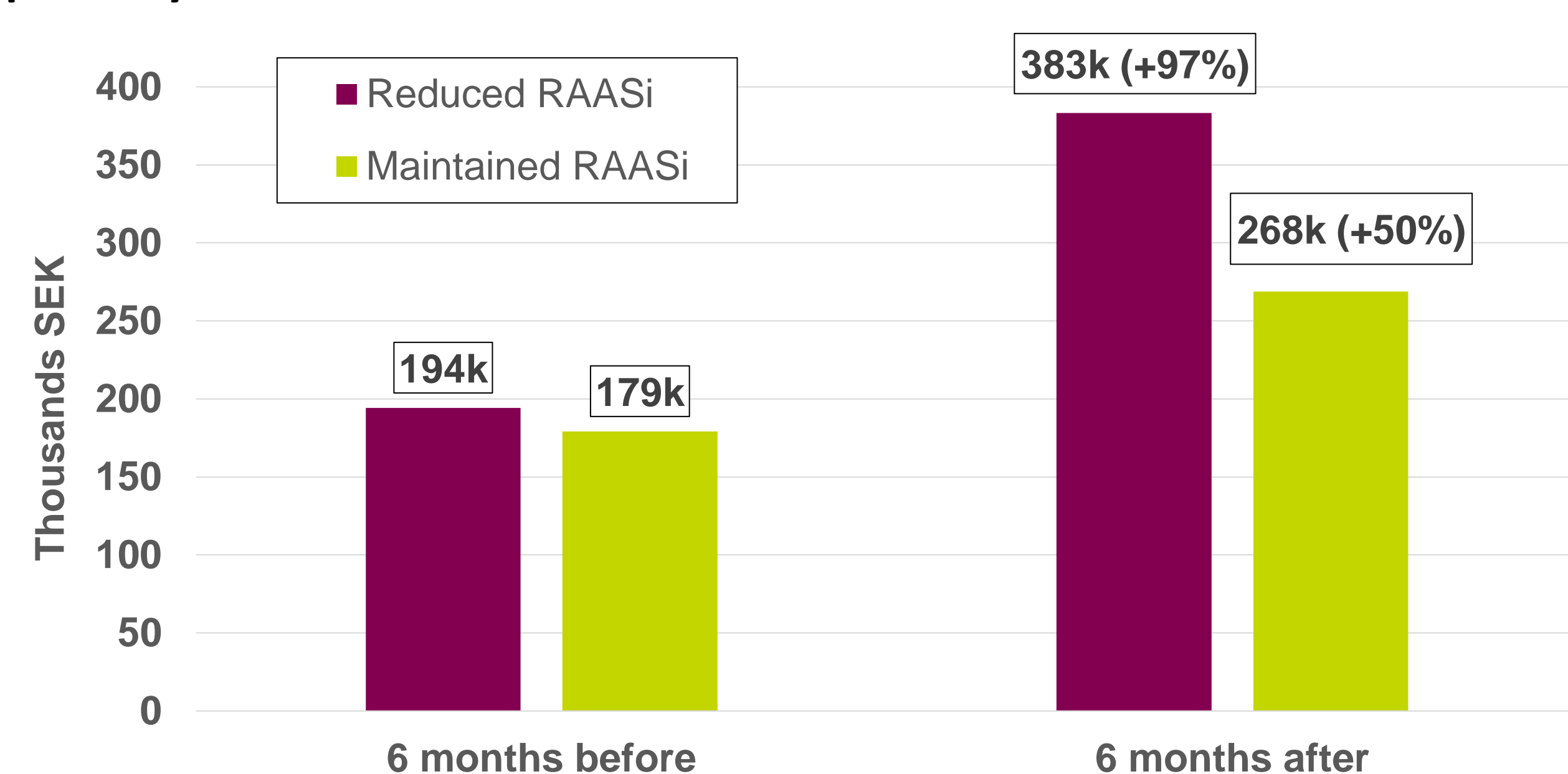
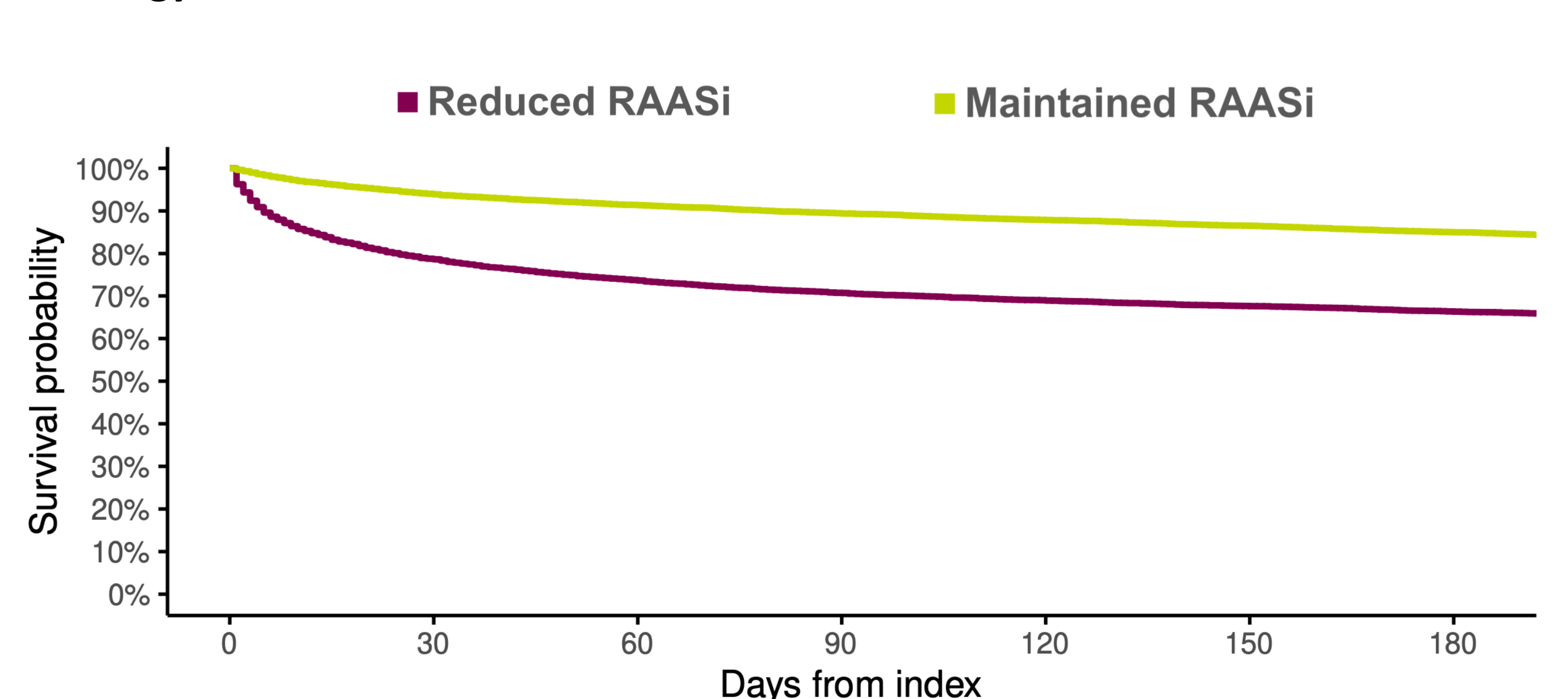


Figure 3. Survival within 6 months from the hyperkalaemia event, by RAASi treatment strategy



- Following matching, 6998 patients remained in each group (maintained vs. reduced RAASi) and the cohorts were well balanced (e.g. mean age 76.7 vs 76.8 years, female 42.9 vs 42.4%, mild HK 63.8 vs 61.0%, CKD stage 3 50.9 vs 51.5%, HF 58.9 vs 57.7% and diabetes 44.8 vs 45.1%).
- During the six months following the index HK event, all-cause inpatient bed-days increased by 117% among patients who reduced their RAASi treatment and by 62% among those who maintained compared to 6 months before the hyperkalaemia event (Figure 1A). Similarly, all-cause outpatient visits per person-year showed an increase of 27% in those who reduced RAASi vs. 12% in those who maintained (Figure 1B).
- The healthcare costs increased from 6 months before to 6 months after the hyperkalaemia event by 97% in those who reduced and with 50% in those who maintained RAASi treatment, corresponding to a 47% higher absolute difference (Figure 2).
- Patients who reduced RAASi were on average alive 121.5 (SD 75.0) days outside the hospital, compared to 154.0 (SD 51.3) in those who maintained RAASi.
- The survival rate at 6 months was 0.663 (CI: 0.652-0.674) among those who reduced their RAASi therapy compared to 0.850 (CI: 0.842-0.858) among those that maintained (Figure 3).

Conclusion

In clinical practice, reducing RAASi therapy after a hyperkalaemia event is associated with increased healthcare costs and an increased risk of all-cause mortality. These findings indicate a need for improved guidelines adherence in targeted management of HK to facilitate maintained RAASi therapy and achieve optimal treatment outcomes for patients with CKD or HF.

References

- de Boer IH et al. *Kidney Int.* 2020;98(4S):S1–S115.
- KDIGO: Diabetes Work Group. *Kidney Int.* 2022;102(5S):S1–S127.
- McDonagh TA et al. *Eur Heart J.* 2021;42(36):3599–3726.
- Heidenreich PA et al. *J Am Coll Cardiol.* 2022;79(17):e263–421.
- Kanda E et al. *BMC Nephrol.* 2023;24(1):18.

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Contact information: Krister.jarbrink1@astrazeneca.com