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Introduction

- Faricimab is a bispecific antibody targeting ANG-2 and VEGF for the treatment of neovascular age-related macular degeneration (nAMD).
- In the TENAYA and LUCERNE trials (2-year long Phase III trials),¹ patients treated with individualized dosing of faricimab followed by a Treat & Extend (T&E) regime required less frequent treatments compared to Aflibercept given every eight weeks (Q8W) and achieved similar vision gains.
- However, clinical practice in nAMD in Canada is typically characterized by T&E regimens. This research aims to assess the cost-effectiveness of faricimab vs. anti-VEGF treatments applied in such regimens.²

Methods

Related Macular Degeneration in Canada

A Markov cohort model based on the NICE guideline review³ was developed in Excel to estimate bilateral visual acuity changes linked to quality of life, injection frequency and associated costs from a Canadian payer as well as a societal perspective.

Cost-Effectiveness of Faricimab in Patients with Neovascular Age-

- Transition probabilities and injection frequency were informed by the TENAYA and LUCERNE trials for faricimab and a network-meta analysis for comparators⁴. Deterministic and probabilistic sensitivity analyses were performed for costs and key model parameters.
- Time horizon was 25 years to reflect a life time horizon. Utility for visual acuity states was modelled using Czoski-Murrey et al. (2009)⁵ including administration and related disutilites.
- Drug prices were based on publicly available list prices in Canada.

Results

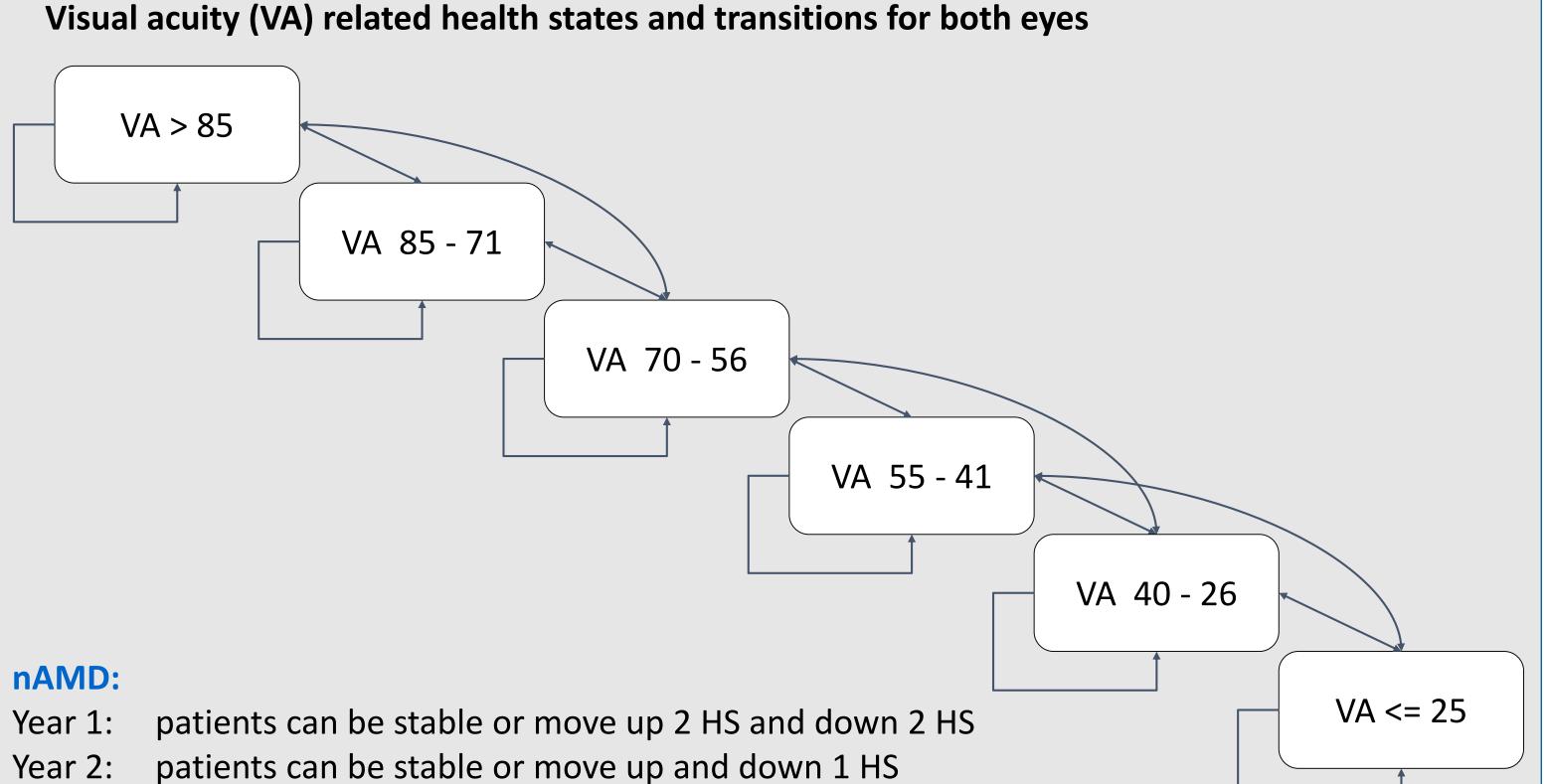
- In the deterministic base case, Faricimab reduced the number of injections by 37%, 21%, 28% and 46% vs. ranibizumab, aflibercept, brolucizumab and bevacizumab respectively using T&E.
- From a payer perspective, faricimab generated lower costs vs. ranibizumab, aflibercept and brolucizumab of CAD 76,496, 29,117 and 38,235 as well as higher costs of CAD 11,987 vs. bevacizumab. From a societal perspective, faricimab was cost saving vs. all anti-VEGF treatments including bevacizumab.
- Faricimab was associated with a mean QALY gain of 0.03, 0.05, 0.06 and 0.05 accordingly, driven by vision gains and disutilities.
- The ICUR vs. bevacizumab was 226,373 while the ICER per injection avoided was 222. Sensitivity analyses were consistent with the base case.

Conclusions

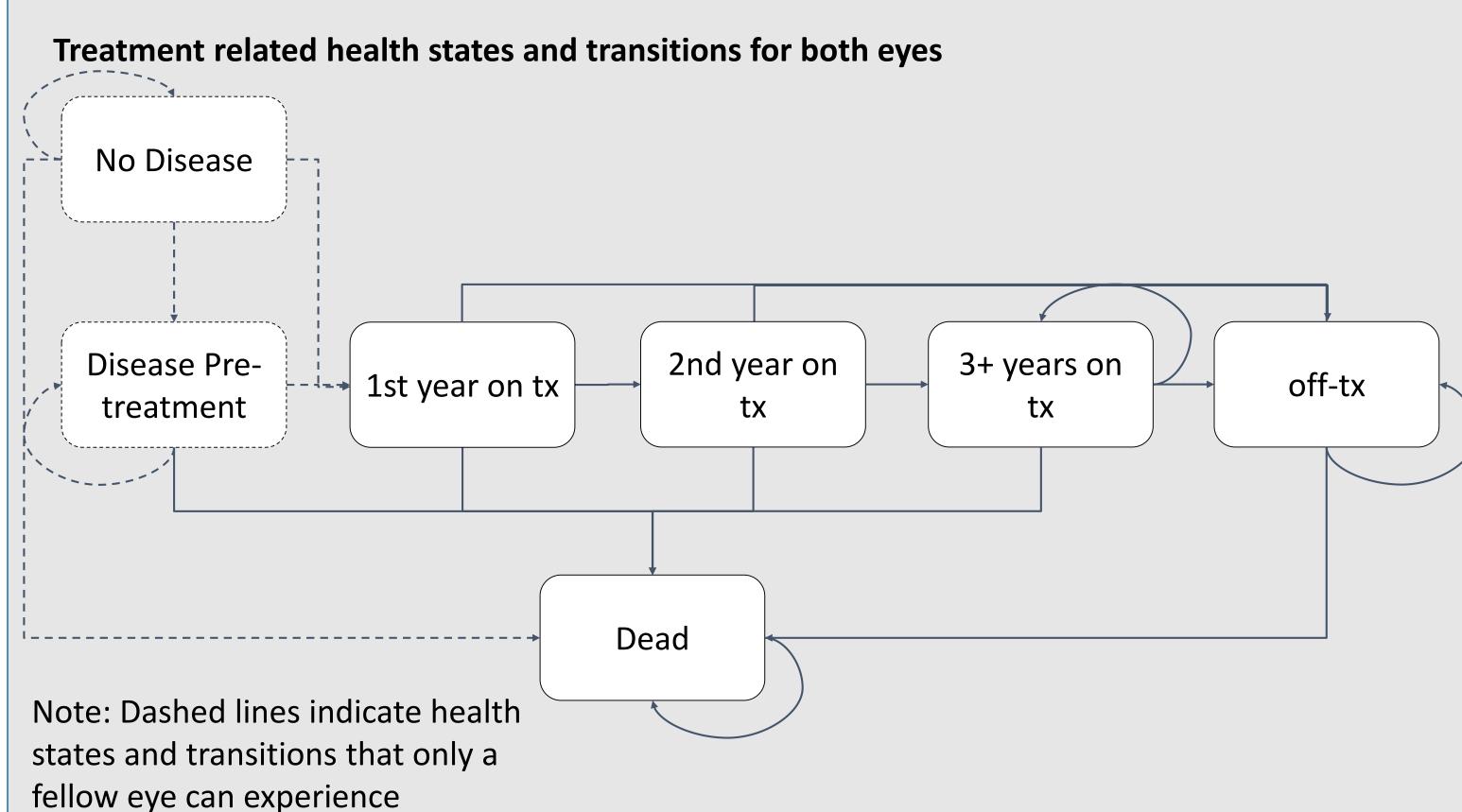
- The results indicate that faricimab dominates ranibizumab, aflibercept and brolucizumab administered in T&E regimes that are typically used in clinical practice.
- From a societal perspective, faricimab is cost saving and thus dominates bevacizumab by cutting injection visits and related costs in half. Savings were mostly driven by a reduced caregiver burden.
- Faricimab offers an innovative option reducing the treatment burden for patients and caregivers, leading to more efficient use of healthcare resources and long term cost savings.
- The results also indicate that societal costs such as informal care represent a substantial economic burden and should be considered when evaluating novel therapeutic options in ophthalmology.

1. Model Structure (Vision Related Health States)

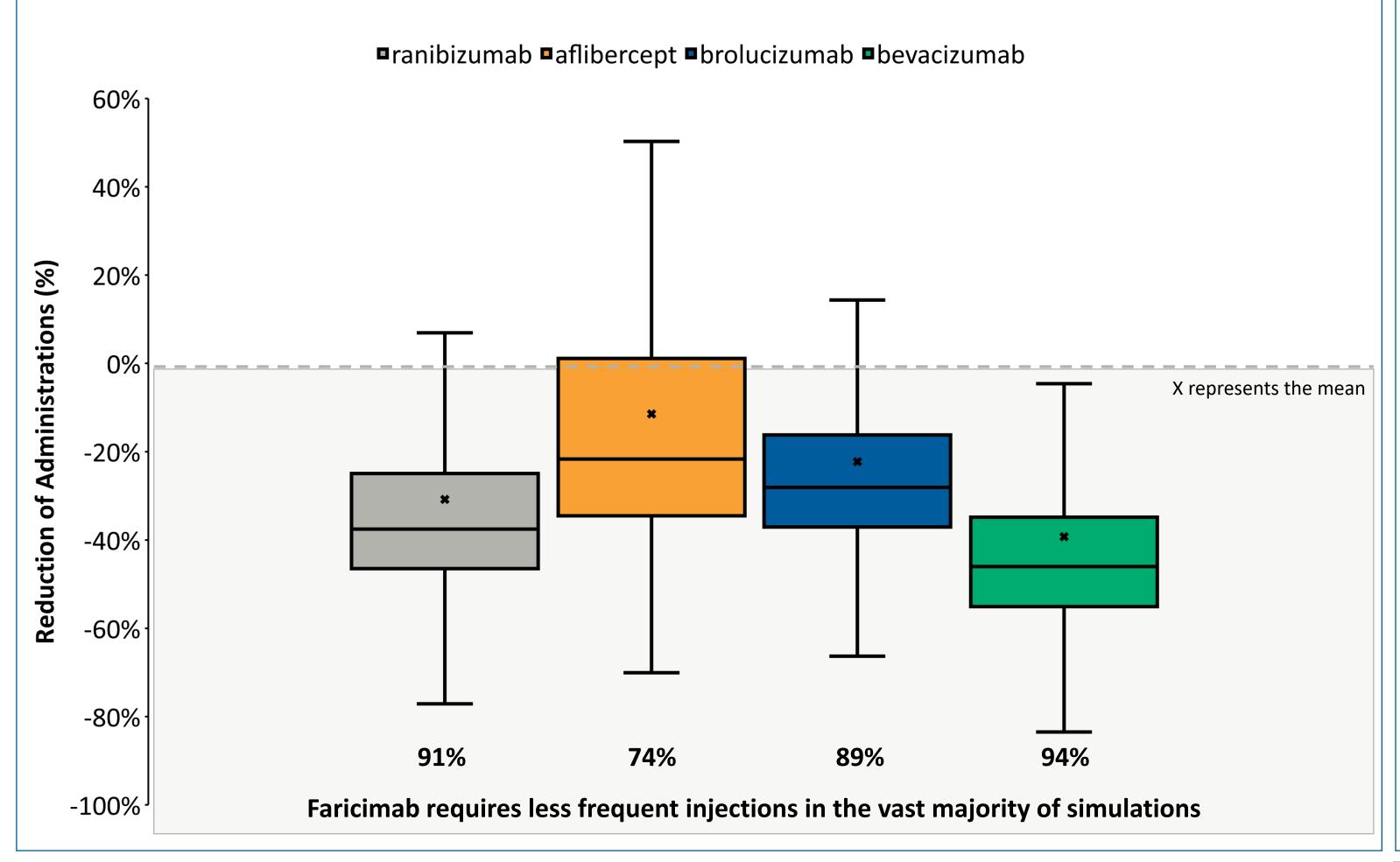
Year 3+: patients can be stable or move down up to 2 HS



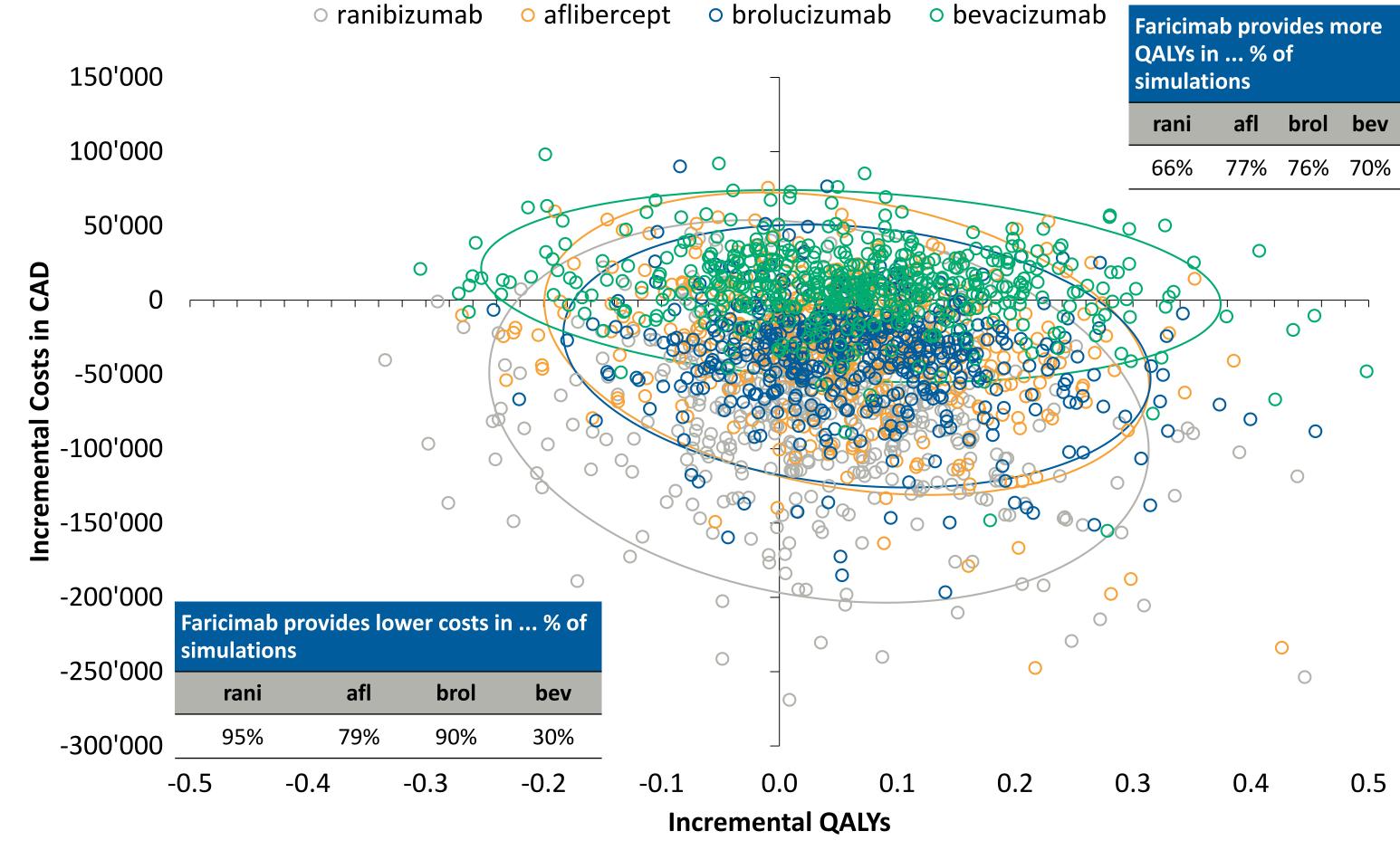
2. Model Structure (Treatment Related Health States)



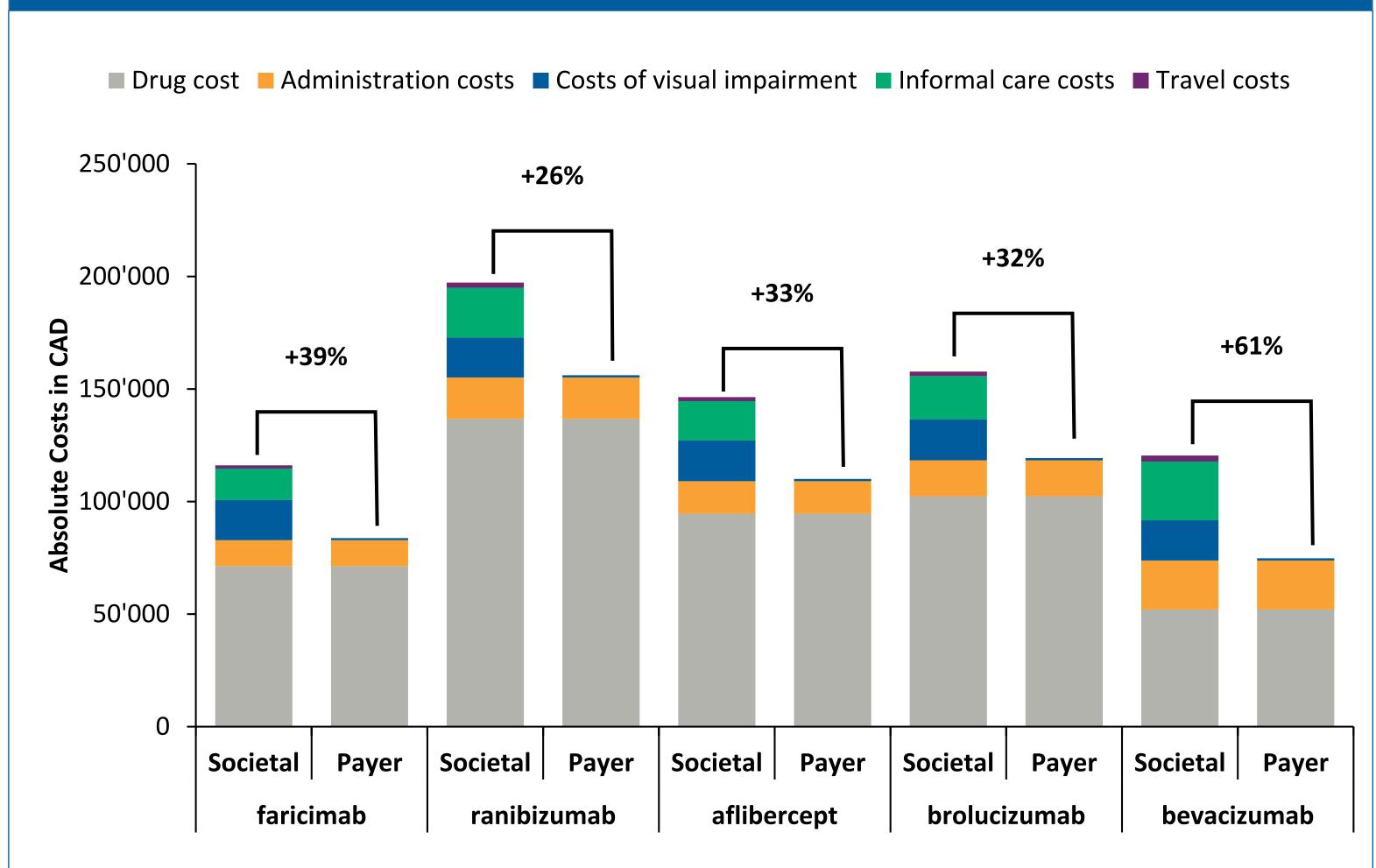
3. Mean Reduction of Administrations of Faricimab vs. Treat & Extend Regimens



4. Incremental Cost-Effectiveness Plane for Faricimab vs. Treat & Extend Regimes – **Payer Perspective**



5. Breakdown of Deterministic Total Costs – Societal vs. Payer Perspective



6. Deterministic Sensitivity Analysis – Societal Perspective

