

Introduction

- Faricimab is a bispecific antibody targeting ANG-2 and VEGF for the treatment of diabetic macular oedema (DMO).
- In the YOSEMITE and RHINE trials (2-year long Phase III trials),¹ patients treated with faricimab in a Treat & Extend (T&E) regime required less frequent treatments compared to aflibercept given every eight weeks (Q8W) and achieved similar vision gains.
- Clinical practice in DMO in Canada is typically characterized by pro re nata (PRN) regimens as well as T&E.²
- This research aims to assess the cost-effectiveness of faricimab vs. anti-VEGF treatments applied in such regimens.

Methods

- 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.
- Transition probabilities and injection frequency were informed by the YOSEMITE and RHINE 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-Murray et al. (2009)⁵ including administration and adverse event related disutilities.
- Drug prices were based on publicly available list prices in Canada.

Results

- The deterministic base case resulted in a mean QALY gain of 0.48, 0.21 and 0.53 vs. ranibizumab, aflibercept and bevacizumab respectively using PRN regimens and 0.55 vs. ranibizumab using a T&E regimen (data was only available for ranibizumab). QALY gains were mostly driven by improved quality of life due to better vision.
- From a payer perspective, faricimab generated lower costs vs. ranibizumab (PRN & T&E) and aflibercept of CAD 22,031, 20,600 and 8,480 as well as higher costs of CAD 31,019 vs. bevacizumab enabling patients to spend more than 1.5 additional years without visual impairment. Faricimab was also associated with a better durability profile than current treatment options.
- The ICER for the latter from a payer and societal perspective was 58,637 and 33,516. Sensitivity analyses were consistent with the base case.

Conclusions

- The results indicate that faricimab dominates ranibizumab and aflibercept administered in flexible regimens such as T&E and PRN that are normally used in clinical practice.
- Faricimab is associated with an ICER vs. bevacizumab that is within the typically accepted range especially from a societal perspective.
- Faricimab offers an innovative option enabling patients to retain their vision and independence for longer and contribute to society.
- The results also indicate, that societal costs such as informal care or costs of visual impairment represent a substantial economic burden and should be considered when evaluating novel therapeutic options.

