

Cost-Utility Analysis of Lorlatinib in First-Line Treatment of adult patients with ALK-Positive Advanced Non-Small Cell Lung Cancer (aNSCLC) in Colombia

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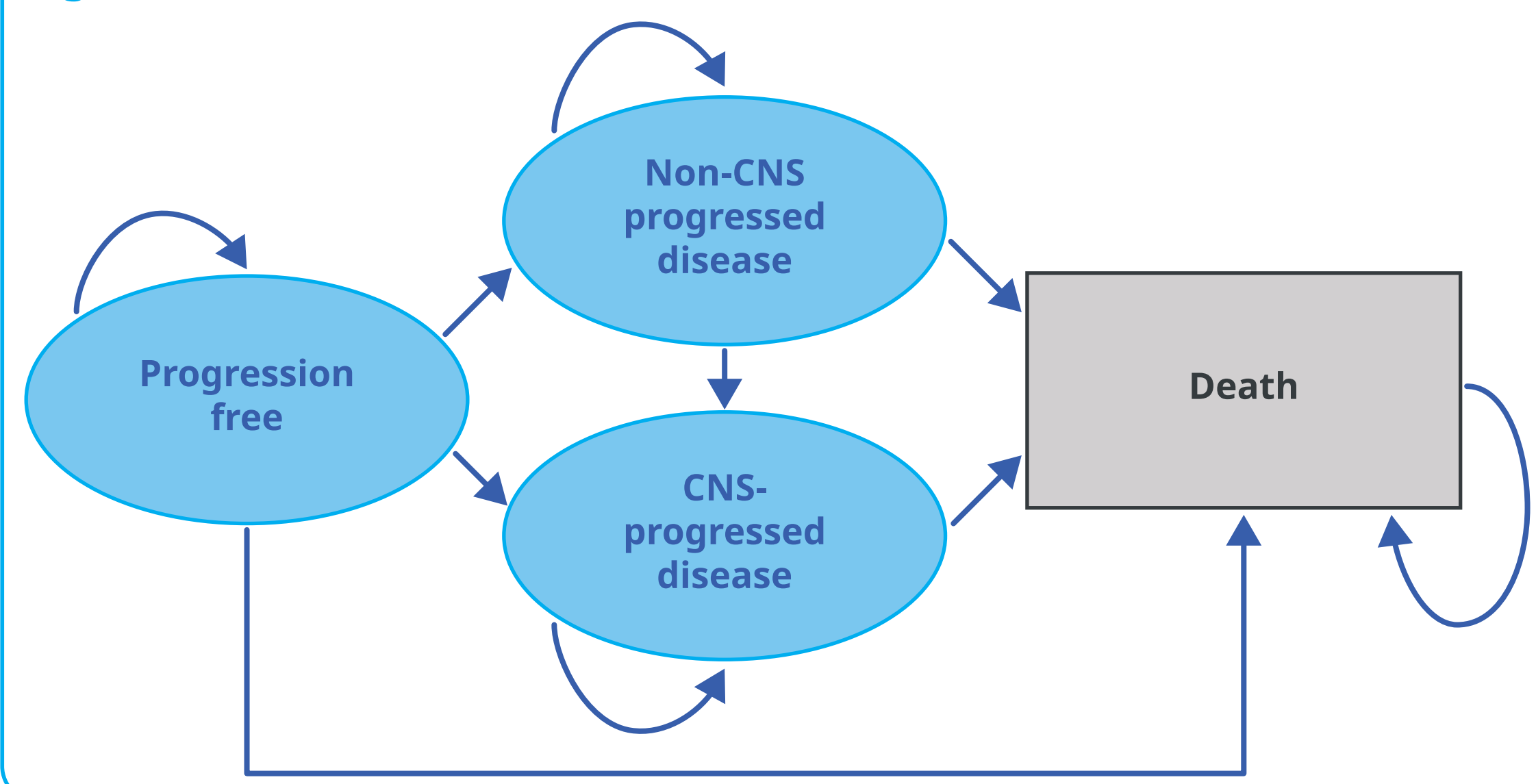
OBJECTIVE

- To evaluate the cost-utility of lorlatinib compared to the available ALK TKIs (alectinib, brigatinib, crizotinib) as first-line treatment option for adult patients with ALK+ aNSCLC from the Colombian healthcare system.

METHODS

- A partitioned survival Markov model with four health states was developed. The defined states included: progression-free, non-central nervous system (CNS) progressed disease, CNS-progressed disease, and death (Figure 1).

Figure 1. Model structure



- The time horizon for the model was set at 10 years with a 30-day cycle, a 5% annual discount rate, and the Colombian healthcare system perspective was adopted.
- Clinical, utility, and cost parameters required for the model were sourced from key clinical trials, published literature, national databases, and clinical experts. Prices were expressed in 2024 COP, and an exchange rate of 4,413.8 COP = 1USD was used¹.
- Due to the absence of head-to-head comparisons, an indirect comparison through a network meta-analysis was conducted to generate corresponding survival curves for lorlatinib against comparators, following Cope et al.²; Achana et al.³; and Guyot et al.⁴ methods. Crizotinib was the common comparator⁵⁻⁸.
- The cost-effectiveness threshold was set at three times the per-capita-gross-domestic-product (~20,500 USD), in line with national recommendations⁹.
- Robustness checks were performed utilizing one-way sensitivity analysis (OWSA) and probabilistic sensitivity analysis (PSA).

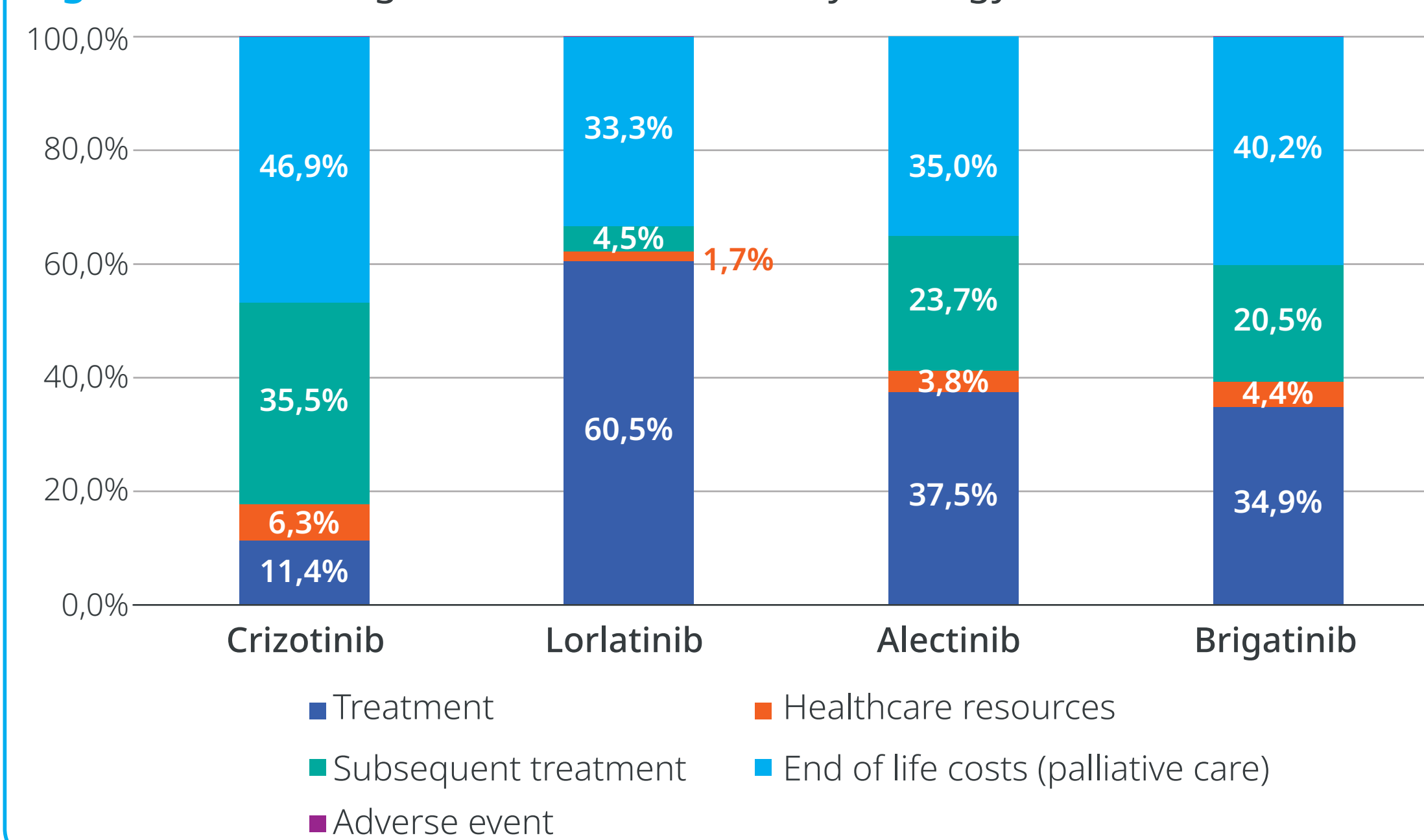
RESULTS

- Lorlatinib led to the highest number of quality adjusted life years with 4.87, followed by alectinib, brigatinib and crizotinib, with 4.18, 3.72 and 2.76, respectively.

RESULTS (cont)

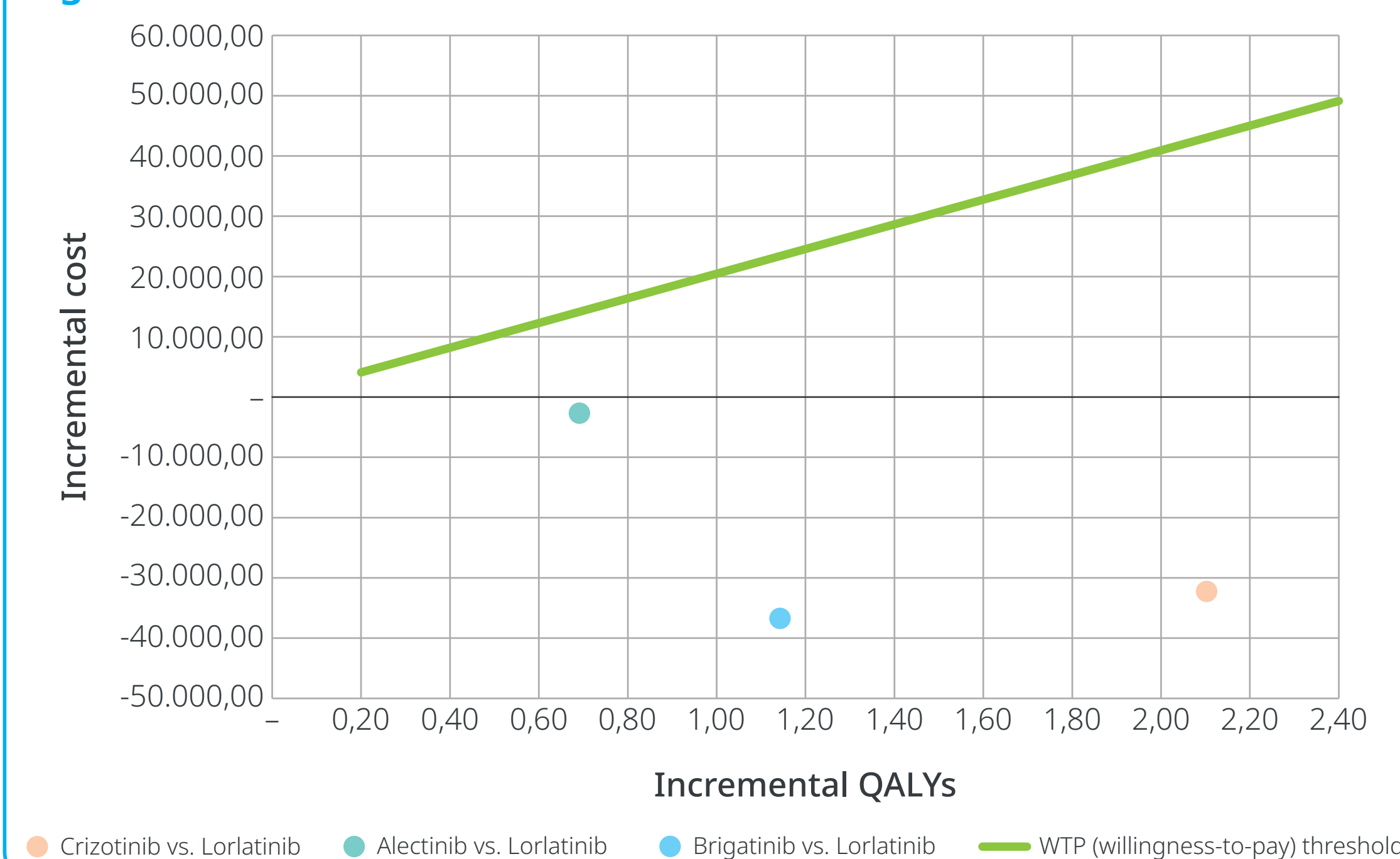
- In addition, lorlatinib occasioned the lowest cost (\$497,653), followed by alectinib (\$500,356), crizotinib (\$529,949) and brigatinib (\$534,388).
- Patients on the lorlatinib pathway presented the lowest end of life, subsequent treatment and healthcare resources (e.g. whole brain radiotherapy, follow-up visits, etc.) costs (Figure 2).
- No more than 0.1% of the costs were for managing adverse events, in the lorlatinib pathway they did not exceed 60 USD (Figure 2).

Figure 2. Percentage breakdown of costs by strategy



- Lorlatinib is a dominant strategy compared to the options available in the country. Lorlatinib savings per QALY compared to alectinib are about 2,703 USD, compared to crizotinib are 32,295 USD and compared to brigatinib are over 36 thousand USD (Figure 3).

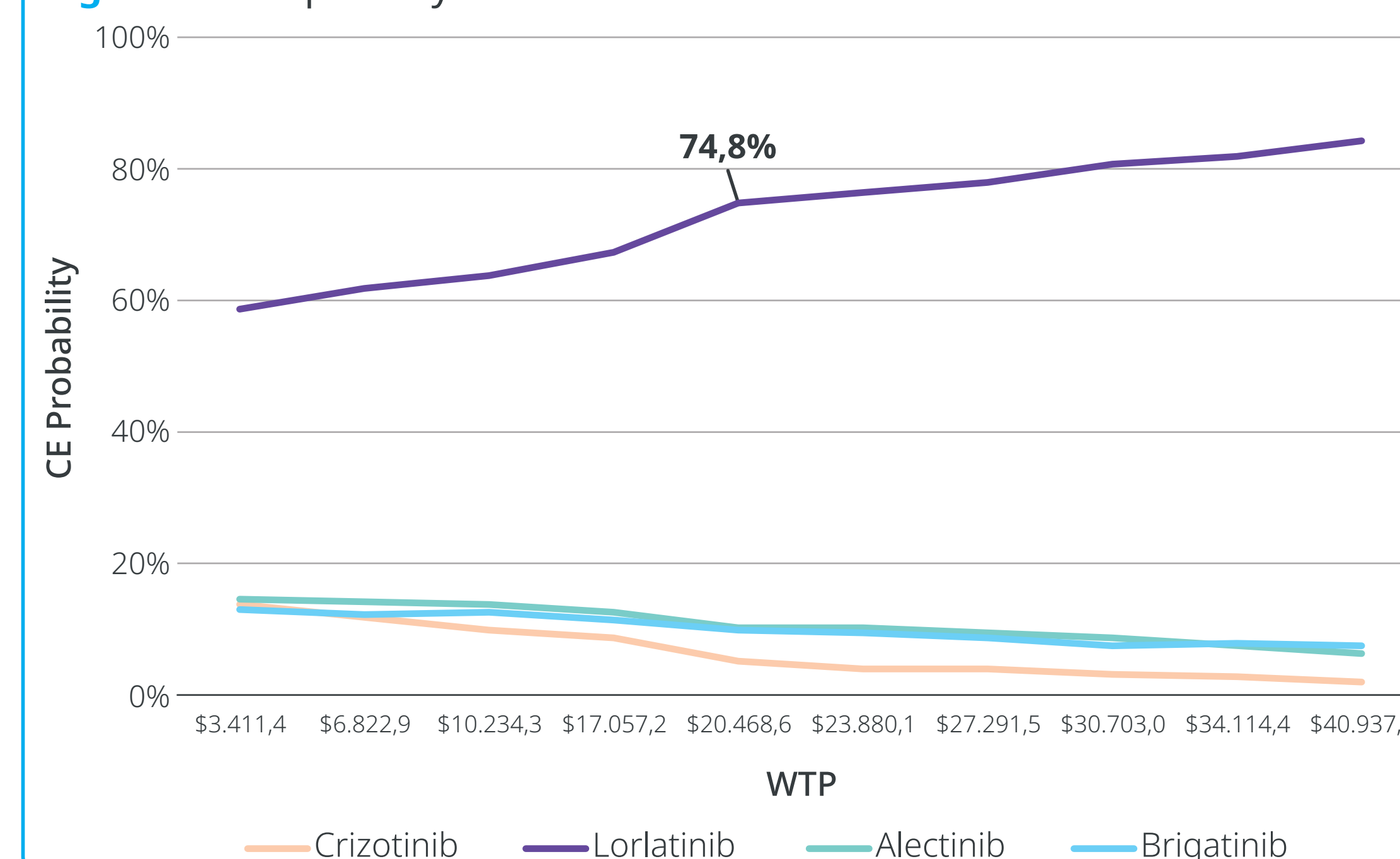
Figure 3. Cost-Effectiveness Plane



RESULTS (cont)

- The OWSA indicated that lorlatinib's price and comparators subsequent treatment costs were key parameters and the PSA showed that lorlatinib was the alternative with the highest probability of being cost-effective, with a probability of approximately 75% (Figure 4).

Figure 4. Acceptability curve



CONCLUSION

- In the Colombian health care system setting, lorlatinib is a cost-effective alternative against alectinib, brigatinib and crizotinib for the first-line treatment of adult patients ALK-positive aNSCLC.

REFERENCE

- BanRep. Tasa Representativa del Mercado (TRM - Peso por dólar) | Banco de la República n.d. <https://www.banrep.gov.co/es/estadisticas/trm>.
- Cope S, Chan K, Jansen JP. Multivariate network meta-analysis of survival function parameters. Res Synth Methods 2020;11:443-56. <https://doi.org/10.1002/jrsm.1405>.
- Achana FA, Cooper NJ, Bujkiewicz S, Hubbard SJ, Kendrick D, Jones DR, et al. Network meta-analysis of multiple outcome measures accounting for borrowing of information across outcomes. BMC Med Res Methodol 2014;14:92. <https://doi.org/10.1186/1471-2288-14-92>.
- Guyot P, Ades A, Ouwens MJ, Welton NJ. Enhanced secondary analysis of survival data: reconstructing the data from published Kaplan-Meier survival curves. BMC Med Res Methodol 2012;12:9. <https://doi.org/10.1186/1471-2288-12-9>.
- Camidge DR, Kim HR, Ahn MJ, Yang JCH, Han J-Y, Hochmair MJ, et al. Brigatinib Versus Crizotinib in ALK Inhibitor-Naïve Advanced ALK-Positive NSCLC: Final Results of Phase 3 ALTA-1L Trial. J Thorac Oncol 2021;16:2091-108. <https://doi.org/10.1016/j.jtho.2021.07.035>.
- Peters S, Camidge DR, Shaw AT, Gadgeel S, Ahn JS, Kim D-W, et al. Alectinib versus Crizotinib in Untreated ALK-Positive Non-Small-Cell Lung Cancer. N Engl J Med 2017;377:829-38. <https://doi.org/10.1056/NEJMoa1704795>.
- Mok T, Camidge DR, Gadgeel SM, Rosell R, Dziadziuszko R, Kim D-W, et al. Updated overall survival and final progression-free survival data for patients with treatment-naïve advanced ALK-positive non-small-cell lung cancer in the ALEX study. Ann Oncol 2020;31:1056-64. <https://doi.org/10.1016/j.annonc.2020.04.478>.
- Solomon BJ, Liu G, Felip E, Mok TSK, Soo RA, Mazieres J, et al. Lorlatinib Versus Crizotinib in Patients With Advanced ALK-Positive Non-Small Cell Lung Cancer: 5-Year Outcomes From the Phase III CROWN Study. J Clin Oncol 2024;42:3400-9. <https://doi.org/10.1200/JCO.24.00581>.
- IETS. Manual para la elaboración de evaluaciones económicas en salud 2014.