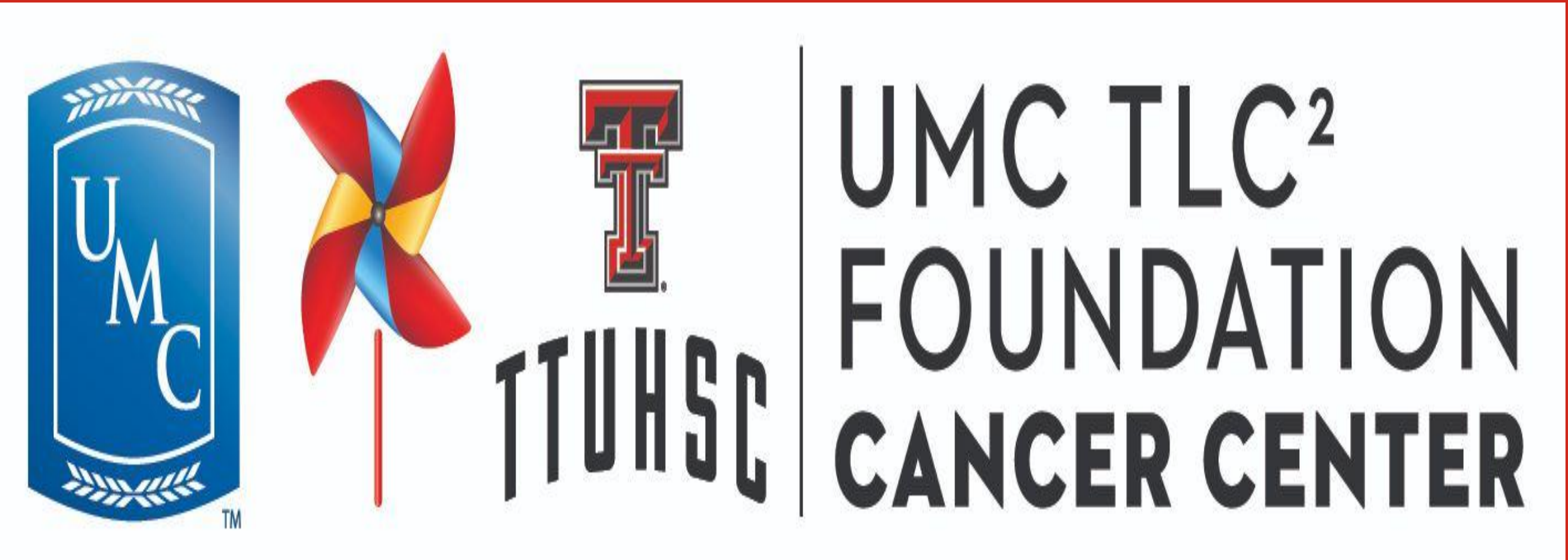


Cost-effectiveness Analysis of Teclistamab and Daratumumab Versus Ciltacabtagene Autoleucel in Relapsed Multiple Myeloma



Sohaib Irfan, MBBS¹; M. Abdullah Jamil, MBBS²; Muhammad Bilal Abid, MD, MS³
1. New York Medical College / Landmark Medical Center, Woonsocket, Rhode Island, USA
2. The Aga Khan University Medical College, Karachi, Pakistan
3. Stem Cell Transplantation and Cellular Immunotherapy Program, Department of Hematology/Oncology, TLC2 Foundation Cancer Center, Texas Tech University Health Sciences Center School of Medicine, Lubbock, Texas, USA

INTRODUCTION

- The phase III MajesTEC-3 trial showed superior progression-free survival (PFS) with teclistamab plus daratumumab (tec-dara) versus standard of care in relapsed/refractory multiple myeloma (RRMM): HR 0.17; 95% CI, 0.12–0.23; P<0.001.
- Ciltacabtagene autoleucel (cilta-cel), a BCMA-directed CAR T-cell therapy, is also approved in this clinical setting.
- The two strategies carry divergent economic profiles: a one-time CAR T-cell infusion versus indefinite bispecific antibody therapy.

AIM

- To evaluate the cost-effectiveness of tec-dara versus cilta-cel in RRMM after 1–3 prior lines of therapy, from a United States payer perspective.

METHODS

- Partitioned survival model: tec-dara (MajesTEC-3) vs cilta-cel (CARTITUDE-4) in RRMM, 1–3 prior lines.
- Three health states (PFS, progressed disease, death); 10-year horizon, monthly cycles.
- Long-term PFS extrapolated using exponential parametric fits to clinical data.
- Teclistamab costs capped at 48 months in the base case, reflecting plausible payer-imposed restrictions.
- Costs in 2025 US dollars, discounted 3% annually; utilities 0.82 (PFS) and 0.64 (progressed disease).
- Deterministic one-way and probabilistic sensitivity analyses; willingness-to-pay (WTP) threshold \$150,000/QALY.

CONCLUSIONS

- Despite superior modeled PFS, tec-dara is not cost-effective compared with cilta-cel at current pricing in RRMM.
- Response-adapted, fixed-duration treatment strategies merit further evaluation to reconcile clinical benefit with cumulative economic burden.
- As the RRMM landscape evolves, accessibility of bispecific antibodies must be balanced against the cost-efficiency of one-time cellular therapies.

RESULTS

Key findings

- Tec-dara delivered 2.094 additional QALYs at an incremental cost of \$565,686.
- The resulting ICER of \$270,103/QALY exceeds the \$150,000/QALY willingness-to-pay threshold, so tec-dara was not cost-effective relative to cilta-cel.
- The teclistamab treatment-duration cap was the single most influential driver of cost.
- One-way sensitivity analysis produced ICERs from \$44,016 to \$367,572/QALY; restricting therapy duration could render tec-dara cost-effective at the standard threshold.
- Probabilistic analysis confirmed cilta-cel as the preferred strategy in the majority of simulations, with tec-dara becoming cost-effective only at WTP thresholds substantially exceeding commonly accepted benchmarks.

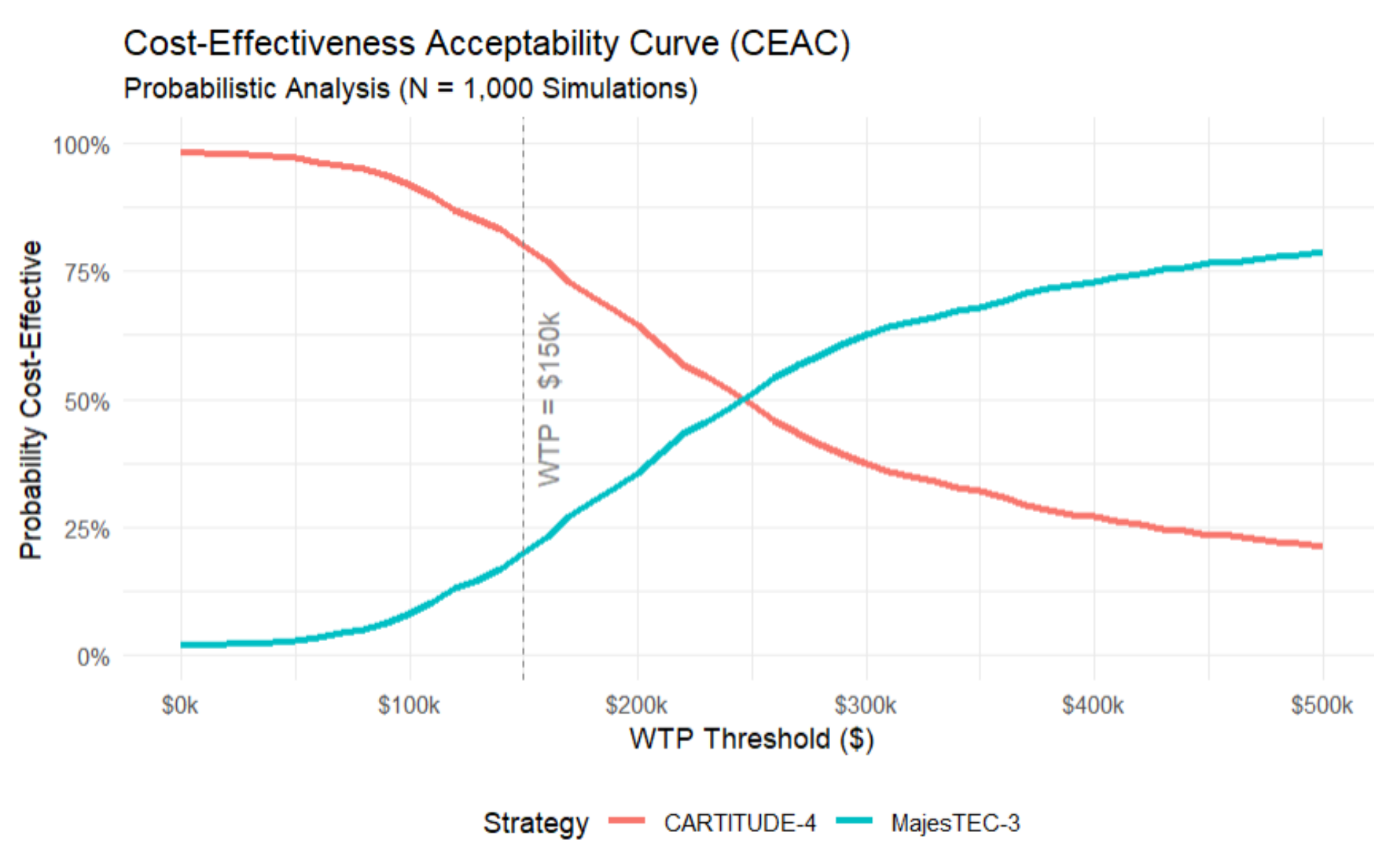


Figure 1. Cost-Effectiveness Acceptability Curve (CEAC)

Outcome	Tec-Dara	Cilta-cel	Incremental
Total cost (2025 USD)	\$1,122,707	\$557,021	+\$565,686
Total QALY	5.219	3.125	+2.094
ICER (\$/QALY)	—	Reference	\$270,103
Cost-effective at WTP \$150,000/QALY	No	Reference	—

Table 1. Base-case cost-effectiveness results, US payer perspective, 10-year horizon.

ICER: \$270,103 per QALY gained

Above the \$150,000/QALY threshold — tec-dara is not cost-effective. One-way sensitivity range: \$44,016–\$367,572/QALY.

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CONTACT INFORMATION

Muhammad Bilal Abid
Stem Cell Transplantation and Cellular Immunotherapy Program, Department of Hematology/Oncology, TLC2 Foundation Cancer Center, Texas Tech University Health Sciences Center School of Medicine, Lubbock
Email: Bilal.Abid@ttuhsc.edu