

Cost-Effectiveness of Tivozanib in Advanced Renal Cell Carcinoma in the US

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Background

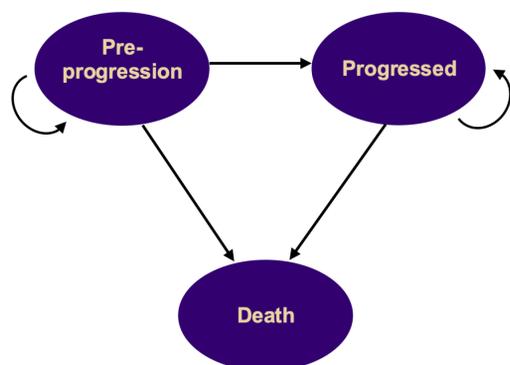
- Advanced renal cell carcinoma (aRCC) is the progressed form of the most common type of urogenital cancer¹
- In the United States (US), there are ~63,000 new cases and ~14,000 deaths due to RCC annually¹
- Annual economic burden of aRCC in the US ranges from \$0.60 to \$5.19 billion annually
- Despite the increasing treatment options for aRCC in recent years, many new drugs' cost-effectiveness and budgetary impact remain unexplored³
- In TIVO-3 trial, tivozanib outperformed sorafenib in terms of progression-free survival, but its higher cost leaves its cost-effectiveness uncertain⁴

Objective

- To evaluate the cost-effectiveness of tivozanib and sorafenib in patients diagnosed with relapsed or refractory aRCC from a US commercial payer perspective using a three-state partitioned survival model

Methods

Figure 1. Three-state Partitioned Survival Model



Methods

Table 1. Model Details

Model Structure	
Cycle Length	1 month
Time Horizon	Life-time
Perspective	US commercial payer
Correction	Half cycle
Input Parameters	
Health State Occupancy	Survival data was derived from the Kaplan Meier curves from TIVO-3
Utilities* (Monthly)	Pre-progression: 0.075 Progressed: 0.068
Disutilities	None. No drug-related adverse event ≥ grade 3 exceeded the 5% incidence threshold
Costs** (Monthly)	Tivozanib: \$36,800 Sorafenib: \$25,300
Discount (Monthly)	Cost: 0.25% Outcomes: 0.25%

*: adjusted to 2024 values, +: applied 20% variation

Population

- Patients with a diagnosis of relapsed or refractory aRCC and failed 2 or more systemic regimens other than sorafenib or tivozanib
- Mean age: 63 years old

Assumptions

- Patients remained on their drug regimen for their remaining life span
- There were no discontinuations or therapy switches

Results

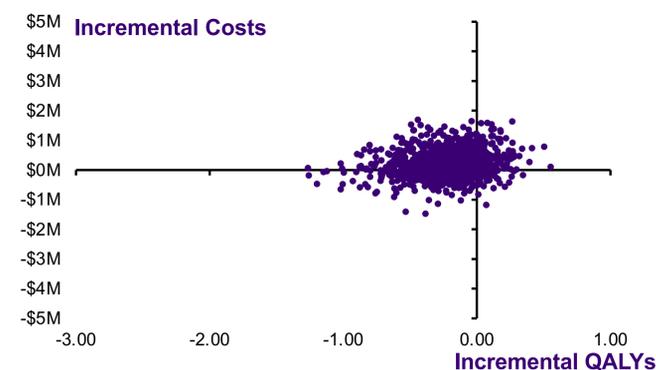
Table 2. Base Case Deterministic Results

	Tivozanib	Sorafenib	Incremental
Cost	\$483,325	\$375,781	\$107,544
Life-Years	25.80	29.24	-3.44
QALYs	1.82	2.05	-0.23
ICER			Dominated

Figure 2. One-Way Sensitivity Analysis



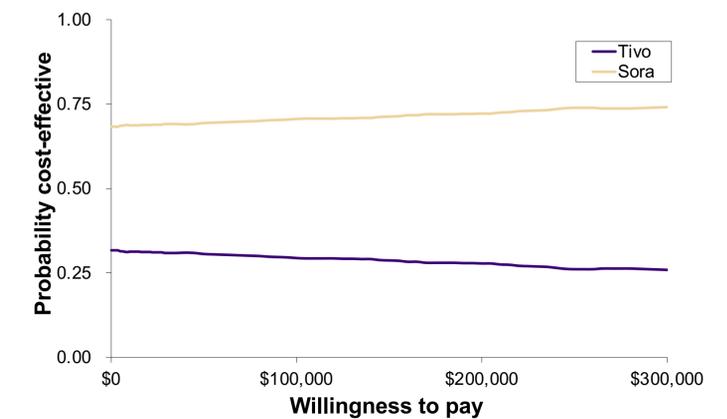
Figure 3. Probabilistic Sensitivity Analysis



- Tivozanib was less effective and more costly 55% of the time, while being less effective and costly 27% of the time compared to sorafenib

Results

Figure 4. Cost-effectiveness Acceptability Curve



- Sorafenib was more likely to be cost-effective than tivozanib by a large margin (70% vs. 30%) between the willingness-to-pay threshold of \$0 to \$300,000

Discussion

- At the willingness-to-pay threshold of \$150,000/QALY, tivozanib was dominated by sorafenib

Limitations

- Model assumed that patients would remain on the therapy throughout their remaining lifespan
- Model did not incorporate costly serious adverse events, such as pulmonary embolism and stroke
- The model did not factor in variations in adherence resulting from different administration frequencies between the two therapies

References

- Bahadoram S, Davoodi M, Hassanzadeh S, Bahadoram M, Barahman M, Mafakher L. Renal cell carcinoma: an overview of the epidemiology, diagnosis, and treatment. *G Ital Nefrol Organo Off Della Soc Ital Nefrol*. 2022;39(3):2022-vol3.
- Shih YC, Chien CR, Xu Y, Pan IW, Smith GL, Buchholz TA. Economic burden of renal cell carcinoma: Part I—an updated review. *Pharmacoeconomics*. 2011;29(4):315-329. doi:10.2165/11596100-000000000-00000
- Chien CR, Geysman DM, Kim B, Xu Y, Shih YCT. Economic Burden of Renal Cell Carcinoma—Part I: An Updated Review. *Pharmacoeconomics*. 2019;37(3):301-331. doi:10.1007/s40273-018-0746-y
- Rini BI, Pal SK, Escudier BJ, et al. Tivozanib versus sorafenib in patients with advanced renal cell carcinoma (TIVO-3): a phase 3, multicentre, randomised, controlled, open-label study. *Lancet Oncol*. 2020;21(1):95-104. doi:10.1016/S1470-2045(19)30735-1