# Cost-Effectiveness and Value of Information Analyses of Sotorasib vs. Docetaxel in Previously Treated Patients With Advanced Non-Small Cell Lung Cancer With KRAS G12C Mutation.

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#### **INTRODUCTION**

 Sotorasib is a first-in-class oral KRAS G12C inhibitor that showed progression-free survival benefit and an improved safety profile over docetaxel in the CodeBreak 200 trial.<sup>1</sup>

#### **OBJECTIVE**

 To evaluate the cost-effectiveness of sotorasib vs. docetaxel and to estimate the cost of uncertainty and the potential value of collecting additional information using value of information analysis.

### **METHODS**

- A 3-state partitioned survival model (progression-free, progressed, death) over a 5-year time horizon and from a US payer perspective was developed using Tree Age Pro.<sup>2</sup>
- Progression-free and overall survival estimates were determined from the Kaplan-Meier curves of the CodeBreak 200 trial using the best-fitting parametric distribution in R.<sup>3</sup>
- Costs of drugs were sourced from Redbook, administration costs from Physician Fee Schedule, cost of adverse events management, utilities, and disutilities from published literature.<sup>4-7</sup>
- One-way and probabilistic sensitivity analyses (PSA) accounted for model uncertainties. Discounting- 3% per year.
- PSA results were used to calculate the net health benefits (NHBs) and net monetary benefits (NMBs) forgone and the population expected value of perfect information (EVPI).8

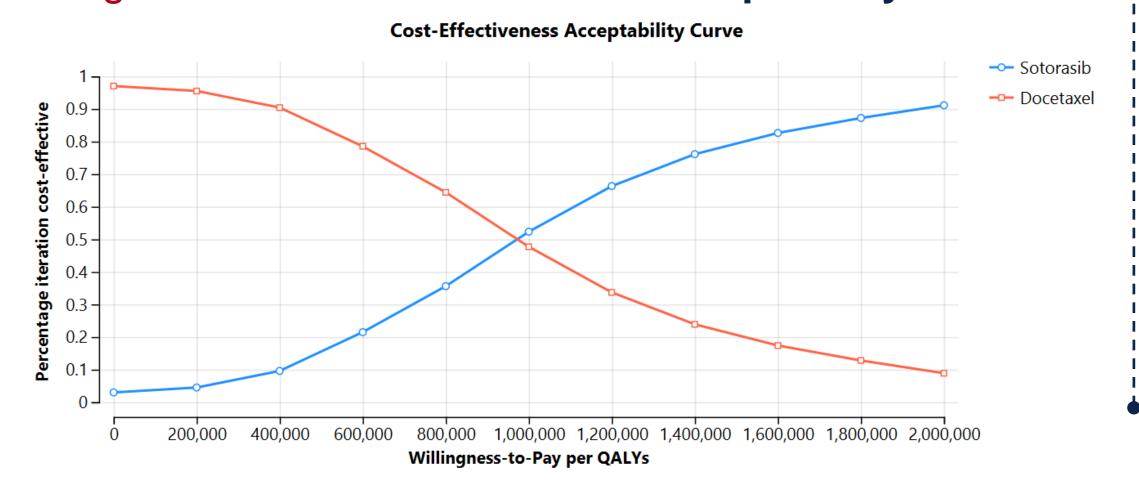
#### **RESULTS**

- Sotorasib yielded an increase of 0.35 QALY at an incremental cost of \$329,619 (Table 1).
- The probability of being cost-effective at a willingness to pay threshold of \$150,000 per QALY was 3% vs 97% for sotorasib and docetaxel respectively (Figure 1).

Table 1. Base-Case Analysis and Probabilistic Sensitivity Analysis For Sotorasib vs. Docetaxel

| Base-Case Analysis (Probabilistic Sensitivity Analysis) |                   |                   |  |  |  |  |
|---|-------------------|-------------------|--|--|--|--|
|   | Docetaxel         | Sotorasib         |  |  |  |  |
| Cost (\$)   | 204,532 (203,227) | 534,151 (535,969) |  |  |  |  |
| QALYg   | 0.70 (0.69)       | 1.05 (1.04)       |  |  |  |  |
| Incremental QALYg                                       | Ref               | 0.35 (0.35)       |  |  |  |  |
| Incremental Cost (\$)                                   | Ref               | 329,619 (332,742) |  |  |  |  |
| ICUR (\$ per QALYg)                                     | Ref               | 941,768 (950,691) |  |  |  |  |

Figure 1. Cost-Effectiveness Acceptability Curve



• The. average per-patient NHBs and NMBs forgone were 0.061QALY and \$5019 respectively. The population EVPI was estimated to be \$627.3 million (Table 2).

Table 2. Value of Information Analysis: Expected Value of Perfect Information (EVPI)

|       | Per person<br>NHB<br>(QALYs) | Per Person<br>NMB<br>(\$) | Population<br>NMB**<br>(\$) | Population<br>NMB (5-year<br>time horizon)<br>(\$) |
|-------|------------------------------|---------------------------|-----------------------------|--|
| EVPI* | 0.061                        | 5,019                     | 125,475,000                 | 627,375,000  |

\*EVPI based on a willingness to pay threshold of \$150,000 \*\*Calculated based on a yearly incidence of NSCLC with KRAS G12C mutation in the US (approximately 25,000 cases)<sup>9</sup>. NHB- Net health benefit NMB- Net monetary benefit.

## **CONCLUSION**

- Sotorasib may require a higher WTP threshold or a reduction in acquisition cost to be considered cost-effective.
- The estimated EVPI exceeds the cost of conducting another trial; future research to acquire additional evidence is considered worthwhile to inform clinical and policy decisions.

# REFERENCES

