IN-HOUSE VERSUS SEND-OUT NGS TESTING FOR METASTATIC NON-SMALL CELL LUNG CANCER: A BUDGET-IMPACT ANALYSIS

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Objectives

- Next generation sequencing (NGS) is used to identify genetic markers of disease, making it important for personalized cancer treatment.
- NGS testing can occur in external laboratories (send-out) or in the hospital (in-house).
- We analyzed the impact on hospital budgets of increasing inhouse NGS for metastatic non-small cell lung cancer (mNSCLC).

Methods

- A cohort-level, decision-tree model (Figure 1) feeding into a Markov model (Figure 2) was used to compare two hospital pathways:
 - Only send-out
 - Mixed in-house (75%) and send-out (25%)



Figure 1 Decision tree modeling movement of patients through standardized testing pathway until therapy decision. The arrows describe the movement of individuals between the different states.



Figure 2 Markov model for disease progression from initiation of therapy until death. The arrows describe the movement of individuals within and between the different states.

- A time horizon of five years was considered from the perspective of a US hospital.
- Costs are in 2021 USD, including costs of NGS capital acquisition associated with the hospital payer.
- The model inputs were derived from a retrospective analysis of real-world data of newly diagnosed stage IV mNSCLC cases and all others from peer-reviewed articles and expert opinions.¹

(Table 1)

Table 1 A selection of key model inputs

| Parameter | Value |
|--|---------------------------------|
| Cost of in-house NGS testing, per test | \$600* |
| Cost of send-out NGS testing, per test | \$300*‡ |
| Cost of single-gene testing per test | \$141 ² |
| Acquisition cost of in-house NGS | \$200,000** |
| Reimbursement for in-house NGS testing, per test | \$580 ³ |
| Revenue per hospital visit | \$124 ⁴ |
| Send-out turnaround time, days | 10.3-27.8 ^{5,1} |
| In-house turnaround time, days | 3 ^{5,6} |

*List price of laboratories, ‡ \$3,000 per send-out test with 10% of invoices funded by the hospital (\$300), **Expert opinion

Results

- For a hospital with 500 mNSCLC cases per year, the model estimated increases in overall testing costs as well as revenue with the use of the mixed approach.
- Compared to send-out, the mixed approach resulted in \$710,060 of increased testing costs and \$1,732,506 of increased revenue over the five-year time horizon.
- The return on investment was \$1,022,446 (95% credible Interval: \$787,903; 1,252,846) with a positive break-even point after 15 (95% credible Interval: 14; 17) months of investment.
- Different combinations of mNSCLC cases per year and proportions of in-house NGS implementation result in different break-even points. (Figure 3)
- More cases per year would require a lower rate of in-house NGS adoption to reach the break-even point and vice versa (Figure 3)
- The one-way sensitivity analysis showed that the proportion of send-out NGS testing and the proportion of patients on targeted therapy had the greatest impact on the return on investment.

| A | | | | | | | | | | | |
|------------|-----------------------------|------------|-----------------------------|------------|------------|------------|-------------|-------------|-------------|--|--|
| | Number of patients per year | | | | | | | | | | |
| | _ | 50 | 100 | 150 | 200 | 300 | 400 | 600 | 800 | | |
| <u>-</u> | 5% | -\$190,156 | -\$180,312 | -\$170,468 | -\$160,624 | -\$140,936 | -\$121,248 | -\$81,872 | -\$42,496 | | |
| | 10% | -\$182,127 | -\$164,255 | -\$146,382 | -\$128,510 | -\$92,764 | -\$57,019 | \$14,471 | \$85,962 | | |
| of GS | 15% | -\$174,099 | -\$148,198 | -\$122,296 | -\$96,395 | -\$44,593 | \$7,210 | \$110,815 | \$214,420 | | |
| ion S | 20% | -\$166,070 | -\$132,140 | -\$98,210 | -\$64,281 | \$3,579 | \$71,439 | \$207,158 | \$342,877 | | |
| ort us(| 40% | -\$133,956 | -\$67,911 | -\$1,867 | \$64,177 | \$196,266 | \$328,355 | \$592,532 | \$856,709 | | |
| Prop | 60% | -\$101,841 | -\$3,682 | \$94,476 | \$192,635 | \$388,953 | \$585,270 | \$977,905 | \$1,370,541 | | |
| | 80% | -\$69,727 | \$60,547 | \$190,820 | \$321,093 | \$581,640 | \$842,186 | \$1,363,279 | \$1,884,372 | | |
| | 100% | -\$37,612 | \$124,775 | \$287,163 | \$449,551 | \$774,326 | \$1,099,102 | \$1,748,653 | \$2,398,204 | | |
| В | | | | | | | | | | | |
| | | | Number of patients per year | | | | | | | | |
| | _ | 50 | 100 | 150 | 200 | 300 | 400 | 600 | 800 | | |
| | 5% | >60 | >60 | >60 | >60 | >60 | >60 | >60 | >60 | | |
| | 10% | >60 | >60 | >60 | >60 | >60 | >60 | >60 | 49 | | |
| of GS | 15% | >60 | >60 | >60 | >60 | >60 | >60 | 45 | 36 | | |
| e Non | 20% | >60 | >60 | >60 | >60 | >60 | 51 | 37 | 29 | | |
| ort us(| 40% | >60 | >60 | >60 | 52 | 37 | 29 | 21 | 17 | | |
| oh | 60% | >60 | >60 | 47 | 37 | 27 | 21 | 16 | 12 | | |
| Pr | 80% | >60 | 52 | 38 | 30 | 21 | 17 | 12 | 10 | | |
| | 4000/ | | 10 | 0.4 | 05 | 10 | | 4.0 | | | |

Figure 3 Scenario analysis A Return on investment after five years. B Payback period (months), dark grey: scenario does not break even withn five year time horizon.

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CONCLUSION

- An in-house NGS system reduces the testing turnaround time, and increases the number of mNSCLC patients on targeted therapy.
- Hospitals with >100 mNSCLC cases per year are expected to find in-house NGS profitable within 5 years.
- The mixed approach reduced the mean test turnaround time by 9.86 (9.21; 10.49) days and led to a +3.38 (2.31; 4.05) %-points increase in patients on targeted therapies (Figure 4).



Figure 4 Effects of a mixed approach **A** Average turn-around time in days **B** Patients on targeted therapy as a percentage of all patients

References

- Robert Smith, MD, Mei Xue, Rhonda Williams, Natalie Dorrow. Retrospective Analysis Using Real-World Data (RWD) in Predominately Newly Diagnosed Stage 4 Non-small Cell Lung Carcinoma (NSCLC-4) to Determine the Effect of Genomic Profiling on Treatment Decisions; 2021; ST59
- Johnston KM, Sheffield BS, Yip S, Lakzadeh P, Qian C, Nam J. Costs of in-house genomic profiling and implications for economic evaluation: a case example of non-small cell lung cancer (NSCLC). J Med Econ. 2020;23(10):1123-1129.
- Sabatini LM, Mathews C, Ptak D, et al. Genomic Sequencing Procedure Microcosting Analysis and Health Economic Cost-Impact Analysis: A Report of the Association for Molecular Pathology. J Mol Diagn. 2016:18(3):319-328.
- Vanderpoel J, Stevens AL, Emond B, et al. Total cost of testing for genomic alterations associated with nextgeneration sequencing versus polymerase chain reaction testing strategies among patients with metastatic non-small cell lung cancer. J Med Econ. 2022;25(1):457-468.
- Sheffield BS, Beharry A, Diep J, et al. Point of Care Molecular Testing: Community-Based Rapid Next-Generation Sequencing to Support Cancer Care. Curr Oncol. 2022;29(3):1326-1334.
- Ilié M, Hofman V, Bontoux C, et al. Setting Up an Ultra-Fast Next-Generation Sequencing Approach as Reflex Testing at Diagnosis of Non-Squamous Non-Small Cell Lung Cancer; Experience of a Single Center (LPCE, Nice, France). Cancers (Basel). 2022;14(9).

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