

# Cost consequence model of the MammaPrint® (70 -gene signature) and 21 -gene signature in Patients with primary HR+ HER2 -, N1 early-stage breast cancer in Germany

<sup>1</sup>Kooperatives Brustzentrum Paderborn, Klinik für Gynäkologie und Geburtshilfe, Frauenklinik St. Louise, Paderborn, Germany <sup>2</sup>Agendia NV, Amsterdam, the Netherlands <sup>3</sup> Kooperatives Brustzentrum Paderborn, Klinik für Hämatologie und Onkologie, Brüderkrankenhaus St. Josef, MVZ im MediCo, Paderborn, Germany

### BACKGROUND

The MammaPrint 70-gene signature helps to identify breast cancer patients who are at high risk of distant metastasis and might benefit from chemotherapy, as opposed to those with low-risk tumors who can safely forego chemotherapy. <sup>1,2</sup> MammaPrint enables physicians to make informed decisions for chemotherapy and as a result will minimize unnecessary exposure to chemotherapy's toxic effects and decrease financial burden for the healthcare system by avoiding overtreatment with associated direct and indirect medical and non-medical costs. A reduction of chemotherapy-related direct and indirect costs can have a significant impact on reducing the overall costs of breast cancer treatment.

This analysis evaluates the economic impact of implementing MammaPrint, the 21-gene signature (Oncotype DX [ODx]), or no gene signature testing for patients with hormone receptor-positive (HR+) HER2negative early breast cancer with 1 to 3 positive lymph nodes (N1) in Germany.

### METHODS

We developed a cost-consequence model to evaluate the budgetary implications of using MammaPrint, ODx, or forgoing gene signature testing altogether. This model considers both perspectives: healthcare payers and society. The model focuses on patients with HR+/HER2-/N1 breast tumors. Input data were sourced from MINDACT, RxPONDER, literature, and German pricing repositories (EBM and DRG system). Key input parameters for the primary analysis are displayed in Table 1 and Table 2.

## METHODS Table 1: Input parameters for no test, MammaPrint and Oncotype DX

| Scenario                 |
|--------------------------|
| No Test                  |
|                          |
|                          |
| Oncotype DX <sup>5</sup> |
|                          |
|                          |
| MammaPrint <sup>2</sup>  |

Chemothera Supportive Supportive Control a Short term a Long term a Loss Sic Transpo

Costs of dista Costs of loca Percentage of DR

**Table 1:** The model considers genomic risk proportions and
 efficacy parameters based on MINDACT and RxPONDER. 
**Table 2:** The model considers chemotherapy related costs based
 on EBM prices and the DRG system. The healthcare payer perspective covers testing, direct/indirect costs of chemotherapy, and disease recurrence costs. The societal perspective adds transport costs and productivity losses.

Lux M.P.<sup>1</sup>, Sandor M. F.<sup>1</sup>, Hofmann V.<sup>1</sup>, Pronin D.<sup>2</sup>, Klinkhamer J.C.<sup>2</sup>, Müller-Huesmann H.<sup>3</sup>

| Cost [€]              | Patients in each risk category<br>[%] |    | Assignment to<br>Treatment | Recurrence<br>Probability |
|-----------------------|---------------------------------------|----|----------------------------|---------------------------|
| 0.00                  | Node positive                         | 37 | No Chemo                   | 0.0850 2,4                |
|                       | disease <sup>3</sup>                  | 63 | Chemo                      | 0.0572 <sup>2</sup>       |
|                       |                                       |    |                            |                           |
| 2,881.80 <sup>7</sup> | RS = 0 - 25                           | 83 | No Chemo                   | 0.0610 <sup>5</sup>       |
|                       | RS = 26-100                           | 17 | Chemo                      | 0.1690 <sup>6</sup>       |
|                       |                                       |    |                            |                           |
| 2,302.77 <sup>7</sup> | Low Risk                              | 73 | No Chemo                   | 0.0402 <sup>2</sup>       |
|                       | High Risk                             | 27 | Chemo                      | 0.1030 <sup>2</sup>       |

### Table 2: Chemotherapy and recurrence associated costs <sup>8,9</sup>

| 1 /                        |               |            |
|----------------------------|---------------|------------|
| osts                       | Type of Costs | Price [€]  |
| apy application            | Direct        | 5,292.22   |
| e medication               | Direct        | 4,932.20   |
| e medication               | Indirect      | 655.64     |
| nd planning                | Direct        | 988.83     |
| adverse events             | Direct        | 660.59     |
| adverse events             | Direct        | 515.84     |
| in GDP                     | Indirect      | 16,752.41  |
| k Pay                      | Indirect      | 2,729.83   |
| oort costs                 | Indirect      | 1,895.94   |
|                            |               |            |
| ant recurrence             |               | 152,925.70 |
| cal recurrence             |               | 32,859.00  |
| having experienced<br>a LR |               | 10%        |
|                            |               |            |

### RESULTS

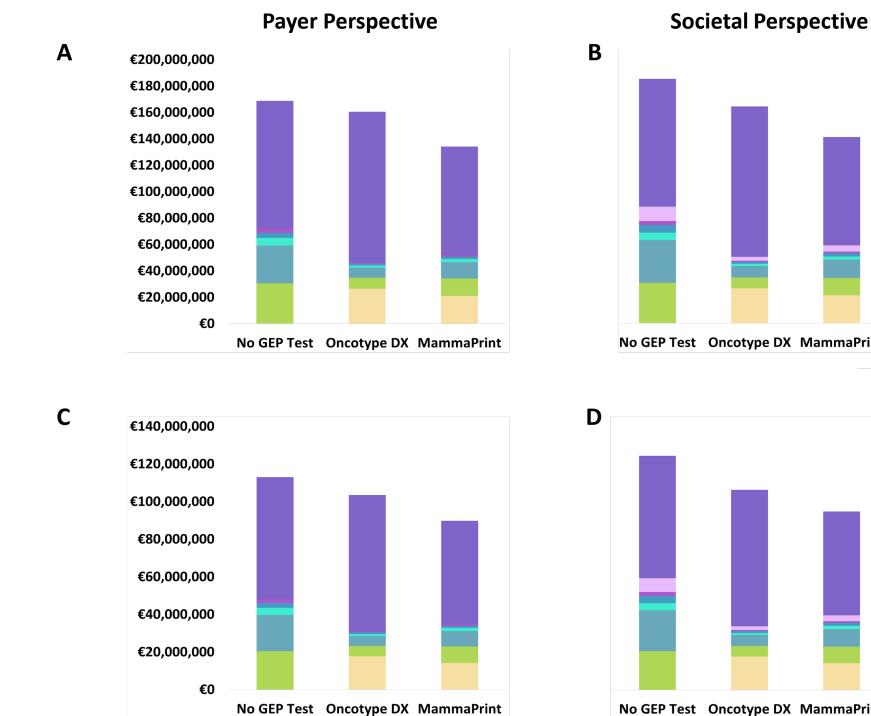
Figure 1A & 1B: The model demonstrated per patient savings from a healthcare payer perspective of €3,766 for MammaPrint and €900 for ODx compared to not using any gene signature for N1 breast cancer patients in Germany. Importantly, the savings are even more substantial from a societal perspective, reaching €11,815 for MammaPrint and €11,243 for ODx.

Figure 1C and 1D: When limiting the population to women >50 years due to available evidence, similar results were observed with per patient savings from a payer perspective of €3,766 and €1,549, and from a societal perspective of €9,826 and €9,349, for MammaPrint and ODx, respectively.

### CONCLUSIONS

- German healthcare budget.





• In HR+/HER2-/N1 breast cancer, the use of MammaPrint leads to a reduction in chemotherapy use and associated costs in Germany compared to the absence of gene expression profiling testing. Although MammaPrint designates a slightly smaller proportion of genomically Low Risk patients compared to ODx, MammaPrint Low Risk patients without chemotherapy demonstrate higher survival outcomes, resulting in MammaPrint being a cost-conscious option. MammaPrint contributes to a personalized treatment plan, achieving a net reduction in the use of chemotherapy, providing savings for the



MammaPrint versus no GEP test in HR+/HER2-/N1 breast cancer



perspective

MammaPrint versus no GEP test, largely driven by fewer chemotherapy related costs



| 0    |   |
|------|---|
| е    |   |
|      |   |
|      |   |
|      |   |
|      | Test costs  |
| rint | <ul> <li>Control</li> <li>Transport</li> </ul>                              |
|      | <ul> <li>Chemotherapy</li> <li>Short Term AE</li> <li>Recurrence</li> </ul> |
|      | <ul> <li>Supportive medication</li> <li>Long Term AE</li> </ul>             |
|      |   |
| rint |   |



**Total savings societal** perspective

MammaPrint versus no GEP test, largely driven by fewer productivity losses

ASCO 2024