EndoPredict® in early hormone receptor-positive, HER2-negative breast cancer

K. Almstedt1· S. Mendoza1· M. Otto2· M. J. Battista1· J. Steetskamp1· A. S. Heimes1· S. Krajnak1· A. Poplawski3· A. Gerhold-Ay3· A. Hasenburg1· C. Denkert4· M. Schmidt1

Received: 30 March 2020 / Accepted: 11 May 2020 / Published online: 20 May 2020
© The Author(s) 2020

Abstract
Purpose Evaluating consecutive early breast cancer patients, we analyzed both the impact of EndoPredict® on clinical decisions as well as clinico-pathological factors influencing the decision to perform this gene expression test.

Methods Hormone receptor (HR)-positive and human epidermal growth factor receptor 2 (HER2)-negative early breast cancer patients treated between 2011 and 2016 were included in this study to investigate the role of EndoPredict® (EPclin) in the treatment of early breast cancer. A main study aim was to analyze the changes in therapy recommendations with and without EPclin. In addition, the impact of clinico-pathological parameters for the decision to perform EPclin was examined by Pearson’s chi-squared test ($\chi^2$-test) and Fisher’s exact test as well as univariate and multivariate logistic regressions.

Results In a cohort of 869 consecutive early HR-positive, HER-negative breast cancer patients, EPclin was utilized in 156 (18.0%) patients. EPclin led to changes in therapy recommendations in 33.3% ($n=52$), with both therapy escalation in 19.2% ($n=30$) and de-escalation in 14.1% ($n=22$). The clinico-pathological factors influencing the use of EPclin were age ($P<0.001$, odds ratio [OR] 0.498), tumor size ($P=0.011$, OR 0.071), nodal status ($P=0.021$, OR 1.674), histological grade ($P=0.043$, OR 0.432), and Ki-67 ($P<0.001$, OR 3.599).

Conclusions EPclin led to a change in therapy recommendations in one third of the patients. Clinico-pathological parameters such as younger age, smaller tumor size, positive nodal status, intermediate histological grade and intermediate Ki-67 had a significant influence on the use of EndoPredict®.

Keywords Early breast cancer · EndoPredict® · Endocrine therapy · Treatment decision · Prognosis · Gene expression

Introduction

The extent of the benefit from adjuvant chemotherapy depends on the patient’s individual risk profile [1]. According to the current recommendations, traditional clinico-pathological parameters such as tumor size, nodal status, histological grading, hormone receptor (HR) status and human epidermal growth factor receptor 2 (HER2) status should be used for risk stratification in women with early stage invasive breast cancer [2, 3]. In clinical practice, breast cancer is subdivided into different subtypes (i.e., luminal A-like, luminal B-like, HER2-like, triple-negative) [3]. The development and use of genomic tests using gene expression profiling has the aim to improve prediction of clinical outcome. Especially for HR-positive, HER2-negative, node-negative tumors, the additional use of multigene assays should help to reliably assess the risk profile in order to avoid both under- and overtreatment.
For clinical use, it is crucial that gene expression analyses are assessed according to clear evidence-based criteria [4]. Recommended commercially available biomarker assays by the St. Gallen International Consensus Conference and the American Society of Clinical Oncology (ASCO) are Oncotype DX®, EndoPredict®, MammaPrint®, PAM50/Prosigna® and Breast Cancer Index [2, 3, 5, 6]. All of these gene expression assays are endorsed for guiding the decision on adjuvant chemotherapy in hormone receptor-positive, node-negative tumors if all other criteria do not allow a therapeutic decision. MammaPrint®, EndoPredict® and Prosigna® might also be used in 1–3 involved axillary lymph nodes. Retrospective evidence exists for the use of OncotypeDX® in node positive disease [7, 8], although the prospective phase 3 RxSPONDER trial (SWOG S1007, https://clinicaltrials.gov/ct2/show/NCT01272037) is still ongoing. For the 21-gene recurrence score (TAILORx trial) [9–11] and the 70-gene signature (MINDACT trial) [12] prospective data are available. There is also a smaller prospective study on EndoPredict® in 200 patients, which addresses the clinical and psychological impact of this multigene assay [13]. In addition, there is ample prospective-retrospective evidence for the approved multigene assays.

The RNA-based multigene score EndoPredict® (EP, Myriad Genetics®, Salt Lake City, USA) is based on the gene expression analysis of 12 genes, eight cancer-related, three reference genes and one control gene for DNA contamination. EndoPredict® integrates genomic and clinical information by including clinico-pathological parameters like tumor size and number of involved lymph nodes (EPclin). Integrating these two clinical parameters to the molecular test (EPclin) increased the prognostic value especially for late-distant recurrence for the multigene assay EndoPredict® [14]. The test result distinguishes EPclin low-risk and EPclin high-risk patients [15–17]. The clinical benefit of EndoPredict® to identify a subgroup with an excellent prognosis with endocrine therapy only is well-established [15, 18–20].

With an increasing use of multigene tests, it is of utmost importance to know whether their use changes treatment decisions. This retrospective study in consecutive patients compared pre- and post-test therapy decisions and changes in treatment recommendations. Furthermore, we evaluated clinico-pathological factors influencing the decision for the additional use of EPclin. These real-world data should be of special interest for indication and evaluation of EndoPredict®.

Methods

Patient cohorts and data collection

The certified breast cancer center of the University Medical Center Mainz has been prospectively documenting clinico-pathological characteristics as well as therapies of all treated breast cancer patients. This database was searched for patients with hormone receptor-positive, HER2-negative early breast cancer, with a maximum of three positive axillary lymph nodes diagnosed between November 2011 and September 2016. Patients with neoadjuvant chemotherapy, distant metastasis or breast cancer recurrence were not eligible for this study (Fig. 1).

Information on patient characteristics (e.g., age at primary breast surgery, menopausal status), histopathological data (tumor size, axillary lymph-node status, histological tumor type, histological grading, estrogen receptor (ER)-, progesterone receptor (PR)-, and HER2-status, proliferation index assessed by Ki-67), therapeutic recommendations as well as the decision to use EndoPredict®, were collected from the patients’ medical records and the documented tumor board decisions. The results of the EndoPredict® test (EP- and EPclin test) were also obtained from this source.

Gene expression analysis (EndoPredict®)

EndoPredict® was performed either in the Institute of Pathology of the Charité, University Medical Center Berlin, Campus Mitte, Charitéplatz 1, 10117 Berlin (n = 8) or in the Molecular Pathology Trier, Max-Planck-Str. 17, 54296 Trier (n = 148) using RNA from formalin-fixed and paraffin-embedded (FFPE) tissue from the surgical specimen. EP was combined with tumor size and nodal status into a comprehensive risk score, EPclin, as previously described [15]. The EPclin-score ranged from 0 to 7. EPclin-score ≥ 3.3 differentiated between high-risk and low-risk patients.

To assess a possible change in the treatment decision, the recommendation of the tumor conference with knowledge of the EPclin result (post-test) was compared with the original recommendation (pre-test).

Statistical analysis

The primary objective was to assess differences in pre- and post-test therapy decisions. The secondary objectives of this retrospective study were (i) to evaluate clinico-pathological factors influencing the decision for the application of EPclin in HR-positive, HER2-negative early breast cancer patients with 0–3 involved axillary lymph nodes as well as the relationship between EndoPredict® and (ii) clinico-pathological variables, and (iii) molecular subtypes. Patient characteristics were presented in absolute and relative numbers. Comparisons between clinico-pathological factors and the results of the EndoPredict® were calculated using the Pearson’s chi-squared test (χ²-test) and Fisher’s exact test. Afterwards, univariate logistic regressions were performed including one of the following independent variables: age at diagnosis, menopausal status (pre-/perimenopausal vs. postmenopausal),
breast tumor size (pT1 vs. pT2 vs. pT3/pT4), axillary nodal status (pN0 vs. pN1), histological grade of differentiation (grade I vs. II vs. III), Ki-67 (<20% vs. 20–40% vs. >40%), ER status (positive vs. negative), PR status (positive vs. negative), and cancer molecular subtype (luminal A-like/Ki-67 <20% vs. luminal B-like/Ki-67 ≥20%). All independent variables were included for the multivariable analysis and the best fitting model was obtained utilizing backward selection by AIC. All associations were reported as odds ratios (OR) including their 95% confidence interval (CI) and $P$ values. All $P$ values are two-sided, and $P<0.05$ was considered significant. As all analyses were explorative and not adjusted for multiple testing, the $P$ values should be interpreted with caution and in connection with the effect estimates. Statistical analyses were performed using R (v. 3.6.2) [R Development Core Team (2008). R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. ISBN 3-900051-07-0, https://www.R-project.org.]. The study was approved by the Research Ethics Committee of the University Medical Center Mainz (Mainz, Germany). Informed consent was obtained from all patients.

Results

Patient population

950 patients with hormone receptor-positive, HER2-negative early breast cancer, with a maximum of three positive axillary lymph nodes, were screened. 869 (91.5%) patients were eligible for evaluation. Patients were excluded due to bilateral breast cancer ($n=18$) or missing clinico-pathological data or missing EndoPredict® result ($n=63$) (Fig. 1). In 156 (18.0%) patients, EndoPredict® was performed based on the clinician’s decision. EndoPredict® test results classified 67 patients (42.9%) as low-risk and 89 patients (57.1%) as high-risk (Fig. 2). In the entire cohort, adjuvant chemotherapy was recommended for 271 patients (31.2%).

Impact of EndoPredict® for adjuvant treatment decisions

According to the EPclin results, chemotherapy was recommended in 57.1% ($n=89$) whilst no chemotherapy was advised in 42.9% ($n=67$) (post-test). Without knowledge...
of EPclin, chemotherapy was recommended in 81 patients (51.9%) and not recommended in 75 patients (48.1%) (pre-test) (Appendix 1, Only Online). In detail, 104 patients (66.7%) received the identical treatment recommendation pre- and post-test. In 33.3% (n = 52), however, there was an adjustment of the pre-test therapy decision after knowledge of EPclin became available. In 30 patients (19.2%), adjuvant chemotherapy was recommended post-test compared to the pre-test “low risk” classification based on clinical-pathological parameters (n = 75). In 45 patients there was no change to an endocrine therapy recommendation. Conversely, in 22 patients (14.1%) adjuvant chemotherapy was omitted (EPclin “low risk”). Based on clinico-pathological classification, adjuvant chemotherapy was advised in 81 patients, i.e., adjuvant chemotherapy was furthermore recommended in 59 patients (Appendix 1, Only Online) (Fig. 3).

Among patients who had not received an EndoPredict® test (n = 713), chemotherapy was recommended to 182 patients (25.5%). For the remaining 531 patients (74.5%) chemotherapy was not advised.

**Influence of clinico-pathological factors on the application of EndoPredict®**

The traditional clinico-pathological prognostic factors were examined in univariate analysis with respect to the decision to perform EPclin (Table 1). Multivariate analysis included all parameters that showed a significant result in the univariate testing: age at primary breast surgery (P < 0.001), tumor size (P = 0.014), nodal status (P < 0.001), histological grade (P < 0.001), Ki-67 (P < 0.001) and molecular subtype (P < 0.001) [21]. Younger age (OR 0.498, 95% CI 0.335–0.734, P < 0.001), smaller tumor size (OR 0.071, 95% CI 0.004–0.354, P = 0.011), positive nodal status (OR 1.674, 95% CI 1.080–2.587, P = 0.021), low/intermediate histological grade (OR 0.432, 95% CI 0.192–0.984, P = 0.043), and intermediate Ki-67 (OR 3.599, 95% CI 2.370–5.508, P < 0.001) retained a higher chance for EPclin (Table 2).

**Relationship between EndoPredict® and clinico-pathological variables**

The common clinico-pathological prognostic factors were examined in all patients with available EPclin (n = 156) using univariate logistic regression with respect to the EndoPredict® result. Only pT2 tumor size (P = 0.007) and positive nodal status (P = 0.031) showed a higher chance for EPclin risk category (Table 3). The sample size for pT3/4 tumor size (n = 1) and for higher Ki-67 (n = 5) were too small for stable estimation.

**Relationship between EndoPredict® and molecular subtypes**

EndoPredict® classified 33 Luminal A-like tumors (49.3% of all low-risk tumors) as low risk and 30 (33.7% of all high-risk tumors) as high risk. In Luminal B-like tumors, EPclin allocated 34 cases (50.7%) in the low risk and 59 cases (66.3%) in the high risk category, respectively (P = 0.051) (Table 3).

**Discussion**

In our analysis of 869 consecutive HR-positive, HER2-negative breast cancer patients, gene expression analysis with EPclin was performed in 18.0% for adjuvant treatment decisions. In 33.3% use of EPclin led to a change in treatment recommendations.

![Fig. 3 Changes in therapy decision](image-url)
Using EPclin, 67 (42.9%) patients were assigned low-risk and 89 (57.1%) high-risk. Remarkably, the distribution between the low and high risk category was different from other studies. In contrast to our results, more low-risk patients were observed in the cohort used for EPclin validation. In this study, 63% of 1702 patients were classified as low-risk and 37% as high-risk [18, 22]. Similarly, the low-risk group of postmenopausal patients from the TransATAC study (58.8%) was also considerably larger than in the present study [14]. However, both studies included patients of a prospective study population for whom mainly endocrine therapy was indicated. Therefore the ABCSG-6/-8 and the TransATAC study cohort represented a completely different study population than the current real-world population in our study. Most likely, a clinical pre-selection to favorable risk might explain this difference. Conversely, another study

Table 1  Associations between EndoPredict® and clinico-pathological characteristics (n = 869)

|                        | Total number of patients (n = 869) | EP test | P value |
|------------------------|------------------------------------|---------|---------|
|                        | Not done (n = 713) | Done (n = 156) |         |
| Age at primary breast surgery | < 0.001 |
| Mean (years) | 61 | 62 | 57 |
| Minimum (years) | 28 | 28 | 32 |
| Maximum (years) | 106 | 106 | 75 |
| Menopausal status | 0.634 |
| Pre-/perimenopausal | 244 (28.1%) | 198 (27.8%) | 46 (29.5%) |
| Postmenopausal | 625 (71.9%) | 515 (72.2%) | 110 (70.5%) |
| Histopathologic classification | 0.394 |
| Invasive carcinoma of no special type (NST) | 678 (78.0%) | 557 (78.1%) | 121 (77.6%) |
| Invasive lobular carcinoma | 120 (13.8%) | 96 (13.5%) | 24 (15.4%) |
| Others | 71 (8.2%) | 60 (8.4%) | 11 (7.1%) |
| Tumor size | 0.014 |
| pT1 | 558 (65.2%) | 466 (65.4%) | 92 (59.0%) |
| pT2 | 276 (31.8%) | 213 (29.9%) | 63 (40.4%) |
| pT3/pT4 | 35 (4.0%) | 34 (4.8%) | 1 (0.6%) |
| Nodal status | < 0.001 |
| pN0 | 686 (78.9%) | 582 (81.6%) | 104 (66.7%) |
| pN1 | 183 (21.1%) | 131 (18.4%) | 52 (33.3%) |
| Histological grade | < 0.001 |
| G1 | 244 (28.1%) | 227 (31.8%) | 17 (10.9%) |
| G2 | 466 (53.6%) | 353 (49.5%) | 113 (72.4%) |
| G3 | 159 (18.3%) | 133 (18.7%) | 26 (16.7%) |
| Estrogen receptor status | 0.934 |
| Positive | 863 (99.3%) | 708 (99.3%) | 155 (99.4%) |
| Negative | 6 (0.7%) | 5 (0.7%) | 1 (0.6%) |
| Progesterone receptor status | 0.825 |
| Positive | 784 (90.2%) | 644 (90.3%) | 140 (89.7%) |
| Negative | 85 (9.8%) | 69 (9.7%) | 16 (10.3%) |
| Ki-67 | < 0.001 |
| Low (< 20%) | 547 (62.9%) | 484 (67.9%) | 63 (40.4%) |
| Intermediate (20–40%) | 268 (30.8%) | 180 (25.2%) | 88 (56.4%) |
| High (> 40%) | 54 (6.2%) | 49 (6.9%) | 5 (3.2%) |
| Molecular subtype | < 0.001 |
| Luminal A-like | 547 (62.9%) | 484 (67.9%) | 63 (40.4%) |
| Luminal B-like | 322 (37.1%) | 229 (32.1%) | 93 (59.6%) |
| EndoPredict® risk category |      |
| Low (≤ 3.3) | 67 | 0 | 67 (42.9%) |
| High (> 3.3) | 89 | 0 | 89 (57.1%) |

Bold indicates P < 0.05
investigating two different chemotherapy regimens in early breast cancer assigned only 25% to the low-risk group [16]. Nonetheless, a risk distribution comparable to our results was shown by Müller et al., who evaluated all EndoPredict® requests in a single pathology institute during one year. 46.4% were classified as low-risk and 53.5% as high-risk [23]. Only a minority of the patients in our study (n = 8) was also included in the aforementioned analysis. However, in contrast to their analysis, our study included all eligible HR-positive, HER2-negative patients with 0–3 involved axillary lymph nodes over a period of five years, providing a more unbiased view on the use of EPclin in early breast cancer. The indication to perform a gene expression assay may also be influenced by the experience of the physicians ordering the test, in particular by their individual threshold for administration of chemotherapy. This selection bias could explain the different risk-profiles in different EndoPredict® cohorts.

In our analysis, the decision to perform EPclin was multivariable associated with younger age, smaller tumor size, positive nodal status, low/intermediate histological grade and intermediate Ki-67 in multivariate testing. The younger median age in the EPclin-tested cohort (57 years) was similar to the observations in other studies [23–25]. The younger age in breast cancer patients is an unfavorable prognostic factor. More aggressive tumor therapy, however, is associated with the increased probability of long-term side effects. Therefore, a reason for the additional performance of a multigene assay in younger patients is primarily whether chemotherapy can be safely avoided. Conversely if a luminal-like tumor is present in elderly patients, endocrine therapy alone is more likely. Histological grade and the proliferative marker Ki-67 are important variables for the risk classification of early breast cancer. Both variables, however, have considerable inter-laboratory and observer variability [26, 27]. Therefore, it is not surprising that, similar to other reports, the additional gene expression analysis is most often used in tumors of intermediate histological grade [23] and intermediate Ki-67. Median Ki-67 was 20% in the EP cohort v. 10% in the untested patient group. Accordingly, in a large study by Nitz and co-workers, patients with intermediate Ki-67 (> 10% to < 40%) were most likely to benefit from a gene expression test [28]. Histological grade as an influencing factor for use of multigene assays has also been observed in other settings with an increased percentage of G2 tumors in cohorts tested with Oncotype DX® [25, 29]. In the assessment of the relationship between EndoPredict® and clinico-pathological variables only T2 tumor size (P = 0.007) and positive nodal status (P = 0.031) showed a higher chance for EPclin risk category (Table 3). Among the multigene tests, the EPclin test is the only one that integrates both clinical parameters (tumor size and nodal status) in addition to molecular genetic parameters. Thus, our results underline the strong influence of these two clinical factors on the patient’s individual risk profile with breast cancer.

We have shown that in 33.3% EPclin led to changes in the therapy recommendation. Comparable rates of therapy changes have been observed in other studies. Penault-Llorca et al. showed in a prospective study that EPclin resulted in a change of therapy recommendation in one third of patients [13]. Similarly, Müller et al. reported a change in 37.7% [23]. In our study, EPclin led in 19.2% to therapy escalation and in 14.1% to de-escalation. Interestingly, both studies described therapy de-escalation considerably more often than escalation, presumably reflecting both differences in the patient population as well as in the willingness to recommend adjuvant chemotherapy in intermediate-risk HR-positive, HER2-negative early breast cancer. In line with the latter argument is the fact that in our patients without gene expression profiling the rate of chemotherapy recommendation was only 25.5%. Differences of 30–50% between pre- and post-test treatment recommendations were also shown in studies using Oncotype DX® and the 70-gene signature (MammaPrint®) [12, 30–32].

Our study has some strengths and limitations. A potential weakness is that our study was retrospective and performed in a single certified breast cancer center. Furthermore, patient preferences were not addressed. An extension of our study endpoints to patient-related questions (e.g., expectations and wishes on a gene signature test, impact of the test result for the patient), could have led to valuable
However, a major strength of our study is the consecutive inclusion of patients which enables the assessment of EPclin in the context of well-established clinic-pathological factors. To the best of our knowledge, this is the first study dealing with this topic using real-world data. In our study post-test results of EndoPredict® were the reason for a change in clinical decisions in more than 1/3 of tested patients. This is a valuable finding in a "real life" use of this multigene test. With an increasing use of multigene assays it is of particular interest for the indication and evaluation of the EPclin results to know if the clinical treatment decision changes.

In conclusion, EPclin was only performed in a subset of consecutive HR-positive, HER2-negative early breast cancer patients. However, EPclin resulted in a considerable change in therapy recommendations in one third of patients, potentially reducing over- and under-treatment in early breast cancer patients.

| Table 3 Association between EndoPredict® result and patient and tumor characteristics (n = 156) |
|-----------------------------------------------|-----------------|-----------------|-----------------|-----------------|
|                                              | EndoPredict® done (n = 156) | EndoPredict® result Low risk (n = 67) | EndoPredict® result High risk (n = 89) | P value [OR] |
| Age at primary breast surgery                |                                              |                                              |                                              |              |
| Mean (years)                                 | 57                                           | 70                                           | 56                                           |              |
| Minimum (years)                              | 32                                           | 68                                           | 35                                           |              |
| Maximum (years)                              | 75                                           | 72                                           | 75                                           |              |
| ≤ 60 years                                    | 90 (57.7%)                                    | 37 (55.2%)                                    | 53 (59.6%)                                    | Ref          |
| > 60 years                                    | 66 (42.3%)                                    | 30 (44.8%)                                    | 36 (40.4%)                                    | 0.588 [0.838]|
| Menopausal status                            |                                              |                                              |                                              |              |
| Pre-/perimenopausal                          | 46 (29.5%)                                    | 17 (25.4%)                                    | 29 (32.6%)                                    | Ref          |
| Postmenopausal                               | 110 (70.5%)                                   | 50 (74.6%)                                   | 60 (67.4%)                                   | 0.329 [0.703]|
| Histopathologic classification               |                                              |                                              |                                              |              |
| Invasive carcinoma of no special type (NST)  | 121 (77.6%)                                   | 53 (79.1%)                                   | 68 (76.4%)                                   | Ref          |
| Invasive lobular carcinoma                   | 24 (15.4%)                                    | 11 (16.4%)                                   | 13 (16.4%)                                   | 0.855 [0.921]|
| Others                                       | 11 (7.1%)                                     | 3 (4.5%)                                     | 8 (9.0%)                                     | 0.297 [2.078]|
| Tumor size                                   |                                              |                                              |                                              |              |
| pT1                                          | 92 (59.0%)                                    | 48 (71.6%)                                   | 44 (49.3%)                                   | Ref          |
| pT2                                          | 63 (40.4%)                                    | 19 (28.4%)                                   | 44 (49.3%)                                   | 0.007 [2.526]|
| pT3/pT4                                      | 1 (0.6%)                                      | 0                                            | 1 (1.1%)                                     |              |
| Nodal status                                 |                                              |                                              |                                              |              |
| pN0                                          | 104 (66.7%)                                   | 51 (76.1%)                                   | 53 (59.6%)                                   | Ref          |
| pN1                                          | 52 (33.3%)                                    | 16 (23.9%)                                   | 36 (40.4%)                                   | 0.031 [2.165]|
| Histological grade                           |                                              |                                              |                                              |              |
| G1                                           | 17 (10.9%)                                    | 9 (13.4%)                                    | 8 (9.0%)                                     | Ref          |
| G2                                           | 113 (72.4%)                                   | 47 (70.1%)                                   | 66 (74.2%)                                   | 0.381 [1.580]|
| G3                                           | 26 (16.7%)                                    | 11 (16.4%)                                   | 15 (16.9%)                                   | 0.495 [1.534]|
| Estrogen receptor status                     |                                              |                                              |                                              |              |
| Positive                                     | 155 (99.4%)                                   | 67 (100%)                                    | 88 (98.9%)                                   | Ref          |
| Negative                                     | 1 (0.6%)                                      | 0                                            | 1 (1.1%)                                     | 0.987        |
| Progesterone receptor status                 |                                              |                                              |                                              |              |
| Positive                                     | 140 (89.7%)                                   | 63 (94.0%)                                   | 77 (86.5%)                                   | Ref          |
| Negative                                     | 16 (10.3%)                                    | 4 (6.0%)                                     | 12 (13.5%)                                   | 0.136 [2.454]|
| Ki-67                                        |                                              |                                              |                                              |              |
| Low (< 20%)                                   | 63 (40.4%)                                    | 33 (49.3%)                                   | 30 (33.7%)                                   | Ref          |
| Intermediate (20–40%)                        | 88 (56.4%)                                    | 33 (49.3%)                                   | 55 (62.8%)                                   | 0.070 [1.833]|
| Higher (> 40%)                                | 5 (3.2%)                                      | 1 (1.5%)                                     | 4 (4.5%)                                     |              |
| Molecular subtype                            |                                              |                                              |                                              |              |
| Luminal A-like                               | 63 (40.4%)                                    | 33 (49.3%)                                   | 30 (33.7%)                                   | Ref          |
| Luminal B-like                               | 93 (59.6%)                                    | 34 (50.7%)                                   | 59 (66.3%)                                   | 0.051 [1.909]|

Bold indicates P < 0.05
Acknowledgements  Open Access funding provided by Projekt DEAL.

Funding No funding was received.

Compliance with ethical standards

Conflict of interest K. Almstedt received speaker honoraria from Roche Pharma AG, Pfizer Pharma GmbH and AstraZeneca. M. Schmidt received honoraria for speaker or consultancy role from AMGEN, AstraZeneca, Eisai, Lilly, Myelo Therapeutics, Novartis, Pantarhei Bioscience, Pfizer, and Roche. He received research funding from AstraZeneca, BioNTech, Eisai, Genentech, Myelo Therapeutics, Novartis, Pantarhei Bioscience, Pfizer, Pierre Fabre, and Roche. He received travel reimbursement from Pfizer and Roche. M. Otto received honoraria for speaker or consultancy role from AstraZeneca, Boehringer-Ingelheim, Roche and Sividon. He received travel reimbursement from AstraZeneca, and Boehringer-Ingelheim. M.J. Battista received honoraria for speaker or consultancy role from AstraZeneca, MSD, PharmaMar, Roche Pharma AG, TEVA, Tesaro. He received travel reimbursement from Celgene, PharmaMar and Pierre Fabre. S. Krajnak received honoraria from Roche. He received research funding from Novartis. He received travel reimbursement from PharmaMar. A. Hasenburg received honoraria from AstraZeneca, Celigen, MedConcept Gm, Med update GmbH, Mediculits, Pfizer, Promedics GmbH, Pierre Fabre, Softconsult, Roche Pharma AG, Streamedup!GmbH and Tesaro Bio Germany GmbH. She is a member of the advisory board of PharmaMar, Promedics GmbH, Pierre Fabre Pharma GmbH, Roche Pharma AG and Tesaro Bio Germany GmbH. She received research funding from Celgene, C. Denkert has been cofounder and shareholder of Sividon Diagnostics (now Myriad), has received speaker honoraria from Teva, Novartis, Pfizer, Roche, Amgen and has been consultant for MSD Oncology, Amgen, Roche and Daiichi-Sankyo. All other authors declare that they have no conflicts of interest.

Ethical approval All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

Informed consent Informed consent was obtained from all individual participants included in the study.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

References

1. Peto R, Davies C, Godwin J, Gray R, Pan HC, Clarke M, Cutter D, Darby S, McGale P, Taylor C, Wang YC, Bergh J, Di Leo A, Albin K, Swain S, Piccart M, Pritchard K (2012) Comparisons between different polychemotherapy regimens for early breast cancer: meta-analyses of long-term outcome among 100,000 women in 123 randomised trials. Lancet 379(9814):432–444. https://doi.org/10.1016/S0140-6736(11)61625-5
2. Harris LN, Ismaila N, McShane LM, Andre F, Collyar DE, Gonzalez-Angulo AM, Hammond EH, Kuderer NM, Liu MC, Mennel RG, Van Poznak C, Bast RC. Hayes DF (2016) Use of biomarkers to guide decisions on adjuvant systemic therapy for women with early-stage invasive breast cancer: American Society of Clinical Oncology oncologic practice guideline. J Clin Oncol 34(10):1134–1150. https://doi.org/10.1200.jco.2015.65.2289
3. Curigliano G, Burstyn HJ, Winer EP, Ganot M, Dubsky P, Loibl S, Colleoni M, Regan MM, Piccart-Gebhart M, Senn HJ, Thurlimann B, Andre F, Baselga J, Bergh J, Bonnello H, Brucker SY, Cardoso F, Carey L, Ciruelos E, Cuzick J, Denkert C, Di-Leo A, Ejlertsen B, Francis P, Galimberti V, Garber J, Gullouglu B, Goodwin P, Harbeck N, Hayes DF, Huang CS, Huober J, Hussein K, Jassem J, Jiang Z, Karlsson P, Morrow M, Orecchia R, Osborne KC, Pagani O, Partridge AH, Pritchard K, Ro J, Rutgers EJT, Sedlmayer F, Semiglazov V, Shao Z, Smith I, Toi M, Tutt A, Viale G, Watanabe T, Whelan TJ, Xu B (2017) De-escalating and escalating treatments for early-stage breast cancer: the St Galen International Expert Consensus Conference on the Primary Therapy of Early Breast Cancer 2017. Annals of Oncology 28 (8):1700–1712
4. Simon RM, Paik S, Hayes DF (2009) Use of archived specimens in evaluation of prognostic and predictive biomarkers. J Natl Cancer Inst 101(21):1446–1452. https://doi.org/10.1093/jnci/djp335
5. Krop I, Ismaila N, Andre F, Bast RC, Barlow W, Collyar DE, Hammond ME, Kuderer NM, Liu MC, Mennel RG, Van Poznak C, Wolff AC, Stearns V (2017) Use of biomarkers to guide decisions on adjuvant systemic therapy for women with early-stage invasive breast cancer: American Society of Clinical Oncology Clinical Practice Guideline Focused Update. J Clin Oncol 35(24):2838–2847. https://doi.org/10.1200.jco.2017.74.0472
6. Andre F, Ismaila N, Henry NL, Somerfield MR, Bast RC, Barlow W, Collyar DE, Hammond ME, Kuderer NM, Liu MC, Van Poznak C, Wolff AC, Stearns V (2019) Use of biomarkers to guide decisions on adjuvant systemic therapy for women with early-stage invasive breast cancer: ASCO Clinical Practice Guideline Update-Integration of Results from TAILORx. J Clin Oncol 37(22):1956–1964. https://doi.org/10.1200/jco.19.00945
7. Albain K, Barlow W, Shk S (2010) Prognostic and predictive value of the 21-gene recurrence score assay in postmenopausal women with node-negative, oestrogen-receptor-positive breast cancer on chemotherapy: a retrospective analysis of a randomised trial. Lancet Oncol 11:55–65
8. Dowsett M, Cuzick J, Wale C, Forbes J, Mallon EA, Salter J, Quinn E, Dunbier A, Baum M, Buzdar A, Howell A, Bugarini R, Baehner FL, Shk S (2010) Prediction of distant recurrence: using the 21-gene recurrence score in node-negative and node-positive postmenopausal patients with breast cancer treated with anastrozole or tamoxifen: a TransATAC study. J Clin Oncol 28(11):1829–1834. https://doi.org/10.1200/jco.2009.24.4798
9. Sparano JA, Gray RJ, Makower DF, Pritchard KL, Albain KS, Hayes DF, Geyer CE Jr, Dees EC, Goetz MP, Olson JA Jr, Lively T, Badve SS, Saphner TJ, Wagner LI, Whelan TJ, Ellis MJ, Paik S, Wood WC, Ravdin PM, Keane MM, Gomez Moreno HL, Reddy PS, Goggins TF, Mayer IA, Brueskyl AM, Labrador Bioscience, Pfizer, Toppmeyer DL, Kalamani VG, Berenberg JL, Abrams J, Sledge GW Jr (2018) Adjuvant therapy for hormone receptor positive, HER2-negative, node-negative early breast cancer: ASCO Clinical Practice Guideline Update-Integration of Results from TAILORx. J Clin Oncol 37(8):1700–1712
10. Sparano JA, Gray RJ, Makower DF, Pritchard KL, Albain KS, Hayes DF, Geyer CE Jr, Dees EC, Perez EA, Olson JA Jr, Zuzewski J, Lively T, Badve SS, Saphner TJ, Wagner LI, Whelan TJ, Ellis MJ, Paik S, Wood WC, Ravdin P, Keane MM, Gomez Moreno HL, Reddy PS, Goggins TF, Mayer IA, Brueskyl AM.
11. Sparano JA, Gray RJ, Ravdin PM, Makower DF, Pritchard KL, Albain KS, Hayes DF, Geyer CE Jr, Dees EC, Goetz MP, Olson JA Jr, Lively T, Badve SS, Saphner TJ, Wagner LI, Whelan TJ, Ellis MJ, Paik S, Wood WC, Keane MM, Gomez Moreno HL, Reddy PS, Goggin TF, Mayer IA, Brufsky AM, Toppmeyer DL, Kaklamani VG, Atkins JN, Berenberg JL, Sledge GW (2015) Prospective validation of a 21-gene expression assay in breast cancer. N Engl J Med 373(21):2005–2014. https://doi.org/10.1056/NEJMoa1510764

12. Cardoso F, Vant-Veer LJ, Bogaerts J, Slaets L, Viale G, Delaloge S, Pierga JY, Brain E, Cauzaert S, DeLorenzi M, Glaz AM, Golfinopoulos V, Gouliioti T, Knox S, Matos E, Meulemans B, Neijenhuis PA, Nitz U, Passalaquara R, Ravdin P, Rubio IT, Saghatchian M, Sotiriou C, Stork L, Strauss JH, Thomas G, Thomson AM, van der Hoeven JM, Vuylsteke P, Bernard N, Tryfonidis K, Rutgers E, Piccatt M (2016) 70-gene signature as an aid to treatment decisions in early-stage breast cancer. N Engl J Med 375(8):717–729. https://doi.org/10.1056/NEJMoa1602253

13. Renault-Llorca F, Kwiatkowski F, Arnaud A, Levy C, Leheur M, Sestak I, Martin M, Dubsky P, Kronenwett R, Filipits M, Rudas M, Jakesz R, Dubsky P, Fitzal F, Singer CF, Buus R, Sestak I, Kronenwett R, Denkert C, Dubsky P, Krappmann K, Ruiz-Borrego M, Golfinopoulos V, Gouliioti T, Knox S, Matos E, Meulemans B, Neijenhuis PA, Nitz U, Passalaquara R, Ravdin P, Rubio IT, Saghatchian M, Smilde TJ, Toppmeyer DL, Kaklamani VG, Atkins JN, Berenberg JL, Abrams J, Sledge GW Jr (2019) Clinical and genomic risk to guide the use of adjuvant therapy for breast cancer. N Engl J Med 380(25):2395–2405. https://doi.org/10.1056/NEJMoa1904819

14. Toppmeyer DL, Kaklamani VG, Atkins JN, Berenberg JL, Sledge GW (2015) Prospective validation of a 21-gene expression assay in breast cancer. N Engl J Med 373(21):2005–2014. https://doi.org/10.1056/NEJMoa1510764

15. Martin M, Brase JC, Calvo L, Krappmann K, Ruiz-Borrego M, Fisch K, Ruiz A, Weber KE, Munarriz B, Petry C, Rodriguez CA, Kronenwett R, Crespo C, Alba E, Carrasco E, Casas M, Caballero R, Rodriguez-Lescure A (2014) Clinical validation of the EndoPredict test in node-negative, chemotherapy-treated ER+/HER2- breast cancer patients: results from the GEICAM 9906 trial. Breast Cancer Research BCR 16(2):R38. https://doi.org/10.1186/bcr3642

16. Fitzal F, Filipits M, Rudas M, Greil R, Dietze O, Samonigg H, Lax S, Herz W, Dubsky P, Bartsch R, Kronenwett R, Gnant M (2015) The genomic expression test EndoPredict is a prognostic tool for identifying risk of local recurrence in postmenopausal endocrine receptor-positive, her2neu-negative breast cancer patients randomised within the prospective ABCSG 8 trial. Br J Cancer 112(8):1405–1410. https://doi.org/10.1038/bjc.2015.98

17. Muller BM, Keil E, Lehmann A, Winzer KJ, Richter-Ehrenstein C, Prinzler J, Bangemann N, Reales A, Stadie S, Schoenevag W, Eucker J, Schmidt M, Lippek F, Johrens K, Pahl S, Sinn BV, Budczies J, Dietel M, Denkert C (2013) The EndoPredict gene-expression assay in clinical practice—performance and impact on clinical decisions. PLoS ONE 8(6):e68252. https://doi.org/10.1371/journal.pone.0068252

18. Ray GT, Mandelblatt J, Habel LA, Ramsey S, Kushi LH, Li Y, Lieu TA (2016) Breast cancer multigene testing trends and impact on chemotherapy use. Am J Manag Care 22(5):e153–160

19. Dinan MA, Mi X, Reed SD, Hirsch BR, Lyman GH, Curtis LH (2015) Initial trends in the use of the 21-gene recurrence score assay for patients with breast cancer in the medicare population, 2005–2009. JAMA Oncol 1(2):158–166. https://doi.org/10.1001/jamaoncol.2015.43

20. Furburn PN, Taub N, Assmann KJ, Banfi G, Cosyns JP, Dormann AM, Hill CM, Kapper SK, Waldherr R, Laurinavicius A, Marcussen N, Martins AP, Nogueira M, Regele H, Seron D, Carrera AM, Hill CM, Kapper SK, Waldherr R (2016) Initial trends in the use of the 21-gene recurrence score assay for patients with breast cancer in the medicare population, 2005–2009. JAMA Oncol 1(2):158–166. https://doi.org/10.1001/jamaoncol.2015.43

21. Wang M, McLaren S, Jeyathasan R, Allison BM, Ireland A, Kang A, Meehan K, Thomas C, Robinson C, Combrinck M, Harvey J, Sterrett G, Dessauvage B (2019) Laboratory validation studies in Ki-67 digital image analysis of breast carcinoma: a pathway to routine quality assurance. Pathology. https://doi.org/10.1016/j.pathol.2018.12.416

22. Nitz U, Gluz O, Christgen M, Kates RE, Clemens M, Malter W, Salehi F, Krabisch P, Just M, Augustin D, Liedtke C, Chao C, Shuk S, Wuerstlein R, Kreipe HH, Harbeck N (2017) Reducing chemotherapy use in clinically high-risk, genomically low-risk ER- and ER+/HER2- breast cancer patients: results from the GEICAM 9906 trial. Breast Cancer Research BCR 16(2):R38. https://doi.org/10.1186/bcr3642
the prospective, randomised phase 3 West German Study Group (WSG) PlanB trial. Breast Cancer Res Treat 165(3):573–583. https://doi.org/10.1007/s10549-017-4358-6

29. Zhu X, Dent S, Paquet L, Zhang T, Graham N, Song X (2014) Factors influencing Oncotype DX use in the management of early breast cancer: a single centre experience. Eur J Cancer 50(15):2544–2549. https://doi.org/10.1016/j.ejca.2014.07.010

30. Lo SS, Mumby PB, Norton J, Rychlik K, Smerage J, Kash J, Chew HK, Gaynor ER, Hayes DF, Epstein A, Albain KS (2010) Prospective multicenter study of the impact of the 21-gene recurrence score assay on medical oncologist and patient adjuvant breast cancer treatment selection. J Clin Oncol 28(10):1671–1676. https://doi.org/10.1200/jco.2008.20.2119

31. Augustovski F, Soto N, Caporale J, Gonzalez L, Gibbons L, Ciapponi A (2015) Decision-making impact on adjuvant chemotherapy allocation in early node-negative breast cancer with a 21-gene assay: systematic review and meta-analysis. Breast Cancer Res Treat 152(3):611–625. https://doi.org/10.1007/s10549-015-3483-3

32. Carlson JJ, Roth JA (2013) The impact of the Oncotype Dx breast cancer assay in clinical practice: a systematic review and meta-analysis. Breast Cancer Res Treat 141(1):13–22. https://doi.org/10.1007/s10549-013-2666-z

Publisher's Note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.