

Distribution of the 21-Gene Breast Recurrence Score in Patients with Primary Breast Cancer in Germany

Verteilung des 21-Gen-Rezidiv-Scores bei Patientinnen mit primärem Mammakarzinom in Deutschland



Authors

Vincent P. Walter¹, Florin-Andrei Taran¹, Markus Wallwiener², Armin Bauer¹, Eva-Maria Grischke¹, Christina Barbara Walter¹, Markus Hahn¹, Sara Y. Brucker¹, Andreas Daniel Hartkopf¹

Affiliations

- 1 Department of Women's Health, University of Tübingen, Tübingen, Germany
- 2 Department of Obstetrics and Gynecology, University of Heidelberg, Heidelberg, Germany

Key words

21-gene breast recurrence score, breast cancer, multigene assays, Oncotype DX

Schlüsselwörter

21-Gen-Rezidiv-Score, Mammakarzinom, Multigen-Assays, Oncotype DX

received 4. 6. 2019

revised 1. 2. 2020

accepted 2. 2. 2020

Bibliography

DOI <https://doi.org/10.1055/a-1111-8734>

Geburtsh Frauenheilk 2020; 80: 619–627 © Georg Thieme Verlag KG Stuttgart · New York | ISSN 0016-5751

Correspondence

Prof. Andreas Daniel Hartkopf
Department of Women's Health, University of Tuebingen
Calwerstraße 7, 72076 Tübingen, Germany
andreas.hartkopf@med.uni-tuebingen.de

ABSTRACT

Background Multigene assays are being used increasingly to aid in decision-making about chemotherapy in breast cancer. Here, we present the 21-gene recurrence score (RS) of patients tested in routine clinical practice in Germany.

Patients and Methods In a retrospective analysis, 4695 patients with hormone receptor-positive and HER2-negative early breast cancer (pT1–3, pN0–1, M0) were included in whom RS testing was conducted in Germany between No-

vember 2015 and July 2018. RS groups as defined in the TAILORx trial (RS result 0–10; 11–25; 26–100) were used.

Results Of these patients, 21% were assigned to the low RS group, 63% to the midrange RS group, and 15% to the high RS group. 1772 (81%) of 2175 node-negative patients over 50 years of age were grouped either into the low RS group or the midrange RS group. The portion of patients with a low or midrange RS was 90% among node-positive patients (1284 of 1432 patients), 79% among patients with Ki-67-high ($\geq 20\%$) tumors (1829 of 2310 patients), 86% vs. 70% among patients with G2 and G3 tumors (3244 of 3762 patients and 368 of 522 patients), respectively, 88% among patients with a tumor size of > 5 cm (140 of 159 patients), and 82% among node-negative patients at high clinical risk (1110 of 1352).

Conclusions The distribution of the 21-gene RS in German patients that were tested in routine clinical practice indicates that, according to the results of the TAILORx trial, chemotherapy may not be beneficial in most of these.

ZUSAMMENFASSUNG

Einleitung Multigen-Assays werden zunehmend als Entscheidungshilfe für eine Chemotherapie beim Mammakarzinom verwendet. Wir stellen hier den 21-Gen-Recurrence-Score (RS) von Patientinnen mit Brustkrebs vor, die routinemäßig in Deutschland untersucht wurden.

Patientinnen und Methoden 4695 Patientinnen mit hormonrezeptorpositivem und HER2-negativem Brustkrebs im Frühstadium (pT1–3, pN0–1, M0) wurden einer retrospektiven Analyse unterzogen. Bei diesen Patientinnen wurde in Deutschland zwischen November 2015 und Juli 2018 der Genexpressionstest Oncotype DX zur Ermittlung des Recurrence-Scores durchgeführt. Die Klassifikation der RS-Gruppen erfolgte gemäß der TAILORx Studie (RS: 0–10; 11–25; 26–100).

Ergebnisse Von diesen Patientinnen wurden 21% in die niedrige RS-Gruppe, 63% in die mittlere RS-Gruppe, und 15% in die hohe RS-Gruppe eingeteilt. 1772 (81%) von 2175 Patientinnen im Alter von über 50 Jahren und ohne Lymphknotenbefall wurden entweder in die niedrige oder die mittlere RS-

Gruppe eingeteilt. Der Prozentsatz an Patientinnen mit einem niedrigen oder mittleren RS betrug 90% bei Patientinnen ohne Lymphknotenbefall (1284 von 1432 Patientinnen), 79% bei Patientinnen mit einem hohen ($\geq 20\%$) Ki-67-Wert (1829 von 2310 Patientinnen), 86% bzw. 70% bei Patientinnen mit G2- bzw. G3-Tumoren (3244 von 3762 Patientinnen bzw. 368 von 522 Patientinnen), 88% bei Patientinnen mit einem Tumordurchmesser von >5 cm (140 von 159 Patientinnen),

und 82% bei Patientinnen ohne Lymphknotenbefall, aber mit einem hohen klinischen Risiko (1110 von 1352).

Ergebnisse Die Verteilung des 21-Gens RS bei deutschen Patientinnen, die in der klinischen Routinepraxis getestet wurden, deutet darauf hin, dass gemäß den Ergebnissen der TAILORx-Studie die Chemotherapie bei den meisten dieser Patientinnen keinen Nutzen hat.

Abbreviations

HER2	human epidermal growth factor 2
HR	hormone receptor
IQR	interquartile range
RS	21-gene recurrence score

Key Message

The 21-gene breast recurrence score classified 83% of node-negative and 90% of node-positive patients tested in routine clinical practice in Germany as low or midrange RS.

Introduction

Breast cancer is the most common cancer and remains the number one cause of cancer-related deaths among women in the US [1] and Europe [2]. With advances in diagnostics and therapy, however, breast cancer mortality has improved remarkably over the last few decades [1]. Amid these developments, increasing efforts are being made to distinguish between patients who are likely to benefit from adjuvant chemotherapy and those who can be spared the toxic side effects while retaining their favorable prognosis [3, 4].

As weighing the advantages and disadvantages of chemotherapy is challenging, especially in hormone receptor (HR)-positive, human epidermal growth factor 2 (HER2)-negative patients [5], a number of multigene assays, such as the 21-gene Oncotype DX breast recurrence score (Oncotype DX[®], RS) [6], the 70-gene signature (MammaPrint[®]) [7], Endopredict[®] [8], and Prosigna[®] [9] are used in routine clinical practice to aid in decision-making.

The TAILORx trial was designed to prospectively validate the ability of the RS to estimate chemotherapy benefit in axillary lymph node negative HR+ HER2- patients [10]. Here, patients with a low RS (≤ 10) were assigned to receive endocrine treatment alone and patients with a high RS (≥ 26) were assigned to receive chemoendocrine treatment [10]. Patients with a midrange RS between 11–25 were randomized to receive either chemoendocrine treatment or endocrine treatment alone [10]. Initially published results show 9-year distant recurrence risks of 5, 8, 7 and 15% for the low RS, midrange RS + endocrine therapy, midrange RS + chemoendocrine therapy and high RS groups, respectively, and non-inferiority of outcome in the midrange RS group not receiving chemotherapy on the basis of the RS result was postulated with some exceptions [11].

These exceptions were described in more detail in a secondary analysis recently published, where it was shown that clinical risk as defined in the MINDACT (Microarray in Node-Negative Disease May Avoid Chemotherapy) trial [12] (low clinical risk if primary tumor ≤ 3 cm & low grade or ≤ 2 cm and intermediate grade or ≤ 1 cm and high grade) provided additional prognostic information in all RS groups [13]. Furthermore, patients ≤ 50 years of age seemed to benefit from chemotherapy if their RS was 21–25 or if they were at high clinical risk with an RS of 16–20 [13].

Retrospective analyses of several prospective trials have suggested that the RS is prognostic and predictive of chemotherapy benefit also in node-positive patients [14–16]. According to the 2018 German S3 breast cancer guidelines, as well as the 2019 AGO breast cancer guidelines multigene assays may be used for patients with HR-positive HER2-negative, node-negative disease (in case of Oncotype DX, Prosigna and Endopredict) or in N0–N1 patients irrespective of hormone receptor and HER2 status (in case of MammaPrint), only if no clear decision regarding the use of adjuvant chemotherapy can be made based on conventional prognostic parameters [17, 18]. The current NCCN guidelines strongly recommend considering the 21-gene assay in HR-positive node-negative patients with tumors >0.5 cm in size and to consider multigene assays in HR-positive node-positive patients with <4 involved lymph nodes [19].

Here, we compare the RS result with clinical parameters in patients in Germany with primary invasive breast cancer for whom Oncotype DX testing was performed in routine clinical practice to aid in treatment decision-making.

Methods

Patients and recurrence score

This is a retrospective analysis of patients with HR-positive and HER2-negative early invasive breast cancer (pT1–3, pN0–1, M0) who received an Oncotype DX test in routine clinical practice between October 2015 and June 2017 in Germany. For this purpose, we obtained retrospective, anonymized data from Genomic Health Inc., Redwood City, USA. Grading and Ki-67 were evaluated by local pathologists and submitted to Genomic Health Inc. alongside the patients' lymph node status, tumor size and age. We could not obtain data about the treating entity or therapeutic regimen. After being comprehensively informed by their treating physicians, patients who underwent RS testing had to sign an Informed Consent Document providing detailed explanations about the purposes and use of personal data comprising

scientific research and related publications. The Informed Consent Document is regularly updated and complies with all applicable data protection laws, regulations and rules; in particular the EU-GDPR and the German Federal Data Protection Act. No ethics vote was required for analyzing the anonymized data according to the ethics commission of the University Hospital Tübingen, Germany.

Statistical analysis

We defined the low, midrange, and high RS groups in accordance with the definitions used for the TAILORx trial as an RS of 0–10, 11–25, and 26–100, respectively. Additionally, Ki-67 values of < 20% were defined as low and ≥ 20% as high [20]. For clinical risk, the definition used in the MINDACT trial [12] (low clinical risk if node-negative & primary tumor ≤ 3 cm & low grade or node-positive & ≤ 2 cm & low grade or node-negative & ≤ 2 cm and intermediate grade or node-negative & ≤ 1 cm and high grade) was deployed.

Correlations between categorical variables were assessed using Pearson’s Chi-Squared Test. The significance level was set at $\alpha < 0.05$. All tests were carried out as two-sided. Statistical analysis was performed using R version 3.5.0 and data visualization using the ggplot2 package version 3.1.0.

Results

Patient characteristics

Data from 4695 HR-positive, HER2-negative patients were available for analysis. The mean age was 56.6 years (standard deviation

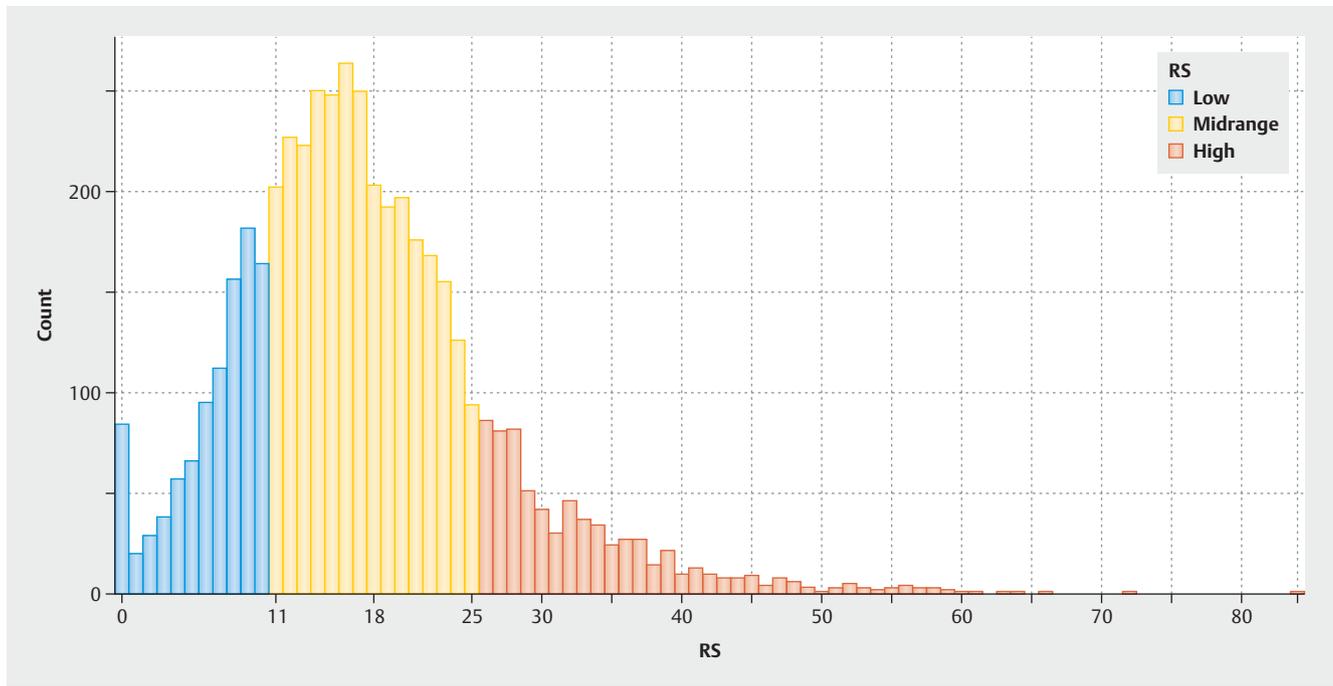
10.4 years) with a median tumor size of 1.8 cm (interquartile range [IQR] 1.3–2.5 cm) and a median Ki-67 of 20% (IQR 10–25%), placing 2177 (49%) into the low Ki-67 and 2310 (51%) into the high Ki-67 group. In all, 3263 (69%) patients had no lymph node involvement, while 1432 (31%) were node positive. Furthermore, 283 (6%), 3762 (82%), and 522 (11%) patients were graded as G1, G2, and G3, respectively and 1792 (40%) were classified as low clinical risk. Compared with node-negative patients, the node-positive patients had a higher proportion of lower grade, Ki-67-low, clinically high-risk, and larger tumors, all $p < 0.001$ (► **Table 1**).

Recurrence score

The RS distribution can be seen in ► **Fig. 1**. The median RS was 16 (IQR 11–22), placing 1003 (21%) patients into the low RS group, 2975 (63%) into the midrange RS group, and 717 (15%) into the high RS group. The distribution of clinicopathological patient characteristics by RS group is illustrated in ► **Table 2**. Patients with high-grade tumors, Ki-67-high tumors, and node-negative patients were more likely to be in the high RS group (all $p < 0.001$) whereas no association was seen for tumor size ($p = 0.265$) and clinical risk ($p = 0.255$) (► **Table 2, Fig. 2**). This was true both for patients older than 50 years and patients 50 years of age or younger. In the subgroups of node-negative and node-positive patients the same associations between clinicopathological features and RS result were observed as in the combined cohort (► **Fig. 3**). In 55% of cases, Ki-67 and RS concordantly classified patients as low – low/midrange or high – high (► **Fig. 4**); 1772 (81%) of 2175 node-negative patients over 50 years of age were assigned a low or midrange RS. The proportion of patients with a low/midrange

► **Table 1** Patient characteristics by axillary lymph node status.

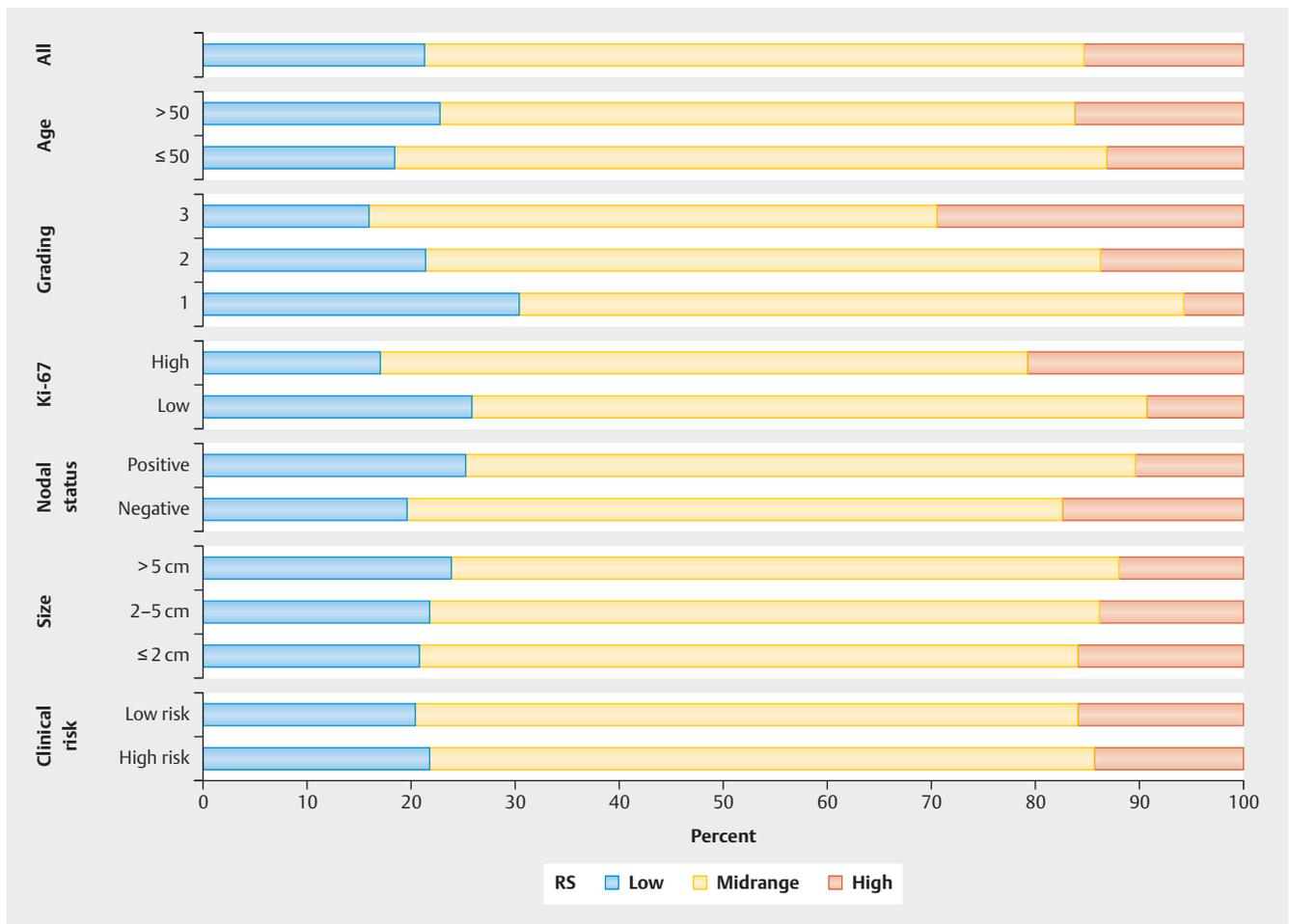
		All patients n	Axillary lymph node		p (χ^2)
			negative n	positive n	
All patients		4695 (100)	3263 (100)	1432 (100)	
Grading	G1 (%)	283 (6)	142 (5)	141 (10)	< 0.001
	G2 (%)	3762 (82)	2558 (81)	1204 (84)	
	G3 (%)	522 (11)	440 (14)	82 (6)	
Ki-67	Low (%)	2177 (49)	1274 (41)	903 (64)	< 0.001
	High (%)	2310 (51)	1812 (59)	498 (36)	
Tumor size	≤ 2 cm (%)	2644 (59)	1912 (62)	726 (51)	< 0.001
	2–5 cm (%)	1703 (38)	1076 (35)	627 (44)	
	> 5 cm (%)	159 (4)	94 (3)	65 (5)	
Age (years)	≤ 50 (%)	1497 (32)	1087 (33)	410 (29)	0.002
	> 50 (%)	3197 (68)	2175 (67)	1022 (71)	
RS group	Low (%)	1003 (21)	641 (20)	362 (25)	< 0.001
	Midrange (%)	2975 (63)	2053 (63)	922 (64)	
	High (%)	717 (15)	569 (17)	148 (10)	
Clinical risk	Low (%)	1792 (40)	1712 (56)	80 (6)	< 0.001
	High (%)	2686 (60)	1352 (44)	1334 (94)	



► Fig. 1 Distribution of RS in clinical routine in Germany.

► Table 2 Patient characteristics by RS group.

		All patients n	RS group			p (χ ²)
			low n (%)	midrange n (%)	high n (%)	
All patients		4695	1003 (21)	2975 (63)	717 (15)	
Nodal status	negative	3263	641 (20)	2053 (63)	569 (17)	< 0.001
	positive	1432	362 (25)	922 (64)	148 (10)	
Grading	G1	283	86 (30)	181 (64)	16 (6)	< 0.001
	G2	3762	805 (21)	2439 (65)	518 (14)	
	G3	522	83 (16)	285 (55)	154 (30)	
Ki-67	low	2177	563 (26)	1413 (65)	201 (9)	< 0.001
	high	2310	394 (17)	1435 (62)	481 (21)	
Tumor size	< 2 cm	2644	549 (21)	1674 (63)	421 (16)	0.265
	2–5 cm	1703	372 (22)	1095 (64)	236 (14)	
	> 5 cm	159	38 (24)	102 (64)	19 (12)	
Age (years)	≤ 50	1497	275 (18)	1025 (68)	197 (13)	< 0.001
	> 50	3197	728 (23)	1949 (61)	520 (16)	
Clinical risk (N0)	Low risk	1712	338 (20)	1093 (64)	281 (16)	0.526
	High risk	1352	256 (19)	854 (63)	242 (18)	
Clinical risk (N+)	Low risk	80	29 (36)	47 (59)	4 (5)	0.034
	High risk	1334	329 (25)	863 (65)	142 (11)	



► Fig. 2 Distribution of RS groups by patient characteristics.

RS was 90% among node-positive patients (1284 of 1432 patients), 79% among patients with Ki-67-high tumors (1829 of 2310 patients), 86% vs. 70% among patients with G2 and G3 tumors (3244 of 3762 patients and 368 of 522 patients), respectively, 88% among patients with a tumor size of > 5 cm (140 of 159 patients) and 82% of node-negative patients at high clinical risk (1110 of 1352).

Discussion

Using the RS in a large cohort of patients with HR-positive/HER2-negative early breast cancer in Germany, we could identify according to the RS a large proportion of patients with clinically high-risk features such as high Ki-67 or high tumor grade, who according to the TAILORx results do not benefit from additional chemotherapy [11, 13].

The RS was initially developed as a prognostic tool, with scores of < 18 defined as low risk, 18–30 as intermediate risk, and > 30 as high risk [6]. Using modified cut-off values the TAILORx trial demonstrated a lack of benefit from chemotherapy in the midrange RS group in a prospective randomized setting [11]. However, subgroup analyses revealed that women ≤ 50 years of age with an RS

of 21–25 regardless of clinical risk and women > 50 with an RS of 16–20 at high clinical risk may in fact derive a benefit from additional chemotherapy [13]. This benefit may likely be attributed to ovarian suppression caused by chemotherapy treatment [13] and should be investigated in future trials. Considering “prospective retrospective” data [21] from the National Surgical Adjuvant Breast and Bowel Project (NSABP) B20 [22] for chemotherapy benefit in patients with a low RS and TAILORx data [11, 13] for midrange RS patients, women > 50 years of age with an RS ≤ 25 and women ≤ 50 years of age with an RS ≤ 15 regardless of clinical risk or ≤ 20 at low clinical risk do not seem to benefit from chemotherapy in addition to endocrine therapy. In this cohort, 2397/3263 (73%) of the node-negative patients were > 50 years of age and had an RS ≤ 25 or were ≤ 50 and had an RS ≤ 15 or were ≤ 50, at low clinical risk and had an RS of ≤ 20.

Retrospective data from prospective trials have been published indicating that node-positive patients with a low recurrence score may also not benefit from chemotherapy [14–16]. In the prospective Plan B study, clinically high-risk patients (including patients with 1–3 involved lymph nodes) with low RS (≤ 11) had an excellent prognosis (94% 5-year DFS), although chemotherapy benefit was not evaluated separately within the node-positive



► Fig. 3 Distribution of RS groups by patient characteristics in node-negative patients (a) and node-positive patients (b).

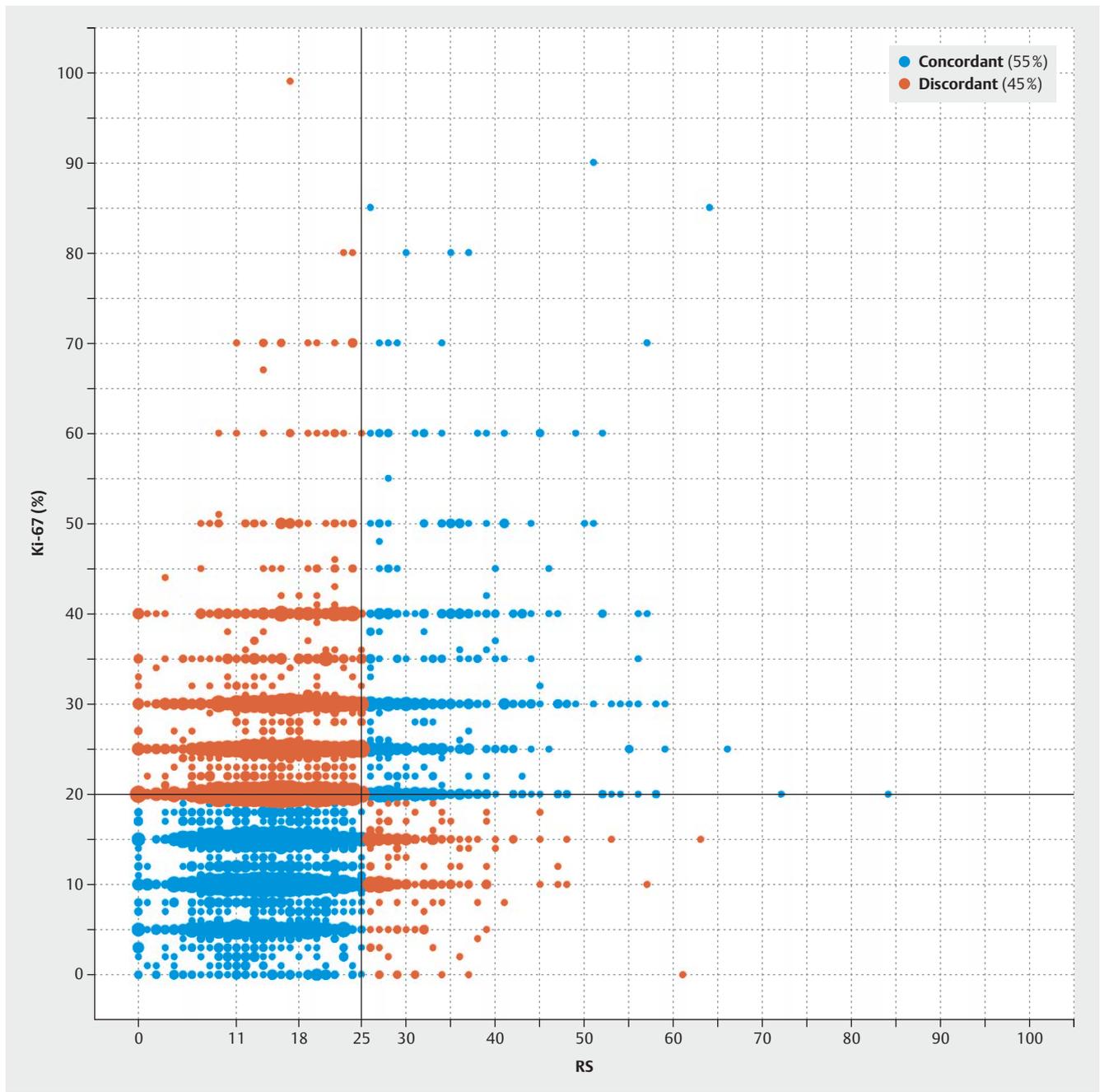
subgroup [23]. Additionally, the MINDACT trial has prospectively shown that multigene assays can be of help in guiding therapy decisions regardless of node status [12]. There, 1404 node-positive patients were included and 912 (65%) were classified as low genomic risk [12]. Of the 709 node-positive patients at high clinical and low genomic risk, 356 received no chemotherapy and no statistically significant difference was seen in distant metastasis free survival when compared with the 353 patients, who received chemotherapy [12]. Results from the RxPONDER study (NCT01272037), which aims to determine the clinical validity of OncotypeDX in node-positive patients, are currently pending. This prospective randomized trial is set to define cut-off values for the possible omission of chemotherapy in node-positive patients [24]. As these are not yet available, we used the TAILORx cutoff values of 0–10, 11–25, and 26–100 for our entire cohort, which includes node-positive patients of whom 90% had an RS \leq 25.

Ki-67 is used clinically to distinguish between luminal A- and B-like subtypes [20]. However, no clear cut-off values have been established; there is also high inter-observer variability and its role in predicting chemotherapy benefit remains unclear [25–27]. Using 20% as cut-off value, 49% of the patients in our cohort would have been classified as luminal A-like; however, 10% of these patients had a high RS, which is in line with earlier results [23]. Although we and others [28] found an association between Ki-67

and RS, the overall concordance rate of patients who would have received chemotherapy or not according to Ki-67 and RS results, respectively, was 55%. What measure to base treatment decisions on, in such cases, where the RS and Ki-67 lead to different conclusions, is a question that has yet to be answered.

In retrospective analyses the RS has been shown to frequently disagree with other molecular tests [29] and in postmenopausal women in the TransATAC trial it was outperformed by other multigene assays as a prognostic tool, even after improving its performance substantially incorporating clinicopathological information in form of the RS-pathology-clinical assessment of distant recurrence risk (RSPC) [30]. However, the only two tests validated to estimate chemotherapy benefit (or rather the lack thereof) in large prospective trials are the 70-gene signature (MammaPrint) [12] and the RS [11, 13].

In the MINDACT trial, patients at low clinical but high genomic risk did not benefit from chemotherapy and patients at high clinical but low genomic risk may have [12]. With the RS, the predictive value in the low and high RS group remains unclear, as these patients were not randomly assigned to treatment groups in the TAILORx trial but treated uniformly [10]. Further research in this area is needed and it therefore remains important even in the age of multigene assays to always take clinical risk into consideration when decisions on treatment are made [31].



► **Fig. 4** Comparison of Ki-67 staining and RS results. Dot size indicates number of observations. Here, 55% were concordantly classified as high Ki-67 and high RS or low Ki-67 and low/midrange RS.

As a limitation of this study, the distribution of tumor characteristics is biased by the decision to use the RS and is therefore not representative for the whole population of HR-positive/HER2-negative patients with early breast cancer. This is most likely why patients in whom axillary lymph nodes were involved more frequently had low-grade tumors and therefore had a higher rate of a low/midrange RS than node-negative patients. Just as observed in the Surveillance, Epidemiology, and End Results (SEER) database, node-positive patients in whom the RS was ordered tended to otherwise have low-risk clinical features when compared to

their node-negative counterparts [32]. Bello et al. recently found that the RS distribution does not differ between node-negative and -positive tumors [33]. Additional limitations include the retrospective design of our study, the lack of treatment information and follow-up data, as well as the fact that no additional clinical risk factors such as menopausal status or progesterone receptor status were provided. We can therefore neither report patient outcome nor treatment efficacy. Furthermore, the records we received were incomplete with missing tumor size in 189 cases, Ki-

67 in 208 cases, grading in 128 cases, and age in 1 case, which is why the clinical risk of 217 patients could not be classified.

Conclusion

In conclusion, the RS was performed in routine practice in both node-negative and node-positive patients. In a large fraction of node-negative patients it indicates that chemotherapy may not be beneficial based on TAILORx results [11, 13]. In node-positive patients, its use was increased when other clinical factors, such as grading, Ki-67, or tumor size, indicated a lower clinical risk. Data from the prospective randomized RxPONDER-trial are awaited to evaluate whether patients with a low RS also might not benefit from chemotherapy and to determine the optimal RS cut-off values for decision-making.

Funding

No billing or payment was made by either party for the performance of this study.

Conflict of Interest

V. P. W. has been reimbursed for travel expenses by Genomic Health Inc. F.-A. T. received consulting fees from Novartis, Tesaro, Genomic Health, Roche, Hexal, Astra Zeneca as well as a research grant from Genomic Health Inc.

S. Y. B. received a research grant from Genomic Health Inc. and speaker fees and honoraria from Pfizer, Roche, Novartis and AstraZeneca.

A. D. H. received a research grant, speaker and consultancy honoraria from Genomic Health Inc. and speaker fees and honoraria from Pfizer, Roche, Novartis, Lilly, MSD and AstraZeneca, Tesaro, Colovis and Eisai. All remaining authors declare that they have no conflict of interest.

References

- [1] Howlader N, Noone A, Krapcho M et al. SEER Cancer Statistics Review, 1975–2016, National Cancer Institute. 2019. Online: https://seer.cancer.gov/csr/1975_2016/; last access: 01.05.2019
- [2] Ferlay J, Colombet M, Soerjomataram I et al. Cancer incidence and mortality patterns in Europe: Estimates for 40 countries and 25 major cancers in 2018. *Eur J Cancer* 2018; 103: 356–387
- [3] Hartkopf A, Müller V, Wöckel A et al. Update Breast Cancer 2019 Part 1 – Implementation of Study Results of Novel Study Designs in Clinical Practice in Patients with Early Breast Cancer. *Geburtsh Frauenheilk* 2019; 79: 256–267
- [4] Untch M, Thomssen C, Bauerfeind I et al. Primary Therapy of Early Breast Cancer: Evidence, Controversies, Consensus: Spectrum of Opinion of German Specialists on the 16th St. Gallen International Breast Cancer Conference (Vienna 2019). *Geburtsh Frauenheilk* 2019; 79: 591–604
- [5] Dowsett M, Goldhirsch A, Hayes DF et al. International Web-based consultation on priorities for translational breast cancer research. *Breast Cancer Res* 2007; 9: R81
- [6] Paik S, Shak S, Tang G et al. A Multigene Assay to Predict Recurrence of Tamoxifen-Treated, Node-Negative Breast Cancer. *N Engl J Med* 2004; 351: 2817–2826
- [7] van de Vijver MJ, He YD, van't Veer LJ et al. A Gene-Expression Signature as a Predictor of Survival in Breast Cancer. *N Engl J Med* 2002; 347: 1999–2009
- [8] Filipits M, Rudas M, Jakesz R et al. A New Molecular Predictor of Distant Recurrence in ER-Positive, HER2-Negative Breast Cancer Adds Independent Information to Conventional Clinical Risk Factors. *Clin Cancer Res* 2011; 17: 6012–6020
- [9] Gnant M, Filipits M, Greil R et al.; on behalf of the Austrian Breast and Colorectal Cancer Study Group. Predicting distant recurrence in receptor-positive breast cancer patients with limited clinicopathological risk: using the PAM50 Risk of Recurrence score in 1478 postmenopausal patients of the ABCSG-8 trial treated with adjuvant endocrine therapy alone. *Ann Oncol* 2014; 25: 339–345
- [10] Sparano JA, Paik S. Development of the 21-Gene Assay and Its Application in Clinical Practice and Clinical Trials. *J Clin Oncol* 2008; 26: 721–728
- [11] Sparano JA, Gray RJ, Makower DF et al. Adjuvant Chemotherapy Guided by a 21-Gene Expression Assay in Breast Cancer. *N Engl J Med* 2018; 379: 111–121
- [12] Cardoso F, van't Veer LJ, Bogaerts J et al. 70-Gene Signature as an Aid to Treatment Decisions in Early-Stage Breast Cancer. *N Engl J Med* 2016; 375: 717–729
- [13] Sparano JA, Gray RJ, Ravdin PM et al. Clinical and Genomic Risk to Guide the Use of Adjuvant Therapy for Breast Cancer. *N Engl J Med* 2019; 380: 2395–2405
- [14] Albain KS, Barlow WE, Shak S et al. Prognostic and predictive value of the 21-gene recurrence score assay in postmenopausal women with node-positive, oestrogen-receptor-positive breast cancer on chemotherapy: a retrospective analysis of a randomised trial. *Lancet Oncol* 2010; 11: 55–65
- [15] Dowsett M, Cuzick J, Wale C et al. Prediction of Risk 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* 2010; 28: 1829–1834
- [16] Penault-Llorca F, Filleron T, Asselain B et al. The 21-gene Recurrence Score[®] assay predicts distant recurrence in lymph node-positive, hormone receptor-positive, breast cancer patients treated with adjuvant sequential epirubicin- and docetaxel-based or epirubicin-based chemotherapy (PACS-01 trial). *BMC Cancer* 2018; 18: 526
- [17] Interdisziplinäre S3-Leitlinie für die Früherkennung, Diagnostik, Therapie und Nachsorge des Mammakarzinoms Langversion 4. 1 – September 2018. 2018. Online: <http://www.leitlinienprogramm-onkologie.de/leitlinien/mammakarzinom>
- [18] Janni W. Diagnostik und Therapie früher und fortgeschrittener Mammakarzinome. Online: https://www.ago-online.de/fileadmin/downloads/leitlinien/mamma/2019-03/DE/Alle_aktuellen_Empfehlungen_2019.pdf; last access: 01.05.2019
- [19] Goetz MP, Gradishar WJ, Anderson BO et al. NCCN Guidelines Insights: Breast Cancer, Version 3.2018. *J Natl Compr Canc Netw* 2019; 17: 118–126
- [20] Goldhirsch A, Winer EP, Coates AS et al. Personalizing the treatment of women with early breast cancer: highlights of the St Gallen International Expert Consensus on the Primary Therapy of Early Breast Cancer 2013. *Ann Oncol* 2013; 24: 2206–2223
- [21] Simon RM, Paik S, Hayes DF. Use of Archived Specimens in Evaluation of Prognostic and Predictive Biomarkers. *J Natl Cancer Inst* 2009; 101: 1446–1452
- [22] Paik S, Tang G, Shak S et al. Gene Expression and Benefit of Chemotherapy in Women With Node-Negative, Estrogen Receptor-Positive Breast Cancer. *J Clin Oncol* 2006; 24: 3726–3734
- [23] Nitz U, Gluz O, Christgen M et al. Reducing chemotherapy use in clinically high-risk, genomically low-risk pN0 and pN1 early breast cancer patients: five-year data from the prospective, randomised phase 3 West German Study Group (WSG) PlanB trial. *Breast Cancer Res Treat* 2017; 165: 573–583

- [24] National Cancer Institute. Tamoxifen Citrate, Letrozole, Anastrozole, or Exemestane With or Without Chemotherapy in Treating Patients With Invasive Breast Cancer – RxPONDER. Online: <https://clinicaltrials.gov/ct2/show/NCT01272037>; last access: 01.05.2019
- [25] Dowsett M, Nielsen TO, A'Hern R et al. Assessment of Ki67 in Breast Cancer: Recommendations from the International Ki67 in Breast Cancer Working Group. *J Natl Cancer Inst* 2011; 103: 1656–1664
- [26] Gluz O, Liedtke C, Huober J et al. Comparison of prognostic and predictive impact of genomic or central grade and immunohistochemical subtypes or IHC4 in HR+/HER2– early breast cancer: WSG-AGO EC-Doc Trial. *Ann Oncol* 2016; 27: 1035–1040
- [27] Viale G, Regan MM, Mastropasqua MG et al.; on the behalf of the International Breast Cancer Study Group. Predictive Value of Tumor Ki-67 Expression in Two Randomized Trials of Adjuvant Chemoendocrine Therapy for Node-Negative Breast Cancer. *J Natl Cancer Inst* 2008; 100: 207–212
- [28] Gluz O, Nitz UA, Christgen M et al. West German Study Group Phase III PlanB Trial: First Prospective Outcome Data for the 21-Gene Recurrence Score Assay and Concordance of Prognostic Markers by Central and Local Pathology Assessment. *J Clin Oncol* 2016; 34: 2341–2349
- [29] Bartlett JMS, Bayani J, Marshall A et al.; on behalf of the OPTIMA TMG. Comparing Breast Cancer Multiparameter Tests in the OPTIMA Prelim Trial: No Test Is More Equal Than the Others. *J Natl Cancer Inst* 2016; 108
- [30] Sestak I, Buus R, Cuzick J et al. Comparison of the Performance of 6 Prognostic Signatures for Estrogen Receptor–Positive Breast Cancer: A Secondary Analysis of a Randomized Clinical Trial. *JAMA Oncol* 2018; 4: 545
- [31] Kolberg H-C, Schneeweiss A, Fehm TN et al. Update Breast Cancer 2019 Part 3 – Current Developments in Early Breast Cancer: Review and Critical Assessment by an International Expert Panel. *Geburtsh Frauenheilk* 2019; 79: 470–482
- [32] Petkov VI, Miller DP, Howlader N et al. Breast-cancer-specific mortality in patients treated based on the 21-gene assay: a SEER population-based study. *NPJ Breast Cancer* 2016; 2: 16017
- [33] Bello DM, Russell C, McCullough D et al. Lymph Node Status in Breast Cancer Does Not Predict Tumor Biology. *Ann Surg Oncol* 2018; 25: 2884–2889