

This is the peer reviewed version of the following article:

Distinct HR expression patterns significantly affect the clinical behavior of metastatic HER2+ breast cancer and degree of benefit from novel anti-HER2 agents in the real world setting / Pizzuti, L; Krasniqi, E; Barchiesi, G; Della Giulia, M; Izzo, F; Sanguineti, G; Marchetti, P; Mazzotta, M; Giusti, R; Botticelli, A; Gamucci, T; Natoli, C; Grassadonia, A; Tinari, N; Iezzi, L; Tomao, S; Tomao, F; Tonini, G; Santini, D; Astone, A; Michelotti, A; De Angelis, C; Mentuccia, L; Vaccaro, A; Magnolfi, E; Gelibter, A; Magri, V; Cortesi, E; D'Onofrio, L; Cassano, A; Rossi, E; Cazzaniga, M; Moscetti, L; Omarini, C; Piacentini, F; Fabbri, Ma; Scinto, Af; Corsi, D; Carbognin, L; Bria, E; La Verde, N; Samaritani, R; Garufi, C; Barni, S; Mirabelli, R; Sarmiento, R; Veltri, E; D'Auria, G; Paris, I; Giotta, F; Lorusso, V; Cardillo, F; Landucci, E; Mauri, M; Ficorella, C; Roselli, M; Adamo, V; Ricciardi, Grr; Russo, A; Berardi, R; Pistelli, M; Fiorio, E; Cannita, K; Sini, V; D'Ostilio, N; Foglietta, J; Greco, F; Zamagni, C; Garrone, O; Di Cocco, B; Baldini, E; Livi, L; Desideri, I; Meattini, I; Sarobba, G; Del Medico, P; De Tursi, M; Generali, D; De Maria, R; Risi, E; Ciliberto, G; Sperduti, I; Villa, A; Barba, M; Di Leo, A; Vici, P.. - In: INTERNATIONAL JOURNAL OF CANCER. - ISSN 0020-7136. - 146:7(2020), pp. 1917-1929. [10.1002/ijc.32585]

[Terms of use](#)

The terms and conditions for the reuse of this version of the manuscript are specified in the publishing policy. For all terms of use and more information see the publisher's website.

09/04/2024 05:32

(Article begins on next page)

09/04/2024 05:32

Distinct HR expression patterns significantly affect the clinical behavior of metastatic HER2+ breast cancer and degree of benefit from novel anti-HER2 agents in the real world setting.

L Pizzuti¹, E Krasniqi^{1*}, G Barchiesi¹, M Della Giulia¹, F Izzo¹, G Sanguineti², P Marchetti^{3,4}, M Mazzotta³, R Giusti³, A Botticelli⁴, T Gamucci⁵, C Natoli⁶, A Grassadonia⁶, N Tinari⁶, L Iezzi⁶, S Tomao⁷, F Tomao⁸, G Tonini⁹, D Santini⁹, A Astone¹⁰, A Michelotti¹¹, C De Angelis¹¹, L Mentuccia¹², A Vaccaro¹², E Magnolfi¹², A Gelibter⁴, V Magri⁴, E Cortesi⁴, L D'Onofrio⁹, A Cassano¹³, E Rossi¹³, M Cazzaniga¹⁴, L Moscetti¹⁵, C Omarini¹⁵, F Piacentini¹⁵, M A Fabbri¹⁶, A F Scinto¹⁷, D Corsi¹⁷, L Carbognin¹⁸, E Bria¹³, N La Verde¹⁹, R Samaritani²⁰, C Garufi²¹, S Barni²², R Mirabelli²³, R Sarmiento²⁴, E Veltri²⁵, G D'Auria⁵, I Paris¹⁸, F Giotto²⁶, V Lorusso²⁶, F Cardillo²⁷, E Landucci¹¹, M Mauri²⁸, C Ficarella²⁹, M Roselli³⁰, V Adamo³¹, G R Ricciardi³¹, A Russo³², R Berardi³³, M Pistelli³³, E Fiorio³⁴, K Cannita³⁵, V Sini³⁶, N D'Ostilio³⁷, J Forrietta³⁸, F Greco³⁹, C Zamagni⁴⁰, O Garrone⁴¹, B Di Cocco²⁵, E Baldini⁴², L Livi⁴³, I Desideri⁴³, I Meattini⁴³, G Sarobba⁴⁴, P Del Medico⁴⁵, M De Tursi⁶, D Generali⁴⁶, R De Maria⁴⁷, E Risi⁴⁸, G Ciliberto⁴⁹, I Sperduti⁵⁰, A Villa⁵¹, M Barba¹, A Di Leo⁴⁸ and P Vici¹.

FOOTNOTES

1. Division of Medical Oncology 2 , IRCCS Regina Elena National Cancer Institute , Rome , Italy.
2. Department of Radiation Oncology , IRCCS Regina Elena National Cancer Institute , Rome , Italy.
3. Medical Oncology Unit, Azienda Ospedaliero Universitaria Sant'Andrea, Rome, Italy
4. Medical Oncology Unit B, Policlinico Umberto I , Rome , Italy.
5. MedicalOncology , Sandro Pertini Hospital , Rome, Italy.
6. Department of Medical, Oral and Biotechnological Sciences, Centro Scienze dell'Invecchiamento e MedicinaTraslazionale -CeSI-MeT , Chieti , Italy.
7. Department of Radiological, Oncological and Anatomo-Pathological Sciences, 'Sapienza' University of Rome, Policlinico Umberto I, Rome, Italy.
8. Department of Gynecology-Obstetrics and Urology, "Sapienza" University of Rome, Rome, Italy.
9. Department of Oncology , University Campus Biomedico of Rome , Rome , Italy.
10. Division of Medical Oncology, Villa San Pietro Hospital , Rome , Italy
11. UO Oncologia Medica I, S. Chiara Hospital, Dipartimento di Oncologia, Dei Trapianti e Delle Nuove Tecnologie, Azienda Ospedaliera Universitaria Pisana, Pisa, Italy.
12. Medical Oncology Unit , ASL Frosinone , Frosinone , Italy.
13. Department of Medical Oncology , Policlinico Universitario "A. Gemelli" , Rome , Italy.
14. Research Unit Phase I trials and Oncology Unit, ASST Monza, Monza, Italy.

This article has been accepted for publication and undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process which may lead to differences between this version and the Version of Record. Please cite this article as doi: 10.1002/ijc.32583

15. Division of Medical Oncology, Department of Oncology and Hematology , University Hospital of Modena , Modena , Italy.
16. Medical Oncology Unit , Belcolle Hospital , Viterbo , Italy.
17. MedicalOncology Unit , Fatebenefratelli Hospital , Rome, Italy.
18. Gynecology Oncology Unit , Catholic University of the Sacred Heart , Rome , Italy.
19. Oncology Unit, ASST Fatebenefratelli Sacco Presidio Ospedaliero Fatebenefratelli, Milano, Italy.
20. Medical Oncology Unit , "Regina Margherita" Hospital , Rome , Italy.
21. Division of Medical Oncology , Pescara Hospital , Pescara , Italy.
22. Department of Oncology, Oncology Unit , ASST Bergamo Ovest , Treviglio , Italy.
23. Department of Hematology & Oncology, Azienda Ospedaliera Pugliese-Ciaccio , Catanzaro , Italy.
24. Oncology Unit , San Filippo Neri Hospital , Rome , Italy.
25. Oncology Unit , S. Maria Goretti Hospital, Latina , Italy.
26. Department of Medical Oncology , "Giovanni Paolo II" Institute, Bari, Italy.
27. Oncology Unit , Formia Hospital, ASL Latina, Italy.
28. Division of Oncology, San Giovanni Hospital, Rome, Italy.
29. Medical Oncology, Department of Biotechnological and Applied Clinical Sciences, University of L'Aquila, L'Aquila, Italy.
30. Department of Systems Medicine, Medical Oncology, University of Rome "Tor Vergata," Rome, Italy.
31. Medical Oncology Unit A.O. Papardo & Department Human Pathology University of Messina
32. Department of Surgical, Oncological and Oral Sciences, Section of Medical Oncology, University of Palermo, Palermo, Italy.
33. Oncology Clinic, Università Politecnica delle Marche, Ospedali Riuniti di Ancona, Ancona, Italy.
34. U.O.C. Oncology, University of Verona, Azienda Ospedaliera Universitaria Integrata, Verona, Italy.
35. Medical Oncology, St. Salvatore Hospital, L'Aquila, Italy.
36. Oncology Unit, ASL Roma 1, Santo Spirito Hospital, Rome, Italy.
37. U.O.C. Lanciano-Vasto, Italy.
38. Medical Oncology, P.O. Narni-Amelia, Terni, Italy.

39. Department of Pathology, Surgery and Oncology, "Mater Salutis" Hospital, ULSS21, Verona, Italy.
40. MedicalOncology Unit, AddariiInstitute of Oncology, S. Orsola-Malpighi Hospital, Bologna, Italy.
41. MedicalOncology, A.O. Ospedale di Insegnamento S. Croce e Carle, Cuneo, Italy.
42. Department of Oncology, Lucca Hospital, Lucca, Italy.
43. Radiation Oncology Unit and Department of Clinical and Experimental Biomedical Sciences "Mario Serio", Azienda Ospedaliera Universitaria Careggi, University of Florence, Florence, Italy
44. Department of Medical Oncology, ASL Nuoro, Nuoro, Italy.
45. Division of Medical Oncology, Reggio Calabria General Hospital, Reggio Calabria, Italy.
46. Breast Cancer Unit & Translational Research Unit, ASST Cremona, Cremona, Italy.
47. Institute of General Pathology , Catholic University of the Sacred Heart , Rome, Italy.
48. Sandro Pitigliani Medical Oncology Department, Hospital of Prato, Prato, Italy
49. Scientific Direction , IRCCS Regina Elena National Cancer Institute , Rome, Italy.
50. Bio-Statistics Unit , IRCCS Regina Elena National Cancer Institute, Rome, Italy.
51. Endocrinology Unit, Fondazione Policlinico Universitario A. Gemelli IRCCS, Roma – Università Cattolica del Sacro Cuore.

Corresponding author:

Isabel Krasniqi. Division of Medical Oncology 2, IRCCS Regina Elena National Cancer Institute, Rome, Italy. Via Elia
Chini, 53 - 00144 Rome (Italy). e-mail: krasniqier@gmail.com; Tel. +390652665036; Fax: +390652665075.

Novelty and Impact: We assessed the impact of HRs expression on tumour behavior and treatment outcomes in 738
HER2+ metastatic breast cancer patients treated with new generation anti-HER2 agents in a real world setting.
Distinct HR expression patterns significantly affected the natural history of disease and the amount of benefit in
terms of progression free and overall survival from pertuzumab-based regimens and T-DM1 treatments, thus giving
some hints on an unmet need for better optimizing their sequential use.

Abbreviations: HER2 = human epidermal growth factor receptor 2; BC = breast cancer; IHC = immunohistochemistry;
PFS = progression free survival; HR = hormone receptor; OS = overall survival; ER = estrogen receptor; PgR =
progesterone receptor; TP = triple positive; FISH = fluorescence in situ hybridization; CISH = chromogenic in situ
hybridization

Conflict of interest:

EK, GB, MDG, FI, GS, PM,MM, RG, AB,AG, NT, LI, ST, CO, FP, MAF, AFS, DC,LC, RS,CG, SB, RM, Ro.Sa, EV,GDA, FT, GT, DS, AA, CDA, LM, AV, EM, Al.Gel., VM, EC, LDO, AC, ER,Fr.Gi., VLR, FC, EL,Ma.Mau, CF, MR, VA, GRRR, AR, RB, MP, EF, KC, VS, NDO, JF, FG, CZ, BDC, EB, LL, ID, IM, Gi.Sar., PDM, MDT, DG, Em.Ri, GC, IS, Al.Vi., MB declare no conflicts of interest.

LP received travel grants from Eisai, Roche, Pfizer, Novartis; speaker fees from Roche, Pfizer, Novartis, Gentili. TG received travel grants from Eisai, Roche, Pfizer, Novartis; speaker fees/advisory boards from Roche, Pfizer, Novartis, Gentili, Lilly. CN received travel grants/personal fees from Pfizer, Eisai, Novartis, Merck Sharp &Dohme, AstraZeneca.An.Mic. received travel grants from Eisai, Celgene, NovartisIpsen; personal fees, advisory boards from Eisai, Novartis, Astra Zeneca, Teva, Pfizer, Celgene. MC received personal fees, advisory boards from Pierre Fabre, Astra Zeneca, Celgene, Eisai, Novartis, Lilly. Lu.Mos. received advisory board from Roche;. Em.Br. is supported by the Italian Association for Cancer Research AIRC-IG 20583; he was supported by the International Association for Lung Cancer (IASLC), the LILT (Legaltaliana per la LottacontroiTumori) and Fondazione Cariverona; he received speakers' and travels' fee from MSD, Astra-Zeneca, Celgene, Pfizer, Helsinn, Eli-Lilly, BMS, Novartis and Roche; consultant's fee from Roche, Pfizer; institutional research grants from Astra-Zeneca, Roche.NLV received personal fees from Eisai and Novartis; research funding from Eisai, travel grants from Pfizer, Roche, Gentili; consulting role from Celldex. IP received personal fees/advisory boards from Roche, Pfizer, Novartis, Italfarmaco, Gentili, Pierre Fabre. OG received personal fees from Celgene, Novartis, Eisai; research funding from Eisai, consulting activities with Celgene, Eisai, Pfizer, Amgen. RDM declares to be a scientific advisory board member at ExosomicsSpA (Siena IT), HiberCell Inc. (New York, US), Kiromic Inc. (Houston TX, US) and at Exiris Inc. (Rome, IT).ADLreceived consulting fees from Novartis and Roche.PV received travel grants from Eisai, Roche, Pfizer, Novartis; speaker fees/advisory boards from Roche, Pfizer, Novartis, Gentili

ABSTRACT

We analyzed data from 738 HER2-positive metastatic breast cancer (mbc) patients treated with pertuzumab-based regimens and/or T-DM1 at 45 Italian centers. Outcomes were explored in relation to tumor subtype assessed by immunohistochemistry (IHC). The median progression free survival at first-line (mPFS1) was 12 months. Pertuzumab as first-line conferred longer mPFS1 compared to other first-line treatments (16 vs 9 months, $p=0.0001$), regardless of IHC subtype. Median PFS in second-line (mPFS2) was 7 months, with no difference by IHC subtype, but it was more favorable with T-DM1 compared to other agents (7 vs 6 months, $p=0.03$). There was no PFS2 gain in patients with tumors expressing both hormonal receptors (HRs) ($p=0.17$), while a trend emerged for tumors with one HR ($p=0.05$). Conversely, PFS2 gain was significant in HRs-negative tumors ($p=0.04$). Median overall survival (mOS) was 74 months, with no significant differences by IHC subtypes. Survival rates at 2 and 3 years in patients treated with T-DM1 in second-line following pertuzumab were significantly lower compared to pertuzumab-naïve patients($p=0.01$). When analyzed by IHC subtype, the outcome was confirmed if both HRs or no HRs were expressed ($p=0.02$ and $p=0.006$, respectively). Our results confirm that HRs expression impacts the clinical behavior and novel treatment-related outcomes of HER2-positive tumors when treatment sequences are considered. Moreover, multivariate analysis showed that HRs expression had no effect on PFS and OS. Further studies are warranted to confirm our

findings and clarify the interplay between HER2 and estrogen receptor (ER) pathways in HER2-positive (mbc) patients.

Key words: advanced breast cancer; HER2 positive; pertuzumab; trastuzumab; T-DM1; real world

Accepted Article

INTRODUCTION

Amplification or over-expression of the human epidermal growth factor receptor 2 (HER2) is observed in approximately 20% of breast cancers (BCs), and is often associated with an aggressive clinical behavior and poor outcomes [1]. When overexpressed, the HER2 oncogene is the dominant driver of BC biology, and treatment strategies targeting HER2 have become the standard of care in all the disease settings since 2006 [2]. Currently, four anti-HER2 agents are licensed in Europe: trastuzumab, lapatinib, pertuzumab and T-DM1. The use of anti-HER2 agents combined or not with chemotherapy has dramatically improved prognosis in all stages of HER2-positive BC [3-7].

About half of HER2-positive BCs also express hormone receptors (HRs)[8]. The coexistence of both estrogen receptors (ER) and progesterone receptors (PgR) expression and overexpression/amplification of HER2 is the hallmark of the subgroup called "triple positive" (TP) breast cancer. Even though treatment with anti-HER2 agents has shown benefit independently on HR status, the cross-talk between the two downstream pathways has an impact on the natural history of the disease and magnitude of treatment-benefit.

In the metastatic setting, recent reports of outcomes by tumor subtype demonstrated that HR-positive/HER2-positive tumors usually exhibit different behaviors and response to trastuzumab-based therapy, as well as distinct timings and patterns of relapse compared to HR-negative/HER2-positive tumors [9,10]. In the CLEOPATRA trial, the addition of pertuzumab to a first-line regimen with trastuzumab and docetaxel did not show any advantage in terms of overall survival (OS) in the subgroup of patients with ER and/or PgR-positive tumors (HR 0.73; 95%CI, 0.50-1.06) [11]. Conversely, in the Emilia and in the TH3RESA trials, patients received a significant PFS and OS benefit from the administration of T-DM1 regardless of HRs expression [7;11]. Yet, no data on novel HER2-blocking agents' outcomes have been specifically reported on tumors overexpressing HER2 and expressing both HRs, i.e., TP tumors. So, we conducted a retrospective analysis on HER2-positive metastatic patients treated with pertuzumab-based regimens and/or T-DM1 according to standard clinical practice, and we specifically focused on whether the expression of HRs defines distinct subtypes with different biological behaviors and patterns of response/resistance to novel HER2-blocking agents in HER2-positive metastatic BCs.

MATERIALS AND METHODS

Our cohort included 738 HER2-positive metastatic BC patients consecutively recruited and treated with pertuzumab-based regimens and/or T-DM1 in any treatment line according to routine practice at 45 Italian oncologic centers from December 2003 through November 2017. The observational study was carried out according to a retrospective approach. Written informed consents were obtained from all patients providing data to our analysis. Information on demographics, clinical, histopathological and immunohistochemical (IHC) features, anti-tumoral therapies and related outcomes were retrieved from the patients' medical records by specifically trained research assistants. All included patients were treated for a metastatic disease (defined as a BC spread over the mammary gland and the pertinent locoregional lymph nodes, including the supraclavicular ones). Each patient was evaluated during treatment according to the follow-up strategies of each center. In all cases clinical evaluation, bone scan, computerized tomography (CT) and/or positron emission tomography (PET) – CT were performed at least every 3-months. Clinical response was evaluated by RECIST criteria. Anonymized data were entered into a dedicated database with a SPSS operating interface. Median follow up was calculated starting from diagnosis of metastatic disease to death or date at the last follow up. Endpoints for efficacy outcome included progression free survival (PFS) and

overall survival (OS). Progression free survival for any specific line of treatment was calculated from the time of treatment start to the time of progression of disease, interruption of treatment for toxicity, death or lost to follow-up. Overall survival was calculated starting from diagnosis of metastatic disease to death or last follow-up. For both PFS and OS the median value, respectively median PFS (mPFS) and median OS (mOS), were calculated using the Kaplan-Meier limit product method. A further assessment of OS in different patient s'subsets was performed by calculating the OS rate at 2 years and 3 years.

Pathology assessment was performed on surgical specimens of primary tumors by dedicated pathologists at the participating centers as per national standards. Estrogen receptor and PgR status were determined at each center by IHC according to local standards. Positivity was considered at a cutoff of ≥ 1 %. A positive HER2 status required an IHC score of 3+ or positive fluorescence in situ hybridization/cromogenic in situ hybridization (FISH/CISH). This study was approved by the institutional ethical committee of the coordinating center (IRCCS Regina Elena National Cancer Institute of Rome) and satellite centers, and was conducted in compliance with Helsinki Declaration.

DATA AVAILABILITY

Raw data were generated at Regina Elena National Cancer Institute - IFO and collected from other 44 Italian cancer centers. Derived data supporting the findings of this study are available from the corresponding author E. K. on request.

STATEMENT ON PREPRINT:

This article has not been posted on a non-commercial preprint server.

STATISTICAL ANALYSIS

Descriptive statistics were used to summarize patient- and disease relevant characteristics. The associations between variables were tested by Chi-square or Fisher's exact test, according to the number and size of the groups compared. Survival estimates were computed by Kaplan-Meier product-limit and compared by log-rank test. Significance was set at $p \leq 0.05$. The associations of interest were evaluated in light of the distinction of the overall study cohort into categories defined by molecular subgroups, with the inherent modalities being set based on the results of IHC analysis and according to the criteria fully reported in the prior paragraph (patients and methods). The impact of the most relevant variables on OS and PFS1 was tested in COX uni/multivariate models. Significance was set at $p \leq 0.05$. The following variables were considered: ICH subgroup, age, PS, metastatic disease at diagnosis, number of metastatic sites, visceral metastasis, Ki-67, pertuzumab-based regimen as first-line of treatment, T-DM1 as a second-line treatment (when testing for OS). The covariates with a significant effect on OS and PFS1 in univariate analysis were further tested in multivariate analysis. The variable "ICH subgroup" was included in the multivariate model independently on the results of univariate analysis. Our choice was due to the relevance of this variable to our study purposes. SPSS software (SPSS version 21.0, SPSS Inc., Chicago, IL) was used for all statistical evaluations.

RESULTS

From 2003 through 2017, 738 HER2-positive metastatic BC patients were retrospectively identified at 45 Italian cancer centers. Recruited patients had received at least one cycle of pertuzumab-based regimen and/or T-DM1. Main patient and tumor characteristics are listed in Table 1. The median age was 54 years, and 63.4% of these patients had an ECOG PS of 0. Three-hundred-nineteen (43.2%) patients had a TP breast cancer. One-hundred-sixty (21.7%) had a tumor expressing only one HR, and the remaining 259 patients (35.1%) had HRs negative tumors. The majority of the patients had visceral metastases (68.7%), bone-exclusive disease was recorded in 51 patients (6.9%), and 239 patients (32.4%) had multiple metastatic sites. As shown in Table 2, after analyzing the distribution of metastatic sites as a function of the IHC characteristics, we observed a significantly higher rate of bone-only disease in TP and ER or PgR positive tumors compared to HRs negative tumors (8.8% and 9.2% versus 3.6%, respectively) ($p=0.02$). Brain metastases were more frequently observed in patients with ER or PgR-positive BCs and in HRs negative tumors than in patients with a TP subtype, even though this difference did not reach statistical significance (20.6% and 27.8%, respectively, versus 20.8%) ($p=0.06$). Conversely, no differences were highlighted in terms of visceral metastases distribution by HR status ($p=0.94$).

Overall, our study cohort included 247 patients (34.3%) with *de novo* metastatic disease, 206 patients (28.6%) with metastasis occurrence within 3 years from diagnosis, and 268 patients (37.2%) with metastatic spread after at least 3 years from initial BC diagnosis (supplementary Table 1). When analyzing our results by IHC subtype, we observed that in the TP subgroup almost half of the patients experienced metastasis after the first 3 years (47%), while only 18.3% of patients experienced relapse within this same time-window, and in 34.1% of patients the disease was metastatic at the onset. Conversely, in patients with HRs negative tumors, we found a higher percentage of early relapses (40.8%) or *de novo* metastatic disease (36.7%), compared to patients who developed metastases after the first 3 years (22.4%). Patients with ER or PgR positive tumors showed an intermediate behavior with respect to the previous two subsets. Their percentage of *de novo* metastatic disease was 28.7%, early relapses (within 3 years) were encountered for 27.3% of patients and the remaining 43.8% had a metastatic recurrence beyond the 3-year period. When we compared the clinical behavior between these three subsets, namely TP, only one HR positive and HRs negative patients, no differences were found regarding the percentage of patients with metastatic disease at the diagnosis ($p=0.27$), while the tendency of TP and one HR positive BC patients to develop less frequently early relapses with respect to HRs negative patients was statistically significant ($p=0.001$), (supplementary Table 1).

Among the 738 patients included in our cohort, 371 (50.3%) received a pertuzumab-trastuzumab-taxane regimen, which was always administered in the first-line setting. Conversely, a total of 517 (70.1%) patients received T-DM1, delivered as any line of treatment. In more detail, the number of patients who received T-DM1 as a first-, second- and third-line was respectively 31 (4.2%), 371 (50.3%) and 96 (13.0%). The remaining 19 (2.6%) patients received T-DM1 beyond the third-line of treatment. All the patients treated with T-DM1 in second-line or beyond had previously received a pertuzumab-trastuzumab-based or trastuzumab-based treatment as first-line. The small cohort of patients (31) who received T-DM1 as first-line regimen had experienced a recurrence while on or within 6 months from trastuzumab-based adjuvant treatment. Overall, a total of 531 (72.0%) patients received a second-line treatment, including or not T-DM1. Among the 371 patients who received T-DM1 as a second-line of treatment, 177 had been previously treated with a pertuzumab-trastuzumab-based regimen as first-line, while 194 had been previously treated with a trastuzumab-based first-line regimen. Among the 531 patients having received second-line treatment, 160 received treatment different from T-DM1 (i.e., lapatinib/capecitabine, trastuzumab/chemotherapy). In these 160 patients first-line treatments were represented by a pertuzumab-based regimen in 109 of them, T-DM1 in 25, and other treatments in 26 patients (Figure 3). Patients with HRs-positive tumors received maintenance endocrine therapy concomitantly with maintenance pertuzumab/trastuzumab, or maintenance trastuzumab after

trastuzumab/chemotherapy regimen, whereas, patients treated with T-DM1, didn't received maintenance endocrine therapy after T-DM1. Moreover, we must take into account that a portion of HRs positive patients after diagnosis of metastatic disease received endocrine therapy plus trastuzumab as first-line, delaying chemotherapy/Her2-block. Overall, median follow-up was 32.9 months (95%CI, 3-256).

First-line treatment

Among the 738 evaluable patients, mPFS at first-line treatment (mPFS1) was 12 months (95%CI, 11-13), with no significant differences among the IHC subtypes, being 12 months (95%CI, 11-13) in TP subtype, 12 months (95%CI, 10-14) in ER or PgR positive, and 12 months (95%CI, 10-14) in HRs negative tumors ($p=0.53$). Among the 371 patients treated with a first-line pertuzumab-trastuzumab-based regimen, the overall mPFS was 16 months (95%CI, 13-19), with a significant improvement with respect to patients who didn't receive pertuzumab-based but had received trastuzumab-based treatments, showing a mPFS of 9 months (95%CI, 8-10) ($p=0.0001$). The advantage in mPFS1 for patients that received pertuzumab is also shown by the PFS1 survival curve in Fig.1 A. The mPFS1 benefit related to pertuzumab was observed in all the IHC subtypes at a statistically significant level, as it can be observed in Figures 1B, 1C and 1 D, respectively for TP, ER or PgR positive, and HRs negative subgroups (see also Table 3). In the multivariate analysis (supplementary Table 2), the only two variables related to a worse PFS1 were: not having received a pertuzumab-based regimen as a first-line treatment (HR 1.7; 95% CI 1.4-2; $p < 0.0001$) and having *de novo* metastatic disease (HR 1.2; 95% CI 1.0 – 1.4; $p = 0.042$). The multivariate analysis showed no significant effect of the IHC subgroup categories on PFS1.

Second-line treatment

Among the 531 patients evaluable for second-line treatment outcome, the median PFS (mPFS2) was 7 months (95%CI, 6-8), with no differences by IHC subtype, being mPFS2 6 months (95%CI, 5-7) in TP, 7 months (95%CI, 5-9) in ER or PgR positive, and 8 months (95%CI, 6-10) in HRs negative tumors ($p=0.44$). Overall, treatment with second-line T-DM1 improved PFS, being mPFS2 7 months (95%CI, 5-9) in 371 patients treated with second-line T-DM1 and 6 months (95%CI, 5-7) in 160 patients who received as second-line other HER-2 based treatments ($p=0.003$) (Table 1 and Table 4). However, T-DM1 advantage in mPFS disappeared when we analyzed the subset of TP patients, which showed a mPFS2 of 6 months (95%CI, 5-7), both with second-line T-DM1 and other second-line treatments ($p=0.17$). In ER or PgR positive BC patients, mPFS2 was 7 months (95%CI, 4-10) if treated with second-line T-DM1, while it was 5 months (95%CI, 3-7) if treated with another regimen ($p=0.05$). In HRs negative patients, second-line T-DM1, compared to other treatments, improved mPFS2, being 10 months (95%CI, 8-12), compared with 7 months of other treatments (95%CI, 6-8), respectively ($p=0.04$) (Table 4). We also addressed the effect of the regimen sequence between first-line and second-line treatments on PFS2 (see Figure 3 for more details), particularly with respect to having received or not a pertuzumab-based regimen in first-line, and having received or not T-DM1 in second-line. We focused especially on the patients who received T-DM1 in second-line, who represent also the major part of subjects who received a second-line treatment (371 patients, 69.9% of patients who received a second-line of treatment). Among them, 177 patients who received pertuzumab-based first-line had a mPFS2 with T-DM1 of 5.6 months (95%CI, 4.5-6.6), while the 194 patients who did not receive pertuzumab, but were treated with a trastuzumab-based first-line, had a mPFS2 with T-DM1 of 8 months (95%CI, 6.6-9.6). This difference was statistically significant in the overall population ($p = 0.02$). The longer mPFS2 in patients who received T-DM1 after a trastuzumab-based first-line compared to those who received it after a pertuzumab-based first-line regimen was maintained in all the three IHC subgroups (TP, ER or PgR +, HRs -), without significant differences among these three

groups ($p=0.44$). Concerning the 160 patients who did not receive T-DM1 in second-line, 109 had received pertuzumab in first-line and showed a mPFS2 of 6 months (95%CI, 4.2-6.8), while the 26 women who were not pretreated with pertuzumab in the first-line had a mPFS2 of 6 months (95%CI, 4.2-7.8), with no difference between the two groups overall and by IHC subtype. However, our study may be underpowered for an adequate comparison between these latter patients' subsets. Among the 31 patients having received T-DM1 as first-line, 25 received a second-line. Unfortunately, data on mPFS2 are unavailable.

Overall survival

Overall, in the 738 patients who contributed data to our analysis, mOS was 74 months (95%CI, 62-87). No differences in mOS emerged when we analyzed our population by IHC subtype, being 73 months (95%CI, 62-87) in TP patients, 76 months (95%CI, 57-98) in ER or PgR positive patients and 76 months (95%CI, 56-95) in HRs negative patients ($p=0.61$). Furthermore, we analyzed data regarding the rates of OS at 2 and 3 years according to the IHC subtype and treatment received for metastatic disease in first and/or second line (Table 5). In the overall population, the administration of pertuzumab-based regimens as first-line followed by T-DM1 in second-line was associated with significantly lower rates of 2 and 3 year survival with respect to the other possible sequences, as it is clearly shown by the relative survival curves in Figure 2A ($p=0.001$). The statistical analysis by IHC subtypes showed that this lower survival rate at both 2 and 3 years was statistically significant in the TP and HRs negative subsets ($p=0.02$ and $p=0.006$, respectively), while no significant differences in OS emerged when we considered the ER or PgR positive patient subgroup ($p=0.57$). We represented these findings also by calculating the associated comparison of survival curves for TP, ER or PgR positive and HRs negative subgroups, respectively, reported in Figures 2B, 2C and 2D. In the multivariate analysis (supplementary Table 2) the only three variables related to a worse OS were: not having received a pertuzumab-based regimen as a first-line treatment (HR 2.0; 95% CI 1.4 - 2.9; $p < 0.0001$), not having received T-DM1 in second-line (HR 1.5; 95% CI 1.1 - 2.0; $p = 0.008$) and a baseline Ki 67 $>20\%$ (HR 1.59; 95% CI 1.1 - 2.4; $p = 0.021$). Moreover, multivariate analysis showed no significant effect of the IHC subgroup categories on OS. Thus, the distinctive effect of the HRs expression patterns on OS observed in the non-parametric test emerges only if treatment sequences are considered separately. This confirms the hypothesis of HRs expression relevance when considering the treatment choice.

DISCUSSION

We conducted an observational study of 738 HER2-positive metastatic BC patients who were treated at 45 Italian Cancer Centers in the time frame between May 2003 and November 2017. All of them had received at least one cycle of pertuzumab-based treatment in first-line and/or at least one T-DM1 cycle in the second or following lines of therapy. Data analysis was performed to assess if the expression of HRs was related to significant differences in terms of key patient-and/or (other than HR) disease-related features and/or treatment outcomes.

Within our study population, we observed some differences in disease features according to HRs expression. In more detail, TP patients had a significant higher prevalence of bone-only disease and a reduced occurrence of brain metastases ($p=0.02$ and 0.06 , respectively). Moreover, in the TP and in patients with tumors with only one HR-positive tumors, there was also a greater proportion of patients who had a late development of metastases, i.e., after the first 3 years from the initial diagnosis, with respect to the HRs negative subgroup ($p<0.0001$).

When we analyzed the overall benefit from the treatment received for metastatic disease, mPFS1 seemed not to be affected by HRs expression ($p=0.53$). However, as shown in Table 3, in patients treated with pertuzumab-trastuzumab-taxane regimen, mPFS1 was significantly higher, being 16 months (95%CI, 13-19), compared with 9

months of other treatments ($p < 0.0001$). Interestingly, this improvement in terms of mPFS1 was recorded across all the three IHC subtypes analyzed, suggesting a benefit of pertuzumab-based treatment independently on HRs expression (Table 3, Figure 1). This was confirmed by the multivariate analysis which also showed that HRs expression pattern did not have a confounding effect on this outcome.

Among the 531 patients evaluable for the second-line treatment outcomes, overall, mPFS2 to T-DM1 treatment was 7 months, in comparison with 6 months for the other second-line treatments ($p = 0.003$). The improvement related to second-line T-DM1 didn't emerge when considering only TP patients, which had a mPFS2 of 6 months for both T-DM1 and for other treatments ($p = 0.17$, Table 4). Conversely, in HRs negative tumors, the benefit of second-line T-DM1 was clearly evident, with 10 months of mPFS2 versus 7 months of other treatments ($p = 0.04$). These findings were based on a calculation performed on all patients who received a second-line, without making any distinction between those who had received a pertuzumab-based regimen in the first-line and those who hadn't. The intriguing result that shows a relatively higher mPFS2 advantage in HER2+ HRs negative BC patients supports the hypothesis according to which HRs negative tumors might have a higher sensitivity to chemotherapy and HER2-blocking agents [12]. Whereas, the absence of a mPFS2 advantage associated with T-DM1 compared to other treatments in TP patients, besides the possible lower sensitivity of this subgroup to HER2-blockers, may be also at least partly attributed to the fact that patients with HRs positive tumors received endocrine maintenance therapy after trastuzumab or trastuzumab/pertuzumab-based therapy. This was not an option for patients who received T-DM1. The additive effect of these two factors may represent a plausible explanation for the lack of an evident advantage of T-DM1 in TP BC patients with respect to other therapeutic options. Moreover, results showed that among the 371 patients who received T-DM1 as second-line, those receiving it after first-line pertuzumab (N:177) had a mPFS2 of 5.6 months, which was significantly lower than the 8 months of mPFS2 for those patients who received T-DM1 in second-line after a first-line regimen not containing pertuzumab (N:194) ($p = 0.02$). This evidence supports a lower efficacy of this agent when delivered immediately after pertuzumab, independently on the IHC subtype. This finding suggests a possible cross-resistance mechanism between the pertuzumab-trastuzumab double-block and T-DM1, which is stronger than a possible similar cross-resistance mechanism existing between trastuzumab and T-DM1.

No differences in OS emerged when we analyzed our entire population by IHC subtype. We also analyzed the clinical outcomes of our study population on the basis of the treatment-sequences received for metastatic disease. Overall, the administration of pertuzumab-based regimens as first-line and subsequently T-DM1 as second-line was significantly associated with a lower rate of 2- and 3-year survival ($p = 0.001$, Table 5). This result is congruent with what we observed when analyzing mPFS2. However, differently than mPFS2 (where this lower benefit was present in all the IHC subtypes), when analyzing OS, the worse outcome of the sequence pertuzumab → T-DM1 in first and second-line did not emerge for all the three IHC subgroups. In fact, these unfavorable OS rates for sequential pertuzumab → T-DM1 treatments were statistically significant both in TP and in HRs negative subsets ($p = 0.02$ and $p = 0.006$, respectively), but not in patients with ER or PgR positive tumors ($p = 0.57$). This finding could be due to some underlying biological mechanism that needs interpretation, although selection bias and confounding factors cannot be excluded, since the effect of the IHC subtype on OS was not confirmed in multivariate analysis.

Among HER2-positive tumors, the expression pattern of HRs apparently defines distinct subtypes. Specifically, the TP subtype could be considered as the subset which most closely resembles the luminal-like tumors, in comparison to HER2 overexpressing tumors that express only one or none of the HRs [9,10]. This hypothesis is supported in our case series by the fact that TP patients had a significant higher prevalence of bone-only disease, a lower prevalence of brain metastases, and showed a greater proportion of "late" metastases, reinforcing the data already published in this regard [13]. Moreover, TP patients showed distinctive characteristics also regarding treatment outcomes, since

the subgroup showed no PFS benefit and lower OS rates, together with HRs negative subset, when receiving T-DM1 in second-line. Therefore, the contemporary expression of HER2 and both HRs may represent a relevant element to be considered when choosing the treatment strategy.

To our knowledge, this is the first study to evaluate the efficacy of new anti-HER2 therapeutic strategies in relation to HRs expression in metastatic BC patients, while making a clear distinction between patients with HER2-positive BC that express both HRs, only one of them, or no HRs at all. A differential sensitivity to combined HER2-blocking agents and chemotherapy according to HR status was consistently reported both in the early and in the advanced setting, giving some clues on the possible role of maintenance endocrine therapy [14-20]. A retrospective study performed in HER2-positive metastatic BC patients suggested that an expression of ER in $\geq 30\%$ of tumor cells was predictive of reduced response to chemotherapy plus trastuzumab, but at the same time it indicated patients that could get a benefit from a maintenance endocrine treatment added to trastuzumab administration after the induction with chemotherapy and trastuzumab [20]. Further data confirm the important role of maintenance endocrine treatment in HER2+ mBC patients also when treated with the pertuzumab-trastuzumab double block.

Since 2012, a paradigm shift was observed in the management of HER2-positive metastatic BC following the results from the CLEOPATRA [21], the EMILIA [7], and the TH3RESA trials [11]. The first study demonstrated a significant increase in both PFS and OS when adding pertuzumab to trastuzumab plus docetaxel. This benefit was less evident in HR-positive BC patients. However, these patients did not receive maintenance hormonal therapy. Recently, our group carried out a retrospective observational study in 264 HER2-positive metastatic BC patients treated with a pertuzumab-based regimen as a first-line. Results were consistent with the findings from the CLEOPATRA trial. Differently from the pivotal trial, in our patient population maintenance endocrine therapy was added to pertuzumab-trastuzumab maintenance in 103 patients, who had the most favorable clinical outcomes in terms of PFS and OS, suggesting that the double-maintenance therapy (HER2 blockade and endocrine treatment) could have a relevant positive clinical impact in patients with HER2-positive/HR-positive BC [22]. In the present analysis, patients treated with pertuzumab/trastuzumab-based regimens experienced a mPFS1 that was comparable to that of the pivotal trial, being 16 months, vs 9 months in those patients who received other first-line treatments ($p < 0.0001$). A clear advantage from pertuzumab/trastuzumab-based regimens was evident in all analyzed IHC subtypes, including TP and one HR positive subgroups, who, in this study cohort, received also endocrine maintenance treatment, which may theoretically “regain” the “PFS loss” in the HRs-positive patients, presumably less responsive to HER2-blocking agents. Besides the proven impact of endocrine treatment received in the metastatic setting, we performed a further analysis to evaluate whether endocrine treatment received in the neoadjuvant or adjuvant setting may have had an impact on mPFS1 and mPFS2. As we show in Supplementary Table 3, among the 491 patients who were initially diagnosed with an early or locally advanced disease, 298 received endocrine treatment as neoadjuvant or adjuvant therapy, while 193 did not. Overall, the patients who had received an endocrine therapy in the early setting had a mPFS1 of 12 months versus 11 months of patients who did not. This difference was not significant ($p = 0.14$). Neither significant differences were found when comparing the mPFS2 of these same two groups ($p = 0.54$). When this same comparison was performed in patients stratified by IHC subgroups, in TP patients, who had received endocrine therapy in the early setting, we verified some impact on the mPFS1 outcome. In fact, mPFS1 for TP patients who had received neoadjuvant/adjuvant endocrine therapy was 11 months compared to 16 months of patients who did not. This difference was at the limit of statistical significance ($p = 0.05$). We may thus hypothesize that the worse outcome in terms of PFS1 for TP patients having received endocrine therapy in the early setting maybe related to the fact that maintenance endocrine treatment for this patients yielded less PFS1 advantage due to the onset of resistance to anti-hormonal agents. At the same time, this result confirms, at least indirectly, the

positive impact of endocrine maintenance therapy in TP patients for PFS1. In this same analysis, no relevant impact on mPFS2 was found for endocrine treatment in the early setting of TP patients ($p=0.26$). Moreover, the TP positive subgroup, as defined based on IHC, was the only one to show some impact of endocrine treatment in early setting on outcomes verified in the metastatic setting. In fact, for both the remaining subgroups, ER or PR positive patients and HRs negative patients, administration of endocrine treatment in the early setting did not impact either mPFS1 (respective p -values were 0.48 and 0.86) or mPFS2 (respective p -values were 0.92 and 0.24). It is noteworthy that the 41 patients in the HRs IHC subgroup, who had received endocrine treatment in the early setting, had initially a HRs positive disease. Overall, this further analysis confirms the distinct nature of TP disease and the particular importance of anti-hormonal treatment in this subgroup.

The EMILIA trial compared T-DM1 to lapatinib and capecitabine for treatment of advanced HER2-positive BC in second-line or beyond [7]. The experimental arm had superior PFS and OS. Subgroup analysis failed to show a differential impact of the treatment based on HRs status. Our group carried out a retrospective, observational analysis of T-DM1 clinical activity in 250 HER2-positive metastatic BC patients [23]. Overall, our results were comparable with those from randomized trials and, similarly to what was found in those studies, no differences in clinical outcomes emerged when we analyzed our population by HRs status ($p=0.29$). However, when analyzed in relation to pertuzumab-pretreatment, patients who received second-line T-DM1 had mPFS2 and mOS of 3 and 12 months respectively ($p=0.0001$) when pertuzumab-pretreated, and 8 and 26 months when pertuzumab-naïve ($p=0.06$). Conversely, in third-line and beyond, mPFS and mOS to T-DM1 were 16 and 18 months in pertuzumab-pretreated ($p=0.05$), and 6 and 17 months in pertuzumab-naïve patients ($p=0.30$). The results of this study regarding the PFS of the second-line of treatment and OS are consistent with those from the present study analyzing a larger population, when pertuzumab pre-treatment is considered. In fact, in the current analysis, patients pre-treated with pertuzumab who received T-DM1 in second-line had a PFS advantage with respect to those not receiving second-line T-DM1 ($p=0.03$). Moreover, in the cohort of the present study, receiving T-DM1 in second-line, immediately after pertuzumab, resulted detrimental in terms of OS with respect to receiving it after a trastuzumab-based first-line. As we mentioned before, this might be related to a possible transient cross-resistance between the two agents, although the outcome could be explained also by a selection bias, considering that patients who received T-DM1 as third or more advanced lines presumably had a more indolent disease which allowed several lines of treatments, compared to patients treated with T-DM1 as second-line. In the current study, a further analysis to explore the OS rates in the different IHC subgroups showed that the aforementioned difference in OS rates according to sequential pertuzumab and T-DM1 treatment was statistically significant in the TP and HRs negative subsets ($p=0.006$). Conversely, it was not recorded in patients with ER or PgR-positive tumors ($p=0.57$). Specifically, TP BC could have more potential to develop chemotherapy resistance related to the activation of the ER pathway [24]. This mechanism could be less evident in HER2-positive tumors that express low levels of HRs or only one of them. At the same time, our results in HRs negative tumors might be related to an intrinsic aggressiveness of the disease in this subset [25].

Overall, our results are consistent with some previously emerged evidences on the reduced activity of T-DM1 when given immediately after pertuzumab-based regimens. A retrospective study investigating the efficacy of T-DM1 after pertuzumab-based combination therapy showed shorter median duration of T-DM1 therapy (4.0 months) in patients who had received prior pertuzumab-based regimens. When discussing this latter evidence, the authors ascribed it at least partially to the retrospective nature of the research and the relatively high percentage of *de novo* stage IV patients (44%)[26]. However, the present study included only 33.5% of *de novo* stage IV patients. Therefore, it is

plausible that the lower efficacy of T-DM1 in patients pretreated with pertuzumab/trastuzumab-based combinations might be the result other mechanisms.

Evidence of lower efficacy of T-DM1 when administered immediately after a pertuzumab-based regimen was also found in prospective studies. The preliminary results of the PERNETTA study conducted in 210 patients with HER2-positive metastatic BC treated with a pertuzumab-containing regimen in first-line and with T-DM1 as second-line, showed that mPFS in second-line treatment was lower (5.3 months) than that recorded in the pivotal trial, where patients treated with second-line T-DM1 were pertuzumab-naïve [27]. Furthermore, a recently published prospective study carried out in 42 Japanese HER2-positive metastatic BC patients has shown that the mPFS following T-DM1 administration was lower in the group pretreated with pertuzumab/trastuzumab-based regimens compared to the group of patients who received trastuzumab-based regimens (2.8 months vs 7.8 months, respectively, $p = 0.0030$) [28].

The present study has some limitations. It has a retrospective design, which *per se* represents a source of confounding and bias. Selection bias may be fueled by the intrinsic characteristics of our patients' disease, which may have affected at some extent treatment outcomes. In more detail, patients who received T-DM1 as third or more advanced lines might have a more indolent disease which allowed them to receive T-DM1 after other treatment-lines. In addition, since we address treatment outcomes by cancer subtype, the lack of centralized evaluation by either IHC or NGS techniques may lessen our confidence in the results observed, although quality controls are in place at the participating centres. In addition, for patients who provided data to this analysis, the follow-up period was relatively short. Our study also has several strengths. This is the first report to address outcomes of treatment sequences in a quite large cohort of HER2-positive metastatic BC patients in light of the biological characteristics of the disease, with a specific focus on HRs expression. In addition, our study was conducted in patients treated outside clinical trials. As such, it faithfully reproduces the current clinical practice. Numerous are the evidences related to the biological differences and response to treatment within the subgroup globally defined as HER2-positive BC. This is the first report that has consistently shown a response to novel anti-HER2 treatments differing by HRs status. Since HER2-positive BC evolves under the selective pressure of targeted agents, it is of paramount importance recognizing resistance pathways related to the exposure to novel treatments according to a given sequence. Our results provides a strong rationale to perform studies aiming at a deeper understanding of the intrinsic biology of HER2-positive disease, in order to optimally combine the available hormonal and anti-HER2 therapy treatments to overcome both endocrine and anti-HER2 resistance. The final goal is the achievement of decisions driving to the most effective therapeutic choice at an individual-patient level.

In conclusion, breast cancer biology is of uttermost relevance for optimizing treatment and better interpreting the clinical outcomes. The central role of the HRs and the HER2 pathways in sustaining BC development and growth has been clearly established. The expression of both HRs concomitantly with the overexpression of HER2 depicts a TP subtype, that may have distinct clinical manifestation and treatment outcomes compared to the other HER2-positive subtypes. Results from this historical cohort support our hypothesis. In more details, in patients with TP tumors, we actually observed clinical manifestations that more closely resemble those of luminal A patients, with higher rates of bone-only disease and lower rate of brain metastasis. No differences emerged in terms of first-line treatment outcomes in TP patients with respect to the other subsets, being the benefit of pertuzumab-based treatment consistent in all subgroups. However, the PFS gain due to the administration of T-DM1 as second-line that was observed in the overall population seems to disappear when analyzing the TP subgroup, giving a hint on a possible additional mechanism of resistance towards HER2-blockers in this subset of patients. On this basis, our data confirm the need for further exploration of TP BC throughout the conduct of ad-hoc designed, appropriately sized,

prospective trials. These latter studies will be aimed to clarify the extent to which treatment outcomes in HER2-positive metastatic BC treated with sequences including hormonal therapy and novel anti-HER2 agents may differ depending on the co-expression of both the HRs vs only one of the two. This research will provide data on “canonical” treatment outcomes, which may be more appropriately interpreted in light of the results from the annexed tasks and experimental designs focused on the underlying mechanisms regulating the development of resistance across the different subtypes of HER2-positive metastatic BC patients. A possible equalizer for outcome differences in TP patients maybe a wider use of concomitant anti-hormonal therapy when possible.

ACKNOWLEDGEMENTS

We thank Ana Maria Edlisca and Dr. Rosa Carbone for editorial assistance and data managing

References

1. Slamon DJ, Clark GM, Wong SG, Levin WJ, Ullrich A, McGuire WL. Human breast cancer: correlation of relapse and survival with amplification of the HER-2/neu oncogene. *Science* 1987; 235: 177-182.
2. O'Sullivan CC, Bradbury I, Campbell C, Spielmann M, Perez EA, Joensuu H, Costantino JP, Delaloge S, Rastogi P, Zardavas D, Ballman KV, Holmes E, de Azambuja E, Piccart-Gebhart M, Zujewski JA, Gelber RD. Efficacy of Adjuvant Trastuzumab for Patients With Human Epidermal Growth Factor Receptor 2-Positive Early Breast Cancer and Tumors ≤ 2 cm: A Meta-Analysis of the Randomized Trastuzumab Trials. *J Clin Oncol*. 2015; 33(24): 2600-2608.
3. Cortés J, Saura C, Bellet M, Muñoz-Couselo E, Ramírez-Merino N, Calvo V, Pérez J, Vidal M. HER2 and hormone receptor-positive breast cancer--blocking the right target. *Nat Rev Clin Oncol*. 2011; 8: 307-311.
4. Untch M, Fasching PA, Konecny GE, Hasmüller S, Lebeau A, Kreienberg R, Camara O, Müller V, du Bois A, Kühn T, Stickeler E, Harbeck N, Höss C, Kahlert S, Beck T, Fett W, Mehta KM, von Minckwitz G, Loibl S. Pathologic complete response after neoadjuvant chemotherapy plus trastuzumab predicts favorable survival in human epidermal growth factor receptor 2-overexpressing breast cancer: results from the TECHNO trial of the AGO and GBG study groups. *J Clin Oncol*. 2011; 29: 3351-3357.
5. Moja L, Tagliabue L, Balduzzi S, Parmelli E, Pistotti V, Guarneri V, D'Amico R. Trastuzumab containing regimens for early breast cancer. *Cochrane Database Syst Rev*. 2012; 4: CD006243.
6. Swain SM, Kim SB, Cortés J, Ro J, Semiglazov V, Campone M, Ciruelos E, Ferrero JM, Schneeweiss A, Knott A, Clark E, Ross G, Benyunes MC, Baselga J. Pertuzumab, trastuzumab, and docetaxel for HER2-positive metastatic breast cancer (CLEOPATRA study): overall survival results from a randomised, double-blind, placebo-controlled, phase 3 study. *Lancet Oncol*. 2013; 14(6):461-71.
7. Verma S, Miles D, Gianni L, Krop IE, Welslau M, Baselga J, Pegram M, Oh DY, Diéras V, Guardino E, Fang L, Lu MW, Olsen S, Blackwell K; EMILIA Study Group. Trastuzumab emtansine for HER2-positive advanced breast cancer. *NEngl J Med*. 2012; 367(19): 1783-91.
8. Lal P, Tan LK, Chen B. Correlation of HER-2 status with estrogen and progesterone receptors and histologic features in 3,655 invasive breast carcinomas. *Am J Clin Pathol*. 2005; 123: 541-546.
9. Yersal O, Barutca S. Biological subtypes of breast cancer: Prognostic and therapeutic implications. *World J Clin Oncol*. 2014; 5(3):412-24.
10. Prat A, Pascual T, Adamo B. Intrinsic molecular subtypes of HER2+ breast cancer. *Oncotarget*. 2017; 8(43):73362-73363.

- Accepted Article
11. Krop IE, Kim SB, Martin AG, LoRusso PM, Ferrero JM, Badovinac-Crnjevic T, Hoersch S, Smitt M, Wildiers H. Trastuzumab emtansine versus treatment of physician's choice in patients with previously treated HER2-positive metastatic breast cancer (TH3RESA): final overall survival results from a randomised open-label phase 3 trial. *Lancet Oncol.* 2017;18(6):743-754.
 12. Llombart-Cussac A, Cortés J, Paré L, Galván P, Bermejo B, Martínez N, Vidal M, Pernas S, López R, Muñoz M, Nuciforo P, Morales S, Oliveira M, de la Peña L, Peláez A, Prat A. HER2-enriched subtype as a predictor of pathological complete response following trastuzumab and lapatinib without chemotherapy in early-stage HER2-positive breast cancer (PAMELA): an open-label, single-group, multicentre, phase 2 trial. *Lancet Oncol.* 2017;18(4):545-554.
 13. Vici P, Pizzuti L, Natoli C, Gamucci T, Di Lauro L, Barba M, Sergi D, Botti C, Michelotti A, Moscetti L, Mariani L, Izzo F, D'Onofrio L, Sperduti I, Conti F, Rossi V, Cassano A, Maugeri-Saccà M, Mottolese M, Marchetti P. Triple positive breast cancer: a distinct subtype? *Cancer Treat Rev.* 2015;41(2):69-76.
 14. Vici P, Pizzuti L, Sperduti I, Frassoldati A, Natoli C, Gamucci T, Silverio T, Michelotti A, Moscetti L, Gori S, Baldini E, Giotto F, Cassano A, Santini D, Giannarelli D, Di Lauro L, Corsi D, Marchetti P, Sini V, Sergi D, Barba M, Maugeri-Saccà M, Russillo M, Mentuccia L, D'Onofrio L, Iezzi L, Scinto AF, Da Ros L, Bertolini I, Basile ML, Rossi I, De Maria R, Montemurro F. "Triple positive" early breast cancer: an observational multicenter retrospective analysis of outcome. *Oncotarget.* 2016;7(14):17932-17944.
 15. Vici P, Pizzuti L, Natoli C, Moscetti L, Mentuccia L, Vaccaro A, Sergi D, Di Lauro L, Trenta P, Seminara P, Santini D, Iezzi L, Tinari N, Bertolini I, Sini V, Mottolese M, Giannarelli D, Giotto F, Maugeri-Saccà M, Barba M, Marchetti P, Michelotti A, Sperduti I, Gamucci T. Outcomes of HER2-positive early breast cancer patients in the pre-trastuzumab and trastuzumab eras: a real-world multicenter observational analysis. The RETROHER study. *Breast Cancer Res Treat.* 2014;147(3):599-607.
 16. Blackwell KL, Burstein HJ, Storniolo AM, Rugo HS, Sledge G, Aktan G, Ellis C, Florance A, Vukelja S, Bischoff J, Baselga J, O'Shaughnessy J. Overall survival benefit with lapatinib in combination with trastuzumab for patients with human epidermal growth factor receptor 2-positive metastatic breast cancer: final results from the EGF104900 study. *J Clin Oncol* 2012;30:2585-92.
 17. Vaz-Luis I, Seah D, Olson EM, Wagle N, Metzger-Filho O, Sohl J, Litsas G, Burstein HJ, Krop IE, Winer EP, Lin NU. Clinicopathological features among patients with advanced human epidermal growth factor-2-positive breast cancer with prolonged clinical benefit to first line trastuzumab-based therapy: a retrospective cohort study. *Clin Breast Cancer* 2013; 13: 254-63.
 18. Tripathy D, Kaufman PA, Brufsky AM, Mayer M, Yood MU, Yoo B, Quah C, Yardley D, Rugo HS. First line treatment patterns and clinical outcomes in patients with HER2-positive and hormone receptor-positive metastatic breast cancer from registHER. *Oncologist* 2013;18:501-10.

- Accepted Article
19. Blackwell KL, Burstein HJ, Storniolo AM, Rugo H, Sledge G, Koehler M, Ellis C, Casey M, Vukelja S, Bischoff J, Baselga J, O'Shaughnessy J. Randomized study of Lapatinib alone or in combination with trastuzumab in women with ErbB2-positive, trastuzumab-refractory metastatic breast cancer. *J Clin Oncol* 2010;28:1124–1130.
 20. Montemurro F, Rossi V, Cossu Rocca M, Martinello R, Verri E, Redana S, Adamoli L, Valabrega G, Sapino A, Aglietta M, Viale G, Goldhirsch A, Nolè F. Hormone receptor expression and activity of trastuzumab with chemotherapy in HER2-positive advanced breast cancer patients. *Cancer* 2012;118:17–26.
 21. Baselga J, Cortés J, Kim SB, Im SA, Hegg R, Im YH, Roman L, Pedrini JL, Pienkowski T, Knott A, Clark E, Benyunes MC, Ross G, Swain SM; CLEOPATRA study group. Pertuzumab plus trastuzumab plus docetaxel for metastatic breast cancer. *N Engl J Med*. 2012;366(2):109-119.
 22. Gamucci T, Pizzuti L, Natoli C, Mentuccia L, Sperduti I, Barba M, Sergi D, Iezzi L, Maugeri-Saccà M, Vaccaro A, Magnolfi E, Gelibter A, Barchiesi G, Magri V, D'Onofrio L, Cassano A, Rossi E, Botticelli A, Moscetti L, Omarini C, Fabbri MA, Scinto AF, Corsi D, Carbognin L, Mazzotta M, Bria E, Foglietta J, Samaritani R, Garufi C, Mariani L, Barni S, Mirabelli R, Sarmiento R, Graziano V, Santini D, Marchetti P, Tonini G, Di Lauro L, Sanguineti G, Paoletti G, Tomao S, De Maria R, Veltri E, Paris I, Giotta F, Latorre A, Giordano A, Ciliberto G, Vici P. A multicenter Retrospective observational study of first-line treatment with PERTuzumab, trastuzumab and taxanes for advanced HER2 positive breast cancer patients. *RePerStudy. Cancer Biol Ther*. 2019;20(2):192-200.
 23. Vici P, Pizzuti L, Michelotti A, Sperduti I, Natoli C, Mentuccia L, Di Lauro L, Sergi D, Marchetti P, Santini D, Magnolfi E, Iezzi L, Moscetti L, Fabbri A, Cassano A, Grassadonia A, Omarini C, Piacentini F, Botticelli A, Bertolini I, Scinto AF, Zampa G, Mauri M, D'Onofrio L, Sini V, Barba M, Maugeri-Saccà M, Rossi E, Landucci E, Tomao S, Alberti AM, Giotta F, Ficorella C, Adamo V, Russo A, Lorusso V, Cannita K, Barni S, Laudadio L, Greco F, Garrone O, Della Giulia M, Marolla P, Sanguineti G, Di Cocco B, Ciliberto G, De Maria R, Gamucci T. A retrospective multicentric observational study of trastuzumab emtansine in HER2 positive metastatic breast cancer: a real-world experience. *Oncotarget*. 2017;8(34):56921-56931.
 24. Risi E, Grilli A, Migliaccio I, Biagioni C, McCartney A, Guarducci C, Bonechi M, Benelli M, Vitale S, Biganzoli L, Biciato S, Di Leo A, Malorni L. A gene expression signature of Retinoblastoma loss-of-function predicts resistance to neoadjuvant chemotherapy in ER-positive/HER2-positive breast cancer patients. *Breast Cancer Res Treat*. 2018;170(2):329-341.
 25. Kennecke H, Yerushalmi R, Woods R, Cheang MC, Voduc D, Speers CH, Nielsen TO, Gelmon K. Metastatic behavior of breast cancer subtypes. *J Clin Oncol*. 2010;28(20):3271-3277.
 26. Dzimitrowicz H, Berger M, Vargo C, Hood A, Abdelghany O, Raghavendra AS, Tripathy D, Valero V, Hatzis C, Pusztai L, Murthy R. T-DM1 activity in metastatic human epidermal growth factor receptor 2-positive

breast cancers that received prior therapy with trastuzumab and pertuzumab. *J Clin Oncol*. 2016;34:3511–3517.

27. Huober J, Weder P, Veyret C, Thürlimann B, Xyrafas A, Vanlemmens L, Guio S, Brain E, Grenier J, Dalenc F, Levy C, Savoye AM, Müller A, Membrez-Antonioli V, Gerard MA, Lemonnier J, Hawle H, Boven E, Bonnefoi H. PERNETTA: A non-comparative randomized open label phase II trial of pertuzumab (P) + trastuzumab (T) with or without chemotherapy both followed by T-DM1 in case of progression, in patients with HER2-positive metastatic breast cancer (MBC). (SAKK 22/10 / UNICANCER UC-0140/1207). *Ann Oncol*, Volume 29, Issue suppl_8.
28. Noda-Narita S, Shimomura A, Kawachi A, Sumiyoshi-Okuma H, Sudo K, Shimoi T, Noguchi E, Yonemori K, Shimizu Ch, Fujiwara Y, Tamura K. Comparison of the efficacy of trastuzumab emtansine between patients with metastatic human epidermal growth factor receptor 2-positive breast cancers previously treated with combination trastuzumab and pertuzumab and with trastuzumab only in Japanese population. *Breast Cancer*. 2019 Jul;26(4):492-498.

Table 1. Main baseline characteristics of the study population (Number: 738)

Characteristics	Patients, Number (%)
Age at diagnosis of metastatic disease	
Median (range)	54 (26-87)
Histology	
Ductal	634 (85.9)
Lobular	32 (4.3)
Other	72 (9.8)
Metastatic at diagnosis	
Yes	247 (33.5)
No	491 (66.5)
ECOG performance status at diagnosis of metastatic disease	
0	468 (63.4)
1	155 (21)
≥2	11 (1.5)
Unknown	104 (14.1)
HER2-positive at initial diagnosis	
Yes	627 (85)
No	111 (15)
Immunohistochemical subtype at diagnosis of metastatic disease	
Triple Positive	319 (43.2)
ER or PgR positive	160 (21.7)
HER2 negative	259 (35.1)
Neo-/adjuvant chemotherapy	
Yes	441 (59.8)
No	297 (40.2)
Neo-/adjuvant trastuzumab	
Yes	261 (35.4)
No	477 (64.6)
Neo-/adjuvant hormonal treatment	
Yes	298 (40.4)
No	193 (26.2)
Perituzumab-based treatment in first line	
Yes	371 (50.3)
No	367 (49.7)
T-DM1 administered as second-line	
Yes	371 (50.3)
No	367 (49.7)
Metastatic sites	
Bone only	51 (6.9)
Visceral	507 (68.7)
Brain	185 (25.1)
Number of metastatic sites	

1	499 (67.6)
2	136 (18.4)
≥3	103 (14.0)

Abbreviations: ER, estrogen receptor; PgR, progesteron receptor

Table 2. Sites of metastases sites according to molecular subtype

Molecular subtype	Bone only	Visceral	Brain
Triple Positive	28 (8.8%)	221 (69.5%)	66 (20.8%)
ER or PgR positive	13 (9.2%)	97 (68.2%)	42 (29.6%)
HRs negative	10 (3.6%)	189 (68.2%)	77 (27.8%)
<i>p</i>	<i>0.02</i>	<i>0.94</i>	<i>0.06</i>

Abbreviations: ER, estrogen receptor; PgR, progesterone receptor

Table 3. Progression Free Survival at first line treatment (PFS1) overall and according to molecular subtype and type of treatment received

Overall	Yes	16 (13-19)	<i><0.0001</i>
	No	9 (8-10)	
Triple Positive	Yes	15 (12-18)	<i><0.0001</i>
	No	8 (6-10)	
ER or PgR Positive	Yes	15 (10-20)	<i>0.0004</i>
	No	9 (7-11)	
HER2 negative	Yes	20 (16-24)	<i><0.0001</i>
	No	9 (8-10)	

Abbreviations: ER, estrogen receptor; PgR, progesterone receptor

Table 4. Progression Free Survival at second line treatment (PFS2) overall and according to molecular subtype and type of treatment received

Overall	Yes	7 (5-9)	<i>0.003</i>
	No	6 (5-7)	
Triple Positive	Yes	6 (5-7)	<i>0.17</i>
	No	6 (5-7)	
ER or PgR Positive	Yes	7 (4-10)	<i>0.05</i>
	No	5 (3-7)	
HER2 negative	Yes	10 (8-12)	<i>0.04</i>
	No	7 (6-8)	

Abbreviations: ER, estrogen receptor; PgR, progesterone receptor

Table 5. Overall Survival at 2 and 3 years according type of treatment received in first and second-line and molecular subtype (Tarone Ware test)

Overall	No	No	88.8	84.7	<i>0.001</i>
	Yes	No	89.1	78.5	
	No	Yes	92.1	82.9	
	Yes	Yes	78	62.7	
Triple Positive	No	No	87.1	83.6	<i>0.02</i>
	Yes	No	89.1	71.1	
	No	Yes	93.4	86.3	
	Yes	Yes	80.2	66.0	
ER or PgR Positive	No	No	88.9	88.9	<i>0.57</i>
	Yes	No	87.3	87.3	
	No	Yes	91.8	83.0	
	Yes	Yes	83.9	73.1	
HER2 negative	No	No	89.9	84.5	<i>0.006</i>
	Yes	No	89.9	84.3	
	No	Yes	90.7	77.9	
	Yes	Yes	72.3	52.5	

Abbreviations: ER, estrogen receptor; PgR, progesterone receptor

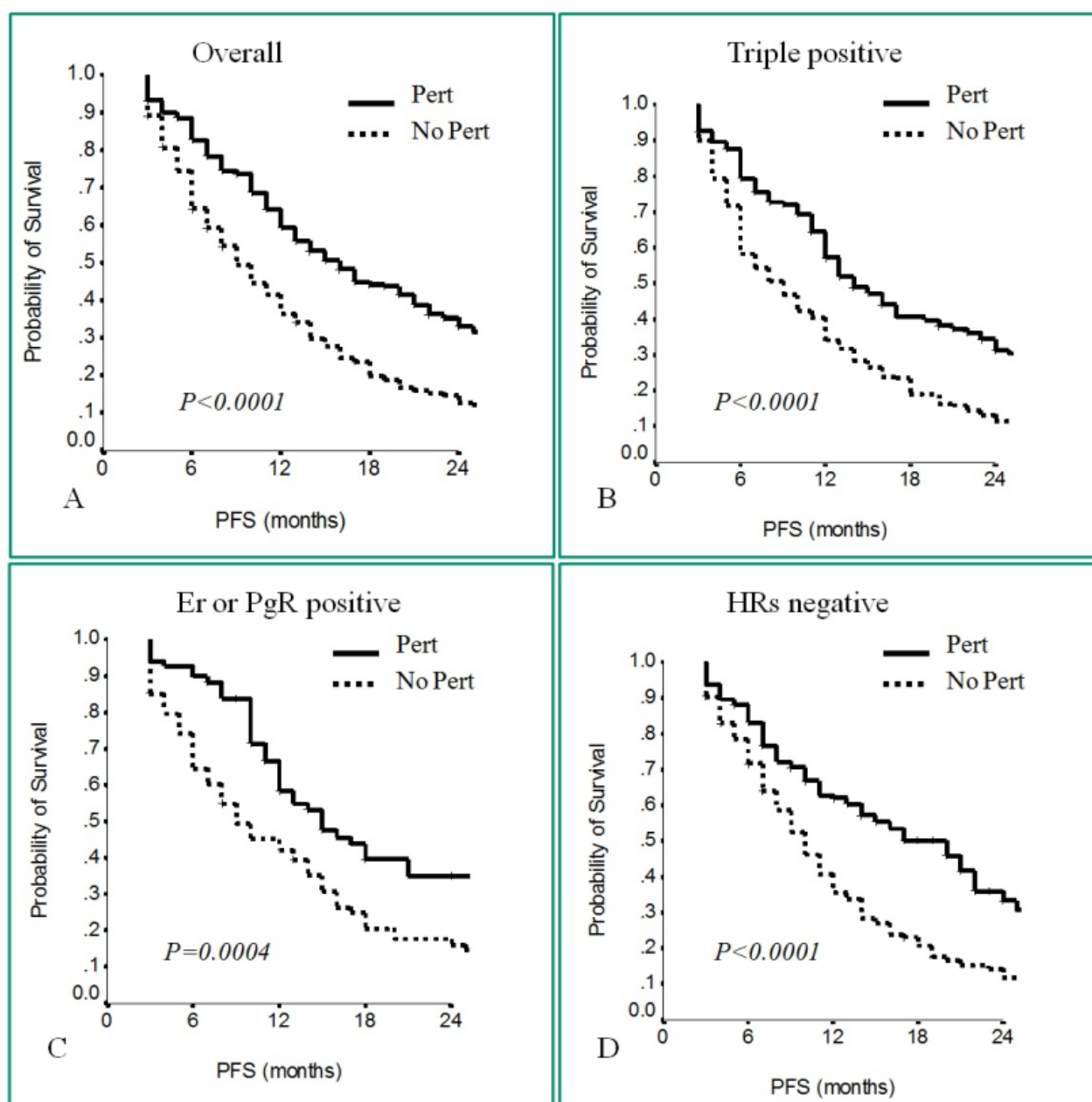


Figure 1. The comparison of progression-free survival in the first-line of treatment (PFS1) between patients that received a pertuzumab-based regimen (Pert) and those who didn't (No Pert) in the overall population (A), in triple positive patients (B), in Er or PgR positive patients (C) and in HRs negative patients (D). Er: Estrogen receptor; PgR: Progesterone receptor; HRs: Hormone Receptors.

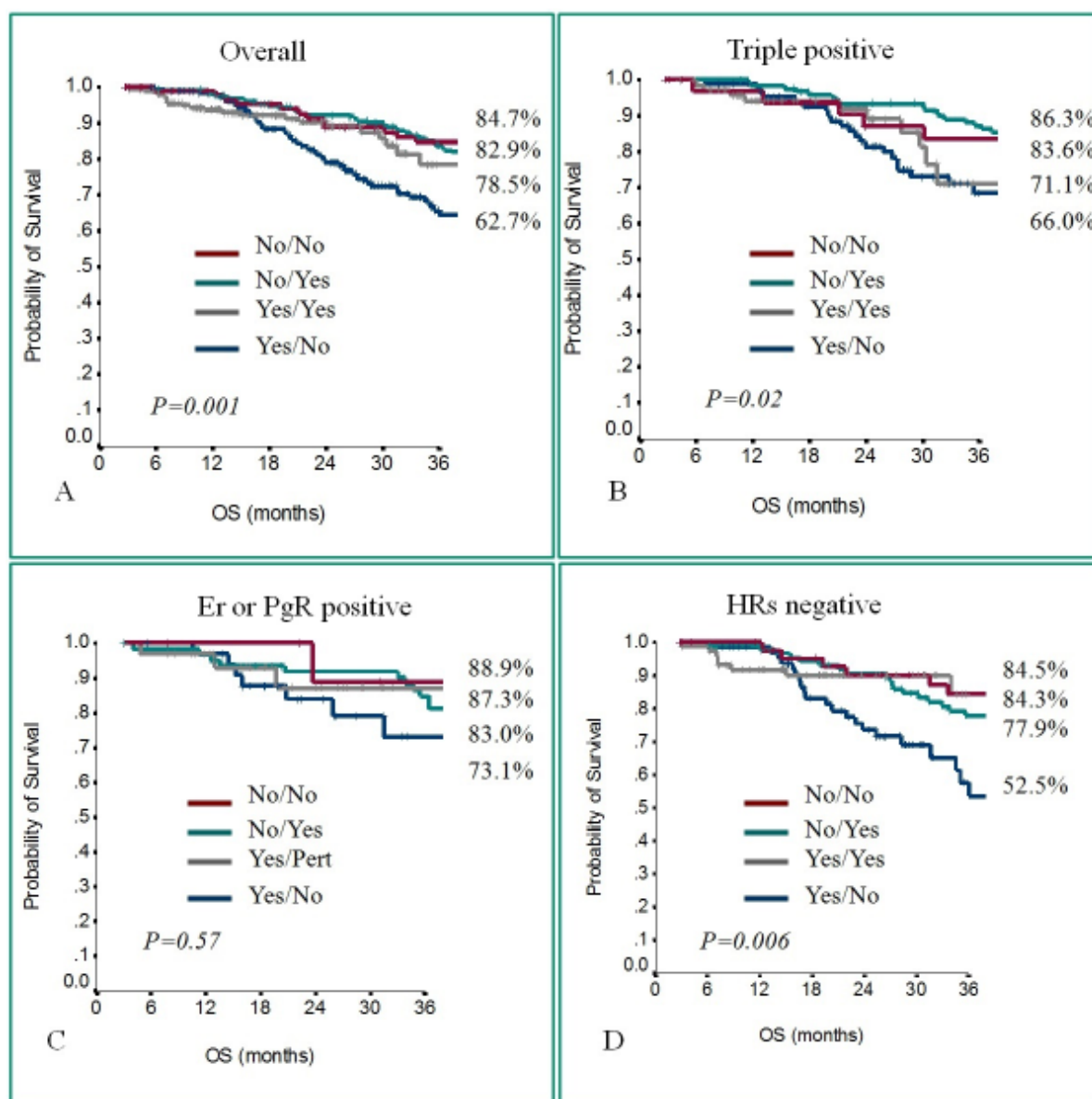


Figure 2. The comparison of overall survival (OS) between patients that didn't receive a pertuzumab-based regimen in first-line and didn't receive T-DM1 as a second-line of treatment (No/No, red colour), patients that didn't receive a pertuzumab-based regimen in first-line and received T-DM1 as a second-line of treatment (No/Yes, green colour), patients that received a pertuzumab-based regimen in first-line and T-DM1 as a second-line of treatment (Yes/Yes, grey colour) and those who received a pertuzumab-based regimen in first-line and didn't receive T-DM1 as a second-line of treatment (Yes/No, blue colour). The comparison was done in the overall population (A), in triple positive patients (B), in Er or PgR positive patients (C) and in HRs negative patients (D). The adjacent percentages to each survival curve refer to the 3-year survival rate.

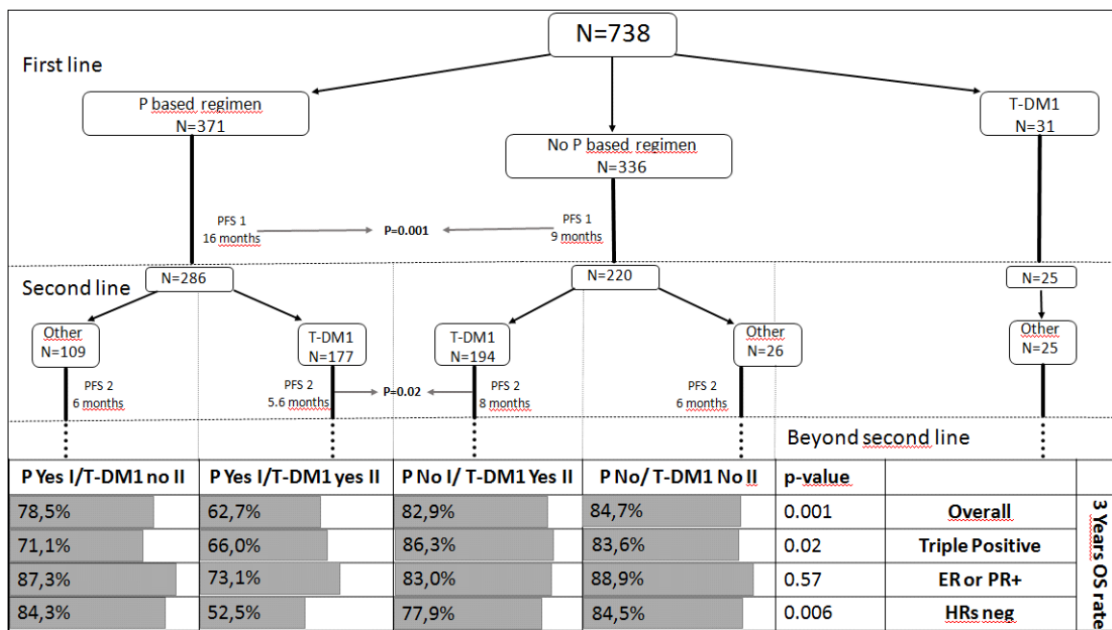


Figure 3: first line and second line of patients divided according to if they received Pertuzumab or not in first line and T-DM1 or not in second line. Correlation between sequences of treatment and 3 years OS rate of patients, divided according to immunohistochemistry subtype.

ER: estroegen receptor; HRS: hormone receptors; OS: Overall survival; P: Pertuzumab; PFS: progression free survival PR: progesteron receptor; P based regimen: Pertuzumab-based regimen; P Yes I: received Pertuzumab in first-line; T-DM1 Yes II: received T-DM1 in secondline; P No I: didn't a Pertuzumab-based regimen in first-line; T-DM1 No II: didn't receive T-DM1 in second-line

About half of breast cancers positive for human epidermal growth factor (HER2) also express hormone receptors but the impact of hormone receptor status on the success of HER2-directed treatments is not fully explored. Here the authors retrospectively assessed tumour behavior and treatment outcomes in 738 women with HER2+ metastatic breast cancer treated with new generation anti-HER2 agents. Distinct hormone receptor expression patterns significantly affected the progression free and overall survival, justifying further studies to define optimal treatment regimens and the interplay between hormone receptor and HER2 signaling.