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Durvalumab with olaparib and paclitaxel for high-risk HER2-negative stage II/III breast cancer: Results from the adaptively randomized I-SPY2 trial.

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AUTHOR CONTRIBUTIONS

All authors have reviewed the data analyses, contributed to data interpretation, contributed to the intellectual content of the manuscript, approved the final version to be published, and agree to be accountable for all aspects of the work. Conceptualization: LE, DB, AD, NH, LV, DY, WFS, JP; Methodology: LE, DB, LV, CY, DW, LP, HH, LD, LS, EP, JW, MC, SB; Formal Analysis: CY, DW, LP; AS, SB, DB; Investigation: LP, CY, DW, HH, LD, AW, ES, JB, AJC, AE, HB, RN, KA, AC, KK, CI, AT, RS, TH, AF, ML, LB, EP, JW, MC, GH, WFS, DY, LV, NH, AD, LE; Data Curation: CY, DW, AW, NO, RS, AS, AA; Original Draft: LP, CY, DW, HH, RN, DY, JM; Visualization: LP, CY, BY, DW, LD, AS; Supervision: LE, DB, AD, NH, LV, DY, WFS, RS, HR, MM, LP, HH; Project Administration: SA, RS, LS, GH, AA, JM; Funding Acquisition: LE, JM.

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L. Pusztai has received consulting fees and honoraria from Pfizer, AstraZeneca, Merck, Novartis, Bristol-Myers Squibb Genentech, Eisai, Pieris, Immunomedics, Seattle Genetics, Clovis, Syndax, H3Bio, Daiichi and Nanostring research support to his institution from AstraZeneca, Pfizer, Merck, Seagen and Bristol Myers Squibb.

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A. Forero-Torres became a Seattle Genetics employee in 2018; and holds stock option from this employment.

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SUPPLEMENTAL INFORMATION

Supplemental information can be found online.

Trial registration: www.clinicaltrials.gov/ct2/show/NCT01042379

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Durvalumab with olaparib and paclitaxel for high-risk HER2-negative stage II/III breast cancer: Results from the adaptively randomized I-SPY2 trial

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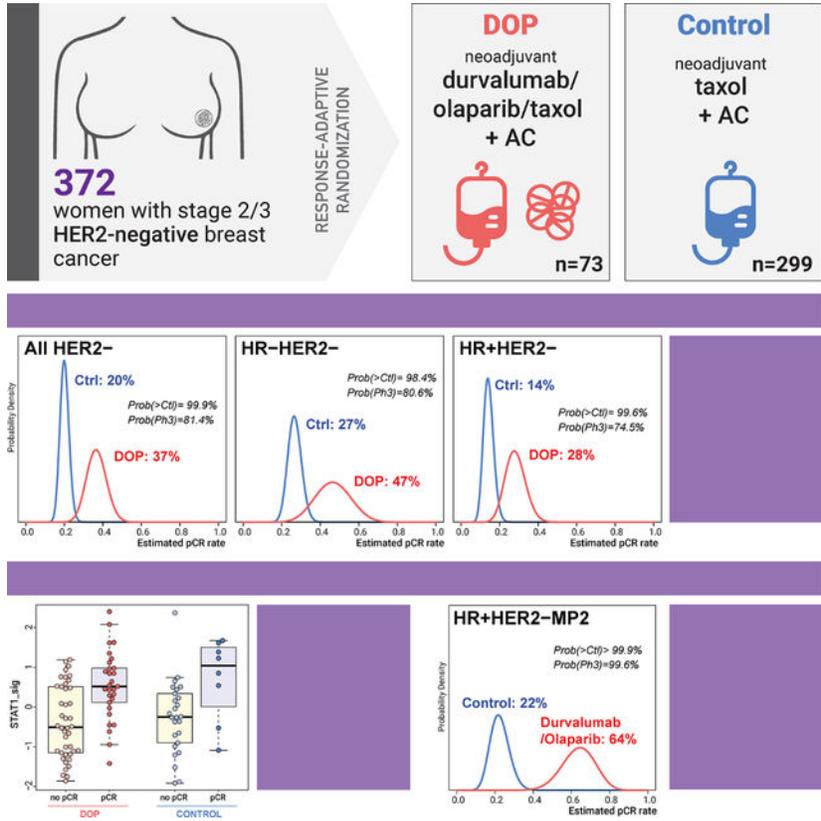
SUMMARY

The combination of PD-L1 inhibitor durvalumab and PARP inhibitor olaparib added to standard paclitaxel neoadjuvant chemotherapy (DOP) was investigated in the phase II I-SPY2 trial of stage II/III HER2-negative breast cancer. Seventy-three participants were randomized to DOP and 299 to standard-of-care (paclitaxel) control. DOP increased pathologic complete response (pCR) rates in all HER2-negative (20 to 37%), hormone receptor (HR)-positive/HER2-negative (14% to 28%), and triple negative (TNBC) (27% to 47%). In HR-positive/HER2-negative cancers, MammaPrint ultra-high (MP2) cases benefited selectively from DOP (pCR 64% vs 22%), no benefit was seen in MP1 cancers (pCR 9% vs 10%). Overall, 12.3% of patients in DOP arm experienced immune-related grade 3 adverse events vs. 1.3% in control. Gene expression signatures associated with immune response were positively associated with pCR in both arms, while a mast cell signature was associated with non-pCR. DOP has superior efficacy over standard neoadjuvant chemotherapy in HER2-negative breast cancer, particularly in a highly sensitive subset of high-risk HR-positive/HER2-negative patients.

eTOC

Pusztai et al. report findings from the I-SPY2 trial showing durvalumab and olaparib administered with paclitaxel improved pathologic complete response (pCR) rate in HER2-negative breast cancers including TNBC and ER-positive cancers. Among the ER-positive/HER2-negative cancers, only the highly proliferative, estrogen receptor low, MammaPrint MP2 subset benefited from the combination therapy.

Graphical Abstract



Keywords

Breast cancer; immunotherapy; DNA repair inhibitor; clinical trial

INTRODUCTION

Several studies indicate that the addition of immune checkpoint inhibitors to standard of care chemotherapy can improve pathologic complete response (pCR) rates in stage II-III triple negative breast cancer (TNBC). In a previous arm of the I-SPY2 trial, 4 cycles of pembrolizumab added to standard paclitaxel-based neoadjuvant chemotherapy improved estimated pCR rate from 22% to 60% in TNBC (Nanda et al., 2020). A subsequent randomized phase III trial confirmed a significantly higher pCR rate (64.8% versus 51.2%) when pembrolizumab was added to chemotherapy; preliminary results suggest improved recurrence free survival (hazard ratio, 0.63) in TNBC (Schmid et al., 2020). Durvalumab, an anti-Programmed Death-Ligand (PD-L1) monoclonal antibody, has received FDA approval for bladder and lung cancers (Stewart et al., 2015). In a single arm, single institution, phase I/II neoadjuvant trial (n=59) of stage I-III TNBC, when durvalumab was administered concurrent with weekly nab-paclitaxel and followed by dose-dense doxorubicin/cyclophosphamide (AC), the final pCR rate was 44%, with no dose-limiting toxicities observed (Foldi et al., 2021). A randomised phase II study in TNBC (n=174), GeparNuevo, demonstrated a numerical, but not statistically significant, increase in pCR rate (53.4% vs. 44.2%, $P = 0.287$) when durvalumab was added to weekly

nab-paclitaxel followed by epirubicin/cyclophosphamide neoadjuvant chemotherapy (Loibl et al., 2019). The safety and efficacy of the combination of durvalumab and olaparib [a polyadenosine 5' diphosphoribose polymerase (PARP) inhibitor] were established in the MEDIOLA (NCT02734004) phase I/II multi-cohort basket trial. Preliminary results of the germline *BRCA* mutant metastatic breast cancer cohort of the trial indicated good tolerability and an objective tumor response rate of 59% (Domchek et al., 2020).

Preclinical studies suggest the potential for synergistic improvement using a combination of immune checkpoint inhibitors and PARP inhibitors (Esteva et al., 2019). Impaired nucleotide and base excision repair caused by PARP inhibition can increase mutation and neoantigen loads that are associated with increased sensitivity to immune checkpoint therapy (Lassen, 2019). DNA fragments generated by impaired DNA repair can also activate the intracellular STING (Stimulator of Interferon Genes) pathway in cancer cells, leading to signaling events that activate immune cells in the tumor microenvironment (Pantelidou et al., 2019).

Here, we report the results of an arm of the neoadjuvant I-SPY2 adaptive platform trial for stage II/III early breast cancer evaluating a combination of durvalumab and olaparib. Our goal was to estimate if the combination of durvalumab and olaparib concurrent with weekly paclitaxel neoadjuvant therapy (DOP) could increase the pCR rate compared to chemotherapy alone in HER2-negative, stage II-III breast cancers. A priori defined and exploratory molecular markers were also assessed to identify predictive markers. The primary endpoint was pathologic complete response (pCR), and a priori defined and exploratory molecular markers were assessed for the purpose of identifying biomarkers predictive of response. I-SPY is a multi-arm, adaptively randomized trial that aims to rapidly identify new drug combinations that induce higher pCR rates than standard of care chemotherapy in various breast cancer biomarker subsets (Barker et al., 2009). The pCR rates are monitored in real-time, along with early imaging response, as they accumulate in each arm and Bayesian adoptive randomization ensures that patients are preferentially accrued to arms with higher pCR rates. A minimum proportion (20%) of patients are continuously allocated to the control arm (chemotherapy alone). This design allows rapid identification of promising new therapies in *a priori* defined biomarker subtypes with a smaller sample size than traditional randomized phase II trials. An arm is considered successful and “graduate” when based on the accumulating responses the statistical engine predicts an 85% probability that the experimental arm would have a statistically significantly higher pCR rate than the control arm in a 300-patient traditional 1:1 randomized trial. An arm could graduate in one or more biomarkers subsets.

RESULTS

Efficacy

Between May 3, 2018 and June 2, 2019, 73 HER2-negative patients (21 TNBC, 52 HR-positive/HER2-negative) were enrolled in the (DOP arm. Three hundred and thirty-six participants were randomized to the control arm (paclitaxel) but 37 did not receive the allocated treatment, resulting in 299 evaluable HER2-negative patients in the control arm (CONSORT diagram, Figure S1). Baseline characteristics were similar between the experimental and control arms (Table 1). Seventeen patients in the control arm and 1 in the

experimental arm did not proceed to surgery (due to patient's choice, leaving the treating institution, or progression) and were considered non-pCR for primary efficacy analysis.

On June 2, 2019, the prespecified Bayesian modelling of surgical pathology results and serial MRI assessments from the first 23 of the 73 patients predicted that the probability of success in a hypothetical phase III trial were 0.944, 0.950 and 0.838 in all HER2-negative, HR-positive/HER2-negative, and TNBC subtypes, respectively. Per protocol, further accrual was therefore stopped and DOP graduated in the HER2-negative, and HR-positive/HER2-negative signatures. After all patients subsequently completed surgery, the final efficacy analysis indicated that durvalumab plus olaparib improved estimated pCR rates (over control) from 20% to 37% in HER2-negative cancers, from 14% to 28% in HR-positive/HER2-negative cancers, and from 27% to 47% in TNBC. Final probability distributions for achieving pCR are shown in Figure 1A including the 95% probability intervals. Final predicted probabilities of phase III success were 0.814, 0.745 and 0.806 in all HER2-negative, HR-positive/HER2-negative, and TNBC signatures, respectively, after all participants had completed surgery. The overall probability that the experimental arm was superior to the control was >98% for all three signatures.

DOP also shifted residual cancer burden categories towards lower values across the entire range of residual cancer burden (RCB) scores (Symmans et al., 2017). The proportion of patients in RCB II and III categories (i.e. moderate and extensive residual disease) were lower in the DOP arm indicating smaller residual cancers across the entire residual disease spectrum in all subsets (Figure 1B).

Median follow-up duration for participants in the experimental arm at the time of submission was 18 months, thus distant metastasis-free and overall survival results are not yet mature.

Safety and Toxicity

All patients who received at least one dose of study-assigned therapy were evaluable for safety and toxicity. In the control arm, 34% (n=102) and in the experimental arm 56% (n=41) of patients experienced at least one grade 3 adverse event. Grade 3 and 4 adverse events are summarized in Table 2 (also see Data S1, "Frequency of all adverse events observed"). Immune-related adverse events were more common in the experimental arm; 9 patients (12%) in the DOP arm experienced immune-related grade 3 or greater adverse events compared to 4 patients (1.3%) in the control arm. Overall, 20 patients (27.4%) experienced an immune related adverse event (grades 1 through 4) in the experimental arm compared to 6 patients (2%) in the control arm. In the experimental arm, immune related adverse events included 5 patients (7%) with hypothyroidism (grade 1 and 2), 4 (5%) pneumonitis (grade 1/2), 4 (5%) adrenal insufficiency (1 patient with grade 1 and 3 with grade 3), and 1 of each non-insulin dependent diabetes (grade 1), thyroiditis (grade 3), pancreatitis (grade 3). Dose reductions and early discontinuation of therapy are also shown on Table 2. The median time from treatment consent to surgery was similar in the experimental (168 days, range: 107–273) and control arms (165 days, range: 85–289).

Biomarker analysis

Seventy-one participants in the DOParm and 34 assigned to control had expression data from pretreatment FFPE biopsies were available for biomarker analysis. Thirteen a priori defined mRNA biomarkers (Table S1), including 10 immune signatures, and one proliferation, DNA repair and estrogen receptor (ER) signature each were examined as predictive markers. In the whole biomarker population, 7 and 5 of the 10 immune mRNA markers studied were significantly associated with higher pCR rate in the experimental and control arms, respectively (Figure 2). The DNA repair deficiency (PARPi7) and mitotic signatures (described in Table S1) were also positively associated with pCR but only in the experimental arm. The T-cell signature and CD68 mRNA expression showed no significant association with pCR in either treatment arm, and the dendritic cell signature was only significant in the experimental arm. The mast cell signature and ESR1/PGR expression were significantly associated with lack of pCR in the experimental arm and showed a similar but non-significant trend for the same direction in the control arm. The marker treatment interaction tests were not significant for nearly all of these markers (9/10), indicating that they cannot identify the patient population that selectively benefited from inclusion of durvalumab and olaparib. We also note that the expression of 8 of the 10 immune markers were highly positively correlated with one another, only the mast cell and macrophage/T-cell ratio signatures showed weak or negative correlation with the other immune markers. ESR1 expression also correlated negatively with all other markers except the mast cell signature (Figure S2). Pairwise comparisons of the 13 mRNA biomarkers between cases of pCR and no-PCR by treatment arm in the whole study population, and in the TNBC and HR+/HER2- subsets, respectively are shown in Supplementary Figures S3–S5. Associations between the 13 mRNA biomarkers and pathologic response in the whole biomarker study population, and in the TNBC and HR+/HER- subsets are shown in Table S2.

In the TNBC subset (N=21 DOP, N=19 control), the STAT1, macrophage/T cell, and mitotic signatures were positively associated with pCR in the experimental arm, and no marker was associated with response in the control arm (Figure 2). However, the marker-treatment interaction tests were not significant. The only immune marker that was associated with lack of pCR in both treatment arms was high expression of the mast cell signature.

In the HR-positive/HER2-negative subset (N=50 durvalumab/olaparib, N=15 control), high ESR1, PGR expression and mast cell signature were associated with lack of pCR, and all other markers except CD68 (that was not associated with response) were positively associated with pCR in the experimental arm (Figure 2). There were too few pCR (N=2) in the control arm in this biomarker cohort to perform analysis. We also examined predicted pCR rates in the two Mammaprint subsets, MP1 (N=109 control, N=24 experimental arm) and MP2 (N=49 control, N=28 experimental). The MP2 subgroup, corresponding to 53% of the Mammaprint high, HR-positive/HER2-negative subset in this trial arm, derived all the benefit from adding durvalumab and olaparib to chemotherapy, in this subset the predicted pCR probabilities were 64% (95%CI: 47%-80%) and 22% (95%CI: 13%-32%) in the experimental and control arms, respectively (Figure 3A–B). Patients in MP1 group had low pCR rates in both arms and showed no improvement with DOP (9% versus 10%), the RCB score distributions in the MP1 and MP2 subsets are shown on Figure

S4. Because in HR-positive cancers, estrogen receptor signaling and proliferation determine endocrine and chemotherapy sensitivities, respectively, we compared expression levels of the proliferation and SET_{ER/PR} gene signatures between the MP1 and MP2 HR-positive/HER2-negative cancers. We also assessed if the MP1 and MP2 subtypes differ in immune gene expression using the tumor inflammation signature as a general marker of immune presence. We selected this signature because it showed the highest correlation with other immune signatures and has also been extensively validated as marker of immunotherapy response in clinical trial tissues (Ayers et al., 2017). The MP2 subset had significantly higher proliferation and significantly lower SET expression, but there was no significant difference in the tumor inflammation signature (Figure 3C–E). We also found that PARPi7 gene signature was significantly higher in MP2 than in MP1 group (LR p=0.007). However, the PARPi7 signature correlated strongly with proliferation. The mRNA expression levels of BRCA1 and BRCA2 genes were not significantly associated with MP status.

DISCUSSION

Durvalumab and olaparib improved pCR rates when added to standard of care neoadjuvant chemotherapy from 20% to 37% in HER2-negative cancers. In TNBC, pCR rate improved from 27 to 47%; and in HR-positive/HER2-negative disease from 14 to 28%. Importantly, as indicated by RCB distributions, even among patients who did not achieve pCR, the immune checkpoint inhibitor/PARP inhibitor combination reduced the amount of residual cancer across the entire spectrum of RCB scores in all HER2-negative subtypes.

We recognize that the pCR rates in I-SPY, particularly in the TNBC subset, are lower, 27%, than reported by other trials using taxane/anthracycline neoadjuvant chemotherapy (35–40%). We believe that this is largely attributable to patients switching to non-protocol therapy, prompted by no-response/stable disease on imaging at 3 and 12 weeks, 11% of patients in the control arm discontinued early due to “progression/lack of sufficient response”; these patients are considered non-pCR per protocol, yet some ends up with pCR at the time of surgery. From larger randomized trials (CALGB 40603, GeparSixto, GeparQuinto), that did not routinely assess response with imaging during therapy, we know that clinical progression on taxane/anthracycline regimens is < 5%. Early imaging is integral to I-SPY, but it also turned out to be a liability because we lose patients who are concerned about lack of imaging response early on during treatment.

Two other trials evaluated the addition of durvalumab to neoadjuvant chemotherapy in TNBC. A single arm phase I/II trial at Yale Cancer Center assessed durvalumab (10 mg/kg every two weeks) administered concurrently with weekly paclitaxel (80 mg/m²) and dose dense AC and reported 44% pCR rate (Foldi et al., 2021). The randomized, GeparNuevo, phase II trial compared durvalumab (1500 mg every 4 weeks) concurrent with weekly nab-paclitaxel and epirubicin/cyclophosphamide to the same chemotherapy alone and demonstrated a numerical, but not statistically significant increase in pCR rate from 44% to 53% (p=0.287) (Loibl et al., 2019). The 95% confidence intervals of the pCR estimates in the experimental arms of these 3 trials 47% (95%CI: 29–64%) in I-SPY2, 44% (95%CI: 30–57%) in Yale phase I/II, and 53% (95%CI: 42–61%) in GeparNuevo, respectively, overlap substantially despite the differences in durvalumab administration,

chemotherapy back bones, and having a third drug, olaparib, in the I-SPY2 trial. Synergy between sequential taxane/platinum anthracycline neoadjuvant chemotherapy and PARP inhibitors were examined in two large neoadjuvant trials (BrighTNess, GeparOla), and both failed to demonstrate improved pCR rate with the addition of a PARP inhibitor in either germline *BRCA* mutant or wild type breast cancers (Pusztai et al., 2020). Clinical trial data also remain inconclusive about synergy between PARP inhibitors and immune checkpoint inhibitors (Peyraud and Italiano, 2020). Both the MEDIOLA (durvalumab + olaparib) and TOPACIO (niraparib + pembrolizumab) basket trials reported efficacy mostly in germline *BRCA* mutant cancers, where PARP inhibitors alone are expected to be effective. Overall, these results, suggest relatively modest, if any, contribution from olaparib to the increased pCR rate in the experimental arm of I-SPY2.

The DOParm of I-SPY2 is the 4th randomized trial to demonstrate improvement in pCR rate with the inclusion of an immune checkpoint inhibitor in neoadjuvant chemotherapy for TNBC. A previously reported arm of I-SPY2, randomized patients to 4 cycles of pembrolizumab with weekly paclitaxel followed by AC (without pembrolizumab) and reported an increase in pCR from 22% to 60% (Nanda et al., 2020). This finding was confirmed in a large randomized trial, KN-522, that administered pembrolizumab concurrent with paclitaxel plus carboplatin followed by anthracycline/cyclophosphamide also with pembrolizumab and demonstrated a statistically significant increase in pCR from 51% to 65% ($p=0.0005$) (Schmid et al., 2020). The recently presented Impassion-031 study also showed that atezolizumab in combination with albumin-bound paclitaxel followed by AC improved pCR rates (41% vs 58%, $p=0.004$) compared to the same chemotherapy alone in TNBC (F.Hoffman-LaRoche, 2020) There are less data on the efficacy of neoadjuvant immune checkpoint therapy in HR-positive cancers. The only published result from a randomized trial is from the HR-positive/HER-negative cohort in the pembrolizumab arm of I-SPY2 that demonstrated improvement in pCR rate from 13% to 30% (Nanda et al., 2020) which is very similar to our current result showing that DOP increased pCR rate from 14% to 28%.

In this, and all previously reported neoadjuvant trials, no unexpected safety signals, or perioperative complications were seen. However, there was an increase in known, immune-related adverse events in the experimental arms. The most common was thyroid function abnormalities. In the current study 27.3% of patients in the experimental arm experienced some immune-related adverse events, and 12.3% had a grade 3 or greater event, a rate very similar reported by the KN-522 trial using pembrolizumab.

Despite extensive efforts, it has been difficult to identify predictive biomarkers that identify stage II/III TNBC that selectively benefit from inclusion of immune checkpoint therapy with neoadjuvant chemotherapy. High tumor infiltrating lymphocyte count, high expression of PD-L1 protein and a broad range of immune-related genes predict for higher pCR rate with both chemotherapy alone and with chemotherapy plus immune checkpoint therapy (F.Hoffman-LaRoche, 2020; Gianni et al., 2020; Loibl et al., 2019; Schmid et al., 2020). Unlike in metastatic TNBC, in all neoadjuvant randomized trials, PD-L1 protein or mRNA expression did not define the population that selectively benefit from immune checkpoint therapy. We also observed that most immune signatures were positively associated with

pCR in both treatment arms. At the individual marker level, some associations reached significance in the experimental arm but not the control arm; however, given the small and unequal cohort sizes and multiple tests, it is unclear if these are truly differentially predictive of response. In formal statistical testing, none of the immune markers showed significant marker-treatment interaction in TNBC.

HR-positive/HER2-negative patients with Mammprint classified as ‘ultra high’ (MP2) showed selective benefit from durvalumab/olaparib. In MP2 patients, the pCR rate was 64% in the experimental arm compared to 22% in control; no benefit from the combination was seen in the MP1 group (pCR 9% vs 10%). Previously, we reported that MP2 status was significantly associated with pCR in HR-positive/HER2-negative patients randomized to the pembrolizumab arm of I-SPY (Yau et al., 2018). We also reported that the HR-positive/HER2-negative MP2 group benefited from adding carboplatin and veliparib to paclitaxel in an earlier arm of I-SPY2, where pCR rates were 33% versus 18%, whereas no improvement in pCR was seen in the MP1 group (Rugo et al., 2016). These findings suggest that the MP2 tumors are more chemotherapy sensitive than MP1 and this sensitivity is further enhanced by adding an immune checkpoint inhibitor. The molecular characteristics of the MP2 group, showing high proliferation and low ER and ER-related gene expression are consistent with greater chemotherapy sensitivity and lower endocrine therapy sensitivity of these cancers. These features also suggest that this subset of HR-positive cancers might derive substantial survival benefit from more effective chemotherapy that induces higher pCR. Benefit from adjuvant endocrine therapy, that could improve outcome in cases with residual cancer, is likely small in this subset due to low level of ER expression.

In conclusion, durvalumab and olaparib concurrent with weekly paclitaxel significantly improved pCR rate in all HER2-negative cancers including TNBC and HR-positive cancers. Among the HR-positive/HER2-negative cancers, only the highly proliferative, estrogen receptor low, Mammprint MP2 subset benefitted from the combination therapy.

STAR * METHODS

RESOURCE AVAILABILITY

Lead Contact: Further information and requests for resources or data should be directed to the I-SPY Data Access and Publications Committee coordinator (ispy2dapc@quantumleaphealth.org).

Materials availability: Requests for sharing of materials should be directed to the Lead Contact.

Data and code availability: Transcriptomic and clinical data used in this study is available at <https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE173839>. Additional de-identified subject level data may be requested by qualified investigators. Details of the trial, data, contact information, proposal forms, and review and approval process are available at the following website: <https://www.ISPYtrials.org/collaborate/proposal-submissions>.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

I-SPY 2 TRIAL Overview—I-SPY2 is an ongoing, open-label, adaptive, randomized phase II, multicenter trial of neoadjuvant therapy for early-stage breast cancer (NCT01042379). It is a platform trial evaluating multiple investigational arms in parallel against a common standard of care control arm. The primary endpoint is pCR (ypT0/is, ypN0), defined as the absence of invasive cancer in the breast and regional nodes at the time of surgery. As I-SPY2 is modified intent-to-treat, patients receiving any dose of study therapy are considered evaluable; those who switch to non-protocol therapy, progress, forgo surgery, or withdraw are deemed ‘non-pCR’. Secondary endpoints include residual cancer burden (RCB) and event-free and distant relapse-free survival (EFS and DRFS)(Symmans et al., 2017).

Trial Design—Assessments at screening establish eligibility and classify participants into subtypes defined by hormone receptor (HR) status, HER2, and 70-gene signature (Mammaprint®) status (Cardoso et al., 2016). Adaptive randomization in I-SPY2 preferentially assigns patients to trial arms according to continuously updated Bayesian probabilities of pCR rates within each biomarker signature; 20% of patients are randomly assigned to the control arm.(Berry, 2011) While accrual is ongoing, a statistical engine assesses the accumulating pathologic and MRI responses at weeks 3 and 12 and continuously re-estimates the probabilities of an experimental arm being superior to the control in each defined biomarker signature. An arm can be dropped for futility if the predicted probability of success in a future 300-patient, 1:1 randomized, phase 3 trial drops below 10%, or graduate for efficacy if the probability of success reaches 85% or greater in any biomarker signature. The clinical control arm for the efficacy analysis uses patients randomized throughout the entire trial. Experimental arms have variable sample sizes: highly effective therapies graduate with fewer patients in the experimental arm; arms that are equal to, or marginally better than, the control arm accrue slower and are stopped if they have not graduated, or terminated for lack of efficacy, before reaching a sample size of 75. During the design of each new experimental arm the investigators together with the pharmaceutical sponsor decide in which of the 10 a priori defined biomarker signatures the drug will be tested. Upon entry to the trial, participants are dichotomized into hormone receptor (HR) negative versus positive, HER2 positive versus negative, and MammaPrint High1 [MP1] versus High2 [MP2] status. From these 8 biomarker combinations (2×2×2) I-SPY has created 10 biomarker signatures that represent the disease subsets of interest (e.g. all patients, all HR+, all HER2+, HR+/HER2-, etc, for complete list see reference (Berry, 2011)) in which a drug can be tested for efficacy.

The DOP arm was studied in three subtypes including (i) all HER2-negative, (ii) TNBC, and (iii) HR positive/HER2-negative cancers. HER2 positive patients were not eligible for randomization into this arm. Efficacy is monitored in each of these biomarker signatures separately and an arm could graduate in any or all biomarker signature of interest. When graduation occurs, accrual to the arm stops, final efficacy results are updated when all pathology results are complete. The final estimated pCR results therefore may differ from the predicted pCR rate at the time of graduation. Additional details on the study design have been published elsewhere (Park et al., 2016; Rugo et al., 2016).

Eligibility—Participants eligible for I-SPY2 are women >18 years of age with stage II or III breast cancer with a minimum tumor size of >2.5 cm by clinical exam, or >2.0 cm by imaging, and Eastern Cooperative Oncology Group performance status of 0 or 1 (Oken et al., 1982). HR-positive/HER2-negative cancers assessed as low risk by the 70-gene MammaPrint test are ineligible as they receive little benefit from systemic chemotherapy. Only HER2-negative patients were eligible for randomization to the olaparib/durvalumab arm. Additional exclusion criteria for this arm included prior PARP inhibitor or immune checkpoint inhibitor therapy, use of immunosuppressive medications, or history of autoimmune disease.

Treatment—All HER2-negative participants in I-SPY2 receive standard of care neoadjuvant chemotherapy, which also served as the control arm, consisting of intravenous paclitaxel 80 mg/m² weekly for 12 weeks, followed by four cycles of 60 mg/m² doxorubicin and 600 mg/m² cyclophosphamide (AC) every 2–3 weeks. Participants in the experimental arm also received 3 cycles of intravenous durvalumab 1500 mg every 4 weeks and 100mg oral olaparib twice a day from weeks 1 through 11, concurrent with paclitaxel. For the first paclitaxel infusion, 20 mg dexamethasone was given and if no infusion reaction occurred, dexamethasone was reduced to 10 mg for week two, if no infusion reaction was observed with the first two treatments, dexamethasone was discontinued. Dose reductions and toxicity management were specified in the protocol. Adverse events were collected according to the NCI Common Terminology Criteria for Adverse Events (CTCAE) version 4.0.

After completion of AC, patients underwent lumpectomy or mastectomy and nodal sampling, with choice of surgery at the discretion of the treating surgeon. All patients were screened for potential adrenal insufficiency before surgery with a morning serum cortisol level.

Trial Oversight—I-SPY2 is conducted in accordance with the guidelines for Good Clinical Practice and the Declaration of Helsinki, with approval for the study protocol and associated amendments obtained from independent ethics committees at each site. Written, informed consent was obtained from each participant prior to screening and again prior to treatment. The I-SPY2 Data Safety Monitoring Board meets monthly to review patient safety.

METHOD DETAILS

Pretreatment Biopsy Processing and gene expression profiling—Core needle biopsies of 16-gauge were taken from the primary breast tumor at screening, before treatment. I-SPY2 changed biopsy requirements from fresh frozen (FF) tissue to formalin fixed paraffin embedded (FFPE) tissues in May 2018; therefore, all DOP patients had RNA extracted from FFPE biopsies. The collected tissue samples are immediately placed in 10% Neutral Buffered Formalin for between 6 and 72 hours before dehydration and processing to FFPE blocks. A 5µM section is stained with hematoxylin and eosin (H&E) and pathologic evaluation performed to confirm the tissue contains at least 30% tumor. A tissue sample meeting the 30% or greater tumor cellularity requirement is further sectioned centrally at the I-SPY 2 laboratory to produce between ten to thirty 5 µM sections collected on charged slides microarray profiling. The sections are processed at Agendia, Inc., for RNA extraction

and gene expression profiling on Agilent 44K microarrays. For each array, the green channel mean signal is log₂-transformed and centered within array to its 75th quantile as per the manufacturer's data processing recommendations. A fixed value of 9.5 is added to avoid negative values.

Biomarker analysis cohort definition—For the first 7 years of the I-SPY 2 TRIAL, only fresh frozen tumor (FF) tissues were collected and assayed for gene expression using Agilent microarrays. In 2018, Agendia switched to analyzing only the FFPE samples. The DOP arm opened in 2018 and therefore these patients have FFPE-derived biomarker data. The clinical control arm for the DOP efficacy analysis used patients randomized throughout the entire trial. Therefore, samples that match to the clinical control arm include expression data derived from FF (n=265) and FFPE (n=34) tissues. During a pilot study of 74 patients where both FF and FFPE samples were assayed for mRNA expression to assess interchangeability of the tissue sources we noted that for several of the immune gene signatures that we evaluated, the correlation coefficient ranged from 0.35 to 0.77. Because of the low concordance, we decided to only include the 34 FFPE-derived expression data from the control arm in the gene signature analysis to provide comparable results to the 71 samples from the DOP arm. The MammaPrint scores are required for I-SPY 2 trial eligibility that is performed with an FDA approved assay, and therefore, the MP 1 and MP1 class are available for all patients.

Continuous Gene Expression Biomarkers Assessed—Nine gene signatures were examined as predictors of response to durvalumab/olaparib, including a proliferation signature (mitotic score)(Bianchini et al., 2013), a DNA repair deficiency signature (PARPi7)(Daemen et al., 2012; Wolf et al., 2017), and 7 immune signatures corresponding to various immune cell types (Danaher et al., 2017) (T cell, B cell, dendritic cell, mast cell), STAT1 cytokine signaling (Rody et al., 2009), macrophage/T cell ratio (Campbell et al., 2013) and an immunotherapy response signature (Ayers et al., 2017) (tumor inflammatory signature). These gene signatures were taken from the literature and each has been associated with response to chemotherapy, targeted therapy, or prognosis. mRNA expression of single genes ESR1 (ER), PGR (PR), CD274 (PDL1), CD279 (PD1) and CD68 (macrophage marker) were also assessed.

Continuous gene expression biomarkers—Table S2 lists the 13 gene expression biomarkers evaluated as predictors of response to durvalumab/olaparib, along with their type (pre-specified or exploratory), pathway designation, member genes, implementation method, and PubMed ID of references.

***FFPE-revised PARPi-7 signature**—The PARPi-7 DNA repair deficiency signature as published was developed and tested on gene expression data from fresh frozen (FF) samples (Daemen et al., 2012; Wolf et al., 2017). As PARPi-7 contains genes that are not highly correlated across FF and FFPE, we replaced poorly translating genes by others in the same DNA repair pathway that (1) correlate to the original genes in FF, (2) associate with response to veliparib/carboplatin, and (3) are correlated in expression data generated from FF and FFPE tissues. We evaluated this revised PARPi-7 by 1) extracting revised predictor

genes BRCA1 (A_23_P207400), CHEK2 (A_23_P109452), MAPKAPK2 (A_23_P201483), XRCC4 (A_23_P122174), RAD17 (A_24_P97836), POLB (A_32_P34552), and CIRBP (AG_0673); 2) combining probesets for normalization genes RPL24, ABI2, GGA1, E2F4, IPO8, CXXC1, and RPS10 into gene level summaries by averaging; 3) dividing each PARPi-7 predictor gene level by the geometric mean of the normalization genes; 4) log₂-transforming each ratio and median centering across the population; and 5) calculating signature scores using the published weights and boundaries: Weights<-c(-0.5320, 0.5806, 0.0713, -0.1396, -0.1976, -0.3937, -0.2335), Boundaries<-c(-0.0153, 0.006, 0.0031, -0.0044, 0.0014, -0.0165, -0.0126), Score = Weights*(Genes -Boundaries). The score is then standardized to sd=1; and used for our qualifying biomarker evaluation.

MP1 and MP2 categorization—In I-SPY2, patients were classified as MammaPrint High risk (MP1) or MammaPrint ultra-high risk (MP2) by Agendia, Inc., using a pre-defined threshold applied to the MP 70-gene risk score evaluated on Agilent 44K arrays. The threshold used is equivalent to the median cut point of I-SPY 1 participants (-0.154 in the original I-SPY 1 dataset) who fit the eligibility criteria for I-SPY 2.

SET Index—The ‘sensitivity to endocrine therapy’ (SET_{ER/PR}) gene signature includes the ESR1 gene and 17 other estrogen (ER) and progesterone (PR) receptor regulated transcripts. (Sinn et al., 2019) The SET_{ER/PR} signature is predictive of progression-free and overall survival in both metastatic and early-stage HR-positive breast cancers treated with endocrine therapy. (Sinn et al., 2019; Symmans et al., 2010) We evaluated the distribution of the SET index in the MP1 and MP2 subtypes of the HR-positive/HER2-negative cohort. All HR-positive patients included in I-SPY2 are MammaPrint high, but these patients can be further subdivided into high risk (MP1) and ultra-high risk (MP2) groups as described above and as published previously.

QUANTIFICATION AND STATISTICAL ANALYSIS

I-SPY 2 Efficacy Analysis—The I-SPY 2 efficacy analysis model have two main components: (1) a core covariate-adjusted Bayesian logistic regression and (2) an adjustment for time trends as described below.

The Core Bayesian Logistic Regression Model—Probability distributions of pCR are calculated using a Bayesian covariate-adjusted logistic model with HR, HER2 and MammaPrint statuses as covariates for each eligible signature. Let $y_i \in \{0,1\}$ be the indicator for the pCR response of patient i ($i = 1, \dots, N$). Covariates $x_{1i}, x_{2i}, x_{3i} \in \{0,1\}$ represent the HR, HER2 and MP statuses of patient i (with 1 indicating positive and 0 negative for HR and HER2 and ‘High1’ and ‘High2’ for MP). Label A_i the treatment arm assigned to patient i

The model is

$$y_i \sim \text{Bernoulli}(p_i)$$

$$\text{logit}(p_i) = \beta_0 + \beta_1 x_{1i} + \beta_2 x_{2i} + \beta_3 x_{3i} + \theta_{A_i} + \gamma_{1, A_i} x_{1i} + \gamma_{2, A_i} x_{2i} + \gamma_{3, A_i} x_{3i}$$

The model’s components are as follows:

- The $\beta_1x_{1i} + \beta_2x_{2i} + \beta_3x_{3i}$ terms capture the effect of being in a particular subtype defined by (HR, HER2, MP) status.
- The $\gamma_{1,A_i}x_{1i}, \gamma_{2,A_i}x_{2i}, \gamma_{3,A_i}x_{3i}$ terms are the treatment effects within each of the (HR, HER2, MP) subtype.
- We set $\theta_0 = \gamma_{1,0} = \gamma_{2,0} = \gamma_{3,0} = 0$ to ensure parameter identifiability.
- The θ_{A_i} represent the effect of being on a particular treatment arm for all patients.

For each of the coefficients Θ in the regression model, we assume independent normal prior distributions: $\Theta \sim N(\mu_\Theta, \sigma_\Theta^2) = N(0,1)$.

We also include the data from the control arm of the I-SPY 1 trial as historical prior information. However, the likelihood function evaluation corresponding to the I-SPY1 data is raised to the power of 0.2 to reflect a discounting or weak borrowing. In effect the historical results count the approximately 200 patients from I-SPY 1 as the equivalent of 40 I-SPY 2 patients.

Adjusting for Time Trends—In addition to the core logistic regression structure of the above terms, we include time trend parameters to capture the effect of possible drifting pCR rates over time.

Motivation: The initial statistical analyses in I-SPY 2 compared investigational arms with concurrently randomized controls. (Park et al., 2016; Rugo et al., 2016) However, in September 2013, the FDA granted accelerated approval for pertuzumab + trastuzumab + docetaxel as neoadjuvant therapy for high risk HER2+ breast cancer. Our investigators and DSMB required dropping the I-SPY 2 control arm for HER2+ subtypes (trastuzumab + docetaxel) because it did not contain pertuzumab, which we did by amendment in early 2014. Going forward, we wanted to be able to use the results for the original control arm (trastuzumab plus docetaxel for HER2+, and docetaxel for HER2–) to assess the efficacy of new experimental arms but were concerned about the possibility of a drift in the pCR rates of the patient population over time and within patient subtype.

In response we modified the above Bayesian model with a time adjustment resulting in an overall “time machine” model that adjusts for any pCR drift that may occur over time within each arm and subtype. Having multiple arms in the trial with different time periods during which they are accruing patients enabled bridging across the different eras of trial accrual. The “time machine” discounts results from the past, with more discounting if they are further in the past. The mathematical basis and motivation for our approach to time adjustment is a statistical model for bridging eras in sports (Berry et al., 2012).

This model, described below, enables use of all available controls over all time periods in assessing efficacy of an experimental agent. We have used the time machine in all analyses in I-SPY 2 since Amendment #11 in July 2014, including for durvalumab + olaparib.

Time adjusted model (Time Machine): We explicitly incorporate terms in the model to account for time trends in pCR response. This is done by using a set of time-dependent

offset terms in the above Bayesian logistic model. Time is set to 0 at each analysis. We partition time in the past into bins of 90 days each. The index of the most recent bin, 0–90 days, is 1. The index of the bin 91–180 days in the past is 2. And so on.

Let t_i be the index of the bin for the randomization time of patient i . We model time-trend parameters $\delta(t)$ within each bin t . These are additive parameters in the model for the log-odds ratio of pCR rate. We use two sets of time-trend parameters, $\delta_+(t)$ for HER2-positive and $\delta_-(t)$ for HER2-negative. Consider patient i who has subtype (HR–, HER2+, MP–) and was randomized 750 days before present. Her bin t_i is 9 and her time-trend offset is $\delta_+(9)$.

We set $\delta_+(t) = \delta_-(t) = 0$ for $t = 1, 2, 3, 4$ which means that for a one-year period from the time of the analysis the pCR rate is assumed to be constant. Beyond a year in the past, we model the $\{\delta(t)\}$ as a second-order Normal Dynamic Linear Model (NDLM) (West and Harrison, 1997). This enables us to fit a ‘smooth’ effect over time. Suppressing the + and – subscripts, it has the following structure:

$$\delta(1) = \delta(2) = \dots = \delta(4) = 0$$

$$\delta(5) \sim N(\mu_0, \tau_0^2)$$

$$\delta(6) - \delta(5) \sim N(\mu_1, \tau_1^2)$$

$$\delta(t) - 2\delta(t-1) + \delta(t-2) \sim N(0, \tau^2) \text{ for } t > 6$$

$$\tau^2 \sim IG(\alpha, \beta)$$

For the priors in the model described above, the Normal priors are $N(\mu_0, \tau_0^2) = N(\mu_1, \tau_1^2) = N(0, 0.1)$ and $\alpha = 1$, $\beta = 0.001$ for the Gamma priors.

After including the time trend component, the full Bayesian logistic regression model is:

$$\begin{aligned} y_i &\sim \text{Bernoulli}(p_i) \\ \text{logit}(p_i) &= \beta_0 + \beta_1 x_{1i} + \beta_2 x_{2i} + \beta_3 x_{3i} + \theta_i + \gamma_{1,t_i} x_{1i} + \gamma_{2,t_i} x_{2i} + \gamma_{3,t_i} x_{3i} + \delta_-(t_i) I(x_{2i} = 0) \\ &\quad + \delta_+(t_i) I(x_{2i} = 1) \end{aligned}$$

(Where $I()$ is the indicator function.)

Though in principle the time machine model can be applied at any time to the totality of results in I-SPY 2, the final efficacy analysis for any investigational arm includes only data available up until pathology results are complete for all patients in that arm (from all I-SPY 2 patients enrolled prior to arm closure). Although the model uses data from

multiple investigational arms along with control, we only report the probability distribution of the specific investigational arm of interest and controls, and compute the probability that the pCR rate of that arm is greater than control and the predictive probabilities of success in a future trial (against control) for each biomarker signature. Comparisons between investigational arms are not reported or otherwise announced.

Biomarker Analysis—Biomarker analysis in I-SPY2 employs a predefined 3-step Qualifying Biomarker Evaluation method (Wolf et al., 2017) First, we assess association of the biomarker with response in the experimental and control arms using a logistic model (likelihood ratio (LR) test $p < 0.05$). Relative performance between arms is assessed using a logistic model (biomarker \times treatment interaction, likelihood ratio $p < 0.05$). If the biomarker \times treatment interaction term coefficient is significant (LR $p < 0.05$), and if the predictive marker is significantly associated with response in the experimental arm (LR $p < 0.05$), the marker succeeds as a qualifying biomarker (QB). Analysis is also performed adjusting for HR status as a covariate, and numbers permitting, within receptor subsets. Our statistics are descriptive rather than inferential and do not adjust for multiplicities. pCR rates within MP1/2 classes are estimated using Bayesian logistic modeling. Analyses were performed in the computing environment R (v.3.6.3) using R Packages ‘stats’ (v.3.6.3), ‘lmtest’ (v.0.9–37), ‘rjags’ (v.4–10). Scripts are available upon request.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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ROLE OF FUNDING SOURCE

The trial was designed by the I-SPY2 study investigators. AstraZeneca provided study drug and funding, but played no role in the study design, collection/analysis of data or in manuscript preparation. The authors of the manuscript vouch for the accuracy and completeness of the data.

INCLUSION AND DIVERSITY

We worked to ensure gender balance in the recruitment of human subjects. We worked to ensure ethnic or other types of diversity in the recruitment of human subjects. We worked to

ensure that the study questionnaires were prepared in an inclusive way. One or more of the authors of this paper self-identifies as an underrepresented ethnic minority in science. While citing references scientifically relevant for this work, we also actively worked to promote gender balance in our reference list. The author list of this paper includes contributors from the location where the research was conducted who participated in the data collection, design, analysis, and/or interpretation of the work.

REFERENCES

- Ayers M, Lunceford J, Nebozhyn M, Murphy E, Loboda A, Kaufman DR, Albright A, Cheng JD, Kang SP, Shankaran V, et al. (2017). IFN- γ -related mRNA profile predicts clinical response to PD-1 blockade. *J Clin Invest* 127, 2930–2940. [PubMed: 28650338]
- Barker A, Sigman C, Kelloff G, Hylton N, Berry D, and Esserman L (2009). ISPY 2: An Adaptive Breast Cancer Trial Design in the Setting of Neoadjuvant Chemotherapy. *Clin Pharmacol Ther* 86, 97–100. [PubMed: 19440188]
- Berry DA (2011). Adaptive clinical trials in oncology. *Nature Reviews Clinical Oncology* 9, 199–207.
- Berry SM, Reese CS, and Larkey PD (2012). Bridging Different Eras in Sports. *J Am Stat Assoc* 94, 661–676.
- Bianchini G, Pusztai L, Karn T, Iwamoto T, Rody A, Kelly CM, Müller V, Schmidt M, Qi Y, Holtrich U, et al. (2013). Proliferation and estrogen signaling can distinguish patients at risk for early versus late relapse among estrogen receptor positive breast cancers. *Breast Cancer Res* 15, R86. [PubMed: 24060333]
- Campbell MJ, Wolf D, Mukhtar RA, Tandon V, Yau C, Au A, Baehner F, Veer L van't, Berry D, and Esserman LJ. (2013). The Prognostic Implications of Macrophages Expressing Proliferating Cell Nuclear Antigen in Breast Cancer Depend on Immune Context. *PLOS ONE* 8, e79114. [PubMed: 24205370]
- Cardoso F, Veer LJ van't, Bogaerts J, Slaets L, Viale G, Delaloge S, Pierga J-Y, Brain E, Causeret S, DeLorenzi M, et al. (2016). 70-Gene Signature as an Aid to Treatment Decisions in Early-Stage Breast Cancer. *New England Journal of Medicine* 375, 717–729. [PubMed: 27557300]
- Daemen A, Wolf DM, Korkola JE, Griffith OL, Frankum JR, Brough R, Jakkula LR, Wang NJ, Natrajan R, Reis-Filho JS, et al. (2012). Cross-platform pathway-based analysis identifies markers of response to the PARP inhibitor olaparib. *Breast Cancer Res Tr* 135, 505–517.
- Danaher P, Warren S, Dennis L, D'Amico L, White A, Disis ML, Geller MA, Odunsi K, Beechem J, and Fling SP (2017). Gene expression markers of Tumor Infiltrating Leukocytes. *J Immunother Cancer* 5, 18. [PubMed: 28239471]
- Domchek SM, Postel-Vinay S, Im S-A, Park YH, Delord J-P, Italiano A, Alexandre J, You B, Bastian S, Krebs MG, et al. (2020). Olaparib and durvalumab in patients with germline BRCA-mutated metastatic breast cancer (MEDIOLA): an open-label, multicentre, phase 1/2, basket study. *Lancet Oncol*.
- Esteva FJ, Hubbard-Lucey VM, Tang J, and Pusztai L (2019). Immunotherapy and targeted therapy combinations in metastatic breast cancer. *Lancet Oncol* 20, e175–e186. [PubMed: 30842061]
- Hoffman-LaRoche F (2020). Roche's Tecentriq in combination with chemotherapy (including Abraxane) meets primary endpoint of improved pathological complete response, regardless of PD-L1 status, as initial treatment for people with early triple-negative breast cancer.
- Foldi J, Silber A, Reisenbichler E, Singh K, Fischbach N, Persico J, Adelson K, Katoch A, Horowitz N, Lannin D, et al. (2021). Neoadjuvant durvalumab plus weekly nab-paclitaxel and dose-dense doxorubicin/cyclophosphamide in triple-negative breast cancer. *Npj Breast Cancer* 7, 9. [PubMed: 33558513]
- Gianni L, Huang C-S, Egle D, Bermejo B, Zamagni C, Thill M, Anton A, Zambelli S, Bianchini G, Russo S, et al. (2020). Abstract GS3-04: Pathologic complete response (pCR) to neoadjuvant treatment with or without atezolizumab in triple negative, early high-risk and locally advanced breast cancer. NeoTRIPaPDL1 Michelangelo randomized study. *Cancer Res* 80, GS3-04–GS3-04.

- Lassen U (2019). Combining PARP inhibition with PD-1 inhibitors. *Lancet Oncol* 20, 1196–1198. [PubMed: 31378457]
- Loibl S, Untch M, Burchardi N, Huober J, Sinn BV, Blohmer JU, Grischke EM, Furlanetto J, Tesch H, Hanusch C, et al. (2019). A randomised phase II study investigating durvalumab in addition to an anthracycline taxane-based neoadjuvant therapy in early triple-negative breast cancer: clinical results and biomarker analysis of GeparNuevo study. *Annals of Oncology* 30, 1279–1288. [PubMed: 31095287]
- Nanda R, Liu MC, Yau C, Shatsky R, Pusztai L, Wallace A, Chien AJ, Forero-Torres A, Ellis E, Han H, et al. (2020). Effect of Pembrolizumab Plus Neoadjuvant Chemotherapy on Pathologic Complete Response in Women With Early-Stage Breast Cancer: An Analysis of the Ongoing Phase 2 Adaptively Randomized I-SPY2 Trial. *Jama Oncol* 6, 676–684. [PubMed: 32053137]
- Oken MM, Creech RH, Tormey DC, Horton J, Davis TE, McFadden ET, and Carbone PP (1982). Toxicity and Response Criteria of the Eastern-Cooperative-Oncology-Group. *American Journal of Clinical Oncology-Cancer Clinical Trials* 5, 649–655.
- Pantelidou C, Sonzogni O, Taveira MDO, Mehta AK, Kothari A, Wang D, Visal T, Li MK, Pinto J, Castrillon JA, et al. (2019). PARP Inhibitor Efficacy Depends on CD8+ T-cell Recruitment via Intratumoral STING Pathway Activation in BRCA-Deficient Models of Triple-Negative Breast Cancer. *Cancer Discov* 9, 722–737. [PubMed: 31015319]
- Park JW, Liu MC, Yee D, Yau C, Veer LJ van t, Symmans WF, Paoloni M, Perlmutter J, Hylton NM, Hogarth M, et al. (2016). Adaptive Randomization of Neratinib in Early Breast Cancer. *New England Journal of Medicine* 375, 11–22. [PubMed: 27406346]
- Peyraud F, and Italiano A (2020). Combined PARP Inhibition and Immune Checkpoint Therapy in Solid Tumors. *Cancers* 12, 1502. [PubMed: 32526888]
- Pusztai L, Reisenbichler E, Bai Y, Fischbach N, Persico J, Adelson K, Katoch A, Horowitz N, Lannin D, Killelea B, et al. (2020). Abstract PD1–01: Durvalumab (MEDI4736) concurrent with nab-paclitaxel and dose dense doxorubicin cyclophosphamide (ddAC) as neoadjuvant therapy for triple negative breast cancer (TNBC). *Cancer Res* 80, PD1-01–PD1-01.
- Rody A, Holtrich U, Pusztai L, Liedtke C, Gaetje R, Ruckhaeberle E, Solbach C, Hanker L, Ahr A, Metzler D, et al. (2009). T-cell metagene predicts a favorable prognosis in estrogen receptor-negative and HER2-positive breast cancers. *Breast Cancer Res* 11, R15. [PubMed: 19272155]
- Rugo HS, Olopade OI, DeMichele A, Yau C, Veer LJ van t, Buxton MB, Hogarth M, Hylton NM, Paoloni M, Perlmutter J, et al. (2016). Adaptive Randomization of Veliparib–Carboplatin Treatment in Breast Cancer. *New England Journal of Medicine* 375, 23–34. [PubMed: 27406347]
- Schmid P, Cortés J, Pusztai L, McArthur H, Kümmel S, Bergh J, Denkert C, Park YH, Hui R, Harbeck N, et al. (2020). Pembrolizumab for Early Triple-Negative Breast Cancer. *New England Journal of Medicine* 382, 810–821. [PubMed: 32101663]
- Sinn BV, Fu C, Lau R, Litton J, Tsai T-H, Murthy R, Tam A, Andreopoulou E, Gong Y, Murthy R, et al. (2019). SETER/PR: a robust 18-gene predictor for sensitivity to endocrine therapy for metastatic breast cancer. *Npj Breast Cancer* 5, 16. [PubMed: 31231679]
- Stewart R, Morrow M, Hammond SA, Mulgrew K, Marcus D, Poon E, Watkins A, Mullins S, Chodorge M, Andrews J, et al. (2015). Identification and Characterization of MEDI4736, an Antagonistic Anti–PD-L1 Monoclonal Antibody. *Cancer Immunol Res* 3, 1052–1062. [PubMed: 25943534]
- Symmans WF, Hatzis C, Sotiriou C, André F, Peintinger F, Regitnig P, Daxenbichler G, Desmedt C, Domont J, Marth C, et al. (2010). Genomic index of sensitivity to endocrine therapy for breast cancer. *Journal of Clinical Oncology* 28, 4111–4119. [PubMed: 20697068]
- Symmans WF, Wei C, Gould R, Yu X, Zhang Y, Liu M, Walls A, Bousamra A, Ramineni M, Sinn B, et al. (2017). Long-Term Prognostic Risk After Neoadjuvant Chemotherapy Associated With Residual Cancer Burden and Breast Cancer Subtype. *Journal of Clinical Oncology* 35, 1049–1060. [PubMed: 28135148]
- West M, and Harrison J (1997). *Bayesian Forecasting and Dynamic Models* (New York, NY: Springer).
- Wolf DM, Yau C, Sanil A, Glas A, Petricoin E, Wulfkuhle J, Severson TM, Linn S, Brown-Swigart L, Hirst G, et al. (2017). DNA repair deficiency biomarkers and the 70-gene ultra-high risk signature

as predictors of veliparib/carboplatin response in the I-SPY 2 breast cancer trial. *Npj Breast Cancer* 3, 31. [PubMed: 28948212]

Yau C, Wolf D, Brown-Swigart L, Hirst G, Sanil A, Singhrao R, Investigators I-S 2 T., Asare, S., DeMichele, A., Berry, D., et al. (2018). Abstract PD6-14: Analysis of DNA repair deficiency biomarkers as predictors of response to the PD1 inhibitor pembrolizumab: Results from the neoadjuvant I-SPY 2 trial for stage II-III high-risk breast cancer. *Cancer Res* 78, PD6-14.

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HIGHLIGHTS

Durvalumab plus olaparib improved chemotherapy efficacy in HER2-negative breast cancer

Immune rich tumors had greater sensitivity to therapy

Among ER+ cancer, only Mammaprint MP2 cancers benefited from immune checkpoint therapy

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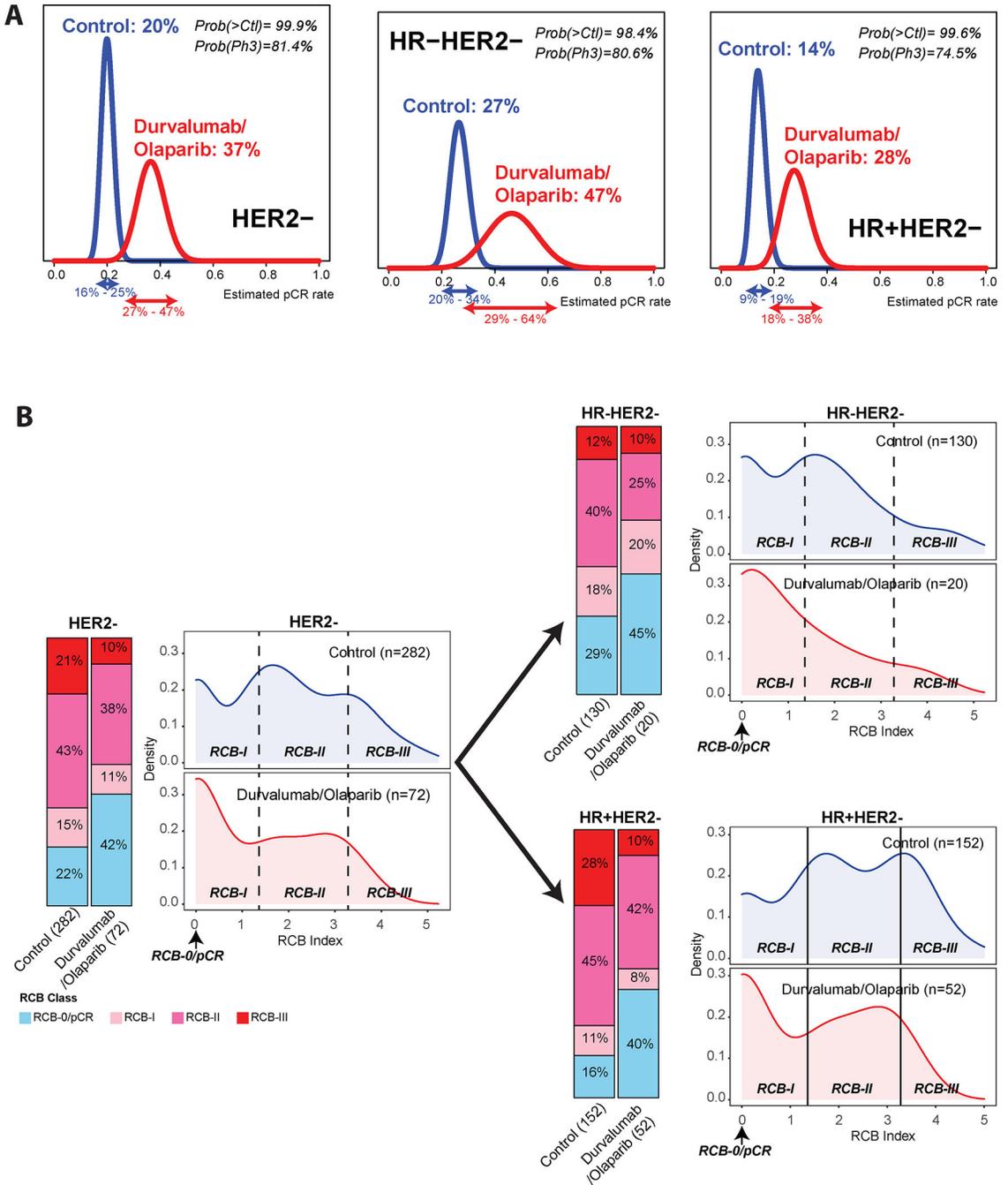


Figure 1. Bayesian probability distributions for achieving pathologic complete response (pCR) and residual cancer burden (RCB) distributions by treatment arm in the three a priori defined biomarker subsets.

A. Estimated pCR rates and the corresponding 95% probability intervals (X -axes) for the experimental and control arms. Final predictive probabilities of success in a future 300 patient phase 3 trial (Prob(Ph3)), and the probabilities that the experimental arm is superior to the control (Prob(>Ctl)) are also shown. **B.** RCB categories and distribution of RCB scores by treatment arm in the 3 biomarker subsets.

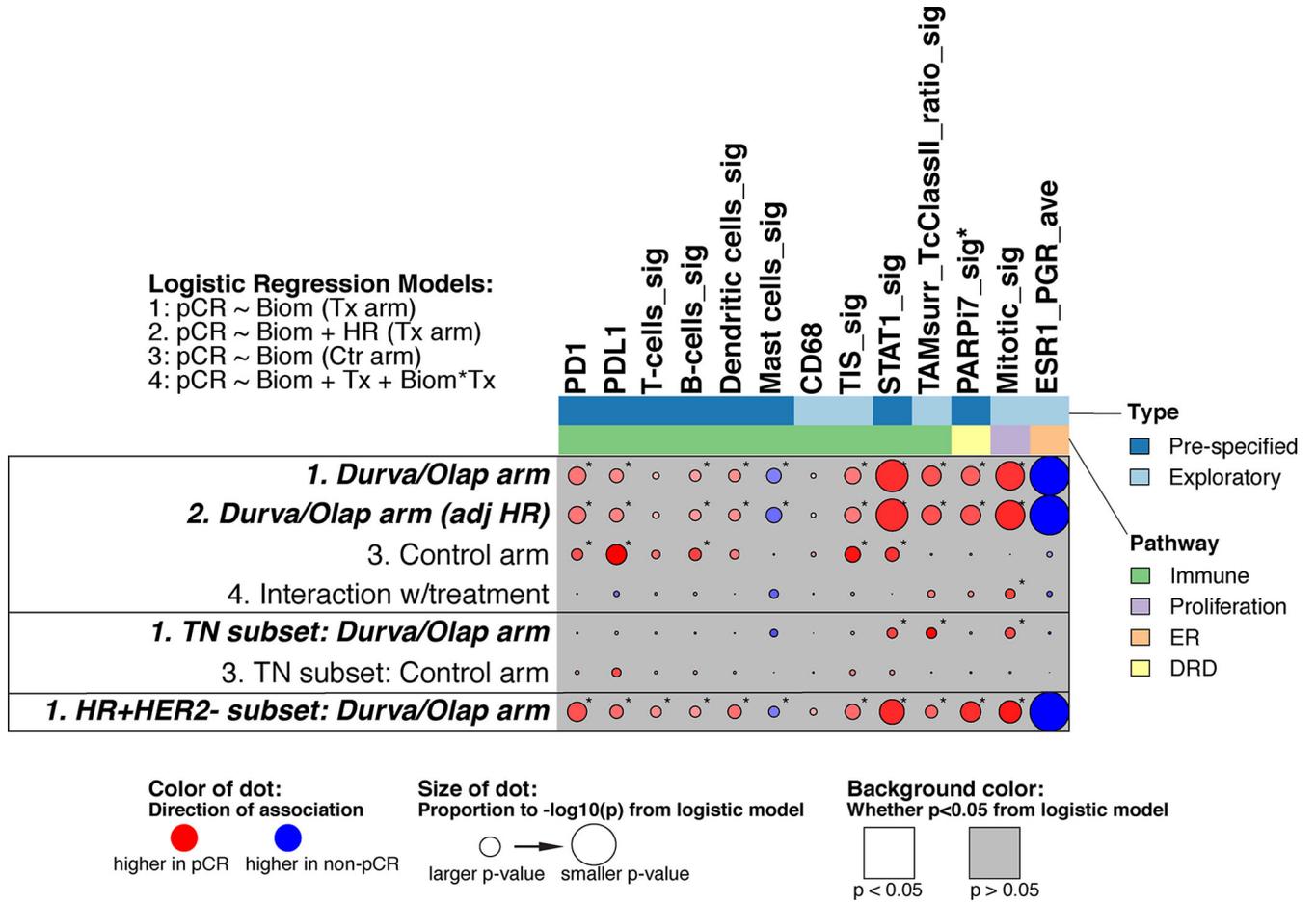


Figure 2. Predictive function of 13 mRNA markers in all HER2-negative, triple negative cancers, and HER2 negative/hormone receptor positive cancers.

Red dots indicate a positive association with pathologic complete response (pCR), blue indicates association with residual disease. The size of the dots is inversely proportional to the p-values; statistically significant associations are highlighted by the white rectangles. P-values and associations were derived from logistic regression models as indicated by the numbers (1 through 4) on the figure.

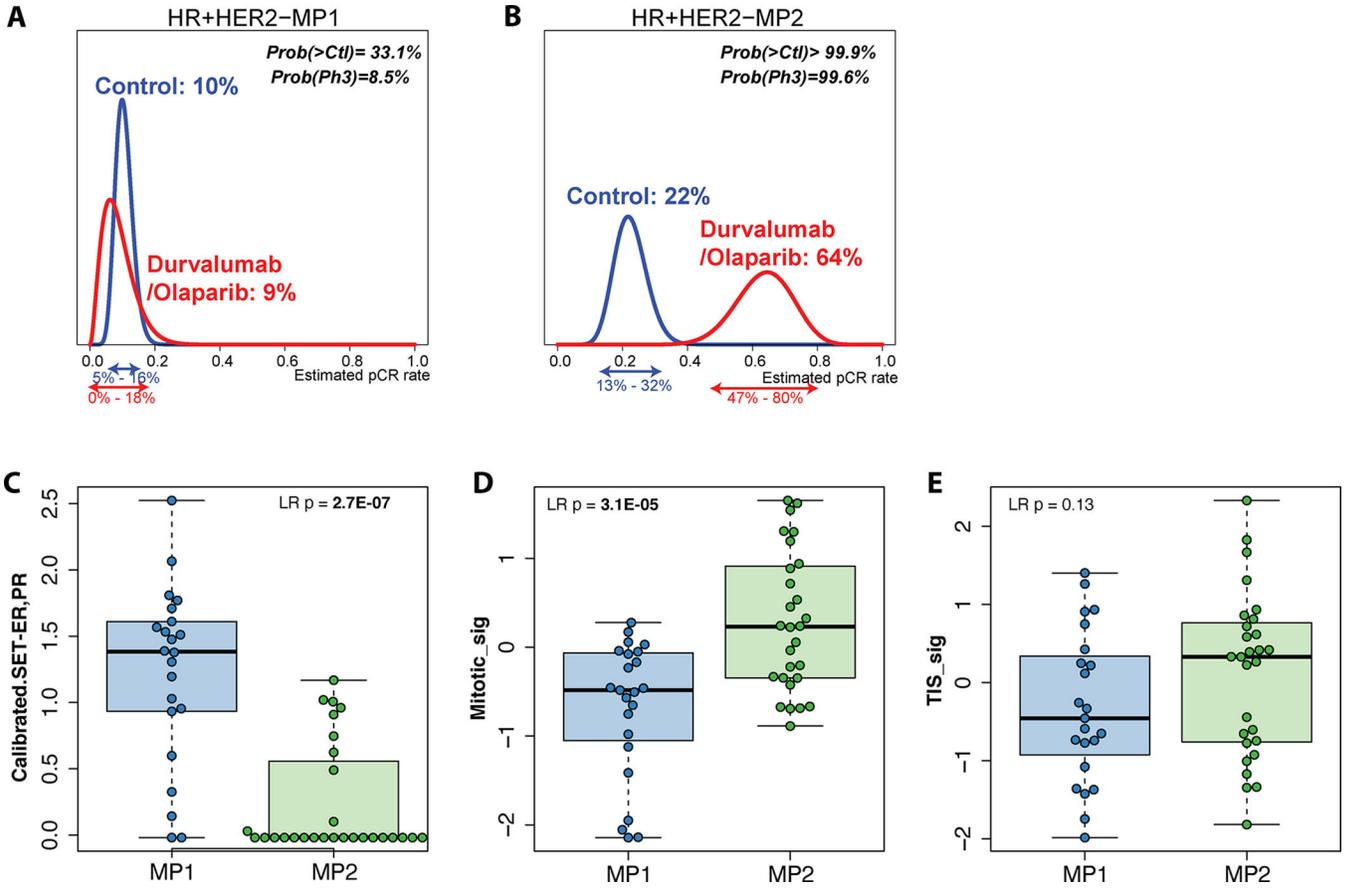


Figure 3. Bayesian probability distribution of pCR, and expression of proliferation, tumor inflammation, and sensitivity to endocrine therapy (SET) gene signatures in MP1 and MP2 ER+/HER2- cancers.

Estimated pCR rates and the corresponding 95% probability intervals (X -axes) for the experimental and control arms in MP1 (A) and MP2 (B) cancers. Final predictive probabilities of success in a future 300 patient phase 3 trial (Prob(Ph3)), and the probabilities that the experimental arm is superior to the control (Prob(>Ctl)) are also shown. Expression of the SET gene signature (C), and proliferation (D), and tumor inflammation (E) signatures in MP1 and MP2 cancers.

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Table 1:

Demographics and baseline characteristics of participants

Characteristic	duvalumab/olaparib (n=73)	Control (n=299)
Median age, yr (range)	46 (28–71)	48 (24–80)
Ethnicity, n (%)		
White	59 (81%)	234 (78%)
African American	8 (11%)	40 (13%)
Asian	6 (8%)	22 (7%)
Other/Mixed	0 (0%)	3 (1%)
HR Status, n (%)		
Positive	52 (71%)	157 (53%)
Negative	21 (29%)	142 (47%)
Mammprint, n (%)		
MP1	26 (36%)	133 (44%)
MP2	47 (64%)	166 (56%)
Median Tumor Size, cm (range)	3.7 (1.9–13)	3.8 (1.2 – 15)
Baseline node status, n (%)		
Palpable	21 (29%)	109 (36%)
Non-palpable	19 (26%)	129 (43%)
N/A	33 (45%)	61 (20%)

Table 2.
Grade 3 adverse events, dose reduction and discontinuations rates.

Grade 3 adverse events reported during the entire treatment period (durvalumab/olaparib/paclitaxel [DOP] followed by doxorubicin/cyclophosphamide [AC] for the investigational arm and paclitaxel followed by AC for the control arm) are shown. Dose reductions and discontinuations are shown by treatment period.

Adverse Event	Agent (n=73)		Control (n=299)	
	Grade 1–2	Grade 3–4	Grade 1–2	Grade 3–4
Neutropenia	10 (13.7%)	9 (12.3%)	17 (5.7%)	27 (9%)
Febrile neutropenia	0 (0%)	7 (9.6%)	0 (0%)	21 (7%)
Fatigue	63 (86.3%)	4 (5.5%)	202 (67.6%)	4 (1.3%)
Anaemia	20 (27.4%)	3 (4.1%)	44 (14.7%)	13 (4.3%)
Neuropathy	50 (68.5%)	3 (4.1%)	176 (58.9%)	6 (2%)
Diarrhoea	43 (58.9%)	3 (4.1%)	97 (32.4%)	4 (1.3%)
Adrenal insufficiency	1 (1.4%)	3 (4.1%)	0 (0%)	0 (0%)
White blood cell count decr.	8 (11%)	2 (2.7%)	14 (4.7%)	9 (3%)
Hypertension	6 (8.2%)	2 (2.7%)	36 (12%)	6 (2%)
Alanine aminotransferase incr.	11 (15.1%)	2 (2.7%)	26 (8.7%)	4 (1.3%)
Stomatitis	27 (37%)	2 (2.7%)	77 (25.8%)	3 (1%)
Dehydration	5 (6.8%)	2 (2.7%)	13 (4.3%)	2 (0.7%)
Dyspnoea	19 (26%)	2 (2.7%)	46 (15.4%)	1 (0.3%)
Rash maculo-papular	13 (17.8%)	2 (2.7%)	48 (16.1%)	0 (0%)
Vaginal infection	3 (4.1%)	2 (2.7%)	4 (1.3%)	0 (0%)
Headache	31 (42.5%)	1 (1.4%)	88 (29.4%)	3 (1%)
Abdominal pain	23 (31.5%)	1 (1.4%)	33 (11%)	2 (0.7%)
Aspartate aminotransferase incr.	10 (13.7%)	1 (1.4%)	19 (6.4%)	2 (0.7%)
Back pain	21 (28.8%)	1 (1.4%)	40 (13.4%)	2 (0.7%)
Nausea	53 (72.6%)	1 (1.4%)	176 (58.9%)	1 (0.3%)
Non-cardiac chest pain	4 (5.5%)	1 (1.4%)	14 (4.7%)	1 (0.3%)
Hypokalaemia	5 (6.8%)	0 (0%)	14 (4.7%)	8 (2.7%)
Arthralgia/myalgia	46 (63%)	0 (0%)	149 (49.8%)	5 (1.7%)
Hyperglycaemia	5 (6.8%)	0 (0%)	9 (3%)	3 (1%)
Pneumonitis	4 (5.5%)	0 (0%)	2 (0.7%)	2 (0.7%)
Vomiting	28 (38.4%)	0 (0%)	50 (16.7%)	2 (0.7%)
Onychalgia	3 (4.1%)	0 (0%)	17 (5.7%)	1 (0.3%)
Pruritus	11 (15.1%)	0 (0%)	32 (10.7%)	1 (0.3%)
Pyrexia	23 (31.5%)	0 (0%)	36 (12%)	1 (0.3%)
Urinary tract infection	4 (5.5%)	0 (0%)	11 (3.7%)	1 (0.3%)
Dose Reductions, n (%)	10 (13.7%)		23 (7.7%)	
Early Discontinuation, n (%)				

	Agent (n=73)	Agent (n=73)	Control (n=299)	Control (n=299)
All	19 (26%)		90 (30%)	
Toxicity	9 (12%)		20 (7%)	
Progression/lack of response	0 (0%)		33 (11%)	
Other	10 (14%)		37 (12%)	
Time from Treatment Consent to Surgery (days)				
Median (range)	168 (107 – 273)		165 (85 – 289)	

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KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Biological samples		
Tumor biopsy before treatment	I-SPY 2 TRIAL	https://clinicaltrials.gov/ct2/show/NCT01042379
Critical commercial assays		
Custom Agilent 44K expression arrays	Agendia, Inc	https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GPL20078
MammaPrint	Agendia, Inc	https://agendia.com/mammaprint/
Deposited data		
Raw and processed transcriptomic data	This study	GEO accession number: GSE173839 https://www.ncbi.nlm.nih.gov/geo/auev/acc.cgi?acc=GSE173839
Patient-level expression signature and clinical data	This study	GEO accession number: GSE173839 https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE173839
Software and algorithms		
stats R package (v.3.6.3)	R Core Team (2020)	https://stat.ethz.ch/R-manual/R-devel/library/stats/html/stats-package.html
lmtest R package (v.0.9–37)	Zeileis A, Hothorn T (2002). “Diagnostic Checking in Regression Relationships.” R News, 2(3), 7–10.	https://CRAN.R-project.org/package=lmtest
rjags R package (v.4–10)	Martyn Plummer (2019). rjags: Bayesian Graphical Models using MCMC. R package v4–10.	https://CRAN.R-project.org/package=rjags