ABSTRACT

Triple-negative breast cancer (TNBC) remains the most challenging breast cancer subtype to treat. To date, therapies directed to specific molecular targets have rarely achieved clinically meaningful improvements in outcomes of patients with TNBC, and chemotherapy remains the standard of care. Here, we seek to review the most recent efforts to classify TNBC based on the comprehensive profiling of tumors for cellular composition and molecular features. Technologic advances allow for tumor characterization at ever-increasing depth, generating data that, if integrated with clinical–pathologic features, may help improve risk stratification of patients, guide treatment decisions and surveillance, and help identify new targets for drug development.

Significance: TNBC is characterized by higher rates of relapse, greater metastatic potential, and shorter overall survival compared with other major breast cancer subtypes. The identification of biomarkers that can help guide treatment decisions in TNBC remains a clinically unmet need. Understanding the mechanisms that drive resistance is key to the design of novel therapeutic strategies to help prevent the development of metastatic disease and, ultimately, to improve survival in this patient population.

INTRODUCTION

Breast cancer is the most frequently diagnosed cancer and the second most common cause of cancer mortality in women worldwide (1). Breast tumors that are immunohistochemically characterized by lack of estrogen receptor (ER), progesterone receptor (PR), and HER2 (also defined by lack of HER2 amplification by FISH) are classified as triple-negative breast cancer (TNBC) and account for approximately 15% to 20% of all breast carcinomas (2). Compared with hormone receptor–positive or HER2-positive disease, TNBC has a highly aggressive clinical course, with earlier age of onset, greater metastatic potential, and poorer clinical outcomes as shown by the higher relapse and lower survival rates (2, 3). The molecular mechanisms that drive TNBC recurrence have not been fully elucidated. Consequently, to date, targeted therapies have not significantly improved survival in patients with TNBC, and chemotherapy remains the standard of care. Although many patients with early stages of TNBC are cured with chemotherapy, in those who develop metastatic disease, median overall survival (OS) with current treatment options is 13 to 18 months (4).

Major effort has been devoted over the past decade to classify TNBC into distinct clinical and molecular subtypes that could guide treatment decisions. Characterization of genomic, transcriptomic, proteomic, epigenomic, and micro-environmental alterations has expanded our knowledge of TNBC. Here, we review the most recent innovations in TNBC molecular taxonomy, the complex interaction between these classifications (Fig. 1), and their potential therapeutic implications.

TNBC AND INTRINSIC BREAST CANCER SUBTYPES

Early transcriptomic profiling of breast cancer using microarrays classified tumors into five intrinsic subtypes: luminal-A, luminal-B, HER2-enriched, basal-like, and a normal breast-like group (5, 6). Although all intrinsic subtypes can be found within immunohistochemically defined triple-negative disease, basal-like tumors exhibit the greatest overlap with TNBC. Between 50% and 75% of TNBC have basal phenotype, and approximately 80% of basal-like tumors are ER-negative/HER2-negative (Fig. 2; refs. 7, 8). Characterization
Heterogeneity of Triple-Negative Breast Cancer

Figure 1. Overview of the complex interactions among molecular classifications of TNBC based on genomic, transcriptomic, proteomic, epigenomic, and immune characterization of the tumor and its microenvironment. ER, estrogen receptor; PR, progesterone receptor; Mut, mutant; RTK, receptor tyrosine kinase; MMR, mismatch repair; CNA, copy-number alteration; AR, androgen receptor; HRD, homologous recombination deficiency; IHC, immunohistochemistry.

of intrinsic subtypes using a 50-gene assay (established as the PAM50 subtype predictor) has provided independent predictive information of pathologic complete response (pCR) to neoadjuvant therapy across all subtypes (9), but when restricting analyses to TNBC, none of the PAM50 signatures at the time of diagnosis have significantly correlated with pCR (10). In basal-like TNBC, low expression of the luminal-A signature and high expression of the proliferation score were both significantly associated with pCR (10). High expression of cell cycle–related genes (e.g., CCNE and FANCA) and low levels of estrogen signaling–related genes (e.g., FOXA1 and PGR) were associated with pCR, whereas high expression of epithelial–mesenchymal transition (EMT) genes (e.g., TWIST1 and ZEB1) was significantly enriched in residual disease (10). Again, in the adjuvant setting, no significant gene-signature predictors of disease-free survival (DFS) have been found in TNBC (10). However, in basal-like TNBC in GEICAM/9906, and in basal-like tumors treated with adjuvant chemotherapy in the METABRIC data set and in CALGB/9741, the two previously identified signatures (low luminal-A and high proliferation score) predicted improved DFS and recurrence-free survival (RFS).

PAM50-defined subtypes have not yet been validated as predictors of benefit to individual chemotherapeutic agents in TNBC. An increase in pCR rates from 47% to 61% was noted with the addition of carboplatin to neoadjuvant therapy in patients with basal-like TNBC in CALGB/40603 (11), although this improvement did not differ from that observed in the overall population after incorporating the small number of non–basal-like tumors. In the metastatic setting, carboplatin and docetaxel achieved comparable objective response rates (ORR) in basal-like tumors in the TNT trial (32.5% vs. 31.0%, respectively; \( P = 0.87 \); ref. 12). Of note, though a significant interaction was observed between PAM50 subgroups and treatment arm, this was driven by the unexpected finding of greater efficacy of docetaxel compared with carboplatin in non–basal-like tumors (ORR, 72.2% vs. 16.7%; \( P = 0.002 \); ref. 12). Further studies prospectively evaluating taxanes and other agents in predefined subgroups are needed to confirm any differential activity in non–basal-like TNBC.

Additional gene-expression analyses later revealed the presence of another intrinsic subtype, claudin-low, present in 7% to 14% of all breast cancers (6). Approximately 70% of claudin-low tumors are TNBC, with high representation of metaplastic and medullary breast carcinomas. Although claudin-low and basal-like subtypes share low luminal and HER2 gene expression, claudin-low tumors do not highly express proliferation genes. They are uniquely characterized by low levels of cell adhesion proteins and elevated expression of immune-related genes (e.g., CD4 and CD79a). These mesenchymal features (including elevated expression of CD44, vimentin, and N-cadherin) and low epithelial differentiation (low CD24 gene expression) resemble a mammary stem cell–like phenotype (CD44+/CD24−) that can be acquired by breast cancer cells. This stem cell–like phenotype is associated with lower (39%) pCR rates compared with basal-like subtype (73%), and worse prognosis than luminal-A tumors but similar survival as luminal-B, HER2-enriched, or basal-like tumors (6). Formation of cancer stem cells is induced by TGFβ in claudin-low cell lines (13), and in chemotherapy-resistant TNBC, TGFβ signaling and other stem cell markers are overexpressed (14). Thus, inhibition of TGFβ...
signaling may represent a potential therapeutic strategy to help prevent the development of chemorefractory disease, particularly in the claudin-low subtype.

### MOLECULAR DEFINITION OF TNBC HETEROGENEITY

With evolving transcriptomic studies, the heterogeneity of TNBC has been further dissected. Lehmann and colleagues analyzed 21 public microarray data sets filtered for TNBC based on ESR1, PGR, and ERBB2 expression and identified seven clusters within TNBC: basal-like 1 (BL1), basal-like 2 (BL2), immunomodulatory (IM), mesenchymal (M), mesenchymal-stem-like (MSL), luminal androgen receptor (LAR), and an unstable cluster (UNS; ref. 15). These subtypes are characterized by distinct patterns of molecular alterations, in terms of RNA expression, somatic mutations, and copy-number variations, that tend to cluster in genes implicated in specific pathways. The BL1 subtype, enriched in genes involved in DNA-damage response and cell-cycle regulation (including the highest rate of TP53 mutations (92%), high gain/amplifications of MYC, CDK6, or CCNE1, and deletions in BRCA2, PTEN, MDM2, and RB1; ref. 16), and the BL2 subtype, with high levels of growth factor signaling and metabolic pathway activity, share a highly proliferative phenotype that correlates with improved pCR with mitotic inhibitors, such as taxanes. Genes involved in antigen processing and presentation, immune cell and cytokine signaling (e.g., JAK/STAT, TNF, and NFκB) pathways are highly expressed in the IM subtype. Mesenchymal-like TNBC subtypes, M and MSL, display similar expression profiles related to cell motility, differentiation, and EMT, but are discernible by the unique enrichment in MSL of angiogenesis- and stem cell–associated genes, and low claudin expression. Finally, despite ER negativity, the LAR subtype displays a luminal pattern of gene expression (e.g., high levels of FOXA1, GATA3, SPDEF, and XBPI), with elevated

**Figure 2.** Distribution of intrinsic subtypes among TNBC and distribution of TNBC among basal-like breast cancer. A, Comparison of distribution of intrinsic subtypes defined by PAM50 and PAM50 + claudin-low in The Cancer Genome Atlas (TCGA) and METABRIC data sets in TNBC. TNBC was defined as clinical ER-, PR-, and HER2-negative testing per IHC. In TCGA, 88 TNBC samples had available PAM50 data. The distribution of intrinsic subtypes was basal-like (86%), HER2-enriched (6%), luminal A (5%), luminal B (1%), and normal-like (2%). In METABRIC, 320 TNBC samples had available intrinsic subtype data. When including claudin-low in the PAM50 predictor, the distribution of subtypes was basal-like (49%), claudin-low (37%), HER2-enriched (9%), normal-like (4%), luminal A (1%), and luminal B (0%). When excluding the 119 samples with claudin-low subtype, the distribution of subtypes was basal-like (78%), HER2-enriched (15%), normal-like (5%), luminal A (2%), and luminal B (0%). B, Comparison of distribution of breast cancer subtype according to receptor status defined by IHC in TCGA and METABRIC data sets in basal-like breast cancer. Of 98 basal-like breast cancers in TCGA, 78% were TNBC per IHC. Of 209 basal-like breast cancers (PAM50 + claudin-low classifier) in METABRIC, 75% were TNBC. Figures generated by reanalysis of publicly available studies (refs. 22, 36, 37) using cBioPortal (refs. 150, 151).
mRNA and protein levels of androgen receptor (AR), overlapping in 82% of cases with luminal-A- or luminal-B-intrinsic subtypes. Thus, not surprisingly, LAR tumors are enriched in mutations in PIK3CA (55%), KMT2C (19%), CDH1 (13%, in conjunction with a higher prevalence of inactivating lobular histology), NFI (13%), and AKTI (13% ref. 16). The seven-subtype classification independently predicted pCR, but not distant metastasis-free or overall survival in a retrospective analysis of patients with TNBC treated with neoadjuvant chemotherapy (17). Median OS was highest in the LAR and BL1 subtypes, despite low pCR rate in the LAR group. Follow-up in vitro studies with representative cell lines of TNBC subtypes demonstrated differential drug sensitivity that, if validated, may have clinically relevant implications (15). Of note, all seven clusters were not detected in an independent analysis of five data sets of IHC-identified TNBC, as opposed to gene expression–defined TNBC (15). Even across other studies in which TNBC was identified using mRNA expression, reproducibility of the BL2 and UNS subtypes has not been consistent (16, 17).

In a follow-up study, by performing histologic assessment and laser microdissection prior to RNA isolation and gene-expression analysis, Lehmann and colleagues confirmed that the presence of stromal cells in tumor specimens—such as infiltrating lymphocytes and tumor-associated mesenchymal cells—influences the definition of the IM and MSL subtypes, respectively (18). This led to a revised classification, TNBC-type 4, into four stable transcriptional subtypes (BL1, BL2, M, and LAR) that significantly differ not only in prognosis and response to chemotherapy, but also in initial presentation and patterns of recurrence, where regional nodal involvement is more common in LAR TNBC and metastatic recurrences have tropism to the lung in M subtypes and to the bone in LAR subtypes. Similar to the seven-subtype classification, response to neoadjuvant chemotherapy (platinum- and taxane-based regimen) is significantly associated with TNBC-type 4 subtypes ($p \geq 0.027$), with the highest and lowest pCR rates reported in BL1 (65.6%) and LAR (21.4%), respectively (19). These findings highlight a major limitation of classifiers defined based on the profiling of bulk tumors that cannot distinguish between tumor and stromal cells and support the increasing use of single-cell techniques to improve the characterization of the tumor and its microenvironment. In fact, single-cell RNA sequencing has demonstrated the presence of multiple subtypes within most primary TNBC tumors, suggesting that the dominant signature identified through bulk sequencing may not accurately inform underlying biological processes, including interactions between malignant and normal stromal cell types (20). Differences in the prevalence of intratumoral heterogeneity between TNBC and ER-positive breast cancer could partly explain the challenges to date to apply commercially available gene-expression assays in routine clinical practice to demonstrate differential drug sensitivity that, if validated, may have clinically relevant implications (15). Of note, all seven clusters were not detected in an independent analysis of five data sets of IHC-identified TNBC, as opposed to gene expression–defined TNBC (15). Even across other studies in which TNBC was identified using mRNA expression, reproducibility of the BL2 and UNS subtypes has not been consistent (16, 17).

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**ANDROGEN RECEPTOR–POSITIVE TNBC**

As described above, a luminal phenotype, characterized by expression of the AR and luminal lineage-driving transcription factors, has been consistently identified across several studies in TNBC. In core-basal tumors, the prevalence of AR positivity defined by ≥1% of tumor cell nuclei IHC staining has been reported to be 52% (24). Interestingly, other studies have suggested that LAR tumors are characterized by a quiescent cell state (25), as opposed to rapidly proliferative basal tumors, raising the question of the optimal method of testing for AR positivity and possibly lack of a robust approach due to limited sample size. Altogether, this has prompted interest in exploring the role of antiandrogens in this subgroup. In *in vitro* studies have shown that tumors derived from LAR cell lines (e.g., MDA-MB-453, SUM158PE, and CAL-148) are highly sensitive to the AR antagonist bicalutamide (15). In phase II single-arm trials conducted in patients with metastatic AR-positive, ER/PR-negative breast cancer, bicalutamide and enzalutamide demonstrated stable disease at 6 months of 19% and 28%, respectively, though no objective responses were observed (26, 27). Abiraterone acetate and prednisone achieved a similar 20% clinical benefit rate (CBR) at 6 months, and although the study failed to meet the prespecified >25% cutoff necessary to reject the null hypothesis, prolonged responses were observed (range, 6.4–23.4 months; ref. 28). An androgen-driven genomic signature, Dx, predicted improved OS with enzalutamide (29), and this led to the design of a phase III trial comparing enzalutamide, paclitaxel, and the combination in selected Dx-positive advanced TNBC (NCT02929576).

Similar to luminal tumors, strategies to enhance the effectiveness of hormone receptor blockade have been pursued in AR-positive TNBC. Enrichment in PIK3CA mutations has been described in triple-negative tumors that are AR-positive (36%–40%) by IHC compared with AR-negative (4%–9%; refs. 30, 31), the majority of which are located in the kinase domain.
H1047 mutational hotspot and co-occur with amplification of the PIK3CA locus (30). Combination of PI3K/mTOR inhibition and AR antagonism has demonstrated synergistic activity in AR-positive TNBC preclinical models, and a phase I trial is planned to explore enzalutamide plus alpelisib, an α-specific PI3K inhibitor, in patients with AR-positive, PTEN<sup>in</sup> (IHC 0%) TNBC (NCT03207529). Additional studies have revealed that, in contrast to basal-like and mesenchymal subtypes, LAR TNBC cell lines are highly sensitive to CDK4/6 inhibitors, with comparable sensitivity to that observed in the ER-positive MCF7 cell line (25). LAR cell lines exhibit lower transcriptomic levels of CCNE1 and CDK2 compared with basal-like TNBC and, thus, are dependent on CDK4/6 to phosphorylate RB1 and reenter the cell cycle. In vitro PI3K inhibition decreases postmitotic CDK2 activity in PIK3CA-mutant TNBC, suggesting potential sensitization to CDK4/6 inhibitors, including in non-LAR TNBC (25); this has provided the rationale for the ongoing clinical trial testing palbociclib combined with either tareculisib or pictilisib in PIK3CA-mutant ER-negative breast cancer (NCT02389842).

**PROTEIN MARKERS IN TNBC FOR TARGETED ANTIBODY–DRUG CONJUGATES**

Isolation of glycoproteins on the surface of epithelial cancer cells has triggered the development of antibody–drug conjugates (ADC) designed to improve delivery of elevated concentrations of cytotoxic drugs to cells expressing these molecules. Many of these targets are not necessarily cancer drivers or specific to breast cancer; instead, they require differential protein expression in malignant versus normal cells. Interestingly, several ADC have demonstrated encouraging activity in TNBC. Saccituzumab govitancan (IMMU-132) is an antibody–SN-38 conjugate targeting TROP2, which is expressed in almost 90% of TNBC (32). In patients with heavily pretreated metastatic TNBC, IMMU-132 achieved an ORR of 30%, and median progression-free survival (PFS) and OS were 6.0 and 16.6 months, respectively. Liv-1 is a transmembrane protein with metalloprotease activity expressed in 68% of metastatic TNBC samples. Ladratuzumab vedotin (SGN-LIV1A), with monomethyl auristatin-E (MMAE) as the payload, yielded a 25% ORR in a similar population of patients with TNBC, and median PFS was 11 months (33). Significant expression of glycoprotein-NMB (gpNMB), defined as staining ≥25% of tumor epithelial cells, is present in approximately 40% of TNBC, and in this subgroup, glembatumumab vedotin (CDX-011, an ADC that binds to gpNMB to deliver MMAE) achieved 40% ORR versus 0% with investigator’s choice of therapy (34). However, when compared with capcetibamine in preselected gpNMB-overexpressing metastatic TNBC in the METRIC phase II trial, glembatumumab vedotin failed to demonstrate improved PFS, ORR, or OS, leading to discontinuation of the development of this ADC (Cellnex’s METRIC Study Press release, April 16, 2018; https://globenewswire.com/news-release/2018/04/16/1471890/0/en/Cellnex-s-METRIC-Study-in-Metastatic-Triple-negative-Breast-Cancer-Does-Not-Meet-Primary-Endpoint.html). SGN-LIV1A is currently being evaluated in phase II trials, and IMMU-132 has advanced to phase III development (ASCENT: NCT02574455). Given the high prevalence of many of these markers in TNBC, IHC confirmation may not be necessary prior to starting therapy, but other proteins overexpressed less frequently may require prescreening efforts to help identify patients who are more likely to benefit from ADC.

**SOMATIC GENETIC ALTERATIONS IN TNBC**

Cancers harbor numerous somatic genetic alterations, though only a small proportion of them confer clear fitness advantage, also known as “cancer drivers” (35). Large-scale exome and targeted sequencing studies in primary breast tumors have revealed the presence of many alterations in putative cancer-driver genes in TNBC (36–38). The average mutation rate in basal-like breast cancer is among the highest in breast tumors, 1.68 mutations per megabase (Mb); tumors that reach rates greater than three standard deviations above the mean (>4.68 mutations/Mb) are considered hypermutated (36). Different genomic classifications in breast cancer have been proposed by grouping next-generation sequencing (NGS)-detected alterations in known cancer-driver genes according to the intracellular pathways in which they are involved, such as PI3K/AKT and RAS/MAPK signaling, DNA damage repair, and cell-cycle or transcriptional regulation (Table 1; refs. 36, 37, 39).

Most somatic mutations in TNBC occur in tumor suppressor genes (e.g., TP53, RB1, and PTEN), which have not been successfully targeted therapeutically to date. Although less prevalent, oncogenic alterations in the PI3K/AKT pathway have also been described in basal-like breast cancer (PIK3CA mutation, 7%; AKT3 amplification, 28%; PTEN mutation or loss, 35%; ref. 36), potentially qualifying patients for clinical trials with matched therapies. Consistent with findings in untreated triple-negative tumors, targeted sequencing of residual disease post-neoadjuvant chemotherapy showed that >90% of patients had at least one altered pathway (39). However, only three alterations were found to be significantly prognostic for OS (JAK2 amplification, BRCA1 truncation or mutation: predicted poor OS; PTEN alteration: better OS). Drugs that inhibit these pathways have been explored in clinical trials in TNBC, mostly in combination with other therapies due to limited single-agent activity (Table 2).

Considering the underlying complexity of the genomic landscape of TNBC, analysis of single mutations in a putative driver or known oncogenic pathway is likely insufficient (40). Different processes, such as age, exposure to carcinogens, DNA replication errors, defects in DNA repair, and the family of APOBEC cytidine deaminases, imprint patterns of mutations known as mutational signatures on the cancer genome. Whole-genome sequencing of 21 breast tumors initially showed the presence of five different mutational signatures in breast cancer, including focal hypermutation and APOBEC (40). More recently, the expanded analysis of 560 breast cancers revealed somatic base substitutions, indels, rearrangements, and copy-number alterations in 93 candidate driver genes (41). Of the 10 most frequently mutated genes that accounted for 62% of drivers in the overall set, TP53, MYC, PTEN, ERBB2, and RB1 appeared enriched in the ER-negative cohort. Application of mathematical algorithms discriminated 12 base-substitution signatures (including the five previously identified signatures), two indel signatures, and six rearrangement signatures. Large
Heterogeneity of Triple-Negative Breast Cancer

**Table 1.** Classifications according to potentially targetable pathways based on exome or targeted sequencing

<table>
<thead>
<tr>
<th>TCGA (basal-like; ref. 36)</th>
<th>Genomic alteration (frequency, %)</th>
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<tbody>
<tr>
<td>p53 pathway</td>
<td>TP53 mut (84), gain of MDM2 (14)</td>
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<tr>
<td>PI3K/PTEN pathway</td>
<td>PTEN mut/loss (35), INPP4B loss (30), PIK3CA mut (7)</td>
</tr>
<tr>
<td>RB1 pathway</td>
<td>RB1 mut/loss (20), CCNE1 amp (9), high expression of CDKN2A, low RB1 expression</td>
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**METABRIC (ER-negative; ref. 37)**

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<tr>
<th>Mutated gene (frequency, %)</th>
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<tbody>
<tr>
<td>AKT signaling</td>
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<td>Cell-cycle regulation</td>
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<tr>
<td>Chromatin function</td>
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<tr>
<td>DNA damage and apoptosis</td>
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<tr>
<td>MAPK signaling</td>
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<tr>
<td>Tissue organization</td>
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<tr>
<td>Transcription regulation</td>
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<tr>
<td>Ubiquitination</td>
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<tr>
<td>Other</td>
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**Residual disease post-neoadjuvant chemotherapy (triple-negative; ref. 39)**

<table>
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<tr>
<th>Genomic alteration (frequency, %)</th>
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<tr>
<td>Cell cycle</td>
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<tr>
<td>PI3K/mTOR pathway</td>
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<tr>
<td>Growth factor receptor</td>
</tr>
<tr>
<td>RAS/MAPK pathway</td>
</tr>
<tr>
<td>DNA repair</td>
</tr>
<tr>
<td>JAK2/STAT3 pathway</td>
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</table>

**NOTE:** Mut, gene mutation; gain, gene copy-number gain (<5 but more than 2 copies); amp, gene amplification (≥5 copies and/or gene-specific and centromeric probe ratio ≥2). The definition of copy-number gain vs. amplification is partly platform and study dependent. In general, copy-number gain ≥5 is considered an amplification, whereas copy-number gain ≥2 but below 5 is considered a copy-number gain. However, some studies define amplification when gene-specific vs. centromeric probe ratio is ≥2.

Tandem duplications (>100 kb) were associated with rearrangement signature 1, mostly found in TP53-mutated, triple-negative tumors with high homologous recombination-deficiency (HRD) index but without BRCA1/2 mutations or BRCA1 promoter hypermethylation. In contrast, 91% of cases with BRCA1 mutation or promoter hypermethylation fell into rearrangement signature 3, characterized predominantly by small tandem duplications (<10 kb). Additional research is required to fully understand the prognostic and therapeutic implications of these signatures.

**TARGETING GENETICALLY ALTERED SIGNALING PATHWAYS IN TNBC**

Tumors with genetic alterations that promote activation of the PI3K pathway, found at a higher frequency in TNBC cell lines classified as LAR and mesenchymal-like, demonstrate in vitro and in vivo sensitivity to BEZ235 (a dual PI3K and mTOR inhibitor; ref. 15). Loss of PTEN and INPP4B, which also sensitizes cell lines to PI3K inhibition (42), is more common in basal-like tumors (36). Oral pan-PI3K inhibitors, such as buparlisib (BKM120), or selective p110α-PI3K inhibitors, including alpelisib (BYL719) or telsisib (GDC-0032), have shown enhanced clinical activity in ER-positive PIK3CA-mutant breast cancer, though fewer studies have been conducted in TNBC. In the BELLE-4 trial, patients with locally advanced or metastatic HER2-negative breast cancer were randomized to buparlisib or placebo in combination with paclitaxel as first-line therapy (43). Stratification was performed according to PI3K pathway activation, defined as PIK3CA mutation (detected by Sanger sequencing in exons 1, 7, 9, or 20) and/or low PTEN expression (1+ in ≤10% tumor cells). Approximately 25% of all enrolled patients (99/416) had hormone-receptor–negative disease (i.e., TNBC), and of these, 36 (36.4%) had tumors considered to be PI3K-pathway activated. The addition of buparlisib to paclitaxel failed to demonstrate a significant improvement in PFS in the overall population or in those with PI3K-activated tumors. In patients with TNBC, there was a trend toward shorter median PFS with buparlisib compared with placebo (5.5 vs. 9.3 months, respectively).

Ipatasertib, a highly selective AKT inhibitor, was evaluated in the phase II randomized trial LOTUS in combination with paclitaxel as first-line metastatic treatment for unselected TNBC (44). Ipatasertib improved PFS in the intent-to-treat (ITT) population, and a similar trend was also noted in patients with TNBC-low tumors (IHC 0 in ≥50% tumor cells). In a prespecified analysis in patients with PIK3CA/ AKT/PTEN-altered tumors (presence of activating PIK3CA/ AKT1 mutations or PTEN-inactivating alterations using targeted NGS), median PFS with ipatasertib plus paclitaxel was...
Table 2. Efficacy of genomic-based targeted therapies in clinical trials in TNBC

<table>
<thead>
<tr>
<th>Pathway</th>
<th>Drug</th>
<th>Mechanism</th>
<th>Patient population</th>
<th>Trial design (total N patients)</th>
<th>Intervention</th>
<th>Exploratory biomarker</th>
<th>Efficacy</th>
<th>Clinical trials.gov identifier</th>
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<tr>
<td>PI3K/ AKT/ mTOR</td>
<td>Buparlisib</td>
<td>PI3K inhibitor</td>
<td>Locally advanced/metastatic HER2-negative</td>
<td>Randomized phase II (n = 416; ref. 43)</td>
<td>Buparlisib + paclitaxel vs. placebo + paclitaxel</td>
<td>Stratification by PI3K pathway activation</td>
<td>PFS (full population): 8.0 vs. 9.2 (HR, 1.18; 95% CI, 0.82–1.68)</td>
<td>NCT01572727</td>
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<td>PFS (PI3K-activated): 9.1 vs. 9.2 (HR, 1.17; 95% CI, 0.63–2.17)</td>
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<td>PFS (TNBC): 5.5 vs. 9.3 (HR, 1.86; 95% CI, 0.91–3.79)</td>
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<td>Ipatasertib</td>
<td>AKT inhibitor</td>
<td>Locally advanced/metastatic TNBC</td>
<td>Randomized phase II (n = 124; ref. 44)</td>
<td>Ipatasertib + paclitaxel vs. placebo + paclitaxel</td>
<td>Stratification by tumor PTEN status</td>
<td>PFS (intent-to-treat): 6.2 vs. 4.9 (HR, 0.60; 95% CI, 0.37–0.98; P = 0.037)</td>
<td>NCT02162719</td>
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<td>PFS (PTEN-low): 6.2 vs. 3.7 (HR, 0.59; 95% CI, 0.26–1.32; P = 0.18)</td>
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<td>PFS (PI3KCA/AKT1/PTEN-altered): 9.0 vs. 4.9 (HR, 0.44; 95% CI, 0.20–0.99; P = 0.041)</td>
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<tr>
<td>MK2206</td>
<td></td>
<td>AKT inhibitor</td>
<td>Neoadjuvant stage II–III breast cancer (any subtype)</td>
<td>Randomized phase II (n = 149; ref. 45)</td>
<td>Paclitaxel ± MK2206 (followed by AC)</td>
<td>NA</td>
<td>pCR (all): 35.2 vs. 21.1 (P = 0.01)</td>
<td>NCT0142379</td>
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<td></td>
<td>pCR (TNBC): 40.2 vs. 22.4</td>
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<tr>
<td>Temsirolimus, everolimus</td>
<td></td>
<td>mTORC1 inhibitor</td>
<td>Metastatic metaplastic TNBC</td>
<td>Phase I dose expansion (n = 52; ref. 46)</td>
<td>Liposomal doxorubicin + bevacizumab + temsirolimus or everolimus</td>
<td>Exploratory analysis by PI3K pathway activation</td>
<td>ORR (all): 21 (95% CI, 11–35)</td>
<td>NCT00761644</td>
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<td>ORR (PI3K-activated): 31 (95% CI, 16–50)</td>
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<tr>
<td>Everolimus</td>
<td></td>
<td>mTORC1 inhibitor</td>
<td>Neoadjuvant stage II–III TNBC</td>
<td>Randomized phase II (n = 145; ref. 47)</td>
<td>Cisplatin + paclitaxel + everolimus vs. cisplatin + paclitaxel + placebo</td>
<td>Exploratory analysis of mutated genes, TNBC subtype, Ki67, AR, and TILs</td>
<td>pCR (all): 36 vs. 48 (P = 0.41)</td>
<td>NCT00930930</td>
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<tr>
<td><strong>EGFR</strong></td>
<td>Panitumumab</td>
<td>EGFR monoclonal antibody</td>
<td>Locally advanced/metastatic TNBC</td>
<td>Nonrandomized phase II (n = 71; ref. 49)</td>
<td>Panitumumab + carboplatin + gemcitabine</td>
<td>EGFR amp, p53 loss, PTEN loss, PIK3CA mut</td>
<td>PFS (all): 4.4 (95% CI, 3.2–5.5)</td>
<td>NCT00894504</td>
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<tr>
<td>Cetuximab</td>
<td>EGFR monoclonal antibody</td>
<td>Neoadjuvant stage II–IIIA TNBC</td>
<td>Nonrandomized phase II (n = 28; ref. 50)</td>
<td>Cetuximab + docetaxel</td>
<td>EGFR, Ki67, cytokeratins, CD8/FOXP3</td>
<td>pCR (intent-to-treat): 25 (95% CI 9–41)</td>
<td>EFS (TNBC): 4.6 vs. 4.8 (HR: 1.25; 95% CI, 0.85–1.83)</td>
<td>NCT00600249</td>
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<tr>
<td>Lapatinib</td>
<td>EGFR/HER2 inhibitor</td>
<td>Locally advanced/metastatic HER2-negative</td>
<td>Randomized phase III (n = 580; refs. 51, 149)</td>
<td>Lapatinib + paclitaxel vs. placebo + paclitaxel</td>
<td>EGFR</td>
<td>EFS (TNBC EGFR+): 4.2 vs. 4.9</td>
<td>EFS (TNBC EGFR−): 5.2 vs. 4.3</td>
<td>NCT0075270</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th><strong>RAS/MAPK</strong></th>
<th>Cobimetinib</th>
<th>MEK1/2 inhibitor</th>
<th>Locally advanced/metastatic TNBC</th>
<th>Open-label safety run-in (n = 16), randomized phase II (n = 90; ref. 55)</th>
<th>Cobimetinib + paclitaxel vs. placebo + paclitaxel</th>
<th>TNBC subtype, genetic alterations, PD-L1 expression</th>
<th>PFS (intent-to-treat): 5.5 vs. 3.8 (HR: 0.73; 95% CI, 0.43–1.24; P = 0.25)</th>
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<tbody>
<tr>
<td><strong>JAK/STAT</strong></td>
<td>Ruxolitinib</td>
<td>JAK1/2 inhibitor</td>
<td>Metastatic TNBC or IBC of any subtype</td>
<td>Nonrandomized phase II (n = 21; ref. 66)</td>
<td>Ruxolitinib</td>
<td>JAK2 amplification, pSTAT3</td>
<td>PFS (all): 1.2 (95% CI, 0.97–1.84)</td>
<td>NCT01562873</td>
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<tr>
<td><strong>NOTCH</strong></td>
<td>PF-03084014</td>
<td>Gamma-secretase inhibitor</td>
<td>Metastatic HER2-negative breast cancer</td>
<td>Phase I dose-finding/dose expansion (n = 29; ref. 67)</td>
<td>PF-03084014 + docetaxel</td>
<td>NA</td>
<td>ORR: 16 (95% CI 4.5–36.1)</td>
<td>NCT01876251</td>
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</tbody>
</table>

**NOTE:** Main efficacy analyses of biomarker-selected subgroups of interest are highlighted. HR, 95% CI, and P values are included when available.
Abbreviations: TNBC, triple-negative breast cancer; PFS, progression free survival (months); pCR, pathologic complete response (%); ORR, objective response rate (%); IBC, inflammatory breast cancer; AR, androgen receptor; AC, adriamycin/cyclophosphamide; HR, hazard ratio; CI, confidence interval; N, number; NA, data not available; NR, not reached, amp, amplification; EFS, event-free survival (months); TIL, tumor-infiltrating lymphocyte.
9 months versus 4.9 months in the placebo plus paclitaxel group, suggesting that the pathway may drive oncogenesis in a subset of patients with TNBC and providing the rationale for the ongoing randomized phase III IPATunity130 trial assessing the combination in preselected patients with activation of the PI3K pathway (NCT03337724). In addition, results from I-SPY 2, an adaptive-design trial testing novel agents in the neoadjuvant setting, showed an improvement in pCR with the addition of an allosteric AKT inhibitor, MK-2206, to standard chemotherapy in TNBC (40.2% vs. 22.4% in the control group), with a predicted 75.9% probability of success in a phase III trial (45).

Considering the higher prevalence of PI3K pathway aberrations in mesenchymal TNBC, of which 10% to 30% are metastatic, a phase I study was conducted in this histologic subgroup to evaluate the combination of mTOR inhibition (temsirolimus or everolimus) with liposomal doxorubicin and bevacizumab (46). Responses were limited to patients with NGS aberrations in PIK3CA, AKT, or PTEN. In the neoadjuvant setting, the addition of everolimus to cisplatin and paclitaxel did not increase pCR in molecularly unselected TNBC, and exploratory analyses showed that those who achieved pCR were not enriched for mutations in the PI3K/AKT/mTOR pathway (47).

Although alterations in genes encoding components of the RAS–MAPK pathway, such as KRAS, HRAS, BRAF, and MEK1/2, are not observed as frequently in treatment-naive TNBC as in other cancer types, EGFR is highly expressed in TNBC and can lead to upregulation of RAS–MAPK signaling (48). Across phase II and III trials, EGFR overexpression has not selected patients with TNBC who are more likely to derive benefit from EGFR-targeting monoclonal antibodies (e.g., cetuximab and panitumumab) or tyrosine kinase inhibitors (e.g., lapatinib; refs. 49–52). Synergistic effects of combined RAF and MEK inhibition have been observed in MDA-MB-231 and MDA-MB-468 TNBC cell lines (53), likely due to the presence of an activating mutation in KRAS, EGFR, or PIK3CA, AKT, PTEN, or PIK3R1 (54, 55). Activation of JAK2/STAT5 has been observed in TNBC cell lines, regardless of the PI3K/AKT/mTOR pathway (47).

With increasing understanding of the PI3K/AKT/mTOR pathway, the development of drugs targeting the pathway has accelerated. Inhibitors of AKT (e.g., perifosine; refs. 56, 57) and mTOR (e.g., everolimus, temsirolimus, and ridaforolimus; refs. 58, 59) have been investigated in preclinical models. Preclinical data have suggested that cotargeting both pathways (65). In addition, amplifications at the JAK2 locus (9p24) have been detected at a higher frequency in post-neoadjuvant TNBC samples compared with basal-like untreated tumors in TCGA, suggesting possible clonal selection after acquired chemotherapy resistance (39, 66). Selective inhibition of JAK2 with NVP-BSK-805 (20-fold selectivity of JAK2 over JAK1), administered with paclitaxel, significantly reduced pSTAT3 levels and tumor volume in vivo and in vitro compared with paclitaxel alone (66). In contrast, this effect was not observed with ruxolitinib (oral JAK1 and JAK2 inhibitor, with more limited activity against JAK1 and JAK2/STAT3) plus chemotherapy in JAK2-amplified TNBC cell lines. In a phase II trial in patients with metastatic TNBC, despite on-target inhibition and decreased pSTAT3 after two cycles of treatment, no responses were observed with single-agent ruxolitinib (67).

The NOTCH signaling pathway has been implicated in the differentiation and survival of stem cell–like tumor cells and resistance to cytotoxic chemotherapy (68). Neutralizing antibodies targeting NOTCH1 significantly inhibit tumor growth in CD44+/CD24− cells and enhance the activity of docetaxel (69). This synergistic effect with taxane-based therapy is also seen with PF-03084014, a reversible selective gamma-secretase inhibitor that blocks NOTCH signaling, in patient-derived TNBC xenograft models (70). NOTCH receptor mutations and focal amplifications are enriched in the triple-negative subtype, with most mutations either clustering in the heterodimerization domain or causing disruption of the PEST-negative regulatory domain (71). These aberrations show evidence of pathway activation in TNBC and exhibit sensitivity to PF-03084014. In cell lines expressing NOTCH1 fusion alleles, gamma-secretase inhibition also downregulates expression of MYC and CCND1, two targets whose oncogenic role has been well established in murine breast cancer (39). The presence of MYC amplification in 40% of basal-like tumors (36) suggests that MEK inhibition may be an attractive strategy in this selected population. Recently reported results from COLET, a randomized trial evaluating the MEK1/2 inhibitor cobimetinib with paclitaxel versus placebo and paclitaxel as first-line treatment for advanced TNBC, showed a modest but not statistically significant increase in PFS (56). Selumetinib (MEK1/2 inhibitor) is also being tested in combination with vistusertib (mTORC1/2 inhibitor) in treatment-refractory solid tumors (NCT02583542). Although no objective responses were observed in the phase I trial, stable disease for >16 weeks was confirmed across tumor types, including TNBC (57).

As previously described, elevated expression of MYC has been identified across breast cancer types, with a strong association observed in triple-negative and basal-like tumors (58). Downregulation of MYC alone is insufficient to induce synthetic lethality, and several combinatorial approaches have been investigated in preclinical models (59, 60). Activation of the MYC pathway sensitizes TNBC cell lines to CDK inhibition, possibly by promoting cellular apoptosis through upregulation of BIM, a proapoptotic BCL2 family member (58). CDK inhibitors, such as dinaciclib, downregulate MYC, and a synergistic effect has been observed in combination with PARP inhibitors in MYC-driven TNBC cell lines, regardless of BRCA status (59). Other strategies focus on epigenetic modulation of gene transcription, such as inhibition and/or degradation of BET bromodomain proteins. BET inhibitors/degraders also induce downstream suppression of MYC and an apoptotic effect that is significantly enhanced when combined with small-molecule BCL-2 inhibitors (61, 62). Altogether, these studies encourage further clinical research targeting MYC and exploring BET inhibitors in TNBC, and several clinical trials are ongoing in this area.
Heterogeneity of Triple-Negative Breast Cancer

NOTCH-driven tumors (72). It is estimated that 13% of TNBC may be driven by these NOTCH-oncogenic alterations. In a phase Ib trial, 29 patients with molecularly unselected treatment-refractory HER2-negative breast cancer (TNBC; n = 26) were treated with PF-03084014 plus docetaxel. An ORR of 16% was confirmed among evaluable patients, and median PFS was 4.1 months in the expansion cohort (68).

As illustrated by the variable efficacy across clinical trials, the role that many of these genes play as potential oncogenic drivers in TNBC remains unclear. Many of these trials have not yielded clinically relevant improvements in outcomes. Although some of these studies show promising preliminary data for targeted therapies, many have yet to be explored either in larger, randomized studies or in populations enriched for molecular alterations. Also, up to 12% of TNBC carry low mutational burden and do not harbor mutations in known candidate driver or cytoskeletal genes (73), further highlighting the heterogeneity in the mutational landscape of TNBC and the need to improve our understanding of the functional implications of many of these alterations.

GERMLINE BRCA-ASSOCIATED TNBC

Cancers that lack functional BRCA1 or BRCA2 have a deficiency in homologous recombination (HR) repair of DNA double-strand breaks (DSB), leading to dependence on alternative mechanisms to repair these lesions, and genomic instability (74, 75). Drugs that generate DSBs, such as alkylating agents (e.g., platinum, mitomycin C) or PARP inhibitors, cause persistent DNA damage in HR-deficient cells and, consequently, induction of cell-cycle arrest and apoptosis (76, 77). Germline mutations in BRCA1 or BRCA2 (BRCA1/2) are present in approximately 10% of patients with TNBC, and confer sensitivity to these drugs (78). In the previously mentioned TNT trial, despite failure to show a significant difference in activity between treatments in the overall population (n = 376), in the 43 patients with deleterious BRCA1/2 germline mutations, carboplatin significantly improved ORR compared with docetaxel (68% vs. 33.3%, P = 0.03) and PFS (6.8 vs. 4.4 months, interaction P = 0.002; ref. 12). In the neoadjuvant setting, elevated pCR rates (61%–65%) have been observed with platinum agents in germline-associated TNBC, albeit BRCA-mutant patients in the GeparSixto trial obtained high pCR regardless of the addition of carboplatin (79, 80).

Recently, PARP inhibitors (e.g., olaparib and talazoparib) have been compared with standard nonplatinum chemotherapy in two phase III trials, OlympiAD and EMBRACA, respectively, in germline BRCA1-associated metastatic HER2-negative breast cancer (81, 82). Eligibility criteria included receipt of no more than two to three previous lines of chemotherapy for metastatic disease, and receipt of an anthracycline and a taxane whether in the neoadjuvant, adjuvant, or metastatic setting. Neoadjuvant or adjuvant platinum was allowed if the time that had elapsed since the last dose was 12 months in OlympiAD and 6 months in EMBRACA. Both trials enrolled a similar patient population, with some differences including the distribution of germline mutations (57% BRCA1 in OlympiAD; 54.5% BRCA2 in EMBRACA) and, concordantly, a slightly greater proportion of patients with hormone receptor–positive disease in EMBRACA (55.9%) than OlympiAD (50.3%). Results of both studies were positive, with improvements in ORR, PFS, and quality of life, favoring the PARP inhibitor. Compared with standard chemotherapy, a significant increase in median PFS was observed with olaparib (7 months vs. 4.2 months, HR 0.58; P < 0.001) and with talazoparib (8.6 months vs. 5.6 months, HR 0.54; P < 0.001). Safety profiles were also comparable across trials, and hematologic toxicity was the most common cause of dose modifications with PARP inhibition. An adjuvant trial (OlympiA, NCT02032823) in patients with germline BRCA-associated breast cancer is currently accruing. Of note, the reported response rates in the metastatic phase III trials of olaparib and talazoparib (59.9% and 62.6%, respectively) were similar to those previously reported with carboplatin, and platinum agents were not allowed in the chemotherapy control arm. At the present time, the comparative efficacy and optimal sequencing (given potentially overlapping resistance mechanisms) of PARP inhibitors versus platinum agents is unknown. In addition, whether PARP inhibitors may have activity in patients with other germline DNA-repair defects (e.g., PALB2), or in patients with acquired somatic BRCA1/2 deleterious mutations, is unknown but is being tested in an ongoing clinical trial (NCT03344965).

Multiple mechanisms underlie the development of primary and acquired resistance to both platinum agents and PARP inhibitors, many of which have also been well characterized in ovarian or prostate cancer. Molecular alterations leading to therapeutic resistance include, for example, small insertions/deletions that result in frameshift mutations and synthesis of truncated proteins (e.g., inherited founder mutation BRCA14510delAT, ref. 83); secondary BRCA reversion mutations that reinstate HR proficiency through restoration of the open reading frame and BRCA reexpression (84); exon 11 deletion splice variants that produce truncated, hypomorphic proteins (85); or point mutations in PARP1 that alter PARP trapping (86). In addition to genomic alterations, epigenetic changes such as loss of BRCA1 promoter hypermethylation via BRCA1 locus fusion rearrangements, with subsequent BRCA1 reexpression, have also been described after acquired resistance to DNA-damaging drugs, including platinum or olaparib (87).

Several strategies to exploit potential synthetic lethality in HR-deficient tumors are being explored across solid tumors, including clinical trials combining PARP inhibitors with PI3K/AKT inhibitors (NCT02208375), immune-checkpoint inhibition (NCT02657889), and HSP90 inhibitors (NCT02898207). HSP90 is a chaperone that assists in intracellular protein homeostasis by mediating protein folding and stabilization. HSP90 inhibitors block adequate protein folding, leaving the “client” protein (e.g., BRCA1) in the cytoplasm to be degraded by the proteasome. In vitro, HSP90 inhibition results in loss of BRCA1 expression and function and impaired DSB repair, sensitizing tumors to DNA-damaging agents (88). Stabilizers of G-quadruplex DNAs such as CX-5641 bind to G4 DNA structures, interfering with progression of DNA replication complexes and inducing single-strand breaks that require HR for repair; thus, in BRCA-deficient tumors, failure to repair DNA damage leads to lethality, including in taxane-resistant BRCA1/2-deficient
TNBC patient-derived xenograft models (89). Given its promising in vivo activity, CX-5461 is currently being explored in a phase I trial, with an expansion phase for unresectable breast cancer in patients with known BRCA1/2 or HRD germline aberrations (NCT02719977).

“BRCAAness” in Sporadic TNBC

Somatic mutations and epigenetic alterations that inactivate BRCA1/2 and other DNA-repair genes have been identified in sporadic cancers (90). Given that HR deficiency exposes specific therapeutic vulnerabilities, the detection of sporadic tumors with this so-called “BRCAAness” phenotype could have clinical implications. Most BRCA1-related tumors are basal-like (91), and there is a marked resemblance in phenotype and biology between sporadic basal-like tumors and BRCA-associated cancers (90). Despite these similarities, targeting the HR pathway in sporadic basal-like cancer has revealed conflicting data in the metastatic and neoadjuvant settings. High HRD score or basal phenotype (by PAM50 or IHC) did not predict greater benefit from carboplatin in settings. High HRD score or basal phenotype (by PAM50 revealed conflicting data in the metastatic and neoadjuvant settings. High HRD score or basal phenotype (by PAM50 or IHC) did not predict greater benefit from carboplatin in settings. High HRD score or basal phenotype (by PAM50 or IHC) did not predict greater benefit from carboplatin in settings.

In GeparSixto, the addition of carboplatin to paclitaxel/liposomal doxorubicin improved pCR in HR-deficient tumors (64.9% vs. 45.2%, P = 0.006), but not in HR-proficient tumors (40.7% vs. 20%, P = 0.146; ref. 94). Discrepancies across trials may be explained by significantly less methylated BRCA1/2 in metastases than in primary tumors, leading to potential loss of HR deficiency (96). Treatment exposure to alkylating agents commonly used in early-stage TNBC could drive clonal selection of HR-proficient cells less likely to respond to platinum in the metastatic setting. However, another explanation for these observed differences could be the robustness of the genomic metrics used to calculate HRD scores. With advances in sequencing technologies, an algorithm using whole-genome sequencing, also known as the HRDetect model, identified six mutational signatures present in germline BRCA1/2-mutated tumors that were then found to also predict HR deficiency in sporadic tumors in the Sanger data set (97). This aggregated BRCAAness score was independently associated with benefit from platinum-based chemotherapy after adjusting for germline BRCA status and treatment timing, although the relatively small sample size (33 patients with metastatic breast cancer treated with either carboplatin or cisplatin as a single agent or in combination regimens) and the retrospective nature of the study (clouding the ability to establish a causal relationship) are limitations to be considered (98). Direct comparisons of these different measures should be further evaluated in ongoing prospective trials in HR-deficient breast cancer.

Currently, we lack predictive biomarkers to guide the choice of chemotherapy in sporadic basal-like TNBC, which comprises the majority of TNBC. Beyond germline BRCA mutations and the recent approval of olaparib and talazoparib in these patients, much remains unknown about the BRCAAness features that may confer sensitivity to PARP inhibitors and DNA-damaging agents. Trials assessing these drugs are ongoing both in unselected and biomarker-selected populations (Table 3). In addition, preclinical data have demonstrated upregulation of PD-L1 expression after exposure to PARP inhibition in triple-negative MDA-MB-231 cells, with subsequent resensitization to a PARP inhibitor when combined with a PD-L1 antibody (99). Furthermore, the accumulation of cytosolic damaged DNA induced by PARP inhibition activates the STING pathway, which in turn increases the expression of type-I IFN signaling and immune cell infiltration, regardless of BRCA mutational status (100). Altogether, this has provided the rationale to explore the combination of niraparib, a PARP inhibitor, and pembrolizumab, a PD-1 inhibitor, in the phase II clinical trial TOPACIO. Results from the TNBC cohort showed promising activity with an ORR of 28% in the 46 evaluable patients, and durable responses irrespective of tumor BRCA status, PD-L1 status, or prior platinum exposure, although the highest ORR was observed in patients with tumor BRCA1 or BRCA2 mutations (60%; ref. 101). A randomized phase II trial comparing olaparib in combination with the PD-L1 inhibitor atezolizumab versus olaparib alone in patients with BRCA-associated metastatic TNBC is currently ongoing (NCT02849496).

Epigenetic Markers and Therapies in TNBC

Epigenetic alterations, including changes in DNA methylation of gene promoter regions and posttranslational modification of histone proteins, are a recognized hallmark of cancer. Approximately 60% to 80% of basal-like and claudin-low breast cancers have aberrant DNA hypermethylation (102). Compared with luminal and HER2-positive cancers, TNBC exhibits extensive Cpg methylation of the promoter regions of nine epigenetic biomarker genes (CDH1, CECAM6, CST6, GNA11, ESR1, MUC1, MYB, SCN11A, and TFF3). DNA hypermethylation–dependent silencing of these genes is associated with worse RFS across all molecular subtypes and stages, compared with breast cancers unmethylated for these genes (40% RFS at 70 and 30 months, respectively). A nonsignificant trend toward RFS disadvantage has also been described among basal-like and claudin-low tumors that have this 9-gene methylation signature (102). In addition, promoter hypomethylation of three breast cancer stem cell–related genes (CD44, CD133, and MSH1), which strongly correlates with positive IHC staining and thus gene activation, has been shown to predict triple-negative status (103). Differences in histone modifications are also associated with differences in the expression of breast cancer genes across subtypes, separating luminal tumors, enriched with H3K27me3-modified genes, from nonluminal tumors (TNBC/HER2-positive), enriched with H3K9ac-regulated genes (104).

Therapies targeting epigenetic modifications, such as inhibitors of DNA methyltransferases (DNMT; 5-azacitidine, decitabine) and histone deacetylases (HDAC; entinostat, vorinostat), have yielded disappointing results to date in TNBC. The combination of 5-azacitidine and entinostat did not achieve any responses among 13 women with...
### Table 3. Ongoing clinical trials in BRCA-mutant or BRCAness-associated TNBC

<table>
<thead>
<tr>
<th>Clinicaltrials.gov identifier</th>
<th>Title</th>
<th>BRCA status eligibility criteria</th>
<th>Phase</th>
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<tbody>
<tr>
<td>NCT03109080</td>
<td>A Phase I of Olaparib with Radiation Therapy in Patients with Inflammatory, Locoregionally Advanced or Metastatic TNBC or Patient with Operated TNBC with Residual Disease</td>
<td>BRCA mutation not required.</td>
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<tr>
<td>NCT03329937</td>
<td>An Open-Label, Single-Arm Pilot Study Evaluating the Antitumor Activity and Safety of Niraparib as Neoadjuvant Treatment in Localized, HER2-Negative, BRCA-Mutant Breast Cancer Patients</td>
<td>Deleterious or suspected deleterious BRCA1 or BRCA2 mutation (germline or somatic).</td>
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<tr>
<td>NCT02978495</td>
<td>Neoadjuvant Carboplatin in Triple-Negative Breast Cancer—A Prospective Phase II Study (NACATINE Trial)</td>
<td>BRCA mutation not required. Includes BRCA-mutant–specific cohorts.</td>
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<tr>
<td>NCT02789332</td>
<td>A Randomized Phase II Trial to Assess the Efficacy of Paclitaxel and Olaparib in Comparison with Paclitaxel/Carboplatin Followed by Epirubicin/Cyclophosphamide as Neoadjuvant Chemotherapy in Patients with HER2-Negative Early Breast Cancer and Homologous Recombination Deficiency</td>
<td>BRCA deleterious tumor or germline mutation and/or high HRD score.</td>
<td>II</td>
</tr>
<tr>
<td>NCT03150576</td>
<td>Randomized, Phase II/III, 3 Stage Trial to Evaluate the Safety and Efficacy of the Addition of Olaparib to Platinum-Based Neoadjuvant Chemotherapy in Breast Cancer Patients with TNBC and/or gBRCA</td>
<td>TNBC or germline BRCA mutation HER2-negative breast cancer.</td>
<td>II/III</td>
</tr>
<tr>
<td>NCT02032823</td>
<td>A Randomized, Double-Blind, Parallel Group, Placebo-Controlled Multicenter Phase III Study to Assess the Efficacy and Safety of Olaparib versus Placebo as Adjuvant Treatment in Patients with gBRCA1/2 Mutations and High-Risk HER2-Negative Primary Breast Cancer Who Have Completed Definitive Local Treatment and Neoadjuvant or Adjuvant Chemotherapy</td>
<td>Suspected deleterious or deleterious BRCA1 and/or BRCA2 germline mutation.</td>
<td>III</td>
</tr>
<tr>
<td>NCT02950064</td>
<td>Escalation Study of BTP-114 in Patients with Advanced Solid Tumors and BRCA or DNA-Repair Mutation</td>
<td>Deleterious germline or somatic BRCA mutation or DNA-repair mutation. Abnormal HRD tests are also allowed.</td>
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<tr>
<td>NCT00576654</td>
<td>A Phase I Dose-Escalation Study of Oral ABT-888 (NSC #737664) plus Intravenous Irinotecan (CPT-11, NSC#616348) Administered in Patients with Advanced Solid Tumors Olaparib Dose Escalation in Combination with High Dose Radiotherapy to the Breast and Regional Lymph Nodes</td>
<td>BRCA mutation not required. Includes BRCA-mutant–specific cohort.</td>
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<tr>
<td>NCT02227082</td>
<td>A Phase I Study of PARP Inhibitor Olaparib and HS599 Inhibitor AT13387 for Treatment of Advanced Solid Tumors with Expansion in Patients with Recurrent Epithelial Ovarian, Fallopian Tube, Peritoneal Cancer or Recurrent TNBC</td>
<td>BRCA mutation not required. Dose expansion excludes germline BRCA1 or BRCA2 mutations.</td>
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<tr>
<td>NCT03075462</td>
<td>An Open, Nonrandomized, Multicenter Phase I Study to Assess the Safety and Efficacy of Fluzoparib Given in Combination with Apatinib in Patients with Recurrent Ovarian Cancer or TNBC</td>
<td>BRCA mutation not required.</td>
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<tr>
<td>NCT03109080</td>
<td>A Phase I of Olaparib with Radiation Therapy in Patients with Inflammatory, Locoregionally Advanced or Metastatic TNBC or Patients with Operated TNBC with Residual Disease</td>
<td>BRCA mutation not required.</td>
<td>I</td>
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<tr>
<td>NCT03101280</td>
<td>A Phase IB Combination Study of Rucaparib (CO/338) and Atezolizumab (MPDL3280A) in Participants with Advanced Gynecologic Cancers and TNBC</td>
<td>Part 1: All comers; part 2: deleterious germline or somatic BRCA mutation, or wild-type tumor BRCA but high levels of LOH.</td>
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(continued)
advanced TNBC treated in a phase II study (105). No significant changes in gene expression in paired biopsies before and after 2 months of treatment were observed, possibly due to absent ER promoter DNA methylation at baseline. Novel approaches in epigenetic modulation include BET bromodomain inhibitors that bind to acetylated lysine residues in histones, displacing bromodomain proteins from chromatin and inhibiting transcriptional activity (106). BET inhibitors achieve potent suppression of tumor growth in TNBC cell lines characterized by more basal-like and claudin-low/ stem cell–like features (61). Several BET inhibitors are currently in early stages of clinical testing as single agents

Table 3. Ongoing clinical trials in BRCA-mutant or BRCAness-associated TNBC (Continued)
Heterogeneity of Triple-Negative Breast Cancer

or in combination with immunotherapy (NCT01587703, NCT02391480, and NCT02771137).

**IMMUNE SUBTYPES OF TNBC**

Increasing data suggest that the immune system is critical for disease outcome in TNBC. Analyses from neoadjuvant and adjuvant TNBC trials have shown that tumor-infiltrating lymphocytes (TIL), assessed by hematoxylin–eosin staining, are predictive of response to therapy and strongly associated with improved survival (107, 108). Stratification of TNBC based on quantitative TIL evaluation has distinguished immune “hot” (high-TIL) and “cold” (low-TIL) tumors, which also appear to correlate with response to immune-checkpoint inhibitors in the metastatic setting (109). Paired biopsies pre- and post-neoadjuvant therapy have shown that the immune microenvironment can be modulated by chemotherapy, converting tumors from “cold” to “hot,” and these cases with highly infiltrated residual TNBC have improved survival (110). Phenotypic TIL characterization has also provided further insight into the populations of immune cells (e.g., CD8+ T cells; elevated CD8+/FOXP3 ratio) that may be responsible for this positive effect (111). Elevated expression in TNBC of immune markers of tumor evasion PD-L1/PD-L1 has prompted clinical assessment of inhibitors of these checkpoints, with modest efficacy as monotherapy and encouraging results in combination with chemotherapy (Table 4; refs. 109, 112–118).

Recently, results from a large phase III trial (iMPassion130) that randomized patients in the first-line TNBC metastatic setting to receive nab-paclitaxel combined with either atezolizumab (PD-L1 inhibitor) or placebo were reported (118). Although the absolute difference in median PFS in the PD-L1-positive population (2.5 months) was not strikingly different from that seen in the ITT cohort (1.7 months), at a median follow-up of 12.9 months, a 9.5-month clinically meaningful improvement in median OS was noted in patients with PD-L1-positive tumors, in contrast to a 3.7-month difference in the ITT population (118). No PFS or OS differences were noted in the subset of patients with PD-L1-negative tumors (119). Several other randomized trials have completed accrual and are awaiting data maturity to report. Whether similar results may be achieved with chemotherapy plus immunotherapy in later lines is unknown at this time. Of note, increased ORR have been observed in patients with previously untreated metastatic TNBC with monotherapy PD-L1/PD-L1 inhibitors, suggesting that these agents may be more active in less heavily pretreated metastatic disease (120).

Efforts to identify patients with tumors that are more or less likely to benefit from immunotherapy-based approaches are ongoing. As evidenced in the iMPassion130 trial, not all patients with PD-L1 tumors (defined by the presence of ≥1% IHC staining on immune cells) respond to PD-L1 inhibition and, contrarily, there are patients who, despite negative PD-L1 staining, appear to derive benefit from treatment. Beyond IHC classifications, genetic alterations of immune-regulatory genes have also segregated TNBC into subgroups with different prognostic and possibly therapeutic implications. CD274 (encoding PD-L1) and PDCD1LG2 (encoding PD-L2) genes localize to the 9p24 locus, adjacent to JAK2, constituting the PDJ amplicon. Overexpression of PD-L1 is observed in 88% of tumors with amplifications in the 9p24/JAK2 locus, which are found at higher frequency in post-neoadjuvant residual TNBC (66). In TNBC, the PDJ amplicon identified a subset of patients at significantly greater risk of recurrence (121), and could be a potential biomarker for selection of high-risk patients who may benefit from PD-1/PD-L1 blockade. Activating mutations in the RAS/MAPK pathway, present in 15% of residual disease, correlated with reduced TIL, inhibition of MEK upregulated PD-L1 expression, synergizing with PD-1/PD-L1 antibodies in murine models (122). Furthermore, high tumor mutational burden has been associated with improved outcomes with PD-1 inhibition in other cancer types (123), and may represent an independent biomarker of response.

Transcriptomic analysis of tumor-associated stroma in TNBC has revealed the presence of four axes, each with differential expression of genes related to T-cell, B-cell, epithelial (E), and desmoplasmia (D) markers. The E-axis inversely correlated with LAR Lehmann subtype, and the D-axis was positively associated with MSL while also determining the prognostic value of the T-, B-, and E-axes (124). Furthermore, these axes strongly influenced the location of CD8+ TIL (125), which may affect antitumoral response to immune-checkpoint inhibitors. Similarly, when analyzing the tumor compartment, the presence of the immunomodulatory signature (associated with elevated lymphocytic infiltration and increased expression of immune-checkpoint regulators, e.g., PD-1/PD-L1; ref. 18), significantly differs across refined TNBC types, with the highest rates observed in BL1 (48%) and the lowest in M (0%; ref. 126). Whether transcriptomic profiling could be incorporated to routine clinical practice to help select patients with TNBC with a greater likelihood of responding to immune-checkpoint inhibitors, similar to the applicability of gene-expression assays (e.g., 21-gene recurrence score, oncootype) to predict chemotherapy benefit in ER-positive breast cancer (127), remains to be seen.

To date, one single marker has not been proven to effectively select patients who are more likely to respond to immunotherapies. Recently, the development of multiplexed imaging techniques has enabled analysis of the spatial distribution and interaction between tumor and immune cells, showing that in TNBC there is high intratumor topologic heterogeneity for the expression of PD-1 on cytotoxic CD8+ and helper CD4+ T cells (128). Tumors with immune cells that are spatially separated from tumor cells, also defined as compartmentalized (as opposed to mixed immune cells with tumor cells), predominantly express PD-1 on CD4+ T cells and are independently associated with improved survival. Given the complexity of these interactions, integration of comprehensive omics analyses of samples with detailed clinical data annotation will be needed to better understand how the relationship between the tumor and its microenvironment affects response to treatment.

**EVOLUTIONARY PATHS OF TNBC**

Analyses of paired primary and metastatic TNBC samples are also needed to better understand the drivers of disease progression. Clonal frequencies vary significantly across TNBC at the time of diagnosis, suggesting their occurrence at different stages of tumorigenesis (73). There are limited sequencing data in metastatic triple-negative tumors, and
Table 4. Results of PD-1/PD-L1 inhibition in advanced TNBC

<table>
<thead>
<tr>
<th>Tumor characteristics</th>
<th>Single-agent immunotherapy</th>
<th>Combination with chemotherapy</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Anti–PD-1</td>
<td>Anti–PD-L1</td>
</tr>
<tr>
<td></td>
<td>Pembrolizumab in PD-L1−</td>
<td>Pembrolizumab in metastatic TNBC (KEYNOTE-086; ref. 112, 113)</td>
</tr>
<tr>
<td>Tumor characteristics</td>
<td>Defnition of PD-L1 positivity</td>
<td>≥1% TC or any staining in stroma</td>
</tr>
<tr>
<td></td>
<td>PD-L1 status inclusion criteria</td>
<td>Positive</td>
</tr>
<tr>
<td></td>
<td>Frequency of PD-L1 positivity among evaluable cases (%)</td>
<td>65/111 (58.6)</td>
</tr>
<tr>
<td>Patient characteristics</td>
<td>Total number of patients enrolled</td>
<td>32</td>
</tr>
<tr>
<td></td>
<td>Total number of patients included in efficacy analysis</td>
<td>27</td>
</tr>
<tr>
<td></td>
<td>Median prior lines of therapy in metastatic setting (range)</td>
<td>2 (0–9)</td>
</tr>
</tbody>
</table>
### Efficacy

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Arm 1</th>
<th>Arm 2</th>
<th>Overall</th>
<th>ITT: Arm 1 vs. Arm 2</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>ORR, %</strong></td>
<td>18.5</td>
<td>18.5</td>
<td>9.8</td>
<td>5.2</td>
</tr>
<tr>
<td><strong>First line</strong></td>
<td>4.7</td>
<td>4.8</td>
<td>5.2</td>
<td>2.0</td>
</tr>
<tr>
<td><strong>Second line</strong></td>
<td>12.7</td>
<td>22.2</td>
<td>22.2</td>
<td>9.8</td>
</tr>
<tr>
<td><strong>Third/+ line</strong></td>
<td>NA</td>
<td>31</td>
<td>NA</td>
<td>31</td>
</tr>
<tr>
<td><strong>Overall</strong></td>
<td>26.4</td>
<td>36.8</td>
<td>37.5</td>
<td>40</td>
</tr>
<tr>
<td><strong>First line</strong></td>
<td>29.2</td>
<td>36.8</td>
<td>37.5</td>
<td>40</td>
</tr>
<tr>
<td><strong>Second line</strong></td>
<td>22.6</td>
<td>1.5</td>
<td>1.5</td>
<td>4.2</td>
</tr>
<tr>
<td><strong>Third/+ line</strong></td>
<td>7</td>
<td></td>
<td>7.7</td>
<td>40</td>
</tr>
<tr>
<td><strong>ORR in PD-L1+ cohort, %</strong></td>
<td>18.5</td>
<td>18.5</td>
<td>18.5</td>
<td>18.5</td>
</tr>
<tr>
<td><strong>CBR, %</strong></td>
<td>25.9</td>
<td>25.9</td>
<td>25.9</td>
<td>25.9</td>
</tr>
<tr>
<td><strong>Median PFS, mo.</strong></td>
<td>1.9</td>
<td>2.1</td>
<td>1.9</td>
<td>2.1</td>
</tr>
<tr>
<td><strong>(95% CI)</strong></td>
<td>(1.7–5.5)</td>
<td>(2.0–2.3)</td>
<td>(1.4–1.7)</td>
<td>(4.1–6.1)</td>
</tr>
<tr>
<td><strong>Median OS, mo.</strong></td>
<td>11.2</td>
<td>8.9</td>
<td>9.3</td>
<td>9.2</td>
</tr>
<tr>
<td><strong>(95% CI)</strong></td>
<td>(7.2–11.2)</td>
<td>(7.0–12.6)</td>
<td>(7.0–12.6)</td>
<td>(4.3–NE)</td>
</tr>
</tbody>
</table>

**NOTE:** Response rates are per RECIST 1.1 criteria.

**Abbreviations:** TNBC, triple-negative breast cancer; TC, tumor cells; IC, immune cells; ORR, objective response rate; CBR, clinical benefit rate (defined as complete response, partial response, or stable disease for ≥24 weeks); mo., months; NR, not reached; NE, not estimable; NA, not available; PFS, progression-free survival; OS, overall survival; CI, confidence interval; ITT, intent-to-treat population; HR, hazard ratio.

*DCR is defined as confirmed complete, partial response, or stable disease as best response.

*Number of patients considered objective response–evaluable.

*In the overall population, the median number of prior lines of therapy in any setting was 4 (range, 1–10). In the TNBC cohort, 50% had received ≥2 prior lines of therapy for metastatic disease.

*According to PD-L1 positivity in IC.
much remains unknown about the differences in the molecular landscape of TNBC over its natural history. Multiclonal seeding from different cell populations in the primary tumor to the metastasis has been reported in two cases of basal-like TNBC, where, in addition, most putative driver mutations were shared, rather than acquired, between primary and metastatic lesions (129, 130). Also, most TNBC primary tumors and metastases are polyclonal, with overlapping clones, suggesting that polyclonal metastasis is common in TNBC. Phylogenetic analysis has the potential to distinguish local recurrences from second primary tumors and to help determine the origin of a metastatic lesion in a patient with a history of independent primary tumors (131). Given the differences in management of primary and recurrent tumors, sequencing of longitudinal samples could affect treatment decisions.

Receptor status, according to IHC, and also intrinsic subtype, can change at time of recurrence (132), but the clinical relevance of molecular phenotype switch remains unclear, and IHC subtypes largely drive current treatment decisions in breast cancer. Loss of ER and PR expression occurs in approximately 10% to 12% of asynchronous recurrences, inducing a switch to TNBC in the metastasis (133), and has been associated with worse survival compared with cases with concordant hormone receptor-positive-recurrent positive (134). To date, we do not fully understand the mechanisms that cause this conversion, and whether there are special considerations that should be made when treating this patient population. Of note, there are also breast tumors that express low levels (1%-9%) of ER and PR, and it remains unclear whether these cases derive significant benefit from endocrine therapy (135). Retrospective studies have shown that almost half of tumors with 1% to 9% ER staining are basal-like (136), suggesting that we should consider these tumors similar to TNBC and apply treatment algorithms, including enrollment onto clinical trials, for TNBC in these patients.

The extent of residual disease post–neoadjuvant chemotherapy, quantified per residual cancer burden index, is a well-established risk factor for recurrence (137). Residual disease has been used as a marker to select patients for escalation of adjuvant therapy, particularly in TNBC, based on the significant absolute improvement observed in patients treated with versus without capecitabine in terms of 3-year DFS (69.8% vs. 56.1%, respectively; HR, 0.58; 95% CI, 0.39–0.87) and OS (78.8% vs. 70.3%, respectively; HR, 0.52; 95% CI, 0.30–0.90; ref. 138). However, not all patients with residual disease will recur. Distinguishing between the molecular mechanisms of chemoresistance and those that drive the development of metastatic disease remains a challenge. Intratumor genetic heterogeneity has been widely described in TNBC and may be associated with a decreased likelihood of achieving a pCR (139, 140). Bulk exome and single-cell sequencing in a small number of pre- and post-neoadjuvant therapy samples suggest the occurrence of adaptive clonal extinction or persistence and acquired transcriptional reprogramming as potential models of chemoresistance (140). Other single-cell resolution studies support the hypothesis that most mutation and copy-number events occur in early stages of tumor evolution, rather than develop gradually over time, implying punctuated evolution (141). Validation of these findings in larger sets of tumors with associated long-term outcome data is key to understand the impact of genomic and phenotypic evolution of triple-negative cell.

CONCLUSIONS

In summary, TNBC is composed of a broad spectrum of biologically distinct subtypes with overlapping alterations. Despite advances in tumor characterization, separately, each classification has not yet translated to specific treatments or choices of treatments, with the exception of PARP inhibitors or platinum agents in germline BRCA1/2 carriers, and potentially in the near future immune-checkpoint inhibition in tumors with PD-L1–positive immune cells. Comprehensive integrated analysis of data generated from different “omics” technologies may provide more insight into the etiology, evolution of TNBC, and, possibly, prevention and new treatment strategies. Nonetheless, as the volume of information exponentially increases, identifying alterations that are critical for tumor growth and survival continues to be a challenge. In addition, the utility of these profiles is largely limited by genetic and epigenetic heterogeneity within the tumor. There have been several large-scale efforts to find new targets, including shRNA/CRISPR screens (64, 142–144). Using loss-of-function RNAs-based screens across more than 500 cancer cell lines, biocomputational algorithms have been developed to help predict cancer dependencies (143), and novel potently selective inhibitors, as single agents or in combination, will be needed to effectively block these targets (61, 145, 146). Similarly to cell lines and organoids, patient-derived xenografts enable high-throughput drug screening, but with the potential advantages of analyzing tumor growth metrics and characterizing drug response in models that retain the histopathologic features and intratumor and intratumor genomic heterogeneity of the explanted tumor (147). Given the complexity of these techniques and sample size of individual cohorts, institutional collaborations should be forged to create biobanks that will provide a platform to help answer questions of interest in specific subsets of patients with TNBC.

Most trials to date have been performed in unselected TNBC, hoping to find a signal of efficacy in subgroup analyses. Prospective validation of biomarker-driven approaches has been widely considered a necessary step for approval of targeted therapies over the past years. Only recently were results published from the first trial in TNBC to prospectively stratify patients by the presence of a tumor gene signature (148). In this neoadjuvant study, patients were randomized to receive paclitaxel with or without LCL161, a small-molecule antagonist of inhibitor of apoptosis proteins. LCL161 induces tumor necrosis factor (TNF)–mediated apoptosis, and preclinical work identified a three-gene signature (elevated TNFα, elevated RIPK1, and reduced STK39) that was associated with sensitivity to LCL161. In patients with signature-positive tumors, the pCR rate was higher in the combination versus the control arm (38.2% vs. 17.2%, respectively), as opposed to lower pCR in those who were negative for the signature (5.6% vs. 16.4%, respectively), albeit with significant toxicity that led to treatment discontinuation in almost one fifth of patients treated with LCL161 and paclitaxel (148). Of the total of 312 patients who signed consent for molecular prescreening for this trial, 207 had a valid signature score...
and were treated on study [of which 63 (30.4%) were found to be positive for the signature]. Enrollment was completed in approximately 25 months but required participation of 47 international sites across 11 countries. Inability to ship samples for testing (4.2%) and assay failure (7.1%) were among the reasons for exclusion of patients, highlighting the challenges of prospectively implementing molecular testing in clinical trials, including those evaluating biomarkers with a prevalence as high as the 30% rate observed in this trial.

Another limitation of conducting single oncogene-driven clinical trials is that there are complex interactions and overlap between different genomic alterations (e.g., comparable prognosis between PI3K-activated, TP53 wild-type TNBC, and ER-positive breast cancer) with consequences that are not clearly understood to date nor taken into consideration in study designs. As the field of genomics in TNBC evolves and new insights are gained, these factors may need to be incorporated into trial designs, particularly when post hoc stratification by various forms of analysis may be needed to interpret and demonstrate subgroup effects.

As NGS, immune-profiling, and other technologies become widely available, biomarker-selected basket trials across multiple cancer types are of particular importance to evaluate the efficacy of matched targeted therapies. Considering the multiple molecular hypotheses for treatment, dynamic biomarker-adjusted platforms, such as WSG-ADAPT or I-SPY, aim to improve the efficiency of early drug development by predicting the probability of success in phase III clinical trials (45, 149). However, given smaller and smaller potential subsets of interest, the success of biomarker-enriched designs will increasingly depend on more effective strategies to ensure that a larger pool of potentially eligible patients have the opportunity to be offered participation in such trials. Furthermore, given the evident heterogeneity in the molecular landscape of TNBC and current efforts to integrate omics data to better understand the underlying biological processes, the combinations of features in the tumor and its microenvironment that may be identified, and potentially targeted, seem endless. Conduction of randomized studies that require a large number of patients, aiming to test each individual hypothesis and demonstrate superiority of novel drugs to current standard-of-care regimens, is simply not feasible. As subsets of patients with rare, potentially actionable targets are identified, exploring multiple treatment options in these select populations is becoming more challenging, and this will likely translate into an increasing need to mindfully extrapolate results from subgroup analyses. Optimization of trial designs, including umbrella trials in TNBC (in which patients are assigned to an intervention based on genomic and/or immune profiling of the tumor at baseline), “pick-the-winner” strategies (with smaller sample sizes), and incorporation of comprehensive fresh biospecimen collection for correlative substudies, may provide proof-of-concept to help select therapies that are more likely to succeed in larger trials.

To overcome the challenges of limited, single-institution studies, multiple genomic data sharing initiatives such as Project GENIE (American Association for Cancer Research), Genomic Data Commons (National Cancer Institute), or eBioPortal have been developed as large repositories of sequencing data. Despite these efforts, one of the major limitations of these large-scale studies is the lack of detailed clinical annotation, making it difficult to answer specific questions such as the association between genomic features and prior exposure to therapy, changes in receptor subtype over time (i.e., due to absent ER, PR, and HER2 status at different time points) or clinical outcomes (e.g., response and survival). Another limitation is the heterogeneity in the utilization of exome/gene versus targeted panel sequencing across cancer centers, which limits the ability to perform in-depth analyses to genes that are common to all panels. In coming years, we anticipate standardization of clinical sequencing across institutions and implementation of machine-learning tools that will help extract clinical data from electronic medical records, facilitating a seamless integration of genomic and clinical information in both private and public data sets.

Furthermore, to address the need for advances in drug development and biomarker discovery in TNBC, the elaboration of prospective, large-scale, longitudinal multicenter cohort studies in TNBC that have the ability to capture a patient’s clinical course and collect fresh–frozen tissue, blood, and other biospecimens over a longer time frame, over multiple treatments, regardless of trial participation, and across a larger number of patients has the potential to vastly improve our knowledge of the dynamic changes in tumor biology, and the markers of response or resistance to treatment. These platforms may also be utilized to effectively communicate, offer, and expand clinical trial participation to patients across collaborating institutions in order to help answer clinically relevant questions in a timely manner and, ultimately, improve outcomes in patients diagnosed with TNBC.

Disclosure of Potential Conflicts of Interest

K. Polyak is a consultant/advisory board member for Mitra Biotech. N.U. Lin reports receiving commercial research grants from Pfizer, Genentech, Novartis, Array Biopharma, and Seattle Genetics and is a consultant/advisory board member for Shionogi, Genentech, Novartis, Seattle Genetics, and Puma. No potential conflicts of interest were disclosed by the other author.

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REFERENCES

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Insights into Molecular Classifications of Triple-Negative Breast Cancer: Improving Patient Selection for Treatment

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