TBCRC 019: A phase II trial of nanoparticle albumin-bound paclitaxel with or without the anti-death receptor 5 monoclonal antibody tigatuzumab in patients with triple negative breast cancer

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Presentation / Publications: Data from this work have been partially reported at the ASCO Annual Meeting in 2011 and 2013 (Abstract # TPS 128 and 1052 respectively).

Running Head: Phase 2 trial of Tigatuzumab/nab-PAC in Patients with Triple Negative Breast Cancer

Keywords: Tigatuzumab, nab-PAC, Monoclonal, Antibody, Triple negative, Breast Cancer

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Statement of Translational Relevance

The DR-5 tumor cell receptor is a promising target for an antibody-based therapy as it is expressed in solid tumors including breast cancer. Activation of DR-5 triggers apoptosis of tumor cells through activation of the extrinsic apoptotic pathway. Tigatuzumab is a novel agonistic humanized monoclonal antibody against DR5. In preclinical studies the antibody demonstrated strong in vitro (cell lines) and in vivo (xenograft models) activity against basal-like breast cancer that is enhanced by chemotherapy agents including albumin-bound paclitaxel (nab-PAC). Other types of breast cancer (hormone receptor and HER2 positive cancers) were resistant to Tigatuzumab alone or in combination with chemotherapy. Consequently, a clinical trial with this antibody in combination with nab-PAC in patients with triple negative breast cancer was conducted with signs of efficacy in a subset of patients. A single arm with nab-PAC was included as there was no prior prospective experience with this agent in this patient population.
Abstract

**Purpose:** Tigatuzumab (TIG), an agonistic anti-DR5 antibody, triggers apoptosis in DR5+ human tumor cells without crosslinking. TIG has strong in vitro/in vivo activity against basal-like breast cancer cells enhanced by chemotherapy agents. This study evaluates activity of TIG and chemotherapy in patients with metastatic triple negative breast cancer (TNBC).

**Experimental Design:** Randomized 2:1 phase II trial of albumin-bound paclitaxel (nab-PAC) + TIG in patients with TNBC stratified by prior chemotherapy. Patients received nab-PAC weekly x 3 + TIG every other week, every 28 days. Primary objective was within-arm objective response rate (ORR). Secondary objectives were safety, progression free survival (PFS), clinical benefit, and TIG immunogenicity. Metastatic research biopsies were required.

**Results:** Among 64 patients (60 treated; TIG/nab-PAC n=39 and nab-PAC n=21), there were 3 complete remissions (CRs), 8 partial remissions (PRs; 1 almost CR), 11 stable diseases (SDs) and 17 progressive diseases (PDs) in the TIG/nab-PAC arm (ORR=28%), and no CRs, 8 PRs, 4 SDs and 9 PDs in nab-PAC arm (ORR=38%). There was a numerical increase in CRs and several patients had prolonged PFS (1025+, 781, 672, 460, 334) in the TIG/nab-PAC arm. Grade 3 toxicities were 28% and 29% respectively with...
no grade 4-5. Exploratory analysis suggests an association of ROCK1 gene pathway activation with efficacy in the TIG/nab-PAC arm.

**Conclusions:** ORR and PFS were similar in both. Preclinical activity of TIG in basal-like breast cancer and prolonged PFS in few patients in the combination arm support further investigation of anti-DR-5 agents. ROCK pathway activation merits further evaluation.
**Introduction**

Triple-negative breast cancer (TNBC) is defined by the absence of estrogen and progesterone receptors (ER/PR), and HER-2 amplification; further sub-classification is being evaluated\(^1\). TNBC represents 15 to 20% of all breast cancers\(^2-5\) and is more frequent in younger patients, BRCA1 mutation carriers, and in specific ethnic groups such as African American women.\(^6, 7\) TNBC tumors are generally invasive ductal carcinomas and often have unfavorable features such as higher histologic grade, larger tumor size, and positive lymph nodes.\(^8\) The metastatic potential in TNBC is similar to that of other subtypes, but these tumors are associated with a shorter median time to relapse and death.\(^9, 10\) TNBC represents a significant clinical challenge as there are no targeted drugs available; however, chemotherapy remains the mainstay of treatment, but important limitations still need to be overcome in the next few years if any significant clinical strides are to be made.

Tumor necrosis factor-related apoptosis-inducing ligand (TRAIL), a member of the TNF superfamily of cytokines, is a type 2 membrane protein expressed in the majority of normal tissues and can undergo protease cleavage, resulting in a soluble form able to bind to TRAIL death receptors (DRs).\(^11\) TRAIL induces apoptosis of cancer cells in vitro and has potent tumor activity against tumor xenografts of various cancers in vivo via DRs.\(^11\) Although five receptors for TRAIL have been identified, only two (DR4 and DR5) are able to trigger apoptosis of tumor cells through activation of the extrinsic apoptotic pathway (caspase mediated).\(^11-14\) High expression of DR5 is frequently observed in
various human cancers including breast cancer. Our group has recently evaluated the phenotypic expression of DR5 in different subtypes of breast cancer; expression of DR5 was present in all triple negative ductal breast cancer tested, including primary and metastatic tumors (data not shown).

Tigatuzumab (TIG) is the humanized version of the agonistic anti-DR5 murine monoclonal antibody TRA-8. It is composed of the complementarity-determining region of the murine antibody and the variable region framework and constant regions of human immunoglobulin IgG-1 mAb58’CL. TIG is able to trigger apoptosis in DR5-positive human tumor cells without the aid of crosslinking. In preclinical studies, the antibody has demonstrated strong \textit{in vitro} and \textit{in vivo} activity against basal-like breast cancer cells that is enhanced by chemotherapy agents like paclitaxel and albumin-bound paclitaxel (nab-PAC).

A phase 1, dose-escalation study of TIG in patients with relapsed or refractory carcinomas was conducted to determine the maximal tolerated dose (MTD), pharmacokinetics, immunogenicity, and safety. Seventeen patients were enrolled in 4 cohorts (1, 2, 4 and 8 mg/kg). TIG was well tolerated with no infusion reactions or grade 3-4-5 toxicity; the MTD was not reached. Plasma half-life was 6–10 days, and no anti-TIG responses were detected. Seven patients had stable disease (SD), with the duration of response ranging from 81 to 798 days. Phase 2 studies in other solid tumors using TIG in combination with chemotherapy demonstrated the safety of the combination.
Thus, based on the preclinical data showing the remarkable sensitivity of basal-like breast cancer to TIG in combination with nab-PAC and the safety of TIG as single agent and in combination with chemotherapy, we conducted a randomized, phase II clinical trial, of nab-PAC with or without TIG in patients with TNBC.
Materials and Methods

Patients

Patients older than 18 years of age with histologically confirmed metastatic TNBC were enrolled. A tumor was considered triple negative if HER-2-neu was negative (0 or 1+ staining by IHC or gene amplification ratio < 2.0 by FISH), and the ER and PR were negative (<1% of the tumor cells by IHC). There was no restriction as to the number of prior chemotherapy regimens for metastatic disease but patients had to have prior exposure to anthracyclines and taxanes in the neoadjuvant or adjuvant settings. Patients with no prior chemotherapy for metastatic disease and patients who have received prior therapy with taxanes for metastatic disease (paclitaxel or docetaxel) were eligible. All patients had to have measurable disease per Response Evaluation Criteria in Solid Tumors (RECIST Version 1.1), an ECOG ≤ to 2, and adequate organ and bone marrow function (Supplemental Material). Patients previously treated with nab-PAC or with active central nervous involvement were excluded.

Study Design and Treatment Schedule

This study was a randomized (2:1) phase 2 multicenter trial of nab-PAC with or without TIG in patients with metastatic TNBC. The trial was conducted through the Translational Breast Cancer Research Consortium (TBCRC); 13 sites activated the study. A treatment cycle was defined as 4 weeks. Patients received intravenous nab-PAC on days 1, 8, and 15 (100 mg/m²) at 28-days interval with or without TIG intravenously on days 1 and 15.
of every cycle (10 mg/kg loading dose followed by 5 mg/kg every other week). Response to therapy was assessed every two cycles (every 8 weeks). Treatment continued without interruption in patients with a complete response (CR) or partial response (PR) or SD until progressive disease (PD) or unacceptable toxicity. Patients with tumor progression on the nab-PAC arm were allowed to rollover to the TIG/nab-PAC arm. All patients gave informed consent to participate in the study, which was approved by local Institutional Review Boards and conducted in accordance with the ethical principles of the Declaration of Helsinki, International Conference on Harmonization Guideline E6 for Good Clinical Practice and applicable local regulatory requirements.

**Study End Points**

The primary efficacy end point was objective response rate (ORR) based on RECIST 1.1 criteria. Secondary efficacy end points were progression free survival (PFS), duration of response, clinical benefit ratio (CBR) and safety of the combination. The ORR was defined as the proportion of patients who achieved best overall response of confirmed CRs and PRs. PFS was defined as the time from the date of initial treatment to the date of the first objective documentation of PD or death. The duration of response was defined as the time from the date of the first documentation of CR or PR to the date of the first documentation of PD. CBR for this protocol was defined as the percentage of patients who have achieved CR, and PR and SD for > 4 cycles. Treatment-emergent adverse events (TEAEs) were collected and reported from the time of the first dose administration of the study drugs to 30 days after the last dose administration. Toxicities were graded
according to National Cancer Institute Common Terminology Criteria for Adverse events (CTCAE) Version 3.0.

Human anti-human antibody measurements (HAHA) were conducted before the infusion of TIG, at 4 and 8 weeks after the TIG infusion and every 8 weeks thereafter during active treatment, and 3 months after the end of treatment. HAHA analysis was performed using a qualitative solid-phase assay as previously described.26

Biopsy of a reasonably accessible metastatic lesion (chest wall, breast, skin, subcutaneous, superficial lymph nodes, bones and liver metastases) was required for participation in the trial. Lung and brain metastasis were not considered reasonably accessible lesions. Biopsy samples were obtained using a 14-18 gauge core needle; at least two core biopsies were obtained and snap frozen individually and a third one for the preparation of paraffin-embedded blocks. Frozen tissues were used for high-throughput genomic analyses after macro-dissection and data related to treatment response is presented in this manuscript.

Circulating tumors cells (CTCs) were collected and the overall results are reported by Paoletti, et.al, in a companion manuscript.42

**Tumor sample processing**

De-identified fresh frozen tumor tissue biopsy specimens were obtained from the University of Alabama at Birmingham’s Comprehensive Cancer Center Tissue
Procurement Shared Facility. The specimens were macro-dissected by a board certified pathologist at the Tissue Procurement Shared Facility to enrich for tumor cell content and remove adjacent normal tissue. The dissected specimens were weighed, transferred to a 15 mL conical tube containing ceramic beads, and RLT Buffer (Qiagen) plus 1% BME was added so that the tube contained 35 uL of buffer for each milligram of tissue. The conical tubes containing tissue, ceramic beads and buffer were agitated in a MP Biomedicals FastPrep machine at 6.5 meters per second for 90 seconds to homogenize the tissue. The homogenized tissue was stored at -80°C. Total RNA was extracted from 350 uL of tissue homogenate (equivalent to 10 mg of tissue) using the Norgen Animal Tissue RNA Purification Kit (Norgen Biotek Corporation). Cell lysate was treated with Proteinase K before it was applied to the column and on-column DNase treatment was performed according to the manufacturer’s instructions. Total RNA was eluted from the columns and quantified using the Qubit RNA Assay Kit and the Qubit 2.0 fluorometer (Invitrogen). RNA-seq libraries for each sample were constructed from 250 ng total RNA using the polyA selection and transposase-based non-stranded library construction (Tn-RNA-seq) described previously. RNA-seq libraries were barcoded during PCR using Nextera barcoded primers according to the manufacturer (Epicentre). The RNA-seq libraries were quantified using the Qubit dsDNA HS Assay Kit and the Qubit 2.0 fluorometer (Invitrogen) and four barcoded libraries were pooled in equimolar quantities for sequencing. The pooled libraries were sequenced on an Illumina HiSeq 2000 sequencing machine using paired-end 50 bp reads and a 6 bp index read, and we obtained at least 50 million read pairs from each library. TopHat v1.4.1 was used with the options -r 100 -mate-std-dev 75 to align 50 million RNA-seq read pairs, and used...
GENCODE version 9\textsuperscript{30} as a transcript reference. Gene expression values (Fragments Per Kilobase of transcript per Million reads, FPKMs) were calculated for each GENCODE transcript using Cufflinks 1.3.0 with the -u option.\textsuperscript{31}

Statistics

There were no prior data on ORR of nab-PAC in this patient population although a trial of nab-PAC in patients with therapy resistant tumors had a 14\% ORR in the TNBC patient subgroup\textsuperscript{32}. Therefore, the sample size calculation was based on estimation of ORR. With an accrual of 40 patients to the TIG/nab-PAC regimen, the ORR estimation would have a standard error of less than 7.5\% if one assumes the ORR is between 20\%-35\%; the estimated two-sided 95\% confidence intervals (CI) would be 21.2\%-51.7\% for an ORR of 35\% with Blythe-Still-Casella Exact Method and 9.4\% - 34.4\% if the ORR were 20\%. In the single agent arm with 20 patients, the ORR would have a standard error of 8.9\%; two-sided 95\% CI would be 7.1\% - 41.1\% for an ORR of 20\% using the same method.

Patients were randomized in the trial as 2:1 ratio and stratified by patients’ prior chemotherapy. All randomly assigned patients were included in the intent-to-treat efficacy analysis and safety analysis. Descriptive analysis for patients demographic and clinical characteristics such as means, medians, and ranges were used to describe continuous variables. Frequency and proportion were used to describe categorical variables. Fisher’s exact test was used to examine two portions in 2 by 2 contingency
table. Survival distributions for PFS was estimated using the Kaplan-Meier method and were compared with long-rank tests stratified by stratification factors. Two-sided 95% confidence intervals for the median survival time were constructed using a nonparametric method.33

A modified Gehan’s two-stage design was used in the trial34 to minimize exposure to ineffective therapy; at least one patient in the first 11 patients enrolled per arm had to have a CR or PR in order to complete enrollment in that arm. A safety interim analysis was scheduled to be done after the first 6 patients enrolled in the TIG/nab-PAC arm (Supplemental Material).

RNA-seq Gene Expression Analysis of Tumor Biopsy Tissue: DESeq235 was used to analyze gene count data to identify genes whose expression was significantly associated with response to therapy. The DESeq2 nbinomLRT function was used to identify genes that were significantly differentially expressed between two classes: Class 1 contained patients who achieved CR or PR, Class 2 contained patients who had SD or PD. We also identified genes that were significantly associated with response criteria when response was represented as a quantitative variable ranging from CR (1) to PR (2) to SD (3) to PD (4). The significant genes (FDR< 0.05) were filtered to identify genes whose maximum FPKM expression value across samples was greater than or equal to 1.
Results

Patients

Sixty-four patients were enrolled; 42 in the TIG/nab-PAC arm and 22 in the nab-PAC arm (Table 1). All patients gave signed informed consent, and 60 patients received at least 1 cycle of therapy. In the TIG/nab-PAC arm the median age for the patients was 51 years (range, 32 to 72), 33% were African American, 33% had no prior chemotherapy in the metastatic setting, and the median number of prior therapy regimens was 2 (range, 0-5). The nab-PAC arm had similar characteristics; in those patients the median age was 51 (range 34-75), 27% were African American, 32% had no prior chemotherapy in the metastatic setting, and the median number of prior chemotherapy regimens was 1 (range, 0 to 4). All patients had an ECOG of ≤ 2.

Efficacy

Of the 42 patients in the TIG/nab-PAC arm, 39 received at least one course of therapy and were eligible for evaluation of response (3 patients had PD before initiation of therapy); of the 22 patients in the nab-PAC arm, 21 patients were treated and were eligible for evaluation of response (1 patient had PD before initiation of therapy). At least one PR was seen in the first 11 patients treated in each arm and accrual continued to completion. Eleven patients progressed before the first protocol-specified evaluation of response.
In the TIG/nab-PAC arm, there were 3 CRs, 8 PRs (1 patient had a near CR with 96% reduction in the index lesions) with an ORR of 28% (95% exact CI 14.9% to 45.0%). The median PFS for the patients enrolled in the TIG/nab-PAC arm was 2.8 months (95% CI 1.9-3.6) (Table 2 and Figure 1A) and 3.8 months in patients with objective response (95% CI 2.8-19.7). Sixteen of the 39 patients (41%) in the TIG/nab-PAC arm achieved clinical benefit. There were 5 patients in the TIG/nab-PAC arm with long PFS including 3 CR patients (1025+, 781, and 672 days), 1 near CR (460 days) and 1 SD (334 days). Four of the 11 patients that achieved CR or PR in the TIG/nab-PAC arm had progression in the brain but no systemic progression.

Although the study was not designed for statistical comparison of the two treatment arms, the control arm (single agent nab-PAC) had similar overall efficacy as combination therapy with an ORR of 38% (95% CI exact 18-61.1%), no CRs and 8 PRs. Clinical benefit was noted in 11 patients (52%) enrolled in the nab-PAC arm (Table 2). The median PFS for patients enrolled in the nab-PAC arm was 3.7 months (95% CI 2.3-5.7) (Figure 1A), and long term PFS occurred in 1 patient (1004+ days). Two additional patients had PFS for 224 and 220 days. Thus, proportionally more patients in the combination arm experienced prolonged clinical benefit (5 out of 39 [13%] versus 1 out of 21 [5%] patients). No objective responders in the nab-PAC arm had progression in the brain without progression of index lesions. Only 8 patients crossover to the TIG/nab-PAC after progression in the nab-PAC arm; no responses were seen in those patients.
Patient Demographics and Efficacy

We examined the effect of patient demographics and prior therapy on the whole patient population since outcomes were similar in the two arms (Table 3). Chemotherapy naïve patients had an increased ORR (53% [95% CI exact 31-76.3%] vs. 22% [95% CI exact 10.5-40.1%] respectively) and decreased PD rate (26 vs. 51% respectively). PFS was not significantly greater (3.6 vs. 2.5 months; Figure 1B) while the median duration of the response was 137 days (range, 84-1025+ days) and 174 days (range, 111-1004+ days) respectively. Among the 19 patients who were chemotherapy naïve in the metastatic setting, 53% had objective response, 68% had clinical benefit and PFS of 3.6 months (95% CI 2.8-5.6) compared with 22%, 34% and 2.5 months (95% CI 1.9-3.7) for those patients that received prior chemotherapy in the metastatic setting. We found no differences in efficacy for other factors including race (white vs. black), age (less than or greater than 50), tumor behavior (less than or greater than 2 years between primary tumor and relapse), or superficial extent (breast, soft tissue, lymph nodes) vs. systemic metastasis (liver, lung, bone).

Safety

Thirty nine patients in the TIG/nab-PAC arm and 21 in the nab-PAC arm received at least one cycle of therapy and were eligible for toxicity evaluation (Table 4). No adverse or serious adverse events (AEs/SAEs) related with the research agent were seen in the first 6 patients treated in the TIG/nab-PAC arm and accrual continued to completion.
Therapy in both arms was well tolerated; the majority of the AEs were grade 1-2 with very few grade 3 events and no grade 4/5 toxicity. There were no AEs or SAEs associated with TIG infusions. The most common AEs observed in at least 10% of all patients enrolled in the trial deemed by the investigators to be possibly related with the protocol therapy were fatigue (54%), alopecia (49%), peripheral sensory neuropathy (44%), anemia (41%), neutropenia (38%), nausea (23%), thrombocytopenia (10%), anorexia (10%), diarrhea (10%), and vomiting (10%). As expected, due to the use of nab-PAC, the most frequent grade 3 AEs were neutropenia (15%), fatigue (10%), anemia (2%) and peripheral sensory neuropathy (2%). The addition of TIG did not change the safety profile nab-PAC. The most frequent AE seen in the TIG/nab-PAC arm, excluding alopecia, was fatigue while the most frequently seen in the nab-PAC arm was peripheral sensory neuropathy.

Forty two SAEs were reported; 4 were classified as possibly related with the protocol therapy and 38 associated with PD. The 2 SAEs related in the TIG/nab-PAC arm were fever/neutropenia G4 and empyema/neutropenia G3; the 2 SAEs related in the nab-PAC arm were fever/neutropenia G2 and pulmonary thromboembolism. No deaths associated with the treatment agents were seen in the trial. None of the patients enrolled in the TIG/nab-PAC arm developed HAHA.

**Biopsies**
A successful biopsy was defined as one in which a patient had successful dual biopsies of any metastatic lesion (snap frozen), adequate tumor on macro-dissection to assure > 50% tumor cell nuclei and adequate DNA/RNA yield from the macro-dissected tissue. Of the 64 patients enrolled, 38 (59%) were successfully biopsied, 31 (48%) were judged adequate by macro-dissection and 28 (44%) had appropriate DNA/RNA yield for the study. Of the 28 samples, two were from patients who had PD prior to therapy, 20 received combination therapy (5 patients with CR/PR, 7 patients with SD and 8 patients with PD) and 6 received single agent nab-PAC (3 patients with PR and 3 with PD). Seventeen of the 28 patients had only a single tissue sample adequate for DNA/RNA analysis while 11/28 had multiple adequate DNA/RNA samples. The most common biopsy sites for tissue inadequacy were nodes and soft tissue. The reason for tissue macro-dissection failure was extensive necrosis in 50% and absence of tumor cells (benign tissue) in 50% of the specimens. This 40% yield of tissue analysis in treated patients limits the genomic analysis but the 28 metastatic tissues will be extremely valuable in studies relevant to metastatic TNBC.

RNA-seq\textsuperscript{28} was used to measure gene expression in the tumors. Each tumor was classified as belonging to one of the six Vanderbilt TNBC subtypes;\textsuperscript{36, 37} There was no significant association between subtypes and response to therapy. Expression of all genes was examined and seven were significantly associated with response in the patients enrolled in the TIG/nab-PAC (False Discovery Rate < 0.05): ACTA2, DNM3, FBXO32, IFFO2, LIMK2, MYLK, and ZNF469. All seven genes were expressed at a higher level in tumors from patients who responded to the combination therapy compared to those
who did not. Several of these genes are involved in apoptotic membrane blebbing through DR5/Casp-3/ROCK1 signaling (Figure 2 and Figure 3 in Supplemental Data). Activation of DR5 leads to activation of Caspase-3, which cleaves and activates ROCK1. ROCK1 phosphorylates and inhibits MLCP leading to unopposed MYLK phosphorylation of MLC, which catalyzes the generation of actin-myosin contractile force that causes blebbing. ROCK1 also phosphorylates and activates LIMK2 which leads to the accumulation and stabilization of actin filaments, such as those composed of ACTA2, involved in constriction of the cytoskeleton and apoptotic membrane blebbing. DNM3 is a member of the dynamin family that interacts with actin membrane processes and is responsible for constricting and releasing membrane vesicles. Thus, four of the seven genes significantly associated with response to TIG/nab-PAC are associated with the membrane blebbing process. This enrichment suggests that higher expression of this apoptotic pathway could be related to sensitivity to one or both of these drugs. Although the number of cases in the nab-PAC arm were very limited (6 patients), the expression of these seven genes was examined in tumors from those patients; these genes were not positively correlated with response to nab-PAC.
Discussion

This trial was undertaken based on the pre-clinical studies which indicated that basal like breast cancer cells were highly sensitive to anti-DR5, that the combination of an anti-DR5 monoclonal antibody and chemotherapy were quite effective in murine models of basal type breast cancer in vivo and that basal type breast tumor stem cells were killed by anti-DR5. Similar studies in hormone dependent and HER2 positive breast cancer demonstrated resistance to anti-DR5 therapy.

At the time of this protocol design, it was not feasible to use platinum compounds as the chemotherapy backbone of our study in view of the expanded access program that was available then for the combination of carboplatin, gemcitabine and iniparib for patients with newly diagnosed metastatic TNBC following a promising phase 2 randomized trial of chemotherapy with/without iniparib. In addition, there was no prior prospective experience with nab-PAC in this patient population although heavily pretreated TNBC patients appeared to have a 14% response rate in retrospective analysis of nab-PAC as single agent in patients with metastatic breast cancer. Thus, we designed a randomized Phase 2 to obtain a reasonable measure of efficacy with the combination arm (objective response rate in 40 patients with a standard error < 7.5%). In addition, we included a single agent nab-PAC arm as a frame of reference for this patient population.

The outcome of the trial was that the combination arm had an ORR of 28% and PFS of 2.8 months. The experience was similar in the concurrent single arm with ORR of 38%
and PFS of 3.7 months. This experience did not support moving forward with this current combination regimen in the same population of patients. Despite the negative overall trial findings, we did note that the combination arm included 3 CRs and 1 near CR while no CRs occurred in the single agent arm. In addition, proportionally more patients in the combination arm experienced prolonged clinical benefit (5 out of 39 [13%] versus 1 out of 21 [5%] patients). nab-PAC was associated with an unexpectedly high rate of objective response in patients with TNBC, reinforcing the need for a reference arm in our trial design; unfortunately, as with other agents evaluated in this patient population, responses were often not durable. A new anti-DR5 monoclonal antibody (DS8273 from Daiichi Sankyo) has shown better preclinical activity than TIG as a single agent or in combination with chemotherapy and is now being evaluated in a phase 1 trial (NCT02076451).

Metastatic TNBC is an aggressive disease as illustrated in our trial with 4 enrolled patients having progression prior to initiation of therapy and 26/60 (43%) of treated patients had progression prior to or at their initial evaluation (8 weeks). Patients with no prior therapy for metastatic disease experienced a higher ORR and clinical benefit rate.

Our experience with core needle biopsies for genomic studies is informative in designing future correlative studies within trials. First, trials should be designed for patients with accessible metastases and biopsies should be required (100% biopsies). Second, duplicate biopsies would increase the yield of appropriate tissue samples and third, incisional biopsies on superficial metastatic sites (chest wall, breast and lymphatic nodes) should be
considered. Also, standard for needle biopsies should be considered (e.g., ≥ 1.0 cm in length). CTCs were collected and the complete analysis of the data is presented in a companion manuscript by Paoletti et al. CTCs were detected in approximately one-third of the patients. Elevated CTCs at baseline and days 15 and 29 were prognostic, and reductions in CTC levels reflected response.

Finally, our genomic analysis (RNA seq) relating to therapeutic response was limited due to small numbers of patients tissues with 20 samples in the combination therapy arm and 6 samples in the single agent nab-PAC arm. In the combination arm, efficacy was significantly associated with elevated levels of seven genes including 4 of which participate in DR5 mediated ROCK1 activation of apoptosis associated membrane blebbing. This is an important and interesting observation; interpretation is tempered by limited patient samples.

In conclusion, the high degree of anti-DR5 sensitivity of basal-like breast cancer cell lines compared to other tumor cell lines and the prolonged PFS in a few patients in the TIG/nab-PAC suggest that DR5-mediated therapy deserves further investigation with novel, more efficacious, anti-DR5 agents.
References


**Table 1. Patient Demographics**

<table>
<thead>
<tr>
<th></th>
<th>TIG/nab-PAC (N=42)</th>
<th>nab-PAC (N=22)</th>
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<tbody>
<tr>
<td><strong>Race</strong></td>
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<tr>
<td>White</td>
<td>26 (63%)</td>
<td>16 (73%)</td>
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<td>Black</td>
<td>14 (33%)</td>
<td>6 (27%)</td>
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<td>American Indian or Alaska native</td>
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<td>1 (2%)</td>
<td>0 (0%)</td>
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<tr>
<td><strong>Age (years)</strong></td>
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<tr>
<td>Median (min-max)</td>
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<td><strong>Prior treatment in metastatic setting</strong></td>
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<td>No prior Chemotherapy</td>
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<td><strong>Median # of Chemotherapy Regimens</strong></td>
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<td>1 (range, 0-4)</td>
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</table>

*Chemotherapy in the metastatic setting*
Table 2. Efficacy Data

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<th>Best Response</th>
<th>TIG/nab-PAC (n=39)</th>
<th>nab-PAC (n=21)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Complete Response</td>
<td>3 (8%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Partial Response</td>
<td>8 (21%)</td>
<td>8 (38%)</td>
</tr>
<tr>
<td>Objective Response</td>
<td>11 (28%) (95% CI 14.9-45%)</td>
<td>8 (38%) (95% CI 18-61.1%)</td>
</tr>
<tr>
<td>Stable Disease</td>
<td>11 (28%)</td>
<td>4 (19%)</td>
</tr>
<tr>
<td>Clinical Benefit Rate (&gt; 4 cycles)</td>
<td>16 (41%)</td>
<td>11 (52%)</td>
</tr>
<tr>
<td>Progressive Disease</td>
<td>17 (44%)</td>
<td>9 (43%)</td>
</tr>
<tr>
<td>Median Duration of response – Days (Range)</td>
<td>118+ (84 to 1025+)</td>
<td>167+ (91 to 1004+)</td>
</tr>
<tr>
<td>Median Progression Free Survival - months</td>
<td>2.8 (95% CI 1.9-3.6)</td>
<td>3.7 (95% CI 2.3-5.7)</td>
</tr>
</tbody>
</table>
Table 3. Prior Therapy effect on Efficacy Data

<table>
<thead>
<tr>
<th></th>
<th>Chemo naïve (n=19)</th>
<th>Prior Chemo (n=41)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Complete Response</td>
<td>2 (11%)</td>
<td>1 (2%)</td>
</tr>
<tr>
<td>Partial Response</td>
<td>8 (38%)</td>
<td>8 (28%)</td>
</tr>
<tr>
<td>Objective Response</td>
<td>10 (53%)*** (95% CI 31-76.3%)</td>
<td>9 (22%) (95% CI 10.5-40.1%)</td>
</tr>
<tr>
<td>Stable Disease*</td>
<td>4 (21%)</td>
<td>11 (27%)</td>
</tr>
<tr>
<td>Clinical Benefit**</td>
<td>13 (68%)</td>
<td>14 (34%)</td>
</tr>
<tr>
<td>Progressive Disease</td>
<td>5 (26%)</td>
<td>21 (51%)</td>
</tr>
<tr>
<td>Median Duration of Response – Days (range)</td>
<td>137+ (84 to 1025+)</td>
<td>174+ (111 to 1004+)</td>
</tr>
<tr>
<td>Median Progression Free Survival – months</td>
<td>3.6 (95% CI 2.8-5.6)</td>
<td>2.5 (95% CI 1.9-3.7)</td>
</tr>
</tbody>
</table>

* Initial evaluation at day 56 (2 cycles of therapy)
** CR and PR and Stable disease greater than 4 cycles of therapy
*** p < 0.0347 (Fisher Exact Test)
### Table 4. Adverse Events Related with Protocol-Therapy

<table>
<thead>
<tr>
<th>Adverse Event</th>
<th>All Patients</th>
<th>TIG / nab-PAC (39)</th>
<th>nab-PAC(n=21)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Toxicity Grade</td>
<td>1</td>
<td>2</td>
</tr>
<tr>
<td>Fatigue</td>
<td></td>
<td>33 (54)</td>
<td>14 (36)</td>
</tr>
<tr>
<td>Alopecia</td>
<td></td>
<td>30 (49)</td>
<td>11 (28)</td>
</tr>
<tr>
<td>Peripheral Sensory Neuropathy</td>
<td></td>
<td>27 (44)</td>
<td>13 (33)</td>
</tr>
<tr>
<td>Anemia</td>
<td></td>
<td>25 (41)</td>
<td>8 (21)</td>
</tr>
<tr>
<td>Neutropenia</td>
<td></td>
<td>23 (38)</td>
<td>5 (13)</td>
</tr>
<tr>
<td>Nausea</td>
<td></td>
<td>14 (23)</td>
<td>6 (15)</td>
</tr>
<tr>
<td>Thrombocytopenia</td>
<td></td>
<td>6 (10)</td>
<td>2 (5)</td>
</tr>
<tr>
<td>Anorexia</td>
<td></td>
<td>6 (10)</td>
<td>1 (3)</td>
</tr>
<tr>
<td>Diarrhea</td>
<td></td>
<td>6 (10)</td>
<td>3 (8)</td>
</tr>
<tr>
<td>Vomit</td>
<td></td>
<td>6 (10)</td>
<td>2 (5)</td>
</tr>
</tbody>
</table>
Figure 1.

![Kaplan-Meier Progression Free Survival Estimates](chart1.png)

**A**
- Kaplan-Meier Progression Free Survival Estimates
- $P_{strat}=0.3152$
- Median Survival (95% CI)
  - Abraxane 3.7 (2.3-5.7)
  - Tigatuzumab_Abraxane 2.8 (1.9-3.6)
- Number at risk:
  - Abraxane: 21, Tigatuzumab_Abraxane: 39
- Progression From The First Treatment (Month)

![Kaplan-Meier Progression Free Survival Estimate](chart2.png)

**B**
- Kaplan-Meier Progression Free Survival Estimate
- $P_{strat}=0.1071$
- Median Survival (95% CI)
  - No_Prior_Cho 3.6 (2.8-5.6)
  - Other 2.5 (1.9-3.7)
- Number at risk:
  - No_Prior_Cho: 19, Other: 41
- Progression From The First Treatment (Month)
Figure 2
Legends to Figures

**Figure 1.** A. Progression Free Survival for each arm of the trial. B. Progression Free Survival According to Prior Therapy for all Patients Enrolled in the Trial.

**Figure 2.** Apoptotic membrane blebbing through DR5/Casp-3/ROCK1 signaling pathway. Genes associated with response to treatment with nab-PAC and TIG are highlighted in orange.
TBCRC 019: phase II trial of nab-PAC with/without the anti-death receptor 5 monoclonal antibody tigatuzumab in patients with triple negative breast cancer

Andres Forero-Torres, Katherine E Varley, Vandana Abramson, et al.

Clin Cancer Res  Published OnlineFirst March 16, 2015.