The treatment patterns, efficacy, and safety of nab®-paclitaxel for the treatment of metastatic breast cancer in the United States: results from health insurance claims analysis

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Abstract

Background: nab-Paclitaxel is an albumin-bound formulation of paclitaxel approved for the treatment of metastatic breast cancer (MBC). This analysis was designed to characterize the treatment patterns, efficacy, and safety of nab-paclitaxel for MBC treatment using health claims data from US health plans associated with Optum.

Methods: Women aged ≥ 18 years who initiated nab-paclitaxel for MBC treatment from January 1, 2005, to September 30, 2012, and who met eligibility criteria were selected from the Optum Research Database for this analysis. Patients were required to have complete medical coverage and pharmacy benefits, ≥ 6 months of continuous enrollment, and a diagnosis of MBC prior to nab-paclitaxel initiation. The pattern of use for nab-paclitaxel (e.g., regimen, schedule, duration, and administration) and claims-captured toxicities were characterized by line of therapy. Overall survival (OS) and time to next therapy or death (TNTD) were described by line of therapy, regimen, and schedule.

Results: Of the 664 nab-paclitaxel patients, 172 (25.9%) received it as first-line therapy, 211 (31.8%) as second-line therapy, and 281 (42.3%) as third-line or later therapy. Overall, the majority of patients received monotherapy (61%) and followed a weekly (71%) rather than an every 3 weeks treatment schedule. nab-Paclitaxel was often (31.7%) combined with targeted therapy (57.5% with bevacizumab and 23.9% with trastuzumab or lapatinib). The median duration of therapy was 128 days (4.2 months). For the overall population, median OS was 17.4 months (22.7, 17.4, and 15.1 months in first-, second-, and third-line or later therapy, respectively). Median TNTD was 6.1 months (7.1, 6.6, and 5.3 months in first-, second-, and third-line or later therapy, respectively). For patients aged ≤ 50 years or with ≥ 3 metastatic sites, median OS was 15.6 months. No new safety signal was identified.

Conclusions: In this US healthcare system, the majority of patients received nab-paclitaxel as second-line or later therapy, monotherapy, and weekly treatment. The efficacy and safety outcomes of nab-paclitaxel observed in this real-world setting appear consistent with those from clinical trial data.

Keywords: Metastatic breast cancer, nab-Paclitaxel, Claims analysis

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Background

Taxanes are some of the most active chemotherapeutic agents in the treatment of breast cancer [1–3]. However, sensory neuropathy, neutropenia, and significant toxicities—such as severe hypersensitivity reactions, which require substantial premedication with high doses of steroids and antihistamines—have been reported in patients treated with solvent-based (sb) taxanes (ie, paclitaxel and docetaxel) [4–6]. An albumin-bound formulation of paclitaxel (Abraxane®, nab-paclitaxel) was developed in an effort to overcome the toxicities associated with sb-paclitaxel and improve efficacy [7]. Preclinical studies have shown that nab-paclitaxel delivers a 33% higher paclitaxel concentration to tumors and demonstrates enhanced transport across endothelial cell monolayers compared with sb-paclitaxel [7]. Recently published population pharmacokinetic data on nab-paclitaxel compared with sb-paclitaxel demonstrated more rapid and greater tissue penetration and slower elimination of paclitaxel [8]. nab-Paclitaxel was approved by the US Food and Drug Administration in January 2005 for the treatment of breast cancer after failure of combination chemotherapy, including anthracyclines, for metastatic disease or relapse within 6 months of adjuvant chemotherapy [9].

The safety and efficacy of single-agent nab-paclitaxel have been well established in clinical trials of patients with metastatic breast cancer (MBC) (Table 1) [10–13]. In a phase three trial [10], nab-paclitaxel dosed at 260 mg/m² every 3 weeks (q3w) vs sb-paclitaxel dosed at 175 mg/m² q3w demonstrated a significantly higher overall response rate (33% vs 19%; P = 0.001) and a significantly longer time to tumor progression (5.3 vs 3.9 months; P = 0.006). The incidence of grade 4 neutropenia was significantly lower with nab-paclitaxel compared with sb-paclitaxel. Although the incidence of grade 3 sensory neuropathy was significantly higher with nab-paclitaxel compared with sb-paclitaxel, it was manageable with dose modifications and treatment interruptions and improved to grade ≤2 in a median of 22 days. Although the 260 mg/m² q3w nab-paclitaxel monotherapy regimen is indicated for the treatment of patients with MBC, other doses and schedules of nab-paclitaxel have been explored in clinical trials. In a phase two trial, three different nab-paclitaxel regimens (300 mg/m² q3w, 100 mg/m², or 150 mg/m² given weekly for the first 3 of 4 weeks [qw 3/4]) were compared with docetaxel 100 mg/m² qw for the treatment of chemotherapy-naïve patients with MBC [12, 13]. Results from this trial indicated that the 150 mg/m² qw 3/4 dose of nab-paclitaxel was a significantly more effective regimen than docetaxel [13]. Median overall survival (OS) was 33.8 months compared with 22.2, 27.7, and 26.6 months for nab-paclitaxel 100 mg/m² qw 3/4, nab-paclitaxel 300 mg/m² q3w, and docetaxel, respectively [13]. The frequency of grade 3/4 neutropenia, febrile neutropenia, and fatigue was lower in all nab-paclitaxel arms compared with docetaxel. The incidence of grade 3 sensory neuropathy was higher for the 300 mg/m² q3w and 150 mg/m² qw 3/4 nab-paclitaxel regimens vs docetaxel, which may be related to the higher median dose intensities associated with these two nab-paclitaxel dose regimens (100 and 101 mg/m²/week, respectively), compared with docetaxel (33 mg/m²/week) [13]. The median time to improvement of sensory neuropathy to ≤ grade 2 was 20 to 22 days for nab-paclitaxel compared with 41 days for docetaxel [13].

nab-Paclitaxel has also been studied in combination with other cytotoxic or targeted agents for the treatment of MBC (Table 1) [14–19]. Results of phase two trials of nab-paclitaxel in combination with gemcitabine and oral capecitabine have demonstrated efficacy and favorable tolerability. The results of other clinical trials have shown that nab-paclitaxel is a reasonable substitution for sb-taxanes in combination with targeted agents such as bevacizumab, trastuzumab, and lapatinib for the treatment of MBC [16–21].

Clinical trials target highly selected patients with restrictive eligibility criteria, limiting the generalizability of outcomes. Therefore, we conducted an observational study based on US health insurance claims data to characterize the therapeutic context (line of therapy, monotherapy vs combination therapy, and dosing schedule) and to estimate the OS and time to next therapy or death (TNTD) among patients who received nab-paclitaxel for the treatment of MBC.

Methods

Data source

In this retrospective cohort study, health insurance claims data were extracted from the Optum Research Database, which contains eligibility, pharmacy claims, medical claims, and other information, such as mortality data, from health plans associated with Optum. The health claims are linked to enrollment information with data covering the period from 1993 to present. The information in the claims database includes over 12 million individuals from geographically diverse locations across the United States who have both medical and pharmacy benefit coverage. Medical claims or encounter data were collected from all available healthcare sites (inpatient hospital, outpatient hospital, emergency room, physician’s office, surgery center, etc.) for virtually all types of provided services, including specialty, preventive, and office-based treatments. Diagnoses on the claims are recorded using International Classification of Disease, Ninth Revision Clinical Modification (ICD-9-CM) codes. Procedures map to ICD-9-CM, Current Procedural Terminology, and Healthcare Common Procedure
Table 1: Select clinical trials of nab-P in metastatic breast cancer

| Trial | Phase | Patient population | Regimen | Efficacy | Select Grade ≥ 3 AEs, % |
|-------|-------|--------------------|---------|----------|------------------------|
|       |       |                    |         | PFS, mo  | OS, mo                 |
|       |       |                    |         | Neutropenia | Neuropathy            |
| **Monotherapy** |       |                    |         |                       |                        |
| Ibrahim et al. 2005 [11] | 2     | First line (n = 15) | nab-P 300 mg/m² q3w | TTP 6.1 | 51 | 11^b |
| | | Second line or later (n = 48) | | | | |
| Gradishar et al. 2009 [12] & 2012 [13] | 2     | First line | nab-P 300 mg/m² q3w (n = 76) | 11.0 | 27.7 | 43 | 21^b |
| | | | nab-P 100 mg/m² qw 3/4 (n = 76) | 12.8 | 22.2 | 25 | 9^b |
| | | | nab-P 150 mg/m² qw 3/4 (n = 74) | 12.9 | 33.8 | 45 | 22^b |
| Gradishar et al. 2005 [10] | 3     | First line (n = 97) | nab-P 260 mg/m² q3w | TTP 5.3 | 15.0 | 30 | 10^b |
| | | Second line or later (n = 132) | | | | |
| **Combination therapy with cytotoxic agents** | | | | | | |
| Roy et al. 2009 [14] | 2     | First line (N = 50) | nab-P 125 mg/m² + gemcitabine 1000 mg/m² qw 2/3 | 7.9 | Median not reached; 6-mo OS 92% | 54 | 8^b |
| Schwartzberg et al. 2012 [15] | 2     | First line (N = 50) | nab-P 125 mg/m² qw 2/3 + oral capecitabine 825 mg/m² twice daily on days 1 and 15 of a 21-day cycle | 10.6 | 19.9 | 10 | 2^b |
| **Combination therapy with targeted agents** | | | | | | |
| Seidman et al. 2013 [16] | 2     | First line, HER2 negative | nab-P 130 mg/m² qw + bev 10 mg/kg qw (n = 79) | 8.8 | 23.7 | 33 | 46 |
| | | | nab-P 260 mg/m² qw 3/4 + bev 10 mg/kg qw (n = 54) | 5.8 | 19.0 | 6 | 56 |
| | | | nab-P 260 mg/m² qw 3/4 + bev 15 mg/kg qw (n = 75) | 7.7 | 21.3 | 16 | 33 |
| Rugo et al. 2015 [17] | 3     | First line, predominantly HER2 negative | nab-P 150 mg/m² qw 3/4 + bev 10 mg/kg qw (n = 271) | 9.3 | 23.5 | 51 | 27 |
| Mirtsching et al. 2011 [18] | 2     | First line (N = 72) | nab-P 125 mg/m² qw 3/4 + trastuzumab 4 mg/kg bolus then 2 mg/kg qw (HER2 positive only) | 14.5 | 29.0 | 11^b | 8^b |
| Yardley et al. 2013 [19] | 2     | First/second line (N = 60) | nab-P 125 mg/m² qw 3/4 + oral lapatinib 1250 mg daily | 9.1 | Median not reached | 22^b | 3^b |

AE adverse event, bev bevacizumab, HER2 human epidermal growth factor receptor, nab-P nab-paclitaxel, OS overall survival, PFS progression-free survival, TTP time to tumor progression, qw every week, q2w every 2 weeks, q3w every 3 weeks, qw 2/3 weekly for the first 2 of 3 weeks; qw 3/4 weekly for the first 3 of 4 weeks

^No grade 4 events

Coding System codes. Pharmacy claims data include drug name, dosage form, drug strength, fill date, days of supply, financial information, and de-identified patient and prescriber codes, allowing for longitudinal tracking of medication refill patterns and changes in medications.

**Study population**

The study population consisted of women aged ≥ 18 years with a diagnosis of MBC who received nab-paclitaxel treatment. Patients were eligible for the study if they had complete medical coverage and pharmacy benefits; had ≥ 6 months of continuous enrollment in a US health plan from January 1, 2005 to September 30, 2012; and had a diagnosis of MBC prior to the initiation of nab-paclitaxel. Patients were selected for the claims analysis based on the criteria listed in Fig. 1. Diagnosis codes appearing on claims suggesting a laboratory or diagnostic service were not considered when these criteria were applied, because these claims often reflect a “rule-out” diagnosis that has not yet been confirmed.
Lines of therapy

New users of nab-paclitaxel were defined as those with a first dispensing of nab-paclitaxel within the study period (January 1, 2005 to September 30, 2012), with no dispensing of nab-paclitaxel during the 6 months prior to the first dispensing (baseline period). Patients who received neoadjuvant (≤4 months prior to surgery) or adjuvant (≤90 days after surgery) therapy with nab-paclitaxel were excluded. The index date was defined as the date of nab-paclitaxel initiation. Patients who met the cohort entry eligibility criteria were further categorized into 3 subgroups by line of therapy (first-, second-, or third-line or later) with nab-paclitaxel.

A patient must have received ≥1 cycle (defined as 30 days) of treatment prior to being defined as switching to a greater line of therapy. Any switching or addition of agents within 30 days of the start of each line of therapy was considered to be the same line of therapy.

First-line therapy

First-line therapy with nab-paclitaxel was defined as initial dispensing of nab-paclitaxel as the first chemotherapy received after a diagnosis of metastatic disease. All agents received within 30 days following nab-paclitaxel were considered part of first-line therapy.

Second-line therapy

Second-line therapy with nab-paclitaxel was defined as dispensing of nab-paclitaxel as part of second-line therapy, defined as additional treatment different from the first-line therapy and initiated ≥30 days after the first chemotherapy or after a large gap (eg, 90 days) in therapy.

Third-line or later therapy

Third-line or later therapy with nab-paclitaxel was defined as a first dispensing of nab-paclitaxel as part of third-line therapy, defined as any additional treatment different from any initiated first- or second-line therapy and after 60 days of the first chemotherapy or after a large gap (eg, 90 days) in therapy. Similar methods were used to identify later lines of therapy.

Outcome identification

The study outcomes included all-cause death, TNTD, and major toxicities following nab-paclitaxel initiation. All-cause death was identified using Social Security Administration data linked to claims data. TNTD was used as a surrogate of progression-free survival (PFS). Major toxicities were identified following each line of therapy and determined by tabulating the 25 most frequent ICD-9-CM diagnoses codes. The toxicities of interest included select adverse events consistent with the known safety profile of nab-paclitaxel: neutropenia, anemia, thrombocytopenia, infections, peripheral neuropathy,

Fig. 1 Flowchart of study patients. Patients in the Optum Research Database who met the criteria outlined in the flowchart were included in the claims analysis. *xx indicates any subcode.
asthenia, nausea, vomiting, diarrhea, fluid retention, myalgia/arthralgia, and alopecia.

Statistical analysis
A descriptive analysis was conducted to identify the background characteristics, nab-paclitaxel treatment patterns, and nab-paclitaxel toxicities of interest in patients with MBC by line of therapy. The background characteristics, including demographics and breast cancer risk factors, were ascertained during the 6-month baseline period. Treatment patterns of nab-paclitaxel were described in terms of treatment regimen (monotherapy or combination therapy), treatment schedule (weekly or q3w), duration of line of therapy, number of administrations, intervals between dispensings, and dose. The occurrence of toxicity claims of interest following nab-paclitaxel treatment was also summarized.

A survival analysis using an intent-to-treat approach was performed to evaluate the OS and TNTD. Each patient was followed from nab-paclitaxel initiation in each line of therapy until the first occurrence of a study endpoint (all-cause death and TNTD, separately), disenrollment from the health plan (eg, a gap of > 32 days in membership), or the end of the study period (September 30, 2012). OS was defined as the interval between the first dispensing of nab-paclitaxel and death. TNTD was defined as the interval between the first dispensing of nab-paclitaxel and switching of line of therapy or death. Kaplan-Meier plots were used to depict the cumulative probability of OS and TNTD by line of therapy. The median OS and median TNTD as well as their 95% CIs were also estimated. These survival analyses were conducted overall, by line of therapy, by regimen, and by schedule. A subgroup analysis was also performed among patients aged ≤ 50 years or with ≥ 3 metastatic sites.

Results
Patient characteristics
There were 2637 nab-paclitaxel initiators identified during the study period. After the eligibility criteria were applied, a total of 664 patients remained in the final analysis (Fig. 1). The 664 eligible patients were predominantly aged 50 to 69 years and were from the southern region of the United States (Table 2). There were sparse data recorded in the claims for family history of breast cancer, oral contraceptive use, hormone replacement therapy, alcohol use, obesity, and smoking. All patients had physician visits during the 6-month baseline period, 38.3% of patients visited an emergency department, and 31.8% of patients were admitted to a hospital. The median duration of hospitalization was 5 days (Table 2). The median length of health plan membership was 2.4 years prior to initiation of nab-paclitaxel.

nab-Paclitaxel treatment patterns
Treatment patterns by line of therapy are summarized in Table 3. There were 172 (25.9%) patients who received nab-paclitaxel as first-line therapy, 211 (31.8%) as second-line therapy, and 281 (42.3%) as third-line or later therapy. Overall, there were 405 (61.0%) users who had nab-paclitaxel administered as monotherapy and 259 (39.0%) who had nab-paclitaxel administered as combination therapy. When nab-paclitaxel was given as a combination therapy, targeted agents were often used (57.5% bevacizumab and 23.9% trastuzumab or lapatinib). Bevacizumab combination was more often prescribed in first-line therapy with nab-paclitaxel vs trastuzumab or lapatinib (81.0 vs 4.8%). Trastuzumab combination therapy was more often given in the third line or later (39.5%) compared with first-line (4.8%) or second-line (17.1%) therapy. Of the 605 users whose treatment schedules could be determined, a majority (n = 428 [70.7%]) received weekly treatment and 177 (29.3%) received q3w treatment. The median durations that patients received nab-paclitaxel as first-line, second-line, and third-line or later therapy were 159 days (5.2 months), 119 days (3.9 months), and 122 days (4.0 months), respectively (Table 3).

nab-Paclitaxel safety outcomes
Table 4 shows the claims of the major toxicities of interest among patients without corresponding events during the baseline period. Anemia (26.3%), nausea and vomiting (24.5%), neutropenia (17.5%), and asthenia (15.6%) were the most common incident claims. This study also found that 14.5% of claims were for peripheral neuropathy. These events were more frequently recorded in patients with first-line therapy compared with patients receiving nab-paclitaxel in later lines of therapy.

nab-Paclitaxel efficacy outcomes
Patients who received first-line nab-paclitaxel–based therapy appeared to have longer median survival vs second- and third-line or later therapy (Fig. 2): 22.7, 17.4, and 15.1 months, respectively (Table 5; Fig. 2). Median TNTD values were 7.1, 6.6, and 5.3 months by first-, second-, and third-line or later therapy, respectively (Table 5; Fig. 3). In the subgroup of patients aged ≤ 50 years or who had ≥ 3 metastases (n = 400), the median OS was 15.6 months (95% CI, 12.9–17.4 months), and the median TNTD was 5.7 months (95% CI, 4.9–6.4 months). Patients who received nab-paclitaxel combination therapy had a median survival time of 18.7 months compared with 16.8 months for those who received nab-paclitaxel monotherapy (Table 5); the respective values for median TNTD were 6.5 and 5.8 months.
Median OS and TNTD values stratified by line of therapy and treatment schedule are shown in Table 6. Median OS was 18.6 months for weekly and 17.4 months for q3w nab-paclitaxel, and median TNTD was 6.5 months for weekly and 6.0 months for q3w nab-paclitaxel, respectively.

**Discussion**

Breast cancer is a heterogeneous disease with various clinical and biological features [22]. Multiple molecular alterations and cellular pathway dysregulations may occur during disease development and progression [23]. Some types of breast cancers are more aggressive than others, and sensitivity to treatment may differ [24, 25]. To get a real-world look at nab-paclitaxel treatment patterns, efficacy, and safety since market approval, we carried out a claims-based retrospective analysis using a large US commercial health insurance database and selected women undergoing treatment with nab-paclitaxel for MBC.

Consistent with the National Comprehensive Cancer Network guidelines [1], our analysis indicated that nab-paclitaxel was most often prescribed as second-line or later therapy and administered as monotherapy. When nab-paclitaxel was used in combination, the targeted agents (e.g., bevacizumab, trastuzumab, or lapatinib) were most often prescribed. Patients treated with nab-paclitaxel in the first line appeared to have favorable survival relative to patients treated in later lines of therapy. However, the treatment effect of nab-paclitaxel as first-line therapy may have been overestimated because the criteria for first-line therapy required patients to be treated for at least 30 days. Therefore, patients who discontinued...
treatment early may not have been captured. Overall, the safety and efficacy profiles of nab-paclitaxel in this setting of US women with MBC were consistent with clinical trial experience (Table 1) [10–19].

Our analysis showed a median OS of 17.4 months for the overall population of patients with MBC who received various doses, schedules, and regimens of nab-paclitaxel across all lines of therapy. These results are in line with those of a phase two trial [11] and the pivotal phase three trial [10], which showed a median OS of 14.6 and 15.0 months, respectively, in patients receiving nab-paclitaxel monotherapy (260–300 mg/m² q3w) for ≥ first-line treatment of MBC (Table 1). In a phase two trial of chemotherapy-naive patients with MBC, median

| Variable | First line (n = 172) | Second line (n = 211) | Third or later line (n = 281) | All (N = 664) |
|----------|----------------------|----------------------|-----------------------------|--------------|
| n%       | n%                   | n%                   | n%                          | n%           |
| Treatment regimen |                      |                      |                             |              |
| Monotherapy          | 109 (63.4)           | 129 (61.1)           | 167 (59.4)                  | 405 (61.0)   |
| Combination chemotherapy | 63 (36.6)           | 82 (38.9)           | 114 (40.6)                  | 259 (39.0)   |
| Bevacizumab          | 51 (81.0)            | 53 (64.6)            | 45 (39.5)                   | 149 (57.5)   |
| Trastuzumab or lapatinib | 3 (4.8)              | 14 (17.1)           | 45 (39.5)                   | 62 (23.9)    |
| Gemcitabine, carboplatin, pegylated liposomal doxorubicin/doxorubicin, docetaxel, doxorubicin, paclitaxel, irinotecan, vinorelbine, or 5-fluorouracil | 3 (4.8)              | 8 (9.8)              | 14 (12.3)                   | 25 (9.7)     |
| ≥ 2 agents from above list | 6 (9.5)              | 7 (8.5)              | 10 (8.8)                    | 23 (8.9)     |
| Treatment scheduleb |                      |                      |                             |              |
| Weekly              | 152 (93.7)           | 193 (91.4)           | 260 (92.5)                  | 605 (91.7)   |
| q3w                 | 28 (16.3)            | 46 (22.6)            | 89 (30.6)                   | 163 (24.3)   |
| Duration of line treatment, d | 159.0 (83.0–241.0)  | 119.0 (65.0–191.0)  | 122.0 (76.0–199.0)          | 128.0 (76.0–199.0) |
| Initial dose, unitc | 200.0 (200.0–429.0)  | 200.0 (200.0–400.0)  | 200.0 (200.0–400.0)         | 200.0 (200.0–400.0) |
| Average dose, unitc | 227.5 (200.0–394.4)  | 220.0 (200.0–400.0)  | 214.9 (200.0–400.0)         | 218.5 (200.0–400.0) |
| IQR interquartile range, q3w every 3 weeks |
| aData are from the Optum Research Database, January 1, 2005 to September 30, 2012 |
| b59 patients (20 in first-line, 18 in second-line, and 21 in third-line or later therapy) could not be classified into a weekly or q3w treatment schedule |
| cEach unit is equivalent to 1 mg. The dosage calculated may not reflect the exact dose dispensed or received |

Table 3: Treatment patterns of nab-paclitaxel initiators by line of therapya

| Adverse event | First line (n = 172) | Second line (n = 211) | Third or later line (n = 281) | All (N = 664) |
|---------------|----------------------|----------------------|-----------------------------|--------------|
| n%            | n%                   | n%                   | n%                          | n%           |
| Neutropenia   | 165 (33.7)           | 190 (26.3)           | 222 (23.3)                  | 577 (17.5)   |
| Anemia        | 123 (43.7)           | 133 (39.4)           | 147 (38.9)                  | 403 (106.3)  |
| Thrombocytopenia | 163 (47.1)           | 199 (52.5)           | 264 (53.9)                  | 626 (24.3)   |
| Infections    | 152 (20.0)           | 184 (24.0)           | 242 (27.1)                  | 578 (41.3)   |
| Peripheral neuropathy | 129 (26.5)           | 155 (30.0)           | 200 (31.5)                  | 484 (70.1)   |
| Atenol    | 141 (31.7)           | 178 (37.7)           | 232 (33.7)                  | 551 (37.7)   |
| Nausea and vomiting | 143 (32.9)           | 159 (30.9)           | 179 (30.6)                  | 481 (31.8)   |
| Diarrhea      | 162 (30.0)           | 199 (36.7)           | 267 (45.3)                  | 628 (32.1)   |
| Fluid retention | 160 (30.0)           | 196 (37.7)           | 257 (45.0)                  | 613 (37.1)   |
| Myalgia/arthralgia | 133 (27.9)           | 159 (30.9)           | 230 (40.6)                  | 522 (30.1)   |
| Alopecia      | 171 (30.0)           | 209 (35.0)           | 280 (40.0)                  | 660 (10.1)   |
| IQR interquartile range |
| aData are from the Optum Research Database, January 1, 2005 to September 30, 2012 |
| bFollow-up time was calculated from index date until disenrollment from the health plan, death (or treatment discontinuation), or the end of the study period (September 30, 2012) |
| cTotal n refers to the total number of patients without baseline events for its respective subgroup |
OS was 22.2 to 33.8 months for weekly and 27.7 months for q3w nab-paclitaxel [13]. The OS claims for first-line therapy (monotherapy; median of 20.8 months; weekly median of 21.6 months) are similar to the median OS values reported in the phase two trial for the nab-paclitaxel 100 mg/m² weekly arm (22.2 months). Notably, treatment duration with nab-paclitaxel was much longer in the phase two trial (6.9 months), which may account for the longer OS results vs this claims analysis.

For the patients who received nab-paclitaxel monotherapy, the median TNTD was 5.8 months, similar to the reported median time to disease progression of patients with MBC who received nab-paclitaxel monotherapy (260–300 mg/m² q3w) for ≥ first-line treatment in the

**Table 5** OS and TNTD among nab-paclitaxel initiators by line of therapy and treatment regimen

| Variable                      | Total no. | OS Median, mo (95% CI) | TNTD Median, mo (95% CI) |
|-------------------------------|-----------|------------------------|--------------------------|
| Overall                       | 664       | 305 17.4 (16.1–19.0)   | 491 6.1 (5.6–6.7)        |
| Monotherapy                   | 405       | 184 16.8 (14.5–18.5)   | 293 5.8 (5.0–6.7)        |
| Combination therapy           | 259       | 121 18.7 (15.9–24.0)   | 198 6.5 (5.8–7.8)        |
| First line                    | 172       | 68 22.7 (18.6–34.3)    | 127 7.1 (6.2–9.0)        |
| Monotherapy                   | 109       | 37 20.8 (16.5–39.2)    | 76 6.7 (5.0–7.4)         |
| Combination therapy           | 63        | 31 23.0 (18.7–37.3)    | 51 8.5 (6.5–13.1)        |
| Second line                   | 211       | 99 17.4 (14.1–22.3)    | 151 6.6 (5.3–8.1)        |
| Monotherapy                   | 129       | 67 15.6 (12.8–21.4)    | 97 5.8 (4.7–7.9)         |
| Combination therapy           | 82        | 32 >18.7 –          | 54 7.3 (5.1–9.7)         |
| Third or later line           | 281       | 138 15.1 (12.7–16.9)   | 213 5.3 (4.6–6.0)        |
| Monotherapy                   | 167       | 80 15.6 (11.6–17.8)    | 120 5.3 (4.2–6.1)        |
| Combination therapy           | 114       | 58 14.3 (11.9–17.4)    | 93 5.5 (4.6–6.4)         |

OS overall survival. TNTD time to next therapy or death.

*Data are from the Optum Research Database, January 1, 2005 to September 30, 2012.

*Follow-up time was calculated from index date until disenrollment from the health plan, death (or treatment discontinuation), or the end of the study period (September 30, 2012).*

*The 95% CI of median survival time is missing because more than half of the patients survived during the study period.*

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**Fig. 2** Overall survival by line of therapy. Kaplan-Meier plot depicting the cumulative probability of overall survival by line of therapy.
phase two and phase three trials: 6.1 and 5.3 months, respectively [10, 11]. However, the median TNTD in the claims analysis was shorter than the median PFS reported in the phase two trial of patients receiving first-line nab-paclitaxel monotherapy: median PFS of 11.1 months with 300 mg/m² q3w nab-paclitaxel and 12.8 to 12.9 months with 100 to 150 mg/m² weekly nab-paclitaxel [12].

This analysis also supports clinical trial data indicating that patients with poor prognostic characteristics derive a clinical benefit from nab-paclitaxel therapy. Patients treated with nab-paclitaxel who were aged ≤ 50 years or had ≥ 3 metastases had outcomes comparable with those of the overall population (median OS: 15.6 months). These results are also similar to those from a retrospective analysis of patients from the pivotal phase three trial who received therapy later than first line and had ≥ 3 metastases (median OS: 13.0 months) [26]. Furthermore, in a separate retrospective analysis of patients with poor

![Fig. 3 Time to next therapy or death (TNTD) by line of therapy. Kaplan-Meier plot depicting the cumulative probability of TNTD by line of therapy.](image-url)

| Variable                  | Total No. | OS Events | Median, mo | 95 % CI       | TNTD Events | Median, mo | 95 % CI       |
|---------------------------|-----------|-----------|------------|---------------|-------------|------------|---------------|
| Overall                   | 605       | 274       | 18.1       | (16.7–20.8)   | 452         | 6.4        | (5.8–6.8)     |
| Weekly                    | 428       | 191       | 18.6       | (16.1–21.6)   | 316         | 6.5        | (5.8–7.1)     |
| q3w                       | 177       | 83        | 17.4       | (14.0–21.7)   | 136         | 6.0        | (5.0–7.1)     |
| First line                | 152       | 60        | 23.0       | (19.0–37.2)   | 115         | 7.1        | (6.2–9.2)     |
| Weekly                    | 114       | 49        | 21.6       | (18.0–29.3)   | 85          | 7.4        | (6.2–10.5)    |
| q3w²                      | 38        | 11        | >22.7      | –             | 30          | 6.6        | (4.5–10.5)    |
| Second line               | 193       | 86        | 19.4       | (14.4–22.6)   | 138         | 6.9        | (5.7–8.6)     |
| Weekly                    | 134       | 55        | 19.8       | (14.4–23.8)   | 94          | 6.9        | (5.7–8.8)     |
| q3w                       | 59        | 31        | 17.4       | (12.0–26.1)   | 44          | 6.9        | (5.0–9.9)     |
| Third or later line       | 260       | 128       | 15.3       | (12.8–17.0)   | 199         | 5.5        | (4.9–6.2)     |
| Weekly                    | 180       | 87        | 15.9       | (12.9–18.1)   | 137         | 5.8        | (4.9–6.5)     |
| q3w³                      | 80        | 41        | 13.5       | (8.9–17.4)    | 62          | 5.0        | (4.0–6.4)     |

OS overall survival, q3w every 3 weeks, TNTD time to next therapy or death

*Data are from the Optum Research Database, January 1, 2005 to September 30, 2012

*Follow-up time was calculated from index date until disenrollment from the health plan, death (or treatment discontinuation), or the end of the study period (September 30, 2012)

*59 patients (20 in first line, 18 in second line, and 21 in third or later line) could not be classified into a weekly or every 3 weeks treatment schedule

*The 95 % CI of median survival time is missing because more than half of the subjects survived during the study period
prognosis, a favorable survival benefit was demonstrated in patients with visceral dominant metastases (OS: 15.1–32.1 months) or a short disease-free interval (OS: 14.6–19.1 months) who received nab-paclitaxel as first-line therapy [27]. Results from this claims analysis are similar to those for the intent-to-treat population of those trials as well (Table 1) [10, 13].

Median TNTD and OS for patients who received nab-paclitaxel–based combination therapy were in line with results of clinical trials of nab-paclitaxel–based combination therapy (Table 1) [14–19]. Our analysis showed that, when used in combination, nab-paclitaxel was most often given with bevacizumab (58 %), and bevacizumab combination therapy was more often initiated in the first line (81 %). It is noted that longer survival and TNTD were observed in the first line for combination therapy vs monotherapy (23.0 vs 20.8 months and 8.5 vs 6.7 months, respectively). In 2011, bevacizumab for the treatment of breast cancer was revoked by the US Food and Drug Administration [28]. The effect of this was reflected in a marked decrease (nearly 70 %) in the rate of nab-paclitaxel combination therapy use after 2011 and was likely due to bevacizumab being revoked (data not shown). At this time it is unclear what the optimal combination partner is for nab-paclitaxel in patients with human epidermal growth factor receptor 2–negative MBC. Currently a phase two/three trial is under way to determine the efficacy and safety of nab-paclitaxel in combination with gemcitabine or carboplatin in patients with triple-negative MBC [29].

The common adverse events identified in the clinical trials were also explored in this claims-based study. The occurrence of select known nab-paclitaxel toxicities (eg, neutropenia, peripheral neuropathy, anemia, infections, and nausea and vomiting) ranged from 15 % to 26 % and was lower than that noted in clinical trials (Table 1) [10–19]. In particular, the frequency of reported neuropathy was relatively low (<15 %) compared with that reported in clinical trials for nab-paclitaxel therapy [10–12], indicating that this adverse event may have been underrepresented in the claims database. This is likely explained in part by the more robust patient monitoring and collection of safety data in the clinical trial setting. In addition, claims data for analysis tend to bias toward underreporting in comparison with prospective National Cancer Institute Common Terminology Criteria for Adverse Events documentation.

Although claims analyses are extremely valuable for the efficient and effective examination of healthcare outcomes, treatment patterns, and healthcare resource utilization, it is challenging to compare our study findings with those from clinical trials. Historical trials of nab-paclitaxel recruited patients according to highly restrictive criteria, and the patients received a specific line of nab-paclitaxel therapy, treatment regimen, or treatment schedule during the study period. For example, the phase two trials often targeted first-line therapy with various doses and schedules, whereas the pivotal phase three trial mixed lines of therapy at a q3w dose/schedule (Table 1). In addition, claims analyses are unable to estimate disease progression. TNTD may be perceived as a weak surrogate for PFS because a potential time lapse between disease progression and initiation of a new line of therapy is not captured. This in effect could overestimate a benefit of treatment. Estimating an overall response rate and determining the grade of toxicities are also not feasible using claims data.

Furthermore, because claims are collected for the purpose of payment and not research, inherent limitations in our claims analysis included the potential for incorrect reporting of diagnosis codes, missing patients with early-stage breast cancer with patients with MBC, missing information on hormone receptor status, misinterpreting disease-onset dates, misclassifying the line of therapy, and inaccurately estimating actual drug dosages and schedules. However, the application of a well-defined algorithm, including the combination of diagnoses, procedures, and medications, reduced the potential for false-positive cases and the misclassification of line of therapy.

Conclusions

nab-Paclitaxel is administered more frequently as a single agent on a weekly schedule and as second-line or later therapy to patients with MBC in a US healthcare system. This analysis demonstrates the use of nab-paclitaxel weekly or q3w and its use for the treatment of patients aged ≤ 50 years or with ≥ 3 metastatic sites. The benefit observed in this US healthcare system is consistent with that from previously reported clinical trials. No new safety signals were identified. Furthermore, our analysis showed that, when used in combination, nab-paclitaxel was most often combined with bevacizumab in first-line therapy. However, because the accelerated approval of bevacizumab for MBC was withdrawn due to the lack of an OS advantage in the RIBBON-1 and AVADO trials [28], bevacizumab is no longer used as standard therapy in MBC. Additional nab-paclitaxel combination partners are being evaluated in patients with MBC, including gemcitabine or carboplatin in patients with triple-negative MBC [29]. Identification of an optimal nab-paclitaxel combination regimen may provide additional options for patients with MBC. Finally, outcomes of this real-world claims analysis are consistent with the data demonstrated in key clinical trials, affirming the effectiveness and manageable safety profile of nab-paclitaxel across all lines of therapy in patients with MBC.
Abbreviations
ICD-9-CM: International Classification of Diseases, Ninth Revision, Clinical Modification; MBC: Metastatic breast cancer; OS: Overall survival; PFS: Progression-free survival; q2w: Every 2 weeks; q3w: Every 3 weeks; qw 2/3: Weekly for the first 2 of 3 weeks; qw 3/4: Weekly for the first 3 of 4 weeks; qw: Weekly; sb: Solvent based; TNTD: Time to next therapy or death.

Competing interests
Calhia Liang: research funding, Celgene; Ling Li: nothing to disclose; Cindy Duval Fraser: employment or leadership position and stock ownership, Celgene; Amy Ko: employment or leadership position and stock ownership, Celgene; Deyanira Cozzo: employment or leadership position, research funding, and stock ownership, Celgene; Cheryl Enger: nothing to disclose; Debra Patt: nothing to disclose.

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CL drafted the initial manuscript. LL, CDF, AK, DC, CE, and DP revised and approved the final manuscript. All authors read and approved the final manuscript.

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References
1. NCCN Clinical Practice Guidelines in Oncology. Breast Cancer. V2.2015. 2015. http://www.nccn.org/professionals/physician_gls/gl水墨_guidelines.asp. Accessed May 7, 2015.
2. De Laurentis M, Cancellio G, D’Agostino D, Giuliano M, Giordano A, Montagna E, et al. Taxane-based combinations as adjuvant chemotherapy of early breast cancer: a meta-analysis of randomized trials. J Clin Oncol. 2008;26(1):44–53.
3. Cardoso F, Bedard PL, Winer EP, Pagani O, Senkus-Konefa E, Fallowfield LJ, et al. International guidelines for management of metastatic breast cancer: combination vs sequential single-agent chemotherapy. J Natl Cancer Inst. 2002;104(17):1174–81.
4. Syrigou E, Dannios I, Kottees E, Makrilia N, Tourkantonis I, Dilana K, et al. Hypersensitivity reactions to docetaxel: retrospective evaluation and development of a desensitization protocol. Int Arch Allergy Immunol. 2011;153(3):320–4.
5. Bristol-Myers Squibb Co. Taxol® (paclitaxel) [prescribing information]. Princeton, NJ USA; Bristol-Myers Squibb Co, 2011.
6. sanofi-aventis US LLC. Taxotere® (docetaxel) [prescribing information]. Bridgewater, NJ USA; sanofi-aventis US LLC, 2013.
7. Desai N, Trieu V, Yao Z, Louie L, Ci S, Yang A, et al. Increased antitumor activity, intratumor paclitaxel concentrations, and endothelial cell transport of cremophor-free, albumin-bound paclitaxel. Abi-007, compared with cremophor-based paclitaxel. Clin Cancer Res. 2006;12(4):1317–24.
8. Chen N, Li Y, Ye Y, Palmisano M, Choppa R, Zhou S. Pharmacokinetics and pharmacodynamics of nab-paclitaxel in patients with solid tumors: disposition kinetics and pharmacology distinct from solvent-based paclitaxel. J Clin Pharmacol. 2014;54(10):1097–107.
9. Celgene Corporation. Abraxane® (paclitaxel protein-bound particles for injectable suspension) [prescribing information]. Summit, NJ Celgene Corporation, 2013.
10. Gradishar WJ, Tjulandin S, Davidson N, Shaw H, Desai N, Bhar P, et al. Phase III trial of nanoparticle albumin-bound paclitaxel compared with polyethylated castor oil-based paclitaxel in women with breast cancer. J Clin Oncol. 2005; 23(31):7794–803.
11. Ibrahim NK, Samuels B, Page R, Doval D, Patel KM, Rao SC, et al. Multicenter phase II trial of ABI-007, an albumin-bound paclitaxel, in women with metastatic breast cancer. J Clin Oncol. 2005;23(25):6019–26.
12. Gradishar WJ, Krasnodin D, Cheporov S, Malkhos AN, Manikhas GM, Clawsaw A, et al. Significantly longer progression-free survival with nab-paclitaxel compared with docetaxel as first-line therapy for metastatic breast cancer. J Clin Oncol. 2009;27(23):3611–19.
13. Gradishar WJ, Krasnodin D, Cheporov S, Malkhos AN, Manikhas GM, Clawsaw A, et al. Phase II trial of nab-paclitaxel compared with docetaxel as first-line chemotherapy in patients with metastatic breast cancer: final analysis of overall survival. Clin Breast Cancer. 2012;12(5):S13–21.
14. Roy V, LaPlant BR, Gross G, Bane CL, Palmieri FM. For the North Central Treatment Group. Phase II trial of weekly nab (nanoparticle albuminbound)-paclitaxel (nab-paclitaxel) (Abraxane) in combination with gemcitabine in patients with metastatic breast cancer (N0531I). Ann Oncol. 2009;20(3):449–53.
15. Schwartzberg LS, Arena FP, Minterz KM, Epperson AL, Walker MS. Phase II multicenter trial of albumin-bound paclitaxel and carboplatin in first-line treatment of patients with metastatic breast cancer. Clin Breast Cancer. 2012;12(6):87–93.
16. Seidman AD, Conlin AK, Bach A, Moynahan ME, Lake D, Fororo A, et al. Randomized phase II trial of weekly vs. every 2 weeks vs. every 3 weeks nanoparticle albumin-bound paclitaxel with bevacizumab as first-line chemotherapy for metastatic breast cancer. Clin Breast Cancer. 2013;13(4):239–46.e1.
17. Rugo HS, Barry WT, Moreno-Aspita A, Lys A, Connine C, Leung E, et al. Randomized phase III trial of paclitaxel once per week compared with nanoparticle albumin bound nab-paclitaxel once per week or bevacizumab as first-line chemotherapy for locally recurrent or metastatic breast cancer: CALGB 40502/NCTCG N063H (Alliance). J Clin Oncol. 2015;33:2361–9.
18. Mitschang B, Cosgiff T, Harker G, Keaton M, Chidac T, Min M. A phase II study of weekly nanoparticle albumin-bound paclitaxel with or without trastuzumab in metastatic breast cancer. Breast Cancer. 2011;11(2):121–8.
19. Yardley DA, Hart L, Bosserman L, Salleh MN, Waterhouse DM, Hagan MK, et al. Phase II study evaluating lapatinib in combination with nab-paclitaxel in HER2-overexpressing metastatic breast cancer patients who have received no more than one prior chemotherapy regimen. Breast Cancer Res Treat. 2013;137(2):457–64.
20. Chirquin J, Chu SL. Management of breast cancer with nanoparticle albumin-bound (nab)-paclitaxel combination regimens: a clinical review. Breast. 2011;20(5):394–406.
21. Montero AJ, Adams B, Diaz-Montero CM, Gucluck S. nab-Paclitaxel in the treatment of metastatic breast cancer: a comprehensive review. Expert Rev Clin Pharmacol. 2011;4(3):329–34.
22. Onitilo AA, Engel JM, Greenlee RT, Mukes BN. Breast cancer subtypes based on ER/PR and Her2 expression: comparison of clinicopathologic features and survival. Clin Med Res. 2009;7(1):2–4–13.
23. The Cancer Genome Atlas Network. Comprehensive molecular portraits of human breast tumours. Nature. 2012;490(7418):61–70.
24. Rouzier R, Perou CM, Symmans WF, Ibrahim N, Cristofanilli M, Andersson K, et al. Breast cancer molecular subtypes respond differently to preoperative chemotherapy. Cancer Res. 2005;65(16):5678–85.
25. Esserman LJ, Berry DA, De Michele A, Carey L, Davis SE, Buxton M, et al. Pathologic complete response predicts recurrence-free survival more effectively by cancer subset: results from the I-SPY 1 TRIAL—CALGB 150007/150012, ACRN 6657. J Clin Oncol. 2012;30(26):3242–9.
26. Patt D, Liang C, Li L, Ko A, Fraser CQ, Corzo D, et al. Real-world efficacy and safety outcomes of nab-paclitaxel in patients with metastatic breast cancer: results from a US health insurance database. San Antonio: Poster presented at: San Antonio Breast Cancer Symposium; 2014. TX [P3-10-06].
27. O’Suaghhnessy J, Gradishar WJ, Bhar P, Iglesias J. nab-Paclitaxel for first-line treatment of patients with metastatic breast cancer and poor prognostic factors: a retrospective analysis. Breast Cancer Res Treat. 2013;138(3):629–37.
28. FDA approval for bevacizumab. http://www.cancer.gov/cancertopics/druginfo/fda-bevacizumab. Accessed September 5, 2014.
29. Yardley D, Brufsky A, Coleman R, Conte P, Cortes J, Gluck S, et al. tnAcity: a phase II/III trial of weekly nab-paclitaxel plus gemcitabine or carboplatin vs gemcitabine/carboplatin as first-line treatment for triple-negative metastatic breast cancer. Chicago, IL: Poster presented at: American Society of Clinical Oncology Annual Meeting; 2014. [TPS1146].