Ixabepilone: a new chemotherapeutic option for refractory metastatic breast cancer

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Abstract: Taxane therapy is commonly used in the treatment of metastatic breast cancer. However, most patients will eventually become refractory to these agents. Ixabepilone is a newly approved chemotherapeutic agent for the treatment of metastatic breast cancer. Although it targets microtubules similarly to docetaxel and paclitaxel, ixabepilone has activity in patients that are refractory to taxanes. This review summarizes the pharmacology of ixabepilone and clinical trials with the drug both as a single agent and in combination. Data were obtained using searches of PubMed and abstracts of the annual meetings of the American Society of Clinical Oncology and the San Antonio Breast Cancer Symposium from 1995 to 2008. Ixabepilone is a semi-synthetic analog of epothilone B that acts to induce apoptosis of cancer cells via the stabilization of microtubules. Phase I clinical trials have employed various dosing schedules ranging from daily to weekly to 3-weekly. Dose-limiting toxicities included neuropathy and neutropenia. Responses were seen in a variety of tumor types. Phase II studies verified activity in taxane-refractory metastatic breast cancer. The FDA has approved ixabepilone for use as monotherapy and in combination with capecitabine for the treatment of metastatic breast cancer. Ixabepilone is an efficacious option for patients with refractory metastatic breast cancer. The safety profile is similar to that of taxanes, with neuropathy and neutropenia being dose-limiting. Studies are ongoing with the use of both iv and oral formulations and in combination with other chemotherapeutic and biologic agents.

Keywords: ixabepilone, epothilone, metastatic breast cancer, taxane-refractory

Introduction
Although great strides have been made in the treatment of breast cancer, once metastatic, cure is unlikely. There are many agents available to treat metastatic breast cancer (MBC), and decisions on treatment must take into account the balance between efficacy and tolerability. Although, there are many new biological agents in clinical trials, chemotherapy is still an important backbone of the treatment. The recent approval of new agents has added to the repertoire available for the treatment of MBC (Gradishar et al 2005; Perez et al 2007; Thomas et al 2007).

The taxanes are the mainstays of chemotherapy for MBC. However, because all patients will eventually become refractory to these, new strategies are needed for taxane-refractory MBC. One drug that has shown efficacy both preclinically and clinically in patients resistant to docetaxel or paclitaxel is ixabepilone (BMS-247550). This drug fills an important void in chemotherapeutic options for patients.

Pharmacology of ixabepilone
Ixabepilone is in the epothilone class of chemotherapeutic drugs. Epothilones were initially discovered when fungicidal activity was observed by the myxobacterium *Sorangium cellolosum* and later these agents were found to interact with and stabilize microtubules (Bollag et al 1995; Gerth et al 1996). Microtubules help form...
the cytoskeleton and have vital roles in various cellular processes including cell signaling and mitosis. Microtubules are made up of 2 subunits, α and β. Microtubules have been an attractive target for many chemotherapeutic drugs, which exert their cytotoxic effects via microtubule stabilization or destabilization (Jordan et al 2004). Ixabepilone is a semi-synthetic analog of epothilone B, created by the substitution of an azide group for oxygen in the macrolide ring (Fumoleau et al 2007) (Figure 1). Similar to taxanes, epothilones exert their effect by binding to and subsequently polymerizing and stabilizing microtubules, thus preventing mitosis and resulting in apoptosis (Bollag et al 1995). Epothilones specifically bind to β-tubulin, although the exact binding site is not known (Giannakakou et al 2000). Despite the fact that epothilones bind close to or at the paclitaxel binding site on tubulin (Nogales et al 1995; Ojima et al 1999; Giannakakou et al 2000), there are also data from yeast studies to suggest that the binding of epothilones to tubulin and the subsequent microtubule formation may be distinct from that of paclitaxel (Bode et al 2002). Another important difference is that epothilones have been shown to have activity both preclinically and clinically in taxane resistance, likely due to the fact that the epothilones are not substrates for P-glycoprotein or MRP-1 and are active in β-tubulin mutations (He et al 2001a; Lee et al 2006). MDR-1 drug transporter expressing tumors are also sensitive to epothilones (McDaid et al 2002). In breast cancer cell lines and xenograft models, including those with multidrug resistance, ixabepilone has been shown to have cytotoxicity values ranging from 1.4 to 45 nM and has improved cytotoxicity over paclitaxel (Lee et al 2001, 2006). Both an iv and oral formulation of ixabepilone have undergone clinical studies, although it is the iv form that has been approved for the treatment of breast cancer. Further trials with the oral formulation are awaited.

Cell lines resistant to epothilones have been reported (Giannakakou et al 2000; He et al 2001b). The resistance is felt to be secondary to point mutations in the β-tubulin subunit, rather than MDR1 expression. These cells lines of various tumor types including ovarian and nonsmall-cell lung cancers have been to shown to exhibit resistance to other taxanes as well. It has been shown that these point mutations lead to less endogenous microtubule stabilization (He et al 2001a). These cell lines resistant to epothilones have also been found to have slower growth, which could be secondary to the endogenously more stable microtubules or could be due to an additional mechanism.

Pharmacokinetic studies with ixabepilone given on a daily dosing schedule have demonstrated a rapid 1-log decline after drug infusion, later followed by a prolonged elimination phase, not unlike taxanes (Abraham et al 2003).

**Figure 1** Chemical structures of the epothilones under clinical development. (A) Structure of epothilone B, ixabepilone, and BMS-310705. (B) Structure of epothilone D. (C) Structure of epothilone D second generation. From Fumoleau P, Coudert B, Isambert N, et al. 2007. Novel tubulin-targeting agents: anticancer activity and pharmacologic profile of epothilones and related analogues. Ann Oncol, 18(Suppl 5):v9–15, by permission of Oxford University Press © 2007.
The mean half-life of the drug was 16.8 hours. Clearance was rapid, dose-independent, and not associated with body weight or surface area. Dosing every 21 days showed a similar plasma concentration curve and independence of clearance, elimination, and dose (Mani et al 2004; Gadgeel et al 2005). The mean half-life was 38.5 hours at a dose of 40 mg/m² given every 21 days (Mani et al 2004). Pharmacodynamic studies with daily dosing did not show any correlation between area under the curve (AUC) and neutropenia (Abraham et al 2003). Similarly, there was no correlation with AUC and neutropenia in the 21-day dosing schedule (Mani et al 2004). However, the percentage change in absolute neutrophil count was significantly negatively correlated with increasing doses of ixabepilone (Gadgeel et al 2005). Pharmacodynamic studies exploring microtubule bundle formation and the subsequent effects on plasma ixabepilone concentration and extent of neutropenia have been performed (McDaid et al 2002; Mani et al 2007). Microtubule bundle formation is a surrogate for ixabepilone binding to β-tubulin and drug binding increases with greater ixabepilone concentration. This binding is correlated with AUC. Interestingly, it has been shown that there were more tumor cells with microtubule bundle formation than peripheral blood mononuclear cells after ixabepilone dosing (Mani et al 2004). The degree of neutropenia has been found to be correlated with the extent of microtubule bundle formation in peripheral blood mononuclear cells.

Clinical trials with ixabepilone
Phase I studies (Table 1)
Numerous phase I trials have evaluated the optimum dose and toxicity with ixabepilone. There have been numerous dosing schedules in both oral and iv formulations used in these trials. The earliest phase I trial was reported by Awada et al (2001) and employed a weekly iv dosing schedule. There were various tumor types in this study including breast cancer. At the time when preliminary results were presented, the current dose level was 30 mg/m²/week. Responses included stable disease in patients who were previously treated with a taxane with toxicities including fatigue, anorexia, arthralgia/myalgia, neuropathy, and myelosuppression. Another phase I study using weekly dosing has been presented (Burris et al 2002). In this study the maximum tolerated dose (MTD) was 25 mg/m²/week. Although there was minimal neutropenia, grade 3 toxicities included fatigue, nausea, diarrhea, myalgia/arthralgia, and neuropathy. This study initially used a 30-minute infusion, but was later amended to allow for a 1-hour infusion time, as well as a 3 weeks on, 1 week off dosing schedule in attempts to reduce neuropathy. Responses were seen in patients who had previously received taxane therapy. Hao et al (2002) explored continuous weekly dosing in another phase I trial. In this study, dose-limiting toxicities were grade 3 fatigue and grade 4 neutropenia at the 20 mg/m² and 30 mg/m² dose levels. Neuropathy was more frequent in patients who were heavily pretreated. Decreases in tumor markers were seen in taxane-refractory patients.

The use of 3-weekly dosing was also evaluated in the phase I setting. The earliest report was the preliminary data of Spriggs et al (2001). In this trial, a 1-hour infusion of ixabepilone was given to patients in dose levels ranging from 7.4 mg/m² to 65 mg/m². MTD was determined to be 50 mg/m². Dose-limiting toxicities included grade 3 arthralgias and myalgias, grade 3 neuropathy, grade 4 neutropenia, febrile neutropenia, pneumococcal sepsis, and 1 death. Antitumor activity was seen again in taxane pretreated patients and a complete response was seen in a patient with ovarian cancer. Another phase I study using a 1-hour infusion every 3 weeks also determined the MTD to be 50 mg/m² (Aghajanian et al 2007). In this study, the most common toxicities limiting dose were neutropenia, stomatitis, myalgia, and arthralgia. A 3-hour infusion time was also investigated in this trial and no dose-limiting toxicities (DLTs) were seen at doses less than 50 mg/m² when the drug was administered over 3 hours. Eight patients attained an objective response, many of whom had previously received taxane therapy. Mani et al (2004) performed a phase I study using 3-weekly dosing with 1-hour infusion times. The recommended phase II dose of ixabepilone was determined to be 40 mg/m². Similarly to previous trials, the dose-limiting toxicities included grade 4 and febrile neutropenia; however, grade 3 abdominal pain, nausea, and vomiting were also seen. A Japanese phase I trial using a 3-hour infusion every 3 weeks also determined the tolerable dose of ixabepilone to be 40 mg/m² (Shimizu et al 2008); DLT was grade 4 neutropenia. There were no objective responses, but over two-thirds of patients had stable disease. Linear pharmacokinetics were observed.

A dose of 40 mg/m² dose was again verified as the MTD in a phase I study in patients with refractory, advanced solid tumors and the DLT was neutropenia (Gadgeel et al 2005). Grade 1 and 2 neuropathy was observed in 3 out of 5 patients who received greater than 28.6 mg/m² for at least 2 cycles. Fatigue was also commonly seen. No patients met criteria for partial responses; however, there were minor responses and stable disease was seen. Pharmacokinetics were linear and there was evidence to suggest that there was a correlation...
| Study                | Dose/dosing schedule | Maximum tolerated dose | Dose-limiting or grade 3/4 toxicities | Activity                                                                 |
|---------------------|----------------------|------------------------|--------------------------------------|--------------------------------------------------------------------------|
| Awada et al 2001    | Weekly               | Not yet determined; Enrolling at 30 mg/m²/week | Grade 3 sensory neuropathy           | Stable disease seen in multiple tumor types                               |
| Burris et al 2002   | Weekly               | 25 mg/m²/week          | Cumulative sensory neuropathy; Grades 3/4 fatigue, nausea, diarrhea, myalgia/arthritis | Partial and minor responses seen in multiple tumor types                  |
| Hao et al 2002      | Weekly               | Enrolling at 20 mg/m²/week in heavily pretreated patients and 30 mg/m²/week in minimally pretreated patients | Grade 3 fatigue, Grade 4 neutropenia | Minor response in melanoma patient, tumor marker decline in taxane-refractory ovarian and prostate cancer patients |
| Spriggs et al 2001  | Q 3 weeks; 1-hour infusion | 50 mg/m² | Grade 3 neuropathy, Grade 4 neutropenia with sepsis and death, Grade 3 myalgia/arthritis | Complete response in patient with paclitaxel-refractory ovarian cancer, also partial responses and stable disease seen in multiple tumor types |
| Mani et al 2004     | Q 3 weeks; 1-hour infusion | 50 mg/m²; recommended phase II dose 40 mg/m² | Sepsis and death, Grade 4 neutropenia, Grade 3 abdominal pain/nausea/vomiting, Grade 3 neuropathy | 4 patients with objective partial and minor responses, responses include taxane-refractory disease |
| Gadgeel et al 2005  | Q 3 weeks; 1-hour infusion | 40 mg/m² | Grade 4 neutropenia, neutropenic sepsis, grade 3 nausea/emesis, grade 4 myalgias, grade 3 fatigue, grade 4 diarrhea, grade 3 and 4 thrombocytopenia | Tumor shrinkage not meeting criteria for partial response, stable disease in multiple tumor types |
| Shimizu et al 2008  | Q 3 weeks; 3-hour infusion | 40 mg/m² | Grade 4 neutropenia, grade 3 nausea, vomiting, and dehydration | Stable disease |
| Aghajanian et al 2007 | Q 3 weeks; 1-hour and 3-hour infusion | 50 mg/m² | Neutropenic infection and death, death from pneumonia and sepsis, grade 3 sensory neuropathy, grade 3/4 nausea/vomiting, grade 3/4 fatigue, grade 3/4 myalgia/arthritis | Complete and partial responses in multiple tumor types and taxane-pretreated patients |
| Thambi et al 2003   | Daily for 3 days every 21 days | Recommended phase II dose: 8 mg/m²/day x 3 days | Grade 3 leukopenia, grade 4 neutropenia, grade 3 thrombocytopenia, grade 3 nausea, grade 3/4 vomiting, grade 3 anorexia | Stable disease, nonconfirmed tumor shrinkage |
| Abraham et al 2003  | Daily for 5 days every 21 days | Recommended phase II dose: 6 mg/m²/day x 5 days | Grade 4 neutropenia, grade 3 fatigue, grade 3 mucositis, grade 3 anorexia | Partial responses including taxane-refractory patients, reduction in CA-125 in ovarian cancer patients |
| Zhuang et al 2005   | Daily for 3 days every 21 days | Recommended phase II dose: 8-10 mg/m²/day x 3 days | Grade 4 neutropenia, grade 3 fatigue, grade 3 hyponatremia, grade 3 anorexia, grade 3 ileus, grade 3 stomatitis, grade 3 emesis | Prolonged stable disease in multiple tumor types |
between the change in neutrophil count and the time at which the ixabepilone plasma concentration was > 15 ng/mL.

In attempts to augment the degree of neurotoxicity observed with 3-weekly dosing, daily administration of ixabepilone was evaluated in the phase I setting. Less neuropathy has been observed in daily compared with 3-weekly dosing. Abraham et al (2003) examined a dosing schedule of treatment daily for 5 consecutive days, with 21-day dosing intervals (Abraham et al 2003). Ixabepilone was given as a 1-hour infusion. Most patients had received prior taxane therapy and half had received ≥ 6 prior lines of treatment. MTD was 6 mg/m². Higher doses (8 mg/m²) were complicated with neutropenia despite filgrastim support. Neurotoxicity was generally mild and no grade 3 or 4 neuropathy was observed. Five out of 27 patients enrolled achieved a partial response, and these patients had received prior taxane therapy. Preliminary data presented explored a daily dosing schedule given every 3 days where the recommended phase II dose was 8 mg/m² for 3 days, with cycles every 21 days (Thambi et al 2003). A published report of a daily dosing schedule for 3 days also determined the MTD to be at a dose of 8 mg/m²/day. This was given as a 1-hour infusion and similarly to the Abraham study, the DLT was neutropenia and there were no cases of dose-limiting neurotoxicity (Zhuang et al 2005). Seventeen of the 26 enrolled patients had received prior taxane therapy and the median number of prior therapies received was ≥ 4. Although no responses were observed in this trial, patients with renal cell carcinoma, ovarian cancer, and primary peritoneal mesothelioma had prolonged stable disease up to 28 months.

Numerous phase I studies have also evaluated the toxicities associated with ixabepilone given in combination with other chemotherapeutic and biologic agents (Plummer et al 2002; Anderson et al 2004; Bunnell et al 2006; Hensley et al 2007). Preliminary results of a study exploring the use of ixabepilone in combination with carboplatin revealed partial responses in patients with breast cancer as well as neuroendocrine cancer (Plummer et al 2002). Some patients also achieved stable disease. Patients were relatively chemotherapy-naïve as only up to 2 prior therapies were allowed. DLT was reached at 40 mg/m² of ixabepilone and an AUC of 5 for carboplatin. A phase I/II study conducted in patients with MBC evaluated the combination of ixabepilone with capecitabine (Bunnell et al 2006). Eligible patients had received prior taxane and anthracycline-based treatment in the adjuvant or metastatic but could not have received greater than 3 prior regimens for metastatic disease. Two treatment schedules were evaluated; a single 3-hour infusion or daily therapy given over 1 hour for 3 days every 21 days. The recommend phase II dose for ixabepilone was determined to be 40 mg/m² as a single infusion and for capecitabine was 2000 mg/m² given on days 1–14 every 3 weeks. Nearly half of the patients had 2 or more therapies in the metastatic setting. Fifteen of 50 enrolled patients achieved a complete (1 patient) or partial response (14 patients) yielding an overall response rate of 30%. Responses had a median duration of nearly 7 months.

**Phase II studies in breast cancer (Table 2)**

Based on ample evidence of response and safety in early trials, various phase II studies were designed for patients with varying levels of previous therapies. Three published phase II trials used a daily dosing schedule. Low et al (2005) used a 6 mg/m²/day dosing schedule on days 1–5 every 3 weeks in patients with taxane-refractory breast cancer. A total of 37 patients were enrolled and 43% had received between 3 and 9 prior chemotherapy regimens in the metastatic setting. The objective response rate (ORR) was 22% and patients received a median of 4 cycles (range 1–11 cycles) with a median time to progression of 80 days for all subjects. An additional 35% of patients had stable disease for at least 6 weeks. Toxicities were generally mild. Twelve patients required dose reductions secondary to toxicities including neuropathy, diarrhea, fatigue, neutropenia, and myalgia. Only 1 patient developed grade 3 neuropathy, although mild neuropathy (grades 1 or 2) was frequent (83% of total subjects).

Another phase II study examined a daily dosing schedule for 5 consecutive days at a dose of 6 mg/m²/day given every 21 days (Denduluri et al 2007a). Patients in this trial were taxane-naïve; however, any number of non-taxane prior therapies was allowed. Of the 23 patients enrolled, 70% had prior chemotherapy, which was primarily adjuvant anthracycline. ORR was 57%. Stable disease for at least 6 weeks was achieved in 26%. Median time to progression was 5.5 months, and for partial responders, the median duration of response was 5.6 months. Four patients had toxicity necessitating removal from study (grade 3 weight loss, grade 3 motor neuropathy, prolonged autonomic neuropathy, and grade 2 fatigue) and dose reductions occurred in 4 patients (prolonged neutropenia, neuropathy, and fatigue). Severe neuropathy was infrequent with 13% having grade 2 sensory neuropathy, 4% grade 2 motor neuropathy, no patients with grade 3 sensory neuropathy, and 1 patient with grade 3 motor neuropathy. The same group performed a small study using daily dosing but with 3 consecutive days of therapy at initial...
doses of 8 mg/m²/day which was titrated up to 10 mg/m²/day if tolerated (Denduluri et al 2007b). Of the 12 patients enrolled, all had received previous taxanes. Median number of previous therapies in the metastatic setting was 3.5. The treatment had an acceptable safety profile; however, no complete or partial responses were seen and the study was stopped. Possible explanations for the lack of response may be decreased dose intensity in comparison with other studies with daily dosing.

There have also been numerous phase II studies with 3-weekly dosing in patients with MBC, in both taxane-naïve and taxane-refractory patients. Perez et al (2007) examined the efficacy of ixabepilone in a heavily pretreated patient population which had received prior anthracycline, taxane, and capecitabine therapies. Dose was 40 mg/m² given as a 3-hour infusion every 3 weeks. This was the third line of treatment in the metastatic setting for 88% of the 126 patients enrolled. An independent radiology facility assessed ORR at

| Study | Number of patients | Dosing schedule | Objective response rate | Survival endpoints | Selected grade 3/4 toxicities (%) |
|-------|--------------------|-----------------|-------------------------|--------------------|---------------------------------|
| Low et al 2005; Prior taxane | 37 | 6 mg/m²/day on days 1–5 every 3 weeks | 22% (95% CI, 9.8%–38.2%) | Median time to progression 80 days | Neutropenia (35%) Febrile neutropenia (14%) Thrombocytopenia (8%) Fatigue (13%) Sensory neuropathy (3%) Myalgia/arthralgia (3%) Nausea (5%) Constipation (11%) Diarrhea (5%) |
| Denduluri et al 2006; Prior taxane | 12 | 8 mg/m²/day on days 1–3 every 3 weeks (increased to 10 mg/m²/day if no toxicity) | No objective responses; 10 patients with stable disease for ≥6 weeks | Median time to progression 2.7 months | Neutropenia (17%) Thrombocytopenia (8%) Allergic reaction (8%) |
| Denduluri et al 2007; No prior taxane | 23 | 6 mg/m²/day on days 1–5 every 3 weeks | 57% (95% CI 34.5%–76.8%) | Median time to progression 5.5 months | Neutropenia (22%) Thrombocytopenia (4%) Myalgia/arthralgia (4%) Diarrhea (4%) Fatigue (13%) Motor neuropathy (4%) Nausea (9%) |
| Perez et al 2007; Prior taxane | 126 | 40 mg/m² every 21 days | Independent radiology facility 11.5% (95% CI 6.3%–18.9%) Investigator assessed 18.3% (95% CI 11.9%–26.1%) | Median progression-free survival 3.1 months; median overall survival 8.6 months | Neutropenia (54%) Febrile neutropenia (3%) Thrombocytopenia (8%) Sensory neuropathy (14%) Fatigue/asthenia (14%) Myalgia/arthralgia (8%) Nausea (2%) |
| Thomas et al 2007; Prior taxane | 49 | 40 mg/m² every 3 weeks | 12% (95% CI 4.7%–26.5%) | Median time to progression 2.2 months, median survival 7.9 months | Febrile neutropenia (6%) Sensory neuropathy (12%) Fatigue (27%) Nausea/vomiting (26%) Myalgia/arthralgia (12%) |
| Roche et al 2007; Prior adjuvant taxane allowed if ≥1 year since treatment | 93 | 40 mg/m² every 3 weeks | 41.5% (95% CI 29.4%–54.4%) | Median time to progression 9.3 months, median survival 29.9 months | Neutropenia (58%) Febrile neutropenia (6%) Severe neuropathy (20%) Motor neuropathy (5%) Myalgia/arthralgia (13%) Nausea/vomiting (7%) |
of patients receiving daily dosing (Vansteenkiste et al 2007). The toxicity profile was acceptable and included a 6% rate of grade 3 sensory neuropathy, and neutropenia was seen in 28% of patients receiving 3-weekly dosing and in 17% of patients receiving daily dosing (Vansteenkiste et al 2007).

Accepted response rates of 15%–48% with and without estramustine phosphate were seen in hormone-refractory prostate cancer. Grade 3 sensory neuropathy was seen in 13%–17% of patients and rates of grade 3/4 neutropenia were 17%–29% (Galsky et al 2005; Hussain et al 2005). Modest activity was also seen in patients with prostate cancer in the second-line setting who had progressed on first-line docetaxel; however, half the patients developed grade 3/4 neutropenia (Rosenberg et al 2007). Final as well as preliminary phase II data with promising activity have also been presented in renal cell cancer (Fojo et al 2005), gynecologic cancers (Chen et al 2004), hepatobiliary cancer (Singh et al 2003), gastric cancer (Ajani et al 2006), pancreatic cancer (Whitehead et al 2006), and non-Hodgkin’s lymphoma (O’Connor et al 2005; Smith et al 2005).

Despite some suggestions of phase I activity, there has not been any meaningful efficacy in phase II studies in melanoma, colorectal cancer, and sarcoma (Eng et al 2004; Pavlick et al 2004; Okuno et al 2005).

Activity in other tumor types
Based on the activity of ixabepilone in a wide variety of tumor types in phase I studies, a number of phase II studies were designed to assess disease specific activity. In patients with nonsmall-cell lung cancer refractory to platinum drugs, ORR was 14.3% with 3-weekly dosing and 11.6% with daily dosing for 5 days, which is similar to other second-line treatments in this disease. The toxicity profile was acceptable and included a 6% rate of grade 3 sensory neuropathy, and neutropenia was seen in 28% of patients receiving 3-weekly dosing and in 17% of patients receiving daily dosing (Vansteenkiste et al 2007).
common in the combination arm, and any neutropenia was frequent in that arm (89%). In the single-agent capecitabine arm 43% of patients had any level of neutropenia.

Neoadjuvant trials
Ixabepilone has been used in the neoadjuvant setting in breast cancer in addition to the metastatic setting. A preliminary report using a dose of 40 mg/m² every 21 days revealed a pathologic complete response rate (pCR) of 21% of patients. Approximately 50% of patients were ER+. Grade 3 or 4 neutropenia was seen in 21% and 13% of patients, respectively. Grade 2 neuropathy was seen in 11% of patients, whereas grade 3 neuropathy was seen in 2% (Baselga et al 2005). Correlative studies revealed a 6 gene predictive model that was able to predict benefit to ixabepilone (Llmobart-Cussac et al 2005). Other preliminary data suggest a higher response rate in the triple negative (ER, PR, and HER2 negative) subgroup of breast cancer patients, with a pCR rate of 26% of the 42 triple negative patients and 15% in the 119 other patients given neoadjuvant ixabepilone (Roche et al 2006).

Safety
As evidenced in phase I and II trials, neuropathy and neutropenia are important potential toxicities. The neuropathy is primarily sensory and generally reversible in most patients with proper dose reductions and drug discontinuation (Perez et al 2007; Thomas et al 2007). Patients with diabetes have been shown to be at increased risk for the development of neuropathy and it is important to note that because patients with pre-existing neuropathy were excluded from clinical trials, their potential severity of neuropathy is unknown. Numerically it appears that daily dosing (Abraham et al 2003; Low et al 2005; Denduluri et al 2007a; Denduluri et al 2007b) may be associated with less severe neuropathy (grade 3 neuropathy in 3%-4%) than in 3-weekly dosing (grade 3 in 13%-23%) (Perez et al 2007; Roche et al 2007; Thomas et al 2007). However, a study in nonsmall-lung cancer did not show any differences in neuropathy between 3-weekly and daily dosing (Vansteenkiste et al 2007). The use of specific neurologic function tests to characterize the neuropathy associated with ixabepilone has been described (Lee et al 2006). The patients enrolled in a monotherapy trial of ixabepilone in breast cancer (Low et al 2005) had a number of neurologic function tests performed, including Semmes-Weinstein monofilament testing and modified Romberg stance tests, which are used to assess diabetic neuropathy, and the Jebsen Test of Hand Function (JTH) and Grooved Pegboard Test (GPT), which are used to assess hand functions particularly in patients with stroke and rheumatoid arthritis. Twenty-three percent of patients developed ≥grade 2 neuropathy (including decreased hand function, decreased sensory function, paresthesias, and motor weakness), 9 patients developing grade 2 neuropathy and 2 patients developing grade 3 neuropathy. Three patients had refractory neuropathy that did not resolve to grade 1 up to 2 years after development. The results of JTH and GPT scores were found to be identifiable with onset of grade 12 or higher peripheral neuropathy.

The most common overall adverse reactions seen in the monotherapy registration trial are summarized in Table 3 (Perez et al 2007). When combined with capecitabine, toxicities commonly encountered with capecitabine such as palmar-plantar erythrodysesthesia and diarrhea were not significantly increased. It is recommended that patients with known severe hypersensitivity to Cremophor® EL or derivatives, baseline ANC <1500 cells/mm³, platelets <100,000 cells/mm³, or abnormal liver function (transaminases >2.5 × ULN or bilirubin >1 × ULN) do not receive ixabepilone.

Conclusions
Ixabepilone is an important addition to chemotherapeutic options for patients with MBC. The activity in patients who have failed previous taxanes makes this drug applicable for virtually all patients with MBC at some time in the course of their treatment. The high level of activity in the first-line setting makes it an attractive option in that setting as well. The results of ongoing trials exploring the additional of various biologic agents, such as trastuzumab and bevacizumab, are eagerly awaited (Cancer.gov 2008a, b, c). The strategy of combining chemotherapy with antiangiogenic agents is increasingly being used especially in the first-line setting for

| Table 3 Nonhematologic adverse events seen in monotherapy registration trial |
|-----------------------------|-----------------|-----------------|
| Nonhematologic adverse event | Total (%) (n = 126) | Grade 3/4 (%) |
| Peripheral sensory neuropathy | 60 | 14 |
| Fatigue/asthenia | 56 | 13 |
| Myalgia/arthralgia | 49 | 8 |
| Alopecia | 48 | 0 |
| Nausea | 42 | 2 |
| Stomatitis/mucositis | 29 | 6 |
| Vomiting | 29 | 1 |
| Diarrhea | 22 | 1 |
| Musculoskeletal pain | 20 | 3 |
MBC and the potential for improved activity exists with the other novel microtubule agents (Miller et al 2007). The taxanes in particular may potentiate the anti-angiogenic effects of these agents (Miller et al 2001; Sweeney et al 2001). A phase III trial conducted by the Cancer and Leukemia Group B (CALGB) and the North Central Cancer Treatment Group (NCCTG) is currently underway comparing the activity of various microtubule stabilizing agents with the vascular endothelial growth factor antibody bevacizumab. This trial is comparing weekly paclitaxel (90 mg/m²), nab-paclitaxel (100 mg/m²), and ixabepilone (16 mg/m²) in combination with bevacizumab. This trial will also supply important information on the activity of these agents given in a “dose-dense” fashion, in which paclitaxel and arguably nab-paclitaxel have been found to have the greatest activity (Gradishar et al 2006; Seidman et al 2008; Sparano et al 2008). There are a number of other agents that target the angiogenic as well as other important pathways with recent promising data (Rugo et al 2007) (Burstein et al 2008), and these agents may provide other rational combinations with ixabepilone. With the advent of new and possibly improved taxane agents, the question will be in what setting are these agents best used and whether they would replace traditional taxanes as the preference for first-line therapy; or whether, perhaps because of activity in docetaxel- and paclitaxel-resistant patients, their activity may be best reserved for patient with refractory disease. The role of ixabepilone will be best ascertained when it is directly compared with the other available taxanes in a variety of dosing schedules and lines of therapy. These questions will be best answered in the setting of a phase III randomized trial such as the CALGB/NCCTG one described above.

Disclosures

Neither author has any financial disclosures related to any product discussed in this review.

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