Plitidepsin: design, development, and potential place in therapy

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Abstract: Plitidepsin is a cyclic depsipeptide that was first isolated from a Mediterranean marine tunicate (Aplidium albicans) and, at present, is manufactured by total synthesis and commercialized as Aplidin®. Its antitumor activity, observed in preclinical in vitro and in vivo studies has prompted numerous clinical trials to be conducted over the last 17 years, alone or in combination with other anticancer agents. Single-agent plitidepsin has shown limited antitumor activity and a tolerable safety profile in several malignancies, such as noncutaneous peripheral T-cell lymphoma, melanoma, and multiple myeloma. In patients with relapsed or refractory multiple myeloma, plitidepsin activity seems to be enhanced after addition of dexamethasone while remaining well tolerated, and a Phase III trial comparing plitidepsin plus dexamethasone vs dexamethasone alone is underway. Additional studies are required to better define the role of plitidepsin in combination with other active agents in these indications. Results of plitidepsin activity in other hematological malignancies or solid tumors have been disappointing so far. Further studies analyzing its mechanisms of action and potential biomarkers will help select patients who may benefit most from this drug. In this review, we critically analyze the published studies on plitidepsin in hematological malignancies and solid tumors and discuss its current role and future perspectives in treating these malignancies. We also review its design, pharmaceutical data, and mechanism of action.

Keywords: aplidin, plitidepsin, lymphoma, myeloma, melanoma

Design

Pharmaceutical data
Aplidin® (plitidepsin; PharmaMar, S.A., Madrid, Spain), the trade name for the active substance plitidepsin, is a synthetically produced anticancer agent. It is a cyclic depsipeptide that is structurally related to didemmins (Figure 1) and was originally isolated from a Mediterranean marine tunicate, Aplidium albicans. Its molecular formula is C₇₅H₁₂₁N₄O₃⁵ and it has a molecular weight of 1,110.34.¹⁻⁴

Plitidepsin was developed in the form of a lyophilized dosage form due to its substantial degradation under heat and light stress testing conditions when solubilized. The optimal reconstitution solution was found to be 15/15/70% (v/v/v) polyoxyl 35 castor oil/ethanol/water for injection. The reconstituted product and its dilutions with normal saline (up to 1:100 v/v) appeared to be stable for at least 24 hours after preparation.⁵

The plitidepsin drug product is supplied as a powder to be made into a concentrated solution for infusion at a concentration of 2 mg/vial. It is recommended that vials be reconstituted with 4 mL of the appropriate solution. The resulting solution must be clear and essentially free of visible particles. It may be colorless, but can turn slightly yellow. The reconstituted solution contains 0.5 mg/mL of plitidepsin (Table 1) and...
requires further dilution with 9 mg/mL sodium chloride (0.9%) or 50 mg/mL glucose (5%) infusion solution for administration as an intravenous infusion.

Plitidepsin and the reconstitution solution should be stored under the same conditions, between 2°C and 8°C. Once reconstituted, the solution is physically and chemically stable for 28 hours up to 25°C under normal light conditions. However, it is recommended that it be diluted and used immediately. After dilution, the solution is stable for only 24 hours at temperatures of up to 25°C under normal light conditions.

**Mechanism of action**

The mechanism of action of plitidepsin IN various tumor cell types has been reviewed already and is represented in Figure 2. Plitidepsin induces dose-dependent cell-cycle arrest and an acute apoptotic process. These effects rely on the induction of early oxidative stress, the rapid activation of Rac1 GTPase, and the sustained activation of c-Jun N-terminal kinase (JNK) and p38 mitogen-activated protein kinases (p38/MAPK), which finally result in caspase-dependent apoptosis. JNK phosphorylation can be seen as early as 5–10 minutes after exposure to the compound. The activation of JNK and p38/MAPK is associated with an increase in the number of reactive oxygen species and a reduction in GSH.

**Table 1** Composition of reconstituted solution

<table>
<thead>
<tr>
<th>Active substance</th>
<th>Amount per mL</th>
</tr>
</thead>
<tbody>
<tr>
<td>Plitidepsin</td>
<td>0.5 mg</td>
</tr>
<tr>
<td>Excipients</td>
<td></td>
</tr>
<tr>
<td>D-Mannitol</td>
<td>25.0 mg</td>
</tr>
<tr>
<td>Polyoxyl 35 castor oil</td>
<td>0.15 mL</td>
</tr>
<tr>
<td>Absolute ethanol</td>
<td>0.15 mL</td>
</tr>
<tr>
<td>Water for injection</td>
<td>0.70 mL</td>
</tr>
</tbody>
</table>

In addition, plitidepsin modifies the microenvironment that facilitates tumor growth. This was clearly observed in chronic lymphocytic leukemia (CLL) models. Plitidepsin not only directly affected the malignant B-CLL clone, but also had potent activity against monocytes and nurse-like cells (NLCs), a subset of cells differentiated from monocytes that favors leukemic cell progression. This effect is induced by triggering apoptosis, as evidenced by early exposure of phosphatidylserine in the outer leaflet of the plasma membrane, the activation of caspase-3, and the subsequent cleavage of PARP.

Antiangiogenic effects have been also reported for this drug. Broggin et al showed that plitidepsin blocks the secretion of the angiogenic factor vascular endothelial growth factor (VEGF) by the human leukemia cells MOLT-4, suggesting a possible effect on tumor angiogenesis. In another study, the same group showed that plitidepsin has antiangiogenic activity in vivo, in the chick embryo allantoic membrane assay, and inhibits endothelial cell functional responses to angiogenic stimuli in vitro. This effect might contribute to the antineoplastic activity of plitidepsin. In addition, a recent study have shown that the plitidepsin analogs PM01215 and PM02781 inhibit angiogenesis in vitro and in vivo.

Recent studies have led researchers to hypothesize that the primary target of plitidepsin could be the eukaryotic Elongation Factor 1A2 (eEF1A2), which is overexpressed in tumors, including multiple myeloma (MM) and plasmacytoma, prostate, pancreas, and ovarian cancers, and that it exhibits oncogenic behavior by favoring tumor cell proliferation while inhibiting apoptosis. eEF1A2 is therefore an interesting target for cancer treatment and may also be a biomarker predicting drug sensitivity. The existence of an interaction between plitidepsin and eEF1A2 is suggested by several findings: 1) plitidepsin interacts with GTP-bound eEF1A2 with a Kd in the nanomolar range, compatible with the concentrations at which the compound shows its proapoptotic effect in tumor cells; 2) eEF1A2 is the only protein retrieved from a tumor cell lysate through a plitidepsin-binding fractionation process; 3) eEF1A2 is commonly depleted in plitidepsin-resistant cells, and its restoration to normal levels resensitizes them to the compound. These results suggest that eEF1A2 is the primary target of plitidepsin, which acts by mediating its antitumor activity.

**Preclinical studies**

In vitro studies showed that plitidepsin has cytotoxic activity against a broad range of cell lines showing IC50 values

**Figure 1** Plitidepsin chemical structure.
of ≤1 nM. Its effects are particularly active against hematological malignancies such as MM, lymphoma and leukemia, and solid tumors including those of non-small-cell lung, pancreas, breast, melanoma, sarcoma, gastric, ovarian, bladder, and colon cancer. In addition, in vitro combination studies revealed a synergism of plitidepsin when combined with other agents like melphalan, dexamethasone, lenalidomide and bortezomib in MM cell lines.

In vivo studies showed antitumor effect of plitidepsin in xenograft models of MM (as a single agent or in combination with dexamethasone), T-cell lymphoma, diffuse large B-cell lymphoma (DLBCL), Burkitt lymphoma (alone or in combination with rituximab), and pancreatic cancer (synergistic antitumor activity in combination with gemcitabine). The preclinical toxicological profile of plitidepsin is characteristic of a cytotoxic drug for which the liver, gastrointestinal tract, spleen, and bone marrow are the main target organs.

Presently, the hydrophobic drug plitidepsin is formulated using Cremophor®, an adjuvant associated with unwanted hypersensitivity reactions. In search of alternatives, two nanoparticle-based formulations of plitidepsin have been tested in a renal xenograft tumor model, showing equivalent antitumor activity, compared to the Cremophor® formulation, but improved biodistribution profiles, presenting novel tools for future plitidepsin-based therapies. Also, in colorectal cancer cell lines, the use of epidermal growth factor receptor (EGFR) targeted polymersomes greatly reduced plitidepsin cytotoxicity as well as the cellular uptake, indicating that the use of this targeted nanocarrier is a promising approach to tackle colorectal cancer disease and avoid the undesired effects of the usual treatment.

**Pharmacokinetics and metabolism**

Several Phase I dose-escalation studies in subjects with advanced malignancies have described the pharmacokinetics of intravenous plitidepsin used as a single agent. Doses ranging from 0.13 to 8.00 mg/m² were administered as a 1- or 24-hour infusion weekly, 3- or 24-hour infusion biweekly or, 1-hour infusion on 5 consecutive days every 3 weeks. In these studies, linear and time-independent pharmacokinetics in plasma were observed after several cycles of administration.

After intravenous administration, plitidepsin was found to have a relatively long half-life, low clearance, and a high volume of distribution in plasma. In plasma and whole blood, mean half-lives ranged from 21 to 44 hours. Plitidepsin is widely distributed, with apparent volumes of distribution at steady state (Vss) ranging from 500 to 1,350 L. Approximately 97.3% of the human plasma protein was bound.
Mean clearance values in plasma range from 45 to 49 L/h. Urinary excretion of the unchanged compound is a minor elimination route. Preliminary results of metabolism studies indicated that plitidepsin undergoes moderate microsomal-mediated metabolism in several animal species and humans.

**Development**

**Phase I trials**

Phase I clinical trials with plitidepsin were started in 1998. These studies, summarized in Table 2, were conducted according to classic Phase I study design standards. Fortnightly, weekly, or daily (days 1–5) schedules were used. Included patients were diagnosed with various solid tumors or non-Hodgkin lymphoma with relapsed or refractory disease. Plitidepsin was evaluated as a single agent in six trials (one of them in children, in which it consistently showed limited efficacy (Table 2). Overall, in 252 patients, two confirmed partial responses (PRs) and one unconfirmed PR were found from all Phase I studies of plitidepsin. Clinical benefit, defined as stable disease (SD) lasting more than 3 months (time to progression [TTP] >3 months) was observed in 30 adult patients (range of clinical benefit, 3.0–9.7 months) and six children (range, 3.5–13.0 months). APL-A-005-02 was a multicenter, open-label, Phase I–II study to determine the maximum tolerated dose and recommended dose of plitidepsin as a single agent in 38 children with refractory or relapsed malignant tumors. Most (n=31) discontinued treatment due to disease progression, but six children showed clinical benefit (Table 2).

Plitidepsin in combination with other antineoplastic drugs has been evaluated in six Phase I studies (Table 2). Plitidepsin plus carboplatin (APL-A-006-05) resulted in an SD >3 months (range, 3.0–7.1 months) in six of 20 patients with advanced malignant solid tumors or lymphoma. Plitidepsin in combination with cytarabine has been tested in a Phase I/II trial in patients with relapsed/refractory leukemia, but the study was concluded early due to low recruitment. Plitidepsin and dacarbazine (Phase I stage of APL-B-016-05) resulted in one confirmed PR, two unconfirmed PRs, and four SDs >3 months in 28 patients with unresectable advanced melanoma as frontline therapy. Bortezomib and dexamethasone have been added to plitidepsin in a Phase I/II trial (APL-A-012-13) in patients with relapsed and/or refractory MM. The recruitment has been completed (N=20) and the results are encouraging, with 55% overall response rate (ORR) in a heavily pretreated population in which 88% were bortezomib exposed and 61% were refractory to the last line of therapy. In addition, toxicity profile was manageable, so the Phase II is ongoing.

**Phase II trials**

Results of plitidepsin Phase II trials are summarized in Table 3. Two different schedules were used, based on the findings from previous Phase I studies. The most common scheme, which was used in 10 of the existing Phase II trials, was the fortnightly scheme, consisting of plitidepsin 5 mg/m² administered as a 3-hour intravenous infusion on days 1 and 15, every 4 weeks. A 24-hour infusion was used in the first Phase II study performed (APL-B-001-01), with a higher dose (up to 7 mg/m²) of plitidepsin as a single agent or in combination with L-carnitine. The second scheme was the weekly schedule, used in four Phase II studies, consisting of plitidepsin 3.2 mg/m² administered as a 1-hour intravenous infusion on days 1, 8, and 15, every 4 weeks. One study (APL-B-016-05) used lower doses of plitidepsin (2.4 mg/m²) and was given in combination with dacarbazine at a dose of 800 mg/m².

With respect to hematological malignancies, a Phase II trial (APL-B-020-10) of 11 patients with primary or secondary myelofibrosis was concluded early due to lack of efficacy. Antitumor activity seems to be greater in lymphoid neoplasms. Fifty one patients with relapsed or refractory MM were included in a Phase II trial (APL-B-014-03) with plitidepsin at 5 mg/m² as a 3-hour intravenous (IV) infusion every 2 weeks. The protocol was amended to allow patients with a suboptimal response to single-agent plitidepsin to add 20 mg/d of oral dexamethasone on days 1–4, every 2 weeks. From 47 patients evaluable for efficacy, the ORR (complete remission plus PR plus minimal response) was 13% with plitidepsin alone, and 22% in the cohort of patients with the addition of dexamethasone (n=19, 18 evaluable). Plitidepsin alone and with dexamethasone were feasible and well tolerated in this heavily pretreated population. The activity observed with plitidepsin after dexamethasone addition merits further study.

In another Phase II trial (APL-B-013-02), 67 patients with relapsed or refractory aggressive non-Hodgkin lymphoma received plitidepsin according to the weekly scheme. Patients were divided into two cohorts: those with noncutaneous peripheral T-cell lymphoma (PTCL) (n=34) who received a median of two cycles (range, 1–8 cycles) per patient, and those with other lymphomas (n=33) who received a median of 1.5 cycles (range, 1–4 cycles). Efficacy was evaluated using the International Working Group criteria (1999). Of the 29 evaluable patients with noncutaneous PTCL, six had a response (ORR 20.7%), including two complete remissions (both angioimmunoblastic T-cell lymphomas pretreated with...
Table 2 Phase I studies of plitidepsin as a single-agent or in combination

<table>
<thead>
<tr>
<th>Design and histology</th>
<th>Schedule (infusion)</th>
<th>N</th>
<th>TTP &gt; 3 months (range of TTP in months)</th>
<th>Response</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Single-agent studies</strong></td>
<td></td>
<td></td>
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<tr>
<td>APL-A-001b-98</td>
<td>Solid tumors and lymphoma</td>
<td>Plitidepsin d1,15 q4wk (3-h IV)</td>
<td>27</td>
<td>5 (3.5–8.8)</td>
</tr>
<tr>
<td>APL-A-003-98</td>
<td>Solid tumors and lymphoma</td>
<td>Plitidepsin d1,15 q4wk (24-h IV)</td>
<td>67</td>
<td>10 (4.4–6.7)</td>
</tr>
<tr>
<td>APL-A-001a-98</td>
<td>Solid tumors and lymphoma</td>
<td>Plitidepsin d1,8,15 q4wk (24-h IV)</td>
<td>35</td>
<td>7 (3.9–5.9)</td>
</tr>
<tr>
<td>APL-A-002-98</td>
<td>Solid tumors and lymphoma</td>
<td>Plitidepsin d1,8,15 q4wk (1-h IV)</td>
<td>48</td>
<td>3 (3–3.8)</td>
</tr>
<tr>
<td>APL-A-004-98</td>
<td>Solid tumors and lymphoma</td>
<td>Plitidepsin d1–5 q3wk (1-h IV)</td>
<td>37</td>
<td>8 (3.9–9.7)</td>
</tr>
<tr>
<td>APL-A-005-02</td>
<td>Refractory or relapsed malignant tumors (children)</td>
<td>Plitidepsin d1,15 q4wk (3-h IV)</td>
<td>38</td>
<td>6 (3.5–13)</td>
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<tr>
<td><strong>Combination studies</strong></td>
<td></td>
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<tr>
<td>APL-A-006-05</td>
<td>Plitidepsin + carboplatin, in subjects with advanced malignant solid tumors or lymphoma</td>
<td>Plitidepsin d1,8,15 q4wk (1-h IV) + carboplatin d1 q4wk (1-h IV)</td>
<td>20</td>
<td>6 (3–7.1)</td>
</tr>
<tr>
<td>APL-A-007-06</td>
<td>Phase I/II study of plitidepsin + cytarabine in patients with relapsed/refractory leukemia</td>
<td>Plitidepsin d1,15 q4wk (3-h IV) + cytarabine d1–5 q4wk (2-h IV)</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>APL-A-008-08</td>
<td>Plitidepsin + sorafenib or plitidepsin + gemcitabine in patients with advanced solid tumors or lymphomas</td>
<td>Plitidepsin d1,8,15 q4wk (1-h IV) + gemcitabine d1,8,15 q4wk (30-min IV). Plitidepsin d1,8,15 q4wk (1-h IV) + sorafenib po bid continuous</td>
<td><a href="https://clinicaltrials.gov/ct2/show/NCT00788099">https://clinicaltrials.gov/ct2/show/NCT00788099</a></td>
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<tr>
<td>APL-A-01-08</td>
<td>Plitidepsin + bevacizumab in patients with advanced and refractory solid tumors</td>
<td>Plitidepsin d1,15 q4wk (3-h IV) + bevacizumab d1,15 q4wk (1.5-h IV)</td>
<td>13</td>
<td>TTP =9 (3.5)</td>
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<td></td>
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<td></td>
<td>TTP =5.2 months</td>
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<td></td>
<td></td>
<td></td>
<td>TTP =3.2 months</td>
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Table 2 (Continued)

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<tr>
<th>Design and histology</th>
<th>Schedule (infusion)</th>
<th>N</th>
<th>TTP &gt;3 months (range of TTP in months)</th>
<th>Response</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phase I–II, Plitidepsin ± dacarbazine, as frontline therapy to subjects with unresectable advanced melanoma</td>
<td>Plitidepsin d1,8,15 q4wk (1-h IV) + DTIC d1 q4wk (1-h IV)</td>
<td>28</td>
<td>7</td>
<td>1 PR + 2 PRu + 4 SD</td>
</tr>
<tr>
<td>Phase I–II, Plitidepsin in combination with ortezomib and dexamethasone in patients with relapsed and/or refractory MM</td>
<td>Plitidepsin d1,15 q4wk (3-h IV) + Bortezomib d1,4,8,11 q4wk (3–5 sec bolus SC injection) + dexamethasone d1,8,15,22 q4wk (oral, 1-h before plitidepsin infusion)</td>
<td>20</td>
<td>Median PFS: 8.3 months; median DOR 6 months</td>
<td>56% ORR, including 33% vGPR</td>
</tr>
</tbody>
</table>

Notes: *Studies terminated early following decision by sponsor, *Ongoing clinical trials or unpublished results.

Abbreviations: d, day; DOR, duration of response; h, hour; IV, intravenous; MM, multiple myeloma; mo, months; po, orally; PR, partial response; ORR, overall response rate; PFS, progression-free survival; PRu, unconfirmed partial response; q4wk, every 4 weeks; SC, subcutaneous; SD, stable disease; TTP, time to progression; vGPR, very good partial response; q3wk, every 3 weeks; DTI, dacarbazine.

Table 3 Phase II studies of plitidepsin as single-agent or in combination

<table>
<thead>
<tr>
<th>Diagnosis</th>
<th>Schedule</th>
<th>N (evaluable)</th>
<th>Median survival (months)</th>
<th>Response</th>
</tr>
</thead>
<tbody>
<tr>
<td>Renal cancer</td>
<td>Plitidepsin 5 mg/m² (24-h IV) d1,15 q4wk</td>
<td>19</td>
<td>PFI =2.1; OS =7.0</td>
<td>Renal cancer: 2 PR, 1 SD</td>
</tr>
<tr>
<td>Medullary thyroid carcinoma</td>
<td>Plitidepsin 5 mg/m² (3-h IV) d1,15 q4wk</td>
<td>16</td>
<td>TTP =5.3; PFS =5.5</td>
<td>11 SD</td>
</tr>
<tr>
<td>Non-small-cell lung cancer</td>
<td>Plitidepsin 5 mg/m² (3-h IV) d1,15 q4wk</td>
<td>17</td>
<td>TTP/PFS =1.2; OS =4.3</td>
<td>3 SD</td>
</tr>
<tr>
<td>Transitional cell carcinoma of the urothelium</td>
<td>Plitidepsin 5 mg/m² (3-h IV) d1,15 q4wk</td>
<td>18</td>
<td>TTP/PFS =1.4; OS =2.3</td>
<td>1 SD</td>
</tr>
<tr>
<td>Melanoma</td>
<td>Plitidepsin 5 mg/m² (3-h IV) d1,15 q4wk</td>
<td>35</td>
<td>TTP/PFS =1.3; OS =3.5</td>
<td>2 PR, 3 SD</td>
</tr>
<tr>
<td>Prostate adenocarcinoma</td>
<td>Plitidepsin 5 mg/m² (3-h IV) d1,15 q4wk</td>
<td>47</td>
<td>NA</td>
<td>2 PR, 4 MR (IWG-MRT)</td>
</tr>
<tr>
<td>Myelofibrosis (primary and secondary)</td>
<td>Plitidepsin 5 mg/m² (3-h IV) d1,15 q4wk</td>
<td>12</td>
<td>PFS =4.6</td>
<td>1 clinical improvement</td>
</tr>
<tr>
<td>Liposarcoma</td>
<td>Plitidepsin 5 mg/m² (3-h IV) d1,15 q4wk</td>
<td>14</td>
<td>PFS =1.6; OS =9.2</td>
<td>2 SD</td>
</tr>
<tr>
<td>Small-cell lung cancer</td>
<td>Plitidepsin 3.2 mg/m² (1-h IV) d1,8,15 q4wk</td>
<td>17</td>
<td>TTP/PFS =1.3; OS =4.8</td>
<td>1 SD</td>
</tr>
<tr>
<td>Indolent NHL</td>
<td>Plitidepsin 3.2 mg/m² (1-h IV) d1,8,15 q4wk</td>
<td></td>
<td></td>
<td><a href="http://www.clinicaltrialsregister.eu/ctr-search/trial/2004-002692-16/ES">http://www.clinicaltrialsregister.eu/ctr-search/trial/2004-002692-16/ES</a></td>
</tr>
<tr>
<td>Drug</td>
<td>Dosage (mg/m²)</td>
<td>Schedule</td>
<td>Outcome</td>
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<tr>
<td>Plitidepsin</td>
<td>2.4, 3.2, 5.0</td>
<td>q4wk</td>
<td>PR, MR (iwG-MRT)</td>
<td></td>
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<tr>
<td></td>
<td></td>
<td></td>
<td>Renal or advanced lung cancer</td>
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<td></td>
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<td></td>
<td>PTCL: CR, PR</td>
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<td></td>
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<td>Melanoma: PR, SD</td>
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</tbody>
</table>

**Note:** Ongoing clinical trials or unpublished results.

**Abbreviations:** APL, acute promyelocytic leukemia; MM, multiple myeloma; NHL, non-Hodgkin lymphoma; OS, overall survival; PFS, progression-free survival; PTCL, peripheral T-cell lymphoma.

**Combination studies**

- Plitidepsin 2.4 mg/m² (1-h iv) (d1,8,15 q4wk) + dacarbazine 800 mg/m² d1 q4wk (1-h iv)
  - Median PFS 4.3 months in those with baseline normal LDH. No responses with plitidepsin/DTIC was 3.3 months in all patients and 4.2 months in responders with normal serum lactate dehydrogenase (LDH) levels and performance status ≤1 at baseline. Median PFS with plitidepsin/DTIC was 3.3 months in all patients and 4.3 months in those with baseline normal LDH. No responses occurred with single-agent plitidepsin – median PFS was 1.5 months. Both regimens were well tolerated. Further evaluation of plitidepsin 2.4 mg/m² fortnightly and DTIC 800 mg/m² q4wk was recommended by the authors.32

Plitidepsin as a single agent has shown very limited or no antitumor activity in other solid tumors. Results from other malignancies comprised: two PRs and one SD among 19 patients with unresectable advanced renal cell carcinoma, with a 2.1-month progression-free interval (PFI) (APL-B-005-02).11 11 SDs out of 16 patients with unresectable advanced medullary thyroid carcinoma, with a clinical benefit noted in only one patient (APL-B-002-02);21 three SDs in 17 patients with locally advanced or metastatic non-small-cell lung cancer, with a TTP of 1.2 months (APL-B-004-02);43 no responses in another trial of 19 patients with small-cell lung cancer pretreated with one line (APL-B-006-02);44 one SD as second-line therapy for 18 patients with advanced or metastatic transitional cell carcinoma of the urothelium (APL-B-005-02),45 two SDs among 22 assessable patients with advanced dedifferentiated liposarcomas, with a median stem-cell transplantation) and four PRs. The median duration of response (DOR) and TTP were 2.2 and 1.6 months, respectively. No responses occurred in the 30 evaluable patients with other lymphomas (including 27 B-cell lymphomas).39

With respect to solid tumors, plitidepsin has shown a low level of antitumor activity in patients with malignant melanoma. In a Phase II study, 37 patients with advanced malignant melanoma who relapsed or progressed after one line of systemic therapy received plitidepsin according to the fortnightly scheme. The objective response rate (primary efficacy end point) was evaluated according to the Response Evaluation Criteria In Solid Tumors. Of the 35 evaluable patients, two dacarbazine-resistant patients (5.7%) with metastatic cutaneous melanoma achieved PR. Five other patients (14.3%) reported SD (median SD duration, 3.5 months; range, 2.2–15.8 months). Therefore, the rate of tumor growth control was 20.0%. With a median follow-up of 11.0 months, the median progression-free survival (PFS) was 1.3 months and the median overall survival (OS) was 3.5 months.40 These results suggest that further evaluation of plitidepsin in combination schedules is warranted. A Phase I/II trial compared plitidepsin alone or combined with dacarbazine (DTIC) as frontline therapy for advanced melanoma. The ORR with plitidepsin/DTIC was 21.4%; all responders had normal serum lactate dehydrogenase (LDH) levels and performance status ≤1 at baseline. Median PFS with plitidepsin/DTIC was 3.3 months in all patients and 4.3 months in those with baseline normal LDH. No responses occurred with single-agent plitidepsin – median PFS was 1.5 months. Both regimens were well tolerated. Further evaluation of plitidepsin 2.4 mg/m² fortnightly and DTIC 800 mg/m² q4wk was recommended by the authors.32
PFS of 1.6 months (APIPLO study).46 Coadministration of L-carnitine did not increase treatment efficacy and did not prevent the muscular toxicity or CPK-elevation associated with plitidepsin in patients with unresectable advanced renal cell carcinoma.41

Phase III trials
According to the encouraging preliminary results with plitidepsin plus dexamethasone in MM, a multicenter, open-label, randomized, Phase III clinical trial (ADMYRE), which started in June 2010 and finished recruitment in June 2015, has compared the efficacy and safety of plitidepsin combined with dexamethasone vs dexamethasone alone in patients with relapsed/refractory MM previously treated with between three and six therapeutic regimens. Efficacy data are currently being analyzed. A total of 255 patients received either plitidepsin 5 mg/m² IV as a 3-hour infusion on days 1 and 15 q4wk plus dexamethasone 40 mg orally on days 1, 8, 15 and 22 q4wk (Arm A), or dexamethasone alone at the same dose and schedule (Arm B). A press release has recently reported the significant benefit of plitidepsin plus dexamethasone over dexamethasone alone (a 35% lower risk of disease progression or death), and we await the final results.

Toxicities in Phase I studies
The primary dose-limiting toxicities (DLTs) found in the dose-finding Phase I studies with single-agent plitidepsin were musculoskeletal adverse events (AEs).26-30,36,41 The most common were myalgia, muscle weakness, and increase in serum CPK (noncardiac fraction) levels. Myalgia usually started proximally (shoulders, neck or thighs) at about 3 weeks after treatment onset, whereas muscle weakness and increases in serum CPK levels usually started about 2 weeks later. Myalgia and muscle weakness both generally resolved within 1–3 weeks. Increases in asymptomatic CPK may occur, as well as myalgia or muscle cramps, with no increase in CPK levels. No deaths or life-threatening consequences occurred due to these events. No rhabdomyolysis, increase of blood myoglobin levels, or myoglobinuria were reported from the administration of plitidepsin either as a single-agent or in combinations. Increased blood CPK levels were the only serious laboratory abnormality related to muscular disorders reported in single-agent Phase I studies.

The other relevant DLT in adult patients was hepatic toxicity, which manifested most frequently as transient and reversible increases in transaminase (particularly ALT).32,36 In Phase II studies, these abnormalities were asymptomatic, reversible, and noncumulative.

Hematological abnormalities were not DLTs in any of the schedules used in Phase I studies. As a single-agent, plitidepsin did not induce clinically significant bone marrow toxicity, stomatitis, or alopecia within the dose range explored. It is of note that neither grade 3/4 neutropenia nor thrombocytopenia were reported in any Phase I trial with plitidepsin as a single agent.

Grade 1/2 vomiting occurred in about half of the patients treated at the recommended dose. The use of prophylactic treatment for emesis resulted in a low incidence of grade 3/4 vomiting in Phase II studies (2%). General symptoms (eg, fatigue) were also reported during treatment with plitidepsin. Mild infusion site reactions were found, particularly (although not exclusively) when plitidepsin was administered through a peripheral line.

The most common DLTs found with the plitidepsin combination schedules32,36 were transaminase increases. DLTs consisting of hematological abnormalities were uncommon.

In the single-agent Phase I trial in pediatric patients (APL-A-005-02),30 the safety profile was not strikingly different from that in the adult Phase I program, although a slightly higher incidence of grade 3/4 hypersensitivity reactions was observed despite mandatory prophylactic medication.

Safety in completed Phase II trials
Treatment exposure
Four hundred and forty patients were treated in Phase II trials (288 patients diagnosed with solid tumors and 152 with hematological malignancies) with 1,646 cycles administered. Two hundred and seventy six patients (1,283 cycles) were treated with the fortnightly schedule, and 164 patients (363 cycles) with the weekly schedule.32,37-45

Muscular adverse events
Muscular events were described in about 50% of patients treated in Phase II studies, sharing similar characteristics to those seen in Phase I studies. Globally, muscle toxicity induced by plitidepsin is generally mild to moderate, manageable, and reversible upon treatment discontinuation or dose reduction, without requiring additional therapeutic measures. Myalgia occurred in about 40% of patients with solid tumors and 17% of those with hematological malignancies. Muscular weakness was the second most common muscular AE. Shorter infusion times may help reduce the incidence of
some types of drug-related events, including CPK elevations, gastrointestinal (anorexia, nausea, and vomiting), constitutional (fatigue), and injection-site reactions.

**Hematological abnormalities**

Most hematological abnormalities were mild to moderate. As expected, grade 3/4 toxicities occurred more often in patients with hematological malignancies than in patients with solid tumors. Most importantly, no grade 3/4 neutropenia occurred in solid tumor patients, in contrast to 16% of patients with hematological malignancies (fortnightly schedule). Bone marrow toxicity has been more common in studies evaluating the 3-hour and 1-hour infusion schedules, due to the higher proportion of patients with hematological malignancies in these studies.

**Biochemical abnormalities**

Most biochemical abnormalities were mild or moderate. Grade 3/4 abnormalities were more frequent in patients with hematological malignancies than in those with solid tumors, including increases in ALT, AST, CPK, and creatinine. In contrast, increases in amylase and bilirubin were more common in patients with solid tumors.

**Metabolic abnormalities**

No substantial differences were found between schedules, or between patients with solid tumors and patients with hematological malignancies. Grade 1/2 hyperglycemia and hypoalbuminemia were common in association either with premedication (dexamethasone) or with the underlying advanced neoplastic disease, respectively. As expected, hypercalcemia was common in MM patients and was related to the underlying malignancy.

**Gastrointestinal adverse events**

Nausea, vomiting, and diarrhea were the most frequent gastrointestinal AEs. Most episodes were mild or moderate and reached grade 3/4 in fewer than 2% of patients on the fortnightly schedule, but were more frequent among patients with solid tumors and those on the weekly schedule. Plitidepsin is defined as a treatment with high (non-cisplatin) emetogenic risk and appropriate primary prophylaxis is indicated.

**Constitutional symptoms**

Fatigue has been the most frequently reported constitutional adverse event in Phase II studies (41%). It is more common (49%) in patients with solid tumors than in those with hematological malignancies (21%–38%). Grade 3–4 fatigue was found less often (1%–18%). Noninfectious pyrexia has been documented in 7%–11% of patients.

**Cardiac adverse events**

The most frequent type of AE was rhythm alterations, while the other events were relatively infrequent. No life-threatening ventricular arrhythmias occurred. No fatal outcome was reported as a consequence of these cardiac AEs in Phase II trials.

**Hypersensitivity reactions**

Drug-related hypersensitivity reactions were found in 10.7% of patients treated with plitidepsin in Phase I and II trials. The incidence of hypersensitivity reactions was similar between the different tumor types. Grade 3/4 reactions occurred in 2.7%, with or without previous prophylaxis. Due to these reactions, all subsequent Phase II trials since February 2004 have included premedication with both anti-H1 and anti-H2 type antihistamines and glucocorticoids. Despite this prophylaxis, the rate of occurrence of serious reactions in all patients has remained unchanged overall, and only grade 1 reactions have decreased. In a phase II study using plitidepsin and dacarbazine as first-line therapy for advanced melanoma, hypersensitivity reactions were increased when using the combination therapy compared to plitidepsin as single agent. Globally, serious reactions tend to occur slightly more frequently in younger patients, females, and possibly those on weekly schedules in subsequent exposures (these reactions were rarely seen in the first infusion). These reactions are rapidly reversible and, to date, none has had a fatal outcome.

**Other adverse events**

The incidence of other AEs, such as neurological or respiratory AEs, was low or very low.

**Deaths**

Almost all deaths that occurred during treatment or follow-up were due to progression of the underlying malignancy and were unrelated to plitidepsin. In Phase I studies only one death was drug related. No deaths due to toxicity have been reported in children.

**Potential place in therapy**

Plitidepsin has shown mild-to-moderate antitumor activity, mainly in patients with MM, PTCL, and melanoma. Adverse
events are manageable and predictable. The mild hematological toxicity associated with plitidepsin makes this drug an attractive option for combination with chemotherapy schedules. As previously mentioned, recent studies suggest that eEF1A2 is the primary target of plitidepsin mediating its antitumor activity and might be a biomarker predicting drug sensitivity, although further studies are needed to confirm this.

Multiple myeloma

Plitidepsin has shown limited, although reproducible, single-agent activity in relapsed/refractory MM, which seems to be enhanced after dexamethasone addition while remaining well tolerated. The results of the Phase I trial combining plitidepsin with bortezomib and dexamethasone support the continuous development of additional studies to better define the role of plitidepsin in combination with other active agents in this indication. After the results of the Phase II trial (NCT00229203, APL-B-014-03) evaluating plitidepsin alone or in combination with dexamethasone, a new registry Phase III, multicenter, open-label, randomized trial (NCT 01102426, ADMYRE, APL-C-001-09) was started in June 2010 with recruitment being completed in June 2015. This trial evaluated the efficacy and safety of plitidepsin combined with dexamethasone vs dexamethasone alone in 255 patients with relapsed/refractory MM previously treated with between three and six therapeutic regimens (bortezomib and lenalidomide–thalidomide included). Efficacy data from ADMYRE will support valuable information regarding which patients could benefit from this drug. A press release has recently reported that plitidepsin plus dexamethasone had a significantly 35% lower risk of disease progression or death compared with dexamethasone. Based on these results, plitidepsin will be put forward for approval for use in Europe in this combination for the treatment of patients with MM.

Lymphoma

Single-agent plitidepsin has shown antitumor activity and a tolerable safety profile in patients with relapsed/refractory noncutaneous PTCL in a Phase II trial (APL-B-013-02). These results support the conduction of further clinical trials to evaluate plitidepsin-containing combined therapies in this disease setting. The absence of antitumor activity found in aggressive B-cell lymphomas precludes further studies of plitidepsin in this indication. A Phase II study of plitidepsin as a single-agent in indolent lymphomas is ongoing and the results are awaited.

Solid tumors

Plitidepsin is a cyclic depsipeptide that was first isolated from a Mediterranean marine tunicate (Aplidium albicans) and, at present, is manufactured by total synthesis and commercialized as plitidepsin. Its antitumor activity, observed in preclinical in vitro and in vivo studies, has prompted numerous clinical trials during the last 17 years involving plitidepsin alone or in combination with other anticancer agents. It has a favorable and predictable toxicity profile and has limited although reproducible single-agent activity in relapsed/refractory MM, which seems to be enhanced after dexamethasone addition, while remaining well tolerated. A Phase III trial comparing plitidepsin plus dexamethasone vs dexamethasone alone is underway, and additional studies are awaited that better define its role in combination with other active agents in this indication. Single-agent plitidepsin shows moderate antitumor activity and a tolerable safety profile in patients with relapsed/refractory noncutaneous PTCL and in patients with melanoma in Phase II trials. Further clinical trials to evaluate combined therapies containing plitidepsin in these settings are warranted. Results of the use of plitidepsin in the treatment of other hematologic malignancies or solid tumors have been disappointing. Further studies analyzing its mechanisms of action and potential biomarkers will help identify those patients who might benefit most from this drug.

Conclusion

Plitidepsin is a cyclic depsipeptide that was first isolated from a Mediterranean marine tunicate (Aplidium albicans) and, at present, is manufactured by total synthesis and commercialized as plitidepsin. Its antitumor activity, observed in preclinical in vitro and in vivo studies, has prompted numerous clinical trials during the last 17 years involving plitidepsin alone or in combination with other anticancer agents. It has a favorable and predictable toxicity profile and has limited although reproducible single-agent activity in relapsed/refractory MM, which seems to be enhanced after dexamethasone addition, while remaining well tolerated. A Phase III trial comparing plitidepsin plus dexamethasone vs dexamethasone alone is underway, and additional studies are awaited that better define its role in combination with other active agents in this indication. Single-agent plitidepsin shows moderate antitumor activity and a tolerable safety profile in patients with relapsed/refractory noncutaneous PTCL and in patients with melanoma in Phase II trials. Further clinical trials to evaluate combined therapies containing plitidepsin in these settings are warranted. Results of the use of plitidepsin in the treatment of other hematologic malignancies or solid tumors have been disappointing. Further studies analyzing its mechanisms of action and potential biomarkers will help identify those patients who might benefit most from this drug.

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Disclosure

The authors report no conflicts of interest in this work.
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