The novel melphalan prodrug J1 inhibits neuroblastoma growth in vitro and in vivo

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Abstract

Neuroblastoma is the most common extracranial solid tumor of childhood. The activity of J1 (L-melphalanyl-p-L-fluorophenylalanine ethyl ester), an enzymatically activated melphalan prodrug, was evaluated in neuroblastoma models in vitro and in vivo. Seven neuroblastoma cell lines with various levels of drug resistance were screened for cytotoxicity of J1 alone or in combination with standard cytotoxic drugs, using a fluorometric cytotoxicity assay. J1 displayed high cytotoxic activity in vitro against all neuroblastoma cell lines, with IC50 values in the submicromolar range, significantly more potent than melphalan. The cytotoxicity of J1, but not melphalan, could be significantly inhibited by the aminopeptidase inhibitor bestatin. J1 induced caspase-3 cleavage and apoptotic morphology, had additive effects in combination with doxorubicin, cyclophosphamide, carboplatin, and vincristine, and synergistically killed otherwise drug-resistant cells when combined with etoposide. Athymic nude mice carrying neuroblastoma xenografts [SH-SY5Y, SK-N-EB(2)] were treated with equimolar doses of melphalan, J1, or no drug, and effects on tumor growth and tissue morphology were analyzed. Tumor growth in vivo was significantly inhibited by J1 compared with untreated controls. Compared with melphalan, J1 more effectively inhibited the growth of mice with SH-SY5Y xenografts, was associated with higher caspase-3 activation, fewer proliferating tumor cells, and significantly decreased mean vascular density. In conclusion, the melphalan prodrug J1 is highly active in models of neuroblastoma in vitro and in vivo, encouraging further clinical development in this patient group. [Mol Cancer Ther 2007;6(9):2409–17]

Introduction

Neuroblastoma is the most common and deadly tumor of childhood often associated with therapy-resistance. For children with metastatic disease at diagnosis or certain genetic tumor features such as MYCN amplification or 1p deletion, the survival remains poor despite intensive multimodal treatment protocols (1).

The alkylating drug melphalan is routinely used in high-dose protocols for children with advanced neuroblastoma. Some decades ago, a mixture of six oligopeptides containing the meta-isomer of melphalan (m-L-sarcolysin), designated Peptichemio, was developed by Italian researchers and initial clinical trials seemed promising with significant response rates in several common diagnoses; among them, childhood neuroblastoma (2–5).

As a result of a screening of peptides, based on pharmacologic data on Peptichemio and melphalan, a highly effective drug candidate, J1 (L-melphalanyl-p-L-fluorophenylalanine ethyl ester), exhibiting higher in vitro and in vivo cytotoxicity than melphalan, was synthesized (6–8). J1 is rapidly incorporated into the cytoplasm followed by intracellular hydrolysis, which results in the release of melphalan. The enzymes responsible for the activation have been identified to be aminopeptidases (9), which may provide an attractive cancer target because the activity of several aminopeptidases is elevated in plasma and effusions from patients with cancer (10–12). A phase I trial with J1 in adult patients with advanced cancers is currently ongoing in Sweden.

There is a need for novel therapies able to bypass drug resistance of high-risk neuroblastoma, and based on previous experience with melphalan and Peptichemio in this diagnosis, it seemed adequate to investigate if J1 is active as a single agent or in combination with standard drugs in experimental models of neuroblastoma in vitro and in vivo. This is the first study showing that J1 effectively inhibits neuroblastoma cell growth in vitro and in vivo by the induction of apoptosis, being significantly more effective than melphalan and with additive or synergistic effects in combination with cytotoxic drugs routinely used for children with advanced neuroblastoma. Furthermore, this is the first report of J1 efficacy using the xenograft model including immunohistochemical examination of the tumors.
Materials and Methods

Chemicals and Reagents

The molecular structures of J1 and melphalan are shown in Fig. 1. J1 was synthesized as described previously (6) and dissolved in DMSO (Sigma-Aldrich) and further diluted with sterile water or PBS (Sigma-Aldrich). The DMSO concentration did not exceed 1% v/v in any experiment. For the in vivo studies, J1 was dissolved in N,N-dimethyl acetamide (Sigma-Aldrich) and further diluted in glucose solution (50 mg/mL; Apoteket AB). Fluorescein diacetate (Sigma-Aldrich) was dissolved in DMSO and kept frozen (−20°C) as a stock solution protected from light. Carboplatin, doxorubicin, etoposide, melphalan, and vincristine were obtained from the Swedish Pharmacy (Aopoteket AB) and dissolved according to guidelines from the manufacturer and further diluted in sterile PBS (except in the xenograft studies, where melphalan was further diluted in 50 mg/mL glucose). 4-Hydroxycyclophosphamide (the active metabolite of cyclophosphamide) was a kind gift from S. Ludeman (Duke Comprehensive Cancer Center, Duke University Medical Center, Durham, NC). The aminopeptidase inhibitor bestatin (Sigma-Aldrich) was dissolved in 50% ethanol to a 4.0 mmol/L stock solution and further diluted in PBS. FAM-DEVD-FMK, supplied as part of the CaspaTag kit (Chemicon) and chloromethyl-X-rosamine (MitoTracker Red CMXRos; Molecular Probes) were dissolved in DMSO and further diluted in PBS to its final concentration. Hoechst 33342 (Sigma-Aldrich) was dissolved in water.

Human Tumor Cell Lines

Seven neuroblastoma cell lines, with different drug sensitivities, were grown in Eagle Minimal Essential Medium (SH-SY5Y; Sigma-Aldrich) or RPMI 1640 [SK-N-BE(2), SK-N-AS, SK-N-FI, SK-N-SH, SK-N-DZ, and IMR-32; Sigma-Aldrich] medium supplemented with 10% fetal bovine serum (Sigma-Aldrich) and 100 units/mL of penicillin (Sigma-Aldrich) at 37°C humidified 5% CO2 atmosphere.

Cytotoxicity Assay

The fluorometric microculture cytotoxicity assay was used (13, 14) to investigate the effect of J1 and the combination of J1 with other cytostatic drugs in vitro. Briefly, 96- or 384-well microtiter plates (NUNC Brand Products) were pre-prepared in duplicate with drug solutions at 10 times the desired final drug concentration. Cells were seeded into the drug-prepared microtiter plates at a cell density of 0.1 × 104 cells/mL and incubated for 72 h at 37°C in a humidified 5% CO2 atmosphere. Fluorometric microculture cytotoxicity assay was done using an automated Optimized Robot for Chemical Analysis (Beckman Coulter) programmed through the software SAMI (Beckman Coulter). The plates were washed, fluorescein diacetate was added, and the fluorescence generated was measured at 485/520 nm using Fluoroscan Optima (BMG Labtech) after 50 min incubation. The resulting fluorescence is proportional to the number of intact cells in each well. A successful assay required a ratio of >5 between the signal in the control wells and the blank wells and a coefficient of variation of <30% in the control wells. Cell survival is presented as a survival index. For the combination studies, fixed concentration ratios of the drugs were used with 5-fold serial dilutions in five steps. All concentrations were tested in duplicate and the experiments were repeated thrice.

Bestatin was used to evaluate the effect of aminopeptidases on the cytotoxic effect of J1 in all seven neuroblastoma cell lines. In these analyses, cells were preincubated with 10 μmol/L of bestatin (nontoxic) for 60 min, seeded into drug-containing microtiter plates, incubated for another 30 min before the drug-containing medium was removed and fresh medium added. The activities were measured after 72 h using a fluorometric microculture cytotoxicity assay.

Apoptosis Analysis

A multiparametric high-content screening assay (15) was used for the measurement of apoptosis in the three neuroblastoma cell lines; SH-SY5Y, SK-N-AS, and SK-N-BE (2). Briefly, cells were seeded into 96-well plates (Perkin-Elmer, Inc.) and incubated with the indicated concentrations of drugs for 8 and 24 h. The FLICA probe FAM-DEVDFMK (carboxyfluorescein-labeled fluoromethyl ketone peptide inhibitor of caspase-3; at a final concentration of 20 μmol/L) was added 1 h before the end of the drug exposure to stain activated caspase-3/7, and on the last 30 min chloromethyl-X-rosamine was included (at a final concentration of 100 nmol/mL) to evaluate mitochondrial membrane potential. The plates were washed and stained with 10 μmol/L of Hoechst 33342 in 3.7% formaldehyde. Plates were analyzed using the ArrayScan high-content screening system (Cellomics®, Inc.). Images were acquired for each fluorescence channel, using suitable filters with a 20× objective, and in each well, at least 800 cells were analyzed. J1 and melphalan were tested in four different concentrations obtained by a 5-fold serial dilution. Each experiment was done thrice.

Treatment with J1 and Melphalan In vivo

Male nude rats (HsdHan: RNU-nu; Harlan) at age 5 to 10 weeks with a weight of 150 to 275 g, and female nude mice (NMRI nu/nu; Taconic) at age 6 weeks with a weight of 18 to 25 g were used for the experiments. The animals were housed and maintained in laminar flow cabinets under specific pathogen–free conditions and given sterile conditions of specific pathogen–free conditions and given sterile conditions.
water and food ad libitum. The animal experiments were approved by the regional ethics committee for animal research (N234-05 and N75-05) in accordance with the Animal Protection Law (SFS 1988:534), the Animal Protection Regulation (SFS 1988:539), and the Regulation for the Swedish National Board for Laboratory Animals (SFS 1988:541). Establishment of neuroblastoma xenografts in nude rats was done as previously described (16). For the establishment of neuroblastoma xenografts in nude mice, animals were engrafted with \( 3 \times 10^6 \) SH-SY5Y cells (in 0.1 mL of medium) s.c. in the flank of the right hind leg using a 23-gauge needle.

Three independent experiments were carried out. One using the multidrug-resistant, MYCN-amplified, p53-mutated cell line SK-N-BE(2) in nude rats and two experiments were done using the less drug-resistant SH-SY5Y cell line, one in rats and one in mice. Initial doses were selected based on previously determined LD50 values from other strains of rats/mice, which could not separate the toxic effects of melphalan and J1 at equimolecular doses.\(^5\) In the first experiment, nude rats (\( n = 16 \)) carrying SK-N-BE(2) xenograft tumors were randomly assigned to receive one dose of 10 \( \mu \)mol/kg J1 i.v. in the tail vein (\( n = 5 \)), 10 \( \mu \)mol/kg of melphalan i.v. (\( n = 5 \)) at day 0 or no treatment (\( n = 6 \)), respectively. In the second experiment, nude rats (\( n = 16 \)) carrying SH-SY5Y xenografts were randomized to receive 0.50 \( \mu \)mol/kg of J1 i.v. (\( n = 5 \)), 0.50 \( \mu \)mol/kg of melphalan i.v. (\( n = 5 \)) at day 0, or no treatment (\( n = 6 \)), respectively. In the third experiment, nude mice (\( n = 25 \)) engrafted with SH-SY5Y cells were randomly assigned to receive 0.50 \( \mu \)mol/kg of J1 i.v. (\( n = 9 \)), 0.50 \( \mu \)mol/kg of melphalan i.v. (\( n = 8 \)) at days 0 and 6, or no treatment (\( n = 8 \)), respectively. Treatment continued for 12 days, starting when a tumor had reached a volume of 0.20 to 0.30 mL (mean tumor volume, 0.30 mL in experiments one and two, 0.26 mL in experiment three). Tumor volume was measured every other day as described previously (16). All animals were monitored for signs of toxicity including weight loss during the experiment.

**Immunohistochemistry**

Apoptosis, proliferation, and angiogenesis in vivo were evaluated in deparaffinized tumor sections of neuroblastoma xenografts. For identification of apoptosis, sections were incubated overnight at 4°C with a rabbit monoclonal anti–active caspase-3 antibody (1:100; R&D Systems) and detected using anti-rabbit HRP-conjugated Superpicture Polymer Kit (Zymed Laboratories, Inc.). For assessment of proliferative activity, sections were incubated with a monoclonal rabbit anti–Ki-67 antibody (1:200; Neomarkers, Inc.) overnight at 4°C and detected using an anti-rabbit HRP SuperPicture Polymer Kit (Zymed). As a control for nonspecific background staining, sections were incubated with rabbit IgG isotype control (Zymed). Biotinylated *Bandiera simplicifolia* (BS-1, L3759; Sigma-Aldrich) lectin was used for highlighting endothelial cells as described previously (17). Apoptosis and proliferation were assessed by counting the number of positively stained nuclei and the total number of tumor cells in three representative regions in three tumors from each treatment group at \( \times 400 \) magnification. The results are expressed as the proportion of positively stained cells. Sections were quantified for vessel density at \( \times 200 \) magnification, and fields were chosen randomly with the inclusion criteria that it had to consist of viable tissue. The results are expressed as an average number of vessels per field.

**Statistical Analysis**

The cytotoxic IC\(_{50}\) values for drugs in the cell lines in vitro were determined from log concentration-effect (survival index %) curves in GraphPad Prism (GraphPad Software, Inc.) using nonlinear regression analysis. For the in vitro results, comparison of activity between two groups was made with a two-sided \( t \) test. To test combination effects in vitro, data were analyzed using the median effect method of Chou and Talalay (18) using the software CalcuSyn Version 2 (Biosoft). Each dose-response curve (individual agents as well as combinations) were fit to a linear model using the median effect equation, allowing calculation of a median effect value \( D \) (corresponding to the IC\(_{50}\)) and slope. Goodness-of-fit was assessed using the linear correlation coefficient, \( r \), and \( r > 0.85 \) was required for a successful analysis. The extent of interaction between the drugs was expressed using the combination index (CI) for mutually exclusive drugs: \( CI = D_1/D_1 + D_2/D_2 \), where \( D_1 \) and \( D_2 \) represent the concentrations of drugs 1 and 2 alone, required to produce a certain effect, and \( d_1 \) and \( d_2 \) are the concentrations of drugs 1 and 2 in combination required to produce the same effect. Different CI values are obtained when solving the equation for different effect levels and a 70% effect was chosen for presentation. A CI lying on one indicates additivity; CI significantly lower was defined as synergy and CI significantly higher as antagonism. One-sample \( t \) tests were used to determine if the CIs differed from one (1.0). For the in vivo experiments, one-way ANOVA with Tukey multiple comparisons tests were used to compare the three treatment groups (GraphPad InStat). \( P < 0.05 \) was considered significant.

**Results**

**The J1 Prodrug Is a Potent Inducer of Neuroblastoma Cell Death and Is Superior to Melphalan In vitro**

The cytotoxic activity of J1, melphalan, and five cytotoxic drugs commonly used in the treatment of patients with neuroblastoma was investigated in seven neuroblastoma cell lines. In two of the neuroblastoma cell lines (SH-SY5Y and SK-N-SH), J1 was found to be the most effective drug with IC\(_{50}\) values of 0.0028 and 0.0051 \( \mu \)mol/L, respectively. In the remaining cell lines, only vincristine was more effective than J1 in inducing cell death (Fig. 2). J1 exhibited a significant increase (mean, 270-fold; range, 35- to 810-fold) in cytotoxicity compared with melphalan in all neuroblastoma cell lines investigated (\( t \) test, \( P < 0.001 \); Fig. 2).

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\(^5\) Unpublished data.
Figure 2. The IC₅₀ (with 95% confidence interval) of J1, melphalan, and five cytotoxic drugs commonly used in the treatment of neuroblastoma in seven neuroblastoma cell lines. The increase in activity of J1 compared with melphalan (defined as IC₅₀ melphalan/IC₅₀ J1, indicated by numbers) was significant (t test, P < 0.001) for all cell lines.
Table 1. Effect of J1 in combination with chemotherapeutic drugs in neuroblastoma cells in vitro

<table>
<thead>
<tr>
<th>Combination</th>
<th>SK-N-AS CI at IC70</th>
<th>Effect</th>
<th>SK-N-BE(2) CI at IC70</th>
<th>Effect</th>
<th>SH-SY5Y CI at IC70</th>
<th>Effect</th>
</tr>
</thead>
<tbody>
<tr>
<td>J1 + doxorubicin</td>
<td>0.86 (0.63–1.11)</td>
<td>Additive</td>
<td>0.88 (0.57–1.2)</td>
<td>Additive</td>
<td>0.78 (0.64–0.92)</td>
<td>Synergistic</td>
</tr>
<tr>
<td>J1 + carboplatin</td>
<td>0.69 (0.38–0.99)</td>
<td>Synergistic</td>
<td>1.1 (0.84–1.4)</td>
<td>Additive</td>
<td>0.96 (0.33–1.6)</td>
<td>Additive</td>
</tr>
<tr>
<td>J1 + cyclophosphamide</td>
<td>0.82 (0.37–1.3)</td>
<td>Additive</td>
<td>1.3 (0.58–2.0)</td>
<td>Additive</td>
<td>1.0 (0.69–1.3)</td>
<td>Additive</td>
</tr>
<tr>
<td>J1 + etoposide</td>
<td>0.47 (0.44–0.49)</td>
<td>Synergistic</td>
<td>0.75 (0.49–0.99)</td>
<td>Synergistic</td>
<td>0.63 (0.25–0.99)</td>
<td>Synergistic</td>
</tr>
<tr>
<td>J1 + vincristine</td>
<td>1.4 (0.34–2.5)</td>
<td>Additive</td>
<td>1.2 (0.46–1.9)</td>
<td>Additive</td>
<td>0.78 (0.30–1.3)</td>
<td>Additive</td>
</tr>
</tbody>
</table>

NOTE: Mean of CI at IC70 with 95% confidence interval. Mutual exclusivity is assumed. Synergism and antagonism are defined as a CI mean statistically significantly lower/higher than one with one-sample t test (P < 0.05).

To confirm that J1 acts as an enzymatic activated prodrug, as previously described (9), cells were pretreated with bestatin, a potent inhibitor of aminopeptidases (19). Pre-exposure to nontoxic concentration of bestatin resulted in a decreased activity of J1 in six of seven cell lines (t test, P < 0.05). In the IMR-32 cell line, the observed difference, however, did not reach statistical significance using this experimental setup. In general, bestatin pretreatment increased the IC50 by 5.5 times (paired t test, P = 0.0006). Bestatin did not affect neuroblastoma cell response to melphalan in any of the cell lines (t test, P > 0.05; paired t test P = 0.30).

**J1 Potentiates Standard Neuroblastoma Chemotherapy In vitro**

Combination chemotherapy, often supported with stem cell infusions, is considered as standard treatment in many clinical situations, and melphalan has been employed in several different protocols. Synergistic, additive, or antagonistic in vitro effects of J1 in combination with relevant chemotherapeutic drugs were evaluated in three neuroblastoma cell lines with different levels of drug resistance [SH-SY5Y, SK-N-AS, and SK-N-BE(2)]. As summarized in Table 1, which shows the CI at IC70, the addition of J1 to standard drugs induced significant synergistic or additive cytotoxic effects in all tested cell lines. Synergism was observed when J1 was combined with etoposide (all cell lines), and for carboplatin (SK-N-AS), and doxorubicin (SH-SY5Y). The other combinations were found to be additive (Table 1).

**J1 Triggers Apoptosis in Neuroblastoma**

In order to examine if J1 can trigger the apoptotic machinery, mitochondrial transmembrane potential and caspase-3 activity were measured, and nuclear DNA was assessed for apoptotic fragmentation in three neuroblastoma cell lines (Fig. 3). Mitochondrial depolarization was observed 8 hours after the addition of J1 (data not shown). At 24 h post-drug addition, J1 and melphalan were both associated with caspase-3 activation and nuclear DNA fragmentation. Taken together, these observations indicate that J1 induces the apoptosis of neuroblastoma cells by activation of the intrinsic apoptotic pathway. Notably, the proapoptotic effects of J1 were significantly more potent compared with melphalan (t test, P < 0.001; Fig. 3).

**J1 Significantly Inhibits the Growth of Established Neuroblastoma Xenografts**

To investigate the effect of J1 and melphalan on neuroblastoma growth in vivo, nude rats and mice carrying either SK-N-BE(2) or SH-SY5Y xenografts were used. In the first experiment, nude rats carrying xenografts from the drug-resistant neuroblastoma cell line SK-N-BE(2) were treated with a single dose of 10 μmol/kg of J1 or melphalan, a dose reported to be close to 50% of LD50. Significant inhibition of tumor growth was detected 2 days after J1 treatment (ANOVA, P = 0.016; Tukey, P < 0.05) and 4 days after melphalan treatment (ANOVA, P = 0.0003; Tukey, P < 0.01), respectively, compared with untreated tumors and throughout the observation period (ANOVA, P > 0.001; Fig. 4A). Using this dose scheduling, no significant difference in tumor growth was observed in rats treated with J1 compared with melphalan (Tukey, P > 0.05).

In the second experiment, nude rats carrying SH-SY5Y xenografts were treated with a single dose of 0.5 μmol/kg of J1 or melphalan, a dose corresponding to ~25% of LD50 (again estimated in Sprague-Dawley strain); thus, considerably lower than in the first experiment. Again, both treatments significantly inhibited tumor growth compared with untreated control tumors (ANOVA, P < 0.001; Fig. 4B). However, J1 was significantly more effective in tumor growth inhibition compared with melphalan using this lower concentration of drugs (Tukey, P < 0.05). Inhibitions were significant after 2 days for both J1 (ANOVA, P = 0.0029; Tukey, P < 0.01) and melphalan (Tukey, P < 0.05), compared with untreated tumors.

Previous in vivo studies with J1 were made with the hollow fiber method in mice only (8), showing the increased potency of J1 but a similar toxicity compared with equimolar doses of melphalan. This is the first presentation of J1’s activity in a xenograft model, as well as its effect in a rat model for comparison, grafts were also studied in mice (having potentially different metabolisms and different maximum tolerated doses in previous studies). Tumor-bearing mice were treated i.v. with two doses of 0.5 μmol/kg, corresponding to ~0.6% of LD50, of

* Unpublished data, estimation in Sprague-Dawley rats.

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either J1 or melphalan. The drugs were given when the tumor volume had reached >0.20 mL (mean, 0.26 mL) and at 6 days after tumor establishment (Fig. 5A). Tumor growth was significantly inhibited 2 days after J1 injection compared with both melphalan and the untreated control (ANOVA, \( P = 0.0073 \); Tukey, \( P < 0.05 \)) and throughout the observation period (ANOVA, \( P = 0.0002 \); Tukey, \( P > 0.01 \) J1 versus melphalan; \( P > 0.001 \) J1 versus control; Fig. 5A). No significant reduction in tumor volume was observed in animals treated with melphalan (Tukey, \( P > 0.05 \)). Animal weight and other signs of toxicity were recorded in all the in vivo experiments. In rats treated with a single dose of 10 \( \mu \)mol/kg of J1 or melphalan, a transient reduction in weight gain was observed on the first 4 days after treatment compared with the untreated control animals (ANOVA, \( P = 0.011 \); Tukey, \( P < 0.05 \); Fig. 4B). No significant differences in weight gain were observed at the end of the experiments (ANOVA, \( P = 0.25 \); Fig. 4B). In rats treated with 0.50 \( \mu \)mol/kg of J1 or melphalan, a temporary reduction in weight gain was observed in rats treated with melphalan on the first 4 days after treatment (ANOVA, \( P = 0.014 \); Tukey, \( P < 0.05 \); Fig. 4D). No weight loss was observed in rats treated with 0.50 \( \mu \)mol/kg of J1 compared with untreated control rats throughout the experiment (Fig. 4D). In mice treated with two doses of 0.50 \( \mu \)mol/kg of J1 compared with untreated control rats at days 1 and 6, no weight loss was observed in the treated mice compared with untreated controls (Fig. 5B). No other signs of toxicity were observed in any of the treatments (data not shown).

**Treatment with J1 Significantly Reduces Proliferation, Inhibits Angiogenesis, and Induces Apoptosis in Neuroblastoma In vivo**

Having established that J1 seems more effective than melphalan in the treatment of established neuroblastoma xenografts, we investigated the effect of the drugs on the induction of apoptosis, tumor cell proliferation, and angiogenesis in SH-SY5Y xenografts.

A significantly increased level of cleaved caspase-3–positive cells was found in tumors from rats treated with J1 compared with untreated tumors (ANOVA, \( P = 0.0021 \); Tukey, \( P < 0.01 \); Fig. 4C). Furthermore, caspase-3 activity was significantly higher in J1-treated tumors than in xenografts from rats treated with melphalan (Tukey, \( P < 0.01 \); Fig. 4C). Cell proliferation as measured by Ki-67 was significantly inhibited after J1 treatment, but not after melphalan treatment compared with untreated tumors (ANOVA, \( P = 0.042 \); Tukey, \( P < 0.05 \) J1 versus control; \( P > 0.05 \) melphalan versus control; Fig. 4D).

A significantly elevated expression of cleaved caspase-3 was detected in SH-SY5Y xenografts from nude mice treated with J1 compared with melphalan-treated tumors (ANOVA, \( P = 0.0021 \); Tukey, \( P < 0.01 \); Fig. 4C). Furthermore, caspase-3 activity was significantly higher in J1-treated tumors than in xenografts from rats treated with melphalan (Tukey, \( P < 0.01 \); Fig. 4C). Cell proliferation as measured by Ki-67 was significantly inhibited after J1 treatment, but not after melphalan treatment compared with untreated tumors (ANOVA, \( P = 0.0021 \); Tukey, \( P < 0.01 \) J1 versus control; \( P > 0.05 \) melphalan versus control; Fig. 4D).

A significantly elevated expression of cleaved caspase-3 was detected in SH-SY5Y xenografts from nude mice treated with J1 compared with melphalan-treated tumors (ANOVA, \( P = 0.017 \); Tukey, \( P < 0.05 \); Fig. 5B). Moreover, a significant reduction of tumor cell proliferation was also shown in J1-treated tumors compared with untreated tumors (ANOVA, \( P = 0.042 \); Tukey, \( P < 0.05 \) J1 versus control; \( P > 0.05 \) melphalan versus control; Fig. 5B). Because inhibition of angiogenesis could contribute to reduced cell proliferation and increased apoptosis, we also determined the effect of J1 and melphalan on mean vessel density. Treatment with J1 or melphalan induced a 49% (95% confidence interval, 27–72%; ANOVA, \( P < 0.001 \); Tukey, \( P < 0.001 \)) or 20% (−0.35% to 41%; Tukey, \( P > 0.05 \))

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7 Unpublished data, estimation in specific pathogen-free Swiss mice.
reduction in mean vessel density compared with untreated tumors. Mean vessel density was also significantly lower (36% reduction, 7.9–65%) in tumors treated with J1 compared with melphalan (Tukey, P < 0.05; Fig. 5D).

**Discussion**

Melphalan could prolong the survival of children with advanced neuroblastoma and is currently used in many high-dose protocols for this patient group (20). The present study shows for the first time that J1, a prodrug of melphalan, is highly active against human neuroblastoma cell lines *in vitro* and *in vivo*. The cytotoxic activity of J1 *in vitro* was found to be superior compared with melphalan, and statistical comparisons also favored J1 in terms of antiproliferative, proapoptotic, and antiangiogenic activity *in vivo*. These findings are particularly interesting in view of the demonstrated clinical activity of Peptichemo in neuroblastoma (4, 5).

Despite different levels of absolute sensitivity, the seven human neuroblastoma cell lines displayed a similar pattern of sensitivity against the tested drugs (Pearson correlation of log IC_{50}, 0.85–0.99), reflecting the importance of a common cytopathologic origin. The *in vitro* activity of J1 was, on average, 270-fold that of melphalan, with a range from 35- to 810-fold. There is a clear tendency toward a greater difference in the generally more sensitive cell lines (i.e., SH-SY5Y and its parental cell line SK-N-SH) compared with the more resistant cell lines (Fig. 2). Thus, one or more drug targets specific for J1 may be more abundantly expressed in SH-SY5Y/SK-N-SH and the observation warrants further mechanistic studies. No obvious differences in IC_{50} of J1 between MYCN and non-MYCN–amplified cell lines could be observed. Both melphalan and J1 activated the intrinsic apoptotic pathway in neuroblastoma cells. However, a reduction in the levels of activated caspase-3/7 was observed for high concentrations of melphalan, which suggests that melphalan may induce neuroblastoma cell death by other mechanisms at higher concentrations (Fig. 3).

The major use of melphalan in neuroblastoma therapy is in high-dose regimens with stem cell support. For this purpose, melphalan has been evaluated in combination with...
with, for example, cyclophosphamide (21), etoposide or carboplatin (22), and etoposide (23). In this work, we chose to study combinations with common neuroblastoma drugs representing different structural classes or mechanisms, and several interesting observations were made; most notably, J1 and the topoisomerase II inhibitor etoposide induced neuroblastoma cell death in a synergistic manner, the mean CI<sub>70</sub> being significantly lower than one in all three cell lines. Notably, synergistic interactions between melphalan and topoisomerase inhibitors in vitro have been reported (24), and high-dose regimens of melphalan and etoposide have been tested clinically in patients with stage IV neuroblastoma and were found to be well-tolerated (23).

Previous in vivo studies were done in mice and included dose-finding toxicity and antitumor efficacy in the hollow fiber model. Monitored variables included behavior, weight change, blood cell analysis, autopsy, and histopathologic examinations of sternal bone marrow and the intestines. Early weight loss comparisons favored J1 over melphalan, but differences in all other variables were not significant (significant effects on blood cell count verses control for both drugs; ref. 8). Despite being simple, elegant, and robust, the hollow fiber method suffers from some disadvantages. For example, an almost complete lack of tumor-matrix interaction and no simple histopathologic or immunohistochemical evaluation of treatment results. The in vivo findings presented in this article showed that J1 is highly active, both against drug-sensitive (SH-SY5Y) and multidrug-resistant [SK-N-BE(2)] xenografts, at doses giving no observed toxicity except for a transient reduced weight gain using high doses of J1 in rats. Compared with melphalan, J1 seems to induce a more potent proapoptotic as well as the antiangiogenic activity in neuroblastoma in vivo. That J1, in addition to its direct antitumoral effects, is able to significantly reduce the number of blood vessels in neuroblastoma, is a novel finding. The metalloproteinase aminopeptidase N is highly expressed in vascular endothelial cells and has been shown to play multiple roles in angiogenesis (25). Preliminary findings in our lab show that aminopeptidase N is a target of J1, which may

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* Wickström et al., unpublished observation.
be relevant in view of the antiangiogenic effect of J1 presented here. In their recent work, Pastorino et al. showed that doxorubicin-containing liposomes coupled to aminopeptidase N–directed peptides successfully targets the vasculature of neuroblastoma xenografts (26). Our finding that the aminopeptidase inhibitor bestatin partly inhibits the direct antitumoral effects of J1 against neuroblastoma cells further supports a role for this enzyme as an important target of J1. Studies to further unravel the involvement of aminopeptidase N as a target of J1 are now ongoing.

One possible explanation for the somewhat moderate activity differences between J1 and melphanal compared with the in vitro data in the neuroblastoma xenograft studies in rodents may be related to the enzymatic differences between rodents and humans. The produg has two hydrolysis susceptible bonds, one peptide and one ester bond. Cleaving of the peptide bond (i.e., by peptidases) leads to melphalan, and in a previous publication, we presented evidence that this cleavage results in the increased activity of J1 compared with melphalan (8). Reports of extended esterase activity in rodents compared with humans have been published previously (27). This has been shown to be important for other produgs like CPT-11 (irinotecan) efficacy and toxicity in rodent experiments in vivo (28). With the rapid de-esterification of J1, some of the advantages compared with melphalan may be lost, e.g., because the de-esterified J1 is less lipophilic than J1, passive transportation into the cells may decrease. Indeed, de-esterified J1 also possesses reduced cytotoxic activity in vitro.

In summary, these studies show significant activity of J1, a novel alkylating dipeptide and produg of melphalan, in neuroblastoma in vitro and in vivo, involving multiple parallel effects on tumor growth. Furthermore, additive or synergistic in vitro interactions with standard chemotherapy-apoetases, particularly etoposide, are shown. These novel findings should encourage further preclinical studies to support clinical studies in children with neuroblastoma refractory to standard treatment.

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References


* Unpublished data.
The novel melphalan prodrug J1 inhibits neuroblastoma growth \textit{in vitro} and \textit{in vivo}

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