Preclinical Development

Cabozantinib (XL184), a Novel MET and VEGFR2 Inhibitor, Simultaneously Suppresses Metastasis, Angiogenesis, and Tumor Growth


Abstract
The signaling pathway of the receptor tyrosine kinase MET and its ligand hepatocyte growth factor (HGF) is important for cell growth, survival, and motility and is functionally linked to the signaling pathway of VEGF, which is widely recognized as a key effector in angiogenesis and cancer progression. Dysregulation of the MET/VEGF axis is found in a number of human malignancies and has been associated with tumorigenesis. Cabozantinib (XL184) is a small-molecule kinase inhibitor with potent activity toward MET and VEGF receptor 2 (VEGFR2), as well as a number of other receptor tyrosine kinases that have also been implicated in tumor pathobiology, including RET, KIT, AXL, and FLT3. Treatment with cabozantinib inhibited MET and VEGFR2 phosphorylation in vitro and in tumor models in vivo and led to significant reductions in cell invasion in vitro. In mouse models, cabozantinib dramatically altered tumor pathology, resulting in decreased tumor and endothelial cell proliferation coupled with increased apoptosis and dose-dependent inhibition of tumor growth in breast, lung, and glioma tumor models. Importantly, treatment with cabozantinib did not increase lung tumor burden in an experimental model of metastasis, which has been observed with inhibitors of VEGF signaling that do not target MET. Collectively, these data suggest that cabozantinib is a promising agent for inhibiting tumor angiogenesis and metastasis in cancers with dysregulated MET and VEGFR signaling.

Mol Cancer Ther; 10(12); 2298–308. ©2011 AACR.

Introduction
The receptor tyrosine kinase MET is the only known receptor for hepatocyte growth factor (HGF), and its signaling activity is required for embryogenesis, cell proliferation, survival, and motility (1). While MET and HGF are found in low levels in normal adult tissues, their expression is frequently dysregulated in a broad spectrum of human tumors. Ectopic activation of MET promotes tumor cell survival, growth, angiogenesis, invasion, and metastasis (2, 3). In preclinical animal models, overexpression of MET and/or HGF has been shown to stimulate tumorigenesis and metastasis (2, 4), whereas downregulation of MET or HGF expression resulted in increased apoptosis and decreased tumor growth and blood vessel density (5, 6). Clinically, dysregulated expression of HGF and/or MET has been observed in many different tumor types, including glioma, melanoma, hepatocellular, renal, gastric, pancreatic, prostate, ovarian, breast, and lung cancers, and is often correlated with poor prognosis (1, 7). Collectively, these observations strongly implicate MET/HGF signaling in the pathogenesis of human cancer.

Induction of angiogenesis has been increasingly recognized as a crucial step in tumor progression and is one of the hallmarks of cancerous growth. Correspondingly, the proangiogenic signaling molecule VEGF and its receptors VEGFR1, VEGFR2, and VEGFR3 play key roles in tumor development. HGF is also a potent angiogenic factor and acts synergistically with VEGF to induce angiogenesis (8, 9). The U.S. Food and Drug Administration-approved VEGF pathway inhibitors bevacizumab, sunitinib, and sorafenib have proven to be clinically important in the treatment of many types of cancer, but their effects are characterized by initial clinical benefit lasting weeks or months before the resumption of tumor growth and progression (10–12). Preclinical studies have found that while continuous VEGF pathway inhibition initially slows tumor growth and trims tumor vasculature, this is quickly followed by rapid revascularization and increased tumor invasiveness (13). These observations show that cancers have the capacity to develop resistance to VEGF pathway–targeted inhibition.
Tumor cell evasion of VEGF pathway–targeted inhibition may occur as a response to hypoxia. Under hypoxic conditions, hypoxia-inducible factor 1α is upregulated, resulting in increased expression of both VEGF and MET (14, 15). These responses apparently allow the tumor cells to compensate for the hypoxic environment through stimulation of angiogenesis or migration away from the hypoxic zone. Because VEGF pathway inhibition can result in induction of hypoxia, it may also trigger upregulation of MET expression, which may then stimulate tumor invasion. Indeed, Shojaei and colleagues recently determined that the MET pathway plays an important role in the development of resistance to VEGF pathway inhibition by sunitinib treatment (16). Recent studies have also showed that the use of VEGFR inhibitors, such as sunitinib, sorafenib, cediranib, or a VEGFR2-targeting antibody, can result in the development of an aggressive tumor phenotype characterized by increased invasiveness and metastasis and, in patients treated with cediranib, higher MET expression levels (17–19). Therefore, targeting both arms of the MET/VEGF axis simultaneously may critically disrupt angiogenesis, tumorigenesis, and cancer progression. Cabozantinib (XL184) is a potent inhibitor of MET and VEGFR2 that also inhibits RET, KIT, AXL, and FLT3, all of which have been implicated in tumor pathogenesis (20, 21). In this study, we report on the consequences of cabozantinib treatment on angiogenesis, cellular invasion, tumor growth, and metastasis. Collectively, these data suggest that cabozantinib may provide enhanced efficacy by simultaneously targeting key pathways important to tumor survival, metastasis, and angiogenesis.

Materials and Methods

Compounds

Cabozantinib (Fig. 1A) was synthesized as described in US Patent 7579473 (WO 2005030140 A2; ref. 22). For in vitro assays, 10 mmol/L cabozantinib was prepared in dimethyl sulfoxide and serially diluted in the appropriate assay media. For in vitro studies, cabozantinib was formulated in sterile saline/10 mmol/L HCl or in water and administered via oral gavage at 10 mL/kg (mice) or 2 mL/kg (rats).

Kinase inhibition assays

The inhibition profile of cabozantinib against a broad panel of 270 human kinases was determined using luciferase-coupled chemiluminescence, 32P-phosphorylation transfer, or AlphaScreen technology (PerkinElmer). Recombinant human full-length, glutathione S-transferase tag, or histidine tag fusion proteins were used, and half maximal inhibitory concentration (IC50) values were determined by measuring phosphorylation of peptide substrate polypeptide Glu, Thr, Tyr or Hsp70 at ATP concentrations 0 or below the Ki, for each respective kinase. The mechanism of kinase inhibition was evaluated using the AlphaScreen Assay by determining the IC50 values over a range of ATP concentrations.

Cell lines

A431, B16F10, C6, H441, H69, Hs746T, HT1080, MS1, PC3, SNU-1, SNU-5, SNU-16, and U87MG cells were purchased from the American Type Culture Collection, BaF3 cells were purchased from the German Resource Center for Biological Material (DSMZ), and HMVEC-L cells were purchased from BioWhittaker Technologies. MDA-MB-231 cells were a gift from Georgetown University School of Medicine, Washington, DC. No further authentication was done on these lines. Cell lines were passaged for less than 3 months and maintained in either Dulbecco’s Modified Eagle’s Media (DMEM) or RPMI-1640 media supplemented with 10% FBS, 1% penicillin-streptomycin, and 1% nonessential amino acids at 37°C in 5% CO2. Where indicated, assays were conducted using parental cell lines stably transfected with expression vectors encoding kinases of interest.

Inhibition of receptor phosphorylation

Receptor phosphorylation of MET, VEGFR2, AXL, FLT3, and KIT were, respectively, assessed in PC3, HUVEC, MDA-MB-231, FLT3-transfected BaF3, and KIT-transfected MDA-MB-231 cells. Cells were serum starved for 3 to 24 hours, then incubated for 1 to 3 hours in serum-free medium with serially diluted cabozantinib before 10-minute stimulation with ligand: HGF (100 ng/mL), VEGF (20 ng/mL), SCF (100 ng/mL), or ANG1 (300 ng/mL). All ligands were purchased from R&D Systems. Receptor phosphorylation was determined either by ELISA using specific capture antibodies and quantitation of total phosphotyrosine (Zymed, Inc.) or immunoprecipitation and Western blotting with specific antibodies and quantitation of total phosphotyrosine (4G10; UpState Biotechnology). Total protein served as loading controls.

Endothelial cell tubule formation assays

Human microvascular endothelial cells (HMVEC) were mixed with serially diluted cabozantinib, then immediately added to cultures of normal human diploid fibroblasts in the presence or absence of 60 ng/mL VEGF for 7 days. Conditioned media from MDA-MB-231, A431, HT1080, and B16F10 cells were also used as a source of secreted growth factors. Following fixation, tubule formation was detected using the endothelial cell marker CD31 (clone WM59; BD Biosciences) and visualized with an immunoperoxidase detection system (Artisan AEC Detection Kit; Dako). Digital images were captured using an inverted microscope (×20), and total tube length was quantitated with Metamorph (Molecular Devices).

Invasion and migration assays

B16F10 cells (2 × 105) were seeded onto 0.8 μm membranes in the upper chambers of a 96-well Transwell plate in the absence or presence of Matrigel (BD Pharmingen; BD Biosciences). Cells were incubated with serially diluted cabozantinib in DMEM/0.1% FBS. The lower chambers contained HGF (50 ng/mL) and the appropriate concentration of cabozantinib. After 24 hours, cells were
recovered from the lower chamber using Accutase (Innovative Cell Technologies). Cell suspensions were incubated with calcein AM (Molecular Probes; Invitrogen) and digital images captured with a fluorescence microscope.

MS1 mouse endothelial cells were plated in 96-well plates coated with fibronectin. A cell-free zone was created that was followed by treatment with serially diluted cabozantinib and 30 ng/mL VEGF or 30 ng/mL HGF. After 20 hours, cell migration was determined from digitally captured images.

In vivo inhibition of receptor phosphorylation
Female nu/nu mice (Taconic) were housed according to the Exelixis Institutional Animal Care and Use Committee guidelines. H441 cells (3 × 10⁶) were implanted intradermally into the hind flank and when tumors reached approximately 150 mg, tumor weight was calculated using the formula: (tumor volume = length (mm) × width² (mm²))/2, mice were randomized (n = 5 per group) and orally administered a single 100 mg/kg dose of cabozantinib or vehicle. Tumors were collected at the indicated time points. Pooled tumor lysates were subjected to immunoprecipitation with anti-MET (SC161; Santa Cruz Biotechnology) and Western blotting with anti-phosphotyrosine MET (pY1230/34/35; BioSource International). After blot stripping, total MET was quantitated as a loading control. In a separate experiment, naive mice (n = 5 per group) were administered a single...
100 mg/kg dose of cabozaantinib or vehicle, followed by intravenous administration of HGF (10 μg per mouse) 10 minutes before liver collection. Analysis of MET phosphorylation in liver lysates was as described above. In a separate experiment, naive mice (n = 5 per group) were administered a single 100 mg/kg dose of cabozaantinib or vehicle, followed by intravenous administration of VEGF (10 μg per mouse) 30 minutes before lung collection. Pooled lung lysates were subjected to immunoprecipitation with FLK1 (SC6251; Santa Cruz Biotechnology) and Western blotting with anti-phosphotyrosine (4G10, Upstate Biotechnology). After blot stripping, total FLK1 concentrations were quantitated as a loading control.

Immunohistochemistry

MDA-MB-231 cells (1 × 10⁶) were implanted subcutaneously into the uncleared mammary fat pad of nu/nu mice. When the tumors reached approximately 100 mg (12 days after implantation), mice were randomized (n = 8 per group) and orally administered cabozaantinib (100 mg/kg) or saline vehicle once daily for 4 days. Body weights were collected daily. Four hours before tumor harvest, animals were injected intraperitoneally with 60 mg/kg pimonidazole to facilitate visualization of tumor hypoxia. Tumors were resected at 4 and 8 hours after the first dose and 4 hours after the second, third, and fourth doses, respectively, snap frozen in optimum cutting temperature (Miles, Inc.) and processed for staining for hypoxia (Hypoxyprobe-1; Chemicon), vascularity [endothelial cell marker MECA-32 (#550563, BD Biosciences)], and proliferation [Ki67 (#RM-9106-S; Lab Vision)]. Apoptosis [terminal deoxynucleotidyl transferase-mediated dUTP nick end labeling (TUNEL)] determination was carried out using an In Situ Cell Death Detection Kit (Roche). Digital fluorescent images were captured using a Zeiss Axioskop 2 with AxioVision (Carl Zeiss MicroImaging LLC) image analysis software. One to 5 nonoverlapping representative fields were captured at ×5 (hypoxia, TUNEL, Ki67), ×100 (MECA-32), and ×400 (endothelial cell death) magnification and quantified using Metamorph (Molecular Devices).

Solid tumor efficacy studies

On day 0 in nu/nu mice, cells were inoculated intradermally into the hind flank (3 × 10⁶ cells, H441) or subcutaneously into a noncleared mammary fat pad (1 × 10⁶ cells, MBA-MB-231). When tumors reached approximately 100 mg (12–14 days after implantation), mice were randomized (n = 10 per group) and treated orally once daily for 14 days with cabozaantinib or saline vehicle. In a separate study, MDA-MB-231 tumors were allowed to reach approximately 500 or 1,000 mg before randomization and initiation of cabozaantinib treatment. On day 0 in female Wistar rats (Charles River Laboratories), C6 cells (5 × 10⁶) were inoculated subcutaneously into the hind flank. When the tumors reached approximately 250 mg (3–4 days postimplantation), rats were randomized (n = 8 per group) and treated orally once daily for 12 days with cabozaantinib or water vehicle. Body weights were collected daily, and tumor weights were collected twice weekly. Percentage of tumor growth inhibition/regression values were expressed as follows: 1 – ([mean treated tumor weight on the final day – mean tumor weight on day 0]/mean vehicle tumor weight on the final day – mean tumor weight on day 0) × 100. Statistical analysis of cabozaantinib-treated tumors versus vehicle-treated tumors or versus predose tumors was done by one-way ANOVA with significance defined as P < 0.05. Blood was collected 4 hours after the final dose, and plasma was prepared to determine cabozaantinib concentrations.

Experimental metastasis

On day 1, MDA-MB-231 cells (1 × 10⁶) were implanted via tail vein injection into nu/nu mice that was immediately followed by randomization (n = 12–15 per group) and oral treatment with cabozaantinib (60 mg/kg), sunitinib (120 mg/kg), or vehicle for 28 days. On day 28, lungs were perfused in situ with India ink solution (15% in PBS), resected, and lung wet weights immediately captured. Lungs were destained and fixed in Fekete’s buffer (70% ethanol, 4% formaldehyde, glacial acetic acid; 20:2:1) for 72 hours, then transferred to 70% ethanol until image acquisition and analysis as previously described (23). Briefly, each lung was separated into 5 lobes, and ventral and dorsal surface aspects were digitally captured with lobes submerged under water. Images were analyzed by Definiens Developer XD to assess lung tumor foci counts. The number of tumor foci for ventral and dorsal lung surface aspects were obtained after applying size threshold parameters to exclude foci with diameters below 4 pixels from analysis (23). As the tumor foci data did not pass the normality and Gaussian distribution tests, statistical analysis was conducted via nonparametric ANOVA (Kruskal–Wallis analysis).

Results

Cabozaantinib inhibits kinase activity

The selectivity of cabozaantinib (Fig. 1A) was profiled against a protein kinase panel of approximately 270 human kinases. Cabozaantinib is a potent inhibitor of MET and VEGFR2 with IC₅₀ values of 1.3 and 0.035 nmol/L, respectively (Table 1). MET-activating kinase domain mutations Y1248H, D1246N, or K1262R were also inhibited by cabozaantinib (IC₅₀ = 3.8, 11.8, and 14.6 nmol/L, respectively). Cabozaantinib displayed strong inhibition of several kinases that have also been implicated in tumor pathobiology, including KIT, RET, AXL, TIE2, and FLT3 (IC₅₀ = 4.6, 5.2, 7, 14.3, and 11.3 nmol/L, respectively). Kinases not potently inhibited include RON, EGFR, IGFR1, and EphA4/B4. In cellular assays, cabozaantinib inhibited phosphorylation of MET and VEGFR2, as well as KIT, FLT3, and AXL with IC₅₀ values of 7.8, 1.9, 5.0, 7.5, and 42 μmol/L, respectively.

IC₅₀ determinations conducted at various concentrations of ATP revealed that cabozaantinib is an
ATP-competitive inhibitor of MET, VEGFR2, TIE2, and FLT3 (data not shown). Kinetic constants for binding to MET and VEGFR2 show tight but reversible binding (Supplementary Table S1).

Cabozantinib inhibits endothelial cell tubule formation in vitro

HMVEC cells were incubated with VEGF in the presence of cabozantinib and tubule formation visualized by immunostaining for CD31. Cabozantinib inhibited tubule formation with an IC50 value of 6.7 nmol/L (Fig. 1B) with no evidence of cytotoxicity, showing that cabozantinib exerts an antiangiogenic rather than cytotoxic effect. Cabozantinib also inhibited tubule formation in response to conditioned media derived from cultures of MDA-MB-231 (IC50 = 5.1 nmol/L), A431 (IC50 = 4.1 nmol/L), HT1080 (IC50 = 7.7 nmol/L), and B16F10 (IC50 = 4.7 nmol/L) cells, suggesting that secreted tumor cell–derived proangiogenic growth factors are unable to circumvent inhibition of tubule formation by cabozantinib.

Cabozantinib inhibits cellular migration and invasion

We have previously shown that HGF promotes MET phosphorylation in B16F10 cells and stimulates migration and invasion (24). Cabozantinib potently inhibited HGF-induced migration (IC50 = 31 nmol/L; Fig. 1C) and invasion (IC50 = 9 nmol/L; Fig. 1D) of B16F10 cells. VEGF- and HGF-mediated migration of proangiogenic murine MS1 endothelial cells (25) were also sensitive to cabozantinib with IC50 values of 5.8 and 41 nmol/L, respectively.

Cabozantinib inhibits tumor cell proliferation in a variety of tumor types

The effect of cabozantinib on proliferation was evaluated in a number of human tumor cell lines. SNU-5 and Hs746T cells harboring amplified MET (26) were the most sensitive to cabozantinib (IC50 = 19 and 9.9 nmol/L, respectively); however, SNU-1 and SNU-16 cells lacking MET amplification were more resistant (IC50 = 5,223 and 1,149 nmol/L, respectively). MDA-MB-231 and U87MG cells exhibited comparable levels of sensitivity to cabozantinib (IC50 = 6,421 and 1,851 nmol/L, respectively), whereas H441, H69, and PC3 cell lines were the least sensitive to cabozantinib with IC50 values of 21,700, 20,200, and 10,800 nmol/L, respectively. In addition, BaF3 cells expressing human FLT3-ITD, an activating mutation in acute myelogenous leukemia (27), were sensitive to cabozantinib (IC50 = 15 nmol/L) when compared with wild-type BaF3 cells (IC50 = 9,641 nmol/L).

Cabozantinib inhibits MET and VEGFR2 phosphorylation in vivo

A single 100 mg/kg oral dose of cabozantinib resulted in inhibition of phosphorylation of MET 2 to 8 hours postdose in H441 tumors that harbor constitutively phosphorylated MET (Fig. 2A). This result is consistent with data showing the sensitivity of these cells to inhibitors selective for MET and MET knockdown by siRNA (28, 29). This effect was reversible, as MET phosphorylation returned to basal levels by 48 hours after treatment. In separate experiments, cabozantinib inhibited in vivo stimulation of MET phosphorylation by HGF in liver hepatocytes (Fig. 2B) and VEGF-stimulated phosphorylation of FLK1 (Fig. 2C) with inhibition of both targets sustained through 8 hours postdose. Furthermore, cabozantinib eliminated endogenous levels of phosphorylated FLK1 that are present in the absence of VEGF stimulation. Plasma concentrations of cabozantinib (Supplementary Table S2) associated with maximal and sustained inhibition of MET and FLK1 were 17 to 34 μmol/L, greater than 3-fold above the MET cellular phosphorylation, VEGF tubule formation, and HGF invasion IC50 values described above.

Table 1. In vitro kinase inhibition profile of cabozantinib

<table>
<thead>
<tr>
<th>Kinase</th>
<th>IC50 ± SD,a nmol/L</th>
<th>Enzyme concentration, nmol/L</th>
<th>ATP concentration, μmol/L</th>
<th>Assay</th>
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<tr>
<td>VEGFR2</td>
<td>0.035 ± 0.01</td>
<td>0.05</td>
<td>3</td>
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<tr>
<td>MET</td>
<td>1.3 ± 1.2</td>
<td>10</td>
<td>1</td>
<td>C</td>
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<tr>
<td>MET (Y1248H)</td>
<td>3.8</td>
<td>13</td>
<td>1</td>
<td>C</td>
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<tr>
<td>MET (D1246N)</td>
<td>11.8</td>
<td>12</td>
<td>1</td>
<td>C</td>
</tr>
<tr>
<td>MET (K1262R)</td>
<td>14.6</td>
<td>12</td>
<td>1</td>
<td>C</td>
</tr>
<tr>
<td>RET</td>
<td>5.2 ± 4.3</td>
<td>15</td>
<td>2</td>
<td>C</td>
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<tr>
<td>TIE2</td>
<td>14.3 ± 1.1</td>
<td>15</td>
<td>5</td>
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<td>AXL</td>
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<td>TBD</td>
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<tr>
<td>FLT3</td>
<td>11.3 ± 1.8</td>
<td>0.5</td>
<td>1</td>
<td>C</td>
</tr>
<tr>
<td>KIT</td>
<td>4.6 ± 0.5</td>
<td>1</td>
<td>3</td>
<td>A</td>
</tr>
<tr>
<td>RON</td>
<td>124 ± 1.2</td>
<td>60</td>
<td>1</td>
<td>C</td>
</tr>
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Abbreviations: A, AlphaScreen; C, Coupled luciferase; R, radiometric; TBD, to be determined.

aMean ± SD of at least 3 independent determinations.
Cabozantinib disrupts tumor vasculature and promotes tumor and endothelial cell death

To examine the effect of cabozantinib on tumor vasculature, antiangiogenic-sensitive MDA-MB-231 cells expressing MET and VEGF were used (28, 30). Tumor-bearing animals were administered a single oral dose of cabozantinib (100 mg/kg) or water vehicle, and levels of phosphorylated and total MET in tumor lysates were determined. B, naive mice were administered a single dose of cabozantinib (100 mg/kg) or water vehicle. Ten minutes before harvest, mice were intravenously administered HGF (10 μg per animal), and levels of phosphorylated and total MET in liver lysates were determined. C, naive mice were administered a single dose of cabozantinib (100 mg/kg) or water vehicle. Thirty minutes before harvest, mice were intravenously administered VEGF (10 μg per animal), and levels of phosphorylated and total FLK1 in lung lysates were determined. Representative Western blot images are shown.

Figure 2. Cabozantinib treatment results in inhibition of phosphorylation of MET in tumor and liver tissue and phosphorylation of FLK1 in lung tissue following a single oral dose. A, mice bearing H441 tumors were administered a single oral dose of cabozantinib (100 mg/kg) or water vehicle, and levels of phosphorylated and total MET in tumor lysates were determined. B, naive mice were administered a single dose of cabozantinib (100 mg/kg) or water vehicle. Ten minutes before harvest, mice were intravenously administered HGF (10 μg per animal), and levels of phosphorylated and total MET in liver lysates were determined. C, naive mice were administered a single dose of cabozantinib (100 mg/kg) or water vehicle. Thirty minutes before harvest, mice were intravenously administered VEGF (10 μg per animal), and levels of phosphorylated and total FLK1 in lung lysates were determined. Representative Western blot images are shown.
together, these results show that cabozantinib disrupts tumor vasculature by inducing endothelial cell death that negatively impacts tumor viability.

**Cabozantinib inhibits tumor growth in a dose-dependent manner**

The *in vivo* efficacy of cabozantinib was evaluated in human tumor models in rodents over a period of time that corresponded to exponential tumor growth of each model. Cabozantinib treatment resulted in significant tumor growth inhibition of MDA-MB-231 tumors (*P* < 0.001) for all doses when compared with vehicle-treated tumors (Fig. 4A). Dose-dependent inhibition was observed for the 3- and 10-mg/kg doses (*P* < 0.01). At the 30- and 60-mg/kg doses, cabozantinib induced stable disease. Continuous treatment at these doses was associated with plasma concentrations of 9,000 to 16,000 nmol/L, which was ~2-fold above IC₅₀ values for cellular proliferation and tubule formation with MDA-MB-231 conditioned media. A single 100 mg/kg dose resulted in sustained MDA-MB-231 tumor growth inhibition for approximately 8 days after which tumors began growing at a rate similar to vehicle-treated control tumors (Supplementary Fig. S1A). In a similar but separate study with larger MDA-MB-231 tumors, a 10-mg/kg dose resulted in induction of stable disease that was independent of initial tumor size (Fig. 4B), whereas a 60-mg/kg dose resulted in significant tumor regression of both 500-mg (37%, *P* = 0.0003) and 1,000-mg (45%, *P* = 0.0001) tumors. Cabozantinib inhibited growth of H441 tumors at all doses (*P* < 0.001), with dose-dependent inhibition observed for the 10- and 30-mg/kg doses (Fig. 4C). The 60-mg/kg dose resulted in significant tumor regression (33%, *P* < 0.005) when compared with predose tumor weights. In a separate experiment, a single 100 mg/kg dose resulted in C6 tumor growth inhibition...
that was sustained during a 15-day observation period (Supplementary Fig. S1B). On a body weight dosage basis, cabozantinib plasma exposures ranged from 6- to 10-fold higher in rats than in mice (Supplementary Table S4), which accounts for lower doses inducing tumor growth inhibition/regression in rats than in mice. Subchronic administration of cabozantinib was well tolerated in mice and rats with no signs of toxicity, as determined by stable and/or increasing body weights during the treatment period (Supplementary Fig. S1C and S1D).

**Cabozantinib does not promote metastasis following intravenous tumor cell inoculation**

The impact of cabozantinib on metastasis to that of VEGFR2-targeting therapies, known to promote metastasis in preclinical models (17), was investigated. Immediately following intravenous injection of MDA-MB-231 cells into mice, oral cabozantinib or sunitinib treatment was initiated and continued once daily for 28 days. Examination of intact lungs from cabozantinib-treated mice revealed no apparent difference in lung surface tumor burden when compared with lungs from vehicle-treated control animals (Fig. 5A); however, lungs from sunitinib-treated animals displayed an apparent increase in tumor burden. Quantitation of tumor foci per lobe surface revealed no statistical difference in the number of foci for vehicle- and cabozantinib-treated mice (56 vs. 64 foci/lobe surface, P > 0.05) and a 2-fold increase in tumor foci for sunitinib-treated mice (109 foci/lobe surface, P < 0.001; Fig. 5B). Additional evidence of inhibition of metastasis was confirmed by lack of a significant increase (P > 0.05) in whole lung wet weights in cabozantinib-treated animals (Fig. 5C). Cabozantinib treatment was well tolerated as determined by stable body weights throughout the 28-day treatment period.

**Discussion**

The MET signaling pathway has been shown to be important in tumor growth, survival, and metastasis and acts synergistically with VEGF to promote angiogenesis (8, 9). Both MET and VEGF are found to be dysregulated in many tumor types, resulting in tumor angiogenesis and tumor cell proliferation and invasion. Because of the synergistic effects of MET and VEGFR signaling, inhibiting both arms of the MET/VEGF axis may offer significant benefit over targeting either pathway individually.
In tumor cells, inappropriate activation of MET occurs through overexpression of wild-type MET or its ligand HGF or as a result of activating mutations in the gene encoding MET. Importantly, cabozantinib potently inhibits both wild-type MET and MET with activating mutations, such as those frequently found in hereditary papillary renal and hepatocellular carcinomas (33, 34). Further evidence for the potent activity of cabozantinib comes from cell proliferation studies, where it robustly inhibited cell lines known to be dependent on MET but not cell lines known to be independent of MET.

As shown in this report, cabozantinib exhibits potent and reversible inhibition of its targets, leading to disruption of cellular processes that have been implicated in angiogenesis and tumorigenesis, including migration and tubule formation. This translates into profound changes in tumor physiology, including widespread endothelial and tumor cell apoptosis, disruptions in tumor vasculature, and increased hypoxia. Subchronic treatment with cabozantinib results in broad antitumor efficacy, and in some models, tumor regression; however, a single dose exhibits antitumor efficacy for approximately 8 to 15 days in the models examined.

We used a variety of tumor models that are driven by diverse mechanisms. H441 cells overexpress constitutively active MET as well as activated EGFR, HER2, and HER3 (35). C6 rat glioma cells overexpress MET and are PTEN deficient (32, 36). While MDA-MB-231 cells do not overexpress MET, they are responsive to HGF and also harbor mutationally activated forms of RAS and RAF (37, 38). None of these cell lines are known to be strictly addicted to or dependent on signaling through MET for proliferation, consistent with their relative resistance to cabozantinib in vitro. However, our studies with cabozantinib suggest that the observed antitumor efficacy is the result of mechanisms affecting tumor angiogenesis and the blockade of invasive tumor growth rather than the result of directly targeting cellular proliferation.

One of the limitations of VEGF-targeted therapies has been a lack of enduring clinical benefit. Bergers and Hanahan recently theorized that tumor cells may develop resistance to antiangiogenic therapies through adaptive mechanisms such as upregulation of alternative proangiogenic signaling pathways and enhancement of metastasis (39). Indeed, VEGFR inhibitors such as sunitinib, sorafenib, cediranib, VEGFR2-targeting antibodies, and neutralizing antimurine VEGF antibodies have all been observed to increase the invasiveness and/or metastatic potential of tumors (17, 18, 19, 40). Ebos and colleagues hypothesized that treatment with sunitinib may lead to upregulation of cytokines that promote metastasis (17), and recent studies have suggested that the MET/HGF pathway is important for proangiogenic and prometastatic signaling in the context of VEGF inhibition (16, 19, 40). In light of these observations, it is interesting that our results show that cabozantinib produces potent antitumor and antiangiogenic effects without increasing tumor metastatic potential, unlike treatment with sunitinib, suggesting that targeting MET and VEGFR2 simultaneously may cut off metastatic escape pathways. A
recent study using RIP-Tag2 transgenic mice similarly showed that treatment with cabozantinib resulted in more extensive tumor shrinkage and decreased tumor invasiveness and metastasis than treatment with vehicle or anti-VEGF antibody. Strikingly, in that study, all of the mice treated with cabozantinib survived until the experiment ended at 20 weeks, whereas none of the vehicle- or anti-VEGF antibody–treated mice survived to that endpoint (40). Together, these data suggest that inhibiting MET and VEGFR2 with cabozantinib effectively blocks the development of MET-driven evasive resistance seen with agents targeting the VEGF pathway alone, thereby providing a more sustained antitumor effect.

Our results provide support for testing cabozantinib in a broad variety of tumors where MET activation has been implicated and/or VEGF pathway inhibitors have shown efficacy. Cabozantinib is currently being studied in clinical trials in a number of tumor types, including medullary and differentiated thyroid, prostate, ovarian, non–small cell lung, hepatocellular, renal cell, and breast cancers, as well as melanoma and glioblastoma (41). In clinical trials, cabozantinib was generally well tolerated, with promising clinical activity and response in multiple tumor types (42–49). The data presented here show that cabozantinib has potent antimetastatic, antitumor, and antiangiogenic activity in preclinical models, and they support the ongoing evaluation of the clinical activity of cabozantinib in patients with a variety of cancers.

Disclosure of Potential Conflicts of Interest

All authors are employees of Exelixis, Inc.

Acknowledgments

The authors thank Dana Aftab and Peter Lamb for critical reading of the manuscript, Barbara Semnini for assistance with the migration images, and Cheryl Chun of BlueMomentum.

Grant Support

The financial support was provided by Exelixis, Inc. The funding for BlueMomentum was provided by Exelixis.

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Received April 7, 2011; revised September 9, 2011; accepted September 13, 2011; published OnlineFirst September 16, 2011.

References


www.aacrjournals.org Mol Cancer Ther; 10(12) December 2011 2307

Published OnlineFirst September 16, 2011; DOI: 10.1158/1535-7163.MCT-11-0264

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Molecular Cancer Therapeutics

Cabozatinib (XL184), a Novel MET and VEGFR2 Inhibitor, Simultaneously Suppresses Metastasis, Angiogenesis, and Tumor Growth

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Mol Cancer Ther 2011;10:2298-2308. Published OnlineFirst September 16, 2011.

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