The Selective Tie2 Inhibitor Rebastinib Blocks Recruitment and Function of Tie2\textsuperscript{Hi} Macrophages in Breast Cancer and Pancreatic Neuroendocrine Tumors

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Abstract

Tumor-infiltrating myeloid cells promote tumor progression by mediating angiogenesis, tumor cell intravasation, and metastasis, which can offset the effects of chemotherapy, radiation, and antiangiogenic therapy. Here, we show that the kinase switch control inhibitor rebastinib inhibits Tie2, a tyrosine kinase receptor expressed on endothelial cells and protumoral Tie2-expressing macrophages in mouse models of metastatic cancer. Rebastinib reduces tumor growth and metastasis in an orthotopic mouse model of metastatic mammary carcinoma through reduction of Tie2\textsuperscript{Hi} myeloid cell infiltration, antiangiogenic effects, and blockade of tumor cell intravasation mediated by perivascular Tie2\textsuperscript{Hi}/Vegf-A\textsuperscript{Hi} macrophages in the tumor microenvironment of metastasis (TMEM). The antitumor effects of rebastinib enhance the efficacy of microtubule inhibiting chemotherapeutic agents, either eribulin or paclitaxel, by reducing tumor volume, metastasis, and improving overall survival. Rebastinib inhibition of angiopoietin/Tie2 signaling impairs multiple pathways in tumor progression mediated by protumoral Tie2\textsuperscript{+} macrophages, including TMEM-dependent dissemination and angiopoietin/Tie2-dependent angiogenesis. Rebastinib is a promising therapy for achieving Tie2 inhibition in cancer patients.

Introduction

The angiopoietin (Ang)/Tie2 kinase signaling pathway is a pivotal angiogenic signaling axis in endothelial cells (1, 2), linked with poor outcome and recurrence in cancer patients (3, 4). Ang/Tie2 signaling is central to the initiation of angiogenesis through vascular remodeling by disrupting endothelial cell interactions. While Ang1 is a Tie2 agonist and has a higher binding affinity to Tie2 than Ang2, Ang2 can act as a context-dependent agonist, although initially described as a Tie2 antagonist (5). Thus, the Ang/Tie2 kinase signaling pathway is an attractive antivascular target (1, 2).

Tie2 is also expressed on a subset of proangiogenic macrophages (i.e., Tie2\textsuperscript{+} macrophages) that are involved in tumor angiogenesis and lymphangiogenesis, as well as in cancer cell intravasation and metastasis (6–11). While antivascular agents (such as bevacizumab and other Vegf-A pathway inhibitors) have shown efficacy in decreasing tumor angiogenesis and disease burden in both preclinical and clinical settings (12, 13), one of the mechanisms of tumor resistance or recurrence after antiangiogenic therapy has been attributed to tumor-infiltrating myeloid cells in response to cell death and hypoxia after vascular regression (14). Of note, Tie2\textsuperscript{+} macrophages are involved in supporting angiogenesis during antiangiogenic therapies (2, 15). Thus, the Ang/Tie2 axis has become an attractive target for inhibiting protumoral functions of Tie2\textsuperscript{+} myeloid cells.

Recently, the paracrine interactions between tumor cells and associated stromal cells, such as fibroblasts, mesenchymal stem cells, and myeloid-derived immune cells among others, have been implicated in a form of transient drug resistance, which has been termed as "environment-mediated drug resistance" (EMDR; refs. 16–18). In particular, tumor-associated macrophages (TAM)
not only promote cancer growth, cancer cell survival, and motility (7, 19–23), but can limit the efficacy of the tumor response to chemotherapy or radiotherapy (24–28). Tie2 activities are known to be proangiogenic, prometastatic, and immunosuppressive in the tumor microenvironment (2, 15, 22, 29). In preclinical studies of mammary carcinoma, Ang2 blockade impeded the association of Tie2 macrophages with the nascent tumor vasculature, thereby suppressing their proangiogenic activity (8) and their prometastatic potential (8, 30).

In mammary carcinoma, cancer cell intravasation and dissemination occur at microanatomical structures on blood vessels of the tumors, called Tumor MicroEnvironment of Metastasis (TMEM). Each functional TMEM is composed of three different cell types in direct physical contact: a tumor cell expressing the actin-regulatory protein Mammalian-enabled (Mena), a perivascular Tie2/Vegf hi macrophage, and an endothelial cell (7, 31). TMEM have been identified in mouse and human mammary carcinomas, and their density correlates with metastatic outcome in breast cancer patients (32–34). High-resolution intravital imaging (IVI) of murine primary breast tumors revealed that TMEM sites induce local and transient dissociation of endothelial cell junctions through which migratory cancer cells intravasate and disseminate to secondary sites (7). TMEM-dependent vascular permeability is particularly localized and is mediated by vascular endothelial growth factor-A (Vegf-A) release from the TMEM-bound Tie2/Vegf macrophage (7).

Biologics that inhibit Ang/Tie2 signaling have been developed, notably angiopoietin-sequestering biologics such as the dual Ang1/Ang2 pepobody AMG-386 (trebananib) and the Ang2-specific monoclonal antibodies MEHD3167 and LC06 (35, 36). In clinical studies, angiopoietin-sequestering biologics increase progression-free survival in patients with metastatic breast cancer, ovarian cancer, and other solid cancers (37, 38). While biologics that sequester Tie2 ligands Ang1 or Ang2 may find clinical utility, there are additional ligands, including Ang4, which activate Tie2 receptors and escape capture by Ang1/Ang2 sequestering biologics (39, 40). Additionally, extracellular signals, including integrins (41, 42) and lysyl oxidase (43, 44), may also activate Tie2-mediated signaling and internalized Tie2 signals to the DNA damage response (44). A selective small molecule inhibitor of Tie2 kinase would be capable of intercepting all of the above activating mechanisms, including those not blocked by the Ang-sequestering biologics.

Herein we report the effects of rebastinib, a potent and selective picomolar inhibitor of the Tie2 receptor tyrosine kinase that inhibits kinase activity by an allosteric “switch control” mechanism, on mammary tumor progression. The goals of this study were to evaluate the cellular effects of rebastinib on both endothelial and macrophage cell populations, its efficacy, both as a single agent and in combination with chemotherapy, in cancer models characterized by Tie2 macrophage involvement, and to further characterize the in vivo effects on myeloid cell composition and vascularization in the tumor microenvironment.

Materials and Methods

Cell lines

CHO-K1 (year of purchase, 2009), EA.hy926 (2012), and HUVECs (various lots from 2012 to 2015) were purchased from the ATCC. All cell lines were cultured as recommended by the supplier, unless otherwise indicated. Media reagents were purchased from Life Technologies or Lonza, Inc. CHO and EA.hy926 cell lines were expanded upon receipt and then frozen at an early passage number in aliquots in liquid N2. Cells were then passaged fewer than 6 months after resuscitation. HUVECs were passaged <10 times upon resuscitation. The ATCC performs STR analysis for characterization. Further STR characterization on CHO-K1 and EA.hy926 was not performed. Mycoplasma testing was performed on a monthly basis for all cell lines (except primary cells such as HUVECs), using the MycoAlert Detection Kit from Lonza, Inc.

Tie2 kinase assay and determination of inhibitor potency

Kinase activity was determined by following the production of ADP from the kinase reaction through coupling with the pyruvate kinase/lactate dehydrogenase system as detailed in Supplementary Data. Percent inhibition values were obtained by comparison of reaction rates with DMSO controls. IC50 values were calculated from a series of percent inhibition values determined at a range of inhibitor concentrations using Prism software (GraphPad). Using this pyruvate kinase/lactate dehydrogenase assay, various concentrations of rebastinib were added to an assay mixture. The dissociation rate constant, koff, was calculated as detailed in Supplementary Data.

Crystallization, data collection, and structure refinement of rebastinib with Tie2

Rebastinib was provided by Deciphera Pharmaceuticals. Purified Tie2 (808-1124) crystals were grown as detailed in Supplementary Data. Diffraction data of Tie2-rebastinib crystals were collected at the Advanced Light Source (ALS) Beamline 5.0.1 at a wavelength of 0.9774 Å using an ADSC Q310r detector. Data were reduced with XDS and scaled to 2.05 Å resolution using XSCALE (45). Tie2-rebastinib crystallizes in space group P4(1) with unit cell dimensions a = b = 63.81 Å, c = 177.45, α = β = γ = 90° with two molecules in the asymmetric unit. The structure was solved using Phaser (46) and the PDB entry 2008 (47) as the search model. The model was completed with COOT (48) and refined with REFMAC (49) using ligand restraint generated with JLigand (50) and one TLS group per chain to Rwork = 0.148 and Rfree = 0.166. Data set and refinement statistics are summarized in Supplementary Table S1.

Western blot assays

Tie2 hi iBMM were generated by immortalizing mouse bone marrow macrophages with an SFFV.TAG.WPRE lentivector, as described (51), and by transducing a mouse Tie2 cDNA under the control of a human PGK promoter. Western blot assays to detect total and phospho-Tie2 in HUVECs, EA.hy926, and Tie2 iBMM were detailed in the Supplementary Data. Antibodies against phospho-Tie2 Tyr992 (catalog #4221) and rabbit IgG (HRP-conjugated; catalog #7074) were obtained from Cell Signaling Technology. The antibody against total Tie2 (catalog #sc-324) was obtained from Santa Cruz Biotechnology, Inc.

HUVEC transwell migration assay

Human umbilical vein endothelial cells (HUVEC; Comin 354151) were maintained in EGM medium (Lonza CC-3124) and used within the first 6 passages for transwell assays detailed in Supplementary Data.
Mice
All studies involving mice were carried out in accordance with the National Institutes of Health regulations concerning the care and use of experimental animals and with the approval of the Animal Care and Use Committee of Molecular Imaging, Inc., an AAALAC accredited facility or with the approval by the Albert Einstein College of Medicine Animal Care and Use Committee. Transgenic mice expressing the Polyoma Middle T (PyMT) oncogene under the control of the mammary tumor virus long terminal repeat (MMTV-LTR) were bred in house at the Albert Einstein College of Medicine and maintained on the FVB background.

Transgenic male C57Bl6/6J/RIP1-Tag2 mice heterozygous for the oncogene were bred with wild-type females. Pups were genotyped for the SV40 large Tag by Transnetx (http://www.transne tx.com). Starting from 12 weeks of age, RIP1-Tag2 mice were maintained on a sucrose-enriched diet and monitored daily. All procedures involving RIP1-Tag2 mice were performed according to protocols approved by the Veterinary Authorities of the Canton Vaud according to the Swiss Law (license 2574 and 2574/a).

PyMT syngeneic breast cancer implant model
Female FVB/NJ mice (JAXWEST:RB05) were implanted in the fourth mammary fat pad on the left side with 1 million cells in serum-free media that had been dissociated from tumor fragments from MMTV-PyMT donor mice. Treatments began on day 31 when the mean tumor burden for all groups in the experiment was 843 mg or on day 52 when the mean tumor burden for all groups in the experiment was 930 mg as detailed in Supplementary Data. For the eribulin survival study in combination with the vehicle group was 1,288 mg. Study progression and necropsy were resected on day 45 when the mean tumor burden for all cohorts are detailed in the Supplementary Data.

FACS analysis
The diverse immune cell populations infiltrating vehicle- or rebastinib-treated PyMT tumors were compared by FACS-based quantification. PyMT tumors were surgically isolated, enzymatically digested and resulting single cell suspensions were stained for flow cytometry analysis. Only viable cells (Indo-1-violet negative) were selected for further analysis from a single-cell population, which was initially sorted based on forward scattering (FSC) to exclude clustered cells. The immune cell compartment of all tumors was then gated based on CD45 expression, while lymphocyte versus non-lymphocyte populations were further discriminated based on side scattering (SSC) of CD45+ cells. A gating strategy was then used in the lymphocyte compartment with quantification selective subpopulations, including cytotoxic T-cells (CD45+ CD8+), T-helper (CD45+ CD4+), T-regulatory (Treg; CD45+ CD4+ FoxP3+ CD25+), and natural killer (NK) cells (CD45+ CD3+ CD16+ CD56+). A gating strategy was used in the nonlymphocyte compartment with quantification the myeloid-derived suppressor cells (MDSC; CD45+ CD11b+ Ly6G+ Ly6C+ Gr1+), as well as macrophages (CD45+ CD11b+ F4/80+ Gr1+). Further gating strategy was deployed to identify M2 macrophage polarization (CD45+ CD11b+ F4/80+ Gr1+ MRC1+) in the macrophage population. Furthermore, Tie2 expression was quantified in both the entire (CD45+ CD11b+ F4/80+ Gr1+ MRC1+) and the M2-only macrophage subpopulation (CD45+ CD11b+ F4/80+ Gr1+ CD206+ MRC1+). The entire gating strategy is outlined in Supplementary Fig. S1.

RIP1-Tag2 transgenic mouse model
Male RIP1-Tag2 mice (11.5- to 12.5-week-old) were treated daily by oral gavage with rebastinib at 10 mg/kg or vehicle control 0.4% HPMC for 4 weeks. Before euthanasia, the mice were retroorbitally injected with FITC-labeled lectin (to reveal perfused blood vessels). Pancreata and livers were harvested at necropsy for analysis. Immunofluorescence staining, lectin labeling of vasculature imaging and quantification of the RIP1-Tag2 (PNET) model is detailed in the Supplementary Data. To quantify liver micrometastases in PNET bearing mice livers were harvested and fixed overnight in 4% PFA solution at 4 °C. The left lobe of each liver was dissected, processed and embedded in optimal cutting temperature compound, as described for pancreata. Analysis of cryosections is detailed in the Supplementary Data.

In vitro intravasation (iTEM) assay
The iTEM assay was performed as described previously (52, 53) and detailed in the Supplementary Data. Transwells were imaged using a Leica SP5 confocal microscope using a 60× 1.4 numerical aperture objective and processed using Imagel (NIH) and IMARIS programs. Quantitation was performed by counting the number of tumor cells that had crossed the endothelium within the same field of view (60×, 10 random fields).

Immunochemistry. Tumor sections were prepared, stained, and quantified as previously described (32).

Quantification of the F4/80+ and Tie2+ percent area staining. After F4/80 and Tie2 immunostaining, whole slides were digitized on a Pannoramic P250 Flash II digital whole slide scanner at 20× magnification. Digital slides were then analyzed in Visiomorph DP (Visiopharm) with a custom developed app specific to mammary gland morphology, as detailed in the Supplementary Data.

Labeling of tumor vasculature and extravasation of 155 kDa dextran-TMR and measuring circulating tumor cells (CTC). One hour after the termination of the experiments with rebastinib, 3 μg of 155 kDa TMR-dextran was administered by tail vein i.v. to label sites of vascular permeability. In addition, CTCs were isolated from anesthetized mice from blood drawn from the right ventricle of the heart and scored. Both were done as described previously (7) and detailed in Supplementary Data.

Immunofluorescence. Tumor sections were prepared and stained as described previously (7) and detailed in Supplementary data. The following primary antibodies were used for immunostaining of mouse tumor tissues: rat anti-mouse CD68 (clone F4-11, Serotec), or Alexafluor488-conjugated CD68 (eBioscience), mouse anti-Mena (NB1-87914, Novus Biologicals), rat anti-ZO-1 (clone R40.76, Millipore), rabbit anti-CD31 (77699s; Cell Signaling Technology), rat anti-Tie2 (16-5987-82; eBioscience), rabbit anti-Idl (019-19741; Wako). Sections were washed with PBS, and the primary antibodies were detected with Alexafluor488, 555, or 647 secondary antibody conjugates (Molecular Probes/Invitrogen) and nuclei stained with 4,6-diamidino-2-phenylindole (DAPI).

TME immunohistochemistry. Tumor sections were prepared, stained, and quantified as previously described (32).
Plasma Ang2 levels in rebastinib-treated patients

Post-dose (C_{max}) and pre-dose trough plasma samples on cycle 1 day 8 and pre-dose trough plasma samples on cycle 1 day 22 were analyzed for levels of Ang2 and rebastinib concentration detailed in the Supplementary Data.

Statistical analysis

Individual animals in each cohort are presented as individual points on a dot plot. A horizontal line indicates the mean value, and the error bars represent the standard error of the mean. Statistical significance was determined by the comparison of the means of two groups using an unpaired, two-sided t test using Prism (Graph Pad Inc.). Data sets were checked for normality (D’Agostino–Pearson omnibus normality test or Shapiro–Wilk normality test) and unequal variance using Prism (Graph Pad Inc.). Welch’s correction was applied to t tests as needed. P values of less than 0.05 were deemed statistically significant.

Results

X-ray cocrystal structure of rebastinib with Tie2 kinase exhibits unique attributes of Type II switch control binding

A cocrystal structure of rebastinib was obtained in complex with the unphosphorylated human Tie2 kinase domain (residues K808–A1124) at 2.05 Å resolution (Supplementary Table S1; Fig. 1A and B and Supplementary Fig. S2A and S2B). Analysis revealed a Type II binding mode in which rebastinib induces Tie2 into a DFG-out inactive enzymatic conformation, making key interactions with regions of the activation loop.
conformation-controlling switch (yellow ribbon, Fig. 1A), regions of Tie2 which serve as the cognate switch pocket (dashed red oval, Fig. 1A), and the ATP hinge region.

For Tie2 conformational activation, the switch is required to bind into its cognate switch pocket. Rebastinib outcompetes the switch for binding into this pocket. Thus, in Fig. 1A and B, rebastinib is shown to occupy the Tie2 switch pocket (dashed red oval), and the activating switch (yellow) is displaced into an inactive conformation, precluding the ability of Tie2 to activate. An annotated description of molecular interactions of rebastinib with Tie2 is shown in Supplementary Fig. S2A and S2B. Rebastinib binding outcompetes the switch by the inhibitor t-butyl moiety (A) displacing switch residue Phe983 (labeled as 1). The fluoro-phenyl ring of the inhibitor (B) additionally stabilizes Phe983 in this inactive state through π-stacking interactions. Inhibitor urea moiety (D) forms hydrogen bonds with both the conserved catalytic salt bridge Glu872 (9)/Lys855 (10) and the switch residue Asp982 (6). This binding modality further nucleates a hydrogen bonding network comprising residues His962, Asp964, Arg968, Asn969, Asp982, and Gly984 (residues 2–7), collapsing these residues into an inactive conformational state incompatible with enzymatic shuttling of phosphate from ATP to a protein substrate. A unique electrostatic π-stacking interaction is formed between switch residue Arg987 (8), the inhibitor quinolinyl ring (C) and Glu872 (9), highlighted with double-tipped red arrows. In composite, rebastinib directly forms 5 hydrogen bonds with Tie2, with overall binding further nucleating an additional 25 hydrogen bonds inducing the switch, switch pocket, and catalytic residues into an inactive state. The hydrogen bond network additionally is consistent with stabilizing the C-terminal inhibitory motif (green) in position for occluding the substrate binding pocket of Tie2. This conformational state of Tie2 induced by rebastinib is incapable of loading a protein substrate or ATP cofactor and provides structural determinants of the picomolar potency of rebastinib (Fig. 1C). Biophysical thermal melt studies also confirmed potent binding of rebastinib to Tie2, resulting in a ΔTm of 14.9°C versus apo-Tie2 (Supplementary Fig. S3A–S3C).

Biochemical inhibition of Tie2 and prolonged off-rate

In biochemical assays, rebastinib demonstrated subnanomolar potency as an inhibitor of recombinant human unphosphorylated Tie2 kinase (IC50 = 0.63 nmol/L, Fig. 1C). This IC50 exceeded the limit of further titration based on the concentration of Tie2 required for the assay. Rebastinib retained subnanomolar potency at adenosine triphosphate (ATP) concentrations as high as 4 nmol/L (Fig. 1D). Rebastinib was found to slowly dissociate from Tie2, yielding a dissociation rate constant (koff) value of 0.0012/minutes (Fig. 1C and E). The t1/2 value associated with this koff value for recovery of kinase activity was approximately 10 hours for unphosphorylated Tie2 (Fig. 1C).

In assays with a more complex cellular environment using human umbilical vascular endothelial cells (HUVEC) or EA.hy926 cells, which express Tie2, rebastinib inhibited Ang1-stimulated Tie2 kinase activity with IC50 values of 0.058 and 0.091 nmol/L, respectively (Fig. 1C and Supplementary Fig. S4A–S4D). In bone marrow–derived murine macrophages transduced to express Tie2 kinase (IBMM Tie2E18), rebastinib inhibited Tie2 phosphorylation with an IC50 value of 0.26 nmol/L (Fig. 1C and Supplementary Fig. S4E and S4F). In a functional chemotaxis assay, rebastinib inhibited Ang1-mediated migration of HUVECs with an IC50 of 0.022 nmol/L (Fig. 1C and Supplementary Fig. S5A and SSB). Finally, in transiently transfected Chinese Hamster Ovary (CHO) cells overexpressing constitutively phosphorylated Tie2, rebastinib inhibited Tie2 activity with an IC50 value of 2.0 nmol/L (Fig. 1C). Rebastinib demonstrated a prolonged off rate (>24 hours) against Tie2 kinase in transfected CHO cells after inhibitor washout (Fig. 1F).

Rebastinib was evaluated in a panel of 300 human kinases, confirming its selectivity as a Tie2 inhibitor (Supplementary Table S2 and Fig. 1G). TRKA was identified as the nearest neighbor kinase inhibited by rebastinib (Supplementary Table S2). The cellular IC50 for inhibition of TRKA phosphorylation by rebastinib was 0.17 nmol/L, which is 3-fold higher than the IC50 of Tie2. Related kinases TRKB and TRKC exhibited IC50 values of 0.42 nmol/L and 2.74 nmol/L, respectively, showing lesser inhibition by rebastinib. BCR-ABL and FLT3 inhibition by rebastinib were even less potent with IC50 values 62- and 71-fold, respectively. Rebastinib has previously been shown to inhibit the BCR–ABL fusion oncoprotein (54). Because rebastinib is approximately 62-fold more selective in binding to Tie2 than BCR-ABL, the effect of rebastinib as a selective Tie2 inhibitor has been further examined in mammary carcinoma and pancreatic neuroendocrine tumors (PNET).

Other marketed, clinical stage, or preclinical compounds have been reported to inhibit Tie2 kinase. Supplementary Table S4 lists these representative agents, including Met inhibitors cabozantinib, foretinib, MGCD265 (glestatinib), crizotinib; BCR-ABL inhibitor ponatinib, ALK/RON inhibitor crizotinib; and other Tie2 inhibitors pexmetinib, and the preclinical tool compound SKB-Tie2. Rebastinib is 17–5,672-fold more potent than these comparator inhibitors (column 3, Supplementary Table S4) for inhibiting Tie2. Moreover, rebastinib is the most selective and potent Tie2 inhibitor versus these comparator compounds (columns 4–14, Supplementary Table S4). By way of example, rebastinib is 7,017-fold more potent for Tie2 versus Met, compared with cabozantinib, foretinib, and MGCD265, all of which exhibit inverse selectivity of less than 1-fold, indicating preferential potency for Met. Rebastinib and SKB-Tie2 are the only inhibitors in Supplementary Table S4, which inhibit Tie2 as their primary target, and rebastinib is 5,672 times more potent than SKB-Tie2 for inhibiting Tie2.

Rebastinib inhibits mammary carcinoma growth and extends mouse survival

To examine rebastinib inhibition of Tie2 signaling in vivo, we used the orthotopic mouse mammary cancer implant model in which the polyoma middle T antigen is under the control of the mouse mammary tumor virus long terminal repeat (MMTV-PyMT; refs. 55, 56). PyMT tumors exhibit histology similar to human luminal breast cancer and progress to metastasis (57). Further, PyMT tumors assemble TME, the vascular doorway structures associated with tumor cell intravasation and dissemination in mouse mammary tumors and human breast cancer patients, which contain Tie2/VEGFA/macrophages (7, 32, 34, 58).

Rebastinib significantly reduced primary tumor growth by 75% and, more importantly, when used in combination with paclitaxel, rebastinib had an additive effect in reducing tumor growth by 90% of control (Fig. 2A). Eribulin, like paclitaxel, is an inhibitor of microtubules and is approved for the treatment of
Rebastinib Inhibits TIE2⁺ Macrophages

Figure 2.
Rebastinib blocks primary tumor growth, inhibits metastatic growth and extends survival in combination with chemotherapy. **A**, Mean tumor burden in mice with implanted PyMT tumors in the mammary fat pad and treated with vehicle (black line), paclitaxel alone (red line), rebastinib alone (blue line), or rebastinib and paclitaxel in combination (green line). **B**, Intermittent treatment of PyMT mice with rebastinib extends overall survival in combination with eribulin, following the excision of the primary tumor.Primary mammary tumors were resected on day 42, and treatment was begun on day 45. Dosing was continued for 27 weeks and animals followed for survival. **C**, H&E staining of lung tissue from PyMT-bearing mice for lung metastases (vehicle-treated, left; rebastinib-treated, right). **D**, Quantification of area of lung metastases by H&E in the lungs of animals as compared with the vehicle control for the indicated treatments after 3 weeks of treatment. **E**, Incidence of lung metastases (nodules) determined by μCT in mice at 9 weeks of treatment. **F**, Rebastinib decreases PNET metastasis to liver, as shown in the Rip1-Tag2 mouse model. HPMC n = 11, rebastinib n = 10. *, P < 0.05.

Relapse of refractory metastatic breast cancer. Rebastinib was evaluated in a postsurgical (adjuvant) setting in the PyMT model in combination with eribulin after primary mammary tumor resection (Fig. 2B). In the absence of the primary tumor, the duration of animal survival is a result of overall systemic tumor burden (primarily lung metastases). Eribulin single agent afforded a median survival of 54 days post tumor resection (red curve), less than median survival of 84 days in the vehicle cohort (black curve). Eribulin in combination with rebastinib extended survival of mice to at least 196 to 200 days, depending on the rebastinib dosing schedule (Fig. 2B). Together, these data demonstrate that rebastinib treatment in PyMT mammary tumors either daily, or through intermittent dosing alone or in combination with chemotherapy, reduces primary tumor volume and extends survival time.

Rebastinib reduces lung metastases in mammary carcinoma and liver metastasis in PNETs

TIE2⁺ macrophages are known to support tumor cell dissemination and metastasis in mammary carcinomas (7, 8, 59). Given that lung is the most frequent site of distant metastasis in breast cancer, we hypothesized that inhibition of TIE2⁺ macrophage-mediated dissemination by rebastinib would reduce tumor cell metastasis to the lung. Indeed, in orthotopic PyMT tumors, rebastinib was found to reduce the formation of lung metastases alone by 72% and in combination with paclitaxel by 93% (Fig. 2C and D). Given the efficacy of intermittent dosing of rebastinib in extending survival and reducing lung metastases, the effect of rebastinib in combination with paclitaxel therapy on metastasis was evaluated in the PyMT model by microCT (Supplementary Fig. S6). As shown (Fig. 2E), there was a difference (P < 0.1) in the incidence of lung metastasis afforded by the combination of paclitaxel with rebastinib dosed twice weekly at 10 mg/kg compared with chemotherapy-treated or vehicle-treated mice.

Unlike PyMT mammary carcinoma, daily oral dosing of rebastinib at 10 mg/kg for 4 weeks did not detectably inhibit the growth of pancreatic insulinomas in the transgenic RIP1-Tag2 model of PNET (60; Supplementary Fig. S7). However, we evaluated rebastinib for inhibition of liver metastases. According to a late treatment protocol targeting established PNETs, 11–12-week-old male RIP1-Tag2 mice were treated for 4 weeks with rebastinib and liver micrometastasis were quantified at termination. Rebastinib significantly inhibited liver metastases by 75% compared with control (Fig. 2F).

Rebastinib reduces TIE2⁺ macrophages and microvessel density in mammary carcinoma and PNETs

TIE2⁺ macrophages have been associated with tumor angiogenesis, tumor cell dissemination, and metastasis (10–12). Because treatment with rebastinib reduced metastasis to the lungs...
in the PyMT mammary carcinoma and liver metastasis in the PNET model, we sought to investigate the effects of rebeastinib on cell infiltration into the tumor and, in particular, Tie2⁺ macrophage infiltration. FACSs sorting of whole PyMT mammary tumors and their associated peripheral stroma showed that rebastinib significantly decreased both total and Tie2⁺ macrophages (Fig. 3A-C) but did not significantly affect lymphoid immune cell composition (Supplementary Fig. S8). There was a trend toward reduction of M2-polarized macrophages (Fig. 3B) and the subset of M2 macrophages expressing Tie2 (Supplementary Fig. S8); however, this reduction did not achieve significance. In a separate analysis, multichannel immunofluorescence (IF) staining (Iba1 for macrophages, CD31 for endothelium and Tie2) was used to specifically distinguish the Tie2⁺ macrophage subpopulation (Fig. 3D and E) and demonstrated that the Tie2⁺ macrophage subpopulation was significantly decreased in response to rebastinib treatment (Fig. 3D and E and Supplementary Fig. S9A), confirming the reduction in the Tie2⁺ macrophage subpopulation in response to rebastinib, as also determined by FACSs analysis (Fig. 3C).

In the PNET model, there was a decrease in Tie2 staining in infiltrate (F4/80⁺ area; Fig. 3G). However, rebastinib significantly decreased intratumoral Tie2⁺ macrophages (Fig. 3H) and led to a significant decrease in protumoral Mrc1⁺ F4/80⁺ macrophages at the tumor invasive front (Fig. 3I and J). Taxane-based chemotherapies, such as paclitaxel, mobilize bone marrow–derived mesenchymal and endothelial progenitors,
Rebastinib inhibits Tie2+ macrophages

Due to the role of Tie2+ macrophages in vascular permeability and tumor cell intravasation at TMEM in vitro (Fig. 6A; ref. 7), we studied whether inhibition of Tie2 with rebastinib would impair TMEM–mediated tumor cell transendothelial migration in vitro. We studied this by using the previously established in vitro TMEM-dependent subluminal-to-luminal transendothelial migration (intravasation) assay (Fig. 6B; refs. 52, 53). We found that rebastinib significantly inhibits macrophage-dependent transendothelial migration of breast cancer cells in the intravasation direction with an IC50 between 50 and 100 pmol/L, and to background levels observed in the absence of macrophages (TC alone) at 500 pmol/L (Fig. 6C, left).

To determine if the Tie2+ macrophages are the cells inhibited by rebastinib in this assay, the intravasation assay was conducted using the complete dose–response evaluation of rebastinib but in the absence of macrophages. When the Tie2+ macrophages were selectively excluded from the intravasation assay, rebastinib had no effect on tumor cell intravasation at any concentration (Fig. 6C, right). The requirement of macrophage-specific Tie2 expression for TMEM-mediated transendothelial tumor cell migration was demonstrated by using murine macrophages in the intravasation assay that expressed either low or high levels of Tie2 (Supplementary Fig. S10). As shown in Fig. 6D, using macrophages expressing high levels of Tie2 (Tie2Hi) resulted in a greater than 4-fold increase in tumor cell transendothelial migration compared with background (Bg), whereas using macrophages expressing low levels of Tie2 (Tie2Lo) did not support such a large increase. Rebastinib treatment ablated transendothelial migration induced by the Tie2Hi macrophages (Fig. 6D). Collectively, these data indicate that Tie2 signaling in TMEM-associated macrophages is required for tumor cell transendothelial migration. These data also demonstrate that inhibition of Tie2 in the intravasation assay selectively impairs Tie2Hi macrophages and does not impair endothelial monolayer integrity.

Rebastinib therapy results in an Ang2 biomarker readout in human patients

In order to clinically evaluate Tie2 inhibition, Ang2 levels were examined in 20 consented patients from a first-in-human phase 1 chronic myelogenous leukemia (CML) and acute myelogenous leukemia (AML) study (NCT 00827138) before and after treatment with 100 or 150 mg twice weekly rebastinib (Fig. 7). Clinically, compensatory elevations in circulating growth factor ligands is observed upon inhibition of their cognate receptor tyrosine kinases, including Vgfr2/Vegfr2,
Figure 4.
Rebastinib reduces paclitaxel-induced increases in Tie2⁺ macrophages and microvessel density. A, Quantification of total macrophages and B, Tie2⁺ macrophages in PyMT tumors in response to control (V), paclitaxel (Pxl), and rebastinib + paclitaxel (Rb + Pxl). C, IF staining of vasculature with CD31 (green) in PyMT tumors in mice treated with the vehicle and drug combinations shown. Bar, 50 μm. D, Quantification of microvessel density as a percentage of area from CD31 staining shown in C. **, P < 0.05; *** P < 0.01; **** P < 0.0005. E, IF staining of vasculature with lectin (green) in PNET tumors treated with vehicle or rebastinib. F, Quantification of microvessel density as a percentage of perfused lectin area from tumor tissue stained in E showing that rebastinib significantly reduces vascular density in PNET. Each dot is a mouse in which multiple tumors were analyzed. HPMC n = 12, rebastinib n = 8. ***, P = 0.0008. Paclitaxel at 10 mg/kg i.v., every five days (n = 10); rebastinib at 10 mg/kg orally, daily.
Csf1/Csf1R, and others (62, 63). Thus, the effect of rebastinib on compensatory increases in circulating angiopoietin levels secondary to Tie2 inhibition was evaluated. On day 22 after starting rebastinib treatment, increases in Ang2 were observed in 19 of 20 patients, no change was observed in one patient, and a \( \geq 2 \)-fold increase in Ang2 plasma levels was observed in 14 of 20 (70%) patients. The average fold increase for all 20 patients was 2.6 \( \pm \) 1.2 SD. To assess whether changes in plasma levels of Ang2 were correlated with plasma levels of rebastinib, data were fit to linear regression models. Both \( C_{\text{max}} \) (Supplementary Fig. S12A) and trough exposures of rebastinib on day 8 (Supplementary Fig. S12B) significantly correlated with increased levels of plasma Ang2. Increased plasma Ang levels also correlated with trough rebastinib exposures on day 22 (Supplementary Fig. S12C).

**Discussion**

Here, we report that rebastinib is a novel switch control inhibitor of Tie2 tyrosine kinase with picomolar potency for blocking Tie2 enzymatic and cellular activity in endothelial cells and TEMs and exhibits efficacy in malignant mammary carcinoma and PNETs. In mammary tumors, Tie2 blockade by rebastinib results in inhibition of tumor growth, invasion, and metastasis. Examination of the effects of rebastinib at the cellular level demonstrates that rebastinib reduced tumor

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**Figure 5.**

Inhibition of Tie2 with rebastinib impairs TMEM function and tumor cell dissemination in PyMT mammary carcinoma. A, IHC staining of tumors for TMEM before and after treatment with rebastinib. Vasculature (endomucin, blue), Mena overexpressing tumor cells (Mena, pink) and macrophages (Iba1, brown). TMEM are circled. B, Quantification of TMEM density in 10 high-power fields (HPFs; no significant difference). C, IF imaging of tumor sections. Tumors are stained for blood vessel endothelial cells (CD31), vascular perfusion/leakage (155 kDa dextran-TMR), cell nuclei (DAPI), or endothelial junctions (ZO-1). D–G, Vehicle or treatment with 10 mg/kg rebastinib both for twice weekly for 3 weeks. D, Extravascular 155 kDa dextran-TMR (vehicle, \( n = 7 \); rebastinib \( n = 9 \); \( * \), \( P = 0.042 \)). E, Normalized quantification of CTCs (\( *\), \( P = 0.004 \)). F, Vascular ZO-1 staining intensity relative to CD31 staining (\( *\), \( P = 0.0124 \)). G, Quantification of TMEM activity as extravascular dextran relative to ZO-1 staining intensity (vehicle, \( n = 7 \); rebastinib \( n = 9 \); \( *\), \( P = 0.0177 \)). All panels: V, vehicle control; Rb, rebastinib.
A. Streaming and invasion

B. Intravasation

C. Fold change intravasation

D. Macrophages

E. + Macrophages

- Macrophages

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vascular density, and Tie2⁺ macrophages in the PyMT mammary tumor and its stroma. However, rebastinib did not lead to a change in immune cell composition beyond the changes observed in Tie2⁺ macrophages. Though the mechanism of such decreased Tie2 expression is not clear, we speculate that the cross-activation of Tie2 by endothelial integrins (41) can be potentially blocked by rebastinib-mediated inhibition of Tie2, further leading to downregulation of the Tie2 receptor from the cell surface. Tie2 internalization has also been reported after ionizing radiation therapy, indicating that other treatment modalities can also mediate Tie2 downregulation (43).

We noticed that rebastinib treatment significantly reduced the general macrophage population in the PyMT mouse model of breast cancer, but not in the transgenic mouse model of PNET, Rip1-Tag2 (Fig. 3A and G). Importantly, however, the intratumoral Tie2⁺ macrophages were reduced in both animal models tested (Fig. 3C and H), indicating that rebastinib affects the dynamics of specific macrophage subpopulations in the primary tumor microenvironment in a context-dependent manner, thus without necessarily affecting the total macrophage population, as also evidenced by radiotherapy, chemotherapy, or other drug treatments (24, 27, 64, 65). In addition, in PyMT tumors, although TMEM density was not decreased by rebastinib, TMEM function was dramatically impaired by rebastinib effects on Tie2⁺ macrophages, as evidenced by decreased vascular permeability, CTCs, and metastasis in vivo, and decreased TMEM-associated transendothelial migration in vitro. The effects of rebastinib are further elevated when combined with paclitaxel or eribulin treatment in mice at least in part because rebastinib significantly inhibits the recruitment of Tie2⁺ macrophages in response to paclitaxel, thereby affecting microvessel density and TMEM assembly.

Rebastinib is a potent picomolar inhibitor of Tie2 kinase that inhibits by an allosteric “switch control pocket” mechanism. Analysis of the cocystal structure of rebastinib with Tie2 revealed a Type II binding mode in which rebastinib induces Tie2 into a DFG-out inactive enzymatic conformation, making key interactions with regions of the conformation-controlling switch, with regions of Tie2 that serve as the cognate switch pocket, and with the kinase hinge region. This binding mode engenders picomolar potency for rebastinib inhibition of Tie2 biochemically and in cellular macrophage and endothelial assays. Cellular potencies for inhibiting murine Tie2 (macrophage assays: IC₅₀ values ~0.075 to 0.26 nmol/L) and human Tie2 (endothelial cell assays: IC₅₀ values ~0.058 to 0.91 nmol/L) are comparable. Administration of rebastinib to high interspecies identity (99%) with only 3 differences out of 355 residues in the kinase domain (Supplementary Fig. S13). The Tie2 kinase binding mode of rebastinib results in a long-dissociation off rate (t₁/₂ = 10 hours).

Tie2⁺ macrophages are a subset of highly skewed protumoral macrophages in the tumor microenvironment that are elevated following treatment with chemotherapy, antiangiogenic agents, or vascular disrupting agents that render tumors hypoxic (6, 8, 9, 15). Tie2⁺ macrophage recruitment contributes to evasive revascularization and invasion in these hypoxic tumors (66). Perivascular Tie2⁺ macrophages have been shown to participate in the doorways for intravasation of tumor cells called TMEM (7, 32, 34, 52). Administration of rebastinib to orthotopic models of mammary carcinoma and PNETs resulted in reduced vascular density, reduced Tie2⁺ macrophages, and...
TMEM function. In addition to efficacy as an antivascular agent, rebastinib reduces the proangiogenic Tie2+ macrophage population in the tumor microenvironment, limiting revascularization and tumor growth. In contrast to other antivascular agents, rebastinib has the potential to not only decrease tumor volume in combination with chemotherapy, but may also prevent Tie2+ macrophage-mediated tumor regrowth and increased TMEM function, leading to metastasis. This is further supported by the evidence that rebastinib reduces tumor cell invasion, dissemination, and metastasis in both the PyMT and RIP1-Tag2 models of invasive and metastatic cancers, and reverses paclitaxel-induced increased intratumoral Tie2+ macrophages and eribulin-induced reduction in survival secondary to lung metastases in the PyMT model.

The finding that rebastinib suppresses the infiltration of Tie2+ macrophages into tumors raises secondary questions involving how rebastinib limits recruitment of Tie2+ macrophages and how the function of isolated Tie2+ macrophages may be altered in response to rebastinib. Recent studies have shown that local vascular endothelial production of Ang2 recruits Tie2+ macrophages to the tumor indicating that inhibition of Ang2 stimulation of the Tie2 receptor on macrophages by rebastinib is involved in suppressing Tie2+ macrophage recruitment (67). Furthermore, recent studies have shown that the inhibition of Vegf production by, or its release from, Tie2+ macrophages blocks their ability to support angiogenesis-dependent tumor relapse (9) and tumor cell dissemination by TMEM (7).

Mechanistically, tumor cell dissemination in mammary carcinoma occurs exclusively at Tie2Hi/Vegf-AHi TMEM sites. Motile tumor cells cross the endothelium when Tie2Hi/Vegf-AHi macrophages in TMEM locally dissolve vascular junctions through Vegf-A signaling (7). Rebastinib impaired TMEM function in vivo, resulting in reduced vascular permeability and tumor cell intravasation, and a corresponding dramatic decrease in CTCs. Decreased tumor cell intravasation from the primary tumor prevented tumor cell metastasis, explaining the decrease in metastatic lung nodules with rebastinib treatment and further supporting the potential impact of rebastinib in preventing Tie2+ macrophage-mediated tumor metastasis. It is also noted that the effects of rebastinib in vivo for inhibiting TMEM function phenocopy macrophage ablation in the MaFIA model and selective gene knockout ablation of Vegf in Tie2+ perivascular macrophages (7).

The observation that rebastinib specifically blocks the Tie2Hi macrophages in TMEM, thereby inhibiting tumor cell intravasation has been further confirmed in the in vitro assay replicating transendothelial tumor cell migration at TMEM. In the presence of 0.5 nmol/L rebastinib, tumor cell intravasation was reduced to background levels of tumor cell intravasation as measured in the absence of macrophages. No effect of rebastinib on tumor cell transendothelial migration was demonstrated in the absence of macrophages, indicating a specific effect on macrophages and no effect of rebastinib on either tumor cell or endothelial cell components of this assay. Furthermore, incorporation of Tie2+ macrophages into the intravasation assay led to a significant increase in tumor cell transendothelial migration compared with macrophages with lower Tie2 expression. Rebastinib reversed the enhanced tumor cell migration induced by Tie2+ macrophages. Collectively, these results support a significant role of Tie2+ macrophages in transendothelial migration and demonstrate that rebastinib specifically blocks Tie2+ macrophages in TMEM which impairs an essential mechanism in tumor cell intravasation and metastasis in mammary carcinoma that is unique to TMEM.

Although the studies included herein demonstrate the potent antiangiogenic and antimacrophage properties of rebastinib to blunt tumor growth, invasiveness, metastasis, and improve survival in murine cancer models, it is also known that the angiopoietin–Tie2 signaling axis is immunosuppressive and blockade of this axis could also provide the potential to favorably modulate cancer treatment outcomes via reversal of Tie2 signaling cross-talk to the adaptive immune system. Tie2+ macrophages have been shown to release IL10 and thereby suppress CD8+ cytotoxic T cells, expand immunosuppressive Tregs, and cause dendritic cell anergy (22, 29). It has also been reported that in long-term responding subjects treated with anti-CTLA4 agents, a humoral reaction is generated against angiopoietins (68). Dual blockade of Ang2 and Vegf-A has been demonstrated to induce an antiangiogenic effect that facilitated extravasation and perivascular accumulation of activated, IFN-γ expressing cytotoxic T lymphocytes in various tumor models, and therapeutic benefit was enhanced by combination with PD-1 blockade (69). Recently, it has also been reported that circulating Ang2 levels are predictive and prognostic for treatment outcomes in melanoma patients treated with checkpoint inhibitors. High pretreatment circulating Ang2 was associated with reduced overall survival in patients treated with CTLA4 and PD-1 checkpoint therapies. Ang2 expression was associated with increased CD163+ macrophage tumor infiltration and Ang2 increased PD-L1 expression on M2-polarized macrophages (67).

Clinical effects of rebastinib on Ang2 plasma levels were evaluated in a recently completed first-in-human phase I trial in CML patients (70). Although not a solid tumor clinical study, during the course of this first-in-human trial, informed consent was obtained from 22 patients to explore treatment versus baseline values of circulating Ang2 as a readout of Tie2 target engagement. Rebastinib treatment led to a significant compensatory increase in Ang2, indicative of systemic Tie2 inhibition. Because Ang2 is primarily secreted from endothelial cells lining the vasculature, it is likely that this Tie2 response is due to endothelial cell inhibition of Tie2 by rebastinib. These results indicate that changes in circulating levels of Ang2 may provide a circulating biomarker of rebastinib-induced Tie2 inhibition in future solid tumor trials.

There is now compelling evidence for Ang2 to function as a (partial) Tie2 agonist in tumor angiogenesis (71). This is also supported by the finding that Ang2 is upregulated in many tumor types, including breast cancer, and its levels may indeed exceed those of Ang1 (4, 72–74). In these contexts (high Ang2/low Ang1), Ang2 may function as a Tie2 activator (75). Therefore, we reason that decreasing Tie2 activity in tumors (e.g., by rebastinib) may phenocopy the effects of Ang2 blockade. We and others have shown that Ang2 blockade in several cancer models, including MMTV-PyMT mice, leads to decreased tumor angiogenesis, which is associated with the "normalization" of the remaining blood vessels (2, 8, 69, 76). Therefore, our finding that Tie2 inhibition increases pericyte coverage of the tumor blood vessels (a parameter of vascular normalization) is consistent with the expected outcome of blocking the Ang2–Tie2 axis in an angiogenic microenvironment. This is particularly true in the presence of Vegf-A, which prevents more profound vascular pruning in the absence of Ang2. The increase in serum Ang2 after
Rebastinib may be consistent with the hypothesis (Fig. 7).

In the current study, we described the preclinical validation for the use of rebastinib in two animal models; breast carcinoma (PyMT) and PNET. As rebastinib is intended for use in human patients, we have recently demonstrated in another study (77) the effects of rebastinib on breast cancer patient-derived xenografts. Rebastinib mediated suppression of TMEM-dependent dissemination, with or without the presence of neoadjuvant chemotherapy. Based on data presented in the Karagiannis and colleagues (2017) study (77) along with the current study, rebastinib has been approved (currently in phase Ib) for a clinical trial in breast cancer patients (NCT02824575). Taken together, rebastinib is a promising therapy for the treatment of solid carcinomas, characterized by prominent Tie2+ macrophage presence in the tumor microenvironment.

Disclosure of Potential Conflicts of Interest

B.D. Smith, C.B. Leary, M.D. Kaufman, W.-P. Lu, G. Al-Ani, and D.L. Flynn have ownership interest (including patents) in Deciphera Pharmaceuticals. J.S. Condeelis is a consultant/advisory board member for Deciphera. No potential conflicts of interest were disclosed by the other authors.

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References


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Grant Support

This research was funded by Deciphera Pharmaceuticals, LLC, the Department of Defense Breast Cancer Research Program under award number (W81XWH-13-1-0010 A.S. Harney and W81XWH-14-1-0286 J. Pignatelli), NIH CA100324, CA150344, CA15330-4553, S10 OD019961, and the Integrated Imaging Program at Albert Einstein College of Medicine.

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Received March 17, 2017; revised July 12, 2017; accepted August 10, 2017; published OnlineFirst August 24, 2017.

Published OnlineFirst August 24, 2017; DOI: 10.1158/1535-7163.MCT-17-0241

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