Effect of mini-tyrosyl-tRNA synthetase on ischemic angiogenesis, leukocyte recruitment, and vascular permeability

Gang Cheng,1 Hua Zhang,1 Xianglei Yang,2 Eleni Tzima,1 Karla L. Ewalt,1 Paul Schimmel,2 and James E. Faber1

1Department of Cell and Molecular Physiology, The University of North Carolina at Chapel Hill, Chapel Hill, North Carolina; 2Department of Molecular Biology, The Scripps Research Institute, La Jolla, California; and 3aTyr Pharma, San Diego, California

Submitted 19 June 2008; accepted in final form 14 August 2008

Cheng G, Zhang H, Yang X, Tzima E, Ewalt KL, Schimmel P, Faber JE. Effect of mini-tyrosyl-tRNA synthetase on ischemic angiogenesis, leukocyte recruitment, and vascular permeability. Am J Physiol Regul Integr Comp Physiol 295: R1138–R1146, 2008. First published August 27, 2008; doi:10.1152/ajpregu.90519.2008.—Mini-TyrRS has recently been shown to possess cytokine-like actions, leading to its inclusion in a growing family of aminocytic tRNA synthetase (AARS) multifunction cytokine-like proteins and peptides (22, 32–35). Mini-TyrRS stimulates neutrophil activation and chemotaxis in vitro and is angiogenic in endothelial cell cultures and in chick chorioallantoic membrane (CAM) and mouse matrigel implants (22, 32–35). Like CXC chemokines, such as IL-8, mini-TyrRS has an ELR motif (Glu-Leu-Arg) that confers its chemokine and angiogenic activities. Mutation of this motif inhibits binding and abolishes stimulation of leukocytes and induction of angiogenesis. Monocytes/macrophages, T-lymphocytes, and endothelial progenitor cells are important in angiogenesis, where they are recruited into and around new capillary sprouts and secrete growth factors and cytokines that promote endothelial cell proliferation and migration (9, 13).

Despite these intriguing in vitro actions, no studies have examined the mini-TyrRS in physiological or pathological settings in vivo. Therefore, the purpose of this study was to determine whether exogenous mini-TyrRS augments angiogenesis and leukocyte recruitment in ischemic tissue in vivo. We also investigated it for other key actions exhibited by angiogenic factors, namely, the induction of vasodilation and increased vascular permeability. Our findings show that mini-TyrRS augments angiogenesis and leukocyte adhesion in a mouse ear model of ischemic angiogenesis, and it increases permeability but lacks vasoactive actions. Intriguingly, at low concentrations mini-TyrRS had opposite effects, i.e., angiostatic activity, as well as reduced baseline and evoked increases in permeability. These novel bifasic actions of mini-TyrRS have potential therapeutic and physiological implications.

METHODS

Reagents. Rabbit anti-human mini-TyrRS antibody and human recombinant mini-TyrRS were from aTyr Pharma (La Jolla, CA), mFlt-trap (soluble VEGF-A receptor decoy) was kindly provided by Napoleon Ferrara and Stuart Bunting (Genentech, South San Francisco, CA). Bovine coronary venular endothelial cells were a gift from Cynthia Meininger, Texas A&M University. Four- to five-month-old mice were used in ear artery ligation (C57BL/6) and permeability models (sv129); a total of 285 animals were used to obtain the in vivo data shown in the figures. Procedures were performed aseptically and approved by the Institutional Animal Care and Use Committee of the University of North Carolina at Chapel Hill.

Unilateral ear artery ligation. One- to two-millimeter incisions at the base of the pinna were made overlaying the central and peripheral
Values are expressed as means per perfusion was determined for the ear in an anatomically defined region of interest (ROI; see MATERIALS AND METHODS) before and at indicated times after ligation.

Laser-Doppler perfusion imaging. Animals were anesthetized with 1.125% isoflurane supplemented with 2:3 oxygen:air. Perfusion was obtained before ligation, immediately after, and on 1, 3, 5, and 7 days using a scanning laser-Doppler perfusion imager (LDI2-IR; Moor Instruments, Devon, UK) modified for high resolution (10). Regions of interest (ROIs) were drawn by an investigator blind to drug treatment using Moor software to delineate a region between the ear margin and a circle extending from a line drawn to connect the two pinna notches. Unless otherwise indicated, perfusion was measured at 38°C rectal temperature to reduce vasomotor tone.

Vascular albumin leakage. Albumin leakage (permeability) was measured with Evans blue dye (T1824) (18) in the ear or dorsolateral back skin. Mini-TyrRS, mutant mini-TyrRS or PBS was injected (10 µl, 32-gauge needle here and elsewhere) subcutaneously into the base of ear (“*” in first panel) 12 h apart. #P < 0.001 vs. sham ligation; *P < 0.01, **P < 0.01 vs. PBS. Recovery of ear perfusion was inhibited by low- and augmented by high-dose mini-TyrRS.

VEGF-induced permeability was examined in the shaved back skin. Twenty microliters of PBS containing mini-TyrRS or PBS alone was injected subcutaneously. Thirty minutes later, T1824 was injected intravenously as above, followed by VEGF-A165 (100 ng in 20 µl PBS; R&D Systems, Minneapolis, MN) or PBS injected subcutaneously at the same location. Thirty minutes later, a skin circle, identified with a stereomicroscope circumscribing the extent of blue dye, was excised and T1824 content was determined as above.

Bovine coronary venule endothelial cell culture and monolayer permeability. Bovine coronary venule endothelial cells (BCVECs) (passages 10–15) were seeded onto culture dishes or onto 0.4-µm transwell inserts (Corning, Corning, NY) (3 × 10⁵ cells/insert), both precoated with 1.5% gelatin, and maintained in Dulbecco’s modified Eagle’s medium (DMEM) with 20% FBS at 10% CO₂ until a tight confluent monolayer was achieved. Cells were then pretreated with mini-TyrRS for 10 min, followed by 100 ng/ml VEGF for 30 min. Monolayers were then treated with Evans blue–BSA complex (0.67 g/l and 40 g/l) in HEPES-buffered saline for 30 min. Evans blue-albumin...
in the lower well was measured at 610-nm absorbance. Transendothelial albumin flux is expressed as percent clearance of albumin, compared with untreated controls.

**Thoracic aorta explants.** Rat thoracic aortas were isolated and maintained in serum-free medium composed of DMEM/F12, 10 mg/l insulin, 5 µg/l selenium and 5.5 mg/l transferrin in 21% or 1% O<sub>2</sub>. After exposure to VEGF for 4 days, samples were frozen in liquid nitrogen for immunoblot assay.

**Immunohistochemistry.** For capillary density, ears were perfusion-fixed with 4% PFA in PBS (pH 7.4) at 120 mmHg and then postfixed in 4% PFA for 24 h and embedded in paraffin. Eight-micrometer-thick sections located 5500 µm from the distal tip of the pinna were quantified for capillary density after staining for CD31 (sc-1506, 1:50; Santa Cruz Biotechnology, Santa Cruz, CA), followed by Cy3-conjugated secondary antibody (1:600). Vessels were imaged in eight different fields (×200 magnification) that covered the entire ear (cartilage, skin surface, and hair follicles with autofluorescence were excluded in digital images). Capillary density was derived from mean intensity of CD31 immunofluorescence (Image-J Software, National Institutes of Health, Bethesda, MD). T-cells and leukocytes were stained in adjacent sections with rat anti-mouse CD4 antibody (1:50, sc-13573; Santa Cruz Biotechnology) and CD45 antibody (1:200, 30-F11, BD Pharmingen, Franklin Lakes, NJ), respectively, followed with Cy3-conjugated secondary antibody (1:400–600). CD4- and CD45-positive cells were counted for the entire ear cross-section at ×400 magnification.

**Immunoblot.** Tissues were powdered in a liquid nitrogen-cooled pulverizer. Tissues and cultured cells were lysed in 1.5% Triton-X100 lysis buffer containing protease inhibitors (30 µg/ml aprotinin, 1 mM phenylmethylsulfonyl fluoride, 1 µg/ml leupeptin, and 1 µg/ml pepstatin) and phosphatase inhibitors (1 mM sodium-orthovanadate, 2.5 mM sodium-pyrophosphate). Samples were electrophoresed through 10% SDS-polyacrylamide and transferred to nitrocellulose membranes. Membranes were probed with antibodies against mini-TyrRS (1:1,000 dilution) and tubulin (1:5,000, ab6160, Abcam) followed by Alexa Fluor-680 (Molecular Probes, Carlsbad, CA) or IRDye 800 (Rockland Immunochemicals, Gilbertsville, PA) conjugated secondary antibodies at 1:5,000. Membranes were scanned and analyzed with the Odyssey system (LI-COR Biosciences, Lincoln, NE).

**Statistics.** Data are given as means ± SE. Differences were subjected to unpaired t-tests (2-tailed) or ANOVA followed by Bonferroni tests for multiple comparisons (2-tailed). P < 0.05 was considered significant.

**RESULTS**

**Mini-TyrRS has biphasic effects on ischemic angiogenesis.** Previous studies have reported that mini-TyrRS induced angiogenesis in cultured endothelial cell, CAM, and mouse matrigel assays (22, 32–35). To determine whether mini-TyrRS has angiogenic activity in vivo in ischemia, we examined its effects in the mouse ear made ischemic by ligation of the peripheral and central ear arteries, leaving the proximal-lateral branch of the central artery intact (Fig. 1). This is the first description of a mouse ear model for quantitative measure of perfusion, angiogenesis, and permeability. We developed it because of its ready access for these measures, plus its amenability to local administration of agents. In the PBS control group, perfusion declined 50% immediately after ligation, followed by recovery within 3–5 days mediated by angiogenesis and growth of collaterals between the above arterial trees (Fig. 1A). Local subcutaneous injection (20 µl, twice daily every 12 h) of 3 µg·kg<sup>-1</sup>·day<sup>-1</sup> mini-TyrRS into the base of the ear inhibited, while 600 µg·kg<sup>-1</sup>·day<sup>-1</sup> augmented, recovery of perfusion. Doses of 0.05 and 30 µg·kg<sup>-1</sup>·day<sup>-1</sup> had no effect (n = 8 mice/dose, data not shown), underscoring the biphasic activity. Doses higher than 600 µg·kg<sup>-1</sup>·day<sup>-1</sup> were not tested for reasons of concentration and cost. Mutant mini-TyrRS (ELR mutated to EYR) had no effect.

We next examined the effective doses in the Fig. 1A experiment for an effect on capillary density measured 7 days after ligation. The biphasic effect of mini-TyrRS on recovery of ear perfusion was accompanied by similar changes in capillary density (Fig. 1B). Capillary density, which increased in ischemia as expected, was inhibited by 3 µg·kg<sup>-1</sup>·day<sup>-1</sup> and augmented by 600 µg·kg<sup>-1</sup>·day<sup>-1</sup> mini-TyrRS, whereas mutant mini-TyrRS had no effect. This biphasic activity is different from in vitro, where only dose-dependent stimulation of angiogenesis occurs. Body weight did not differ between PBS or drug groups at any time point. In addition, daily subcutaneous injection of mini-TyrRS in the ear was not accompanied by erythema or edema.

**Mini-TyrRS increases leukocyte accumulation in ischemia.** Monocytes/macrophages and T-lymphocytes are involved in ischemic angiogenesis in vivo (9, 17). Furthermore, mini-TyrRS stimulates monocyte adhesion and transmigration in vitro (13, 19, 22). We, therefore, examined whether mini-TyrRS affects leukocyte density in cross sections of the ear adjacent to those used for determining capillary density in Fig. 1B. As expected, CD4<sup>+</sup> and CD4<sup>+</sup> cells increased after ligation (Fig. 2). At low-dose mini-TyrRS, which inhibits angiogenesis, no effect on leukocyte density was observed. However, high-dose mini-TyrRS caused a further increase of both cell types, whereas mutant mini-TyrRS was without effect.

**Mini-TyrRS has biphasic effects on baseline and evoked increase in permeability.** Increased permeability of the endothelium is an important step in the initial phase of sprouting angiogenesis, and several angiogenic factors, e.g., VEGF, regulate angiogenesis, in part, through alterations in permeability (2, 7, 9, 20, 25). We thus examined whether mini-TyrRS modulates leakage (“permeability”), using extravasation of Evans blue-conjugated albumin in normal, nonligated ears. Similar to its biphasic effects on angiogenesis (Fig. 1), mini-TyrRS also had biphasic effects on permeability (Figs. 3 and 4). Low-dose mini-TyrRS (30 µg/kg) reduced baseline leakage by 50%, while high-dose (600 µg/kg) increased leakage greater than twofold (Fig. 3A); mutant mini-TyrRS had no effect. We also tested mini-TyrRS on mustard oil-induced increase in permeability. Mini-TyrRS caused dose-dependent inhibition of induced leakage at low doses, with maximal inhibition at 600 µg·kg<sup>-1</sup>·day<sup>-1</sup> (Fig. 3B); mutant mini-TyrRS had no effect. This biphasic activity is different from in vitro, where only dose-dependent stimulation of angiogenesis occurs. Body weight did not differ between PBS or drug groups at any time point. In addition, daily subcutaneous injection of mini-TyrRS in the ear was not accompanied by erythema or edema.

**Mini-TyrRS has biphasic actions on angiogenic factors.** Because some angiogenic factors, such as VEGF and basic fibroblast growth factor (bFGF), exhibit vasodilator and vasoconstrictor activity, respectively (9), and because such activity of mini-TyrRS could alter perfusion in the ear ligation and permeability in the Evans blue experiments, we evaluated the effect of mini-TyrRS on...
perfusion of the normal (nonligated) mouse ear. The following
protocols (conducted in separate animals) reflect two require-
ments in these in vivo experiments: First, 10 min were required
to obtain the laser scanning-Doppler perfusion measurement.
Second, 20–30 min were allowed after a local injection (20
mL for all agents tested) to permit any elevation of interstitial
pressure to subside.

In the first experiment, rectal temperature was lowered to
35°C to increase vascular tone. Baseline perfusion was then
obtained, followed by injection of mini-TyrRS or PBS into the
base of the ear. Rectal temperature was then raised to 37.5°C
over 10 min to cause warming-induced vasodilation, and per-
fusion was obtained again. Mini-TyrRS had no effect on
warming-induced dilation (Fig. 5A). In a second experiment,
baseline perfusion was obtained at 35°C, and mini-TyrRS or
PBS was then injected. Thirty minutes later, the vasodilator
papaverine was injected in the same location, and perfusion
was obtained again 30 min later. Although there was a slight
suggestion that mini-TyrRS might modestly reduce the dilation
(Fig. 5B), the peptide had no significant statistical effect on
papaverine-mediated dilation. If mini-TyrRS has vasoconstric-
tor or vasodilator actions, dilator response to the superimposed
warming period in the first experiment or to papaverine in the
second experiment should have been altered. In a third exper-
iment to investigate possible progressive vasoactive actions
with repeated exposure, mini-TyrRS was administered locally
on six consecutive days. Ear perfusion was measured at 36.5°C
24 h after each administration and just before repeat dosing.
Mini-TyrRS had no effect on perfusion on day 6 (Fig. 5C) nor
at any of the five earlier days (data not shown). Absence of
vasoactive effects in these experiments is consistent with the
absence of erythema noted at any times or dosages, either
immediately after or 24 h after mini-TyrRS administration.
Edema was not evident at the 600 µg/kg dosages in any of the
experiments, even though this dose increased baseline perme-
ability (Fig. 3A). This may reflect low interstitial compliance
and/or efficient lymphatic drainage in the ear.

Ischemia, hypoxia, and VEGF reduce mini-TyrRS in calf and
isolated aorta. Angiogenesis in response to tissue hypoxia and
ischemia is achieved through upregulation of angiogenic fac-
tors such as VEGF, which, in turn, or through other mechan-
isms, downregulate angiostatic factors (9, 30). We thus rea-
soned that if endogenous mini-TyrRS normally exerts angio-
static actions at low concentrations, as suggested by the above
findings showing that low-dose mini-TyrRS inhibited recovery
of flow, angiogenesis, and permeability, then tissue levels of
mini-TyrRS might be regulated negatively in ischemia and in
response to VEGF. We used mouse hindlimb ischemia to
examine this question because it is a widely used model of neovascularization and because it is difficult to extract sufficient protein of high quality from the cartilaginous and fibrous ear. In addition, this model allowed measurement of mini-TyrRS by immunoblot in muscle that experiences ischemia (gastrocnemius) vs. little or no ischemia (adductor) after femoral artery ligation (10, 18). Tissue samples were harvested on days 2, 5, and 10 after ligation of the femoral artery, as detailed elsewhere (7). Mini-TyrRS decreased in the gastrocnemius but not adductor of the ligated leg, compared with the gastrocnemius from sham animals (no surgery) or from the contralateral nonligated leg (Fig. 6, A and B). The specificity of the mini-TyrRS band identified with rabbit anti-human antibody was verified using a second goat anti-rabbit antibody that we raised against a different epitope of mini-TyrRS. To test possible involvement of hypoxia and VEGF in reduced mini-TyrRS, we examined rat thoracic aorta maintained in organ culture. Four days of exposure to VEGF or hypoxia (1% O₂) caused similar reduction of mini-TyrRS (Fig. 6, C and D). Moreover, hypoxic reduction was abolished by VEGF neutralizing trap. These data suggest that hypoxic induction of VEGF may mediate reduced mini-TyrRS in ischemic tissue.

**DISCUSSION**

This is the first characterization of the effects of exogenous mini-TyrRS on ischemic angiogenesis, leukocyte trafficking, permeability, and vasoactivity in vivo. We confirm several properties previously observed in vitro, identify novel biphasic actions on angiogenesis and permeability, and find evidence suggesting that tissue levels of mini-TyrRS may be regulated by hypoxia.
In the mouse ear model of ischemia, low-dose mini-TyrRS (3 μg·kg⁻¹·day⁻¹) inhibited, while high-dose mini-TyrRS (600 μg·kg⁻¹·day⁻¹) augmented angiogenesis. Intermediate doses of 0.05 and 30 μg·kg⁻¹·day⁻¹ had no effect. Mutant mini-TyrRS had no effect, confirming the requirement of an intact ELR. The angiostatic-like action of low-dose mini-TyrRS was not observed in previous in vitro, matrigel and CAM implant studies, in which only angiogenic actions were observed. For example, mini-TyrRS was angiogenic at 2.4–24 μg/ml (60–600 nmol/l) in matrigel implants and induced migration of cultured endothelial cells at 2 μg/ml (50 nmol/l) (33). These differences could arise for several reasons, including the context of ischemia in our study, inherent differences in conditions in vivo, in vitro, and “in matrigel”, and because the concentrations used previously are undoubtedly higher than achieved in our low-dose groups, where dilution and degradation over time would be considerable. For example, assuming mini-TyrRS injected locally into the ear distributes into the extracellular space and that degradation reduces the concentration by 10-fold (actual degradation is likely to be at least 1 order of magnitude more than this), our low- and high-dose regimens would achieve average extracellular concentrations of 0.006 μg/ml and 1.2 μg/ml. Thus, although concentrations in the ear’s extracellular fluid would clearly be higher for some duration after injection, it is likely that time-averaged levels achieved in our low-dose groups were significantly lower than those in previous studies.

Fig. 5. Mini-TyrRS lacks vasoactive actions. A: increase in perfusion (Doppler) induced in nonligated ear by raising rectal temperature from 35°C to 37.5°C was unaffected by mini-TyrRS (20-μl subcutaneous administration into ear immediately after 35°C measurement, followed by measurement 10 min later at 37.5°C). B: increase in perfusion induced by papavarine (adductor area sc) was unaffected by mini-TyrRS injected into same site 30 min earlier. C: baseline (control) ear perfusion was unaffected after 6 days of daily sc mini-TyrRS (daily dose given as two 20-μl injections 12 h apart); n = 4–6/bar.

Fig. 6. Ischemia/hypoxia and VEGF reduce mini-TyrRS levels in vivo and in vitro. A and B: Western blot analysis of mini-TyrRS in gastrocnemius and adductor of sham-surgery mice, in gastrocnemius and adductor of leg with femoral artery ligation (lig), and in gastrocnemius of the contralateral nonligated leg at the indicated days (d) after surgery. Twenty micrograms (A and B) and 30 μg (C and D) protein per lane; normalized to tubulin; n = 4/bar. C and D: Western blot analysis of mini-TyrRS in rat thoracic aorta maintained 4 days in organ culture with 100 ng/ml VEGF or 1% oxygen (hypoxia) ± VEGF trap or IgG control (0.2 mg/ml). *P < 0.05, **P < 0.01 vs. sham or normoxia; n = 3 or 4/bar.
Recruitment of leukocytes and endothelial progenitor cells (EPCs) contributes importantly to angiogenesis in ischemia, inflammation and tumor growth (2, 9, 13, 20). These cells exhibit heterogeneous phenotypes, expressing markers for macrophages, T-cells, smooth muscle cells, fibroblasts, pericytes, and EPCs, and secrete growth factors and cytokines, which act directly or indirectly to augment endothelial cell migration, proliferation, and capillary sprouting. CD4-positive T-lymphocytes play an important role in angiogenesis by secreting angiogenic growth factors, such as VEGF (14) and bFGF (4). In the present study, high-dose mini-TyrRS further increased leukocyte (CD45-positive) accumulation in the ischemic ear by ~2.5-fold compared with PBS control and CD4-positive cells by ~10-fold, while low-dose mini-TyrRS had no effect. This action may contribute to the angiogenic effect of mini-TyrRS.

Besides the novel biphasic effects of mini-TyrRS on angiogenesis, mini-TyrRS also had biphasic effects on both basal and evoked permeability. In most vascular beds, including the skin, permeability to plasma proteins and smaller molecules is normally low. Ischemia, inflammation, and tumor growth are accompanied by increased vascular permeability, which is an important early step in angiogenesis in these conditions (2, 9, 27) The resulting leakage of plasma proteins and other circulating macromolecules helps to convert the normally anti-angiogenic stroma into a proangiogenic provisional stroma (9, 11). Many angiogenic factors, such as VEGF (16), bFGF (7), IL-8 (25), and thrombin (16) increase endothelial permeability (2). On the other hand, antagonism of increased permeability reduces angiogenesis (5, 29, 30). For example, the angiostatic proteins angiostatin (28), caveolin-1 (5) and TNP-470 (26) reduce evoked increases in permeability. However, to our knowledge, no endogenous angiostatic factor has been reported to reduce basal permeability like that observed in the present study for mini-TyrRS. On the other hand, high-dose mini-TyrRS increased basal permeability. We also found that mini-TyrRS at low doses caused dose-dependent inhibition of evoked leakage by mustard oil and VEGF, while high-dose mini-TyrRS tended to augment evoked leakage. This positive regulatory effect of mini-TyrRS on basal and evoked permeability may contribute to the angiogenic action of mini-TyrRS. Although the mechanisms underlying the biphasic effect await future studies, estrogen has similar biphasic effects on permeability in ECs in vitro (36).

The specificity of our findings regarding angiogenesis, leukocyte accumulation, and increased permeability are supported by their dependence on an intact ELR (Glu-Leu-Arg) motif, i.e., mini-TyrRS with this motif mutated had no effects. This motif is required for binding, neutrophil activation/adhesion, and angiogenesis induced by mini-TyrRS in vitro and other CXC ELR-containing chemokines, such as IL-8 (15, 16, 22, 32, 33, 35). However, the ELR motif need not be angiogenic. For example, the ELR motif-containing chemokine GRO-β has antiangiogenic properties (8), and members of the CXC subfamily, induced by interferons, lack the ELR motif but are potent inhibitors of angiogenesis (19). In addition—and of potential relevance to our findings—22–26-amino acid peptides derived from proangiogenic ELR-containing CXC chemokines exhibit potent antiproliferative and antimigratory activity in vitro (19). The mini-TyrRS receptor has not yet been identified, nor is it known whether mini-TyrRS binds to different receptors at low vs. high concentrations. Multiple receptors and/or processing of the protein after export from endothelial and other vascular cells (15, 22, 35) could underlie the biphasic properties we observed. For example, the AARS cytokines mini- and T2-TrpRS, thought to be derived from proteolysis and/or alternative splicing of TrpRS, inhibit VEGF signaling and are angiostatic (21, 22, 34, 35). It has recently been reported in endothelial cells that mini-TyrRS release is induced by TNF-α, undergoes binding, and induces phosphorylation of Src, Akt, ERK, and VEGF-receptor 2, which is required for tube formation (15). These factors are also involved in signaling angiogenesis, angiostatic activity, leukocyte adhesion, and changes in vascular permeability (2, 9, 20, 23, 30) and thus could be involved in the biphasic action of mini-TyrRS on these processes identified herein. Knowledge of the signaling pathways activated at low vs. high concentrations of mini-TyrRS in vivo or in intact vessel organ culture preparations will be required to understand the basis for the biphasic activities identified in this study.

Tissue hypoxia in ischemia and tumor growth induces many of the steps involved in angiogenesis, e.g., increased permeability, inflammation, endothelial cell proliferation and migration, and matrix degradation (2, 3, 6, 9, 12, 20, 23, 24). A number of the proteins that mediate these processes, such as endothelial nitric oxide synthase, VEGF, angiopoietin-2, Akt, and bFGF are regulated by hypoxia. Mini-TyrRS levels detected by immunoblot of tissue lysates were significantly reduced in gastrocnemius muscle when examined at 5 and 10 days after ligation of the femoral artery. Similar reductions were also observed in thoracic aorta explants exposed to either hypoxia or VEGF. Moreover, in the latter model, a VEGF-trap-blocking protein abolished reduction in mini-TyrRS during hypoxia. These findings in explants suggest that the reduction of mini-TyrRS in calf muscle after femoral ligation in vivo may be mediated by hypoxia and VEGF. This supports our hypothesis from the findings in Fig. 1 that the physiological effects of low concentrations of mini-TyrRS are angiostatic. It is possible that the decline in mini-TyrRS detected in tissue extracts of calf and aorta explants reflects a regulated reduction in cleavage of the protein from full-length TyrRS to cause withdrawal of its angiostatic effect produced at low concentrations (Fig. 1). On the other hand, it has recently been shown that mini-TyrRS is released by stimulation of endothelial cells with TNF-α in vitro (15). Thus, it is also possible that export from endothelial cells (and possibly other cell types) in response to ischemia, hypoxia, and VEGF [conditions tested herein; however, VEGF did not induce release of mini-TyrRS in vitro (15)] could have reduced intracellular levels. This plus proteolysis of any released mini-TyrRS may have resulted in lower levels detected in tissue extracts. Additional studies and development of assays to measure released mini-TyrRS will be required to determine whether the decline in mini-TyrRS with the models and stimuli examined herein reflects increased export and achievement of angiogenic concentrations or is due to a regulated reduction in levels to withdraw the angiostatic activity of low mini-TyrRS levels. If evidence for the latter is obtained, the angiogenic and related actions of high-dose mini-TyrRS that we observed may reflect a potential therapeutic use of the protein in ischemia, while the low-dose angiostatic and barrier-increasing actions may reflect physiological activities as well as a potential therapeutic use to inhibit
pathological angiogenesis.” Clearly, more work needs to be done to understand expression/proteolysis of TyrRS under different physiological conditions.

In summary, we report that low-dose mini-TyrRS inhibits basal and evoked permeability and ischemic angiogenesis. High-dose has opposite effects and, in addition, augments recruitment of CD45-positive and CD4-positive cells in ischemic tissue. To our knowledge, mini-TyrRS is the first factor observed to inhibit angiogenesis at low and stimulate it at high concentrations. We also provide in vitro evidence in aorta that mini-TyrRS levels are reduced by VEGF-dependent signaling in hypoxia, suggesting a similar mechanism may underlie its reduction in ischemic tissue in vivo. These findings suggest that endogenous mini-TyrRS may serve an angiostatic function in certain physiological settings. Moreover, administration of it at low vs. high levels could provide a therapeutic approach to limit vs. augment angiogenesis in certain diseases. Important areas of future study include determining whether stimulation of leukocyte accumulation by mini-TyrRS reflects induction of adhesion molecules on endothelial cells, leukocytes, or both, or release of inflammatory cytokines from these or other cell types, and similar questions regarding the biphasic effects of mini-TyrRS on baseline and induced permeability and ischemic angiogenesis. Besides a number of tools that need to be developed, strategies to antagonize endogenous production/clavage/secretion and block the responsible receptor(s) will be needed to test the potential physiological roles suggested by the findings in this study.

ACKNOWLEDGMENTS

The authors thank Young Greenberg for dialyzing mini-TyrRS, K Kirk McNaughton for histological assistance, and John Reader for helpful discussion.

REFERENCES


