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*Running title: *Matrilin-1 Inhibits Neovascularization*

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Background: The relative avascularity of cartilage has made it a promising source of angiogenesis inhibitors.
Results: MATN-1, identified by mass spectrometry, suppresses capillary endothelial cell proliferation and migration.
Conclusion: MATN-1 is a novel inhibitor of neovascularization *in vivo* and *in vitro*.
Significance: This is the first demonstration that MATN-1 is an inhibitor of both normal and pathological neovascularization.

ABSTRACT

In the course of conducting a series of studies whose goal was to discover novel endogenous angiogenesis inhibitors, we have purified Matrilin-1 (MATN-1) and have demonstrated, for the first time, that it inhibits neovascularization both *in vitro* and *in vivo*. Proteins were extracted from cartilage using a 2M NaCl, 0.01M HEPES buffer at 4°C, followed by concentration of the extract. The concentrate was fractionated by size exclusion chromatography and fractions were then screened for their ability to inhibit capillary EC proliferation *in vitro*. Fractions containing endothelial cell (EC) inhibitory activity were pooled and further purified via cation exchange chromatography. The resulting fractions from this step were then screened to isolate the anti-angiogenic activity *in vitro*. This activity was identified via tandem mass spectrometry (MS/MS) as being MATN-1. Human MATN-1 was cloned and expressed in *Pichia pastoris* and purified to homogeneity. Purified recombinant MATN-1, along with purified native protein, was shown to inhibit angiogenesis *in vivo* using the chick chorioallantoic membrane assay via the inhibition of capillary EC proliferation and migration. Finally, using a MATN-1-deficient mouse, we showed that angiogenesis during fracture healing was significantly higher in MATN-1-/- mice in comparison to the wild type mice as demonstrated by *in vivo* imaging and by elevated expression of angiogenesis markers including PECAM1, VEGFR, and VE-cadherin.

INTRODUCTION

Cartilage is an avascular and relatively tumor-resistant tissue that is composed predominantly of proteoglycans, several types of collagen, and non-collagenous matrix proteins (1-3). The resistance of cartilage to capillary invasion has long been postulated to be a function of the presence of endogenous inhibitors of angiogenesis, which inhibit new blood vessel formation from pre-existing vessels. The first of these inhibitors purified to homogeneity was a cartilage-derived tissue inhibitor of metalloproteinases (TIMP), a protein that was demonstrated to be a potent

inhibitor of capillary EC proliferation and migration *in vitro,* and angiogenesis *in vivo* in both the chick chorioallantoic membrane (CAM) assay and the corneal pocket assay (4,5). Shortly thereafter, a chondrocyte-derived angiogenesis inhibitor, a 35.5 kDa protein isolated from chondrocyte primary cultures, was shown to inhibit angiogenesis *in vitro* in EC proliferation and migration assays and *in vivo* in the CAM assay (6). Chondromodulin-I (ChM-I), a 25 kDa protein isolated from bovine epiphyseal cartilage was shown to inhibit capillary tube formation in *in vitro (7-9)* and tenomodulin (TeM), another protein that shares homology with ChM-I, was shown to inhibit angiogenesis *in vitro* as well (9-12) . Troponin I (TnI), a 23 kDa contractile protein typically found in muscle, was isolated from bovine cartilage, and demonstrated to inhibit EC proliferation *in vitro*, angiogenesis in the CAM and mouse corneal pocket assays *in vivo*, as well as tumor metastasis *in vivo* (13).

Non-mammalian cartilage has also been studied as a potential source of angiogenesis inhibitors. Lee and Langer were the first to demonstrate that an extract of shark cartilage could significantly inhibit tumor neovascularization (14). Since then, several laboratories have isolated partially purified fractions from this type of cartilage that have been shown to inhibit at least one of the processes associated with angiogenesis. For example, U995, a fraction containing 10 and 14 kDa peptides isolated from the blue shark, *Prionace glauca*, interfered with HUVEC proliferation and migration (15). SCF2, a glycosaminoglycan whose principal component is keratan sulfate, is a 10 kD proteoglycan that has been reported to inhibit EC proliferation *in vitro,* as well as tumor-induced angiogenesis in the cornea of rabbits and angiogenesis in CAM assays *in vivo* (1). SCAIF80, an 80 kDa protein isolated from shark cartilage, has been shown to significantly suppress EC proliferation and migration in a dose dependent matter *in vitro* (16), and SCP1, a 13.7 kDa protein with sequence similarities to parvalbumin, was reported to inhibit angiogenesis in the rat aortic ring assay (17). AE-941, an 'angiogenic mixture' (18) isolated from *Squalus acanthias* cartilage, has been shown to inhibit angiogenesis *in vitro* in the rat aortic ring vessel

assay (19) and *in vivo* in the CAM assay (20). In addition, the shark tissue inhibitor of metalloproteinase 3 (sTIMP-3), has been cloned and characterized from the cloudy dogfish, *Scyliorhinus torazame* (21). It was later shown to inhibit migration and tube formation in bovine aortic endothelial cells (22).

In this study, we have purified and identified a novel inhibitor of angiogenesis, matrilin-1 (MATN-1). Having demonstrated its ability to inhibit angiogenesis *in vivo*, we determined that MATN-1 exerted this suppression of neovascularization by inhibiting angiogenesis growth factor-driven capillary endothelial cell proliferation and migration. These studies weresupported by both *in vitro* gain of function studies and *in vivo* loss of function experiments using MATN-1 KO mice. MATN-1, formerly known as cartilage matrix protein (CMP), is an abundant component of cartilage (23). A modular protein, MATN-1 mediates interactions between a variety of matrix components (24) and, as a structural protein, it binds to biglycan and decorin (25), collagen (26), cartilage oligomeric matrix protein (27) as well as to itself (28). To our knowledge, this report is the first to document the ability of this structural protein to suppress new capillary growth *in vivo* and suggests that its targeting may be of potential clinical significance.

EXPERIMENTAL PROCEDURES
*Extract preparation--*The chondrocranium cartilage of the spiny dogfish (*Squalus acanthias*) was harvested and scraped free of muscle and connective tissue as described previously (4,14). The prepared cartilage (250 g) was homogenized and extracted in 4L of a 2M NaCl, 0.01M HEPES, 3mM EDTA, 0.02% NaN_3 extraction buffer for 4 days under constant agitation, utilizing a modification of a previous procedure (4). The extraction solution was filtered with gauze, centrifuged at 6,500 x g for 2 hours to remove particulates, then concentrated using a Vivacell 250 (Sartorius Stedim Biotech) to a final volume of approximately 5 ml. The concentrated cartilage extract (CCE) had a final protein concentration of 7.8 mg/ml. All procedures were performed at 4ºC.

*Purification and identification of MATN-1--*A 40 mg sample of CCE, dialyzed against Biogel A-1.5M buffer (4 M guanidine HCl, 20 mM Tris, pH

7.6) overnight, was applied to a Biogel A-1.5M Sepharose size exclusion column (5x50) at a flow rate of 1.0 ml/min; fractions were collected every 5 minutes. Fractions were screened for their ability to inhibit both EC proliferation and matrix metalloproteinase (MMP) activity (Fig. 1A). Given that cartilage had already been reported to contain an anti-angiogenic activity that was identified as being an MMP inhibitor (4), fractions were screened for both of these activities in order to identify a novel inhibitor of neovascularization. Fractions 22-34 from the column were enriched in the ability to inhibit EC proliferation and were pooled, dialyzed overnight to remove the guanidine HCl, and then applied to a Bio-Rex 70 column (Bio-Rad) and fractionated, as previously described by us (4). Fractions 2 and 5 (Fig. 1B), which contained the anti-proliferative activity, were pooled and dialyzed to remove excess salts, then subjected to electrophoresis on 12% SDS-PAGE gels (BioRad) or 12% Bis-Tris NU-PAGE gels (Invitrogen) run under denaturing conditions followed by visualization by either silver or SYPRO Ruby (Invitrogen) staining (29). Protein bands were excised from the gel, subjected to tryptic digestion, and analyzed by MALDI-TOF mass spectrometry (Perceptive STR, Applied Biosystems, Framingham, MA) to determine the molecular weights of the proteins and for peptide mapping of the tryptic digests using a 337-nm wavelength laser for desorption and the reflectron mode of analysis. Using the MSFit (http://www.prospector.ucsf.edu) and SEQUEST (30) search programs, the peptide maps generated were searched against a FASTA data base of public domain proteins constructed of protein entries in the non-redundant data base held by the NCBI and Swiss-Prot. Peptide matches identified by MSFit were filtered according to their MOWSE (molecular weight search) score, percentage of masses matched, molecular weight, and number of observations of peptides and proteins

Cloning, expression and purification of MATN-1--The cMATN-1 gene was cloned using a pcDNA3.1 vector with cMATN-1 insert (31) and primers specific for the mature form of chick MATN-1. The human MATN-1 (hMATN-1) gene was cloned from normal adult human lung and trachea first strand cDNA (Biochain Institute, Inc.; Hayward, CA). For the cloning of both the chick

and human MATN-1 into the pPICZαC expression plasmid, the cDNAs for MATN-1 were amplified by PCR using two primers that covered the translation start codon (primer 1) and the stop codon (primer 2) of CMP cDNA, respectively. The two primers were designed in such a way that primer 1 contained a XhoI site and primer 2 contained a poly His tag (to aid in purification) and a XbaI site (for hMATN-1, primer 1, 5' CCGCTCGAGATGAGGGTCCTCTCTGGC A3'; primer 2, 5' GCTCTAGATCAATGATGATGA TGATGATGTTAGACAACTGTGTTC 3'; for cMATN-1, primer 1, 5' CCGCTCGAGAAA AGACCTCCTCAGCCCAGAGG 3'; primer 2, 5' GCTCTAGATCAATGATGATGATGATGATG GATGATCTTATTCTC 3'). The full-length MATN-1 PCR products were subcloned into pCR4-TOPO vector (Invitrogen). After confirmation of the sequence, the MATN-1 genes were ligated into the yeast expression vector pPICZαC (Invitrogen). Linearized vectors (PmeI) were electroporated into the yeast *Pichia pastoris* for expression (Invitrogen), and integrants were selected by culturing on YPDS (2% peptone, 1% yeast extract, 2% glucose, 1M sorbitol, 2% agar) plates with 100 μg/ml zeocin (Invitrogen) for 3 days. Successful insertion of chick and human genes of interest into the *Pichia* genome were verified by PCR using *Pichia* specific primers, which also verified that recombination occurred at the correct site. Expression of the gene of interest is under the control of the methanol-inducible AOX1 promoter.

Five *Pichia* clones each of cMATN-1 and hMATN-1 were tested for their expression levels and the clones from each set expressing the highest amount of protein were chosen for subsequent studies. The expression conditions were as follows: 25-ml overnight cultures were grown at 30 °C in yeast extract peptone dextrose medium (YPD, 1% yeast extract, 2% peptone, 2% dextrose) containing 100 μg/ml zeocin. Cell pellets were collected after 24 h by centrifugation at 1500 x g. Cultures were induced to express the recombinant proteins by resuspending the cell pellets in 250 ml of buffered methanol-complex medium (BMMY, 2% peptone, 1% yeast extract, 100mM potassium phosphate, pH 6.0, 1.34% yeast nitrogenous base, 1% methanol), and allowed to grow for 24 h at 30 °C with constant shaking.

Medium containing the secreted expressed protein was cleared of cell content by centrifugation at 5000 x g.

Purification of recombinant MATN-1-- Expressed recombinant chicken and human MATN-1 proteins were initially purified from the yeast media using histidine affinity binding to a nickel-nitrilotriacetic acid-agarose (Ni-NTA) resin (Qiagen, Valencia, CA) under native conditions. Expressed protein in 800 ml of pooled, cleared medium was allowed to bind to 5 ml of resin in a pre-packed column by gravity flow at 4 °C. The resin was washed with 50ml (10 volumes) of buffer containing 10mM imidazole (50mM NaH_2PO4, 300mM NaCl, 10mM imidazole, pH 8.0) to reduce nonspecific binding. Protein was then eluted using 25ml (5 volumes) of elution buffer containing 100mM imidazole (50mM NaH_2PO4, 300mM NaCl, 100mM imidazole, pH 8.0) and concentrated by centrifugation using membrane concentrators with 10-kDa molecular mass cutoff (Centriprep, Amicon, Beverly, MA). Purity was confirmed by silver staining of SDS PAGE gels and MS/MS.

*Cell culture--*Capillary EC, isolated from bovine adrenal cortex, were a kind gift of Dr. Judah Folkman and Catherine Butterfield (Boston Children's Hospital). Cells were maintained in Dulbecco's modified Eagle's medium (DMEM; Invitrogen) containing 10% calf serum (HyClone) and 3 ng/ml basic fibroblast growth factor (bFGF) in 10% CO_2 at 37 °C, as previously reported (4).

*Endothelial cell proliferation assay-*Capillary EC proliferation was measured as reported previously (4,6,13). Briefly, capillary EC were plated on pregelatinized 96-well plates at a density of 2000 cells per well in Dulbecco's modified Eagle's medium supplemented with 5% calf serum and allowed to attach for 24 h. The next day, cells were treated with fresh medium with or without 1 ng/ml bFGF and challenged with the test proteins at various concentrations. All samples were tested in duplicate. Control wells contained cell treated with medium alone or medium with bFGF. After 72h, the medium was removed, and the cells were lysed in buffer containing Triton X-100 and the phosphatase substrate p-nitrophenyl phosphate. After 2h incubation at 37 °C, NaOH was added to each well to terminate the reaction and cell density was determined by colorimetric analysis using a

SpectraMax 190 multiwell plate reader (Molecular Devices, Sunnyvale, CA). All samples were tested in duplicate in at least three independent experiments.

Capillary endothelial cell migration-- Capillary EC migration was studied using a modified Boyden chamber as previously described (4,32). The upper half of transwell (8µM pore; Costar) membranes were coated with fibronectin (10µg/ml; Becton Dickinson) overnight at 4 °C to facilitate cell adhesion. Coated membranes were rinsed with PBS and allowed to air dry immediately before use. Cells were detached by trypsinization, and resuspended at a final concentration of 0.5×10^6 cells/ml in serum-free DMEM containing 0.1% BSA. Cells were added to the upper chamber of the transwell and allowed to migrate toward the bottom chamber containing DMEM, or DMEM supplemented with the chemoattractant bFGF for 4h in a humidified incubator containing 5% CO_2. Transwell filters were rinsed once with PBS and fixed and stained using a Diff-Quik kit (Baxter) following the manufacturer's protocol. Stained filters were cut out of the chamber and mounted onto slides using Permount (Sigma). The number of migrated cells were counted (three fields from each membrane were captured using a 10× objective), and images were captured with a CCD camera using SPOT software. Total migration per membrane was quantified from the captured images using Scion Image software (National Institutes of Health). All experiments were run in triplicate.

*MMP inhibitory activity--*MMP inhibitory activity was assessed using a quantitative [C^{14}] collagen film assay, as described previously (4,32). An IC_{50} unit was defined as the amount of protein necessary to inhibit the proteolytic activity of collagenase by 50%.

Angiogenesis inhibitory activity in vivo-- The CAM assay was conducted as reported previously (4,32). Three day old chick embryos were removed from their shells and incubated in plastic Petri dishes for 3 days. On embryonic day 6, samples and controls were mixed into methylcellulose and allowed to dry. The discs were applied to the surfaces of developing CAMs, above the dense subectodermal plexus. After an incubation of 48 h, the eggs were examined for vascular reactions under a dissecting scope and

photographed. All samples were tested in triplicate for each treatment.

Mice tibia fracture model--Animals were studied at Coro Center Facilities of Rhode Island Hospital. NIH guidelines for the care and use of animals were observed and the study was approved by Rhode Island Hospital IACUC. Six to eight week-old male mice were used in the experiments. MATN-1 -/- (n=5) mice and WT mice (n=5) of the same genetic background (C57BL/6J) were genotyped using PCR. The animals were used to create fracture models as previously described (33)(1,33). In brief, animals were anesthetized by intraperitoneal injection of a Ketamine (Bioniche Pharma USA) and Medetomidine (Orion Corporation Espoo, Finland) cocktail (75 + 1 mg/kg). The animals were prepared for surgery by shaving and scrubbing of both hind limbs. A longitudinal short incision was made at the knee, and a 0.5mm hole was drilled above the tibial tuberosity. A 30G stainless-steel needle (Hamilton, Reno, Nevada, USA) was introduced into the intramedullary canal of the tibia. The wound was closed, and the procedure was repeated on the contralateral side. A closed transverse mid-diaphyseal tibia fracture was created by three-point bending in the right tibia, and the animal was allowed to move freely after recovery from anesthesia with the intraperitoneal injection of Antisedan (Orion Corporation Espoo, Finland) (1mg/kg). A preoperative cefadroxil (25mg/kg) was administered subcutaneously to prevent infection. Buprenex (Reckitt Benckiser Healthcare, UK; 0.03 mg/kg) was given once preoperatively and two times per day during the first 3 days after surgery to relieve pain.

Fluorescence molecular tomography measurements of angiogenesis--The commercially available fluorescent probes, AngioSense[TM] (FRFP750, excitation 750t10 nm; emission 780t10 nm) (VisEn Medical Inc., Woburn, MA) was used in this study. The AngioSense[TM]750 is a fluorescent in vivo blood pool imaging agent which enables imaging of blood vessels and angiogenesis. The adult mouse dose of 2 nmoles in 150mL saline (13.3mM) or approximately 80 nmoles/ kg body weight was utilized. The probes were delivered to the animals via IP injection 24 hours before taking the images with Fluorescence Molecular Tomography (FMT) system (VisEn Medical Inc., Woburn, MA) on day 4 post-surgery. Twenty-four hours after the probe injection, mice were anesthetized with a solution of ketamine and medetomidine, and imaged using the VisEn's FMT system, as described previously (34). Three-dimensional regions of interest were drawn around the tibial fracture limbs and control limbs and a threshold was applied equal to 10% of the maximum value of fluorescence in each reconstructed volume. The peak concentration (in nanomoles per liter) and total amount (in picomoles) of fluorochrome were automatically calculated relative to internal standards generated with known concentrations of appropriate dyes.

Callus mRNA expression--The RNeasy Fibrous Tissue Mini kit (Cat.No.74704, Qiagen, Valencia, CA) was used to extract RNA from the fracture calluses 14 days post-surgery with the following modification. After the mice were sacrificed, the fracture calluses were placed in RNA later (Qiagen, Valencia, CA) and stored at −80°C until RNA extraction. The calluses were ground into a fine powder using mortar and pestle with liquid nitrogen, and extracted according to the instructions provided with the kit. Quality and quantity of RNA were determined using a Nano-drop (Ambion, Austin, TX). mRNA was determined by real-time quantitative reverse transcriptase PCR (RT-PCR). Total RNA (1 μg) was transcribed into cDNA using iScripTM (Bio-Rad, Hercules, CA) with (40ng/µl) of the resulting cDNA used as the template to quantify the relative content of mRNA by RT-PCR using Sso Fast EvaGreen Supermix (Bio-Rad, Hercules, CA) with CFX 96 Real Time PCR system (Bio-Rad, Hercules, CA). To normalize the data, mRNA expression of a housekeeping gene, 18S was also determined. The cycle threshold (Ct) values for 18S RNA and that of samples were measured and calculated by Excel (Office 2007, Microsoft, Redmond, WA). Relative transcript levels were calculated as $x = 2^{-\Delta\Delta Ct}$, in which $\Delta\Delta Ct = \Delta E - \Delta C$, and $\Delta E = Ct_{exp} - Ct_{18s}$; $\Delta C = Ct_{ctl} - Ct_{18s}$.

RESULTS

Identification of Matrilin 1--The band migrating at approximately 52 kDa consistent with the molecular mass of MATN-1 was excised from the gel (Fig. 1C) and subjected to mass

spectrometry analysis as described above. The MOWSE score reported by MS-Fit was 547 with 6 matching peptides (40% coverage); the protein was identified as cartilage matrix protein precursor (CMP) or Matrilin-1 (*Gallus gallus*: EMBL-EBI Accession# 115555). SEQUEST also identified the protein as Matrilin-1 (*Gallus gallus*: Accession# P05099) with a reported Xcor value of 3.04 or higher for four peptides; again the protein was identified as a cartilage matrix protein precursor (Matrilin-1; P05099) (Fig. 1C,D,E). Immunoblot analysis using a monoclonal antibody to MATN-1 (1:3000) confirmed the identification of the 52 kDa protein band as being MATN-1. This antibody was a gift from Dr. Paul Goetinck (35) (Fig. 1F).

MATN-1 inhibits angiogenesis in the chick CAM assay--The chick CAM assay was used to determine whether MATN-1 could suppress angiogenesis *in vivo*. We observed significant inhibition of embryonic neovascularization in this assay as evidenced by the avascular zone elicited by representative CAM treatment with 10μg (350 nM) of recombinant chicken MATN-1 or 5μg recombinant human MATN-1 (Fig. 2).

MATN-1 inhibits endothelial cell proliferation and migration--Given its ability to inhibit angiogenesis *in vivo,* we next asked how MATN-1 might be exerting this antiangiogenic effect. We tested MATN-1 in two different assay systems, which measured the inhibition of angiogenic mitogen-driven capillary endothelial cell proliferation and migration, two processes essential for successful angiogenesis. Purified native chick MATN-1 (Fig. 3A), recombinant chick MATN-1 (Fig. 3B) and recombinant human MATN-1 (Fig. 3C) all suppressed EC proliferation. The IC_{50} for each of the three treatments was approximately 175 nM, 275 nM, and 75 nM, respectively. When purified native chick MATN-1, recombinant chick MATN-1, or recombinant human MATN-1 (Fig. 4) were tested for their effect on EC in a transwell migration assay, we found that migration was inhibited in a dose-dependent manner with an IC_{50} of approximately 75 nM, 80 nM, and 10 nM, respectively.

MATN-1-/- mice induce angiogenesis in the early period of fracture healing—We next complemented these *in vitro* gain of function

studies by conducting *in vivo* loss of function experiments using MATN-1 KO mice. The FMT images and tomographic reconstructions showing angiogenesis in the injured site (Fig. 5) provide a clear visualization and quantification of neovascularization in the fracture site and control limb. Figure 5 shows a 3D rendering of the vasculature in WT and MATN-1 -/- mice, respectively, injected with AngioSense™ 24h before FMT analysis taken on Day 4 post fracture. The fractured tibia of MATN-1 -/- mice displayed a significant AngioSense™ signal, which was 1.76-fold higher than that of the WT mice (*p*= 0.0041). There was no significant difference between MATN-1 -/- and WT mice in the control tibia (*p*=0.0970). MATN-1 -/- mice had a 1.87-fold higher AngioSense signal compared to WT mice (*p*=0.0156). These findings from the FMT analysis indicated that there was significantly more vascularization at the early stage of fracture healing in MATN-1 -/- mice.

To confirm these findings, we analyzed the expression of angiogenesis-related gene markers. The animals were sacrificed on Day 14 post-fracture and total RNA was isolated from the fracture calluses for RT-PCR analysis. The mRNA expression of angiogenesis-relevant factors, including PECAM1, VEGFR, and VE-cadherin were examined by RT-PCR (Fig. 6). The results demonstrated that, 14 days after fracture, MATN-1-/- mouse calluses expressed significantly higher PECAM1 (*p*=0.0041), VEGFR (*p*=0.0115), and VE-cadherin (*p*=0.0004) expression levels compared to those of WT mice. Taken together, these results demonstrate that MATN-1 inhibits angiogenesis in the early phase of fracture healing.

DISCUSSION

In the present study, we demonstrate for the first time that MATN-1 is an inhibitor of angiogenesis. We have further demonstrated that the mechanism by which MATN-1 inhibits angiogenesis is through the potent inhibition of growth factor-stimulated capillary endothelial cell proliferation and migration. These two activities have been shown by our laboratory and many others to be necessary mechanisms for the formation of new capillaries from pre-existing vessels, ie., angiogenesis (36). These *in vivo* data are supported by both the *in vitro* gain of function

experiments (Fig. 3 and Fig. 4) and an *in vivo* loss of function study using MATN-1 KO mice (Fig. 5 and Fig. 6).This novel activity of MATN-1 may be due to structural and/or regulatory roles of MATN-1 in cartilage. MATN-1 contains the vWFA domain, also called the I domain in the integrin family, which is responsible for the MATN-1 interaction with ECM and integrins via the MIDAS motif (28). This interaction may affect angiogenesis which critically depends on cell-matrix interaction (4,36). This hypothesis remains to be tested in future.

MATN-1 is a modular matrix protein capable of mediating interactions between a variety of matrix components through von Willebrand Factor type A (vWFA) domains that intervene through adhesion via metal ion-dependent adhesion sites (MIDAS). These domains exhibit high affinity toward collagen which also possesses vWFA-like domains (24), as well as biglycan and decorin (25). The discovery of a non-collagenous matrix protein as an angiogenesis inhibitor is not unprecedented. Decorin, a small proteoglycan composed of a protein core and a covalently linked side chain of chrondroitin/dermatan sulfate, has been shown to inhibit endothelial cell migration, attachment, and the formation of endothelial tube-like structures (37,38). MATN-1 is the newest member of the family of matrix-derived inhibitors of neovascularization. As is the case with thrombospondin-1 and -2 (39-41), it distinguishes itself from other inhibitors in this family such as endostatin (42,43), tumstatin (44,45), arresten (46) and canstatin (47) in that these latter inhibitors are proteolytically-processed products of larger, largely angiogenesis-inert matrix proteins (48,49).

Our group and others have reported the discovery of novel cartilage-derived anti-angiogenic proteins (4,6,13) and a number of other groups have reported the presence of potential inhibitors of neovascularization from non-mammalian sources of cartilage as well. For example, fractionated samples of shark cartilage extracts, such as U-995 (15), SCF2 (1), SCAIF 80 (16), and DCAI (50), have shown preliminary promise in inhibiting angiogenesis, however the active biomolecule has yet to be purified and identified. SCP1, a low molecular weight protein extracted from shark cartilage, with a sequence similar to alpha parvalbumin, a calcium-binding molecule, has been shown to exert some anti-angiogenic activity in a rat aortic ring assay (17). Neovastat, a purified fraction of shark cartilage extract with MMP inhibitory activity (51), was developed for clinical use and demonstrated limited success in early phase I/II clinical trials as treatment for non-small cell lung cancer and renal cell carcinoma (52-54). Ultimately, in phase III trials, when coupled with standard chemotherapy, Neovastat had no additive effect on the improvement of overall survival in patients with unresectable stage III non-small cell lung cancer (55). Neovastat has had limited success in the treatment of renal cell carcinoma (56).

MATN-1 is an inhibitor of both capillary endothelial cell proliferation and migration, processes which, when driven by endothelial cell mitogens, represent critical processes required in order to assess angiogenic potential. It is an inhibitor of angiogenesis in vivo in a model of normal, embryonic angiogenesis such as the CAM, as well as in a model of pathological neovascularization and in the mice tibia fracture model. As such, this protein, either alone or in combination with other inhibitors, may be useful in the treatment of a number of diseases that are characterized by dysregulated neovascularization.

ACKNOWLEDGEMENTS

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FOOTNOTES

The abbreviations used are: CCE, concentrated cartilage extract; MATN-1, Matrilin-1; aFGF, acidic fibroblast growth factor; bFGF, basic fibroblast growth factor,;TIMP-1, tissue inhibitor of metalloproteinases 1; TnI, Troponin I.

FIGURE LEGENDS

Figure 1.   Purification of anti-angiogenic activity in CCE and Identification of anti-angiogenic activity as being MATN-1
Fractions 22-34 of CCE fractionated on a Biogel A-1.5M size-exclusion chromatography significantly inhibited mitogen-stimulated capillary EC proliferation. Fractions 52-75 contained metalloproteinase-inhibiting activity but had no significant effect on EC proliferation (**A**). Fractions 22-34 were pooled, concentrated, dialyzed and subjected to further purification on a Bio-Rex 70 cation exchange column. Fractions 2 and 5 contained inhibitory activity of FGF-stimulated capillary EC proliferation with no significant TIMP activity (**B**). The antiproliferative activity from the Bio-Rex 70 column was concentrated and electrophoresed on a 12% SDS-PAGE gel followed by staining with SYPRO Ruby (**C**). Bands were excised from the gel and analyzed by mass spectrometry with the band highlighted by the red box having two peptide matches identifying it as being matrilin-1 (**C,D,E**). To confirm this identification, a sample of the same fraction from the cation exchange column was electrophoresed on a 12% SDS-PAGE gel and probed with an antibody to matrilin-1(**F**).

Figure 2. MATN-1 suppresses angiogenesis *in vivo*
Recombinant chick MATN-1 and human MATN-1 inhibit angiogenesis *in vivo* as indicated by the avascular zones localizing the zones of new capillary inhibition on the chick chorioallantoic membrane induced by (**B**)10 µg of rcMATN-1 and (**C**) 5 µg of rhMATN-1.

Figure 3.   Inhibition of bFGF-stimulated bovine capillary endothelial cell growth by MATN-1
Purified cMATN-1 (**A**), rcMATN-1 (**B**), and rhMATN-1 (**C**) were tested for their ability to inhibit bFGF-stimulated EC growth. In all cases, MATN-1 was found to inhibit EC proliferation in dose-dependent manner.

Figure 4.   Inhibition of bFGF-stimulated bovine capillary EC migration by MATN-1
Purified cMATN-1 (**A**), rcMATN-1 (**B**), and rhMATN-1 (**C**) were tested for their ability to inhibit bFGF-stimulated EC migration. In all cases, MATN-1 was found to inhibit EC migration in a dose-dependent manner. Cells were labeled as described in Experimental Procedures and allowed to migrate through the transwell. Representative images of migrated cells treated with (**D**) 0 nM, (**E**) 10 nM, (**F**) 100 nM, and (**G**) 200 nM of recombinant human MATN-1.

Figure 5. *In vivo* imaging and quantification of AngioSense signal in WT and MATN-1 KO mice on Day 4 post fracture.
Representative volume renderings taken at the same color gating from WT (**A**) and MATN-1 -/- mice (**B**) injected with AngioSense according to manufacturer's directions. The total amount of fluorescence (pmol) was quantified in specific regions of interest encompassing fracture tibias and control tibias. MATN-1 KO mice showed a significant 1.76 fold increase in AngioSense signal ($p = 0.0041$) in fractured tibia compared to WT mice (**C**). There was no difference between MATN-1 KO and WT mice in control tibias (**D**; $p = 0.0970$). MATN-1 KO mice had 1.87 fold higher AngioSense signal compared to WT mice (**E**), when the signal amount of fractured tibia was subtracted from that of control tibias ($p = 0.0156$).*$p<0.05$.n=5.

Figure 6. mRNA expression of angiogenesis-related genes in the fracture calluses of MATN-1 KO and WT mice on Day 14 determined by real-time RT-PCR analysis
MATN-1 KO mice had significantly higher (**A**) PECAM1 ($p= 0.0041$) (**B**) VEGFR ($p= 0.0115$), and (**C**) VE-cadherin ($p= 0.0004$) expression levels compared to WT mice. *$p<0.05$, n=5.
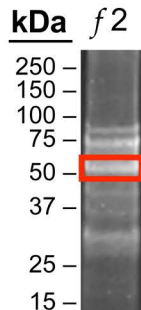
A
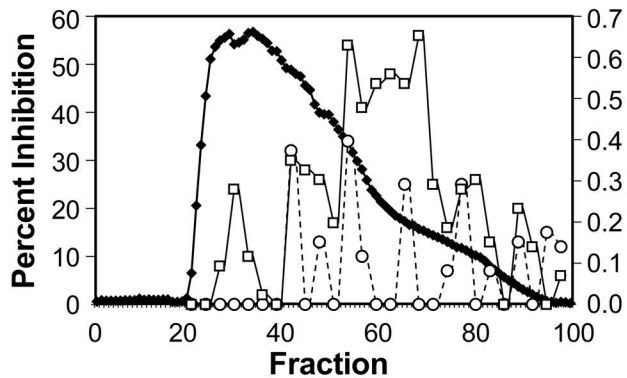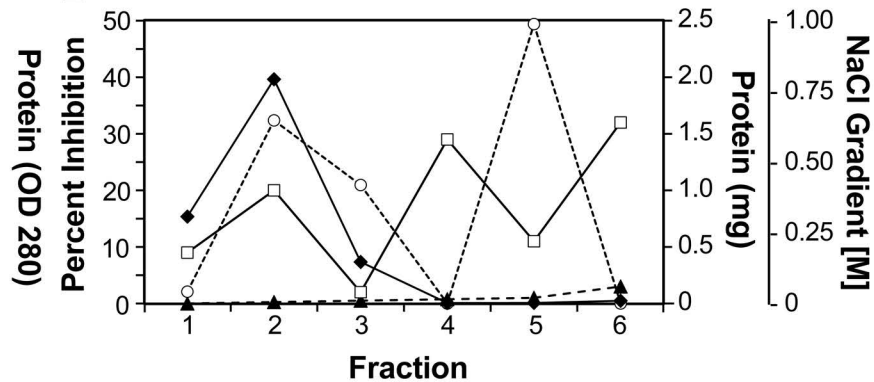
B

C

kDa   ƒ2

D

>GI|115555|SP|P05099, CAMA_CHICK Cartilage matrix protein precursor (Matrilin-1)
MDGIFCALPLSLLLLLQSCGVWGAPPQPRGTLCRTKPTDLVFIIDSSRSVRPQEFEKVKVFLSRVIEGLD
VGPNSTRVGVINYASAVKNEFSLKTHQTKAELLQAVQRIEPLSTGTMTGLAIQFAISRAFSDTEGARLRS
PNINKVAIVVTDGRPQDGVQDVSARARQAGIEIFAIGVGRVDMHTLRQIASEPLDDHVDYVESYSVIEKL
THKFQEAFCVVSDLCATGDHDCEQICISTPGSYKCACKEGFTLNNDGKTCSACSGGSGSALDLVFLIDGS
K**SVRPENFELVKK**FINQIVESLEVSEKQAQVGLVQYSSSVRQEFPLGQFKNKKDIKAAVKKMAYMEKGTM
TGQALKYLVDSSFSIANGARPGVPK**VGIVFTDGR**SQDYITDAAKKAKDLGFRMFAVGVGNAVEDELREIA
SEPVAEHYFYTADFRTISNIGKKLQMKICVEEDPCECKSIVKFQTKVEELINTLQQKLEAVAKRIEALENKII

E

SVRPENFELVKK

Calculated [M+H]⁺ = 1446.69
Calculated [M+2H]²⁺ = 723.85
Experimental [M+3H]³⁺ = 482.96
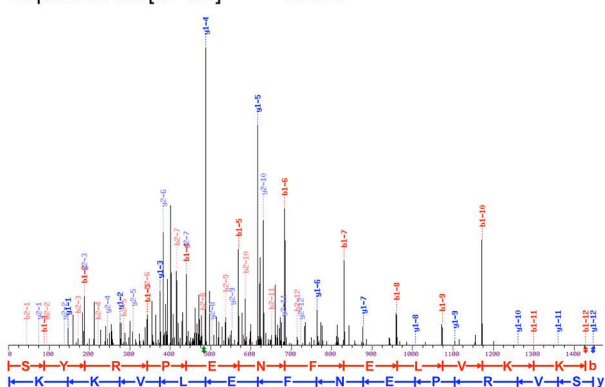
VGIVFTDGR

Calculated [M+H]⁺ = 964.11
Calculated [M+2H]²⁺ = 482.56
Experimental [M+2H]²⁺ = 482.17

F

kDa
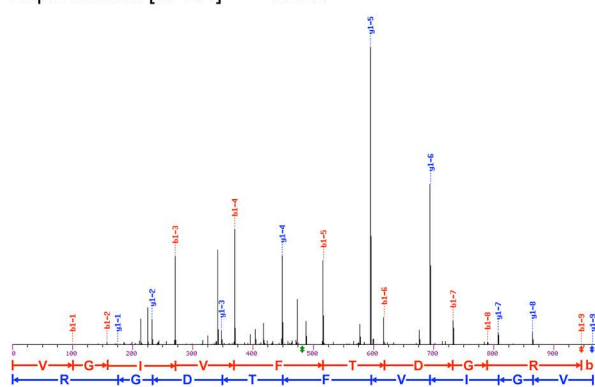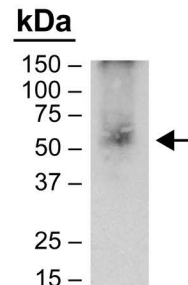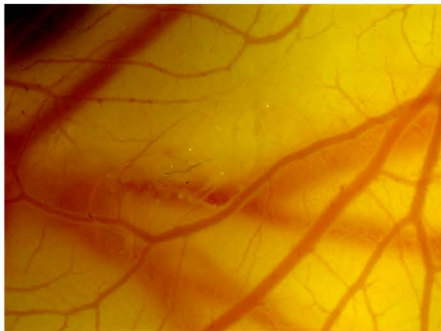
Figure 1

A PBS
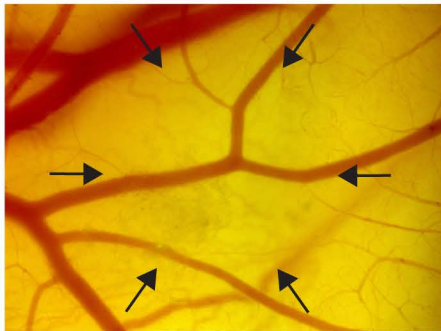
B rcMATN-1
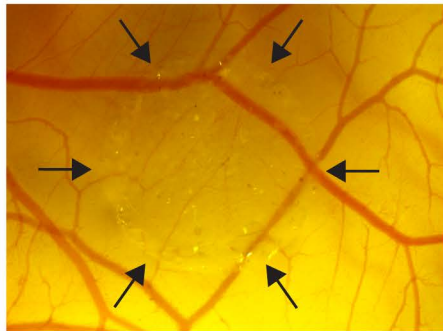
C rhMATN-1

Figure 2
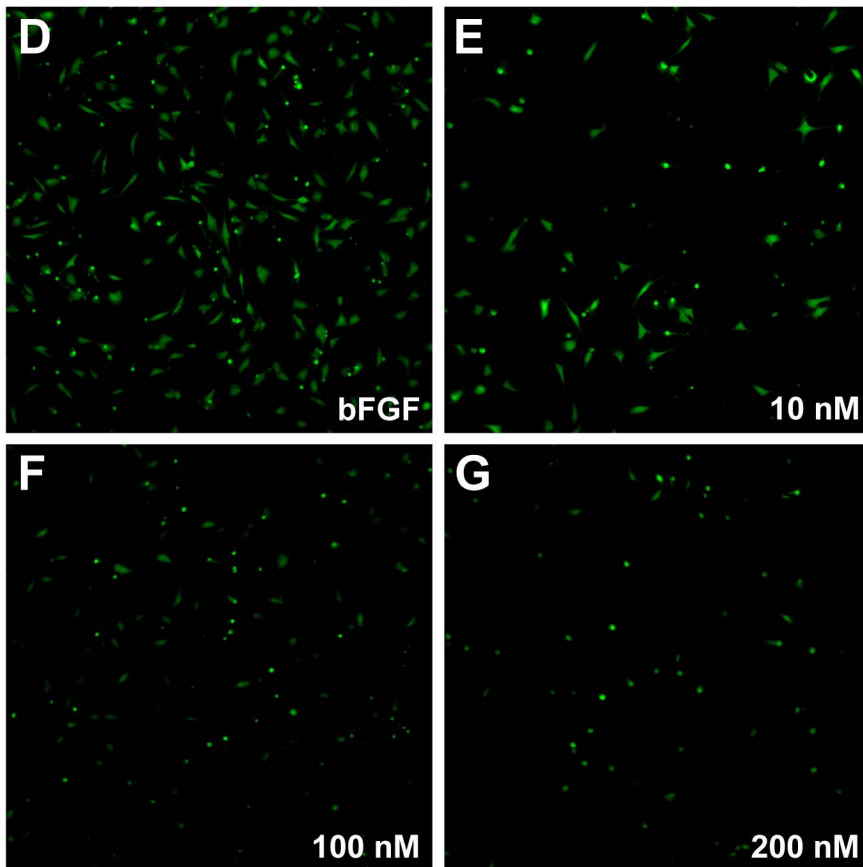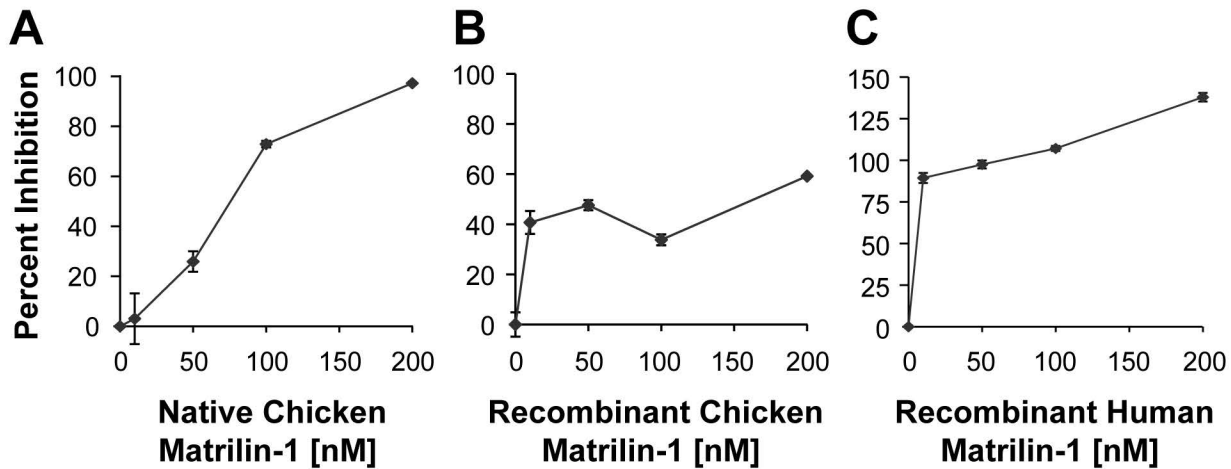
Figure 4
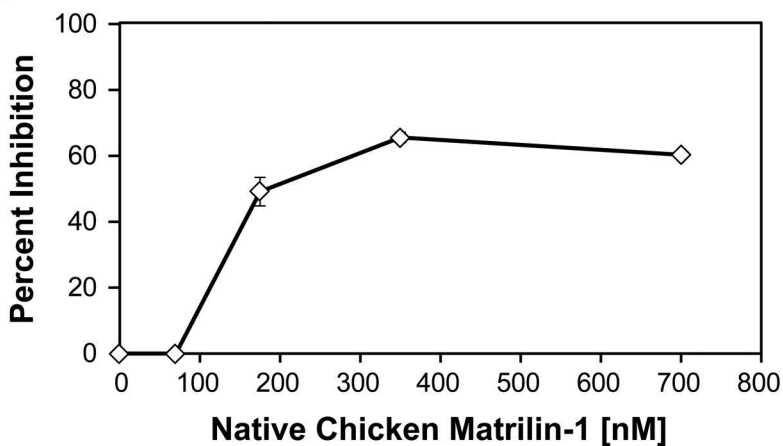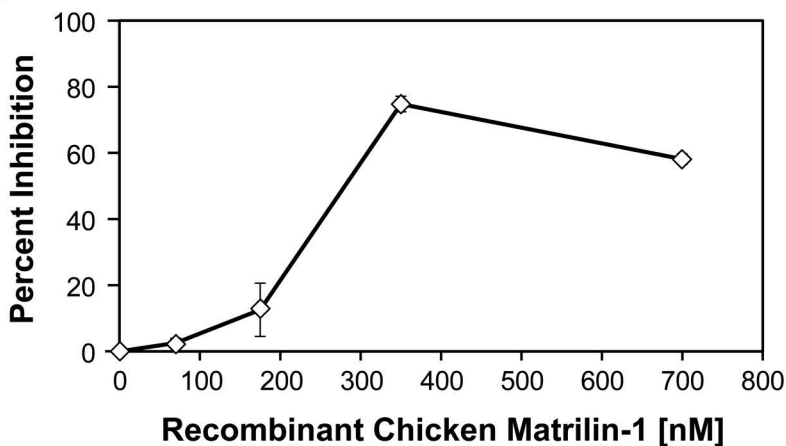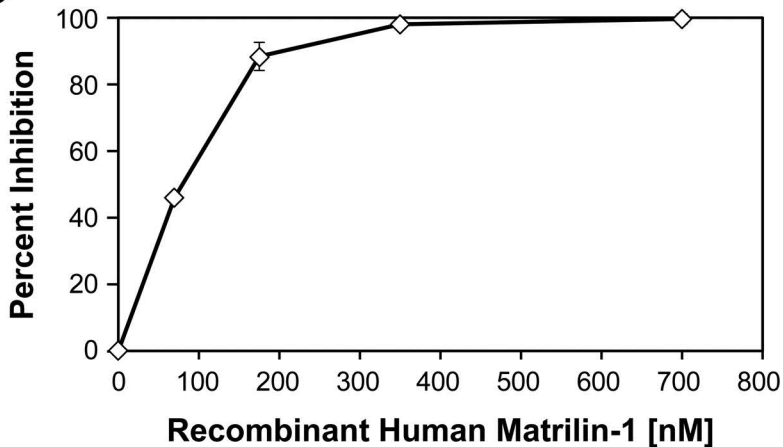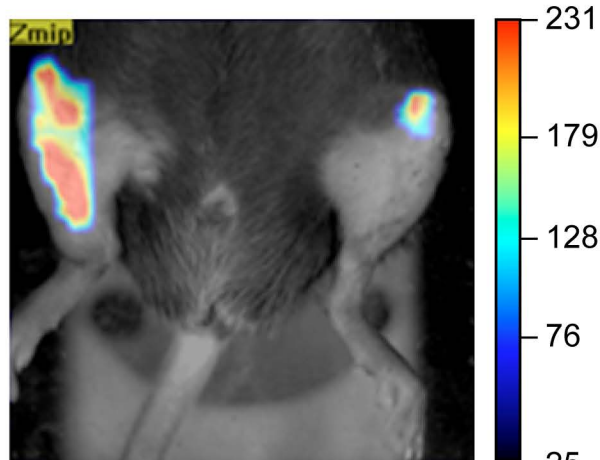
Figure 3

A WT
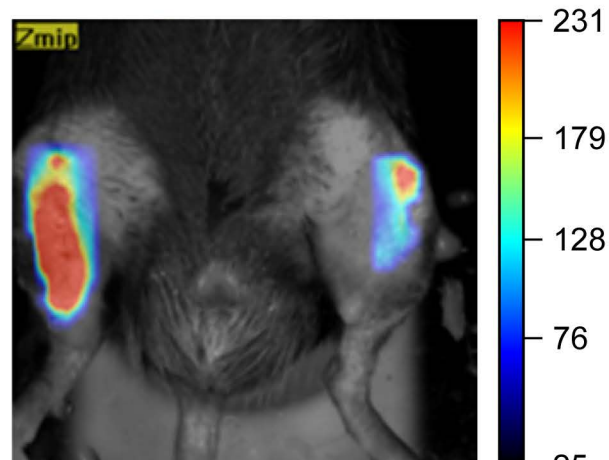
B MATN1−/−

C Fracture side
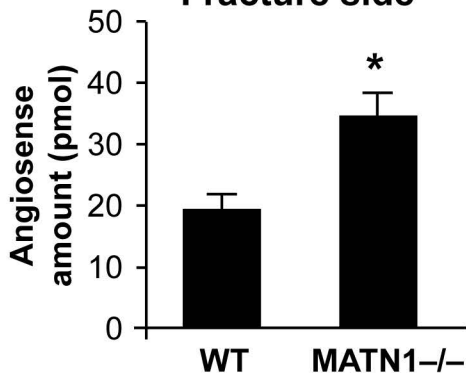
D Control side

E Fracture-Control

Angiosense amount (pmol)

Figure 5

Figure 6