The L6 Protein TM4SF1 Is Critical for Endothelial Cell Function and Tumor Angiogenesis

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Abstract

Transmembrane-4-L-six-family-1 (TM4SF1) was originally described as a cancer cell protein. Here, we show that it is highly expressed in the vascular endothelium of human cancers and in a banded pattern in the filopodia of cultured endothelial cells (EC). TM4SF1 knockdown prevented filopodia formation, inhibited cell mobility, blocked cytokinesis, and rendered EC senescent. Integrin-α5 and integrin-β1 subunits gave a similar staining pattern and interacted constitutively with TM4SF1, whereas integrin subunits often associated with angiogenesis (αV, β3, β5) interacted with TM4SF1 only after vascular endothelial growth factor (VEGF)-A or thrombin stimulation. TM4SF1 knockdown substantially inhibited maturation of VEGF-A164-induced angiogenesis. Thus, TM4SF1 is a key regulator of EC function in vitro and of pathologic angiogenesis in vivo and is potentially an attractive target for antiangiogenesis therapy. [Cancer Res 2009;69(8):3272–7]

Introduction

Transmembrane-4-L-six-family-1 (TM4SF1), also known as L6, was first identified as a protein abundantly expressed in a variety of epithelial cancer cells (1, 2) and weakly expressed in normal vascular endothelium (3, 4). It has the topology of a tetraspanin (5); however, TM4SF1 and five other structurally similar proteins (TM4SF4, 5, 18, 19, and 20) lack overall sequence homology with the 33 genuine tetraspanins, and the characteristic CCG motif in the large extracellular loop (5). Therefore, these proteins have been classified separately as the L6 family (5).

Experiments on tumor cells had previously shown TM4SF1 to be important for growth (1), motility (6), invasion (7), and metastasis (8). However, the role of TM4SF1 in vascular endothelium has not been investigated. We report here that TM4SF1 is overexpressed in the vascular endothelial cells (EC) of several human cancers and is critically important for their function.

Materials and Methods

Antibodies, soluble factors. Antibodies were as follows: mouse anti-human TM4SF1, ITGAS, ITGAV, ITGB1, ITGB3, and ITGB5 (Millipore); Cy3-conjugated mouse anti-human α-smooth muscle actin (αSMA; Sigma); goat anti-human VE-cadherin (Santa Cruz); FITC-labeled anti-mouse IgG; and Texas red-labeled rabbit anti–goat-IgG (Invitrogen). Vascular endothelial growth factor (VEGF)-A, basic fibroblast growth factor (bFGF), transforming growth factor β (TGFβ), hepatocyte growth factor (HGF), and tumor necrosis factor (TNF)α were from R&D Laboratories, and thrombin was from Calbiochem.

Mice, adenoviral vectors. Athymic nude mice were obtained from National Cancer Institute. All studies were performed under protocols approved by the Beth Israel Deaconess Medical Center Institutional Animal Care and Use Committee. Adenoviral vectors expressing mouse VEGF-A164 (Ad-VEGF-A164) and Lac-Z (Ad-Lac-Z) were described previously (9–11).

Cells, flow cytometry, and immunostaining. Human umbilical vein ECs (HUVEC) and other EC were from Clonetics. Cells were grown in growth medium-2-MV (Clonetics) and used at passage 5 to 6. For flow cytometry, EC were stained with propidium iodide and 104 events were collected for each experiment (Becton Dickinson). EC were grown on 8-mm cover glasses, fixed in 4% paraformaldehyde (5 min, 25°C), blocked with PBS/1% fetal bovine serum, and immunostained (12). Filamentous actin (F-actin) was visualized with rhodamine-conjugated phalloidin (Millipore). Crystal violet (50 mg/mL, 2 h, 25°C) and senescent Staining kit (Sigma) were from Sigma. Paraformaldehyde-fixed, paraffin-embedded tissue sections were stained with hematoxylin or were immunostained with TM4SF1 primary Ab (Millipore), mouse IgG; and Texas red–labeled rabbit anti–goat-IgG (Invitrogen). Immunoprecipitation and Western blotting. Human umbilical vein ECs (HUVEC) and other EC were from Clonetics. Cells were grown in growth medium-2-MV (Clonetics) and used at passage 5 to 6. For flow cytometry, EC were stained with propidium iodide and 104 events were collected for each experiment (Becton Dickinson). EC were grown on 8-mm cover glasses, fixed in 4% paraformaldehyde (5 min, 25°C), blocked with PBS/1% fetal bovine serum, and immunostained (12). Filamentous actin (F-actin) was visualized with rhodamine-conjugated phalloidin (Millipore). Crystal violet (50 mg/mL, 2 h, 25°C) and senescent Staining kit (Sigma) were from Sigma. Paraformaldehyde-fixed, paraffin-embedded tissue sections were stained with hematoxylin or were immunostained with TM4SF1 primary Ab (Millipore), mouse IgG; and Texas red–labeled rabbit anti–goat-IgG (Invitrogen). Immunoprecipitation and Western blotting. Human umbilical vein ECs (HUVEC) and other EC were from Clonetics. Cells were grown in growth medium-2-MV (Clonetics) and used at passage 5 to 6. For flow cytometry, EC were stained with propidium iodide and 104 events were collected for each experiment (Becton Dickinson). EC were grown on 8-mm cover glasses, fixed in 4% paraformaldehyde (5 min, 25°C), blocked with PBS/1% fetal bovine serum, and immunostained (12). Filamentous actin (F-actin) was visualized with rhodamine-conjugated phalloidin (Millipore). Crystal violet (50 mg/mL, 2 h, 25°C) and senescent Staining kit (Sigma) were from Sigma. Paraformaldehyde-fixed, paraffin-embedded tissue sections were stained with hematoxylin or were immunostained with TM4SF1 primary Ab (Millipore), mouse IgG; and Texas red–labeled rabbit anti–goat-IgG (Invitrogen). Immunoprecipitation and Western blotting. Human umbilical vein ECs (HUVEC) and other EC were from Clonetics. Cells were grown in growth medium-2-MV (Clonetics) and used at passage 5 to 6. For flow cytometry, EC were stained with propidium iodide and 104 events were collected for each experiment (Becton Dickinson). EC were grown on 8-mm cover glasses, fixed in 4% paraformaldehyde (5 min, 25°C), blocked with PBS/1% fetal bovine serum, and immunostained (12). Filamentous actin (F-actin) was visualized with rhodamine-conjugated phalloidin (Millipore). Crystal violet (50 mg/mL, 2 h, 25°C) and senescent Staining kit (Sigma) were from Sigma. Paraformaldehyde-fixed, paraffin-embedded tissue sections were stained with hematoxylin or were immunostained with TM4SF1 primary Ab (Millipore), mouse IgG; and Texas red–labeled rabbit anti–goat-IgG (Invitrogen).

Virus titration was determined by multiplicity of infection (moi).

Multigene transcriptional profiling. Multigene transcriptional profiling is a quantitative real-time PCR technique that efficiently provides mRNA copies per cell (14). Total mRNA was prepared using the RNeasy kit (Qiagen) and cDNA using random primers and SuperScript III (Invitrogen). For each data point, mean ± SD were calculated from three samples from three separate experiments. PCR reactions for each cDNA sample were performed in duplicate. Transcript abundances were normalized per 105 18S-rRNA copies to approximate number of transcripts per cell (15, 16).

In situ hybridization. In situ hybridization was performed as described (17). Tissues were fixed in 4% paraformaldehyde (5 min, 25°C) and cryostat sections were developed with the ECL system (GE Healthcare).

Angiogenesis quantification. Evans’s Blue dye was injected i.v. into nude mice anesthetized with Avertin (200 mg/kg; ref. 10). Five minutes later, after equilibration in blood but before significant plasma leakage, mice were sacrificed as a cancer cell protein. Here, we show that it is highly expressed in the vascular endothelium of human cancers and in a banded pattern in the filopodia of cultured endothelial cells (EC). TM4SF1 knockdown prevented filopodia formation, inhibited cell mobility, blocked cytokinesis, and rendered EC senescent. Integrin-α5 and integrin-β1 subunits gave a similar staining pattern and interacted constitutively with TM4SF1, whereas integrin subunits often associated with angiogenesis (αV, β3, β5) interacted with TM4SF1 only after vascular endothelial growth factor (VEGF)-A or thrombin stimulation. TM4SF1 knockdown substantially inhibited maturation of VEGF-A164-induced angiogenesis. Thus, TM4SF1 is a key regulator of EC function in vitro and of pathologic angiogenesis in vivo and is potentially an attractive target for antiangiogenesis therapy. [Cancer Res 2009;69(8):3272–7]
euthanized and angiogenic sites were harvested with an 8-mm punch. Dye was extracted at 65°C for 48 h in 0.5 mL formamide. Triplicate 100 μL samples were measured at OD_{620} nm.

Statistical analysis. Statistics were performed with Student's t test.

Results and Discussion

TM4SF1 expression in human cancers. TM4SF1 was found to be highly expressed in the vascular endothelium of smooth muscle cell–coated blood vessels that supply several important human cancers, including those in which the malignant cells themselves did not express TM4SF1 detectably (Fig. 1A and B).

The vasculature supplying nearby normal tissues expressed TM4SF1 weakly (data not shown), consistent with published observations (3).

TM4SF1 expression in cultured EC. TM4SF1 was expressed at consistently high levels (70–90 mRNA copies per cell) in several cultured EC, including umbilical vein (HUVEC), coronary artery (human coronary artery endothelial cell), pulmonary artery (human pulmonary artery endothelial cell), dermal microvascular (human dermal microvascular endothelial cell), and lymphatic (human lymphatic endothelial cell; Fig. 1C, left). In contrast, TM4SF1 expression was low (<5 mRNA copies per cell) in human dermal fibroblasts and smooth muscle cells (data not shown).

Figure 1. TM4SF1 expression in the vascular endothelium of human cancers and in cultured EC. A, in situ hybridization shows strong TM4SF1 expression in the blood vessels of colon, kidney, ovary, and breast carcinomas. Images shown are representative of those taken from four different patients with each type of cancer. Bright field reveals histology and dark field shows mRNA expression. Arrows, TM4SF1-positive blood vessels. B, immunofluorescence staining of a typical colon cancer with antibodies against TM4SF1 (green) and either (i) VE-cadherin or (ii) anti-human α-smooth muscle actin (red). C, TM4SF1 expression in several types of cultured EC (left) and in HUVEC stimulated with indicated growth factors (right). *, P < 0.05. D, TM4SF1 (green) and F-actin (red) double-staining of HUVEC reveals that TM4SF1 is localized to the plasma membrane, perinuclear vesicles (arrow), and particularly to filopodia (a). Filopodia exhibited a banded staining pattern (b and c). Arrows, filopodia branching (c). Scale bars, 100 μm (A and B) and 10 μm (D).
At 6 hours (but not at 1 or 18 hours), VEGF-A (20 ng/mL) and thrombin (1.5 units/mL) stimulated TM4SF1 expression modestly in HUVEC (≈1.5-fold), whereas TGFβ (1 ng/mL), bFGF (50 ng/mL), TNFα (20 ng/mL), and HGF (20 ng/mL) did not significantly alter TM4SF1 expression at any time (Fig. 1C, right).

In subconfluent HUVEC, TM4SF1 was localized to the plasma membrane, to a minor population of perinuclear vesicles, and particularly to filopodia, some of which branched (Fig. 1D). Filopodia exhibited an intermittent, banded pattern of TM4SF1 staining (Fig. 1Db and c). F-actin, a quintessential filopodial marker, was confined to the most proximal portions of filopodia, generally within 10 μm of the cell body (Fig. 1D). Human coronary artery endothelial cell, human pulmonary artery endothelial cell, human dermal microvascular endothelial cell, and human lymphatic endothelial cell showed similar TM4SF1 staining patterns (data not shown).

TM4SF1 knockdown in HUVEC. After transfection of HUVEC with either of two human TM4SF1 shRNAs, TM4SF1 mRNA abundance was reduced by 70% to 95% from days 2 to 15, whereas TM4SF1 expression was unaffected by control shRNA that does not recognize any known human or mouse mRNA (Fig. 2A).

TM4SF1 knockdown (KD) HUVEC underwent striking changes in structure and function. They enlarged greatly, accumulated numerous circumferential stress fibers, and were unable to complete cytokinesis, arresting in G2-M (Fig. 2Bi–iii). Binucleate cells were observed a day after transfection, and by day 5, ≈60% of cells were binucleate and 30% multinucleate. Nonetheless, KD cells remained viable for as long as they were followed in culture (15 days); throughout this period, 18S integrity was maintained and...
Ki-67 expression levels were low (<10% of control cells) as determined by multigene transcriptional profiling. In addition, TM4SF1-KD HUVEC developed high levels of β-galactosidase activity (Fig. 2Biv), resisted trypsin detachment from culture dishes, and, once removed, reattached poorly. Furthermore, KD cells failed to migrate or proliferate after VEGF-A stimulation (data not shown). TM4SF1-KD HUVEC therefore exhibited senescence (17).

Moreover, the occasional HUVEC that underwent spontaneous senescence in culture lost most of their TM4SF1 staining (Fig. 2C).

In addition, TM4SF1-KD HUVEC failed to extend filopodia (Fig. 2Bi and ii), migrated poorly in a wound healing assay, and failed to form tubes in Matrigel (Fig. 2D). Furthermore, although exhibiting extensive lamellipodial activity, KD HUVEC failed to move as much as a quarter cell diameter from their original position over 1.5 hours of time lapse imaging, whereas control shRNA-transfected HUVEC migrated extensively (Supplementary Fig. S1). TM4SF1 staining patterns and KD phenotype were similar in human pulmonary artery endothelial cell, human coronary artery endothelial cell, human dermal microvascular endothelial cell, and human lymphatic endothelial cell. However, cells that expressed TM4SF1 weakly (human dermal fibroblast, human smooth muscle cell) were unaffected by TM4SF1-KD (data not shown).

**Figure 3.** TM4SF1-integrin expression and interactions. A, integrin expression profile in HUVEC ± TM4SF1 KD. B, immunofluorescence staining of integrin α5, αV, and β1 subunits in HUVEC 3 d after 50-moi transfection with either Ad-hTM4-shRNA or Ad-Con-shRNA. Scale bars, 10 μm. C, control HUVEC or HUVEC stimulated for 6 h with VEGF-A (20 ng/mL) or thrombin (THR; 1.5 u/mL). Cell lysates, prepared before and after immunoprecipitation (IP) with anti-TM4SF1 antibody, were immunoblotted with indicated antibodies. IgG and β-actin were loading controls. The anti-TM4SF1 antibody effectively immunoprecipitated native TM4SF1 but reacted poorly with denatured TM4SF1. Expression of β3 and β5 were below detection levels in whole cell lysates.
TM4SF1 interactions with integrins. Classic tetraspanins interact with integrins and participate in a variety of integrin-associated functions (18), but L6 family interactions with integrins have not been investigated. Of the 26 integrin subunits, only 3 (ITGA5, ITGAV, ITGB1) were highly expressed in HUVEC, and their expression was not affected significantly by TM4SF1-KD (Fig. 3A). Antibodies against α5 and -h1 stained filopodia in an intermittent, banded pattern (Fig. 3B), similar to that of TM4SF1 (Fig. 1Db and c). In contrast, αV staining was confined predominantly to focal contacts (Fig. 3B). Enlarged, binucleate TM4SF1-KD HUVEC continued to show normal α5, αV, and β3 cell membrane staining (Fig. 3B). Thus, the failure of TM4SF1-KD HUVEC to form filopodia and migrate is not attributable to changes in the abundances of α5, αV, or their β partners.

VEGF-A and thrombin are known to stimulate integrin expression and EC mobility (19, 20). These agents modestly but significantly increased ITGAV mRNA expression (1.4- to 1.6-fold) in HUVEC without affecting ITGA5 or ITGB1 expression (data not shown). Immunoblots revealed that both α5 and -h1 associated constitutively with TM4SF1 and that these associations were minimally affected by VEGF-A or thrombin (Fig. 3C). However, interaction of TM4SF1 with αV and its β3 and β5 partners was only detected after VEGF-A or thrombin stimulation. Thus, TM4SF1, such as classic tetraspanins (18), interacts with integrins, and its pairing depends on the state of EC activation.

Role of TM4SF1 in VEGF-A–induced angiogenesis. To assess the role of TM4SF1 in vivo, we made use of an adenoviral vector expressing VEGF-A164 (Ad-VEGF-A164) that induces an angiogenic response in mouse ears at indicated times after injection of 2.5 × 10⁷ plaque-forming unit Ad-VEGF-A164. B, in situ hybridization localization of TM4SF1 mRNA to VM in day 11 Ad-VEGF-A164–injected mouse ears (top). Hematoxylin-stained VM illustrating smooth muscle cell coat (green arrows, bottom). C, angiogenic response in mouse ears at indicated times after Ad-VEGF-A164 injection. Moi (10⁹) of Ad-mTM4-shRNA or Ad-Con-shRNA were injected into the right and left ears, respectively, 1 d before Ad-VEGF-A164 injection. The experiment was repeated four times. D, quantification of angiogenesis with the Evan’s Blue dye 19 d after Ad-VEGF-A164 injection.
response in mouse tissues that closely mimics tumor angiogenesis (9, 10). Messenger RNA profiling showed that TM4SF1 expression increased at sites of Ad-VEGF-A164–induced angiogenesis, although not in the initial (1–5 day) phase (Fig. 4A). This earliest phase of VEGF-A–induced angiogenesis is associated with the formation of mother vessels (MV), greatly enlarged, pericyte-poor sinusoids that derive from preexisting normal vessels (11). Instead, TM4SF1 expression increased later, peaking at day 11 as MV differentiated into daughter vascular malformations (VM), MV that have acquired a smooth muscle cell coat (Fig. 4B). Expression of ITGAV, ITGB5, and ITGB1 mRNAs closely tracked that of TM4SF1, whereas ITGA5 and ITGB3 expression increased earlier with MV formation (Fig. 4A). Consistent with these findings, in situ hybridization showed that TM4SF1 was strongly expressed in VM (Fig. 4B).

We knocked down TM4SF1 by injecting an adenovirus expressing mouse TM4SF1–specific shRNA (Ad-mTM4-shRNA) into mouse ears 1 day before injection of Ad-VEGF-A164. Ad-mTM4-shRNA did not affect the initial, MV phase of angiogenesis induced by Ad-VEGF-A164 (Fig. 4C). However, at day 11 and later, MV maturation into VM was substantially blunted by Ad-mTM4-shRNA but not by Ad-Con-shRNA (Fig. 4C). Vascular volume, a surrogate measure of angiogenesis, was significantly decreased in ears injected with Ad-mTM4-shRNA (Fig. 4D). VM formation was also reduced as determined by histology (data not shown). Thus, TM4SF1 plays a critical role in vessel maturation.

In summary, TM4SF1 is overexpressed in the blood vessels of several important human cancers. It is highly expressed in several types of cultured EC and is essential for their maintenance and function. When TM4SF1 was knocked down, HUVEC failed to filipodia, became immobile, and assumed a senescent phenotype. Like classic tetraspansins, TM4SF1 interacts with integrins and may affect cell motility via such interactions. TM4SF1-KD affected vascular maturation in the Ad-VEGF-A164 mouse ear angiogenesis model, perhaps by interacting with integrins to mediate EC migration and promote cell-cell interactions. These data indicate that TM4SF1 is an important new player in EC biology that it could have potential in angiogenesis therapy.

**Disclosure of Potential Conflicts of Interest**

No potential conflicts of interest were disclosed.

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