Effect of tenascin-X together with vascular endothelial growth factor A on cell proliferation in cultured embryonic hearts.
Effect of Tenascin-X Together with Vascular Endothelial Growth Factor A on Cell Proliferation in Cultured Embryonic Hearts

Tomoki IKUTA, Hiroyoshi ARIGA, and Ken-ichi MATSUMOTO*

Department of Molecular Biology, Graduate School of Pharmaceutical Sciences, Hokkaido University, Kita 12, Nishi 6, Kita-ku, Sapporo 060–0812, Japan. Received August 3, 2001; accepted September 3, 2001

Tenascin-X (TNX) is a large glycoprotein that appears in extracellular matrices. Previously, we demonstrated that TNX binds to vascular endothelial growth factors A and B (VEGF-A and -B) and that VEGF-B in combination with TNX induces DNA synthesis in endothelial cells via increased signals mediated by the VEGFR-1 receptor. In this study, we investigated the effect of TNX with VEGF-A on the cell proliferation in embryonic mouse heart explants from either wild-type (TNX+/+) or TNX-deficient (TNX−/−) mice. The addition of VEGF-A to the explants from TNX+/+ mice increased cell proliferation by 1.5 fold compared with that in TNX−/− mice, indicating that TNX with VEGF family member plays an important role in the control of endothelial cell proliferation in vivo.

Key words tenascin-X; vascular endothelial growth factor A; extracellular matrix

Cell differentiation is a continuously regulated process and is dependent upon interactions between the cell and its environment. An important component of the cellular environment is provided by extracellular matrix proteins. Tenascins are a family of extracellular matrix proteins. Tenascin-C (TNC), the first member of the tenascin family to be discovered, has been shown to change its expression pattern during important steps of embryogenesis as well as in pathological conditions, including cancer and wound healing, indicating that TNC plays crucial roles in embryonic development and tissue restructuring. To date, five members of the tenascin family have been characterized: TNC, tenascin-R (TNR), tenasin-X (TNX), tenasin-Y (TNY), and tenascin-W (TNW).

The known tenasin family members share a very distinctive pattern of four domains: heptad repeats, epidermal growth factor (EGF)-like repeats, fibronectin type III-like (FNIII) repeats, and a globular domain shared with fibrinogens.

TNX, the largest member in the tenascin family, is closely related to TNC. It is expressed much more widely than other tenascin family members. TNX is concentrated in the connective tissue of muscle and around blood vessels. In many regions of the body, particularly in the digestive tract and skin, the distributions of TNX and TNC are frequently complementary, suggesting that the genes are regulated by distinctive cis-elements and transcription factors. Recently, we showed that TNX binds to vascular endothelial growth factor B (VEGF-B) and that VEGF-B in combination with TNX promotes DNA synthesis in endothelial cells compared to VEGF-B alone. This effect of TNX and VEGF-B on cell mitogenecity is mediated by an increased signal via a tyrosine kinase receptor, VEGF receptor 1 (VEGFR-1).

In addition to VEGF-B, VEGF-A, VEGF-C, VEGF-D, and placenta growth factor (PIGF) have also been identified as members of VEGF family of growth factors. VEGF-A is known to be the most potent inducer of endothelial cell proliferation through its two high-affinity receptors VEGFR-1 and VEGFR-2/KDR/Flk-1. Recenty, integrin αvβ3 was reported to make a molecular complex with activated VEGFR-2. We have shown that TNX is also capable of binding to VEGF-A. It has also been shown that VEGF-C and VEGF-D stimulate lymphangiogenesis via VEGFR-3/Flt-4 and that they also weakly induce the migration and proliferation of endothelial cells mediated by VEGFR-2.

In an attempt to recapitulate vasculogenic events in vitro, Ratajka et al. demonstrated that VEGF-A stimulates cell proliferation and assembly into cord-like structures in embryonic rat heart explants at 14 d of gestation (E14) cultured on collagen gels. In this study, we isolated heart explants from wild-type or TNX-deficient mouse embryos at E9.5 and cultured them in the absence or presence of VEGF-A and VEGF-B to determine the effects of TNX with these growth factors on the cell proliferation response. We found that cell proliferation outgrowing from E9.5 heart explants was significantly increased by TNX in combination with VEGF-A.

MATERIALS AND METHODS

Materials Biochemical reagents were from Wako Pure Chemical Industries (Osaka, Japan). Oligonucleotides were from Hokkaido System Science (Sapporo, Japan). Collagen gel was from Nitta Gelatin Co., Ltd. (Osaka). Dulbecco’s modified Eagle’s medium (DMEM) and Ham’s F12 medium were from Nissui Pharmaceutical Co., Ltd. (Tokyo, Japan). Eagle’s minimal essential medium (MEM) was from Gibco BRL (Tokyo). Fetal bovine serum (FBS) was from Dainippon Pharmaceutical Co., Ltd. (Osaka). The expressed sequence tag clone (Image clone ID, 1690025; GenBank accession no. AI122772) encoding human basic fibroblast growth factor (bFGF) was purchased from Incyte Genomics, Inc. (St. Louis, U.S.A.).

Animals The generation of tenascin-X-deficient (TNX−/−) mice by homologous recombination using embryonic stem (ES) cells has been described. The genetic background of TNX−/− mice is a mixture of C57BL/6, CBA and ICR strains. The control wild-type ICR (TNX+/+) mice were purchased from Sankyo Labo Service Co., Ltd. (Tokyo, Japan). The mice were housed in laminar flow cabinets under specific pathogen-free conditions and used at 8 weeks of age.

* To whom correspondence should be addressed. e-mail: kematsum@pharm.hokudai.ac.jp

© 2001 Pharmaceutical Society of Japan
Preparation of Recombinant Mouse VEGF-B$_{186}$ and Human bFGF Mouse VEGF-B$_{186}$ cDNA clone was described in a previous paper. Human bFGF cDNA was purchased from Incyte Genomics, Inc. (St. Louis, U.S.A.). Each cDNA was subcloned into the GEX-6p vector. Glutathione S-transferase (GST) fusion proteins were prepared as described previously. For preparation of GST-free proteins, GST was cleaved from GST fusion proteins using PreScission Protease according to specifications from the manufacturer (Amersham Pharmacia Biotech, Tokyo). The purity of recombinant mouse VEGF-B$_{186}$ and human bFGF was assessed by 10% SDS-PAGE and quantified by BCA protein assay reagent (Pierce, Rockford, U.S.A.). Its biological activity was confirmed using a collagen gel tissue culture assay.

Collagen Gel Tissue Culture Pregnant female TNX$^{++}$ or TNX$^{--}$ mice 9.5 d post coitus (E9.5) were sacrificed, and their embryos were removed and placed in a sterilized phosphate-buffered saline (PBS). Hearts of E9.5 embryos were collected. Collagen gel (final concentration, 3.0 mg/ml) was polymerized in 24-well culture plates according to the protocol established by the manufacturer (Nitta Gelatin Co. Ltd., Osaka). After the gel had solidified, serum-free culture medium (HAM F12 and DMEM in a 1:1 ratio) containing VEGF-A (100 ng/ml), VEGF-B (100 ng/ml) or bFGF (50 ng/ml) supplemented with penicillin (100 units/ml) and streptomycin (100 mg/ml) was added to the culture plates, and then the hearts of E9.5 embryos from TNX$^{++}$ or TNX$^{--}$ mice were cultured on the collagen gel. Control explants (mock) were incubated in the same culture medium but in the absence of growth factors. The explants were cultured for 5 d, and the media were changed every 48 h.

Proliferation Area Measurement During the incubation period, a sheath of proliferating cells formed around each explant. The proliferating cells were photographed using an Olympus IX70 inverted microscope containing an Olympus SC35 camera. This was followed by measurements using NIH image Macintosh computer program available at http://rsb.info.nih.gov/nih-image/. At least 11 measurements were performed for each explant. Data were presented as the mean with standard deviation (mean±S.D.). Statistical evaluation of the data was carried out using Student’s t-test.

RNA Purification and Reverse Transcription-Polymerase Chain Reaction (RT-PCR) Total RNA from the explants of E9.5 hearts was isolated as previously described. The RT-PCR conditions used were the same as those described in our previous paper. For the detection of mouse VEGFR-1, the following primer sets were used: forward, 5’-ATGAAAGCCTGCAAGCCGCTGAC-3’; reverse, 5’-AACGGATTAGCCTCTTCTCGTATTCTC-3’. For the detection of mouse TNX, the following primer sets were used: forward, 5’-ATGGCAGCTCAGTGCACCCCGTCTA-3’; reverse, 5’-AAGACACCCGCTGAGGTGCTGAGGC-3’.

RESULTS

We have already shown that VEGF-A as well as VEGF-B binds to TNX and that the association of VEGF-B with TNX enhances endothelial cell proliferation by measuring $[^3]$H]-thymidine incorporation. But we have not yet examined whether VEGF-A also increases the proliferation of endothelial cells when combined with TNX.

In this paper, we investigated the effect of TNX on cell proliferation by VEGF-A using embryonic hearts from wild-type (+/+) or TNX-deficient (−/−) mice on a collagen gel, since the collagen gel culture system is believed to mimic or closely reflect the in vivo environment. Recently, we generated TNX$^{--}$ mice by homologous recombination using ES cells. For the condition of presence or absence of TNX, we used wild-type (TNX$^{++}$) or TNX$^{--}$ mice. TNX$^{--}$ mice were fertile and appeared healthy, with no gross abnormalities observed upon macroscopic inspection; however, we expected different responses to VEGF family members with regard to endothelial cell proliferation on the collagen gel between TNX$^{++}$ and TNX$^{--}$ mice.

When E9.5 heart explants from wild-type mice were incubated in the presence of bFGF, the cell proliferation area was significantly greater than in the controls (mock) (Figs. 1A, B). Furthermore, since the control explants (mock) from both wild-type (TNX$^{++}$) and TNX$^{--}$ mice showed the same extents of cell proliferation, indicating that endogenous growth factors secreted from the explants and their receptors are expressed to almost the same extents in both explants. These results indicate that this assay system is useful for the detection of exogenous growth factor activity. During the 5-d incubation period, mesenchymal cells proliferated from the explants onto the collagen substrate and formed a characteristic cobblestone monolayer. This continuous layer extended further from the explants with time due to the proliferative activities. Occasionally, elongated cell also emerged around the explants irrespective of the presence or absence of bFGF, VEGF-A$_{165}$, or VEGF-B$_{186}$ (Fig. 1A). It is known that most cells growing on the surface of collagen gel are epithelial cells forming an embryonic epicardial mesothelium, which differentiates into endothelial cells. By using this culture system, E9.5 heart explants from TNX$^{++}$ or TNX$^{--}$ mice were incubated with VEGF-A$_{165}$ or VEGF-B$_{186}$ for 5 d.

The addition of VEGF-A$_{165}$ to TNX$^{++}$ mouse explants led to a 1.5-fold greater cell proliferation than that in the case of addition of VEGF-A$_{165}$ to TNX$^{--}$ mouse explants, whereas VEGF-B$_{186}$ had no effect on cell proliferation in TNX$^{++}$ mouse explants compared to that in the case of TNX$^{--}$ mouse explants (Figs. 1A, B) (see Discussion). To try to obtain evidence that the embryonic heart explants respond to VEGF-A$_{165}$, leading to the cell proliferation, we examined whether the heart explants express VEGF-R-1 and VEGF-R-2 by RT-PCR. As expected, both VEGF-R-1 and VEGF-R-2 were expressed to the same extents in the heart explants from either TNX$^{++}$ or TNX$^{--}$ mice (Fig. 1C). These results indicate that a VEGF family member, especially VEGF-A, together with TNX enhances proliferation of cells (progenitors of endothelial cells) compared to that in the case of a VEGF family member alone in an E9.5 heart explant cultured system.

DISCUSSION

In a previous study, we demonstrated that TNX in combination with VEGF-B increases endothelial cell proliferation. In the present study, we obtained evidence that the
proliferation of cells (progenitors of endothelial cells) out-growing from E9.5 heart explants is significantly increased by VEGF-A with TNX. These findings suggest that VEGF-A together with TNX is important for the control of endothelial cell proliferation in vivo.

A collagen gel culture system closely reflects the in vivo environment, but the specific cell responses and interactions in vivo and in vitro differ. Ratajska et al. demonstrated that the addition of VEGF-A to rat E14 atrial explants, but not to ventricles or whole hearts, stimulates the appearance of a cord-like structure. In contrast, treatment of rat E12 atrial and ventricular explants with VEGF-A did not induce cell proliferation, nor the appearance of cord-like structures. In this work, we used mouse E9.5 whole heart explants. Cord-like structures were not formed even during prolonged incubation either in the presence or absence of a VEGF family
member. However, elongated cells occasionally appeared around the explants. The reason why a cord-like structure was not detected in the present study may be because of different embryonic stages of the explants and/or different experimental conditions from previous studies.  

As for signaling pathway by VEGF-A, VEGFR-1 and VEGFR-2 generate the mitotic signals, whereas VEGF-B-elicited signals is mediated by VEGFR-1. In this paper, we demonstrated that VEGF-A together with TNX increases the cell proliferation from E9.5 heart explants more efficiently than in the absence of TNX. Although further study is necessary, we speculate that tyrosine phosphorylations of VEGFR-1 and VEGFR-2 are occurred in TNX