Inhibition of Rat Vascular Smooth Muscle Proliferation In Vitro and In Vivo by Bone Morphogenetic Protein-2

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Abstract

Vascular proliferative disorders are characterized by the proliferation of vascular smooth muscle cells (SMCs) and excessive extracellular matrix synthesis. We found that bone morphogenetic protein-2 (BMP-2) inhibited serum-stimulated increases in DNA synthesis and cell number of cultured rat arterial SMCs in a fashion quite different from that in the case of transforming growth factor-β1 (TGF-β1). In addition, TGF-β1 stimulated collagen synthesis in SMCs, whereas BMP-2 did not. In an in vivo rat carotid artery balloon injury model, the adenovirus-mediated transfer of the BMP-2 gene inhibited injury-induced intimal hyperplasia. These results indicate that BMP-2 has the ability to inhibit SMC proliferation without stimulating extracellular matrix synthesis, and suggest the possibility of therapeutic application of BMP-2 for the prevention of vascular proliferative disorders. (J. Clin. Invest. 1997. 100:2824–2832.) Key words: adenovirus • atherosclerosis • balloon injury • smooth muscle cell • transforming growth factor

Introduction

The proliferation of vascular smooth muscle cells (SMCs) is a common feature associated with vascular proliferative disorders such as atherosclerosis and restenosis after balloon angioplasty. A number of growth factors and growth inhibitors are detected within vascular proliferative lesions. Among them, platelet-derived growth factors, basic fibroblast growth factor, and insulin-like growth factor-I are strongly mitogenic for SMCs, and may play critical roles in the in vivo proliferation of SMCs (1). On the other hand, it has also been reported that atherosclerotic lesions once established are capable of regression with extensive therapies (2). Natriuretic peptides, nitric oxide, and interferon-γ were shown to have growth-suppressive actions on SMCs (3, 4). Therefore, the process of SMC proliferation appears to be a dynamic one, in which SMC proliferation is subject to the concomitant effects of these growth regulatory molecules in atherosclerotic lesions.

TGF-β1 is a bifunctional growth regulator of vascular SMCs. It stimulates or inhibits the growth of SMCs, depending upon the target cells, the cell culture conditions, the presence of other growth-regulatory molecules, the order of their addition, and the TGF-β1 concentration used (5, 6). Bone morphogenetic proteins (BMPs), which belong to the TGF-β superfamily, were originally detected as bone-inducing activity (7). In addition to bone induction, BMPs regulate diverse biological processes, and are involved in the development of nearly all organs and tissues. The action of BMPs as potent ventralizing factors by counteracting dorsalizing factors during gastrulation, which was first revealed in Xenopus embryos, is conserved between insect and vertebrate embryonic development (8). BMPs exhibit antiproliferative activities toward certain cell types (9, 10). Recently, BMP-2 and BMP-6 were reported to be detected in human atherosclerotic coronary arteries (11, 12). They are implicated in the atherosclerotic artery calcification through the analogy between calcification and bone formation; however, their biological activities towards vascular cells remain uncertain.

In this study, we demonstrated that BMP-2 has a potent inhibitory action on the growth of cultured rat aortic SMC. In addition, we showed that the adenovirus-mediated transfer of the BMP-2 gene into an injured artery inhibited SMC proliferation in vivo as well as in vitro, using a rat carotid balloon injury model.

Methods

Materials. Recombinant human BMP-2 and BMP-7/osteogenic protein-1 were kindly provided by Dr. T.K. Sampath, and recombinant human TGF-β1 was purchased from R&D Systems, Inc. (Minneapolis, MN). BMP-2 and TGF-β1 were dissolved in 0.1% BSA in 4 mM HCl and stored at −30°C until use. Recombinant human activin A was provided by Dr. Y. Eto and dissolved in DMEM containing 0.1% BSA for storage. [6−3H]Thymidine and [1-3H]proline were obtained from DuPont NEN (Boston, MA). A replication-deficient adenovirus carrying the Escherichia coli β-galactosidase gene, AxCALacZ, and a cosmid, pAxcw, was kindly provided by Dr. I. Saito (13).

Construction of an adenovirus carrying the BMP-2 gene. A replication-deficient adenovirus carrying the BMP-2 gene, AxCABMP2, was created as follows. EcoRI and SalI digested cDNA (1.26 kbp) containing the complete coding region of human BMP-2 (7) was


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blunt-ended and cloned into pCAGGS (14). Then, the DNA fragment (3.5 kbp) containing the CMV-IE enhancer, the chicken β-actin promoter, an intron, the coding region of BMP-2, and rabbit β-globin polyadenylation signal was blunt-end ligated into SwaI site of a cosm id, pAcw. The resulting cosm id, pAXBMP2, was cotransfected to the 293 embryonic cell line with an EcoT221-digested DNA-TPC (from AdSdix) to generate a replication-deficient adenovirus, Ax-CA BM P2 (15, 16). The obtained viral clones were isolated, screened for the BMP-2 insert, and propagated. For in vitro and in vivo experiments, the viruses were purified through a CsCl cushion, titrated, and stored in PBS containing 10% glycerol at −80°C as described (17).

Cell culture. Rat aortic SMCs were prepared from 8-wk-old Wistar rats by the explant method (18), and were grown in DME supplemented with 10% FBS (Life Technologies Inc., Tokyo, Japan), 100 μg/ml of streptomycin, and 100 U/ml of penicillin at 37°C under a 95% air/5% CO2 atmosphere. Cultured SMCs from passages 4–8 were used for all experiments.

Measurement of 3H]thymidine incorporation into DNA of SMCs. Rat aortic SMCs seeded onto flat-bottomed 24-well tissue culture plates (Falcon Labware, Oxnard, CA) were grown until 70–90% confluence and then made quiescent by culturing in DME containing 0.1% BSA for 60 h. Then, the cells were stimulated with 1 or 10% FBS in the presence of various concentrations of BMP-2, BMP-7, TGF-β1, or activin A, followed by pulse-labeling with 37 kBq/ml of [3H]thymidine in 500 μl of the same medium for 2 h. In some experiments, BMP-2 or TGF-β1 was added at various time points after stimulation with FBS. [3H]Thymidine incorporated into the DNA of the cells was determined by the TCA precipitation method involving liquid scintillation counting as described (19), and expressed as decay per minute (dpm) or percentage of control. To determine the rate of [3H]thymidine incorporation into DNA of SMCs transfected with replication-deficient adenoviruses, SMCs seeded onto 24-well plates were grown until 70–90% confluence, exposed to 200 μl of DME with 0.5% FBS containing the indicated doses of the viruses for 2 h, and then serum-deprived for 60 h. [3H]Thymidine incorporated into DNA was determined at 24 h after the stimulation with 1% FBS. All experiments were performed in triplicate and were repeated at least three times.

Measurement of cell number. SMCs seeded onto 24-well plates at 8 × 104 cells per well were maintained in DME containing 10% FBS for 24 h for their attachment to the plates. Then the medium was replaced, and the cells were grown in DME containing 10% FBS and various concentrations of recombinant BMP-2 or TGF-β1 for 3 d. The medium was changed once at 2 d. At the indicated time points, the wells were washed once with PBS, and then the cells were detached by maintenance in 200 μl of PBS containing 0.05% trypsin and 0.53 mM EDTA at 37°C for 5 min. The cells were counted (four times per well) using a hemocytometer, in quadruplicate. Cell viability was assessed by the trypan blue exclusion method.

Measurement of collagen synthesis. SMCs seeded onto 12-well tissue culture plates were grown until 70–90% confluence, and then serum-deprived in an α-complementation of minimum essential medium (α-MEM) for 24 h. The cells were then treated with the indicated concentrations of TGF-β1 or BMP-2 in the α-MEM containing 50 μg ascorbic acid and 50 μg β-aminopropionitrile for 24 h. Then, the cells were labeled with 185 kBq/ml of [3H]proline in 1 ml of the medium for 3 h. Collagen synthesis by the cells was determined as described (19).

Western blot analysis. The conditioned media of cells infected with adenoviruses were collected and centrifuged at 4°C for 5 min at 10,000 g to remove cellular debris. The supernatants were brought to 62.5 mM Tris-HCl, pH 6.8, 10% glycerol, 2% SDS, 5% β-mercaptoethanol, and 0.002% bromphenol blue and were boiled at 92°C for 3 min. Aliquots of the samples (20 μl) were separated on 10% polyacrylamide gels and proteins were transferred to nylon membranes (Immobilon-P; Millipore Corp., Bedford, MA). Immunostaining of BMP-2 was performed by the avidin–biotin complex method using an ABC kit (Vector Laboratories, Burlingame, CA) and anti-human BMP-2 antibodies prepared by immunizing a rabbit with bacterially expressed human BMP-2. Color was developed with a Konica Immunostaining HRP-1000 (Konica Inc., Tokyo, Japan). In some cases, TCA was added to 1-ml aliquots of the conditioned media to a final concentration of 10% which were cleared of cellular debris, followed by gentle agitation at 4°C for 1 h and then centrifugation at 4°C for 15 min at 10,000 g. The precipitates were washed once with 10% TCA and twice with acetone, and then subjected to Western blots.

Adenovirus-mediated gene transfer to balloon-injured arteries. In vivo gene transfer experiments were performed using 10-wk-old SPF Wistar rats by the method reported previously by Guzman et al. with some modifications (20). All procedures were performed under sterile conditions according to the Guide for Animal Experimentation, Faculty of Medicine, University of Tokyo. General anesthesia was performed by the administration of 90 mg/kg of ketamine intraperitoneally and 15 mg/kg of xylazine intramuscularly. Ketamine was supplemented intraperitoneally as necessary. The left carotid artery was exposed and its branches were ligated using 6-0 nylon. After 75 U/kg of heparin had been injected intravenously, a portion of the external carotid artery and a portion of the internal carotid artery were cross-clipped using a microclip (2v-clip; S&I Inc., Neuhausen, Switzerland). A 2F Fogarty embolectomy catheter (Baxter, Irvine, CA) was introduced into the artery via an ~3 mm longitudinal arteriotomy in the external carotid artery. The common carotid arteries were injured by six passes of an embolectomy catheter inflated with 0.2 ml of air. With an additional clip on the proximal portion of the common carotid artery, the arteriotomy was closed with five to seven stitches, using 10-0 nylon, under a magnification of 30, and then blood flow was resumed by removal of the clips.

At 5 d after the injury, the adenovirus was transferred to the injured common carotid artery. After the rats had been reanesthetized, the balloon-injured carotid was exposed, and the proximal portion of the common carotid artery and the internal carotid were cross-clipped. Then, 50–75 μl of virus fluid (1.0 × 1010 plaque-forming units/ml) was delivered into a 1.5-cm length of the common carotid artery through the incision made in the external carotid artery using a 24-gauge catheter to fill the lumen, and the portion proximal to the incision was ligated with 7-0 nylon. The virus solutions were diluted with a medium (OPTI-MEM I; Life Technologies Inc.). After incubation for 40 min, the cross-clips were released and the common carotid artery was reperfused.

5-Bromo-4-chloro-3-indolyl-β-D-galactopyranoside (X-gal) staining. A portion of the left carotid artery, removed 3 d after the adenoviral infection, was incubated in 5 mM K3Fe(CN)6, 5 mM K3Fe(CN)6, and 2 mM MgCl2 containing X-gal (0.5 mg/ml) at 37°C for 6 h after fixation in 2% formaldehyde and 0.2% glutaraldehyde in PBS at 4°C for 1 h. Then, an X-gal–stained artery was fixed in 4% paraformaldehyde at 4°C for 6–12 h and paraffin-embedded. A section of the paraffin-embedded specimen was stained with hematoxylin and eosin and then analyzed.

Measurement of β-galactosidase activity. Portions of balloon-injured left carotid arteries were removed at 3 d after transfection of an adenovirus, AxCALacZ. After briefly washing in PBS, total β-galactosidase activity of the arteries was measured using a β-galactosidase enzyme assay system according to the manufacturer’s instructions (Promega, Madison, WI). Protein concentrations were determined using a micro BCA protein assay reagent kit (Pierce Chemical Co., Rockford, IL).

Measurement of the intimal and medial cross-sectional areas of a balloon-injured carotid artery. A portion of the left common carotid artery, transfected or untransfected with adenoviruses at 5 d after the balloon injury, was harvested at 14 d after the injury. The artery was fixed in PBS containing 4% paraformaldehyde for 6–12 h at 4°C and then paraffin-embedded. Four round cross-sections per 1.5-cm length of artery specimens stained with hematoxylin and eosin were photographed, and the cross-sectional areas of the intimal and medial regions of the sections were measured using an image analyzing soft-
ware package (NIH Image). The intimal and medial cross-sectional areas and intimal to medial (I/M) area ratio of each injured artery were determined by averaging the values for four sections to evaluate the intimal mass in each artery. Some paraffin-embedded sections were stained with Elastica van Gieson to clearly define the extracellular matrix in the injury-induced neointimal region.

Statistical analysis. All data are expressed as means±SEM. The unpaired Student’s t test was used for comparison of two groups using Statview-J 4.02 (BrainPower, Calabasas, CA). Statistical significance was accepted at P < 0.05.

Results

Effects of BMP-2 and TGF-β1 on the growth of rat aortic SMCs in vitro. To characterize the action of BMP-2 on the growth of cultured SMCs, the effect of BMP-2 on DNA synthesis in SMCs was evaluated and compared with that of TGF-β1. The rate of DNA synthesis (estimated as [3H]thymidine incorporation into DNA) in SMCs that had been made quiescent by serum starvation started to increase at 14–16 h after stimulation with 1% FBS, reached a peak at 24 h, and then decreased (Fig. 1A). TGF-β1 at 100 pM inhibited the serum-stimulated DNA synthesis at 19 and 24 h by 57 and 30%, respectively (P < 0.01). After that, however, the rate of DNA synthesis continued to increase and reached a peak at 32 h, which was comparable with the peak value for serum-stimulated cells without TGF-β1 treatment. Thus, TGF-β1 apparently delayed the peak of serum-stimulated DNA synthesis under these conditions. In contrast, BMP-2 at as low as 0.3 pM strongly inhibited the serum-stimulated DNA synthesis at every time point over 48 h (67% at 19 h and 47% at 24 h, respectively; P < 0.01 at every time point). The inhibitory effect of BMP-2 on DNA synthesis increased dose-dependently up to 0.3 pM, with a threshold concentration of 0.03 pM (Fig. 1B). However, paradoxically, the inhibitory effect of BMP-2 at higher concentrations (3–30 pM) became less pronounced. Inhibitory effect of BMP-2 was observed again at 100 pM.

We next examined the effect of BMP-2 added at various time points after serum addition. As shown in Fig. 1C, even when BMP-2 was added at 20 h after serum addition, i.e., after cells had entered the S phase, BMP-2 still inhibited DNA synthesis by 30% at 24 h (P < 0.01). In sharp contrast, TGF-β1 no longer suppressed the serum-stimulated DNA synthesis when it was added just before (at 13 h) or after (at 20 h) S phase entry (P > 0.05), which is consistent with several previous reports (21–23).

The effect of BMP-2 on the growth of SMCs was evaluated by culturing SMCs in DME containing 10% FBS, since the doubling time of SMCs in 10% FBS was shorter than in 1% FBS.
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FBS. BMP-2 also inhibited the DNA synthesis in SMCs at 24 h after stimulation with 10% FBS (P < 0.01 at 100 pM, Fig. 2 A). The cell number of SMCs grown with 100 pM BMP-2 was decreased, as compared with that of SMCs grown without BMP-2, by 68% on day 2 (P < 0.01) and by 59% on day 3 (P < 0.05), respectively (Fig. 2 B). On the other hand, the increase in the number of the cells treated with TGF-β1 was decreased only by 24%, as compared with the control culture (P < 0.05). Thus, consistent with the inhibition of DNA synthesis, BMP-2 also increased the number of cell number of SMCs grown in the presence of serum. The dose–effect relationship between BMP-2 and inhibition of DNA synthesis stimulated with 10% FBS was dose-dependent up to 3 nM unlike the case with 1% FBS (Fig. 2 C). We also examined the effect of BMP-7 and activin A, members of the TGF-β superfamily, on DNA synthesis stimulated with 10% FBS, however, they did not significantly inhibit DNA synthesis stimulated with 10% FBS.

Effects of BMP-2 and TGF-β1 on collagen synthesis by SMCs. Excessive extracellular matrix synthesis is one of the features of atherosclerosis, and TGF-β1 has the ability to stimulate the synthesis of extracellular matrix components such as collagen and proteoglycan by cultured SMCs (24, 25). Therefore, we examined whether or not BMP-2 has a stimulatory action on collagen synthesis by SMCs similar to that of TGF-β1. TGF-β1 stimulated collagen synthesis by 17% at 10 pM (P < 0.05) and by 40% at 100 pM (P < 0.01). In contrast, BMP-2 at up to 1 nM did not stimulate collagen synthesis by SMCs (Fig. 3).

Effect of adenovirus-mediated transfer of the BMP-2 gene on DNA synthesis of cultured SMCs. To examine the effect of BMP-2 gene transfer into SMCs on their proliferation, we constructed a replication-deficient adenovirus carrying the BMP-2 gene under the control of the CAG promoter, AxCABMP2. The rate of DNA synthesis was evaluated in cultured SMCs which had been transfected with a virus, either AxCABMP2 or AxCALacZ. SMCs that had been infected with AxCABMP2 at 3–24 mois showed a lower rate of DNA synthesis in response to serum stimulation, as compared with mock-infected control cells (lowest at 12 moi of AxCABMP2, 21% of control, P < 0.001) (Fig. 4). In contrast to this, infection of cells with AxCALacZ at the same moi did not inhibit the serum-stimulated DNA synthesis. Inhibition of DNA synthesis in SMCs by AxCABMP2 was reduced by > 24 moi, and the dose–effect relationship between AxCABMP2 and DNA synthesis gave a concave curve.

Figure 2. Inhibition of the increase in cell number together with DNA synthesis in rat aortic SMCs by BMP-2. (A) Serum-starved SMCs were stimulated with 10% FBS in the presence of BMP-2 at various concentrations. [3H]Thymidine incorporation was determined at 24 h after the addition of FBS and presented as a percentage of the serum-stimulated control level. The control value on stimulation with 10% FBS was 49,000 ± 1,300 dpm. (B) SMCs seeded onto 24-well plates at 8 × 10⁴ cells per well were maintained in DME containing 10% FBS for 24 h for their attachment to the plates. Then the medium was replaced, and the cells were grown in 10% FBS alone (open circles) or with either 100 pM BMP-2 (closed circles) or 100 pM TGF-β1 (open triangles) for 3 d. The total cell numbers were determined using a hemocytometer, by harvesting cells at 0, 48, and 72 h after medium replacement and cell viability was > 95% at the end of the experiment. (C) Serum-starved SMCs were stimulated with 10% FBS in the presence of BMP-2 (closed circles), BMP-7 (open circles), or activin A (open triangles) at various concentrations. [3H]Thymidine incorporation was determined at 24 h after the addition of FBS and presented as a percentage of the serum-stimulated control level. The results are representative of three separate experiments.
Production of the BMP-2 protein was examined by Western blot analysis. Anti–BMP-2 antibodies used in this study clearly detected 5 ng recombinant BMP-2, but not 30 ng TGF-β1, activin A, or BMP-7 (data not shown). This suggests that the antibody is highly specific to BMP-2. BMP-2 produced by SMCs is expected to be secreted into the extracellular space, since the BMP-2 gene possesses a signal peptide sequence. A 20-kD protein that reacted with anti–BMP-2 antibody was detected in conditioned medium of SMCs infected with AxCABMP2 at 200 moi, but not in medium of mock-infected cells or cells infected with AxCALacZ (Fig. 5A). Its mobility on SDS-PAGE slightly differed from those of recombinant human BMP-2 purified from the Chinese hamster ovary (CHO) cell line stably producing human BMP-2, which consists of 18- and 22-kD subspecies; 22-kD being a major band (Fig. 5B). When CHO cells were infected with AxCABMP2 and conditioned media were analyzed, we detected a 22-kD band, which was similar to the 22-kD subspecies of recombinant BMP-2 (data not shown). These data indicate that the cells infected with AxCABMP2 might produce BMP-2 proteins with different mobility on SDS-PAGE, depending on the type of the infected cells.

To delineate the amount of BMP-2 secreted by SMCs infected with AxCABMP2, conditioned media of infected cells were treated with TCA and the precipitates were evaluated by Western blot analysis. A 20-kD band was not detected in the conditioned media of SMCs infected with AxCABMP2 at 48 moi or less. Therefore, the concentration of BMP-2 in the conditioned media of SMC infected with AxCABMP2, at mois at which DNA synthesis in infected cells was inhibited, was < 100 pM, since 100 pM BMP-2 in 1 ml of cultured medium should have been detected with this procedure.

Inhibition of balloon injury–induced intimal hyperplasia by AxCABMP2. We next tried to determine whether or not BMP-2 can inhibit in vivo vascular smooth muscle proliferation as well. To this end, we used a rat carotid balloon injury model and examined the effect of the adenovirus-mediated transfer of the BMP-2 gene upon injury-induced intimal SMC hyperplasia. We first determined the efficacy of expression of genes transferred by the adenovirus vector using AxCALacZ. Similar to the previous report (20), nearly 50% or more of the neointimal cells in a balloon-injured carotid artery expressed β-galactosidase, as estimated by X-gal staining of injured arteries transfected with AxCALacZ (Fig. 6A). Tissue β-galactosidase activity was > 50-fold higher in injured carotid

**Figure 3.** Collagen synthesis by rat aortic SMCs is upregulated by TGF-β1 but not by BMP-2. SMCs were treated with the indicated concentrations of BMP-2 (closed circles) or TGF-β1 (open circles), and then labeled with [3H]proline for 3 h. Collagen synthesis by cells was determined and presented as percent difference from control value. The control value for untreated SMCs was 56,300 ± 2,900 dpm.

**Figure 4.** Inhibition of DNA synthesis in vitro on adenovirus-mediated transfer of the BMP-2 gene. Rat aortic SMCs seeded onto a 24-well plate were exposed to 200 μl of DME containing 0.5% FBS and the indicated moi of either AxCABMP2 (closed circles) or AxCALacZ (open circles) for 2 h, and were serum-deprived for 60 h. [3H]Thymidine incorporation into DNA was determined at 24 h after the stimulation with 1% FBS and presented as a percentage of the serum-stimulated control level.
arteries exposed to the virus at 5 d after injury than immediately after the injury (data not shown).

We introduced either AxCABMP2 or AxCALacZ at 10⁶ plaque-forming units/ml into injured rat left carotid arteries at 5 d after balloon injury. 14 d after balloon injury, the injured arteries were harvested to measure the cross-sectional areas of the intima and media of each artery and the I/M area ratio (Fig. 6, B and C). The administration of AxCABMP2 reduced the intimal mass of an injured artery by 41%, as compared with the administration of AxCALacZ (P < 0.01). However, the medial mass was similar in the two groups. As a result, the I/M area ratio was substantially reduced on the administration of AxCABMP2 (P < 0.002). Thus, an adenovirus encoding the BMP-2 gene can effectively limit intimal hyperplasia after balloon injury.

Discussion

This paper is the first to document that BMP-2 is a potent inhibitor of SMC proliferation both in vitro and in vivo. BMP-2 was growth-inhibitory at relatively low concentrations (0.03–1 pM), without inducing delayed DNA synthesis. These concentrations are lower than those required for other biological activities of BMPs, such as the induction of osteoblastic differentiation (26). However, it has been reported that BMP-4, a very closely related homologue of BMP-2, induced the migration of human blood monocytes at far lower concentrations than in our study (i.e. 0.3–3 fM) (27). Since BMPs use multiple receptor systems depending on the cell types and their various biological effects (28), the difference in effective doses of BMPs may be due to receptors expressed on the cells and sensitivities of the assays used in these studies. In our study, the dose-effect relationship of DNA synthesis in 10% FBS was different from that in 1% FBS. It has been reported that the growth inhibition of human embryonic carcinoma cells by BMP-7 is affected by the concentration of FBS in the culture medium (9). It might be that serum contains a counteracting activity towards BMPs, as in the case of follistatin, a binding protein of activin (29). The concentration of BMP-2 required for maximal growth inhibition was far lower than those of previously known growth inhibitors of SMC such as natriuretic peptides and 8-bromo-cyclic guanosine monophosphate (4).

TGF-β1 induces G₁ arrest in some cells such as Mv1Lu cells through its action towards these cells in the late G₁ phase (30). The growth-inhibitory action of BMP-2 on SMCs is different from that of TGF-β1 at least in the time order of their actions, since BMP-2 was active when added later, as shown in Fig. 1 C. Consistent with the suppressive activity of BMP-2 toward serum-stimulated DNA synthesis, BMP-2 inhibited the increase in cell number in the presence of serum. It has been reported that activin A, another member of TGF-β superfamily, has a growth-stimulatory effect on the proliferation of SMCs (31). In our study, however, activin A did not affect DNA synthesis in SMCs, probably due to the different experimental conditions. It is noteworthy that BMP-7 partially mimicked the action of BMP-2 under low serum concentration, since BMP-7 at 100 pM or more inhibited DNA synthesis stimulated with 1% FBS (Nakaoka, T., K. Miyazono, and T. Fujita, unpublished text).
unpublished observation), although BMP-7 did not inhibit DNA synthesis under high serum concentration. It has been reported that Smad1 acts downstream as a transcriptional activator in signaling by members of the BMP subfamily (32). It should be determined whether or not Smad pathways are involved in the BMP-2–induced inhibition of S phase entry and S phase progression.

Furthermore importantly, BMP-2 up to 1 nM did not stimulate collagen synthesis at all. Thus, the action of BMP toward cultured SMCs is different from that of TGF-β1 in the effect on collagen synthesis as well as the mode of growth inhibition. Previous studies implicated TGF-β1 in the increased synthesis of extracellular matrix component in atherosclerotic lesions and in restenotic lesions after balloon angioplasty (33, 34), although TGF-β1 inhibits the serum-stimulated growth of cultured SMCs. The ability of BMP-2 to act as a potent inhibitor of SMC proliferation without stimulating the synthesis of collagen, a major extracellular matrix protein, is considered to be favorable for in vivo application to therapy for vascular proliferative disorders.

Rat aortic SMCs infected with AxCABMP2 produced BMP-2 with different mobility on SDS-PAGE from those of recombinant human BMP-2 purified from the CHO cell line stably producing human BMP-2. It has been reported that mature recombinant BMP-2 protein produced in CHO cells contains high mannose and complex N-linked oligosaccharide and
that 18- and 22-kD subspecies differ by proteolytic processing at their amino termini (35). Moreover, the CHO cells and the SMCs infected with the same adenovirus, AxCABMP2, produced different species of BMP-2 proteins as shown in Fig. 5. Thus, the difference in the mobility in SDS-PAGE might be due to different amino-terminal processing or carbohydrate structure in the mature BMP-2 proteins, which may be dependent on the transfected cells.

BMP-2 was originally identified by its bone formation activity (7). However, histological examination did not reveal any osseous or cartilaginous change in injured arterial tissue infected with AxCABMP2. The expression of a protein through adenovirus-mediated gene transfer is transient in nature. Local expression of BMP-2 for a longer time may predispose an arterial wall to an osseous change. Considering BMP-2 is now recognized as a widely acting differentiation-inducing factor (8), its role as a differentiation factor in vascular proliferative lesions might deserve attention from quite a different viewpoint. It is known that neointimal SMCs show a synthetic phenotype different from the contractile phenotype of medial SMCs (1). Therefore, it would be interesting to determine whether and how the expression of BMP-2 affects the smooth muscle phenotype, especially in the neointima.

Since the proliferation of SMCs is an essential feature of vascular proliferative disorders, considerable effort has been made to develop a therapeutic strategy to effectively suppress SMC proliferation. Previous attempts to suppress SMC proliferation have included the administration of antibodies against growth factors, antagonists for receptors, and antisense oligonucleotides directed to transcription factors and cell cycle regulatory molecules required for cell proliferation (36–38). More recent studies demonstrated that the adenovirus-mediated transfer of a therapeutic gene into the vascular wall could be a promising treatment strategy. For example, it was reported that considerable inhibition of intimal hyperplasia was observed on the administration of an adenovirus containing herpes simplex virus thymidine kinase gene with the prodrug ganciclovir (39–41). An adenovirus containing the gene of the thymbin inhibitor, hirudin (42), and an adenovirus containing the gene of a constitutively active form of the retinoblastoma gene product (43). In the case of the BMP-2 gene, neighboring untransfected cells in the vascular wall may also be affected by diffusible BMP-2, a “bystander effect,” since BMP-2 is a secreted protein. The ability of BMP-2 as a growth inhibitor described above makes this molecule a suitable candidate therapeutic agent for the prevention of vascular proliferative disorders.

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