Improving the therapeutic potential of endostatin by fusing it with the BAX BH3 death domain

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Endostatin (ES) inhibits angiogenesis, reducing tumor growth in animal models. However, it has low therapeutic effect in human clinical trials. BAX is a member of the BCL-2 family of proteins; its proapoptotic (BH3) domain interacts with other members of the family in the cytoplasm, to induce apoptosis. Here, we fused the BAX BH3 domain with murine ES, to enhance ES potency. Endothelial cells specifically internalize the fusion protein ES-BAX. The presence of the BAX domain enhances endothelial cell death by apoptosis by 1.8-fold and diminishes microvessel outgrowth in the rat aortic ring assay by 6.5-fold. Daily injections of 15 μg of ES-BAX/g in tumor-bearing mice reduce tumor weight by 86.9% as compared with ES-treated animals. Co-immunoprecipitation assays confirmed that ES-BAX interacts with members of the BCL-2 family. Also, ES interacts with BCL-2, BCL-XL, and BAK in endothelial cell lysates, suggesting a potential new mechanism for the apoptosis induction by ES. The superiority of the ES-BAX antiangiogenic effect indicates that this fusion protein could be a promising therapeutic alternative to treat cancer.

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Abbreviations: ES, endostatin; BH3, BCL-2 homology 3; BCL-2, B-cell lymphoma 2; BCL-XL, B-cell lymphoma-extra large; BAX, BCL-2-associated X-protein; BAK, BCL-2 antagonist/killer; ES-BAX, ES fused to the BAX BH3 domain; ES-BAK, ES fused to the BAX BH3 domain; ES-BAX-ES, the ES BAX α-helix is substituted by ES α-helix; bFGF, basic fibroblast growth factor
Impaired apoptosis is a critical step in tumor development. Enhanced levels of the prosurvival BCL-2 family members or, alternatively, the loss or inactivation of the pro-death relatives frequently occur in cancer. Therefore, scientists have designed strategies to induce downstream apoptotic events that could overcome the inhibition of tumor cells apoptosis by either delivering proapoptotic BH3 peptides or using compounds that function as cell permeable, small molecular mimics of the BH3 domain. However, there is concern about the therapeutic use of proapoptotic BH3 or its mimetics because of the lack of specificity to tumor cells, possibly prompting to greater toxicity to normal cells. Inducing an imbalance in favor of cell death by raising the levels of the proapoptotic BH3 peptide, is an interesting strategy, especially in cells with normal levels of the antiapoptotic BCL-2 proteins, which is the case of cells of tumor vasculature.

In the present study, we produced three chimerical recombinant proteins based on the core of the ES fused with the BH3 domains of the proapoptotic proteins BAK and BAX as a means to target these proteins. Such proteins display enhanced proapoptotic properties toward the tumor endothelium, avoiding damage to normal tissues. In addition, we determined if ES and ES-BAX interact with members of the BCL-2 family in endothelial cell lysates.

Results

ES, ES-BAX, ES-BAK, and ES-BAX-ES: expression, refolding, and purification. To target tumoral neovascu- lature, we engineered recombinant fusion proteins containing the full-length ES attached to the BAX or BAK death-prompting domains. We named the mutant proteins ES-BAX (ES fused to the BAX BH3 domain) and ES-BAK (ES fused to the BAK BH3 domain), which consisted of ES, two Gly residues (to give flexibility to the peptide), and BAX or BAK amphipathic α helices containing 16 residues mapped to the BH3 domain. In the mutant protein ES-BAX-ES (the ES α-helix is substituted by ES α-helix), we maintained the first two residues of the ES α-helix and substituted the sixteen following residues in the BAX BH3 domain. With this configuration, we retained the Cys residue in ES position C33, because Cys is the natural sixth residue of the BAX peptide (Figure 1a). Figure 1b illustrates the hypothetical model of ES and the hybrid proteins. In ES-BAX, the BAX
BH3 \(\alpha\)-helix appears as an addition at the N-terminal end of the fusion protein; in ES-BAX-ES, the BAX \(\alpha\)-helix substitutes the ES \(\alpha\)-helix. In both configurations, the disulfide bonds are theoretically maintained to mimic the ES.

We expressed high levels of the recombinant proteins ES, ES-BAX, ES-BAK, and ES-BAX-ES as inclusion bodies in *Escherichia coli*, and then we refolded and purified them. ES-BAX-ES presents a molecular weight similar to that of ES (21.5 kDa), while ES-BAX (23.5 kDa) and ES-BAK (23.4 kDa) have slightly higher molecular weight due to the 18 extra amino acids that they contain (Figure 1c). Compared with the other three recombinant proteins, ES-BAX-ES is less stable, which may have contributed to the lower yield observed during its expression and purification (data not shown).

**Endothelial cells internalize ES and the fusion proteins.**

Endothelial cells specifically internalize ES.\(^2\)\(^,\)\(^4\) To determine whether the fusion proteins behave similarly, we incubated C-PAE cells with the biotinylated proteins for 2 h at 37 °C. We viewed cell-associated ES, ES-BAX, ES-BAK, and ES-BAX-ES with Alexa 488-conjugated streptavidin (Molecular Probes, Eugene, OR, USA) in fixed and permeabilized cell preparations. According to Figure 2a, C-PAE cells but not NIH/3T3 fibroblasts internalize the four recombinant proteins,

![Figure 2](image-url)

**Figure 2** Internalization of ES, ES-BAX, ES-BAK, and ES-BAX-ES by endothelial cells. (a) C-PAE or NIH 3T3 cells were incubated at 37 °C for 2 h with 10 μg/ml of the biotinylated proteins: ES, ES-BAX, ES-BAK, or ES-BAX-ES. The cells were then fixed, permeabilized, and incubated with Alexa 488 conjugated with streptavidin. The nucleuses were stained with DAPI. The cells were analyzed under a fluorescence microscope. Scale bar: 10 μm. (b) Western blot of endothelial C-PAE cells incubated with 10 μg/ml of the indicated proteins at 37 °C and detected by anti-ES antibody. Control (+): ES; control (−): lysate of cells incubated with ES that were immediately processed for western blot (t = 0 min).
indicating that the endothelial cells have the ability to specifically internalize the hybrids, similarly to wild-type ES. As described elsewhere,27 ES is internalized through endocytic vesicles and further transported into the nuclei. Apparently, nuclear translocation is more efficient for the variants ES-BAX and ES-BAK as compared with ES (Figure 2a). The presence of granules that possibly indicate endocytic vesicles within the cells treated for 1 h with ES, while the proteins ES-BAX and ES-BAK are more homogeneously distributed within the cytoplasm (Supplementary Figure 1) also suggest that the routes for ES-BAX and ES-BAK within the cells are different from that of the wild-type protein. Internalization of ES and fusion proteins were also observed for HUV-EC-C human endothelial cells (not shown).

We cultivated endothelial cells (C-PAE) in the presence of each protein for 30, 90, or 180 min at 37°C, followed by extensive washings and total cell lysis. We detected recombinant proteins, which represent bound and internalized components, by western blot using the anti-ES antibody. Although the cell-associated recombinant proteins are different at early time points, at 180 min the hybrid proteins exist at levels similar to or higher than those of ES (Figure 2b). The differences in the internalization kinetics may suggest that ES and its hybrid variants use different endocytic routes.

**The presence of the BAX BH3 domain enhances ES potency.** ES induces apoptosis in endothelial cells.28 To verify whether the presence of the BAK or BAX proapoptotic domains enhances the ES ability to induce endothelial cell death, we performed flow cytometry analysis of the propidium iodide-stained HUV-EC-C endothelial cells. Cells treated with the fusion proteins containing the BAX domain present higher degree of apoptosis as compared with cells treated with the same amount of ES. ES-BAX and ES-BAX-ES induce cell death at rates of 63.9% and 61.0%, respectively, although ES only induces cell death at a rate of 35.1% (Figure 3a). Cells treated with ES-BAK behave more similarly to cells treated with ES, at a rate of 44.4%. Importantly, all the recombinant proteins induce caspase-3 activation, as shown in Figure 3b.

To verify whether the presence of BAK or BAX proapoptotic domains enhances the ability of ES to induce endothelial cell viability loss, we incubated C-PAE cells with recombinant proteins and assessed cell viability by the MTS assay (Promega, Madison, WI, USA). A reduction in cell viability was observed for treatment of endothelial cells with 6.25 μg/ml ES and ES-BAX in a dose-response curve and the decrease in cell viability was stabilized with treatment of the cells with 25 μg/ml (Figure 3c). Incubation of the cells with 25 μg/ml ES and ES-BAK diminishes cell viability to the same extent: 76.1% as compared with the untreated control (100% viability). The presence of the BAX domain significantly raises ES activity—the number of viable cells decreases to 50.0% and 65.8% after treatment with ES-BAX and ES-BAX-ES, respectively (Figure 3d). Endothelial cell viability in response to treatment with ES and ES-BAX was also determined for different time points. The cells presented a decline of viability in response to treatment with ES-BAX for 3 h, while the decrease of viability was observed only after 6 h of treatment with ES (Figure 3e). Inhibition of microvessel outgrowth of the ex vivo rat aortic ring in Matrigel (Becton-Dickinson, San Diego, CA, USA) with dose-dependent effects helps to evaluate the biological activity of murine ES.29 We used the aortic ring assay to examine the effect of the recombinant fusion proteins investigated herein. As expected, compared with untreated rings, ES abate outgrowths from the aortic rings. In agreement with our previous results, ES-BAX and ES-BAX-ES induce a more potent antiproliferative effect on microvessels. ES-BAX almost completely prevents the outgrowth of rat aortic endothelial cells (Figure 4a). Quantitative image analysis confirmed that ES-BAX and ES-BAX-ES display increased biological activity as compared with ES (Figure 4b). However, as expected, ES-BAX presents effects similar to those of ES.

**ES-BAX potently suppresses in vivo tumor growth.** Finally, we investigated the effect of in vivo therapy with ES-BAX on a mouse model of allograft adenocarcinoma renal tumor. The dosage of administered ES (15 μg/g mice) was chosen based on the literature30 to induce incomplete tumor growth inhibition. As expected, compared with the untreated group, tumor growth diminishes in the group treated with ES. Moreover, consistent with the results described above, mice treated with ES-BAX experience more pronounced decrease in tumor volume (Figure 5a). In three of the five mice treated with ES-BAX, tumor growth either stabilizes or decreases, which does not occur in any of the animals belonging to the other groups. The average tumor weights were 2.63 ± 1.19 g (untreated), 1.34 ± 0.39 g (ES), and 0.175 ± 0.082 g (ES-BAX; Figure 5b and c), which correspond to a remarkable 93.3% and 86.9% reduction in tumor weight for the ES-BAX-treated group as compared with the untreated control group and the ES-treated group, respectively. To exclude the possibility that ES or ES-BAX can directly affect the renal carcinoma cells, we performed an in vitro viability assay (MTS). We did not detect any loss of cell viability upon incubation with ES or ES-BAX (not shown), as opposed to the results obtained with the endothelial cells (Figure 3c). Hence, as expected, ES-BAX exerts an indirect effect on the tumors.

**ES, ES-BAX, and ES-BAK can bind to cellular BCL-2 and BCL-XL.** The ability of proapoptotic BH3 peptides to induce cell death depends on their capacity to interact with the proapoptotic members of the BCL-2 family of proteins.31 Therefore, we examined whether ES and the hybrid proteins ES-BAX and ES-BAK can bind to the protective proteins BCL-2 and BCL-XL and to the proapoptotic proteins BAX and BAK from an endothelial cell lysate. In agreement with our expectations, the hybrid proteins ES-BAX and ES-BAK co-immunoprecipitate along with BCL-2, BCL-XL, BAX, and BAK present in the lysate of endothelial cells (Figure 6). The binding between ES-BAX, and ES-BAX with the cellular proteins BCL-2 and BCL-XL were confirmed by reciprocal assay. Surprisingly, ES also co-immunoprecipitates with the two antiapoptotic proteins and with BAK (Figure 6). To the best of our knowledge, this is the first observation that ES can directly interact with members of the BCL-2 family. The ES binding to proteins belonging to the BCL-2 family was unexpected and very interesting, possibly opening new perspectives for understanding the ES mechanism of action. Probably the domain of ES responsible for the binding to BAK/BCL-2/BCL-XL diverges from the domain of ES-BAX.
Discussion

In the present study, we circumvented the difficulties related to the lack of specificity of inducers of tumor death by targeting endothelial cells of the tumor vasculature; in other words, we indirectly targeted tumors. The BH3 domains of proapoptotic proteins were targeted to tumor endothelia by the presence of the ES domain, a protein that was previously shown to be able to be specifically internalized by angiogenic endothelial cells. Here, we verified that the endothelial cells specifically internalize the three fusion proteins ES-BAX, ES-BAK, and ES-BAX-ES at least as efficiently as they internalize ES.
Another relevant finding is that unrelated cells do not internalize the hybrid proteins. Enhanced levels of antiapoptotic or decreased levels of proapoptotic Bcl-2 members in tumor cells are common obstacles to cancer chemotherapy. To overcome difficulties pertaining to the imbalance in the tumor cell apoptotic pathway, we therapeutically targeted cells that present normal levels of the BCL-2 relatives, the endothelial cells. The presence of the BAX BH3 domain significantly improves the in vitro activity of the ES recombinant proteins. The ex vivo aorta rings assay reveals a similar effect, showing the antiangiogenic activity of this protein. Most importantly, ES-BAX substantially reduces tumor volumes in mice bearing renal tumors: by 88% and 71% as compared with untreated mice and ES-treated mice, respectively.

ES presents increased activity when the BAX fragment is present, either as a C-terminal fragment or by replacement of the ES native α-helix. A noticeable biochemical difference exists between the effectors of the apoptotic pathways BAK and BAX in healthy cells: BAX is largely cytosolic or loosely associates with mitochondria, whereas BAK is an integral membrane protein on the cytosolic face of the mitochondrion and the endoplasmic reticulum. However, BAX and BAK physiologically appear to be largely redundant in function. The presence of the BAK BH3 domain in the ES-BAK construct does not enhance the cytotoxic effect, indicating that the presence of a peptide with 16 amino acids per se does not suffice to improve ES activity. These results reinforce that the BAX BH3 apoptotic pattern accounts for the enhanced activity.
activity of the hybrid proteins ES-BAX and ES-BAX-ES. However, further studies are necessary to understand the different effects of the BAX and BAK BH3 domains on ES toxicity to endothelial cells.

BAX and BAK present the hydrophobic side chain of the BH3 helix, which is involved in binding to the antiapoptotic family members, facing towards the core of the protein. Upon engagement, which occurs by triggering the BH3 helix, the unstructured loop between α-helices 1 and 2 is displaced, the carboxy-terminal helix 9 is mobilized for membrane translocation, and the BAX BH3 domain is exposed. Consequently, the death signal propagates through an auto-activating interaction that involves the triggering site of inactive BAX monomers. An alternative hypothesis is that exposed proapoptotic BH3 domains of BAX, BAK, or BH3-only proteins can bind to the hydrophobic surface groove of the prosurvival family members and promote apoptosis by releasing the active proapoptotic proteins from their antiapoptotic counterparts. Apoptosis occurs when BAX or BAK suffers from major conformational changes, followed by homo-oligomerization. These alterations can generate the ‘apoptotic pore’ in the mitochondrial outer membrane and release cytochrome c. Obviously, ES-BAX does not participate in BAX oligomerization because it lacks the domains involved in oligomerization or the membrane insertion that exists in the integral BAX protein. However, possibly the proapoptotic BH3 domain exposed in the hybrid protein ES-BAX interacts with intracellular BAX directly and activates it. An alternative is that the proapoptotic domain interacts with the prosurvival proteins, releasing activated BAX. The released BAX then undergoes homo-oligomerization, to form the mitochondrial outer membrane pores that induce cytochrome c release and endothelial cells apoptosis.

It was described that BAX BH3 peptide interfere with BAX/BCL-2 and BAX/BCL-X₅ heterodimerization in vitro and promote cytochrome c release from mitochondria isolated from Jurkat, HL-60, U937, and PC-3 cells. The BAX BH3 peptide also potently interfere with both BAX/BCL-2 and BAX/BCL-X₅ interactions in vitro. This interference correlates with the ability of this peptide to overcome BCL-2 overexpression, as shown by the induction of cytochrome c release from mitochondria of Jurkat T-leukemic cells overexpressing BCL-2. Apart from the length and amino acid composition of the BH3 peptides, their ability to form a stable helical structure is important to their high binding affinity, and there seems to exist a direct correlation between the tendency of BH3 peptides to form a helix and their affinity for the antiapoptotic proteins. However, the BH3 domain peptides

Figure 5 Effect of ES and ES-BAX on tumor growth. BALB/c mice (n = 5 per group) received s.c. injections with 2 × 10⁶ Renca cells. When tumors reached 100 mm³, mice received daily s.c. injections of buffer or 15 μg of either ES or ES-BAX/g of mice. On day 11 after beginning of the treatment of the first set of mice, the animals were euthanized, and the tumors were removed, weighed, and photographed. This procedure was repeated until the treatment of all of the animals was concluded. (a) Kinetics of tumor growth during the 10-day treatment. (b) Photographs from representative tumors withdrawn on day 11 after the beginning of the treatment. (c) Average tumor weight on day 11. Error bars: mean ± S.E.M. Statistical significance was determined using a Student’s t-test (two-tailed), **P < 0.005; *** P < 0.0001. This assay was performed twice.
of proapoptotic proteins are mostly unstructured in aqueous solution\textsuperscript{22,37,38} whereas they adopt a well-defined helical conformation in experimentally determined complex structures\textsuperscript{17,39–41} Therefore, the interaction between the BAX BH3 peptide in the hydrophobic groove of the antiapoptotic members of the BCL-2 family is feasible, even in the case whereupon the ES-BAX BH3 peptide does not present helical conformation in solution.

Many mechanisms of action have been proposed for ES. However, as far as we know, this is the first time that the direct interaction between ES and BCL-2, BCL-X\textsubscript{L}, and BAK has been demonstrated (Figure 6). Further studies are necessary to determine whether the apoptotic effect of wt-ES on proliferating endothelial cells refers to its interaction with members of the BCL-2 family.

BH3-only proteins function either by binding to their prosurvival relatives, which prevents them from inhibiting BAX and BAK, or by directly activating the apoptosis effectors. Although some BH3-only members engage only some of the prosurvival members, BIM and PUMA are potent inducers of cell death because they can engage all the prosurvival members\textsuperscript{38} and directly activate BAX.\textsuperscript{18} It is likely that the BH3 domain of these proteins can serve as efficient effectors of apoptosis. Thus, further studies on hybrid proteins based on the ES core attached to the BH3 domain of the potent killer members of the BH3-only family are relevant and ongoing in our laboratory.

To our knowledge, this is the first time that the BH3 death domain has been examined as a fusion partner to target and induce apoptosis in non-tumor cells. Because BCL-2 overexpression is a common cause of multidrug resistance, our primary goal has been to improve the ES-mediated inhibition of tumor neovasculature. Here, we have provided definitive evidence of enhanced endothelial cell apoptosis and in vivo tumor regression by adding the BAX BH3 domain at the ES C-terminus.

Materials and Methods
Construction of plasmids, expression refolding, and purification of proteins. Hybrid proteins were generated by site-directed mutagenesis using the ES construct pETK1H (ATCC No. 63404), which contains the sequence that codifies for the residues HAHHHHHHHM, followed by the sequence that codifies for murine ES, as template. Coding sequences for two Gly residues, followed by the minimal sequence of the BH3 domain of the proapoptotic peptides BAX or BAK were inserted between the codon for the last amino acid of the ES gene and the stop codon. Site-specific mutagenesis was induced as previously described,\textsuperscript{42} by respectively using the following primers: 5\textsuperscript{-}TGACGGAGGCTCG ATTCTAAGAGTCCAGTTCGTCACCGATACGTTTCAGGCATTCAGACAGTTTTTTACGGATACCGCATGCCT-3\textsuperscript{3}; 5\textsuperscript{-}TAAGAGTCCAGTTCGTCACCGATACGTTTCAGGCATTCAGACAGTTTTTTACGGATACCGCATGCCT-3\textsuperscript{3} was used for the mutagenesis procedure, substituting the DNA coding sequences of the \(\alpha\)-helix of ES by the \(\alpha\)-helix of the BAX BH3 domain. DNA sequencing was performed to confirm the expected mutations. All cDNAs were cloned into the pET28 (Mercur Millipore, Darmstadt, Germany) to generate the expression plasmids pET-ES, pET-ES-BAX, pET-ES-BAK, and pET-ES-BAX-ES.

Vectors encoding ES and the hybrid proteins were transformed into E. coli BL21 (DE3) cells and cultured in rich culture medium (2x-HKSII)\textsuperscript{43} supplemented with 50 \(\mu\)g/ml kanamycin at 37 \(^\circ\)C for 16 h, the cells were harvested by centrifugation at 12 000 \(g\) for 10 min, at 4 \(^\circ\)C. The ES inclusion bodies and the three hybrids produced by ES-BAX, ES-BAK, and ES-BAX-ES expressed as inclusion bodies (Figure 6).

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Internalization assays. Biotinylation of the proteins was achieved using sulfo-NHS-LC-Biotin (EZ-Link sulfo-NHS-LC biotinylation kit, Pierce, Rockford, IL, USA). C-PAE or NIH 3T3 cells (ATCC, CCL-209 and CRL-1658, respectively; 5 \(\times\) 10\(^5\)) were cultured in glass coverslips in EMEM medium (Life Technologies Corporation, Grand Island, NY, USA) supplemented with 2% FBS for 24 h. After 24 h of serum deprivation, the cells were incubated for 12 h in the same medium supplemented with 5 ng/ml biotin-labeled ES, ES-BAX, ES-BAK, or ES-BAX-ES. The cells were washed twice with phosphate-buffered saline (PBS), permeabilized by 6-min incubation in ice-cold methanol, washed twice with PBS, and then blocked with 1% BSA in PBS for 40 min. The cells were then incubated with Alexa
488-conjugated streptavidin (Life Technologies Corporation) for 45 min and washed with PBS five times. Vectashield mounting medium containing DAPI (4’, 6-diamidino-2-phenylindole; Vector Laboratories, Burlingame, CA, USA) was applied to glass slides and coverslips were placed on them. The fields were photographed in a fluorescence microscope.

C-PAE cells (5 × 10^4) were cultured in 12-well plates in EMEM medium supplemented with 20% fetal bovine serum (FBS), 50 μU/ml penicillin, 50 μg/ml streptomycin, and 2 mM l-glutamine. After 24 h, the medium was changed to EMEM supplemented with 2% FBS, 10 ng/ml bFGF (RD systems, Minneapolis, MN, USA), and 10 μg/ml ES, ES-BAX, ES-BAK, or ES-BAX-ES at 37 °C. After different periods of time, the cells were washed three times in PBS, resuspended in SDS-PAGE sample buffer, and heated for 5 min at 95 °C. Protein samples were separated on 12% SDS-polyacrylamide gels under reducing conditions and electroblotted to nitrocellulose membrane (GE Healthcare, Raepagatan, Uppsala, Sweden). The membranes were incubated with rabbit anti-ES polyclonal antibody (AB1880, diluted 1:500, Merck Millipore), followed by incubation with horseradish peroxidase (HRP) conjugated to secondary antibody (Merck Millipore, diluted 1:5000). Immunodetection was carried out with the chemiluminescence reagent (Immobilon, Millck Millipore).

In vitro apoptosis detection. Apoptosis was quantified by analysis of the DNA content using flow cytometry. Briefly, 1.5 × 10^5 C-PAE cells were plated in six-well tissue culture plates in EMEM containing 20% FBS. After cell adhesion, the medium was replaced with medium containing 2% FBS, 10 ng/ml bFGF, and 20 μg/ml ES, ES-BAX, ES-BAK, or ES-BAX-ES. After 16 h of incubation, cells were trypsinized, resuspended in 0.3 ml of a hypotonic solution (50 mg/ml propidium iodide, 0.1% sodium citrate, and 0.1% Triton X-100, Sigma), and kept at 4 °C in the dark until they were examined. The fluorescence of individual nuclei was measured with a FACScalibur flow cytometer (Becton-Dickinson).

Analysis of caspase-3 activation. Caspase-3 activity was measured. Briefly, 4 × 10^4 C-PAE cells were maintained in EMEM medium containing 2% FBS for 24 h. The medium was replaced with fresh medium containing 2% FBS, 10 ng/ml bFGF, and 20 μg/ml ES, ES-BAX, ES-BAK, or ES-BAX-ES. After 24 h, all the cells (adherent or not) were collected and resuspended in SDS-PAGE sample buffer; the proteins were boiled for 5 min, for denaturation. The proteins from the cells (adherent or not) were collected and resuspended in SDS-PAGE sample buffer, and heated for 5 min at 95 °C. After different periods of time, the cells were washed three times in PBS, resuspended in SDS-PAGE sample buffer, and heated for 5 min at 95 °C. Protein samples were separated on 12% SDS-polyacrylamide gels under reducing conditions and electroblotted to nitrocellulose membrane (GE Healthcare, San Diego, CA, USA). The membrane was incubated with anti caspase-3 antibody (Santa Cruz Biotechnology, Santa Cruz, CA, USA) for 16 h, washed, incubated with a secondary antibody conjugated to HRP for 1 h, and washed again. Immunoreactive bands were visualized using an enhanced chemiluminescence system (Immobilon, Merck Millipore).

Cell Viability. C-PAE viability was evaluated by the MTS [3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxyphenyl)-2-(4-sulfophenyl)-2H-tetrazolium] assay (Promega). Briefly, 5 × 10^4 C-PAE cells were plated in each well of a 96-well tissue culture plate in EMEM supplemented with 20% FBS in a final volume of 0.1 ml. After 24 h, the medium was replaced with fresh medium containing 2% FBS, 10 ng/ml bFGF, and 3.12–50 μg/ml ES, ES-BAX, ES-BAK, or ES-BAX-ES and incubated for 45 min to 72 h. Cell viability was measured by addition of 20 μl of an MTS (0.2%), PMS (0.02%), phenazine methosulfate, 20:1 solution and incubation for 2 h at 37 °C in a humidified 5% CO2 incubator. The microplates were read in a spectrophotometer at a wavelength of 495 nm. The absorbance of the control group (without added protein) at 495 nm was set as 100% cell viability. Each sample was analyzed in triplicate.

Rat aortic ring assay. Each well of a tissue culture-grade plate (48 well) was covered with 150 μl of cold Matrigel and allowed 30–45 min to form a gel at 37 °C, 5% CO2. The thoracic aorta was excised from a 9-week-old male Sprague–Dawley rat (Charles River, Wilmington, MA, USA). The aorta was sectioned into ~1-mm long cross-sections, rinsed with EBM-2 medium (Lonza, Walkerills, MD, USA) eight times, and placed on the Matrigel-coated wells, covered with an additional volume of 150 μl Matrigel, and allowed 30–45 min to form a gel in an incubator at 37 °C, 5% CO2. The rings were cultured for 24 h in 1 ml of EBM-2 supplemented with 10 μg/ml gentamycin. After incubation, the medium was removed and replaced with 1 ml of EBM-2 supplemented with 2% FBS, gentamycin, and 10 μg/ml ES, ES-BAX, ES-BAK, or ES-BAX-ES, in triplicate. The medium was replaced every day. The aorta cultures were washed three times in PBS, the medium was replaced with RPMI medium (Life Technologies Corporation) containing 5 mg/ml 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT, Promega), and the culture was incubated for 2 h. The supernatant was discarded, and digital images of the aortic rings were generated. The microvessel outgrowth area was quantified using the ImageJ (http://image.nih.gov/ji) software. Images were analyzed by manually encoding the outgrowth area and computing the square pixels. This assay was repeated three times.

Animals and tumor induction. Male (24–27 g) BALB/c mice (6–7 weeks old, IPEN animal facility, IPEN, São Paulo, Brazil) were used. They were acclimated, caged in groups of five in a barrier care facility, and fed animal chow and water ad libitum. The manipulation of these animals before or during the experiments met the Principles of Laboratory Animal Care (NIH Pub. No. 86–23, revised in 1985), the “Principles of Ethics in Animal Experimentation” (Brazilian College of Animal Experimentation – COBEA), and Project No. 124/CEUA/IPEN/SP Animal Research Ethics Committee authorization. We used a murine renal carcinoma cell line Renca (American Type Culture Collection, Manassas, VA, USA), a line of spontaneous origin derived from a BALB/C mouse kindly donated by Dr. Isaiah Fidler, D.V.M., PhD. (University of Texas MD Anderson Cancer Center, USA). The left hind flank of the mice was subcutaneously (s.c.) injected with 2 × 10^5 Renca cells in 0.1 ml of sterile PBS. Treatment started when each tumor reached 100 mm^3. To this end, the mice received daily s.c. injections of 15 μg of ES or ES-BAX/g of mice into the right flank for 10 days. Tumor dimensions were measured with an electronic caliper, and volumes were calculated as follows: tumor volume = length × width × 0.52. This assay was repeated twice.

Immunoprecipitation. HUV-EC-C cell lysates (10^7 cells) were prepared by incubation with 1 ml of lysis buffer (50 mM Tris-HCl at pH 8.0, 0.15 M NaCl, and 0.5% Triton X-100) containing 10% protease inhibitor cocktail (Sigma) for 30 min at 4 °C. The lysates were centrifuged, and the supernatants were incubated with ES, ES-BAX, or ES-BAX-ES. After 24 h, the proteins were transferred onto nitrocellulose membranes (GE Healthcare, San Diego, CA, USA). The membrane was incubated with anti-ES polyclonal antibody (Santa Cruz Biotechnology, Santa Cruz, CA, USA) for 16 h, washed, incubated with a secondary antibody conjugated to HRP for 1 h, and washed again. Immunoreactive bands were visualized using an enhanced chemiluminescence system (Immobilon, Merck Millipore).

Conflict of Interest

The authors declare no conflict of interest.

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