

Angiogenin and endothelial cells

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Anglogenin is a potent blood vessel-inducing polypeptide that has a unique ribonucleolytic activity. Initially identified from tumor cells conditioned medium through its ability to induce neovascularization in the chick embryo chorioallantoic membrane assay [1], angiogenin has also been purified from normal plasma [2] which suggested that it might be involved in endothelium homeostasis. Angiogenin is highly homologous to pancreatic ribonuclease A with 68% amino acid sequence homology and conserved essential active site residues [3]. However, angiogenin is inactive in standard ribonuclease assays but has a ribonucleolytic specificity toward ribosomal and transfer RNAs [4]. This enzymic activity might be relevant to the process of angiogenesis since a functional active site seems necessary for the expression of its angiogenic property [5], however, it appears not to be sufficient [6].

Interactions of angiogenin with endothelial cells

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Using recombinant human angiogenin produced by Rhône-Poulenc Santé and shown to be angiogenic, in vivo, both in the chick chorioaliantoic membrane and the rabbit cornea [7,8], we have demonstrated the presence of high affinity binding sites on endothelial cells from calf pulmonary artery (CPAE), bovine aorta, cornea and adrenal cortex capillary that could not be detected on Chinese hamster lung fibroblasts [9]. Receptors have also been reported on bovine brain capillary endothelial cells [10]. The interactions on CPAE cells displayed an apparent dissociation constant in the nanomolar range and involved about 200,000 angiogenin molecules per cell [9]. This number, that varies with the cell density, does not necessarily represent a number of receptors. However, it includes receptors since angiogenin, added to cultured cells, triggers off a series of intracellular events in endothelial cells such as a transient increase in 1,2-diacylglycerol [11] and activation of phospholipase A2 [12], and in vascular smooth muscle cells, activation of phospholipase C and a rapid cholesterol esterification [13]. Besides, angiogenin might modulate mitogenic stimuli [10,14].

Angiogenin has also been shown to interact with low-affinity/high capacity specific binding sites on CPAE cells as well as on extracellular

matrix. This second family of interactions, with an apparent dissociation constant of 0.2 μM, involved millions of molecules [9].

An angiogenin-hinding protein has been recently identified on boying

An angiogenin-binding protein has been recently identified on bovine endothelial cells as a dissociable cell-surface component of 42-KDa released by treating the cells with heparan sulfate [15].

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Inhibition of angiogenin properties by Placental Ribonuclease inhibitor

Placental ribonuclease inhibitor (PRI) is a tight binding inhibitor of both the ribonucleolytic and angiogenic activities of angiogenin [16] with an inhibition constant value, Ki, in the femtomolar range [17]. This inhibitor also abolishes angiogenin binding to CPAE cells [9]. However, PRI is an intracellular regulatory protein and although immunoreactivity has been detected in normal human plasma [18], there is not yet any experimental evidence for angiogenin/inhibitor circulating complexes.

Effects of copper metal ions on angiogenin binding to endothelial cells and to ribonuclease inhibitor

Divalent copper, a modulator of angiogenesis [19], has been shown to induce a severalfold increase in specific cell-bound angiogenin and to reverse PRI-induced inhibition of angiogenin binding to CPAE cells [8,9]. Furthermore, divalent copper metal ions inhibits angiogenin ribonucleolytic activity towards tRNA substrate [20].

Discussion

The variety of copper interactions with angiogenin properties is indeed puzzling. It might reflect a tight regulation of angiogenin functions necessary to balance its angiogenic potency with its high intravascular concentration. Copper also interacts with heparin and the angiogenic growth factor FGF [21], and appears, from other reported data, to modulate angiogenesis somehow [19]. At this point, it is worth noting that copper is required for other fundamental events such as central nervous system development, bone and connective tissue biosynthesis, electron transport and antioxidant protection where it

acts as a prosthetic group [22]. Because of angiogenin peculiarities, studies on its mechanism of action should lead the way to a better knowledge of the biochemical events involved in angiogenesis.

In summary, modulation of angiogenin functions might involve interactions with cell-surface receptors, binding protein and extracellular matrix components, tight-binding inhibition of both its ribonucleolytic activity and cell binding by ribonuclease inhibitor under the overall influence of divalent copper, a modulator of angiogenesis.

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