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abolished in the presence of nonradiolabeled angiotensin I. The specific binding of [125I] angiotensin I to the endothelial cells was dose-dependent. The binding was saturable, reversible, and of high affinity. The binding sites were localized to the cell surface and were internalized after binding to the cell surface. The K_d value of the binding was 5.3 nmol/l. ACE activity, as indicated by fluorogenic substrate, was detectable in the medium of pulmonary endothelial cells, but not of fibroblasts. These results suggest that the pulmonary microvascular endothelial cells are distinct from pulmonary fibroblasts in