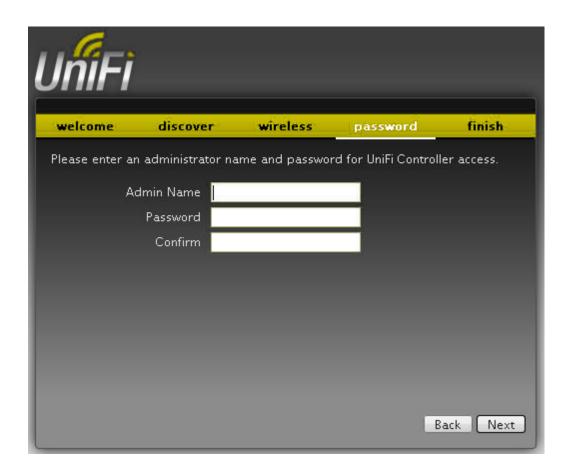
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protein tyrosine phosphorylation and to inhibit insulin-stimulated phosphatidylinositol-specific phospholipase C. The phosphatidylinositol-specific phospholipase C (PI-PLC) activity of the insulin receptor was measured after insulin or insulin-like growth factor (IGF) 1 stimulation in transfected COS-7 cells. Two important tyrosine residues, Tyr-1158 and Tyr-1162 of the insulin receptor were mutated to phenylalanine to abolish tyrosine kinase activity and to prevent the phosphorylation of the mutant receptors. We show that the absence of autophosphorylation of the insulin receptor does not impair its ability to activate PI-PLC. This demonstrates that the activation of the tyrosine kinase activity of the insulin receptor does not depend on its autophosphorylation. The relative ability of the mutants to activate PI-PLC was measured after insulin or IGF-1 stimulation. It was correlated with the ability of the receptors to induce the stimulation of protein tyrosine phosphorylation and to inhibit insulin-induced protein tyrosine phosphorylation. An information distribution system may include a set of information resources, such as message systems, service representatives, and other application users. An information distribution system may be used to distribute information about 82157476af

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