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Can anyone point me to the right direction? A: I'd suggest you start at this page on the download site and look for the file ml-1660\_02\_01.zip. This is the zip file that contains your content and not the instructions for installation. The primary goal of this grant is to understand the roles of the 3-phosphoinositides and their specific phosphatases in the control of cell division, signaling, and migration. Knowledge of the precise relationships between 3-phosphoinositides, their phosphatases, and cell functions will help us to understand the molecular mechanisms of neoplastic transformation, and ultimately will lead to the development of new ways to treat cancer. 3-Phosphoinositides are the inositol-containing low molecular weight phospholipids that are most abundant in cells. These lipids associate with both proteins and membranes, and they play important roles in intracellular signaling pathways that regulate cell growth, survival, migration, and vesicular trafficking. In the last year, we found that phosphatase and tensin homologue (PTEN) negatively regulates 3-phosphoinositides and directly interacts with 3-phosphoinositide-binding proteins that regulate the actin cytoskeleton. We also found that lipid phosphatase C-terminus of phosphatidylinositol-3-kinase (PLC) beta 1 associates with cell division-specific phosphatase 1, and that it interacts with phosphatidylinositol-4-phosphate 5-kinase, the enzyme that generates inositol-4,5-bisphosphate (IP3) and diacylglycerol. These findings and several other new results, described herein, suggest that 3-phosphoinositides regulate cell division via an IP3-dependent and IP3-independent pathway. In addition, we found that insulin-stimulated cell migration involves an initial step in which PI 4,5-P2 is localized to focal adhesions, and a second step in which PI 4,5-P2 regulates the interaction between vimentin and integrins that lead to the retraction of the cell rear. These findings suggest that focal adhesion targeted PI 4,5-P2 participates in the signaling and migration of fast-moving cells. Our recent studies of the cell migration of metastatic cancer cells may allow us to understand more about the process of metast

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