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The centrosome is the major microtubule organizing center in mammalian cells. It functions in the formation of a bipolar spindle that segregates genetic material during cell division. Defects in the centrosome have been shown to be directly related to a number of human developmental diseases, most prominently microcephaly and ciliopathies. Despite the importance of centrosome assembly and function, the underlying mechanisms remain largely unknown. Recently, however, some of the building blocks for the centrosome have begun to emerge. These include not only major proteins that are localized to the centriole but also proteins that are hypothesized to connect the centriole to its surrounding pericentriolar material. Understanding of the biochemistry of centrosome assembly is a necessary first step toward deciphering these poorly understood connections. In the current project, we plan to focus on the relationship of the centrosome to the major microtubule organizing center, the Golgi apparatus. In particular, we are interested in understanding how these two organelles are brought together during the formation of the primary cilium and in understanding the mechanisms that regulate ciliogenesis. For this, we propose to: 1) use in vitro reconstitution assays to understand how centrosome proteins control the assembly of the centrosome, 2) identify the cis-and trans-acting factors that regulate ciliogenesis, and 3

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