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replication resulting in either no or low levels of infection. Our goal was to characterize the cell types that initiate and support the early immune response in these mice. We show that the inflammatory cells, including macrophages, neutrophils, dendritic cells, and T lymphocytes, are recruited to the draining lymph nodes within 24 h of reovirus inoculation. Antigen-presenting cells with dendritic morphology (macrophages and dendritic cells) are the major cellular component of the primary infection sites within 3 days postinoculation. Based on their expression of Class II major histocompatibility complex (MHC), dendritic cells are shown to be antigen-presenting cells (APCs) with an important role in reovirus-induced immunity. In contrast, macrophages are not involved in early immune responses. Reoviruses (arboviruses) have evolved mechanisms to sequester antigen-presenting cells from draining lymph nodes, whereas it is shown that an abundant group of cells expressing antigenic peptides (dendritic cells, macrophages, and B lymphocytes) remain in the primary reovirus infection site.