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squamous cell origin is dependent on CCL2 and CCR2. The recruitment of macrophages is an important feature in mammary tumor progression. Here, we demonstrate that macrophage recruitment can also be induced within different histological subtypes of mammary carcinoma. When using the MMTV-PyMT mouse model of mammary carcinoma, we show that tumor-bearing animals have higher numbers of macrophages in their tumors as compared to wild-type animals. In addition, administration of the CCR2 receptor inhibitor CCX872 during the early phases of PyMT mammary carcinoma development strongly reduces macrophage infiltration and inhibits tumor growth and angiogenesis *in vivo*. Likewise, treatment of MMTV-PyMT tumors grown in CCR2-deficient mice with CCX872 significantly reduces the number of tumor-infiltrating macrophages. A similar pattern of results is observed in two independent mouse models of squamous cell carcinoma, i.e., the DMBA-7