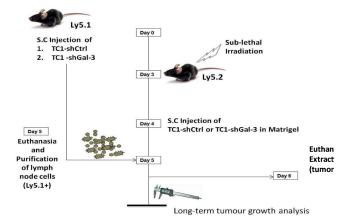
Figure S2



Adoptive transfer method: Lymph node cells from the mice donors (C57BL/6; Ly5.1) were adoptive transferred the correct neo-vascularization of nascent tumor. In all cases where adoptive transfer of lymphocytes was performed or -shGal-3 cells were injected subcutaneously in C57BL/6-Ly5.1 strain donor mice with PBS (without Matrigel). On and cell suspensions were performed as follow. Challenged lymphoid cells were obtained from lymph nodes (super mesenteric lymph nodes), and immediately transferred to cold RPMI medium supplemented with 10% calf serum (organs obtained in R10F medium were homogenized with a concentration of antibiotics twice to the standard concentration of 5 min. The live cells obtained chamber were counted in the Neubauer in the presence of trypan blue, cells was prepared in 100 μ l RPMI medium without supplement. The effect of adoptive transfer of challenged lymp tumorigenic capacity of TC1-shCtrl or -shGal-3 cells in C57BL/6 wild type (Ly5.2) host mice. To decrease endogenou adoptive transfer, the animals used as receptors received sublethal irradiation (1 Gy, radiation source: cesium 137; lymphoid cells transfer. Twenty-four hours before irradiation, antibiotics (Enrofloxacin 85 mg / kg) were given to dri TC1-shCtrl or -shGal-3 cells were injected subcutaneously with Matrigel (100 μ l). The next day, 4x106 challenge lym retro-ocularly injected under anesthesia. Tumor growth was monitored periodically and the pathophysiological par observe exponential tumor growth and the tumor duplication time) were determined.