

### This week in techniques

Approach	Summary	Licensing status	Publication and contact information
<b>Drug platforms</b>			
Depletion of tumor necrosis factor receptor superfamily member 9 (TNFRSF9; 4-1BB; CD137) <sup>+</sup> /CD4 <sup>+</sup> T cells to improve adoptive cell therapy treatment of cancer	<p>Studies in mice and in patient samples suggest depleting CD137<sup>+</sup>/CD4<sup>+</sup> T cells could help increase the efficacy of adoptive cell therapy. In mice immunized with a whole-cell cancer vaccine, levels of CD137<sup>+</sup>/CD4<sup>+</sup> T<sub>reg</sub> cells were higher than those in unimmunized mice. In a mouse model of lymphoma, transplantation of CD137<sup>-</sup>/CD4<sup>+</sup> T cells protected mice from lymphoma cell challenge compared with transplantation of a mixed population of CD137<sup>+</sup>/CD4<sup>+</sup> and CD137<sup>-</sup>/CD4<sup>+</sup> T cells. Next steps include testing antibody-mediated depletion of CD137<sup>+</sup> T cells in transplantation experiments.</p> <p>BMS-663513, an agonistic mAb against CD137 from Bristol-Myers Squibb Co., is in Phase I/II testing to treat cancer.</p> <p><b>SciBX 5(5); doi:10.1038/scibx.2012.139</b>            Published online Feb. 2, 2012</p>	Unpatented; available for licensing	<p>Goldstein, M.J. <i>et al. Cancer Res.</i>; published online Jan. 9, 2012; doi:10.1158/0008-5472.CAN-11-3375</p> <p><b>Contact:</b> Ronald Levy, Stanford University School of Medicine, Stanford, Calif.            e-mail: <a href="mailto:levy@stanford.edu">levy@stanford.edu</a></p>