

### This week in techniques

Approach	Summary	Licensing status	Publication and contact information
<b>Drug platforms</b>			
Nonglycosylated Fc $\gamma$ -receptor I (CD64; FCGR1)-selective IgG antibodies	<p>Nonglycosylated FCGR1-selective IgG antibodies could be useful for treating cancer and other diseases via mechanisms distinct from standard glycosylated mAbs. <i>In vitro</i>, a nonglycosylated variant of Herceptin trastuzumab increased dendritic cell (DC)-mediated killing of a human breast carcinoma cell line compared with no antibody. In contrast, Herceptin and a glycosylated trastuzumab variant did not induce DC-mediated killing of the cancer cells. Next steps could include assessing the ability of nonglycosylated antibodies to induce adaptive immune responses and evaluating their efficacy in a mouse model of human tumors.</p> <p>Herceptin, a humanized mAb against HER2 (ERBB2; neu) from Roche's Genentech Inc. unit, is marketed to treat breast cancer and is under EMEA review for stomach cancer.</p> <p><b>SciBX 3(1); doi:10.1038/scibx.2010.32</b>  <b>Published online Jan. 7, 2010</b></p>	Two patent applications filed; available for licensing from Clayton Biotechnologies Inc.	<p>Jung, S.T. <i>et al. Proc. Natl. Acad. Sci. USA</i>; published online Dec. 14, 2009; doi:10.1073/pnas.0908590107</p> <p><b>Contact:</b> George Georgiou, University of Texas, Austin, Texas            e-mail: <a href="mailto:gg@che.utexas.edu">gg@che.utexas.edu</a></p>