

This week in techniques

Approach	Summary	Licensing status	Publication and contact information
Disease models			
<p>Mouse model for prostate cancer driven by v-ets erythroblastosis virus E26 oncogene homolog (ERG) translocations</p>	<p>Mice with prostate-specific Erg expression could be used as a model to study prostate cancer pathogenesis. Translocations of ETS transcription factors including ERG are common in prostate cancer but have been difficult to model in mice. To develop a mouse model to study the role of ERG in prostate cancer, prostate-specific <i>Erg</i> overexpression was combined with homozygous loss of <i>Pten</i> (<i>Mmac1; Tep1</i>), which led to the development of invasive prostate adenocarcinomas in 80% of the animals at 6 months. In <i>Pten</i>-deficient prostates, <i>Erg</i> overexpression increased the expression of androgen receptor-regulated genes compared with wild-type <i>Erg</i> expression. Next steps could include using the model to identify new therapeutic targets.</p> <p>SciBX 6(29); doi:10.1038/scibx.2013.770 Published online Aug. 1, 2013</p>	<p>Patent and licensing status unavailable</p>	<p>Chen, Y. <i>et al. Nat. Med.</i>; published online June 30, 2013; doi:10.1038/nm.3216 Contact: Charles L. Sawyers, Memorial Sloan-Kettering Cancer Center, New York, N.Y. e-mail: sawyers@mskcc.org Contact: Yu Chen, same affiliation as above e-mail: cheny1@mskcc.org</p>