

## This week in therapeutics

Indication	Target/marker/pathway	Summary	Licensing status	Publication and contact information
<b>Neurology</b>				
Alzheimer's disease (AD)	Inositol 1,4,5-triphosphate receptor (ITPR; IP3R)	<p>Mouse studies suggest antagonizing IP3R could be useful for treating AD. In two mouse models of AD, an engineered loss-of-function mutation that lowered <i>Ip3r</i> expression by 50% improved hippocampal function and decreased pathological intracellular calcium levels compared with wild-type <i>Ip3r</i> expression. In one of the models, lower <i>Ip3r</i> expression decreased levels of the AD markers <math>\beta</math>-amyloid (A<math>\beta</math>) and hyperphosphorylated microtubule-associated protein-<math>\tau</math> (MAPT; tau; FTDP-17) compared with those seen in wild-type controls. Next steps could include examining the effect of <i>Ip3r</i> mutations on neurodegeneration and evaluating IP3R antagonists in AD models.</p> <p><b>SciBX 7(23); doi:10.1038/scibx.2014.682</b>  <b>Published online June 12, 2014</b></p>	Patent and licensing status undisclosed	<p>Shilling, D. <i>et al. J. Neurosci.</i>; published online May 14, 2014;            doi:10.1523/JNEUROSCI.5441-13.2014  <b>Contact:</b> J. Kevin Foskett, University of Pennsylvania, Philadelphia, Pa.            e-mail:  <a href="mailto:foskett@mail.med.upenn.edu">foskett@mail.med.upenn.edu</a></p>