

### This week in therapeutics

Indication	Target/marker/pathway	Summary	Licensing status	Publication and contact information
<b>Neurology</b>				
Cognitive dysfunction	NMDAR	<p><i>In vitro</i> and mouse studies suggest 24(S) hydroxycholesterol (24(S)-HC) could help treat cognitive dysfunction. In screening studies using patch-clamp recordings on mouse hippocampal neurons, 24(S)-HC was selected from a panel of sterol-based compounds as a potent submicromolar potentiator of NMDAR-mediated currents. In mouse hippocampal slices, 24(S)-HC produced long-term potentiation (LTP) from a subthreshold tetanus stimulus and reversed NMDAR antagonist-induced LTP suppression. In behavioral memory tests in mice, a 24(S)-HC analog reversed NMDAR agonist suppression of memory-dependent activity. Next steps are being performed by Sage Therapeutics Inc. and include optimizing 24(S)-HC analogs in models of schizophrenia.</p> <p><b>SciBX 6(48); doi:10.1038/scibx.2013.1395</b>  <b>Published online Dec. 19, 2013</b></p>	<p>Patent application filed by Sage Therapeutics; licensing information available from Sage Therapeutics</p> <p><b>Contact:</b> Jeff Jonas, Sage Therapeutics Inc., Cambridge, Mass.  e-mail: <a href="mailto:jeff@sagerx.com">jeff@sagerx.com</a></p>	<p>Paul, S.M. <i>et al. J. Neurosci.</i>; published online Oct. 30, 2013; doi:10.1523/JNEUROSCI.2619-13.2013</p> <p><b>Contact:</b> Steven M. Paul, Weill Cornell Medical College, New York, N.Y.  e-mail: <a href="mailto:smpaulmd@med.cornell.edu">smpaulmd@med.cornell.edu</a></p>