

This week in therapeutics

Indication	Target/marker/pathway	Summary	Licensing status	Publication and contact information
Cancer				
Acute lymphocytic leukemia (ALL)	Deoxycytidine kinase (DCK)	<i>In vitro</i> and mouse studies suggest simultaneously inhibiting two deoxycytidine triphosphate (dCTP) synthesis pathways could help treat ALL. In human T cell ALL lines, thymidine (dT) decreased dCTP synthesis through the <i>de novo</i> pathway but promoted dCTP synthesis through the nucleotide salvage pathway. In human T cell ALL lines, the combination of dT plus a DCK inhibitor that blocks the nucleotide salvage pathway resulted in increased apoptosis and replication stress compared with either compound alone. In mice, the combination decreased tumor burden in human T cell ALL xenografts and leukemic burden in systemic B cell ALL xenografts compared with either treatment alone. Next steps include assessing safety and toxicology.	Patent application filed for the DCK inhibitors; unlicensed	Nathanson, D.A. <i>et al. J. Exp. Med.</i> ; published online Feb. 24, 2014; doi:10.1084/jem.20131738 Contact: Caius G. Radu, University of California, Los Angeles, Calif. e-mail: cradu@mednet.ucla.edu
<p>SciBX 7(13); doi:10.1038/scibx.2014.364 Published online April 3, 2014</p>				