

### This week in therapeutics

Indication	Target/marker/pathway	Summary	Licensing status	Publication and contact information
<b>Cancer</b>				
Cancer	Casitas B cell lymphoma-b (CBL-B); TYRO3 protein tyrosine kinase (TYRO3; SKY); AXL receptor tyrosine kinase (AXL; UFO); c-Mer proto-oncogene tyrosine kinase (MERTK)	<p><i>In vitro</i> and mouse studies suggest TYRO3, AXL and MERTK receptor (TAM receptor) inhibitors could help prevent metastasis by activating NK cells. In mice, deletion of <i>Cbl-b</i> increased NK cell antitumor activity compared with no deletion.</p> <p><i>In vitro</i> assays identified the TAM receptors as substrates for CBL-B's E3 ubiquitin ligase activity. In multiple mouse models of metastasis, a small molecule inhibitor of the TAM receptors decreased metastasis compared with vehicle. Next steps include developing an siRNA-based clinical candidate to target <i>CBL-B</i>.</p> <p>Rigel Pharmaceuticals Inc. and partner BerGenBio A/S have a small molecule inhibitor of AXL in Phase I or earlier testing to treat various cancers.</p> <p>Tolero Pharmaceuticals Inc. and partner Oribase Pharma have AXL inhibitors in preclinical development to treat cancer.</p> <p>Qurient Co. Ltd. has the AXL inhibitor Q-4 in preclinical development for cancer.</p> <p><b>SciBX 7(10); doi:10.1038/scibx.2014.281</b>  <b>Published online March 13, 2014</b></p>	Patent filed by Lead Discovery Center GmbH and Max Planck Institute of Biochemistry covering a chemical series of TAM receptor inhibitors; licensed to Qurient	Paolino, M. <i>et al. Nature</i> ; published online Feb. 19, 2014; doi:10.1038/nature12998 <b>Contact:</b> Josef M. Penninger, Institute of Molecular Biotechnology of the Austrian Academy of Sciences, Vienna, Austria e-mail: <a href="mailto:josef.penninger@imba.oeaw.ac.at">josef.penninger@imba.oeaw.ac.at</a>