

### This week in therapeutics

| Indication                       | Target/marker/pathway                                | Summary  | Licensing status                         | Publication and contact information  |
|----------------------------------|--|--|--|--|
| <b>Various</b>                   |  |  |  |  |
| Autoimmune disease; inflammation | Phosphoinositide 3-kinase- $\gamma$ (PI3K $\gamma$ ) | <p><i>In vitro</i>, cell culture and mouse studies suggest highly selective inhibitors of PI3K<math>\gamma</math> could help treat inflammation and autoimmune diseases. <i>In vitro</i> screening and optimization led to the development of CZC24832, a small molecule inhibitor that is at least 100 times more selective for PI3K<math>\gamma</math> over the other PI3K isoforms. In cell culture, CZC24832 decreased differentiation of proinflammatory T helper type 17 (Th17) cells compared with vehicle. In a mouse model of collagen-induced arthritis, CZC24832 lowered bone and cartilage destruction by 53% compared with vehicle. Next steps at Cellzome AG include using the <i>in vitro</i> screening platform to identify additional highly selective kinase inhibitors. Cellzome said it discontinued CZC24832 for strategic reasons.</p> <p><b>SciBX 5(20); doi:10.1038/scibx.2012.533</b><br/> <b>Published online May 17, 2012</b></p> | Unpatented; licensing status undisclosed | <p>Bergamini, G. <i>et al. Nat. Chem. Biol.</i>; published online April 29, 2012; doi:10.1038/nchembio.957<br/> <b>Contact:</b> Gitte Neubauer, Cellzome AG, Heidelberg, Germany<br/>                     e-mail: <a href="mailto:gitte.neubauer@cellzome.com">gitte.neubauer@cellzome.com</a></p> |