

## THE DISTILLERY

## This week in therapeutics

| Indication      | Target/marker/<br>pathway   | Summary  | Licensing<br>status                           | Publication and contact<br>information  |
|-----------------|---|--|---|---|
| Hepatic disease |   |  |   |   |
| Liver fibrosis  | Peroxisome<br>proliferation–<br>activated<br>receptor-δ<br>(PPARD; PPARδ) | Cell culture and mouse studies suggest PPARò agonists could<br>help treat liver fibrosis. In mouse models of chemical-induced<br>liver fibrosis and cholestasis-induced liver fibrosis, the PPARò<br>agonist KD3010 decreased levels of liver fibrosis compared with<br>the PPARò agonist GW501516 or vehicle. Next steps could<br>include a Phase II trial of KD3010 to treat liver fibrosis in HCV<br>nonresponders.<br>KD3010 from Kalypsys Inc. has completed Phase I testing in<br>endocrine/metabolic and hepatic indications.<br>GlaxoSmithKline plc discontinued GW501516 after a Phase II<br>trial in dyslipidemia due to safety concerns.<br>At least three other companies have PPARò agonists in Phase I<br>or Phase II trials for endocrine/metabolic or cardiovascular<br>indications. | Patent and<br>licensing status<br>undisclosed | Iwaisako, K. <i>et al. Proc. Natl. Acad.</i><br><i>Sci. USA</i> ; published online April 25,<br>2012;<br>doi:10.1073/pnas.1202464109<br><b>Contact:</b> Bernd Schnabl, University<br>of California, San Diego School of<br>Medicine, La Jolla, Calif.<br>e-mail:<br><b>beschnabl@ucsd.edu</b><br><b>Contact:</b> Ronald M. Evans, Salk<br>Institute for Biological Studies, La<br>Jolla, Calif.<br>e-mail:<br><b>evans@salk.edu</b> |
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