

THE DISTILLERY

This week in therapeutics

| Indication | Target/marker/ pathway | Summary | Licensing status | Publication and contact information |
|--------------------|---|--|---|---|
| Infectious disease | | | | |
| Schistosomiasis | Thioredoxin glutathione reductase (TGR) | A study in cultured worms and worm-infected mice suggests that oxadiazole 2-oxide and phosphinic amide compounds could be useful for treating schistosomiasis. The disease is caused by parasitic worms of the genus <i>Schistosoma</i> that produce TGR, an enzyme that is essential for avoiding host immune response. The compounds showed activity against TGR and against cultured <i>S. mansoni</i> worms. In <i>S. mansoni</i> -infected mice, one of the compounds, furoxan, significantly reduced worm burden and hepatomegaly compared with those in infected, untreated controls (p <0.0001). The researchers are now testing about 40 derivatives for antiparasitic activity. | U.S. patent application filed covering use of furoxan to treat or prevent schistosomiasis; available for licensing | Sayed, A. <i>et al. Nat. Med.</i> ; published online March 16, 2007; doi:10.1038/nm1737 Contact: David Williams, Illinois State University, Normal, Ill. e-mail: dlwilli@ilstu.edu |

EpiCept Corp.'s EP128504, a 3,5-diaryl-oxadiazole, is in preclinical development for cancer.