

### This week in therapeutics

| Indication                | Target/marker/pathway                  | Summary   | Licensing status  | Publication and contact information   |
|---------------------------|--|---|---|---|
| <b>Infectious disease</b> |  |   |   |   |
| Rabies                    | Transmembrane glycoprotein G (RABVgp4) | <p>Studies in mice identified a live attenuated virus vaccine that could help treat and protect against rabies. In mice, intracranial administration of the attenuated triple rabies virus glycoprotein G variant did not cause clinical symptoms of rabies, unlike previous attenuated rabies viruses. In both normal and immunocompromised mice, the new attenuated virus prevented infection after challenge with a pathogenic rabies virus. In mice already infected with pathogenic rabies, the vaccine candidate prevented death in all animals when delivered 4 hours or 48 hours after infection. Next steps include testing the safety and efficacy of the vaccine in dogs and nonhuman primates.</p> <p>At least five companies have rabies vaccines and therapeutics in development stages ranging from preclinical to marketed.</p> <p><b>SciBX 2(28); doi:10.1038/scibx.2009.1109</b><br/> <b>Published online July 23, 2009</b></p> | <p>The vaccine has been patented by Thomas Jefferson University; it has been licensed to undisclosed parties for oral immunization of animals; available for licensing worldwide for vaccination applications excluding oral vaccination of animals</p> | <p>Faber, M. <i>et al. Proc. Natl. Acad. Sci. USA</i>; published online July 6, 2009; doi:10.1073/pnas.0905640106<br/> <b>Contact:</b> Bernhard Dietzschold, Thomas Jefferson University, Philadelphia, Penn.<br/>                     e-mail: <a href="mailto:bernhard.dietzschold@jefferson.edu">bernhard.dietzschold@jefferson.edu</a></p> |