

### This week in therapeutics

Indication	Target/marker/pathway	Summary	Licensing status	Publication and contact information
<b>Infectious disease</b>				
Bacterial infection	Not applicable	<p><i>In vitro</i> and mouse studies suggest that engineered bacteriophages could be used as adjuvants to antibiotics for treating bacterial infection. In a wide spectrum of <i>Escherichia coli</i> strains, including drug-resistant strains, combining an engineered phage targeting the bacterial DNA repair system with an antibiotic of the quinolone, aminoglycoside or <math>\beta</math>-lactam class increased bacterial death by 2–4 orders of magnitude compared with that seen using antibiotic treatment alone. Eighty percent of <i>E. coli</i>-infected mice treated with ofloxacin plus phage survived compared with 20% survival of mice treated only with ofloxacin. Additional <i>in vitro</i> experiments showed that phage targeting non-DNA repair mechanisms were also effective antibiotic adjuvants. Ongoing work includes evaluating the use of this technology for specific infectious diseases and investigating ways to overcome phage-associated hurdles for clinical use. No fewer than 14 companies have antibiotics that are marketed or approved to treat a range of bacterial infections. No fewer than 30 companies have antibiotics in Phase III testing to treat a wide range of bacterial infections. More than 30 companies have antibiotics in preclinical to early-stage clinical testing to treat bacterial infections.</p> <p><b>SciBX 2(9); doi:10.1038/scibx.2009.361</b>  <b>Published online March 5, 2009</b></p>	Patented; available for licensing	<p>Lu, T. &amp; Collins, J. <i>Proc. Natl. Acad. Sci. USA</i>; published online March 2, 2009;            doi:10.1073/pnas.0800442106  <b>Contact:</b> James J. Collins,            Boston University, Boston, Mass.            e-mail:  <a href="mailto:jcollins@bu.edu">jcollins@bu.edu</a></p>