

### This week in therapeutics

Indication	Target/marker/ pathway	Summary	Licensing status	Publication and contact information
<b>Cardiovascular disease</b>				
Myocardial infarction (MI)	MicroRNA-126 (miRNA-126)	Tissue culture and mouse studies suggest that increasing miRNA-126 expression may be useful for treating patients following MI. In the mouse studies, 9 of 11 miRNA-126-deficient mice died at 3 weeks post-MI compared with just 3 of 10 wild-type mice. <i>Ex vivo</i> cardiac tissue from miRNA-126-deficient mice showed significantly less endothelial sprouting at 6 days than tissue from wild-type controls ( $p=0.0001$ ). Next steps include developing strategies to increase miRNA-126 signaling, which may aid in post-MI cardiac repair.	Patent application filed for use in angiogenic therapy; licensed to miRagen Therapeutics Inc.	Wang, S. <i>et al. Dev. Cell</i> ; published online Aug. 11, 2008; doi:10.1016/j.devcel.2008.07.002 <b>Contact:</b> Eric N. Olson, University of Texas Southwestern Medical Center, Dallas, Texas e-mail: <a href="mailto:eric.olson@utsouthwestern.edu">eric.olson@utsouthwestern.edu</a>