

### This week in therapeutics

Indication	Target/marker/pathway	Summary	Licensing status	Publication and contact information
<b>Cancer</b>				
Lung cancer; liver cancer	MicroRNA-221 (miR-221); miR-222; phosphatase and tensin homolog deleted on chromosome ten (PTEN; MMAC1; TEP1); tissue inhibitor of metalloproteinases 3 (TIMP3); tumor necrosis factor-related apoptosis-inducing ligand (TRAIL)	<p>Studies in cell culture suggest that inhibiting miR-221 and miR-222 could help treat lung and liver cancers. In human non-small cell lung cancer (NSCLC) and hepatocellular carcinoma (HCC) cells, increased miR-221 and miR-222 levels were associated with reduced expression of the tumor suppressors PTEN and TIMP3. Cancer cells with low expression of miR-221 and miR-222 had greater TRAIL-induced cell death than those with higher expression of the two miRNAs. Next steps include determining whether liver cancer can be induced via liver-specific overexpression of miR-221 and miR-222.</p> <p><b>SciBX 3(1); doi:10.1038/scibx.2010.10</b>  <b>Published online Jan. 7, 2010</b></p>	Patent application filed; licensing status undisclosed	<p>Garofalo, M. <i>et al. Cancer Cell</i>; published online Dec. 7, 2009; doi:10.1016/j.ccr.2009.10.014</p> <p><b>Contact:</b> Carlo Maria Croce, Ohio State University, Columbus, Ohio  e-mail: <a href="mailto:carlo.croce@osumc.edu">carlo.croce@osumc.edu</a></p> <p><b>Contact:</b> Gerolama Condorelli, University of Naples, Naples, Italy  e-mail: <a href="mailto:gecondor@unina.it">gecondor@unina.it</a></p>