

This week in therapeutics

| Indication | Target/marker/pathway | Summary | Licensing status | Publication and contact information |
|---------------|--|--|---|--|
| Cancer | | | | |
| Solid tumors | VEGF; VEGF receptor 1 (FLT1; VEGFR-1); VEGFR-2 (KDR/Flk-1) | <p>A mouse study suggests prolonged anti-VEGF therapy to treat cancer could compromise endocrine function of the thyroid gland. In healthy mice, antibodies against VEGF and VEGFR-2 decreased vasculature density in the thyroid and other endocrine tissues compared with an antibody against VEGFR-1 or vehicle. Prolonged anti-VEGF treatment also decreased circulating levels of the thyroid hormone thyroxine compared with vehicle. Next steps include determining the functional impact of prolonged anti-VEGF therapy on endocrine tissues in mouse models and assessing endocrine levels in patients receiving anti-VEGF therapies.</p> <p><i>SciBX</i> 6(30); doi:10.1038/scibx.2013.792 Published online Aug. 8, 2013</p> | Patent and licensing status undisclosed | <p>Yang, Y. <i>et al. Proc. Natl. Acad. Sci. USA</i>; published online July 1, 2013; doi:10.1073/pnas.1301331110 Contact: Yihai Cao, Karolinska Institute, Stockholm, Sweden e-mail: yihai.cao@ki.se</p> |