

## IN BRIEF

**CANCER****A new target in Burkitt's lymphoma**

This study identified a new pathway involved in the pathogenesis of Burkitt's lymphoma that is amenable to therapeutic targeting. Using RNA sequencing of patient biopsy samples and Burkitt's lymphoma cell lines, together with re-analysis of published sequence data, the authors identified recurrent mutations in the transcription factor *TCF3* and its negative regulator *ID3*. *TCF3* activated the pro-survival phosphatidylinositol 3-kinase pathway in Burkitt's lymphoma cells, in part by augmenting tonic B cell receptor signalling. Inhibitors of the *TCF3* pathway inhibited growth of Burkitt's lymphoma cells and tumour xenografts.

**ORIGINAL RESEARCH PAPER** Schmitz, R. *et al.* Burkitt lymphoma pathogenesis and therapeutic targets from structural and functional genomics. *Nature* 12 Aug 2012 (doi:10.1038/nature11378)

**EPIGENETIC TARGETS****Towards a male contraceptive?**

Matzuk *et al.* investigated the spermatogenic actions of a small-molecule inhibitor (JQ1) of the testis-specific bromodomain-containing protein BRDT, which is essential for chromatin remodelling during spermatogenesis. Treatment of mice with JQ1 — via intraperitoneal injection — reduced seminiferous tubule area and testis size as well as the number and motility of spermatozoa without affecting hormone levels. Furthermore, JQ1 caused a complete yet reversible contraceptive effect in treated mice. This suggests that BRDT modulators such as JQ1 (which also binds BRD4) could serve as contraceptive agents.

**ORIGINAL RESEARCH PAPER** Matzuk, M. M. *et al.* Small-molecule inhibition of BRDT for male contraception. *Cell* 150, 673–684 (2012)

**ANTICANCER DRUGS****Tumour delivery of RNA interference**

RNA interference is a potentially useful way to study cancer targets *in vivo*, but is hampered by delivery issues. Using the *ID4* oncogene in ovarian cancer as an example, Ren *et al.* developed a tumour-penetrating nanocomplex made up of small interfering RNA (siRNA) complexed with a tumour-penetrating and membrane-translocating peptide, which enabled the specific delivery of siRNA deep into the ovarian tumour parenchyma. Treatment of ovarian tumour-bearing mice with the *ID4*-specific nanocomplex suppressed tumour growth and improved survival.

**ORIGINAL RESEARCH PAPER** Ren Y. *et al.* Targeted tumor-penetrating siRNA nanocomplexes for credentialing the ovarian cancer oncogene *ID4*. *Sci. Transl. Med.* 4, 147ra112 (2012)

**ANTIVIRAL DRUGS****Protection against influenza B infection**

This study reported the identification of three human monoclonal antibodies that protected mice against lethal challenge from two antigenically distinct lineages of influenza B viruses that are responsible for a large part of the annual flu burden. The authors showed that two antibodies (CR8033 and CR8071) recognized distinct conserved epitopes in the head region of influenza B haemagglutinin, whereas CR9114 bound to a conserved epitope in the haemagglutinin stem and protected mice against both influenza A and B viruses.

**ORIGINAL RESEARCH PAPER** Dreyfus, C. *et al.* Highly conserved protective epitopes on influenza B viruses. *Science* 9 Aug 2012 (doi:10.1126/science.1222908)