

This week in therapeutics

Indication	Target/marker/ pathway	Summary	Licensing status	Publication and contact information
Infectious disease				
Malaria	<i>Plasmodium falciparum</i> M17 leucyl aminopeptidase (M17)	<i>In vitro</i> studies identified the crystal structure of M17, which could be useful for rational design of malaria therapeutics. <i>P. falciparum</i> , the parasite that causes malaria, expresses the M17 and M1 leucyl aminopeptidases, which are required for parasite viability in red blood cells. X-ray crystallization studies of M17 with and without inhibitors revealed the structure of the enzyme's active site. Next steps include screening and optimizing additional aminopeptidase inhibitors.	Provisional patent application filed covering the crystal structure of M17 and its use as a structural model; unavailable for licensing	McGowan, S. <i>et al. Proc. Natl. Acad. Sci. USA</i> ; published online Jan. 4, 2010; doi:10.1073/pnas.0911813107 Contact: James C. Whisstock, Monash University, Melbourne, Victoria, Australia e-mail: James.Whisstock@med.monash.edu.au
		SciBX 3(6); doi:10.1038/scibx.2010.190 Published online Feb. 11, 2010		