

## THE DISTILLERY

## This week in therapeutics

Infectious disease   Staphylococcus Staphylococcus aureus   aureusimine (ausA) A study in mice suggests that inhibiting ausA could help   trat S. aureus infection. S. aureus genetic studies identified ausA as a target for   with ausA-deficient S. aureus led to significantly less of ausA inhibitors   weight loss and viral load in the heart, spleen and liver of ausA inhibitors   this infection with a wild-type strain (p<0.001 and p<0.01), patent; licensing staus   ureusing ausA inhibitors ausA inhibitors in animal models.	Indication	Target/marker/ pathway	Summary	Licensing status	Publication and contact information
StaphylococcusStaphylococcus aureus aureusimine (ausA)A study in mice suggests that inhibiting ausA could help treat S. aureus infection. S. aureus genetic studies identified ausAs as peptide secondary metabolites. In mice, infection with ausA-deficient S. aureus led to significantly less weight loss and viral load in the heart, spleen and liver than infection with a wild-type strain (p<0.001 and p<0.01, 	Infectious disease				
	Staphylococcus	Staphylococcus aureus aureusimine (ausA)	A study in mice suggests that inhibiting <i>ausA</i> could help treat <i>S. aureus</i> infection. <i>S. aureus</i> genetic studies identified ausAs as peptide secondary metabolites. In mice, infection with <i>ausA</i> -deficient <i>S. aureus</i> led to significantly less weight loss and viral load in the heart, spleen and liver than infection with a wild-type strain ( $p$ <0.001 and $p$ <0.01, respectively). Next steps include developing and evaluating ausA inhibitors in animal models.	ausA as a target for vaccine development and the development of ausA inhibitors covered by a pending patent; licensing status undisclosed	Wyatt, M.A. <i>et al. Science</i> ; published online June 3, 2010; doi:10.1126/science.1188888 <b>Contact:</b> Nathan A. Magarvey, McMaster University, Hamilton, Ontario, Canada e-mail: magarv@mcmaster.ca

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