



This week in techniques

Summary	Licensing status	Publication and contact information
Human neutralizing mAbs against MERS-CoV could aid the development of new therapeutics to treat or prevent infection. A screen of a single-chain variable domain fragment (scFv) phage library yielded seven unique fragments that bind to the MERS-CoV spike protein. In a nonhuman primate cell line, the most potent human mAbs generated from the identified scFvs neutralized the MERS-CoV with IC ₅₀ values ranging from 1.25 to 2 μg/mL. Next steps could include optimizing the lead neutralizing mAb and evaluating it in models of MERS-CoV infection. SciBX 7(21); doi:10.1038/scibx.2014.629	Patent and licensing status unavailable	Tang, XC. et al. Proc. Natl. Acad. Sci. USA; published online April 28, 2014; doi:10.1073/pnas.1402074111 Contact: Wayne A. Marasco, Dana-Farber Cancer Institute, Boston, Mass. e-mail: wayne_marasco@dfci.harvard.edu
	Human neutralizing mAbs against MERS-CoV could aid the development of new therapeutics to treat or prevent infection. A screen of a single-chain variable domain fragment (scFv) phage library yielded seven unique fragments that bind to the MERS-CoV spike protein. In a nonhuman primate cell line, the most potent human mAbs generated from the identified scFvs neutralized the MERS-CoV with IC $_{\rm 50}$ values ranging from 1.25 to 2 $\mu g/mL$. Next steps could include optimizing the lead neutralizing mAb and evaluating it in models of MERS-CoV infection.	Human neutralizing mAbs against MERS-CoV could aid the development of new therapeutics to treat or prevent infection. A screen of a single-chain variable domain fragment (scFv) phage library yielded seven unique fragments that bind to the MERS-CoV spike protein. In a nonhuman primate cell line, the most potent human mAbs generated from the identified scFvs neutralized the MERS-CoV with IC $_{50}$ values ranging from 1.25 to 2 μ g/mL. Next steps could include optimizing the lead neutralizing mAb and evaluating it in models of MERS-CoV infection.