

This week in techniques

Approach	Summary	Licensing status	Publication and contact information
Drug delivery			
Peptide drug delivery via cleavable albumin-binding peptide conjugates	<p>Cleavable, albumin-binding peptide conjugates could help improve the efficacy of glucagon-like peptide-1 (GLP-1)-targeting therapeutics. In human serum, conjugates consisting of GLP-1 or a GLP-1 analog linked to albumin-binding peptides retained their activity at the GLP-1 receptor (GLP1R; GLP-1R) longer than did unconjugated GLP-1. In a rat model of diabetes, the GLP-1 conjugates had serum half-lives 8–20 times longer than that of the unconjugated analog. Ongoing work includes optimizing the albumin-binding peptide to further increase its half-life and exploring methods of oral delivery for the peptide conjugates.</p> <p>Victoza liraglutide (NN2211), a once-daily GLP-1 analog from Novo Nordisk A/S, is marketed to treat type 2 diabetes.</p> <p>Byetta exenatide, a twice-daily injectable GLP1R agonist from Amylin Pharmaceuticals Inc. and Eli Lilly and Co., is marketed to treat type 2 diabetes.</p> <p>SciBX 3(26); doi:10.1038/scibx.2010.807 Published online July 1, 2010</p>	<p>Patented by Jinan University and Amersino Biodevelop Inc.; available for licensing</p> <p>Contact: Zhengding Su, University of Waterloo, Waterloo, Ontario, Canada e-mail: z2su@uwaterloo.ca</p>	<p>Li, H. <i>et al. Angew. Chem. Int. Ed. Engl.</i>; published online June 10, 2010; doi:10.1002/anie.201000287</p> <p>Contact: Zhengding Su, University of Waterloo, Waterloo, Ontario, Canada e-mail: z2su@uwaterloo.ca</p> <p>Contact: Hongjian Li, Jinan University, Guangdong, China e-mail: tlihj@jnu.edu.cn</p>