



This week in therapeutics

Indication	Target/marker/pathway	Summary	Licensing status	Publication and contact information
Neurology				
Parkinson's disease (PD)	Metabotropic glutamate receptor subtype 4 (mGluR4; GRM4)	In vitro and rat studies suggest mGluR4 positive allosteric modulators (PAMs) could be combined with existing PD therapies to treat PD. In a PD rat model of forelimb asymmetry, a selective mGluR4 PAM plus L-dopa increased forelimb function compared with either compound alone. In a PD rat model of catalepsy, an mGluR4 PAM plus preladenant improved behavior compared with either compound alone. Ongoing work includes investigating whether mGluR4 PAMs can decrease L-dopainduced dyskinesias in animal models and optimizing a new series of mGluR4 PAMs to treat PD. Preladenant (MK-3814; SCH 420814), an adenosine A _{2A} receptor (ADORA _{2A}) antagonist from Merck & Co. Inc., is in Phase III testing to treat PD. Istradefylline (KW-6002), an ADORA _{2A} antagonist from Kyowa Hakko Kirin Co. Ltd., is in Phase III testing to treat PD. SYN115, a selective ADORA _{2A} antagonist from Biotie Therapies Corp. and UCB Group, is in Phase II testing to treat PD.	Unpatented; available for licensing from Vanderbilt University Contact: P. Jeffrey Conn, Vanderbilt Center for Neuroscience Drug Discovery, Nashville, Tenn. e-mail: jeffrey.conn@ vanderbilt.edu Contact: Mary Kosinski, Vanderbilt Center for Technology Transfer and Commercialization, Nashville, Tenn. e-mail: mary.kosinski@ vanderbilt.edu	Jones, C.K. et al. J. Pharmacol. Exp. Ther.; published online Nov. 16, 2011; doi:10.1124/jpet.111.187443 Contact: Colleen M. Niswender, Vanderbilt University, Nashville, Tenn. e-mail: colleen.niswender@vanderbilt.edu

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