



This week in techniques

| Approach | Summary | Licensing status | Publication and contact information |
|---|---|---|--|
| Disease models | | | |
| Fluorescent transgenic zebrafish model of tauopathy | A fluorescent zebrafish model of tauopathy could be used to quickly identify therapeutics for Alzheimer's disease (AD) or frontotemporal dementia. Transgenic expression of fluorescently labeled human τ -protein in zebrafish neurons led to τ -protein accumulation, higher neuronal death and progression of AD behavioral pathology compared with what was seen in controls. Using the model, structure-based design identified a new inhibitor of glycogen synthase kinase 3 β (GSK3B) that blocked τ -kinase more potently than existing inhibitors both in vitro and in the zebrafish. The group also generated transgenic zebrafish expressing other amyloid proteins, such as β -amyloid (A β), TDP-43 and α -synuclein. Next steps could include testing the new inhibitor in additional animal models of neurological disease. Neurim Pharmaceuticals Ltd's Neu-120, a GSK3B inhibitor, is in Phase II testing to treat Parkinson's disease (PD). Noscira S.A.'s NP 12, also a GSK3 inhibitor, is in Phase II testing to treat AD. | Patent and licensing status unavailable | Haass, C. et al. J. Clin. Invest.; published online April 13, 2009; doi:10.1172/JCI37537 Contact: Christian Haass, Ludwig Maximilians University of Munich, Munich, Germany e-mail: chaass@med.uni-muenchen.de |
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