

# Direct Observation of Amyloid Fibril Growth and Propagation

Yuji Goto

Institute for Protein Research, Osaka University, Japan; E-mail: ygoto@protein.osaka-u.ac.jp

Amyloid fibrils form through nucleation and growth. To clarify the mechanism involved, direct observations of both processes are important (1, 2). First, seed-dependent fibril growth of  $\beta$ 2-microglobulin ( $\beta$ 2-m) and amyloid  $\beta$  peptide was visualized in real-time at the single fibril level using total internal reflection fluorescence microscopy combined with the binding of thioflavin T, an amyloid-specific fluorescence dye (3-5). Second, using atomic force microscopy, ultrasonication-induced formation of  $\beta$ 2-m fibrils was shown, indicating that ultrasonication is useful to accelerate the nucleation process (6). Third, with the proteolytic fragment of  $\beta$ 2-m, propagation and a transformation of fibril morphology was demonstrated (7). These direct observations indicate that template-dependent growth and structural diversity are key factors determining the structure and function of amyloid fibrils.

## References

1. Ban, T. *et al.* (2006) *Acc. Chem. Res.* **39**, 663-670.
2. Chatani & Goto (2005) *Biochim. Biophys. Acta* **1753**, 64-75.
3. Ban, T. *et al.* (2003) *J. Biol. Chem.* **278**, 16462-16465.
4. Ban, T. *et al.* (2004) *J. Mol. Biol.* **344**, 757-767.
5. Ban, T. *et al.* (2006) *J. Biol. Chem.* **281**, 33677-33683.
6. Ohhashi, Y. *et al.* (2005) *J. Biol. Chem.* **280**, 32843-32848.
7. Yamaguchi, K. *et al.* (2005) *J. Mol. Biol.* **352**, 952-960.