

Diffusive protofilament switching of kinesin-8 investigated with optical tweezers

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The budding yeast Kinesin-8 *Kip3* is a highly processive motor protein that walks to the end of microtubules and shortens them in a collective manner [1]. Microtubules usually consist of 12 to 15 circularly-arranged tubulin polymer chains, called protofilaments. Left-handed rotations of microtubules in *Kip3* gliding assays indicate sideward motion of *Kip3* perpendicular to the microtubule axis [2], i.e. a switching between single protofilaments. Here, we used a high-resolution optical tweezers setup in a force feedback mode to apply sideward loads on single motor proteins. Our studies show that *Kip3* steps sideward in both directions under alternating sideward loads. In control experiments with immobilized *Kip3* and not protofilament switching kinesin-1, we measured no effective sideward motion. Statistical analysis and comparison with simulations propose a diffusive motion of *Kip3* on the microtubule lattice with a preference to the left with respect to the direction of forward motion. This is consistent with the gliding assays. Protofilament switching has implications for the suggested mechanical signaling role of Kinesin-8 in budding yeast with respect to its ability to bypass obstacles.

References

- [1] V. Varga, C. Leduc, V. Bormuth, S. Diez, J. Howard: *Kinesin-8 motors act cooperatively to mediate length-dependent microtubule depolymerization*. *Cell* **138**, 1174–1183 (2009)
- [2] B. Nitzsche, F. Ruhnaw, S. Diez: *Quantum-dot-assisted characterization of microtubule rotations during cargo transport*. *Nature nanotechnology* **3**, 552–556 (2008)