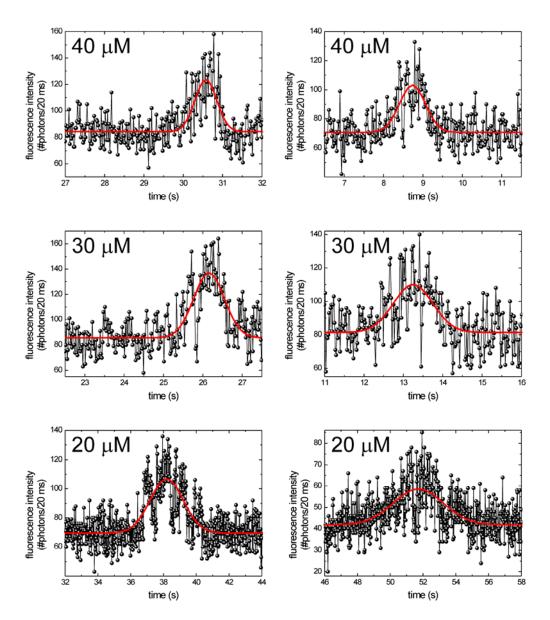
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Supporting Material

Alternating-Site Mechanism of Kinesin-1 Characterized by Single-Molecule FRET Using Fluorescent ATP Analogues

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Supplemental figure 1. Time traces of the donor fluorescence intensity of single-labelled kinesins walking through the confocal spot, in the presence of 20, 30 or 40 μ M fluorescent ATP. Solid red lines represent Gaussian fits.