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Non-Perturbative of Tracking Processive DNA Synthesis with Single-Molecule Fluorescence EVERETT LIPMAN, CHARLES WICKERSHAM¹, Department of Physics, University of California, Santa Barbara — We have demonstrated recently that double-stranded DNA labeled with a periodic series of fluorescent dyes can be used to track a single helicase. Here we describe how this technique can be modified to follow DNA synthesis. By means of a stepwise loss of fluorescence during strand displacement, we monitor processive motion of a single $\phi 29$ DNA polymerase without labeling or altering the enzyme or the template strand, and without applying any force. We observe a wide range of speeds, with the highest exceeding by several times that observed in other singlemolecule experiments. Because this method enables repeated observations of the same polymerase traversing identical segments of DNA, it should prove useful for studying sequence-specific effects in DNA replication and transcription.

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