

# Regulatory ATPase Sites of Cytoplasmic Dynein Affect Processivity and Force Generation\*<sup>§</sup>

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The heavy chain of cytoplasmic dynein contains four nucleotide-binding domains referred to as AAA1–AAA4, with the first domain (AAA1) being the main ATP hydrolytic site. Although previous studies have proposed regulatory roles for AAA3 and AAA4, the role of ATP hydrolysis at these sites remains elusive. Here, we have analyzed the single molecule motility properties of yeast cytoplasmic dynein mutants bearing mutations that prevent ATP hydrolysis at AAA3 or AAA4. Both mutants remain processive, but the AAA4 mutant exhibits a surprising increase in processivity due to its tighter affinity for microtubules. In addition to changes in motility characteristics, AAA3 and AAA4 mutants produce less maximal force than wild-type dynein. These results indicate that the nucleotide binding state at AAA3 and AAA4 can allosterically modulate microtubule binding affinity and affect dynein processivity and force production.

Cytoplasmic dynein is a molecular motor that moves toward the minus-end of microtubules. Underscoring its biological significance, dynein has been implicated in numerous microtubule-related functions, including cargo transport, mitotic spindle positioning, and nuclear segregation (see Ref. 1 for a review). Like many other biological motors, cytoplasmic dynein uses chemical energy derived from ATP hydrolysis to perform mechanical work. However, in contrast to other cytoskeletal motors of the kinesin and myosin superfamilies, dynein has multiple ATP binding sites. This poses the question of how dynein makes use of these multiple ATP sites and whether they might be involved in the regulation of the motor.

Dyneins are AAA+ ATPases, a superfamily of enzymes that have a diverse array of functions ranging from protein unfolding to membrane trafficking (see Refs. 2 and 3) for reviews). Despite their varied functions, AAA+ ATPases all

share a similar core architecture with conserved Walker-A (P loop) and Walker-B (phosphate sensor) motifs in their nucleotide binding domains (4, 5). Most AAA+ proteins oligomerize into hexameric, ringlike structures that act upon their substrates. In some cases, the identical AAA+ subunits may fire stochastically (e.g. ClpX (6)), whereas in other cases, sequential hydrolysis around the ring may occur (e.g. helicases (7)). Dynein is unusual in having multiple AAA+ domains concatenated in a single polypeptide chain that folds into a ringlike structure (8–10). The first four AAA+ domains (AAA1–AAA4) are capable of binding nucleotide (11, 12), whereas the last two AAA+ domains (AAA5–AAA6) are highly divergent, no longer bind nucleotide, and appear to serve a structural role in completing the ring. Between AAA4 and AAA5, an antiparallel coiled-coil stalk emerges with a microtubule binding domain at the tip. NH<sub>2</sub>-terminal to the first AAA domain is a “linker” domain that is thought to swing with respect to the stalk, possibly constituting the dynein power stroke (9).

The roles of the four functional AAA domains have been investigated by biochemical and mutagenesis studies. AAA1, the site of ATP-vanadate photocleavage (13), is generally acknowledged to be the major site of ATP hydrolysis and primary driver of the power stroke. Mutagenesis of this site greatly decreases ATP turnover (14, 15), abolishes motility (14), and eliminates the conformational change of the linker domain (16). The roles of the other sites remain less well understood. Single molecule studies suggest that a single ATP binding event may suffice for dynein to take a step along the microtubule (17). However, mutagenesis of the Walker A domain of AAA3 (predicted to interfere with nucleotide binding) greatly decreases microtubule-stimulated ATPase and microtubule gliding activity and causes “rigor-like” binding with the microtubule (14, 18, 19). Mutagenesis studies of AAA2 and AAA4 suggest they may have more minor roles. Collectively, these results suggest that AAA2–AAA4 assume some sort of regulatory role, but the details of how they participate in the dynein mechanism remain unclear.

Although prior *in vitro* motility studies have been performed on dynein ATP site mutations (14, 18, 19), they have focused upon Walker A mutations that disrupt ATP binding and have not examined the processive movement of a two-headed dynein. Here, we used previously developed single molecule motility assays (17, 20) to investigate the role of ATP hydrolysis at AAA3 and AAA4 on processivity and force production. We find that dynein bearing a Walker B mutation (that specifically disrupts ATP hydrolysis) at AAA3 is still processive, despite a

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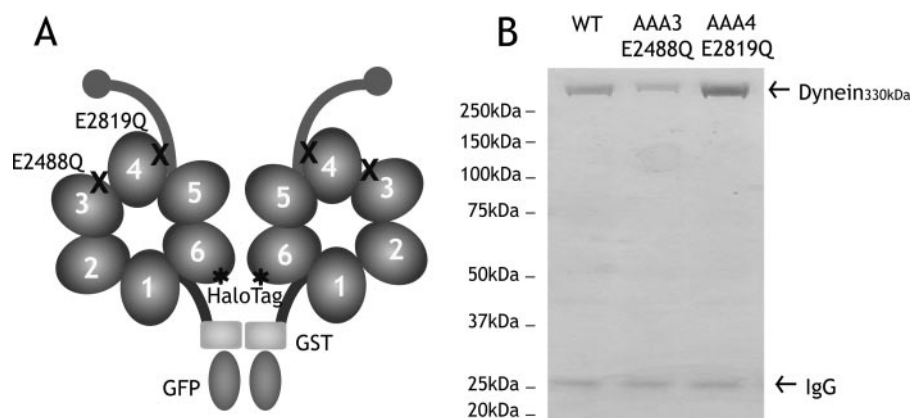
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<sup>§</sup> The on-line version of this article (available at <http://www.jbc.org>) contains supplemental Figs. S1 and S2 and Movies S1–S3.

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## Cytoplasmic Dynein Regulatory ATPase Sites



**FIGURE 1. Construction and purification of dynein ATP hydrolysis mutants.** *A*, a minimal *S. cerevisiae* cytoplasmic dynein that demonstrates processive motility was engineered as described previously (17). A glutathione *S*-transferase tag (*GST*) was incorporated at the  $\text{NH}_2$  terminus of the two heads of cytoplasmic dynein, whereas a HaloTag was fused to the COOH terminus of the two heads. In this paper, this construct is referred to as “wild-type dynein.” Highly conserved glutamate residues in the Walker B motif of domains AAA3 (E2488) and AAA4 (E2819) were mutated to glutamine to block ATP hydrolysis. *B*, Coomassie Blue-stained polyacrylamide gel of recombinant cytoplasmic dynein purified from *S. cerevisiae* by affinity purification. 330-kDa recombinant dynein is purified with minor amounts of 26-kDa IgG from the affinity matrix. *WT*, wild type.

severe effect on ATPase activity and motor velocity. Surprisingly, the AAA4 Walker B mutant displays enhanced processivity that is most likely mediated by an increase in microtubule binding affinity. We also show that AAA3 and AAA4 mutants can only generate 2-fold lower forces than wild-type dynein. Thus, the nucleotide binding state at AAA3 and AAA4 can regulate dynein microtubule affinity, processivity, and force-generating ability.

### EXPERIMENTAL PROCEDURES

**Protein Expression and Preparation**—A 330-kDa artificially dimerized expression construct of cytoplasmic dynein (glutathione *S*-transferase-Dyn1<sub>331kDa</sub>) was prepared and purified from *Saccharomyces cerevisiae* as previously described (17). Single point mutations (E2488Q (AAA3) and E2819Q (AAA4)) were introduced by the QuikChange mutagenesis kit (Stratagene). All constructs contained an  $\text{NH}_2$ -terminal IgG binding domain and a Tev protease cleavage site for purification, a green fluorescent protein tag for specific attachment to surfaces, and a COOH-terminal HaloTag (DHA; Promega) for fluorescent labeling. Prior to single molecule analyses, dynein was further purified by binding 50  $\mu\text{l}$  of  $\sim 300$   $\mu\text{g/ml}$  affinity-purified dynein to 10  $\mu\text{l}$  of 500  $\mu\text{g/ml}$  taxol-stabilized microtubules in the absence of ATP, centrifuging the microtubules, and then releasing from microtubules with 10 mM MgATP.

**Single Molecule Total Internal Reflection Fluorescence Microscopy**—Dynein was labeled with halotetramethylrhodamine (Promega) in the HaloTag domain and assayed for motility on Cy5-labeled axonemes, as previously described (17). Single molecules of dynein were visualized by a custom-built total internal reflection microscope using objective style total internal reflection fluorescence and an argon laser with 514 nm illumination at 3 milliwatts. Images were collected with an intensified CCD camera every 2 s for 5–10 min. Velocities and run lengths were determined by kymo-

graph analysis in ImageJ and corrected for photobleaching rates and axoneme length as previously described (17).

**Measurement of ATPase Activity**—Basal and microtubule-stimulated ATPase activities were measured by the EnzChek phosphate assay kit (Invitrogen). Assays were performed in motility buffer (30 mM Hepes, pH 7.4, 50 mM KAc, 2 mM  $\text{Mg}(\text{Ac})_2$ , and 1 mM EGTA) with 0–15  $\mu\text{M}$  taxol-stabilized microtubules and 5–10 nM dynein. Reactions were initiated with the addition of MgATP to a final concentration of 0–5 mM, and the absorbance at 360 nm was monitored by a spectrophotometer for 5–10 min. Protein concentrations of dynein were determined on SDS-polyacrylamide gels stained with

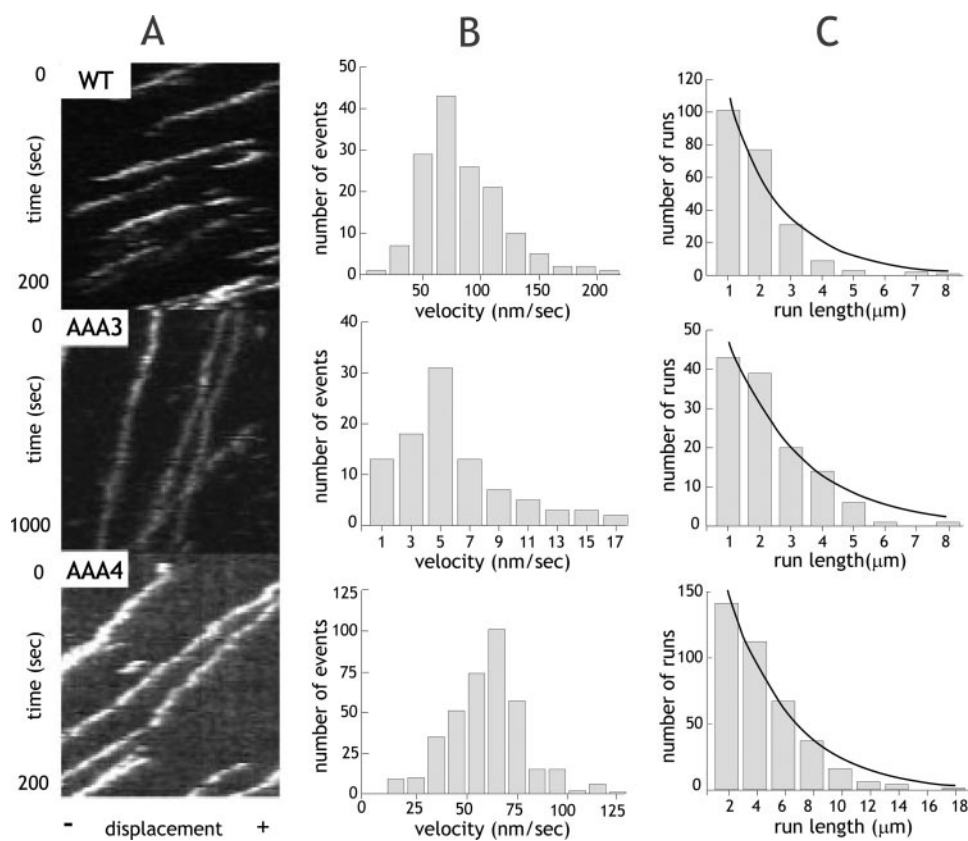
SYPRO-Red (Invitrogen), with a known concentration of  $\beta$ -actin used as a standard.

**Optical Trapping**—All experiments were performed with a custom-built force feedback-enhanced optical trapping microscope, as previously described (20). Carboxylated latex beads (0.92- $\mu\text{m}$  diameter; Invitrogen) coated with anti-green fluorescent protein antibodies were mixed with dynein in an assay solution containing 30 mM HEPES (pH 7.4), 50 mM KAc, 2 mM  $\text{Mg}(\text{Ac})_2$ , 1 mM EGTA, 0.5 mg/ml casein, 4.5 mg/ml glucose, 10 mM dithiothreitol, and an oxygen scavenging system of glucose oxidase and catalase (21). Stall force measurements and nucleotide-dependent movement studies were performed with 1 mM MgATP, whereas nucleotide-independent movement studies were performed in the presence of 10 units/ml apyrase. Dynein-coated beads were flowed into a standard flow chamber with adhered tetramethylrhodamine-labeled sea urchin axonemes, and bead displacements were recorded with a quadrant photodiode at 2 kHz.

### RESULTS

**Construction and Purification of ATPase Mutant Dyneins**—The native dynein heavy chain gene consists of a  $\sim 470$ -kDa polypeptide with  $\text{NH}_2$ -terminal cargo binding and dimerization domains and a COOH-terminal motor domain. We previously engineered a minimal dynein dimer that contains a 330-kDa minimal motor domain fused at its  $\text{NH}_2$  terminus to glutathione *S*-transferase, which self-associates to form a dimer (17). This construct (referred to as “wild-type dynein” in this study), which does not bind the dynein light or intermediate chains and has very low, substoichiometric amounts of the yeast Lis1 homologue, Pac1, shows robust processive movement in a single molecule fluorescence assay.

In order to specifically disrupt ATP hydrolysis at sites AAA3 and AAA4, point mutations changing an essential glutamate to a glutamine were introduced into the Walker B motifs (AAA3-E/Q and AAA4-E/Q; Fig. 1A). This glutamate residue is highly



**FIGURE 2. Single molecule processivity of dynein ATP hydrolysis mutants.** *A*, kymographs of single molecules of wild-type (WT) or ATP hydrolysis mutants. The x axis represents the length of an axoneme, and the y axis shows time. *B*, velocity histograms of wild-type and ATP hydrolysis mutants. The mean velocities  $\pm$  S.D. are  $73.9 \pm 34.2$  nm/s,  $4.6 \pm 3.7$  nm/s, and  $60.6 \pm 18.9$  nm/s ( $n = 221, 117,$  and  $384$ ) for wild type, AAA3-E/Q, and AAA4-E/Q, respectively. *C*, run length histograms of wild-type and ATP hydrolysis mutants are distributed in a single exponential decay. Run lengths were corrected for photobleaching and average axoneme length, and calculations for correct binning were performed as previously described (17). Run lengths ( $\pm$  S.E. as estimated by bootstrapping (17)) are  $2.25 \pm 0.14$ ,  $1.79 \pm 0.18$ , and  $4.38 \pm 0.45$   $\mu$ m for wild type, AAA3-E/Q, and AAA4-E/Q, respectively.

conserved, and glutamate-to-glutamine mutations disrupt nucleotide hydrolysis, but not nucleotide binding, in many other AAA proteins (6, 22). AAA2 does not have the conserved Walker B glutamate and thus was not investigated by mutagenesis in this study. Recombinant dyneins, without or with AAA3 or AAA4 point mutations, were purified from *S. cerevisiae* with an NH<sub>2</sub>-terminal affinity tag and then labeled with tetramethylrhodamine at the COOH-terminal HaloTag (Fig. 1B).

**Single Molecule Motility of Dynein Mutants**—To measure the motility of individual dynein molecules, we observed tetramethylrhodamine-labeled dynein moving along axonemes by total internal reflection fluorescence microscopy (17) (supplemental Movies 1–3). Contrary to bulk gliding assays, this method provides a direct determination of velocities and run lengths by observing single attachment, movement, and detachment events.

At 1 mM ATP, single wild-type, AAA3-E/Q and AAA4-E/Q dynein molecules demonstrated processive movement (*continuous lines* in kymographs in Fig. 2A). However, the velocity of the AAA3-E/Q mutant was substantially decreased ( $4.6 \pm 3.7$  nm/s) compared with wild-type motors ( $73.9 \pm 34.2$  nm/s; Fig. 2B). In contrast, AAA4-E/Q demonstrated only a modest decrease in velocity ( $60.6 \pm 18.9$  nm/s; Fig. 2B). The relative

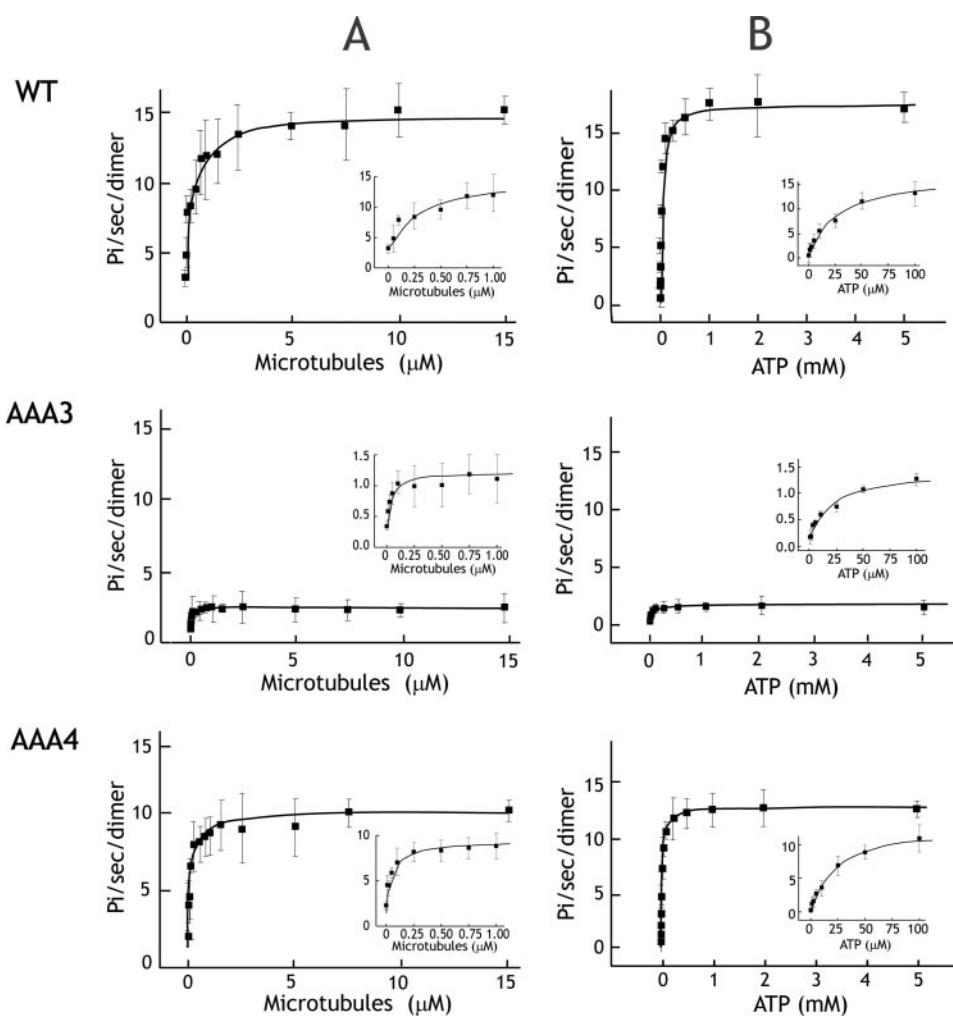
severity of these defects is reflected in the *in vivo* mutant phenotypes of these dyneins, where severe nuclear segregation defects are observed for the AAA3-E/Q but not the AAA4-E/Q mutant (Fig. S1) (18). These velocity decreases are similar to those reported for AAA3 and AAA4 Walker A mutant dynein monomers assayed for microtubule gliding *in vitro* (14), but the results here also demonstrate that these mutants still retain processive behavior.

To further evaluate processivity, run lengths were measured by kymograph analysis (Fig. 2C). The lengths of dynein runs were exponentially distributed, with the exponential decay constant representing the mean run length (23). The AAA3-E/Q mutant displayed a slight decrease in run length ( $1.79 \pm 0.18$   $\mu$ m) compared with wild-type dynein ( $2.25 \pm 0.14$   $\mu$ m). In contrast, AAA4-E/Q mutants had a surprising 2-fold increase in run length ( $4.39 \pm 0.45$   $\mu$ m). The frequency of extremely long runs (10–20  $\mu$ m) further demonstrated the pronounced gain in processivity.

We wished to exclude the possibility that the longer run length of AAA4-E/Q was due to aggregation, since an aggregate might have more

attachments to the microtubule and thereby detach less frequently. To test whether one or multiple dyneins are present in the moving spots, we examined their photobleaching behavior (Fig. S2). We found that all of the moving molecules ( $n = 47, 25,$  and  $45$  for wild-type, AAA3-E/Q, and AAA4-E/Q, respectively) showed only one- or two-step photobleaching, as expected for single dynein dimers. There was no significant difference in photobleaching between wild-type and mutant dyneins, ruling out the possibility that protein aggregation accounts for the increased run length of AAA4-E/Q or decreased velocity of AAA3-E/Q.

**Microtubule-stimulated ATPase Activity of Dynein ATPase Mutants**—To better understand the single molecule behaviors of the AAA3 and AAA4 ATP hydrolysis mutants, we measured their basal and microtubule-stimulated ATPase activities (Fig. 3A). In accordance with velocity reductions in the single molecule assay, the microtubule-stimulated ATPase rates,  $k_{cat}$ , of AAA3 and AAA4 mutants, were reduced 10- and 1.5-fold, respectively, compared with wild-type dynein (Table 1). The basal ATPase rates were reduced by a similar margin. Thus, microtubule stimulation was comparable (3–4-fold) for the mutants and wild-type dynein, implying that the loss of motility in the mutants is not due to a lack of microtubule stimulation.



**FIGURE 3. Microtubule-stimulated ATPase activity of dynein ATP hydrolysis mutants.** A, microtubule-stimulated ATPase activity of wild type and AAA mutants at 2 mM ATP. The insets show detailed views of microtubule-stimulated ATPase activity at low microtubule concentrations.  $K_{m,MT}$  values for wild type (WT), AAA3-E/Q, and AAA4-E/Q are 0.31, 0.03, and 0.069  $\mu\text{M}$ , respectively. B, the ATP dependence of microtubule-stimulated ATPase activity measured with 5  $\mu\text{M}$  taxol-stabilized microtubules. Insets show detailed views of the curve at low ATP concentrations.  $K_{m,ATP}$  values for wild type, AAA3-E/Q, and AAA4-E/Q are 25.2, 24.6, and 24.7  $\mu\text{M}$ , respectively. Each dot represents the mean  $\pm$  S.D. from three measurements of one preparation. Mean values from three preparations are presented in Table 1.

In summary, these results suggest that trapping AAA4 and particularly AAA3 in an ATP state decreases ATP turnover at AAA1, the main hydrolytic site of dynein.

The AAA3 and AAA4 mutants also exhibited a striking increase in their binding affinity for microtubules. A  $\sim 20$ -fold increase in  $K_{m,MT}$ <sup>3</sup> for microtubule-stimulated ATPase activity was observed for AAA3-E/Q ( $K_{m,MT} = 0.03 \mu\text{M}$ ). AAA4-E/Q also exhibited a  $\sim 5$ -fold ( $K_{m,MT} = 0.09 \mu\text{M}$ ) increase in microtubule-binding affinity, and this tighter interaction with the microtubule might explain the increased processivity of the AAA4-E/Q mutant. Interestingly, the relative increase in microtubule affinity for the AAA4-E/Q mutant appeared to be specific for the dimeric dynein construct. In a dynein monomer (lacking the NH<sub>2</sub>-terminal glutathione S-transferase), the  $K_{m,MT}$  for AAA4-E/Q (1.7  $\mu\text{M}$ ) and wild type (2.2  $\mu\text{M}$ ) were comparable (data not shown), suggesting that the mutation

may affect microtubule affinity by increasing coordination between the two heads of dynein.

We also determined the  $K_{m,ATP}$  by measuring microtubule-stimulated ATPase activity at different ATP concentrations (Fig. 3B). If ATP hydrolysis occurs at multiple AAA domains, we would expect the data to show a biphasic fit, with at least two binding constants for ATP, as was suggested motility studies other AAA ATPases (24) and motility studies with mammalian dynein-dynactin complexes (25). However, our data were well fit by a Michaelis-Menten equation (Fig. 3B), implying that a single nucleotide binding site dominates the ATPase reaction. In addition, we find no significant difference between the ATP binding affinities of wild-type and AAA mutant dyneins, implying that blocking ATP hydrolysis at AAA3 and AAA4 does not significantly affect ATP binding at AAA1.

**Stall Force Measurements of ATPase Mutant Dyneins**—We next determined whether ATP hydrolysis at AAA3 and AAA4 contributed to the force generation of dynein. For these experiments, we bound wild-type and mutant green fluorescent protein-tagged dyneins to latex beads, which could be captured by a fixed optical trap (Fig. 4A). To ensure that bead movements were due to a single dynein molecule, the dynein-to-bead ratio was adjusted so that the fraction of moving beads was  $< 0.3$  (representing a  $> 99\%$  probability that movements were due to a single molecule (26)). Dynein mutants exhibited similar behavior to wild-type dynein, moving the bead away from the trap center and then stalling, often for several minutes, under a maximum opposing load (Fig. 4B).

Both AAA3-E/Q and AAA4-E/Q (2.6 and 3.7 pN, respectively; Fig. 4C) exhibited lower stall forces compared with wild-type dynein (4.5 pN;  $p < 0.0001$ , Student's *t* test). These experiments show that the ATP hydrolysis mutants remain processive under load but fail to achieve the same stall forces as wild-type dynein.

**Nucleotide-independent Movement of Dynein Induced by Force**—We have recently shown that a pull from an optical trap will cause dynein to step processively along a microtubule in the absence of nucleotide (20). In this experiment, tension applied from the optical trap causes the rear dynein head to detach from the microtubule and then rebind to a new site further along the

<sup>3</sup> The abbreviations used are: MT, microtubule; pN, piconewtons.

TABLE 1

## Motility and ATPase activity of AAA hydrolysis mutants

Data was collected from three independent preparations of dynein for each construct. Reported values are mean and S.E. for three independent preparations. For velocity and run length data, >100 molecules were measured for each preparation.

	Velocity	Run length	MT-stimulated ATPase			Basal ATPase $k_{cat}$
			$k_{cat}$	$K_{m,MT}$	$K_{m,ATP}$	
	nm/s	$\mu\text{m}$	$s^{-1}$	$\mu\text{M}$	$\mu\text{M}$	$s^{-1}$
Wild type	$72.5 \pm 5.5$	$1.99 \pm 0.16$	$14.1 \pm 0.36$	$0.59 \pm 0.28$	$26.2 \pm 0.8$	$3.74 \pm 0.35$
AAA3 (E2488Q)	$5.1 \pm 0.9$	$1.78 \pm 0.10$	$1.38 \pm 0.14$	$0.03 \pm 0.01$	$23.7 \pm 0.7$	$0.30 \pm 0.05$
AAA4 (E2819Q)	$62.5 \pm 1.6$	$4.55 \pm 0.39$	$10.6 \pm 0.72$	$0.09 \pm 0.03$	$25.3 \pm 2.4$	$3.36 \pm 0.59$

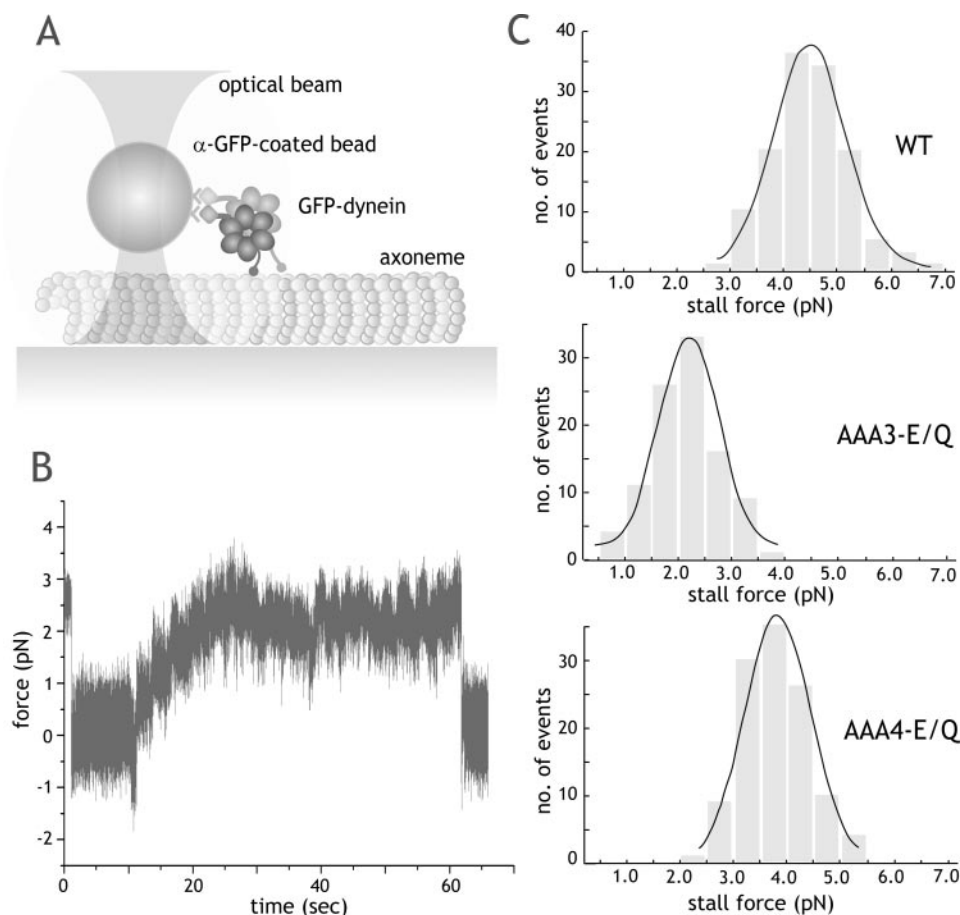


FIGURE 4. Stall force measurements of ATPase mutant dynein in the optical trap. *A*, schematic representation of the fixed optical trap setup used for stall force measurements in this paper. *B*, a representative trace of a single AAA4-E/Q dynein motor moving against force in a fixed optical trap at 1 mM ATP (trap stiffness ( $k$ ) = 0.034 pN/nm). The trace shows a long stall event of ~1 min, followed by release, which is typical of both wild-type and mutant dynein. *C*, stall force distributions of wild-type and ATPase mutant dyneins. Stall forces (mean  $\pm$  S.D.) are  $4.5 \pm 1.3$  pN ( $n = 132$ ),  $2.6 \pm 1.2$  pN ( $n = 100$ ), and  $3.7 \pm 1.2$  pN ( $n = 115$ ) for wild type (WT), AAA3-E/Q, and AAA4-E/Q, respectively. Black lines, Gaussian fit of the data.

microtubule. We applied this assay to gauge the microtubule-binding affinities of AAA mutants under an applied load (Fig. 5A).

Although ATP-driven velocity differed between wild-type, AAA3-E/Q, and AAA4-E/Q dyneins, all three motors behaved in an indistinguishable manner in this nucleotide-independent, force-driven stepping assay. The frequency (Fig. 5C) and velocity (Fig. 5D) were very similar for all three dyneins at different applied loads. Both AAA mutants also showed the same intrinsic asymmetry to force-induced movement as found for wild-type dynein, with only 3 pN of force required to induce robust movement in the forward direction and 10 pN of force to induce robust movement in the backward direction (Fig. 5C).

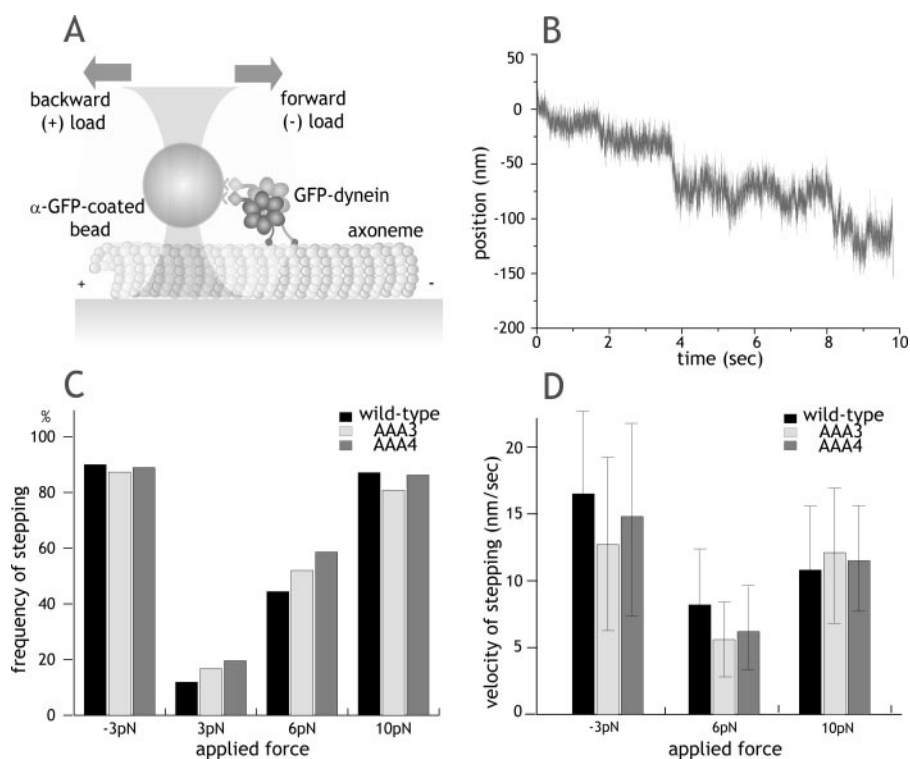
This assay primarily tests the microtubule binding affinity in the apo state, and the results reveal that the Walker B mutations in AAA3 and AAA4 do not affect the microtubule-binding domain under conditions where the motor is not undergoing cycles of ATP hydrolysis.

## DISCUSSION

In this work, we have explored the roles of nucleotide hydrolysis at the dynein "regulatory" AAA domains, AAA3 and AAA4. This study differs from prior biochemical work on the AAA domains (14), which employed nonprocessive dynein monomers and mutated the Walker A motif, which is expected to interfere with nucleotide binding. Our results show that blocking nucleotide hydrolysis at AAA3 and AAA4 significantly affects motor velocity, processivity, and force production but not nucleotide affinity at AAA1 and microtubule binding affinity in the apo state. These studies provide new insight into how the nucleotide states of AAA3 and AAA4 can affect the main hydrolytic site (AAA1) and the mechanics of the motor.

The principal consequence of blocking ATP hydrolysis at AAA4 is on motor processivity, resulting in a 2-fold increase in the run length.

The processivity of molecular motors is thought to be mediated by alternating catalysis of the two heads (27), resulting in hand-over-hand motion (28). A processive run is terminated when both motor domains simultaneously detach from the microtubule, which would be more likely if both heads are in a weak binding state. Here, we show that blocking ATP hydrolysis at the AAA4 domain increases the binding affinity for microtubules in the presence of ATP for the dimeric dynein construct. This higher microtubule binding affinity is probably responsible for the longer run length of AAA4-E/Q, since the tighter interaction would likely equate to a lower probability of dissociation from the track. AAA3-E/Q



**FIGURE 5. Nucleotide-independent movement of dynein induced by force.** *A*, schematic of force-induced stepping experiments. Forward force is defined as the direction in which dynein normally moves (toward the microtubule minus-end). *B*, example trace of nucleotide-free, force-induced stepping for AAA4 mutants in a force feedback trap with 6 pN of backward load.  $k = 0.062$  pN/nm. *C*, frequency of nucleotide-independent dynein movement after applying constant forward ( $-3$  pN) or backward load (3, 6, and 10 pN).  $n > 25$  molecules were tested at each force for each construct. Movement was scored within a 10-s time window of pulling on a dynein-coated bead attached to the microtubule. *D*, velocity of force-induced dynein movement with forward ( $-3$  pN) or backward (6 and 10 pN) load (mean  $\pm$  S.D.). Velocities at 3 pN backward load were not measured due to the small fraction of moving motors. GFP, green fluorescent protein.

has an even higher affinity for microtubules, although its run length is similar to wild-type. However, if one calculates mean attachment times, AAA3-E/Q is attached much longer (360 s) than both wild-type (30 s) and AAA4-E/Q (72 s), showing that higher microtubule affinity is also reflected in the motility characteristics of AAA3-E/Q.

In previous studies, motors with increased processivity have been made by engineering the dimerization domain of the motor (29) or microtubule-interacting elements (30). Our finding highlights a novel example where engineering an ATPase domain, which has no known interactions with microtubules, causes an increase in processivity. ATP hydrolysis at AAA3 and AAA4 could affect microtubule binding affinity either by a direct allosteric effect communicated through the coiled-coil stalk to the microtubule binding domain or by affecting the kinetic cycle of AAA1, such that the motor spends a longer time in nucleotide states associated with tight microtubule binding. In addition, the pronounced difference of microtubule binding affinity between dimeric wild-type and AAA4-E/Q dynein, but not monomeric wild-type and AAA4-E/Q dynein, suggests the possibility that kinetic coupling is increased between the two heads of dynein in AAA4-E/Q mutants. Further studies will be required to distinguish between these mechanisms.

Our data also reveal that blocking ATP hydrolysis at AAA3 or AAA4 affects the catalytic and mechanical force production activities of dynein. This is particularly evident for AAA3-E/Q,

which reduces the overall ATPase rate and movement velocity by an order of magnitude, effects that are most likely mediated by repressing ATP turnover at AAA1. Previous studies have shown that a Walker A mutation (which interferes with nucleotide binding) in AAA3 similarly reduces the ATPase and microtubule gliding velocity of *Dicystostelium* dynein by  $\sim 20$ -fold (14). Taken together, these results suggest that both ATP binding and hydrolysis at AAA3 are important for fast nucleotide turnover at AAA1, implying allosteric communication between different AAA sites.

Although these and other mutagenesis studies show that the nucleotide state of AAA3 and AAA4 can affect overall dynein activity, the predominant nucleotide state of AAA3 and AAA4 has yet to be determined. The nucleotide turnover rate at AAA3 and AAA4 also remains unknown, although prior dwell time analysis (17) and the single site fit of the ATPase data in this study suggest that ATP hydrolysis at AAA3 (and AAA4) does not occur during every cycle of ATP turnover

at AAA1. Thus, AAA3 and AAA4 may function as regulatory sites for AAA1, perhaps analogous to the main (D2) and regulatory (D2) catalytic sites of p97, another AAA+ protein (31–33). To answer such questions and better understand the interplay of the four AAA nucleotide sites, new tools will be needed to directly probe the nucleotide state of each AAA domain in the native dynein enzyme.

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