# ANALYSIS OF NONPROCESSIVE MOLECULAR MOTOR TRANSPORT USING RENEWAL REWARD THEORY\*

CHRISTOPHER E. MILES<sup>†</sup>, SEAN D. LAWLEY<sup>†</sup>, AND JAMES P. KEENER<sup>‡</sup>

Abstract. We propose and analyze a mathematical model of cargo transport by nonprocessive molecular motors. In our model, the motors change states by random discrete events (corresponding to stepping and binding/unbinding), while the cargo position follows a stochastic differential equation (SDE) that depends on the discrete states of the motors. The resulting system for the cargo position is consequently an SDE that randomly switches according to a Markov jump process governing motor dynamics. To study this system we (1) cast the cargo position in a renewal theory framework and generalize the renewal reward theorem and (2) decompose the continuous and discrete sources of stochasticity and exploit a resulting pair of disparate timescales. With these mathematical tools, we obtain explicit formulas for experimentally measurable quantities, such as cargo velocity and run length. Analyzing these formulas then yields some predictions regarding so-called nonprocessive clustering, the phenomenon that a single motor cannot transport cargo, but two or more motors can. We find that having motor stepping, binding, and unbinding rates depend on the number of bound motors, due to geometric effects, is necessary and sufficient to explain recent experimental data on non-processive motors.

Key words. molecular motors, intracellular transport, renewal reward theory, stochastic hybrid systems, switching SDEs

AMS subject classifications. 92C05, 60K20, 60J28, 34F05, 60H10

DOI. 10.1137/17M1156824

1. Introduction. Active intracellular transport of cargo (such as organelles) is critical to cellular function. The primary type of active transport involves molecular motors, which alternate between epochs of active transport (discrete stepping) along a microtubule and epochs of passive diffusion when the motors are unbound from the microtubule. Stochastic modeling of this fundamental process has a rich and fruitful history (see the review [5]).

In both experimental and modeling studies, *processive* motors have received considerable attention. Processive motors are characterized by taking hundreds of steps along a microtubule before unbinding. In contrast, *nonprocessive* motors (such as most members of the kinesin-14 family) take very few (1 to 5) steps before unbinding from a microtubule [6, 11]. Nonprocessive motors are crucial to a number of cellular processes, including directing cytoskeletal filaments [42], driving microtubulemicrotubule sliding during mitosis [14], and retrograde transport along microtubules in plants [48]. Here, we focus on motor behavior during transport.

Some curious properties of nonprocessive motor transport were found in [16]. One nonprocessive (Ncd) motor has extremely limited transport ability, measured by both velocity and run length (distance traveled before detaching from a microtubule).

<sup>\*</sup>Received by the editors November 14, 2017; accepted for publication (in revised form) July 24, 2018; published electronically September 20, 2018.

http://www.siam.org/journals/siap/78-5/M115682.html

**Funding:** This work was supported by NSF grant DMS-RTG 1148230. The first and third authors were also supported by NSF grant DMS 1515130. The second author's work was also supported by NSF grant DMS 1814832.

<sup>&</sup>lt;sup>†</sup>Department of Mathematics, University of Utah, Salt Lake City, UT 84112 (miles@math.utah.edu, lawley@math.utah.edu).

<sup>&</sup>lt;sup>‡</sup>Departments of Mathematics and Bioengineering, University of Utah, Salt Lake City, UT 84112 (keener@math.utah.edu).

However, *two* nonprocessive motors somehow act in unison to produce significant directed motion, a phenomenon termed "clustering." This observation is supported by the subsequent studies [26, 38], where similar experiments were performed creating a mutant of attached nonprocessive kinesin-14 motors, and processivity emerges. Moreover, the authors of [16] note that adding more Ncd motors beyond two further increases transport ability. In contrast, one processive motor (kinesin-1) is sufficient to produce transport, and additional motors do not significantly increase transport ability [16, 45]. Other interesting facets of transport by nonprocessive motors include the emergence of processive transport in the presence of higher microtubule concentration [17] or opposing motors [22].

In this work, we formulate and analyze a mathematical model to investigate the natural question, how do nonprocessive motors cooperate to transport cargo? Our model predicts that nonprocessive motor stepping, binding, and unbinding rates must depend on the number of bound motors, and that this dependence is a key mechanism driving the collective transport of nonprocessive motors. We note that such dependence has been observed in experiments [13, 20] and in simulations of detailed computational models [30, 12, 35, 17], all stemming from geometric effects of cargo/motor configuration.

Nonprocessive motors are notoriously difficult to study experimentally, because they take only a few steps before detaching. For this same reason, it is not clear how to best model nonprocessive motors, or known if existing modeling frameworks, such as mean-field methods [27, 9] or averaging the stepping dynamics into an effective velocity [36], are appropriate. Hence, our model explicitly includes the discrete binding, unbinding, and stepping dynamics of each motor, as well as the continuous tethered motion of the cargo.

Mathematically, our model takes the form of a randomly switching stochastic differential equation (SDE), and thus merges continuous dynamics with discrete events. The continuous SDE dynamics track the cargo position, while the discrete events correspond to motor binding, unbinding, and stepping. Our model is thus a *stochastic hybrid system* [4], which is often a two-component process,  $(J(t), X(t))_{t\geq 0} \in \mathcal{I} \times \mathbb{R}^d$ , where J is a Markov jump process on a finite set  $\mathcal{I}$ , and X evolves continuously by

(1) 
$$dX(t) = F_{J(t)}(X(t)) dt + \sigma dW(t),$$

where  $\{F_j(x)\}_{j\in\mathcal{I}}$  is a given finite family of vector fields,  $\sigma \geq 0$ , and W is a Brownian motion. That is, X follows an SDE whose right-hand side switches according to the process J.

However, our model differs from most previous hybrid systems in some key ways. First, the set of possible continuous dynamics (e.g., possible right-hand sides of (1)) for our model is infinite. Second, the new right-hand side of (1) that is chosen when J jumps depends on the value of X at that jump time, although the rates dictating J are taken to be independent of X.

We employ several techniques to analyze our model and make predictions regarding nonprocessive motor transport. First, we cast our model in a renewal theory framework and generalize the classical renewal reward theorem [47] to apply to our setting, distinct from previous motor applications [31, 23, 24, 25, 44]. Next, we decompose the stochasticity in the system by averaging over the diffusion while conditioning on a realization of the jump process. This effectively turns the randomly switching SDE into a randomly switching ordinary differential equation (ODE) and thus a piecewise deterministic Markov process [8]. Finally, we observe that for biologically reasonable parameter values, the relaxation rate of the continuous cargo dynamics is much faster than the jump rates for the discrete motor behavior. We then exploit this timescale separation to find explicit formulas for key motor transport statistics.

The rest of the paper is organized as follows. We formulate the mathematical model in section 2. In section 3, we generalize the renewal reward theorem to apply to our model. In section 4, we derive explicit formulas to evaluate motor transport. In section 5, we use the model to make biological predictions. We conclude with a brief discussion and an Appendix that collects several proofs.

2. Mathematical model. We model the motion of a single cargo driven by  $M \ge 1$  motors along a single microtubule. These motors are permanently attached to the cargo, but they can bind to and unbind from the microtubule. At any time  $t \ge 0$ , the state of our model is specified by

$$(X(t), \mathbf{Z}(t), \mathbf{J}(t)) \in \mathbb{R} \times \mathbb{R}^M \times \{u, b\}^M$$

where  $X(t) \in \mathbb{R}$  is the location of the center of the cargo,  $\mathbf{Z}(t) = (Z_i(t))_{i=1}^M \in \mathbb{R}^M$ gives the locations of the centers of M motors, and  $\mathbf{J}(t) = (J_i(t))_{i=1}^M$  specifies if each motor is unbound or bound. Spatial locations are measured along the principal axis of the microtubule, which we identify with the real line.

The cargo position evolves continuously in time, while the positions and states of motors change by discrete events, which correspond to binding to the microtubule, stepping along the microtubule, or unbinding from the microtubule. Specifically, in between these discrete motor events, X(t) follows an Ornstein–Uhlenbeck (OU) process centered at the average bound motor position,

(2) 
$$dX(t) = \frac{k}{\gamma} \sum_{i \in I(t)} \left( Z_i(t) - X(t) \right) dt + \sqrt{2k_B T/\gamma} \, dW(t).$$

Here,  $I(t) = \{i : J_i(t) = b\} \subseteq \{1, \ldots, M\}$  gives the indices of motors that are bound at time  $t \ge 0$ , and  $\{W(t)\}_{t\ge 0}$  is a standard Brownian motion. The SDE (2) stems from assuming a viscous (low Reynolds number) regime with drag coefficient  $\gamma > 0$ , and that each bound motor exerts a Hookean force on the cargo with stiffness k > 0. The Stokes–Einstein relation specifies the diffusion coefficient  $k_B T/\gamma$ , where  $k_B$  is Boltzmann's constant and T is the absolute temperature.

The discrete behavior of motors is as follows. Let  $m(t) \in \{0, 1, ..., M\}$  denote the number of bound motors at time  $t \ge 0$ ,

$$m(t) = \sum_{i=1}^{M} \mathbb{1}_{\{J_i(t) \neq u\}} \in \{0, 1, \dots, M\},\$$

where  $1_{\{A\}}$  denotes the indicator function on an event A. Each unbound motor binds to the microtubule at rate  $k_{on}(m(t)) > 0$ . Since unbound motors are tethered to the cargo, if an unbound motor binds at time  $t \ge 0$ , then we assume that it binds to the track at X(t) (motors can bind anywhere along the microtubule, not only at binding sites). We could allow it to bind to a random position, but if the mean binding position is X(t), then our results are unchanged. The position of each bound motor is fixed until it either steps or unbinds. Each bound motor unbinds at rate  $k_{off}(m(t)) > 0$ and steps at rate  $k_{step}(m(t)) > 0$ . When a motor steps, we add  $\delta > 0$  to its position, which is then fixed until it steps again or unbinds. This discrete motor behavior is summarized in Figure 1. We emphasize that the motor binding, unbinding, and stepping rates are allowed to depend on the number of bound motors, m(t), but are otherwise independent of X(t).

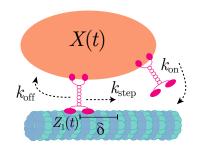


FIG. 1. Schematic describing the binding, unbinding, and stepping of motors. The positions of cargo and bound motors are X(t) and  $Z_i(t)$ , respectively, both measured with respect to the principal axis of the microtubule. The state of the motor can switch between bound or unbound, and while bound, the motor can step, incrementing  $Z_i(t)$  by displacement  $\delta$ .

2.1. Nondimensionalization and assumptions. We now give a dimensionless and more precise formulation of the model described above. First, we nondimensionalize the model by rescaling time by the rate  $k_{\text{off}}(1)$  and space by the inverse length  $\delta^{-1}$ . Next, we note that unbound motors do not affect the cargo position. Hence, for convenience we can take  $Z_i(t) = X(t)$  if the *i*th motor is unbound, meaning that we can include unbound motors in the sum in (2) with zero contribution and make the sum over *all* motors. This yields the simplified dimensionless form

(3) 
$$dX(t) = \varepsilon^{-1} \sum_{i=1}^{M} \left( Z_i(t) - X(t) \right) dt + \sigma dW(t),$$

where

$$\varepsilon := k_{\text{off}}(1)\gamma/k, \qquad \sigma := \sqrt{2k_BT/(\delta^2 k_{\text{off}}(1)\gamma)}$$

and motors bind, unbind, and step at dimensionless rates

(4) 
$$\lambda_{\rm on}(m) := \frac{k_{\rm on}(m)}{k_{\rm off}(1)}, \quad \lambda_{\rm off}(m) := \frac{k_{\rm off}(m)}{k_{\rm off}(1)}, \quad \lambda_{\rm step}(m) := \frac{k_{\rm step}(m)}{k_{\rm off}(1)}.$$

We find it convenient for our analysis to track the number of steps taken by each motor before unbinding, so let us expand the state space of  $\mathbf{J}(t)$  so that its components  $(J_i(t))_{i=1}^M$  each take values in  $\{u, 0, 1, 2, ...\}$  with transition rates

(5) 
$$u \stackrel{\lambda_{\text{on}}(m(t))}{\to} 0, \quad j \stackrel{\lambda_{\text{step}}(m(t))}{\to} j + 1, \quad j \stackrel{\lambda_{\text{off}}(m(t))}{\to} u, \quad j \neq u.$$

The components of  $\mathbf{J}(t)$  are conditionally independent given m(t). At time  $t \ge 0$ , the *i*th motor is unbound if  $J_i(t) = u$ , bound if  $J_i(t) \ge 0$ , and steps when  $J_i(t)$  transitions from j to j + 1 for  $j \ge 0$ . Under these assumptions, m(t) is itself a Markov process on  $\{0, 1, \ldots, M\}$  with transition rates

(6) 
$$0 \stackrel{M\lambda_{\text{on}}(0)}{\rightleftharpoons} 1 \stackrel{(M-1)\lambda_{\text{on}}(1)}{\rightleftharpoons} 2 \rightleftharpoons \cdots \rightleftharpoons M - 2 \stackrel{2\lambda_{\text{on}}(M-2)}{\rightleftharpoons} M - 1 \stackrel{\lambda_{\text{on}}(M-1)}{\rightleftharpoons} M.$$

For simplicity, we assume that the motors are initially unbound and that the cargo and motors start at the origin,

$$J_i(0) = u, \quad X(0) = Z_i(0) = 0, \quad i \in \{1, \dots, M\}.$$

The position of the ith motor is then

7) 
$$Z_i(t) = (X(\tau_i(t)) + J_i(t)) \mathbf{1}_{\{J_i(t) \neq u\}} + X(t) \mathbf{1}_{\{J_i(t) = u\}}, \quad i \in \{1, \dots, M\},$$

where  $\tau_i(t)$  is the most recent binding time of the *i*th motor,

$$\tau_i(t) = \sup\{s < t : J_i(s) = u\}, \quad i \in \{1, \dots, M\}.$$

We assume the Brownian motion  $W = \{W(t)\}_{t \ge 0}$  and the jump process  $\mathbf{J} = \{\mathbf{J}(t)\}_{t \ge 0}$ are independent.

3. Cargo position as a renewal reward process. In order to analyze our model, we first show that X(t) is a renewal reward process with partial rewards (see [47]) and extend the classical renewal reward theorem to our case of partial rewards. This framework has an intuitive interpretation: the net displacement of cargo is determined by the displacement accrued at each epoch of being bound or unbound. However, there is a technical challenge. In the most classical setting, the renewal reward theorem accrues rewards at the end of each epoch, and boundedness of expectation of the rewards is sufficient to apply the renewal reward theorem. In the case of partial rewards (which we have in our model), where rewards are accrued during an epoch, stronger conditions are required, which we prove are satisfied.

First, define the sequence of times in which the cargo completely detaches from the microtubule (off) and subsequently reattaches to the microtubule (on),

$$0 = \tau_{\rm on}^0 = \tau_{\rm off}^0 < \tau_{\rm on}^1 < \tau_{\rm off}^1 < \tau_{\rm on}^2 < \tau_{\rm off}^2 < \cdots$$

by

(8) 
$$\tau_{\text{off}}^{k} := \inf\{t > \tau_{\text{off}}^{k} : m(t) = 0\}, \quad k \ge 1, \\ \tau_{\text{off}}^{k} := \inf\{t > \tau_{\text{off}}^{k-1} : m(t) \ge 1\}, \quad k \ge 1.$$

Next, define the sequence of cargo displacements when the cargo is attached to the microtubule (on) and detached from the microtubule (off),

(9) 
$$R_{\text{on}}^k := X(\tau_{\text{off}}^k) - X(\tau_{\text{on}}^k), \quad R_{\text{off}}^k := X(\tau_{\text{on}}^k) - X(\tau_{\text{off}}^{k-1}), \quad k \ge 1,$$

and the corresponding times spent attached or detached,

(10) 
$$T_{\rm on}^k := \tau_{\rm off}^k - \tau_{\rm on}^k, \quad T_{\rm off}^k := \tau_{\rm on}^k - \tau_{\rm off}^{k-1}, \quad k \ge 1$$

It follows directly from the strong Markov property that  $\{(T_{\text{off}}^k + T_{\text{on}}^k, R_{\text{off}}^k + R_{\text{on}}^k)\}_{k\geq 1}$ is an independent and identically distributed (iid) sequence of random variables. In the language of renewal theory,  $\{T_{\text{off}}^k + T_{\text{on}}^k\}_{k\geq 1}$  are the interarrival times and  $\{R_{\text{off}}^k + R_{\text{on}}^k\}_{k\geq 1}$  are the corresponding rewards. Let N(t) be the renewal process that counts the number of arrivals before time  $t \ge 0$ ,

(11) 
$$N(t) := \sup\{k \ge 0 : \tau_{\text{off}}^k \le t\}.$$

Define the reward function, R(t), and the partial reward function, Y(t), by

(12) 
$$R(t) := \sum_{k=1}^{N(t)} \left( R_{\text{on}}^k + R_{\text{off}}^k \right), \qquad Y(t) := X(t) - X\left( \tau_{\text{off}}^{N(t)} \right),$$

and observe that

$$X(t) = R(t) + Y(t)$$

In other words, R(t) describes rewards accrued during past epochs, and Y(t) is the partial reward accrued during the current epoch. We show below that  $\mathbb{E}[|R_{\rm on}+R_{\rm off}|] < \infty$  and  $\mathbb{E}[T_{\rm on} + T_{\rm off}] < \infty$ , and therefore the classical renewal reward theorem [47] ensures that

(13) 
$$\lim_{t \to \infty} \frac{R(t)}{t} = \lim_{t \to \infty} \frac{\mathbb{E}[R(t)]}{t} = \frac{\mathbb{E}[R_{\text{on}}] + \mathbb{E}[R_{\text{off}}]}{\mathbb{E}[T_{\text{on}}] + \mathbb{E}[T_{\text{off}}]} \quad \text{almost surely.}$$

The following theorem verifies that this convergence actually holds for X(t).

THEOREM 1. The following limit holds:

(14) 
$$\lim_{t \to \infty} \frac{X(t)}{t} = \lim_{t \to \infty} \frac{\mathbb{E}[X(t)]}{t} = \frac{\mathbb{E}[R_{on}] + \mathbb{E}[R_{off}]}{\mathbb{E}[T_{on}] + \mathbb{E}[T_{off}]} \quad almost \ surely.$$

To prove this theorem, we need several lemmas. We collect the proofs of these lemmas in Appendix A. The first lemma bounds the probability that the partial reward function Y(t) in (12) is large when the cargo is detached from the microtubule.

LEMMA 2. Define the sequence of iid random variables  $\{Y_{off}^k\}_{k\geq 1}$  by

$$Y_{off}^k := \sup_{t \in [\tau_{off}^{k-1}, \tau_{on}^k]} \left| X(t) - X(\tau_{off}^k) \right|, \quad k \ge 1.$$

Then for any C > 0 and  $k \ge 1$ , we have that

15) 
$$\mathbb{P}(Y_{off}^k \ge C) \le \sqrt{\pi/x}(2x+1)e^{-x}, \quad where \ x = \frac{C}{\sigma}\sqrt{2M\lambda_{on}(0)} > 0.$$

Similarly, the next lemma bounds the probability that the partial reward function is large when the cargo is attached to the microtubule.

LEMMA 3. Define the sequence of iid random variables  $\{Y_{on}^k\}_{k\geq 1}$  by

$$Y_{on}^k := \sup_{t \in [\tau_{on}^k, \tau_{off}^k]} |X(t) - X(\tau_{on}^k)|, \quad k \ge 1.$$

There exists  $\lambda_0 > 0, \lambda_1 > 0$  so that if  $k \ge 1$ , then  $\mathbb{P}(Y_{on}^k \ge C) \le \lambda_0 \sqrt{C} e^{-\lambda_1 \sqrt{C}}$  for all sufficiently large C > 0.

The next lemma uses Lemmas 2 and 3 to prove that the partial reward function gets large only finitely many times.

LEMMA 4. Define the sequence of iid random variables  $\{Y_k\}_{k\geq 1}$  by

(16) 
$$Y_k := \sup_{t \in [\tau_{off}^{k-1}, \tau_{off}^k]} |X(t) - X(\tau_{off}^k)|, \quad k \ge 1.$$

Then

$$\mathbb{P}\left(\lim_{K\to\infty}\bigcup_{k=K}^{\infty}\left\{Y_k>\sqrt{k}\right\}\right)=0.$$

The last lemma checks that the mean of  $Y_k$  in (16) is finite.

LEMMA 5. Define  $\{Y_k\}_{k\geq 1}$  as in (16). Then  $\mathbb{E}[Y_k] < \infty$  for all  $k \geq 1$ .

With these lemmas in place, we are ready to prove Theorem 1.

Proof of Theorem 1. It follows immediately from Lemma 5 that  $\mathbb{E}[|R_{\text{on}}^k + R_{\text{off}}^k]] < \infty$ . Furthermore,  $T_{\text{off}}^k$  is exponentially distributed with rate  $M\lambda_{\text{on}}(0)$ , and the proof of Lemma 3 shows that  $\mathbb{E}[T_{\text{on}}^k] < \mathbb{E}[S]$  for an exponentially distributed random variable S with some rate  $\lambda > 0$ . Hence,  $\mathbb{E}[T_{\text{on}}^k + T_{\text{off}}^k] < \infty$ , and thus (13) holds by a direct application of the classical renewal reward theorem [47].

Therefore, it remains to check that

(17) 
$$\lim_{t \to \infty} \frac{\mathbb{E}\left[X(t) - X\left(\tau_{\text{off}}^{N(t)}\right)\right]}{t} = 0 = \lim_{t \to \infty} \frac{X(t) - X\left(\tau_{\text{off}}^{N(t)}\right)}{t} \quad \text{almost surely.}$$

The first equality in (17) follows immediately from Lemma 5.

To verify the second equality in (17), we note that Lemma 4 ensures that

$$\limsup_{k \to \infty} \frac{Y_k}{\sqrt{k}} \le 1 \quad \text{almost surely.}$$

Therefore,

$$\lim_{t \to \infty} \frac{\left| X(t) - X\left(\tau_{\text{off}}^{N(t)}\right) \right|}{t} \le \lim_{t \to \infty} \frac{|Y_{N(t)}|}{t} \le \lim_{t \to \infty} \frac{\sqrt{N(t)}}{t} = 0 \quad \text{almost surely,}$$

since

$$\lim_{t \to \infty} \frac{N(t)}{t} = \frac{1}{\mathbb{E}[T_{\rm on} + T_{\rm off}]} \quad \text{almost surely}$$

by the strong law of large numbers for renewal processes [47].

Consequently, the position of the cargo does indeed satisfy a classical renewal reward structure with two different types of epochs: bound and unbound, each of which accrue some net displacement.

4. Mathematical analysis of transport ability. With the framework of renewal theory constructed in section 3, we are ready to analyze the transport ability of the model introduced in section 2. To assess the transport ability of the motor cargo ensemble, we analyze the expected *run length*, expected *run time*, and asymptotic *velocity*. We define the run length to be the distance traveled by the cargo from the first time a motor attaches to the cargo until the next time that all motors are detached from the microtubule, which was defined precisely in (9) and denoted by  $R_{\rm on}$ . The run time is the corresponding time spent attached to a microtubule, which was defined precisely in (10) and denoted by  $T_{\rm on}$ . The asymptotic velocity is

(18) 
$$V := \lim_{t \to \infty} \frac{X(t)}{t}.$$

The velocity V includes the time the cargo is both being transported along the microtubule and diffusing while unattached.

Applying Theorem 1, we have that

(19) 
$$V = \frac{\mathbb{E}[R_{\text{on}}] + \mathbb{E}[R_{\text{off}}]}{\mathbb{E}[T_{\text{on}}] + \mathbb{E}[T_{\text{off}}]} \quad \text{almost surely.}$$

Copyright © by SIAM. Unauthorized reproduction of this article is prohibited.

Now,

2518

(20) 
$$\mathbb{E}[R_{\text{off}}] = 0,$$

since the cargo is freely diffusing when no motors are bound, and since motor binding and unbinding is independent of Brownian motion  $\{W(t)\}_{t\geq 0}$ . Furthermore, when all of the M motors are unbound, each motor binds at rate  $\lambda_{on}(0)$ . Hence,

(21) 
$$\mathbb{E}[T_{\text{off}}] = (M\lambda_{\text{on}}(0))^{-1}$$

It therefore remains to calculate two of the three quantities, V,  $\mathbb{E}[R_{\text{on}}]$ , and  $\mathbb{E}[T_{\text{on}}]$ , since the third is given by (19). We calculate  $\mathbb{E}[T_{\text{on}}]$  first since it is the simplest, as it is a mean first passage time of a continuous-time Markov chain.

**4.1. Expected run time.** As we noted in section 2.1, the number of motors bound m(t) is itself the Markov process (6). To compute the expected run time, we compute the mean time for m(t) to reach state m = 0 starting from m(0) = 1.

Let  $\widetilde{Q} \in \mathbb{R}^{(M+1)\times(M+1)}$  be the generator of the Markov chain m(t) in (6). That is, the (i, j)-entry of  $\widetilde{Q}$  gives the rate that m(t) jumps from state i to state  $j \neq i$ , and the diagonal entries are chosen so that  $\widetilde{Q}$  has zero row sums. Let  $Q \in \mathbb{R}^{M \times M}$ be the matrix obtained from deleting the first row and column of  $\widetilde{Q}$ . The matrix Qis tridiagonal, with the *m*th row containing subdiagonal, diagonal, and superdiagonal entries  $m\lambda_{\text{off}}(m), -(m\lambda_{\text{off}}(m) + (M - m)\lambda_{\text{on}}(m)), (M - m)\lambda_{\text{on}}(m)$ , respectively. The expected run time  $\mathbb{E}[T_{\text{on}}]$  is (by Theorem 3.3.3 in [40])

(22) 
$$\mathbb{E}[T_{\text{on}}] = \mathbf{1}^T \mathbf{t}, \quad \text{where} \quad Q^T \mathbf{t} = -\mathbf{e}_1$$

where  $\mathbf{1} \in \mathbb{R}^M$  is the vector of all 1's and  $\mathbf{e}_1 \in \mathbb{R}^M$  is the standard basis vector.

**4.2. Decomposing stochasticity.** Having calculated  $\mathbb{E}[T_{\text{on}}]$  in (22), we can determine V by determining  $\mathbb{E}[R_{\text{on}}]$  (or vice versa). Two key steps allow us to analyze V and  $\mathbb{E}[R_{\text{on}}]$ : (i) we average over the diffusive dynamics while conditioning on a realization of the jump dynamics, and (ii) we take advantage of a timescale separation between the relaxation rate of the cargo dynamics and the jump rate of the motor dynamics.

**4.2.1. Conditioning on jump realizations.** Observe that the stochasticity in the model can be separated into a *continuous* diffusion part and a *discrete* part controlling motor binding, unbinding, and stepping. Mathematically, the continuous diffusion part is described by the Brownian motion W in (3), and the discrete motor state is described by the Markov jump process **J**. We first average over the diffusion by defining the conditional expectations

23) 
$$x(t) := \mathbb{E}[X(t)|\mathbf{J}], \quad z_i(t) := \mathbb{E}[Z_i(t)|\mathbf{J}], \quad t \ge 0, \ i \in \{1, \dots, M\}.$$

We emphasize that (23) are averages over paths of W given a realization **J**. Thus,  $\{x(t)\}_{t\geq 0}$  and  $\{\{z_i(t)\}_{t\geq 0}\}_{i=1}^M$  are functions of the realization **J**. This definition is convenient, because while X(t) follows the randomly switching SDE (3), the process x(t) follows a randomly switching ODE, whose solution is known explicitly.

PROPOSITION 6. For each t > 0, the expected cargo position x(t) conditioned on a realization of the jump process satisfies

(24) 
$$\frac{d}{dt}x(t) = \varepsilon^{-1} \sum_{i=1}^{M} \left( z_i(t) - x(t) \right) \quad almost \ surrely$$

*Proof.* Using the explicit solution of an OU process, we have that

(25) 
$$X(t) = X(\tau)e^{-\theta(t-\tau)} + \mu(1 - e^{-\theta(t-\tau)}) + \mathcal{M},$$

where  $\tau$  is the most recent jump time of **J**,

$$\tau = \sup \{\{0\} \cup \{s < t : \mathbf{J}(s-) \neq \mathbf{J}(s+)\}\},\$$

 $\theta = m(\tau)\varepsilon^{-1}, \ \mu = \frac{1}{m(\tau)}\sum_{i\in I(\tau)}Z_i(\tau), \ \text{and } \mathcal{M} \ \text{satisfies } \mathbb{E}[\mathcal{M}|\mathbf{J}] = 0.$  We have used the notation  $f(t\pm) := \lim_{s\to t\pm}f(s)$ . Hence, taking the expectation of (25) conditioned on  $\mathbf{J}$  yields

$$\begin{aligned} x(t) &= \mathbb{E}\left[X(\tau)e^{-\theta(t-\tau)}|\mathbf{J}\right] + \mathbb{E}\left[\mu\left(1-e^{-\theta(t-\tau)}\right)|\mathbf{J}\right] \\ &= e^{-\theta(t-\tau)}\mathbb{E}[X(\tau)|\mathbf{J}] + \left(1-e^{-\theta(t-\tau)}\right)\frac{1}{m(\tau)}\sum_{i\in I(\tau)}\mathbb{E}[Z_i(\tau)|\mathbf{J}], \end{aligned}$$

since  $\tau$  are  $\{m(s)\}_{s\geq 0}$  measurable with respect to the  $\sigma$ -algebra generated by **J**.

4.2.2. Separation of timescales. We next make an observation of disparate timescales. After averaging over the diffusive noise W, the model effectively depends on two timescales: the relaxation time of the continuous dynamics (24) (characterized by the dimensionless rate  $\varepsilon^{-1}$ ) and the switching times of the discrete motor dynamics (5) (characterized by the dimensionless rates  $\lambda_{\rm on}, \lambda_{\rm step}, \lambda_{\rm off}$ ). Even for conservative parameter estimates, the continuous timescale is much faster than the discrete switching timescale. For instance, suppose a motor exerts a Hookean force with stiffness k = 0.5 pN/nm [16] on a spherical cargo with radius  $r = 1 \ \mu \text{m}$  in cytosol with viscosity  $\eta$  equal to that of water. It follows that  $k/(6\pi\eta r) \approx 3 \times 10^4 \text{ s}^{-1}$ , whereas  $k_{\text{off}}(1)$  is on the order of  $10^{-1}$  to  $10^1 \text{ s}^{-1}$  [16]. Hence,

(26) 
$$\varepsilon := k_{\text{off}}(1)\gamma/k \approx 3 \times 10^{-4} \ll 1.$$

Further,  $\lambda_{\text{on}}$ ,  $\lambda_{\text{step}}$ ,  $\lambda_{\text{off}}$  are roughly order one since  $k_{\text{on}}$ ,  $k_{\text{step}}$ ,  $k_{\text{off}}$  have similar orders of magnitude [16].

Therefore, compared to the switching timescale, x(t) quickly relaxes to an equilibrium between motor switches. Furthermore, we are interested in studying  $\mathbb{E}[R_{\text{on}}]$  and V, which depend on the behavior of x(t) over the course of several motor switches. Hence, we approximate x(t) by a jump process  $\overline{x}(t)$  obtained from assuming x(t) immediately relaxes to its equilibrium after each motor switch.

More precisely, let  $(\overline{x}(t), \overline{z}_1(t), \dots, \overline{z}_M(t)) \in \mathbb{R}^{M+1}$  be a **J**-measurable, rightcontinuous process,

$$\overline{x}(t) = \overline{x}(t+), \quad \overline{z}_i(t) = \overline{z}_i(t+), \quad t \ge 0, \ i \in \{1, \dots, M\},$$

with  $\overline{x}(0) = x(0)$  and  $\overline{z}_i(0) = z_i(0)$ ,  $i \in \{1, \ldots, M\}$ , that evolves in the following way. In light of (7), we define the effective motor positions by how they are modified through the jump process, binding at  $\tau_i$  and then incrementing from stepping, or staying unbound at the cargo position  $\overline{x}(t)$ ,

(27) 
$$\overline{z}_i(t) = (\overline{x}(\tau_i(t)) + J_i(t)) \mathbf{1}_{\{J_i(t) \neq u\}} + \overline{x}(t) \mathbf{1}_{\{J_i(t) = u\}}, \quad i \in \{1, \dots, M\}.$$

Due to the assumed fast relaxation,  $\overline{x}(t)$  only changes when a motor steps or unbinds, as newly bound motors exert no force. That is, if  $J_i(t+) = J_i(t-)$  for all  $i \in \{1, \ldots, M\}$  satisfying  $J_i(t-) \ge 0$ , then  $\overline{x}(t-) = \overline{x}(t+)$ . Otherwise,  $\overline{x}(t)$ evolves according to the following two rules, which describe how the cargo position  $\overline{x}(t)$  changes when a motor steps or unbinds.

- 1. If the *i*th motor steps  $(J_i(t-) = j \ge 0 \text{ and } J_i(t+) = j+1)$ , then  $\overline{x}(t+) = \overline{x}(t-) + 1/m(t)$ .
- 2. If the *i*th motor unbinds  $(J_i(t-) = j \ge 0 \text{ and } J_i(t+) = u)$ , then  $\overline{x}(t+) = \overline{x}(t-) + \Delta_{i,(\overline{z}_1,\ldots,\overline{z}_{m(t-)})}$ , where  $(\overline{z}_1,\ldots,\overline{z}_{m(t-)})$  gives the positions of the m(t-) bound motors just before time t, and

(28) 
$$\Delta_{i,(\overline{z}_1,\dots,\overline{z}_{m(t-)})} = \frac{1}{m(t-)-1} \sum_{i'=1,i'\neq i}^{m(t-)} \overline{z}_{i'}(t-) - \frac{1}{m(t-)} \sum_{i'=1}^{m(t-)} \overline{z}_{i'}(t-).$$

In other words, if either of these events occurs, the cargo position  $\overline{x}(t)$  relaxes to the mean position of the motors. These two rules describe how the mean motor position changes in the two scenarios. If a single motor steps, incrementing its position by 1, the mean motor position increases by 1/m(t). If a motor unbinds, (28) is the change in the mean motor position from removing that motor.

It follows from these two evolution rules for  $\overline{x}(t)$  that

(29) 
$$\overline{x}(t) = \sum_{m=1}^{M} \frac{1}{m} S_m(t) + \chi(t),$$

where  $S_m(t)$  is the number of steps taken when m motors are bound before time t (each of which modifies the position by 1/m), and  $\chi(t)$  accounts for changes in the cargo position that result from a motor unbinding,

(30) 
$$\chi(t) = \sum_{k=1}^{N_{\text{off}}(t)} \Delta_{j_k, (\bar{z}_1(s_{\text{off}}^k -), \dots, \bar{z}_{m(s_{\text{off}}^k -)})}(s_{\text{off}}^k -)$$

where  $0 = s_{off}^0 < s_{off}^1 < \cdots$  is the sequence of times in which a motor unbinds,

$$s_{\text{off}}^k := \inf \{ t > s_{\text{off}}^{k-1} : J_i(t-) \ge 0 \text{ and } J_i(t) = u \text{ for some } i \in \{1, \dots, M\} \}, k \ge 1,$$

and  $N_{\text{off}}(t) := \sup\{k \ge 0 : s_{\text{off}}^k \le t\}$  is the number of unbindings before time  $t \ge 0$ , and  $j_k \in \{1, \ldots, M\}$  gives the (almost surely unique) index of the motor that unbinds at time  $s_{\text{off}}^k$ . That is,  $j_k$  satisfies  $J_{j_k}(s_{\text{off}}^k -) \ne J_{j_k}(s_{\text{off}}^k) = u$ .

The following proposition checks that x(t) converges almost surely to the jump process  $\overline{x}(t)$  as  $\varepsilon \to 0$ . The proof is in Appendix B.

PROPOSITION 7. If  $T \ge 0$  is an almost surely finite stopping time with respect to  $\{J(t)\}_{t>0}$ , then

$$\lim_{x \to 0} x(T) = \overline{x}(T) \quad almost \ surely.$$

From this proposition, we conclude that studying the mean behavior of the cargo position X(t) can ultimately be reduced to studying the jump process  $\overline{x}(t)$ , where the jumps correspond to motor stepping and unbinding events.

**4.3. Run length and velocity.** Since  $\varepsilon \ll 1$  for biologically relevant parameters, we investigate the run length and velocity of X(t) by investigating the analogous quantities for  $\overline{x}(t)$ ,

(31) 
$$\overline{R} := \overline{x}(\tau_{\text{off}}^1) - \overline{x}(\tau_{\text{on}}^1), \qquad \overline{V} := \lim_{t \to \infty} \frac{\overline{x}(t)}{t}.$$

**4.3.1. Run length.** The following proposition checks that the mean run length of the full process X(t) converges to the mean run length of the jump process  $\overline{x}(t)$  as  $\varepsilon \to 0$ .

PROPOSITION 8.  $\mathbb{E}[R_{on}] \to \mathbb{E}[\overline{R}]$  as  $\varepsilon \to 0$ .

*Proof.* By the tower property of conditional expectation (see Theorem 5.1.6 in [10]), we have that

$$\mathbb{E}[R_{\rm on}] = \mathbb{E}[\mathbb{E}[R_{\rm on}|\mathbf{J}]] = \mathbb{E}[\mathbb{E}[X(\tau_{\rm off}^1) - X(\tau_{\rm on}^1)|\mathbf{J}]] = \mathbb{E}[x(\tau_{\rm off}^1) - x(\tau_{\rm on}^1)].$$

Now, Proposition 7 ensures that

(32) 
$$x(\tau_{\text{off}}^1) - x(\tau_{\text{on}}^1) \to \overline{R} \text{ almost surely as } \varepsilon \to 0.$$

Let  $N \geq 0$  be the number of steps taken between time  $\tau_{\text{on}}^1$  and time  $\tau_{\text{off}}^1$ . Since motors take steps of distance one, we have the almost sure bound,  $x(\tau_{\text{off}}^1) - x(\tau_{\text{on}}^1) \leq N$ . Steps are taken at Poisson rate  $m(t)\lambda_{\text{step}}(m(t)) \leq M\Lambda$ , where  $\Lambda := \max_{m \in \{1,...,M\}} \lambda_{\text{step}}(m)$ . Thus  $\mathbb{E}[N] \leq \Lambda M \mathbb{E}[\tau_{\text{off}}^1 - \tau_{\text{on}}^1] < \infty$ . Thus, (32) and the bounded convergence theorem complete the proof.

**4.3.2.** Velocity. Let us now investigate  $\overline{V}$  in (31), observing that this quantity can be approached in two ways. The first exploits the observation that nonzero mean displacements only occur from motor stepping, so the velocity can be interpreted as the product of how often a step occurs with m motors and the size of the displacement. The second approach is again a renewal reward argument, noting that the only nonzero displacements occur during epochs of bound cargo. The connection between these two approaches provides explicit relationships between the velocity, run lengths, and run times.

Recalling the decomposition of the jump process  $\overline{x}$  in (29), we seek to compute the expected value

$$\mathbb{E}[\overline{x}(t)] = \sum_{m=1}^{m} \frac{1}{m} \mathbb{E}[S_m(t)] + \mathbb{E}[\chi(t)].$$

Using the definition of  $\chi(t)$  in (30), we compute its expectation by summing over all possible displacements from one of m motors unbinding  $\Delta_{j,(\bar{z}_1(t-),...,\bar{z}_{m(s_{off}^k-)}(t-))}$ , which yields

$$\sum_{j=1}^{m} \Delta_{j,(z_1,\dots,z_m)} = \frac{1}{m-1} \sum_{j=1}^{m} \left( -z_j + \sum_{i=1}^{m} z_i \right) - \sum_{i=1}^{m} z_i = 0.$$

Since each of the bound motors is equally likely to unbind, it follows that  $\mathbb{E}[\chi(t)] = 0$ . This can be interpreted as the observation that the arithmetic mean does not change in expectation when removing a randomly (uniformly) chosen element. In other words, the effects of motors unbinding ahead of the cargo are completely offset in the mean by motors unbinding behind the cargo. Therefore, the only long-term influence on  $\overline{x}$  is stepping events.

Given a realization  $\{m(s)\}_{s\geq 0}$ , the number of steps taken with m motors bound before time  $t\geq 0$  is Poisson distributed with mean  $m\lambda_{\text{step}}(m)\int_0^t \mathbf{1}_{m(s)=m} \,\mathrm{d}s$ . Hence,

$$\mathbb{E}[S_m(t)] = m\lambda_{\text{step}}(m)\mathbb{E}\left[\int_0^t \mathbf{1}_{m(s)=m} \,\mathrm{d}s\right].$$

Now,  $\{m(s)\}_{s\geq 0}$  is an ergodic Markov process, so the occupation measure converges almost surely to the stationary measure (see Theorem 3.8.1 in [40])

$$\frac{1}{t} \sum_{n=1}^{M} \int_{0}^{t} \mathbf{1}_{m(s)=m} \, \mathrm{d}s \to p_{m} \quad \text{almost surely as } t \to \infty,$$

where  $p_m := \lim_{t\to\infty} \mathbb{P}(m(t) = m)$  is the stationary probability that m motors are bound. We note that  $p_m$  is the (m+1)st component of the unique probability vector,  $\mathbf{p} \in \mathbb{R}^{1 \times (M+1)}$  satisfying (see Theorem 3.5.2 in [40])

$$\mathbf{p}\widetilde{Q}=0,$$

2522

where  $\widetilde{Q} \in \mathbb{R}^{(M+1)\times(M+1)}$  is the generator matrix defined in section 4.1. Since the occupation measure is bounded above by one, the bounded convergence theorem gives

$$\lim_{t \to \infty} \frac{\mathbb{E}[\overline{x}(t)]}{t} = \lim_{t \to \infty} \sum_{m=1}^{M} \lambda_{\text{step}}(m) \mathbb{E}\left[\frac{1}{t} \int_{0}^{t} \mathbf{1}_{m(s)=m} \, \mathrm{d}s\right] = \sum_{m=1}^{M} \lambda_{\text{step}}(m) p_{m}.$$

It is easy to see that the classical renewal reward theorem applies to  $\overline{x}(t)$  so that

$$\sum_{m=1}^{M} \lambda_{\text{step}}(m) p_m = \lim_{t \to \infty} \frac{\mathbb{E}[\overline{x}(t)]}{t} = \lim_{t \to \infty} \frac{\overline{x}(t)}{t} = \frac{\mathbb{E}[\overline{R}]}{\mathbb{E}[T_{\text{on}}] + \mathbb{E}[T_{\text{on}}]} \quad \text{almost surely}$$

Furthermore, (19), (20), (21), and Proposition 8 yield

$$\lim_{\varepsilon \to 0} V = \lim_{\varepsilon \to 0} \frac{\mathbb{E}[R_{\rm on}]}{(M\lambda_{\rm on}(0))^{-1} + \mathbb{E}[T_{\rm on}]} = \frac{\mathbb{E}[\overline{R}]}{(M\lambda_{\rm on}(0))^{-1} + \mathbb{E}[T_{\rm on}]} = \sum_{m=1}^{M} \lambda_{\rm step}(m) p_m.$$

In summary, we now have explicit formulas for the velocity V and expected run length  $\mathbb{E}[R_{\text{on}}]$  of X(t) in the small  $\varepsilon$  limit,

(34) 
$$\lim_{\varepsilon \to 0} V = \overline{V} = \sum_{m=1}^{M} \lambda_{\text{step}}(m) p_m$$

(35) 
$$\lim_{\varepsilon \to 0} \mathbb{E}[R_{\rm on}] = \mathbb{E}[\overline{R}] = \left( \left( M\lambda_{\rm on}(0) \right)^{-1} + \mathbb{E}[T_{\rm on}] \right) \sum_{m=1}^{M} \lambda_{\rm step}(m) p_m,$$

where  $p_m$  is given by (33) and  $\mathbb{E}[T_{\text{on}}]$  is given by (22). In Figure 2, we compare these formulas for  $\mathbb{E}[\overline{R}]$  and  $\overline{V}$  with estimates of  $\mathbb{E}[R_{\text{on}}]$  and V from simulations of the full process  $(X(t), \mathbf{Z}(t), \mathbf{J}(t))$  (for details on our statistically exact simulation method, see section 4.5).

Furthermore, some experimental works [16, 26] measure the average run velocity,  $\mathbb{E}[\overline{R}/T_{\text{on}}]$ . Now, if  $\sigma(m)$  denotes the  $\sigma$ -algebra generated by  $\{m(t)\}_{t\geq 0}$ , then recalling (8) and (10) and using the tower property of conditional expectation yields

$$\mathbb{E}\left[\frac{\overline{R}}{T_{\text{on}}}\right] = \mathbb{E}\left[\frac{1}{T_{\text{on}}}\mathbb{E}[\overline{R}|\sigma(m)]\right] = \mathbb{E}\left[\frac{1}{T_{\text{on}}^{1}}\sum_{m=1}^{M}\frac{1}{m}\mathbb{E}[S_{m}(\tau_{\text{off}}^{1})|\sigma(m)]\right]$$
$$= \sum_{m=1}^{M}\frac{1}{m}m\lambda_{\text{step}}(m)\mathbb{E}\left[\frac{1}{T_{\text{on}}^{1}}\int_{0}^{\tau_{\text{off}}^{1}}1_{m(s)=m}\,\mathrm{d}s\right].$$

Hence, it follows from (34) that

(36) 
$$\mathcal{V} := \mathbb{E}[\overline{R}/T_{\rm on}] = \overline{V}/p_{\rm on}$$

where  $p_{\text{on}} = \sum_{m=1}^{M} p_m$  is the stationary probability that  $m(t) \ge 1$ .

NONPROCESSIVE MOTOR TRANSPORT

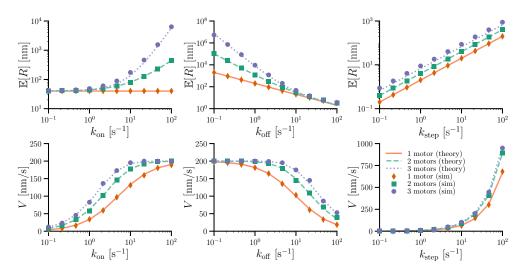


FIG. 2. Expected run lengths  $\mathbb{E}[R]$  and asymptotic velocities V as a function of the parameters  $k_{on}, k_{off}, k_{step}$  for M = 1, 2, 3 total motors. The curves are the analytical formulas (34)–(35) for the  $\varepsilon \to 0$  limit, and the dots are estimates from statistically exact realizations of the full process,  $\{(X(s), \mathbf{Z}(s), \mathbf{J}(s))\}_{s=0}^{t}$ , where the ending time t is such that  $N(t) = 10^{5}$ , where N(t) is defined in (11). Unless noted otherwise,  $k_{on}(m) = 10 [s^{-1}], k_{step}(m) = 20 [s^{-1}], k_{off}(m) = 5 [s^{-1}]$  for each m. Further, k and  $\gamma$  are as in (26), and  $k_BT = 4.1 [pN \cdot nm]$ .

**4.4.** Cases M = 1, M = 2, and M = 3. In this subsection, we collect explicit formulas for the run length  $\mathbb{E}[\overline{R}]$  and run velocity  $\mathcal{V}$  when the total number of motors is M = 1, 2, 3. The run time  $\mathbb{E}[T_{\text{on}}]$  and net velocity V can be easily deduced from these quantities using (34)–(35) but are omitted for brevity.

For M = 1 total motors, the quantities are simply

(37) 
$$\mathbb{E}[\overline{R}] = \lambda_{\text{step}}(1), \quad \mathcal{V} = \lambda_{\text{step}}(1).$$

For M = 2 total motors, we find

(38) 
$$\mathbb{E}[\overline{R}] = \lambda_{\text{step}}(1) + \frac{\lambda_{\text{on}}(1)\lambda_{\text{step}}(2)}{2\lambda_{\text{off}}(2)}, \quad \mathcal{V} = \frac{2\lambda_{\text{off}}(2)\lambda_{\text{step}}(1) + \lambda_{\text{on}}(1)\lambda_{\text{step}}(2)}{2\lambda_{\text{off}}(2) + \lambda_{\text{on}}(1)}.$$

For M = 3 total motors, we find

(39)  
$$\mathbb{E}[\overline{R}] = \lambda_{\text{step}}(1) + \frac{\lambda_{\text{on}}(1)(3\lambda_{\text{off}}(3)\lambda_{\text{step}}(2) + \lambda_{\text{on}}(2)\lambda_{\text{step}}(3))}{3\lambda_{\text{off}}(2)\lambda_{\text{off}}(3)},$$
$$\mathcal{V} = \frac{3\lambda_{\text{off}}(3)\left[\lambda_{\text{off}}(2)\lambda_{\text{step}}(1) + \lambda_{\text{on}}(1)\lambda_{\text{step}}(2)\right] + \lambda_{\text{on}}(1)\lambda_{\text{on}}(2)\lambda_{\text{step}}(3)}{3\lambda_{\text{off}}(3)\left[\lambda_{\text{off}}(2) + \lambda_{\text{on}}(1)\right] + \lambda_{\text{on}}(1)\lambda_{\text{on}}(2)}$$

To put these quantities in dimensional units, recall the jump rates (4) and multiply  $\mathbb{E}[\overline{R}]$  by the dimensional step distance  $\delta > 0$  and multiply  $\mathcal{V}$  by  $\delta k_{\text{off}}(1) > 0$ .

**4.5.** Numerical simulations. To verify our predictions for the expected run lengths and velocities, we compare them to statistically exact numerical simulations of the full process  $(X(t), \mathbf{Z}(t), \mathbf{J}(t))$ . In a given state, we use the classical Gillespie stochastic simulation algorithm to generate the time of the next transition for the Markov chain  $\mathbf{J}(t)$  and to choose which transition occurs. For  $m(t) \geq 1$ , X(t) is an OU process, generically described by

Copyright © by SIAM. Unauthorized reproduction of this article is prohibited.

$$dX(t) = \alpha \left[ \mu - X(t) \right] dt + \beta dW(t).$$

To update X(t) to the next time  $t + \tau$ , we use the statistically exact method described in [18], summarized by

(40) 
$$X(t+\tau) = e^{-\alpha\tau}X(t) + (1 - e^{-\alpha\tau})\mu + \beta\sqrt{\frac{(1 - e^{-2\alpha\tau})}{2\alpha}}\,\mathfrak{n}$$

where **n** is a standard normal random variable. When m(t) = 0, X(t) is a pure diffusion process with  $\alpha = 0$ , so (40) becomes an Euler-Maruyama update. This procedure generates statistically exact sample paths of X(t), sampled at the transition times of  $\mathbf{J}(t)$ . We use this scheme to generate a long realization of  $(X(t), \mathbf{Z}(t), \mathbf{J}(t))$ , thereby providing Monte Carlo estimates for  $\mathbb{E}[R_{\text{on}}]$  and V for a given parameter set.

5. Biological application. We now use the formulas (37)–(39) for run velocity,  $\mathcal{V}$ , and run length,  $\mathbb{E}[\overline{R}]$ , to explore the behavior of nonprocessive motors. The behavior of individual nonprocessive motors is characterized by two observations: (i) short attachment times and (ii) the time it takes to hydrolyze ATP (and consequently, to step) coincides with this attachment time [6, 15, 39]. Concretely, Ncd motors in the kinesin-14 family take 1 to 5 steps before unbinding [2, 11]. In our model,  $\lambda_{\text{step}}(1)$  gives the expected number of steps before unbinding, so we characterize nonprocessive motors as those with  $\lambda_{\text{step}}(1) \in [1, 5]$ .

Using this characterization, we explore the observation made in [16, 26, 48] that nonprocessive motors in the kinesin-14 family cooperate to produce long-range transport. This behavior is reported in [16, 26] in terms of a velocity that is analogous to the run velocity  $\mathcal{V}$  in our model. Specifically, the primary manifestation of cooperativity is that  $\mathcal{V}$  increases substantially when the total number of motors increases from M = 1 to M = 2. For  $M \geq 2$ , the velocity remains relatively constant.

We thus ask the question, what features are necessary to produce this behavior? Now, if the step rate is independent of the number of bound motors, m, then it follows immediately from (34) and (36) that  $\mathcal{V}$  is independent of M. In particular, if the dimensional step rate is  $k_{\text{step}}(m) \equiv k_0$  for all  $m \in \{1, \ldots, M\}$ , then in dimensional units,  $\mathcal{V}$  is simply  $\delta k_0$ , regardless of any other parameter values.

Therefore, our model predicts that the stepping rate must depend on the number of bound motors in order to produce the cooperative behavior seen in run velocities in [16, 26]. This prediction is bolstered by the simulation results of [16]. There, the authors constructed a detailed computational model of motor transport, and they had to improve motor stepping ability when two or more motors are bound in order for simulations of their computational model to match experimental run velocities.

The authors of [16] also describe motor cooperativity in terms of the average distance traveled by a cargo before all of its motors detach from a microtubule, which is analogous to  $\mathbb{E}[\overline{R}]$  in our model. Namely, they find that the run length  $\mathbb{E}[\overline{R}]$  dramatically increases when M increases from 1 to 2. Our model can replicate this cooperativity if and only if we allow the binding rate,  $k_{\text{on}}$ , and/or the unbinding rate,  $k_{\text{off}}$ , to depend on the number of bound motors, m.

To illustrate, we find the parameter values needed for our model to match the measurements from [16]. However, we emphasize the qualitative results rather than the precise quantitative values of our parameters. Indeed, there are issues preventing an exact comparison of our model with the data in [16]. For example, as the authors point out, the length of the microtubules sometimes caused run lengths to be significantly altered (see Figures 1 and S6 in [16]). Furthermore, for a single motor

(M = 1), the authors report average run lengths of approximately 300 [nm], and they note that this value is necessarily an overestimate since they were unable to measure very short runs. Furthermore, this value must also be an overestimate since a single nonprocessive motor takes only a few steps per run (by definition of nonprocessive), and each step is approximately 7 [nm] [11].

We thus assume that  $\lambda_{\text{step}}(1) = 4$ , based on [2, 11] and  $\delta = 7$  [nm]. This gives  $\mathbb{E}[R] = 28$  [nm] for M = 1, which we use instead of the reported value in [16]. We then match the respective approximate run lengths of 1300 [nm] and 3300 [nm] for M = 1, 2 and the respective approximate run velocities of 100, 150, and 150 [nm/s] for M = 1, 2, 3 reported in [16]. Using the formulas in (37)–(39), this uniquely determines the stepping rates,  $k_{\text{step}}(1) \approx 14$  [s<sup>-1</sup>] and  $k_{\text{step}}(2) \approx k_{\text{step}}(3) \approx 21$  [s<sup>-1</sup>], and the unbinding rate  $k_{\text{off}}(1) \approx 3.5$  [s<sup>-1</sup>], which are all within the range of previously reported rates. The other binding/unbinding rates are not uniquely specified but rather must satisfy the relations  $k_{\text{off}}(2) \approx 0.02k_{\text{on}}(1)$  and  $k_{\text{off}}(3) \approx 1.1k_{\text{on}}(2)$ . Hence, if  $k_{\text{off}}$  were constant in m, then  $k_{\text{on}}(1) \approx 200$  [s<sup>-1</sup>] and  $k_{\text{on}}(2) \approx 3$  [s<sup>-1</sup>].

We make two observations about this result: (i) the binding rate  $k_{\rm on}(1)$  is an order of magnitude larger than reported values in [2, 16], and (ii) the binding rate decreases as the number of bound motors increases from 1 to 2. Both of these points can be explained by geometry. First, the value of  $k_{\rm on}(1)$  is enhanced because the single bound motor tethers the unbound motors close to the microtubule and thus allows those motors to bind more easily. This binding enhancement due to geometry has precedent in motor studies. Indeed, in a different family of kinesins, it was shown to be critical for determining run lengths; see [13]. Further, it was shown to play a critical role in enabling dynein processivity (see [20]), and it was posited as an explanation for why myosin motors can become processive when processive kinesin motors are present; see [22]. The authors in [3] report large  $k_{\rm on}$  values in a model of microtubule sliding driven by kinesin-14 motors and also speculate that this is due to tethering effects.

This effect can also be understood in terms of rebinding. If two motors are bound and one unbinds, then that motor can rapidly rebind since the bound motor keeps it near the microtubule. Such rebinding was the mechanism posited in [17] to explain the processive behavior of nonprocessive motors along microtubule bundles. Further, rebinding is very important in enzymatic reactions [46, 19, 33, 34]. In that context, one incorporates rebinding by using an "effective" unbinding rate, which is the intrinsic unbinding rate multiplied by the probability that the particle does not rapidly rebind; see [32]. Hence, this effect could be included in our model by reducing  $k_{\text{off}}(2)$  rather than (or in addition to) increasing  $k_{\text{on}}(1)$ . Importantly, this is exactly what is implied by the relation,  $k_{\text{off}}(2) \approx 0.02k_{\text{on}}(1)$ , derived above.

Second, geometric exclusion effects can explain a decrease in binding rates as the number of bound motors increases from 1 to 2. When more motors are bound, it is more difficult for additional motors to bind because the range of diffusive search is reduced for unbound motors. In numerical investigations of motor transport systems, this exact effect is observed; see [30, 35]. Furthermore this decrease in binding rate can arise due to motors competing for binding sites, a point posited in [29]. Interestingly, these authors find that negative cooperativity has little impact on transport velocity. The same is true in our model, as the value of  $\mathcal{V}$  changes by less than 1 [nm/s] as  $k_{\rm on}(2)$  ranges from 0 to  $\infty$  while keeping the other parameters fixed. However, we note that the run length for M = 3 is greatly affected by  $k_{\rm on}(2)$ , and thus this highlights the importance of using both run velocity and run length to study motor transport.

6. Discussion. In this work, we formulated and analyzed a mathematical model of transport by nonprocessive molecular motors. We deliberately made our model simple enough to enable us to extract explicit formulas for experimentally relevant quantities yet also maintain agreement with detailed computational studies. One such simplification is to assume the motor stepping and unbinding rates are independent of force. The justification for this assumption is that since nonprocessive motors take only a few steps before unbinding (compared to hundreds of steps by processive motors), these motors are unlikely to be stretched long distances and therefore are unlikely to generate large forces. This assumption on the stepping rate has been made in other models involving nonprocessive motors [39] and did not appear to be a necessary feature in that context, and how force affects stepping is not completely clear [41]. However, we note that force-dependent unbinding can be an important characteristic of processive motors, as kinesin-1 and kinesin-2 detach more rapidly under assisting loads than under opposing loads [1], which increases run velocity and run length.

These limitations notwithstanding, our model makes some concrete predictions about motor number-dependent stepping, binding, and unbinding behavior and how these quantities contribute to transport by nonprocessive motors. Specifically, we observe that a complex cooperativity mechanism appears to be a necessary ingredient for nonprocessive motor transport, and these predictions align with several recent experimental and computational works. Furthermore, these predictions can be investigated experimentally. Indeed, we hope that the work here will spur further investigation into how geometry affects nonprocessive motor transport, especially given that kinesin-14 motors are known to transport a wide variety of cargo, including long, cylindrical microtubules [21, 14] and large, spherical vesicles in plants [48].

## Appendix A. Proofs of Lemmas 2–5.

Proof of Lemma 2. Between time  $\tau_{\text{off}}^{k-1}$  and  $\tau_{\text{on}}^{k}$ , the cargo is freely diffusing. Therefore, to control  $Y_{\text{off}}^{k}$ , we need to control the supremum of a Brownian motion. Now, for any fixed T > 0 and C > 0, it follows from Doob's martingale inequality (see Theorem 3.8(i) in [28]) and symmetry of Brownian motion that

(41) 
$$\mathbb{P}\left(\sup_{t\in[0,T]}|W(t)|\geq C\right)\leq 2\exp\left(\frac{-C^2}{2T}\right).$$

Hence, it follows that

(42) 
$$\mathbb{P}\left(Y_{\text{off}}^k \ge C | T_{\text{off}}^k\right) \le 2 \exp\left(\frac{-C^2}{2\sigma^2 T_{\text{off}}^k}\right) \quad \text{almost surely.}$$

Note that (42) is an average over realizations of the diffusion W for fixed realizations of the time  $T_{\text{off}}^k$ . That is, the inequality holds for almost all realizations of  $T_{\text{off}}^k$ .

Now,  $T_{\text{off}}^k$  is exponentially distributed with rate  $M\lambda_{\text{on}}(0)$ . Hence, the tower property of conditional expectation (see Theorem 5.1.6 in [10]) yields

$$\mathbb{P}(Y_{\text{off}}^k \ge C) = \mathbb{E}[\mathbb{P}(Y_{\text{off}}^k \ge C | T_{\text{off}}^k)] \le 2M\lambda_{\text{on}}(0) \int_0^\infty \exp\left(\frac{-C^2}{2\sigma^2 t} - M\lambda_{\text{on}}(0)t\right) \,\mathrm{d}t.$$

Now, we have that

(44) 
$$\int_0^\infty \lambda e^{-\lambda t} e^{-a/t} \, \mathrm{d}t = 2\sqrt{a\lambda} K_1(2\sqrt{a\lambda}) \quad \text{if } a > 0, \, \lambda > 0,$$

where  $K_1(x)$  denotes the modified Bessel function of the second kind. Hence, the proof is complete after combining (43) and (44) and the following bound,

$$K_1(x) \le \sqrt{\pi/x}(1+1/(2x))e^{-x}, \quad x > 0,$$

which was proven in [49].

Proof of Lemma 3. To control  $Y_{\text{on}}^k$ , we note that if  $t \in [\tau_{\text{on}}^k, \tau_{\text{off}}^k]$ , then X(t) is an OU process centered at  $\mu(t) = \frac{1}{m(t)} \sum_{i \in I(t)} Z_i(t)$  with relaxation rate  $\theta(t) = \varepsilon^{-1} m(t)$ . Hence, after shifting time and space so that  $\tau_{\text{on}}^k = 0$  and  $X(\tau_{\text{on}}^k) = 0$ , we have that

(45) 
$$X(t) = \int_0^t \mu(s)\theta(s)e^{-\int_s^t \theta(s')\,\mathrm{d}s'}\,\mathrm{d}s + \sigma \int_0^t e^{-\int_s^t \theta(s')\,\mathrm{d}s'}\,\mathrm{d}W(s).$$

Since each bound motor takes steps of unit length at a Poisson rate, and since a motor binds at the current cargo location, it follows that

(46) 
$$|\mu(t)| \le P(t) + X_{\sup}(t), \text{ where } X_{\sup}(t) := \sup_{t' \in [0,t]} |X(t')|,$$

and  $\{P(s)\}_{s\geq 0}$  is a Poisson process with rate  $\mu := M \max_{m \in \{1,\dots,M\}} \lambda_{\text{step}}(m)$ . Thus,

(47) 
$$|X(t)| \le \sigma \left| \int_0^t e^{-\int_s^t \theta(s') \, \mathrm{d}s'} \, \mathrm{d}W(s) \right| + P(t) + \int_0^t X_{\sup}(s)\theta(s)e^{-\int_s^t \theta(s') \, \mathrm{d}s'} \, \mathrm{d}s$$

where we have used the fact that

(48) 
$$\int_{0}^{t_{0}} f(s)\theta(s)e^{-\int_{s}^{t_{0}}\theta(s')\,\mathrm{d}s'}\,\mathrm{d}s \le \int_{0}^{t} f(s)\theta(s)e^{-\int_{s}^{t}\theta(s')\,\mathrm{d}s'}\,\mathrm{d}s \le f(t), \quad 0 \le t_{0} \le t,$$

if f is any nonnegative and nondecreasing function.

Now, a straightforward calculation using integration by parts shows that

(49) 
$$\sup_{t' \in [0,t]} \left| \int_0^{t'} e^{-\int_s^{t'} \theta(s') \, \mathrm{d}s'} \, \mathrm{d}W(s) \right| \le 2 \sup_{s \in [0,t]} |W(s)|.$$

Therefore, combining (47), (48), and (49) yields

(50) 
$$X_{\sup}(t) \le \alpha(t) + \int_0^t X_{\sup}(s)\theta(s)e^{-\int_s^t \theta(s')\,\mathrm{d}s'}\,\mathrm{d}s,$$

where  $\alpha(t) := 2\sigma \sup_{s \in [0,t]} |W(s)| + P(t)$ . Multiplying (50) by  $e^{\int_0^t \theta(s') \, \mathrm{d}s'}$ , applying Gronwall's inequality, and then dividing by  $e^{\int_0^t \theta(s') \, \mathrm{d}s'}$  yields

$$X_{\sup}(t) \le \alpha(t) + \int_0^t \alpha(s)\theta(s) \,\mathrm{d}s$$

Since  $\alpha(t)$  is nondecreasing and  $\theta(t) \leq \Theta := \varepsilon^{-1}M$ , we obtain  $X_{\sup}(t) \leq \alpha(t)(1 + \Theta t)$ . Therefore, we have the following almost sure inequality for  $\zeta := (1 + \Theta T_{\text{on}}^k)^{-1}$ :

$$\mathbb{P}(Y_{\mathrm{on}}^k \ge C | T_{\mathrm{on}}^k) \le \mathbb{P}\left(P(T_{\mathrm{on}}^k) \ge C\zeta/2 \big| T_{\mathrm{on}}^k\right) + \mathbb{P}\left(\sup_{t \in [0, T_{\mathrm{on}}^k]} |W(t)| \ge C\zeta/(4\sigma) \big| T_{\mathrm{on}}^k\right).$$

Copyright © by SIAM. Unauthorized reproduction of this article is prohibited.

We thus need to control the distribution of  $T_{\text{on}}^k$ . Now, the Markov chain m(t) in (6) is a finite state space birth-death process, and thus there exists (see [7]) a unique quasi-stationary distribution  $\nu \in \mathbb{R}^M$ , which is a probability measure on  $\{1, \ldots, M\}$  so that if  $\mathbb{P}(m(0) = m) = \nu_m$  for  $m \in \{1, \ldots, M\}$ , then

$$\mathbb{P}(m(t) = m | m(s) \neq 0 \text{ for all } s \in [0, t]) = \nu_m, \quad m \in \{1, \dots, M\}.$$

Furthermore, it is known that the first passage time of m(t) to state 0 is exponentially distributed with some rate  $\lambda > 0$  if  $\mathbb{P}(m(0) = m) = \nu_m$  for  $m \in \{1, \ldots, M\}$  [37]. Now,  $T_{\text{on}}$  is the first passage time of m(t) to state m = 0 starting from state m = 1, which must be less than the first passage time to state m = 0 starting from any other state. Therefore, if S is the first passage time to state m = 0 starting from this quasi-stationary distribution, then

$$\mathbb{P}(T_{\text{on}} > T) \le \mathbb{P}(S > T) = 1 - e^{-\lambda T}, \quad T > 0.$$

Thus, since both terms in the upper bound in (51) are increasing functions of the realization  $T_{\rm on}^k > 0$ , the tower property yields for  $\zeta_S := (1 + \Theta S)^{-1}$ ,

(52)  

$$\mathbb{P}(Y_{\text{on}}^{k} \ge C\zeta) = \mathbb{E}\left[\mathbb{P}(Y_{\text{on}}^{k} \ge C\zeta | T_{\text{on}}^{k})\right]$$

$$\leq \mathbb{E}\left[\mathbb{P}(P(S) \ge C\zeta_{S}/2|S)\right] + \mathbb{E}\left[\mathbb{P}\left(\sup_{t \in [0,S]} |W(t)| \ge C\zeta_{S}/(4\sigma)|S\right)\right]$$

Next, if P is Poisson distributed with mean  $\mu_0$ , then Corollary 6 from [43] yields

$$\mathbb{P}(P \ge C_0) \le e^{C_0 - \mu_0} \left(\frac{\mu_0}{C_0}\right)^{C_0} \quad \text{if } C_0 \ge \mu_0$$

Noting that  $\frac{C}{2} < \mu S(1 + \Theta S)$  if and only if  $S > \Sigma := \frac{\sqrt{2C\Theta/\mu + 1} - 1}{2\Theta}$ , we obtain

(53) 
$$\mathbb{P}(P(S) \ge C\zeta_S/2|S) \le 1_{S>\Sigma} + e^{\frac{C}{2(1+\Theta S)} - \mu S} \left(\frac{2\mu S(1+\Theta S)}{C}\right)^{\frac{C}{2(1+\Theta S)}} 1_{S \le \Sigma}.$$

Since  $S \sim \text{Exponential}(\lambda)$ , taking the expectation of (53) gives

$$E[\mathbb{P}(P(S) \ge C\zeta_S/2|S)] \le e^{-\lambda\Sigma} + \int_0^{\Sigma} \lambda e^{\frac{C}{2(1+\Theta_S)} - (\mu+\lambda)s} \left(\frac{2\mu s(1+\Theta_S)}{C}\right)^{\frac{C}{2(1+\Theta_S)}} \,\mathrm{d}s.$$

A quick calculation shows that if  $C \ge (\Theta \varepsilon_0)^{-2}$  for  $\varepsilon_0 := (8e\mu\Theta)^{-1/2}$ , then

$$\lambda e^{\frac{C}{2(1+\Theta s)} - (\mu+\lambda)s} \left(\frac{2\mu s(1+\Theta s)}{C}\right)^{\frac{C}{2(1+\Theta s)}} \leq \begin{cases} \lambda 2^{-\kappa_0\sqrt{C}} & \text{if } s \in [0, \varepsilon_0\sqrt{C}], \\ \lambda e^{-\lambda\varepsilon_0\sqrt{C}} & \text{if } s \in [\varepsilon_0\sqrt{C}, \Sigma] \end{cases}$$

for  $\kappa_0 := (4\Theta \varepsilon_0)^{-1}$ . Hence, if  $C \ge (\Theta \varepsilon_0)^{-2}$ , then

(55) 
$$E[\mathbb{P}(P(S) \ge C\zeta_S/2|S)] \le \lambda e^{-\lambda\Sigma} + \Sigma\lambda \left(2^{-\kappa_0\sqrt{C}} + e^{-\lambda\varepsilon_0\sqrt{C}}\right).$$

Moving to the second term in (52), we have that (41) yields

Downloaded 09/24/18 to 216.165.95.182. Redistribution subject to SIAM license or copyright; see http://www.siam.org/journals/ojsa.php

NONPROCESSIVE MOTOR TRANSPORT

$$\mathbb{E}\left[\mathbb{P}\left(\sup_{t\in[0,S]}|W(t)|\geq C\zeta_S/(4\sigma)|S\right)\right]\leq \int_0^\infty 2\lambda e^{-\lambda s}\exp\left\{-\frac{\kappa C^2}{s(1+\Theta s)^2}\right\}\,\mathrm{d}s$$

where  $\kappa = (32\sigma^2)^{-1}$ . It is straightforward to check that

$$2\lambda e^{-\lambda s} \exp\left\{-\frac{\kappa C^2}{s(1+\Theta s)^2}\right\} \le \begin{cases} 2\lambda \exp\left\{-\frac{\kappa \sqrt{C}}{\Theta^2 + 2\Theta C^{-1/2} + C^{-1}}\right\} & \text{if } s \in [0, \sqrt{C}],\\ 2\lambda e^{-\lambda s} & \text{if } s \in [\sqrt{C}, \infty). \end{cases}$$

Therefore, for sufficiently large C, we have that

(56) 
$$\mathbb{E}\left[\mathbb{P}(\sup_{t\in[0,S]}|W(t)|\geq C\zeta_S/(4\sigma)|S)\right]\leq 2\lambda\sqrt{C}e^{-(2\kappa/\theta^2)\sqrt{C}}+2e^{-\lambda\sqrt{C}}.$$

Combining (52), (55), and (56) completes the proof.

Proof of Lemma 4. Since  $Y_k \leq Y_{off}^k + Y_{on}^k$  for  $k \geq 1$ , we have that

(57) 
$$\sum_{k=1}^{\infty} \mathbb{P}(Y_k > \sqrt{k}) \le \sum_{k=1}^{\infty} \mathbb{P}(Y_{\text{off}}^k > \frac{1}{2}\sqrt{k}) + \sum_{k=1}^{\infty} \mathbb{P}(Y_{\text{on}}^k > \frac{1}{2}\sqrt{k}).$$

Therefore, the upper bounds established in Lemmas 2 and 3 and the integral test show that (57) converges. Applying the Borel–Cantelli lemma (see Theorem 2.3.1 in [10]) completes the proof.

Proof of Lemma 5. Since  $Y_k \ge 0$  almost surely, we have that  $\mathbb{E}[Y_k] = \int_0^\infty \mathbb{P}(Y_k > C) \, \mathrm{d}C$ . Using the bounds in Lemmas 2 and 3 as in the proof of Lemma 4 shows that this integral is finite.

#### Appendix B. Proof of Proposition 7.

*Proof.* Fix a realization **J**. Let  $K \ge 0$  denote the almost surely finite number of jump times of **J** before time T, where t is said to be a jump time if  $\mathbf{J}(t+) \neq \mathbf{J}(t-)$ . Denote these K jump times by  $0 < \tau_1 < \cdots < \tau_K < T$  and let  $\tau_0 = 0$  and  $\tau_{K+1} = T$ .

For ease of notation, define the sequences

$$x_k := x(\tau_k), \quad \overline{x}_k := \overline{x}(\tau_k), \quad z_k^i := z_i(\tau_k), \quad \overline{z}_k^i := \overline{z}_i(\tau_k), \quad m_k := m(\tau_k)$$

for  $k \in \{0, 1, ..., K\}$ . Further, define the time between jumps,  $s_k := \tau_k - \tau_{k-1}$ , for  $k \in \{1, ..., K\}$ . It follows immediately from Proposition 6 that

(58) 
$$x_{k+1} = x_k e^{-s_{k+1}/\varepsilon} + \mu_{k+1} (1 - e^{-s_{k+1}/\varepsilon}), \quad k \in \{0, 1, \dots, K\},$$

where for  $k \in \{0, 1, \dots, K+1\}$  we define

(59) 
$$\mu_{k+1} := \begin{cases} \frac{1}{m_k} \sum_{i \in I(\tau_k)} z_k^i & \text{if } m_k > 0, \\ x_k & \text{if } m_k = 0. \end{cases}$$

Furthermore, it follows from the definition of  $\overline{x}(t)$  that for  $k \in \{0, 1, \dots, K\}$ ,

(60) 
$$\overline{x}_{k+1} := \begin{cases} \frac{1}{m_k} \sum_{i \in I(\tau_k)} \overline{z}_k^i & \text{if } m_k > 0, \\ \overline{x}_k & \text{if } m_k = 0. \end{cases}$$

Now, since motors take steps of size one, it follows that if  $k \in \{0, \ldots, K\}$  and  $i \in \{1, \ldots, M\}$ , then  $0 \le z_k^i \le K + 1$  and  $0 \le x_k \le K + 1$ . Hence, if  $k \in \{0, \ldots, K\}$ , then (59) implies

Copyright © by SIAM. Unauthorized reproduction of this article is prohibited.

2529

C. E. MILES, S. D. LAWLEY, AND J. P. KEENER

(61) 
$$|x_k - \mu_{k+1}| < K + 1.$$

Next, we claim that if  $k \in \{0, \ldots, K\}$  and

(62) 
$$\max_{j \in \{0,...,k\}} \left\{ |x_j - \overline{x}_j|, \max_{i \in \{1,...,M\}} |z_j^i - \overline{z}_j^i| \right\} < \eta,$$

then

2530

(63) 
$$\max\left\{|x_{k+1} - \overline{x}_{k+1}|, \max_{i \in \{1, \dots, M\}} |z_{k+1}^i - \overline{z}_{k+1}^i|\right\} < (K+1)e^{-s_{k+1}/\varepsilon} + \eta.$$

To see this, we use (58) and (61) to obtain

$$|x_{k+1} - \overline{x}_{k+1}| = |x_k e^{-s_{k+1}/\varepsilon} + \mu_{k+1} (1 - e^{-s_{k+1}/\varepsilon}) - \overline{x}_{k+1}|$$
  
$$\leq (K+1) e^{-s_{k+1}/\varepsilon} + |\mu_{k+1} - \overline{x}_{k+1}|.$$

Using (59) and (60), we have that

$$|\mu_{k+1} - \overline{x}_{k+1}| \le \begin{cases} \frac{1}{m_k} \sum_{i \in I(\tau_k)} |z_k^i - \overline{z}_k^i| & \text{if } m_k > 0, \\ |x_k - \overline{x}_k| & \text{if } m_k = 0. \end{cases}$$

Furthermore, it follows from (7) and (27) that

$$|z_{k+1}^i - \overline{z}_{k+1}^i| \le \max_{j \in \{0, \dots, K+1\}} |x_j - \overline{x}_j|, \quad i \in \{1, \dots, M\}.$$

Hence, the claim (63) is verified.

Define the largest time between jumps,  $s := \max_{k \in \{1,...,K\}} s_k$ . Since  $x_0 = \overline{x}_0 = z_0^i = \overline{z}_0^i$  for  $i \in \{1,...,M\}$ , we apply (62) and (63) iteratively to obtain

$$|x_{K+1} - \overline{x}_{K+1}| \le (K+1)^2 e^{-s/\varepsilon}.$$

Taking  $\varepsilon \to 0$  completes the proof.

REFERENCES

- G. ARPAG, S. SHASTRY, W. O. HANCOCK, AND E. TÜZEL, Transport by populations of fast and slow kinesins uncovers novel family-dependent motor characteristics important for in vivo function, Biophys. J., 107 (2014), pp. 1896–1904.
- [2] R. D. ASTUMIAN AND I. DERÉNYI, A chemically reversible Brownian motor: Application to kinesin and Ncd, Biophys. J., 77 (1999), pp. 993–1002.
- [3] M. BRAUN, Z. LANSKY, A. SZUBA, F. W. SCHWARZ, A. MITRA, M. GAO, A. LÜDECKE, P. R. TEN WOLDE, AND S. DIEZ, Changes in microtubule overlap length regulate kinesin-14driven microtubule sliding, Nat. Chem. Biol., 13 (2017), pp. 1245–1252.
- [4] P. C. BRESSLOFF, Stochastic Processes in Cell Biology, Interdiscip. Appl. Math. 41, Springer International Publishing, Cham, Switzerland, 2014.
- [5] P. C. BRESSLOFF AND J. M. NEWBY, Stochastic models of intracellular transport, Rev. Modern Phys., 85 (2013), pp. 135–196.
- [6] R. B. CASE, D. W. PIERCE, N. HOM-BOOHER, C. L. HART, AND R. D. VALE, The directional preference of kinesin motors is specified by an element outside of the motor catalytic domain, Cell, 90 (1997), pp. 959–966.
- [7] J. A. CAVENDER, Quasi-stationary distributions of birth-and-death processes, Adv. in Appl. Probab., 10 (1978), pp. 570–586.
- [8] M. H. A. DAVIS, Piecewise-deterministic Markov processes: A general class of non-diffusion stochastic models, J. R. Stat. Soc. Ser. B. Stat. Methodol., 46 (1984), pp. 353–388.

- T. DUKE, Cooperativity of myosin molecules through strain-dependent chemistry, Philos. Trans. Roy. Soc. London Ser. B. 5, 355 (2000), pp. 529–38.
- [10] R. DURRETT, Probability: Theory and Examples, Cambridge University Press, Cambridge, UK, 2010.
- [11] S. A. ENDOW AND H. HIGUCHI, A mutant of the motor protein kinesin that moves in both directions on microtubules, Nature, 406 (2000), pp. 913–916.
- [12] R. P. ERICKSON, Z. JIA, S. P. GROSS, AND C. C. YU, How molecular motors are arranged on a cargo is important for vesicular transport, PLoS Comput. Biol., 7 (2011), 002032.
- [13] Q. FENG, K. J. MICKOLAJCZYK, G.-Y. CHEN, AND W. O. HANCOCK, Motor reattachment kinetics play a dominant role in multimotor-driven cargo transport, Biophys. J., 114 (2017), pp. 1–12.
- [14] G. FINK, L. HAJDO, K. J. SKOWRONEK, C. REUTHER, A. A. KASPRZAK, AND S. DIEZ, The mitotic kinesin-14 Ncd drives directional microtubule-microtubule sliding, Nat. Cell Biol., 11 (2009), pp. 717–723.
- [15] K. A. FOSTER, A. T. MACKEY, AND S. P. GILBERT, A mechanistic model for Ncd directionality, J. Biol. Chem., 276 (2001), pp. 19259–19266.
- [16] K. FURUTA, A. FURUTA, Y. Y. TOYOSHIMA, M. AMINO, K. OIWA, AND H. KOJIMA, Measuring collective transport by defined numbers of processive and nonprocessive kinesin motors, Proc. Natl. Acad. Sci. USA, 110 (2013), pp. 501–506.
- [17] K. FURUTA AND Y. Y. TOYOSHIMA, Minus-end-directed motor Ncd exhibits processive movement that is enhanced by microtubule bundling in vitro, Current Biol., 18 (2008), pp. 152–157.
- [18] D. T. GILLESPIE, Exact numerical simulation of the Ornstein-Uhlenbeck process and its integral, Phys. Rev. E, 54 (1996), pp. 2084–2091.
- [19] I. V. GOPICH AND A. SZABO, Diffusion modifies the connectivity of kinetic schemes for multisite binding and catalysis, Proc. Natl. Acad. Sci. USA, 110 (2013), pp. 19784–19789.
- [20] D. A. GROTJAHN, S. CHOWDHURY, Y. XU, R. J. MCKENNEY, T. A. SCHROER, AND G. C. LANDER, Cryo-electron tomography reveals that dynactin recruits a team of dyneins for processive motility, Nat. Struct. Mol. Biol., 25 (2018), pp. 203–207.
- [21] M. A. HALLEN, Z.-Y. LIANG, AND S. A. ENDOW, Ncd motor binding and transport in the spindle, J. Cell Sci., 121 (2008), pp. 3834–3841.
- [22] A. R. HODGES, C. S. BOOKWALTER, E. B. KREMENTSOVA, AND K. M. TRYBUS, A nonprocessive class V myosin drives cargo processively when a kinesin-related protein is a passenger, Current Biol., 19 (2009), pp. 2121–2125.
- [23] J. HUGHES, W. O. HANCOCK, AND J. FRICKS, A matrix computational approach to kinesin neck linker extension, J. Theoret. Biol., 269 (2011), pp. 181–194.
- [24] J. HUGHES, W. O. HANCOCK, AND J. FRICKS, Kinesins with extended neck linkers: A chemomechanical model for variable-length stepping, Bull. Math. Biol., 74 (2012), pp. 1066–1097.
- [25] J. HUGHES, S. SHASTRY, W. O. HANCOCK, AND J. FRICKS, Estimating velocity for processive motor proteins with random detachment, J. Agric. Biol. Environ. Stat., 18 (2013), pp. 204–217.
- [26] E. JONSSON, M. YAMADA, R. D. VALE, AND G. GOSHIMA, Clustering of a kinesin-14 motor enables processive retrograde microtubule-based transport in plants, Nat. Plants, 1 (2015), 15087.
- [27] F. JÜLICHER AND J. PROST, Cooperative molecular motors, Phys. Rev. Lett., 75 (1995), pp. 2618–2621.
- [28] I. KARATZAS AND S. SHREVE, Brownian Motion and Stochastic Calculus, Springer Science & Business Media, New York, 2012.
- [29] S. KLUMPP AND R. LIPOWSKY, Cooperative cargo transport by several molecular motors, Proc. Natl. Acad. Sci. USA, 102 (2005), pp. 17284–17289.
- [30] C. B. KORN, S. KLUMPP, R. LIPOWSKY, AND U. S. SCHWARZ, Stochastic simulations of cargo transport by processive molecular motors, J. Chem. Phys., 131 (2009), 245107.
- [31] A. KRISHNAN AND B. I. EPUREANU, Renewal-reward process formulation of motor protein dynamics, Bull. Math. Biol., 73 (2011), pp. 2452–2482.
- [32] D. A. LAUFFENBURGER AND J. LINDERMAN, Receptors: Models for Binding, Trafficking, and Signaling, Oxford University Press, Oxford, UK, 1993.
- [33] S. D. LAWLEY AND J. P. KEENER, Including rebinding reactions in well-mixed models of distributive biochemical reactions, Biophys. J., 111 (2016), pp. 2317–2326.
- [34] S. D. LAWLEY AND J. P. KEENER, Rebinding in biochemical reactions on membranes, Phys. Biol., 14 (2017), 056002.

- [35] A. T. LOMBARDO, S. R. NELSON, M. Y. ALI, G. G. KENNEDY, K. M. TRYBUS, S. WALCOTT, AND D. M. WARSHAW, Myosin Va molecular motors manoeuvre liposome cargo through suspended actin filament intersections in vitro, Nat. Commun., 8 (2017), p. 15692.
- [36] S. A. MCKINLEY, A. ATHREYA, J. FRICKS, AND P. R. KRAMER, Asymptotic analysis of microtubule-based transport by multiple identical molecular motors, J. Theor. Biol., 305 (2012), pp. 54–69.
- [37] S. MÉLÉARD AND D. VILLEMONAIS, Quasi-stationary distributions and population processes, Probab. Surv., 9 (2012), pp. 340–410.
- [38] C. MIECK, M. I. MOLODTSOV, K. DRZEWICKA, B. VAN DER VAART, G. LITOS, G. SCHMAUSS, A. VAZIRI, AND S. WESTERMANN, Non-catalytic motor domains enable processive movement and functional diversification of the kinesin-14 Kar3, eLife, 4 (2015), pp. 1–23.
- [39] B. NITZSCHE, E. DUDEK, L. HAJDO, A. A. KASPRZAK, A. VILFAN, AND S. DIEZ, Working stroke of the kinesin-14, ncd, comprises two substeps of different direction, Proc. Natl. Acad. Sci. USA, 113 (2016), pp. E6582–E6589.
- [40] J. NORRIS, Markov Chains, Camb. Ser. Stat. Probab. Math. 2, Cambridge University Press, Cambridge, UK, 1998.
- [41] E. PECHATNIKOVA AND E. W. TAYLOR, Kinetics processivity and the direction of motion of Ncd., Biophys. J., 77 (1999), pp. 1003–1016.
- [42] P. M. SHAKLEE, T. IDEMA, G. KOSTER, C. STORM, T. SCHMIDT, AND M. DOGTEROM, Bidirectional membrane tube dynamics driven by nonprocessive motors, Proc. Natl. Acad. Sci. USA, 105 (2008), pp. 7993–7997.
- [43] M. SHORT, Improved inequalities for the poisson and binomial distribution and upper tail quantile functions, ISRN Probab. Stat., 2013 (2013), 412958.
- [44] B. SHTYLLA AND J. P. KEENER, Mathematical modeling of bacterial track-altering motors: Track cleaving through burnt-bridge ratchets, Phys. Rev. E, 91 (2015), 042711.
- [45] G. T. SHUBEITA, S. L. TRAN, J. XU, M. VERSHININ, S. CERMELLI, S. L. COTTON, M. A. WELTE, AND S. P. GROSS, Consequences of motor copy number on the intracellular transport of kinesin-1-driven lipid droplets, Cell, 135 (2008), pp. 1098–1107.
- [46] K. TAKAHASHI, S. TANASE-NICOLA, AND P. REIN TEN WOLDE, Spatio-temporal correlations can drastically change the response of a MAPK pathway, Proc. Natl. Acad. Sci. USA, 107 (2010), pp. 2473–2478.
- [47] M. VLASIOU, Regenerative processes, in Wiley Encyclopedia of Operations Research and Management Science, John Wiley and Sons, Hoboken, NJ, 2011.
- [48] M. YAMADA, Y. TANAKA-TAKIGUCHI, M. HAYASHI, M. NISHINA, AND G. GOSHIMA, Multiple kinesin-14 family members drive microtubule minus end-directed transport in plant cells, J. Cell Biol., 216 (2017), pp. 1705–1714.
- [49] Z.-H. YANG AND Y.-M. CHU, On approximating the modified Bessel function of the second kind, J. Inequal. Appl., 2017 (2017), p. 41.