Control of swarming of molecular robots

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Recently we demonstrated swarming of a self-propelled biomolecular motor system microtubule (MT)-kinesin where interactions among thousands of motile MTs were regulated in a highly programmable fashion by using DNA as a processor. However, precise control of this potential system is yet to be achieved to optimize the swarm behavior. In this work, we systematically controlled swarming of MTs on kinesin adhered surface by different physicochemical parameters of MT-kinesin and DNA. Tuning the length of DNA sequences swarming was precisely controlled with thermodynamic and kinetic feasibility. In addition, swarming was regulated using different concentration of DNA crosslinkers. Reversibility of swarming was further controlled by changing the concentration of strand displacement DNA signal allowing dissociation of swarm. The control over the swarm was accompanied by variable stiffness of MTs successfully, providing translational and circular motion. Moreover, the morphology of swarm was also found to be changed not only depending on the stiffness but also body length of MTs. Such detail study of precise control of swarming would provide new insights in developing a promising molecular swarm robotic system with desired functions.

Colonies of bacteria and ants\(^{1,3}\), school of fish\(^4\), or flocks of birds\(^{5,6}\) represent typical examples of swarming in nature\(^7,8\), which emerge through local interactions among the self-propelled living organisms rather than through control by a leader. Swarming offers parallelism, robustness, and flexibility to the organisms\(^10,12\), and inspired attempts to realize swarming in a synthetic environment using natural\(^11,14\), chemically powered\(^16-20\) or mechanical self-propelled objects\(^21-24\). Self-propelled biomolecular motor systems, such as MT-kinesin, actin-myosin have been particularly promising for realizing such artificial swarming because of their small size, efficiency, and scalability\(^13,14\). However, a lack of programmability of local interactions among the self-propelled objects prevents us from establishing precise control over the artificial swarming.

To overcome this drawback, we developed a methodology to program local interactions among a large number of MTs driven by biomolecular motor kinesins\(^25\). By introducing DNA based molecular recognition into the self-propelled system of kinesin driven MTs, we successfully controlled the swarming of thousands of MTs in a reversible fashion. Furthermore, the swarming was programmed using logic gate operations responding to input DNA signals. MT stiffness was shown to be an additional determinant of the swarming behavior. By integrating a photo-responsive compound, azobenzene, to the DNA portion, we have constructed a “molecular swarm robot”, which possesses all the three essential features comprising “robots”, that is the integration of sensors (azobenzene), information processors (DNA), and actuators (MT-kinesin)\(^26-29\). However, alongside with programming the swarming, understanding the effect of physicochemical parameters of the system is equally important to design complex behavior of molecular swarm robots. To utilize the system with controlled swarm behavior, optimization of the key parameters of the system is essential which has been unexplored yet.

Here, we have systematically explored the role of relevant physicochemical parameters in the reversible regulation of swarming of MTs. We demonstrate how the strength of the DNA interaction regulates the swarming of MTs by varying the concentration of receptor DNA (r-DNA), linker DNA (l-DNA), dissociation DNA (d-DNA). We furthermore investigated how the physical properties of MTs influence the swarming of MTs. A detailed understanding of the physicochemical parameters of different components of the swarming system will enable us to expand the controllability and range of swarming behaviors. The knowledge may advance the development of

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complex molecular devices utilizing swarming of biomolecular motor systems in a bottom-up manner giving rise to new emergent functions of molecular robots.

Results

Swarming of DNA conjugated MTs in a dynamic condition. Figure 1a shows a schematic illustration of the present molecular swarm system. The individual swarm units were prepared by conjugating MTs and DNA. MTs of lengths between 2 and 10 μm having the path persistence length, $L_p$, of ~580 μm were conjugated with two different DNA strands, $T_{16}$ and d(TTG)$_5$, termed as receptor DNA ($r$-DNAs), through a copper-free click reaction. Prior to the conjugation to MTs, $T_{16}$ and d(TTG)$_5$ were labeled with a fluorescent dye, TAMRA (red) and FAM (green), respectively in order to allow monitoring of the DNA conjugated MTs under a fluorescence microscope (Supplementary Table S1). The DNA conjugated MTs were propelled by surface-adhered kinesins using the chemical energy of adenosine triphosphate (ATP).

We first investigated how the binding affinity of DNA conjugated MTs promotes swarm formation. We monitored two single MTs conjugated with $r$-DNAs while gliding in the presence of $l$-DNA (d(CAA)$_{16}$A$_{16}$) (Fig. 1b). The $l$-DNA was designed such that it is partially complementary to the $r$-DNAs and may allow crosslinking of the MTs (Supplementary Table S1). Swarming was observed when the red and green MTs collided with each other similar to the snuggling phenomenon. On the other hand, when two MTs approached from totally opposite

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**Figure 1.** Schematic illustration of the system and interaction between red and green MTs in presence of $l$-DNA. (a) MTs are modified with two kinds of fluorescently labeled DNA ($r$-DNAs) and are propelled on a kinesin coated glass slide. Addition of complementary DNA ($l$-DNA) that bridges the two $r$-DNAs triggers swarming of the MTs. (b) Upper part: time lapse images of interaction between red and green MTs modified with $T_{16}$ and d(TTG)$_5$, respectively in presence of $l$-DNA (d(CAA)$_{16}$A$_{16}$) and move together after swarm formation. Association of red and green MTs exhibits yellow color indicating swarming. Scale bar: 5 μm. Lower part: time lapse images of interaction between red and green MTs in presence of $l$-DNA moving in the opposite direction. Swarming was not observed in this case. Scale bar: 10 μm. Images were captured just after the ATP addition. The concentration of kinesin was 0.3 μM.
direction, they passed each other without exhibiting swarming (Fig. 1b, lower part). These results suggest that swarming of MTs depends on the collision angle of the MTs. The swarm of MTs exhibited translational motion with a velocity $(0.52 \pm 0.03 \mu m s^{-1})$ close to that of single MTs $(0.54 \pm 0.05 \mu m s^{-1})$ which coincides well with our previous work.

Along with the binding interaction of DNA, the driving force of the surface adhered kinesins is another factor that should be considered in the swarming of MTs. The motility of the r-DNA conjugated MTs was governed by the kinesins which exert force to propel and assemble the MTs into swarms (Supplementary Fig. S1). By varying the concentration of kinesin in the feed $(0.001–1 \mu M)$, the surface density of kinesin was tuned, which affected the binding of MTs. We have estimated the density of kinesins on substrate for each of the in feed kinesin concentrations as described in the methods section. For the 0.001, 0.01, 0.1, 0.3, 0.5, 0.8, 1 \mu M in feed concentrations the corresponding kinesin density on the substrate were $\sim 14, 140, 1,400, 4,200, 7,000, 11,200, 14,000 \mu m^{-2}$ respectively. At lower concentrations of kinesin $(0.001–0.01 \mu M)$ at a density $14–140 \mu m^{-2}$, the number of MTs bound to the surface was relatively low and motility was rarely observed (Supplementary Fig. S1a). At intermediate concentrations of kinesin (at 0.1–0.3 \mu M) at a density 1,400–4,200 \mu m^{-2}, the motility and interaction between MTs are facilitated. At higher concentration of kinesin $(0.5–1 \mu M)$, the binding of MTs to the surface was found to be further increased but the swarming was decreased, which is evident from the decrease in the association ratio (Supplementary Fig. S1b). Therefore, the swarming of MTs is optimally facilitated at a moderate kinesin concentration of 0.3 \mu M corresponding to the density of $4,000 \mu m^{-2}$ (Supplementary Fig. S1b). We counted the number of individual MTs at initial time and after 30 min of adding ATP buffer and characterized the swarming of MTs by calculating the association ratio (see Methods) at different concentration of kinesin. At $4,000 \mu m^{-2}$ density, the swarming was found to increase at a maximum with association ratio of $\sim 70\%$. Here we estimated the number of kinesins interacting with per unit length of MTs. As already discussed, for the 0.3 \mu M in feed concentration of kinesin the density was estimated to be $\sim 4,000 \mu m^{-2}$, which implies that spacing between two kinesins is $\sim 16 \mu m$. It was reported that kinesin follows a single protofilament of tubulins during its movement along MTs. Considering the width of tubulin as 4.5 nm, we may assume that during motility of MTs on kinesins only one protofilament of MTs is able to interact with kinesins. Therefore, the maximum number of kinesins per \mu m length of a protofilament was estimated to be $\sim 63$. This result indicates that for the 0.3 \mu M in feed concentration of kinesin, there is 1 kinesin for $\sim 2$ tubulin dimers along a protofilament of MTs. In this context, our result suggests that the above ratio of kinesin to tubulin dimer is effective to exert optimum driving forces for maximum swarm formation.

**Control of swarming of MTs by designing the length of the r-DNA sequence.** In order to further confirm that the swarming of the T$_{16}$ and d(TTG)$_{5}$ conjugated MTs is indeed mediated by r-DNAs, we have designed r-DNAs with different length $(d(CAA)_{n}A_{16}; n = 1–5)$ (Supplementary Table S2). Simulations predict how the melting temperature $(T_{m})$ increases with increasing length of r-DNA. Depending on the feasibility of hybridization of complementary DNA strands favored by their $T_{m}$, the length of r-DNA can influence the DNA based self-assembly as found in the previous works. In our system, T$_{16}$ and d(TTG)$_{5}$ conjugated MTs were allowed to move on a kinesin-coated surface at a combined density of $\sim 50,000 \mu m^{-2}$. Upon addition of r-DNA with CAA repeats shorter than 3, no swarming of MTs was observed $(d(CCA)_{16}, d(CAA)_{16})$ (Fig. 2a). In the case of r-DNA where the number of CAA repeats was 3, the r-DNA modified MTs started swarming by interacting with the r-DNA $(d(CAA)_{16})$. While gliding, the red and green MTs came close to each other, associated into clusters and continued moving. For $(d(CAA)_{16}, (28$ base sequences) and $(d(CAA)_{31}, (31$ base sequences), on the other hand, the preferential swarming of the MTs was observed comprising large number of them (Fig. 2a). This may represent that increasing complementary r-DNA interaction provided sufficient interaction energy which resulted in the increasing yellow colored swarms of MTs (Fig. 2a). The MTs swarms in dynamic condition are distinguishable from the unstructured aggregates in static condition formed from freely diffusing MTs in absence of kinesins. Along with the thermodynamic feasibility, the swarm formation rate was also influenced by the length of r-DNA. As the length of r-DNA, i.e. the length of complementary base sequences with r-DNA was increased, the swarm formation rate accordingly increased, which can be easily understood from the change in their association ratio with time (Fig. 2b). For $(d(CAA)_{16}, d(CAA)_{16})$, the swarm formation rate was found maximum up to $\sim 95–98\%$ after reaching a plateau with time.

**Effect of the concentration of r-DNAs and l-DNA on swarming of MTs.** As demonstrated above, the swarming of MTs relies on the interaction between r-DNAs and l-DNA. So, a systematic investigation was performed to determine how the concentration of the r-DNAs and l-DNA affects the swarming. In the investigation, an almost equal density of red and green MTs (1:1 density ratio) was used in the experiments at which swarms with highest association ratio and largest size were formed (Supplementary Fig. S2).

We first investigated the effect of the concentration of r-DNAs on swarming of MTs. Fixing the concentration of l-DNA at 0.6 \mu M, we varied the concentration of r-DNAs (T$_{16}$ and d(TTG)$_{5}$) used for MT modification from 50–1,000 \mu M. The labeling ratio of r-DNA to MTs i.e., tubulin dimers was found to increase as the concentration of r-DNAs increased, and reached $\sim 100\%$ when the concentration of r-DNA was 500 \mu M. Although MTs exhibited swarming with translational motion when the concentration of r-DNA was 200 \mu M, more swarming was observed at 500 \mu M of r-DNA, which is evident from the corresponding association ratio (Fig. 3a, rigid MTs) approximately $\sim 87\%$ within 60 min after the addition of l-DNA. Further increase in r-DNAs concentration $(800–1,000 \mu M)$ brought no considerable change in the association ratio (Fig. 3a). By changing polymerization condition (see experimental section), we changed the rigidity of MTs and found that more MTs exhibited swarming with circular motion upon decreasing their rigidity with $L_{c}$ of $\sim 245 \mu m$ (Fig. 3a, flexible MTs). Even though the rigidity of MTs induced such a change in swarming mode, no difference in the effect of concentration of r-DNA was observed for the two systems. At 500 \mu M of r-DNA, increase in circular swarming was observed which reached a
plateau at 800 µM–1,000 µM r-DNA concentration with association ratio of ~92–95%. Similar trend in association ratio change was obtained as in the case of rigid MTs. Based on these outcomes, a phase diagram was constructed correlating the concentration of r-DNAs with the swarming of MTs (Supplementary Fig. S3a).
We then investigated the effect of the concentration of l-DNA on swarming of MTs. While the concentration of r-DNA1 and r-DNA2 was fixed at 500 µM, the concentration of l-DNA was varied from 0.001 to 1 µM (Fig. 3b). At low concentration of l-DNA (0.001 µM), swarming was observed for neither the rigid nor flexible MTs. Swarming was only observed when the concentration of l-DNA was above 0.05 µM. As the concentration of l-DNA was further increased (0.2–1 µM), the association ratio initially increased and then reached a plateau for both, rigid and flexible MTs with approximate 88% association ratio. The effect of l-DNA concentration on the swarming of MTs is again summarized in a phase diagram (Supplementary Fig. S3b).

**Design of d-DNA and effect of its concentration on the reversible swarming.** DNA strand displacement reactions, in which a DNA strand is displaced by another one, have been proved useful in controlling self-assembly based on DNA interactions. Therefore, we designed a DNA sequence (d-DNA) such that it facilitates dissociation of the swarms into isolated MTs. Strand displacement requires that the binding energy of d-DNA with l-DNA is higher than that between l-DNA and r-DNAs (Supplementary Table S1) which is easily predictable from their Tm. A DNA sequence (d-DNA), which was fully complementary to the l-DNA sequence was chosen to withdraw the l-DNA from MT swarms by the DNA strand displacement reaction. Upon addition of d-DNA (0.05 µM) swarms of MTs with translational and circular motion began to dissociate into single MTs, which was quantified by a significant decrease in the association ratio (Fig. 4). A further increase in the concentration of d-DNA decreased the association ratio further. At 1 µM d-DNA, the association ratio was close to zero and the MT swarms were almost completely dissociated (Fig. 4).

**Effect of the length of MTs on swarming mode.** We previously demonstrated that the rigidity of MTs can be utilized to control the swarming mode of MTs. An open question is if the length of MTs has any influence
on their swarming mode. Flexible MTs with different lengths were prepared by applying shear stress using a micro syringe as reported previously\(^44,45\). The length of MTs was tuned by varying the number of shear treatments (up to ten times). The average lengths of flexible red and green MTs were 22.2 \(\pm\) 12.5 \(\mu\)m (average \(\pm\) standard deviation (s.d.)) and 16.2 \(\pm\) 9.0 \(\mu\)m respectively just after preparation (Fig. 5a). After ten shear treatments the length of red and green MTs were reduced to 4.7 \(\pm\) 2.5 \(\mu\)m and 4.7 \(\pm\) 2.4 \(\mu\)m respectively (Fig. 5b). The changes in the average length of the flexible MTs due to the shear treatment match the results reported in previous studies\(^44,45\).

When the shortened MTs were employed in swarming, the swarming mode was found to change from circular to translational one (Fig. 5a, b). This result indicates that the MT length plays an important role in determining the morphology of MT swarms.

Discussion

In order to develop a robustly controlled molecular swarm system, we systematically investigated the effect of related physicochemical parameters on the DNA assisted swarming of a self-propelled biomolecular motor system, MT-kinesin. We found that the effects are comparable to those discovered in previous studies where self-assembly of MTs was regulated using streptavidin (St)-biotin (Bt) crosslinkers system\(^32–34\). Unlike the self-assembly of MTs using the strong St-Bt interaction, where MTs assembled even when they moved in opposite direction\(^32,33\), almost no sticking of MT swarms on the kinesin-coated surface was noticed in the present system. This observation suggests that swarming of MTs was favored by moderate interaction between \(l\)-DNA and \(r\)-DNAs attached to MTs. The \(r\)-DNAs and \(l\)-DNA are designed such that they can bind in an unzipping manner, a feasible model for DNA interaction\(^46\). DNA hybridization through the unzipping mode was further facilitated when the MTs collided while they were moving in the same direction just before the collision. Consequently, MTs started to form swarm due to DNA hybridization through the unzipping mode. However, for the MTs that collided while moving in the opposite direction, no successful swarm formation was observed. A similar phenomenon was previously reported in which MTs moving in the opposite direction in a crowded condition also failed to form stable bundles\(^30\). The failure in swarm formation by the MTs could be ascribed to the direction of force exerted by the kinesins to the MTs moving in the opposite direction, and energetics of the associated DNA hybridization, confirmation of which requires further investigation.

In addition to that, our study is helpful in controlling the swarm system more efficiently by tuning the interaction not only depending on the concentration of the components but also by designing properties of the crosslinkers such as the length of \(l\)-DNA sequence. This was previously done to tune the self-assembly of particle systems in static conditions\(^39\). Here, we tuned the binding interactions of MTs under dynamic conditions by optimizing the sequence of \(l\)-DNA (Fig. 2). The contributions of the concentrations of \(r\)-DNA, \(l\)-DNA, \(d\)-DNA, and kinesin, and the physical properties of MTs were also found to have a profound influence on the swarming of MTs. Nearly complete association was obtained with DNA constituents above certain concentrations (Fig. 3). The phase diagrams...
predicted from the results indicate similar effects to those found in previous work \cite{34} (Supplementary Fig. S3). The reverse control responding to the \textit{d}-DNA is favored by their elevated \( T_m \) as predicted in the equilibrium condition \cite{25}, which was also successfully tuned by varying the concentration of \textit{d}-DNA. By selective parameter combinations, we are able to regulate not only the thermodynamics and kinetics of MT swarming (Fig. 2) but also the morphology of MT swarms between translational and circular modes corresponding to their stiffness and body length (Fig. 5).

We expect that the present study will enable the precise designing of DNA assisted swarming of self-propelled biomolecular motor systems \cite{47}. Moreover, our observations point towards a way of greatly improving the selectivity of DNA detection systems based upon biomolecular motor systems and also various applications such as active DNA sensors, adaptive actuators or analyte concentrators by optimization of these systems. However, apart from the chemical signal of DNA, physical signal by using light sensor has further advancements such as non-invasiveness, spatiotemporal regularity which are lacking in the present work. Insertion of a light sensor molecule in the system \cite{25}, could provide spatiotemporal control over swarming of MTs in a non-invasive way. Varying different parameters of light sensors such as number of photoresponsive azobenzene molecules in the DNA sequences, intensity of light, regulating localized association and dissociation of MTs swarms responding to the light signal; spatiotemporal control over molecular swarm robots could be further achieved which is expected to present as a future prospect. The knowledge from present and future work would expand potential applications of biomolecular motors with precisely controlled computation and may ultimately benefit molecular robotics \cite{26,27,48}.

**Methods**

**Purification of tubulin and kinesin.** Tubulin was purified from porcine brain using a high-concentration PIPES buffer (1 M PIPES, 20 mM EGTA, and 10 mM MgCl\(_2\)) and stored in BRB80 buffer (80 mM PIPES, 1 mM EGTA, 2 mM MgCl\(_2\), pH adjusted to 6.8 using KOH) \cite{49}. Recombinant kinesin-1 consisting of the first 573 amino acid residues of human kinesin-1 was prepared as described in the literature \cite{50}. Azide labeled tubulin was prepared using N3-PEG4-NHS following the established protocol of labeling tubulin with fluorescent dye \cite{51}. The tubulin concentration was determined by measuring the absorbance at 280 nm using a UV spectrophotometer (Nanodrop 2000c).
Design and preparation of DNA sequences.  r-DNA and l-DNA strands were designed by calculating the melting temperature of different sequences using 'OligoAnalyzer 3.1' software52, and selecting sequences with melting temperatures between 0°C and 50°C for experimental testing. Dibenzocyclooctyne (DBCO) and fluorescein dye labeled strands were chemically synthesized using appropriate CPG columns and a phosphoramidite monomer (Glen Research, VA) on an ABI 3900 automatic DNA synthesizer, purified by reverse phase HPLC and fully characterized by MALDI-TOF/MS (Bruker microflex LRF). The r-DNA was modified at the 3’ end with either 5(-)carboxytetramethylrhodamine (TAMRA) or 5-carboxyfluorescein (FAM) and at the 5’ end with DBCO. l-DNAs and d-DNA were purchased from Eurofins Genomics LLC.

Preparation of MTs. MTs were prepared by adding azide-tubulin to polymerization buffer (80 mM PIPES, 1 mM EGTA, 1 mM MgCl2, 1 mM polymerizing agent, pH~6.8) to a final concentration of 56 µM tubulin and incubating at 37°C for 30 min. The polymerizing agent for flexible MTs was guanosine triphosphate (GTP), and the polymerizing agent for rigid MTs was guanylyl-[α(3)-methylene] diphosphate (GMPCP), a slowly hydrolyzable analogue of GTP. Dimethyl sulfoxide (DMSO) was added to a final concentration of 5% for the polymerization of flexible MTs, but not for rigid MTs. A copper free click reaction was initiated by adding 3.5 µL DBCO conjugated r-DNAs (500 µM) to the 5 µl azide-MTs (56 µM), which allowed an azide-alkyne cycloaddition reaction, and incubated the solution at 37°C for 6hours39, 100 µl of cushion buffer (BRB80 buffer supplemented with 60% glycerol) was used to separate the MTs by centrifugation at 201,000 g (S55A2–0156 rotor, Hitachi) for 1 hour at 37°C. After removing the supernatant, the pellet of r-DNA conjugated MTs was washed once with 100 µl BRB80P (BRB80 supplemented with 1 mM taxol) and dissolved in 15 µl BRB80P.

Flow cells and motility assays for the demonstration of swarming of MTs. A flow cell with dimensions of 9 x 2.5 x 0.45 mm² (L x W x H) was assembled from two cover glasses (MATSUNAMI Inc.) using double-sided tape as spacer. The flow cell was filled with 5 µl casein buffer (BRB80 buffer supplemented with 0.5 mg mL⁻¹ casein). After incubating for 3 min, 0.3 µM kinesin solution was introduced into the flow cell and incubated for 5 min resulting in a kinesin density of 4,000 µm⁻² on the substrate40. After washing the flow cell with 5 µL of wash buffer (BRB80 buffer supplemented with 1 mM DTT, 10 µM taxol), 5 µl of red MTs (TAMRA labeled r-DNA1 MTs) solution were introduced and incubated for 2 min, followed by washing with 10 µL of wash buffer. Subsequently, 5 µL of green MTs solution were introduced and incubated for 2 min, followed by washing with 10 µL of wash buffer. The green MTs (FAM labeled r-DNA2 MTs) were incubated with l-DNA for 15 min at room temperature prior to addition into flow cell. The motility of red and green MTs was initiated by applying 5 µL of ATP buffer (wash buffer supplemented with 5 mM ATP, 4.5 mg mL⁻¹ D-glucose, 50 U mL⁻¹ glucose oxidase, 50 U mL⁻¹ catalase and 0.2% methylcellulose (w/v)). The time of ATP addition was set as 0 min. Soon after the addition of ATP buffer the flow cell was placed in an inert chamber system (ICS)34 and the MTs were monitored using an epi-fluorescence microscope at room temperature (25°C). The experiments were performed at least 3 times for each condition.

Tuning the length of MTs. MTs with different lengths were prepared by applying shear stress, as reported previously44,45, to DNA conjugated MTs using a micro syringe (Hamilton syringe with an inner diameter of 2.06 mm) and a PEEK tube (length: 50 mm, nominal inner diameter: 0.26 mm). 30 µL of MT solution (0.6 µM) were passed back-and-forth through the syringe-mounted PEEK tube by manual operation of the syringe. The length of flexible MTs (GTP-MTs) was tuned by varying the number of passages (up to ten times). The change in length of GTP-MTs before and after shear treatment was manually measured using the image analysis software ‘ImageJ’. The average lengths of red and green GTP-MTs were 22.2 ± 2.4 µm respectively just after preparation. After ten shear treatments, the length of red and green GTP-MTs was reduced to 4.7 ± 2.5 µm and 4.7 ± 2.4 µm respectively. The changes in the average length of the GTP-MTs due to the shear treatment coincide well with the results reported in previous studies44,45.

Estimation of kinesin density. Kinesin density was estimated following a previous report where density of kinesin on a substrate surface was estimated using quartz crystal microbalance (QCM)35. A standard curve was prepared using the previously reported data from which the following relationship between in feed concentration of kinesin and kinesin density on a substrate was obtained: y = 1.4 x 10⁸ x, where x and y are kinesin concentration in feed (µM) and kinesin density on the substrate (µm⁻²) respectively. Using this relationship, we then estimated the kinesin density for each of the in feed concentration of kinesin employed in this study.

Fluorescence microscopy. The samples were illuminated with a 100 W mercury lamp and visualized by an epifluorescence microscope (Eclipse Ti, Nikon) using an oil-coupled Plan Apo 60 × N.A.1.4 objective (Nikon). UV cut-off filter blocks (TRITC: EX 540/25, DM565, BA605/55; GFP: B: EX460-500, DM505, BA510-560; Nikon) were used in the optical path of the microscope. Images were captured using a cooled-CMOS camera (NEO sCMOS, Andor) connected to a PC. Two ND filters (ND4, 25% transmittance for TRITC and ND1, 100% transmittance for GFP-B) were inserted into the illumination light path of the fluorescence microscope to reduce photobleaching of the samples.

Image analysis and measurement of the association ratio of MTs. The length of MTs was measured from images captured by fluorescence microscopy using image analysis software (ImageJ). The association ratio at a given time t was determined by counting the number of single MTs manually and dividing the number at time t by the number present initially (t = 0). The time-dependent association ratio R(t) of red and green MTs was determined as follows
with

\[ R(t) = \frac{N(0) - N(t)}{N(0)} \]

(1)

\[ N(0) = \text{Initial number of single MTs}, \]

\[ N(t) = \text{Number of single MTs after time } t. \]

The mean association ratio was obtained from the average of four regions of interest (126.5 μm × 126.5 μm).

**Data availability.** The data that support this study are available from the corresponding author upon reasonable request.

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**Author Contributions**

A.M.R.K., D.I., K.S., A. Ku and A. Ka conceived and designed the experiments. J.J.K. performed the experiments, and analyzed the experimental results. J.J.K., A.M.R.K., H.H., A. Ku and A. Ka wrote the manuscript.

**Additional Information**

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