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Capturing directed molecular motion in the nuclear pore complex of live cells

Francesco Cardarelli, Luca Lanzano, and Enrico Gratton

Nuclear pore complexes (NPCs) are gateways for nucleocytoplasmic exchange. Intrinsically disordered nucleoporins (Nups) form a selective filter inside the NPC, taking a central role in the vital nucleo-cytoplasmic transport mechanism. How such intricate meshwork relates to function and gives rise to a transport mechanism is still unclear. Here we set out to tackle this issue in intact cells by an established combination of fluorescence correlation spectroscopy and real-time tracking of the center of mass of single NPCs. We find that directed Nup-mediated molecular motion may represent an intrinsic feature of the overall selective gating through intact NPCs.

The authors declare no conflict of interest.

Author contributions: F.C. and E.G. designed research; F.C. and L.L. performed research; F.C. and L.L. analyzed data; and F.C. and E.G. wrote the paper.

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Nup153 is anchored to the nuclear basket of the NPC through its N-terminal domain (18). At the same time it is hypothesized that the long, mobile FG-rich C-terminal domain of Nup153 is able to bind cargo located anywhere within its reach, even close to the cytoplasmic side of the pore (18). By rapidly orbiting around the center of mass of Nup153-GFP distribution, we detect a previously hidden dynamic behavior regulated so as to produce rapid, discrete exchange of the GFP tag between two separate positions within the NPC. Next, we show that this highly regulated vectorial exchange is shared by the Kap β receptor during transport. Finally, by cross-correlation of the two signals, we show that Nup153 activity is compatible with the functional nuclear import of a classical NLS-bearing cargo. Based on these evidences, we propose that Nup-mediated directed motion may contribute to the selective gating of molecules through intact NPCs. We do believe our results pave the way for future investigations of NPC function in intact cells.

**Results and Discussion**

**Analysis of Single Nup153-GFP Dynamics Within the NPC.** To investigate the barrier-like behavior of the FG domains during transport, we use a C-terminally GFP-labeled adduct of nucleoporin Nup153 transiently transfected into live CHO-K1 cells (Fig. 1A). It has been demonstrated by others that GFP-variants of Nup153 are correctly targeted and incorporated into NPCs (19–21). In addition, it was shown that Nup153-GFP dynamically interacts with the pore, with two characteristic residence times in the range of minutes (19, 20). This overall slow turnover defines the lifetime of Nup153 association with the NPC and is well separated from the characteristic timescale (millisecond range) of nucleocytoplasmic transport we want to address here. During transport, as schematically shown in Fig. 1B, the flexible FG-repeat carboxy-terminal domain (here tagged to GFP) extends towards the central channel where it is supposed to interact with soluble transport receptors [i.e., Kap β1, (14)] and promote their translocation to the nucleus (18). A schematic representation of

![Fig. 1](https://www.pnas.org/cgi/doi/10.1073/pnas.1200486109 Cardarelli et al.)

(A) Cell expressing Nup153-GFP (scale bar: 5 μm) imaged at different z positions. Nup153-GFP accumulation on single pores is better shown in the Bottom (scale bar: 1 μm). (B) Schematic representation of the accepted model of Nup153 activity: It binds to the Kap-cargo complex on the cytoplasmic face of the NPC (I), collapses towards the nucleus (II), and is finally released by RanGTP (III). (C) The PSF is scanned along a 64-points orbit (R = 120 nm) around the pore, as described in the text. (D) The ACF is displayed in a pseudo-color carpet in which the x coordinate corresponds to the points along the orbit and the y coordinate to the autocorrelation time (log-scale). (E) The average ACF of columns 8–24 quantitatively describes the hump of the cytoplasmic arc. (F) The pair correlation function at the distance of 32 pixels along the orbit [pCF(+32)] identifies two characteristic times of Nup153 molecular movement at the pore: a fast cytoplasm-to-nucleus collapse (peak position: 3.1 ms; FWHM: 1.8) and a slower nucleus-to-cytoplasm release (region highlighted by the red square in the pCF carpet and in C; correlation peak position: 5 ms; FWHM: 2.4). By contrast, almost no correlation is detected at the same distance but quite perpendicular to the channel (regions highlighted by the green and blue squares in the pCF carpet and in C). (G) The collapse/release components are here compared to the total ACF (dashed line).
the experiment is shown in Fig. 1C: A circular light envelope is formed around the pore by a scanning laser spot [point spread function (PSF)] around the pore in 0.5–1 ms, while the real-time tracking routine keeps the center of mass of the NPC always at the center of the orbit* (see Materials and Methods). The scan start point is known (Fig. S1) and positioned along the nuclear envelope plane (black dot in Fig. 1C), with the first half of the orbit scanning through the cytoplasmic face of the NPC. The orbit radius is usually set to approximately 120 nm, which we found to be optimal for the accuracy of tracking (see Materials and Methods). The intensity along the orbit is measured at 64 points in the NPC reference system and fluctuation analysis is performed to extract information on single Nup153-GFP molecules dynamics within the pore. The autocorrelation function (ACF) carpet unequivocally shows a spatiotemporal regulation of Nup153-GFP dynamics, which produces a peculiar “double-arc” shape of the correlation profile (Fig. 1D). As shown in the plot of Fig. 1E, the hump of the arc (e.g., columns 5–24) depicts a narrow distribution of correlation times in the ACF. Notably, these features are reminiscent of what we observed in the ACF analysis of Kap1 shuttling (16). Thus we are prompted to interpret the hump as a characteristic time of Nup153-GFP movement at the pore, with the same GFP being detected in one position along the orbit at time $t$ and again in the same location with a certain delay $\tau$ that determines the time position of the hump. Such a signature behavior is found in all the measured pores (additional examples in Fig. S2) with an average timing of 7.5 ± 2.2 ms ($N=25$ cells, Table 1). This recalls the molecular model of Nup153 activity (2), described as the result of the controlled collapse and release of its C-terminal FG-rich domain. However, the ACF cannot provide the directionality of the motion, but only the characteristic time of the overall collapse-release cycle. To separate the putative collapsing and releasing components of Nup153 movement we use the pair correlation function (pCF) analysis (22). In this analysis we perform a cross-correlation calculation of the time sequence at a pair of points along the orbit. At the distance of 32 pixels along the orbit, the pCF algorithm detects a sharp distribution of Nup153-GFP transit times in both the nucleus-to-cytoplasm and cytoplasm-to-nucleus directions (Fig. 1F). As a negative control, almost no correlation is detected at the same distance but perpendicular to the NPC channel (Fig. 1F and Fig. S3). The pCF analysis reveals that the directional movement of Nup153-GFP from cytoplasm to nucleus (3.0 ± 0.7 ms, $N=25$; Table 1) is slightly faster compared to the one from nucleus to cytoplasm (5.1 ± 0.9 ms; see also Fig. S4). These data are compatible with the idea that Nup153-GFP acts as a molecular spring alternating a fast collapse into compact molecular conformations and a slightly slower release into extended conformations. Remarkably, atomic force microscopy measurements showed similar properties of Nup153 in an in vitro reconstituted transport assay (14). In the same report, the authors show by immunogold electron microscopy how the FG-carboxy-termi-

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Table 1. Cumulative results from fluorescence correlation analysis along the orbit

<table>
<thead>
<tr>
<th></th>
<th>ACF</th>
<th>pCF(&gt; C)</th>
<th>pCF(&gt; N)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Peak position</td>
<td>FWHM</td>
<td>Peak position</td>
<td>FWHM</td>
</tr>
<tr>
<td>Nup153-GFP</td>
<td>7.5 ± 2.2</td>
<td>5.0 ± 1.8</td>
<td>3.0 ± 0.7</td>
</tr>
<tr>
<td>NLS-mCherry</td>
<td></td>
<td>3.5 ± 1.0</td>
<td>20.0 ± 0.7</td>
</tr>
<tr>
<td>Kap1-GFP*</td>
<td>9.6 ± 3.5</td>
<td>13 ± 6.5</td>
<td>3.3 ± 1.8</td>
</tr>
<tr>
<td>mCherry</td>
<td></td>
<td>35 ± 18</td>
<td>120 ± 54</td>
</tr>
</tbody>
</table>

*In the tracking experiment the PSF is scanned around the distribution of fluorescence of the analyzed protein. Depending on the protein this distribution may not coincide to the real center of mass of the entire NPC (to which we refer for simplicity).

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Kap1-GFP Directed Motion Through the NPC. If the Nup153-GFP dynamics highlighted above truly reflect functional pore activity, then Nup153 vectorial motion should correlate with the motion of the molecules transported by the same mechanism. We recently showed that, similarly to Nup153, the transport of the classical Kap1 receptor at the NPC is regulated so as to produce a peculiar hump in the ACF (16) (Table 1). Here we report results from a two-color data acquisition on cotransfected Kap1-GFP and mCherry (Fig. 2 A and B) in which we concomitantly measure active and passive fluxes at the NPC. The pCF analysis demonstrates that Kap1 movement through the NPC must be directed, as it is univocally described by a single time of transport from cytoplasm to nucleus (Fig. 2 C and D) and from nucleus to cytoplasm (Fig. 2 E and F). Untagged mCherry yields instead a broad distribution of transit times in both directions (Fig. 2 C–F) (i.e., passive diffusion is not able to order molecules in time and space). As a further control, the pair cross-correlation function (pCF) was calculated to look for any possible coupling between translocating Kap1 and mCherry molecules. As expected no (pair cross) correlation was detected in both the cytoplasm-to-nucleus (Fig. 2 C, Right carpet; Fig. 2 D, blue curve) and nucleus-to-cytoplasm direction (Fig. 2 E, Right carpet; Fig. 2 F, blue curve). The individual average transit times for active and passive transport (Table 1) are in agreement with reported values (23–25) and with the pCF-based estimate obtained by sampling over several microns across the nuclear envelope (26). It is worth noting that, contrary to Nup153, Kap1 shows a highly symmetric pCF profile in the two directions of transport (Table 1). In light of what was observed so far, this result suggests that Kap1 may be affected by two independent, although similar, mechanisms: one operating from cytoplasm to nucleus [possibly mediated by Nup153, (14)] and the other from nucleus to cytoplasm (yet to be identified).

Nup153-Mediated Cargo Transport Across the NPC. To probe the involvement of Nup153 in promoting molecular translocations from cytoplasm to nucleus, we performed an experiment in which Nup153-GFP is coexpressed with a mCherry-tagged nuclear localization sequence (NLS) (Fig. 3 A). NLS-mCherry is actively transported into the nucleus by a Kap1-driven process while it moves back to the cytoplasm by passive diffusion (27) (Fig. 3 B). Under these conditions Nup153-GFP behaves as expected in terms of ACF profile (Fig. 3 C), cytoplasm-to-nucleus “-collapse” (Fig. 3 D, Left carpet; Fig. 3 E, green curve), and nucleus-to-cytoplasm “release” (Fig. 3 F, Left carpet; Fig. 3 G, green curve). Concomitantly, NLS-mCherry yields two characteristic distributions of transit times (active + passive) from cytoplasm to nucleus (Fig. 3 D, Middle carpet; Fig. 3 E, red curve), but only one (passive) from nucleus to cytoplasm (Fig. 3 F, Middle carpet; Fig. 3 G, red curve). Remarkably, by pCF we show that NLS-mCherry motion is correlated in time and space with that of Nup153-GFP along the cytoplasm-to-nucleus direction of the NPC (Fig. 3 D, Right...
carpet; Fig. 3E, blue curve). The positive pair-cross-correlation signal suggests that at least a subpopulation of NLS-mCherry molecules may be effectively exploiting Nup153-GFP to move from cytoplasm to nucleus across the NPC. A relevant control for this conclusion is that no pcCF signal can be observed in the opposite direction (Fig. 3F, Right carpet; Fig. 3G, blue curve), where NLS-mCherry passive diffusion is not coupled to Nup153-GFP dynamics.

**Conclusions**

An accurate picture of how selective gating is achieved by the FG-Nups remains unclear due to a general lack of understanding with regard to their behavior within the NPC. The source of this ambiguity stems in part from the difficulty in addressing FG-Nups dynamics in the intact NPC. The thought-provoking idea here is that we can overcome these limitations by a combination of tracking and fluctuation analysis to study the behavior of single pore components with high spatial and temporal resolution in live, minimally perturbed cells. The crucial point in our results is the observation of a previously hidden dynamics of nucleoporin Nup153, characterized by a rapid, discrete exchange between two separate positions within the NPC of intact cells. The nanomechanical mechanism suggested by our results highlights apparent differences with some macroscopic views proposed thus far that deserve consideration. The bulk-like hydrogel meshwork model (5), for instance, places a greater importance on the hydrophobic interactions between neighbor FG-Nups, as they are able to form a sieve-like meshwork in vitro (10). Transport could then occur through binding of transport receptors to FG-repeats, causing a local gel-to-liquid transition and allowing the receptor to catalyze its own diffusion (28). Our data do not exclude that these interactions play a role in transport; however, they suggest that the most kinetically relevant events in the intact NPC have the characteristics of directed transport, not of unbiased diffusion. It must be noted that Nup153 belongs to the FxFG-rich family of nucleoporins, which generally display noncohesive properties (29). At the same time, however, it was recently reported that C-terminal fragments of Nup153 can exist in a collapsed state but are concomitantly able to form a hydrogel in vitro (30). Besides highlighting the multifaceted character of Nup153, these contrasting results may also reflect a more general complexity of the NPC function, with a not yet completely understood balance/interplay between cohesive (i.e., ‘gel-like’) and noncohesive (i.e., “spring-like”) properties, as also emerges from the hybrid model recently proposed by Yamada and coworkers (15). Based on our results on intact cells, we propose that molecular transport through the NPC can be powered, at least in part, by the directed motion of specific nucleoporins (here Nup153). Further experim...
ments on additional constituents of the pore (e.g., more cohesive Nups of the central channel) are needed to achieve more general conclusions about the intimate nature of selective molecular gating through the NPC. Finally, it is worth noting that the observed directed-motion mechanism shares many biochemical and physical characteristics with several spring-like molecular systems present in nature (31). We shall argue that they represent ancient and commonplace eukaryotic molecular engines.

Materials and Methods

Cell Culture, Plasmids, and Treatments. CHO-K1 cells were grown in Ham’s F12K medium supplemented with 10% of fetal bovine serum at 37 °C and in 5% CO_{2}. Freshly split cells were plated on imaging dishes and transiently transfected using Lipofectamine 2000 according to manufacturer’s protocol, 24 h before the experiment. The plasmid encoding for Nup153-GFP was a kind gift from Nathalie Daigle (Cell Biology and Biophysics Unit, European Molecular Biology Laboratory, Heidelberg, Germany) (20). The plasmid encoding for human Kap1-GFP was a generous gift from Marilena Ciciarello (Istituto di Biologia e Patologia Molecolari, Consiglio Nazionale delle Ricerche) (32). The plasmid encoding for NLS-mCherry has been described in a previous publication (27).

Single NPC Tracking Setup. Tracking of single NPCs was performed using a home-built microscope capable of single particle tracking, whose details have been already described (33). Briefly, the microscope is built around an Olympus X71 body. A Chameleon Ultra (tunable) Ti:Sapphire laser (Coherent) tuned at 940 nm was used for two-photon excitation of the GFP (and mCherry) constructs. The scanning of the excitation light was obtained in the x–y plane and in the z axis through two galvano-motor driven mirrors (Cambridge Technology) and a piezo-objective positioner (Phisik Instrument) respectively, both driven by a computer card (three-axis card, ISS). Fluorescence emission was collected by a 1.2-NA water objective (Olympus UplanSApo 60x), split by a dichroic mirror at 570 nm and detected in the 500–550 nm (GFP) and 575–645 nm (mCherry) spectral ranges by two GaAs detectors H7241P (Hamamatsu). During the tracking procedure, the two scanning mirrors are moved independently by π/2-phase shifted sine wave voltages generated in the card so that the laser beam moves in a circular path around the particle. The position of the scanning center is determined by the offset values of the sine waves. The position of the center is updated at each tracking cycle according to the fast Fourier transform (FFT)-based algorithm previously described (34). From the FFT of the intensity trace along the orbit, we get the average intensity or dc as the zeroth term in a Fourier series and the ac as the coefficient of the first harmonic term. The angular coordinate of the particle is given directly by the phase of the ac term, and its distance from the center...
can be calculated from the modulation of the signal, defined as $a = ac/dc$, so that its position can be recovered. The tracking routine changes the coordinates of the center of the scanning orbit in such a way to keep the modulation at a minimum, i.e., to keep the particle always at the center. The tracking procedure started by acquiring a raster scan image of the sample, focusing on the equatorial section of the nuclear envelope. Then we clicked on a location on the image corresponding to an isolated NPC. The fluorescence intensity is collected at 64 points along a circular orbit around the pore, with a period of 0.5 s and a calibrated starting scan point. The orbit radius (R) is usually set to 120 nm, which we found to be optimal for the tracking. The optimal radius for tracking a point-like particle should be on the order of half the size of the PSF (34), but in our case this value is slightly larger due to the finite size of the NPC. The position of the center of the scanning orbit is updated typically every 32–64 orbits (that is defined as the cycle), which is fast enough to follow the NPC movement. The acquisition time for a single NPC tracking measurement typically varied from 15 s up to 100 s. For each NPC we acquired the intensity along the orbit for one or two channels and the trajectory of the center of mass. From the recorded trace of the fluorescence intensity along the orbit the value of the modulation of the signal can be calculated at each cycle. The analysis of the modulation allows to check a posteriori if the particle has been tracked correctly for the entire acquisition or to exclude the portions of the dataset where the particle was temporarily or definitely lost. We selected only the part of the data acquired with the orbit centered and stationary with respect to the center of mass of the NPC.

**Fluctuation Analysis.** The fluctuation analysis of the data was performed with the SimFCS software (www.lfd.uci.edu, University of California at Irvine) using the scanning-FCS analysis tool, as thoroughly described in previous publications (22, 26). Briefly, the ACF, pCFs (pixels), and pCF(pixels) are displayed in pseudo colors in a carpet in which the x coordinate corresponds to the point along the orbit and the vertical coordinate corresponds to the correlation time in a log-scale.

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