Yeast Nuclear Pore Complexes Have a Cytoplasmic Ring and Internal Filaments

Elena Kiseleva¹,², Terence D. Allen¹, Sandra Rutherford¹, Mirella Bucci³*, Susan R. Wente⁴ and Martin W. Goldberg⁵#

¹. Structural Cell Biology
Paterson Institute for Cancer Research
Christie Hospital NHS Trust
Manchester
M20 4BX
U.K.

². Institute of Cytology and Genetics
Russian Academy of Sciences
Novosibirsk-90
630090
Russia

³. Department of Biological Sciences
Stanford University
Stanford, CA 94305-5020
USA

⁴. Department of Cell and Developmental Biology
Vanderbilt University
3120A Medical Research Building III
Nashville, TN 37212
USA

⁵. University of Durham
Dept. of Biological and Biomedical Sciences,
Science Laboratories, South Road
Durham
Co. Durham
DH1 3LE
UK

* current address: Nature, Nature Publishing Group, The Macmillan Building, 4 Crinan, St London, N1 9XW, UK

# corresponding author,
email address: m.w.goldberg@durham.ac.uk
Tel: +44 (0) 191 334 1250
Fax: +44 (0) 191 334 1201
**Key words:** yeast, nuclear pore complex, field emission scanning electron microscopy, structure  
**Running title:** yeast nuclear pore complex structure  
**Abbreviations:** NPC, nuclear pore complex; feSEM, field emission scanning electron microscopy; TEM, transmission electron microscopy; NE, nuclear envelope; FG, phenylalanine-glycine
Abstract
The nuclear pore complex (NPC) controls transport of macromolecules across the nuclear envelope. It is large and complex but appears to consist of only ~30 different proteins despite its mass of >60 MDa. Vertebrate NPC structure has been analyzed by several methods giving a comprehensive architectural model. Despite our knowledge of yeast nucleoporins, structural data is more limited and suggests the basic organization is similar to vertebrates, but may lack some peripheral and other components. Using field emission scanning electron microscopy to probe NPC structure we found that the yeast, like higher eukaryotic, NPCs contain similar peripheral components. We can detect cytoplasmic rings and evidence of nucleoplasmic rings in yeasts. A filamentous basket is present on the nucleoplasmic face and evidence for cytoplasmic filaments is shown. We observe a central structure, possibly the transporter, that which may be linked to the cytoplasmic ring by internal filaments. Immuno-gold labeling suggests that Nup159p may be attached to the cytoplasmic ring, whereas Nup116p may be associated, partly, with the cytoplasmic filaments. Analysis of a Nup57p mutant suggested a role in maintaining the stability of cytoplasmic components of the NPC. We conclude that peripheral NPC components appear similar in yeasts compared to higher organisms and present a revised model for yeast NPC structural composition.
Introduction

Nuclear pore complexes (NPCs) mediate the exchange of macromolecules across the nuclear envelope (NE). Various soluble transport factors carry substrates to the NPC, where they interact with nucleoporins and mediate translocation through the NPC in specific directions (Mattaj and Englmeier, 1998; Adam, 1999; Talcott and Moore, 1999; Rout and Aitchison, 1999; Allen et al., 2000; Vasu and Forbes, 2001).

The NPCs are located in pores created by the fusion of the inner and outer nuclear membranes (Goldbeg et al., 1999; Stoffler et al., 1999; Kiseleva et al., 2000). In vertebrates (Hinshaw et al., 1992; Akey and Radermacher, 1992; Goldberg and Allen, 1992) and invertebrates (Kiseleva et al., 1998) the center of the pore consists of a disc, termed the inner spoke ring, which has a central aperture in which is located a cylindrical structure called the transporter. The inner spoke ring has eight extensions outwards through the pore membrane into the NE lumen. The two known vertebrate integral membrane nucleoporins, POM121 and gp210, may form part of this transmembrane domain. These transmembrane domains join together in the lumen to form the radial arms.

In vertebrates, on the cytoplasmic face, above the level of the membrane, is an eight subunit ring (Hinshaw et al., 1992; Akey and Radermacher, 1992; Goldberg and Allen, 1992). The ring consists of at least three sequential components: (1) the star ring which lies on and is embedded in the membrane; (2) the thin ring which lies on top of the star ring and then (3) eight bipartite subunits that are molded onto the thin ring (Goldberg and Allen, 1992). The star ring has only been detected by field emission scanning electron microscopy (feSEM) (Goldberg and Allen, 1996) and may form part of the cytoplasmic ring and transmembrane vertical domains seen in 3D reconstructions (Akey and Radermacher, 1993), as the tips of its triangular shaped subunits appear to be embedded in the membrane. Filaments extend from the cytoplasmic ring into the cytoplasm and contain the Ran binding protein Nup358 (Wu et al., 1995; Delphin et al., 1997; Walther and Pickersgill et al., 2002). They appear to have controlled dynamics (Goldberg et al., 2000), but are not essential in at least some nuclear transport pathways (Walther and Pickersgill et al., 2002).

Filaments extending from the cytoplasmic ring into the center of the NPC have also been observed by feSEM (Kiseleva et al., 1998; Goldberg and Allen, 1996). These “internal filaments” could link the cytoplasmic ring to the central transporter. Nup214 and Nup62 and their interacting nucleoporins localize to this central region (Grote et al., 1995). These proteins contain FG repeat domains, which interact with transport factors and may play a direct role in guiding the substrates through the NPC (Rout and Aitchison, 2001; Ribbeck and Gorlich, 2001).

The central transporter, which is a cylinder with a central aperture (Akey and Radermacher, 1993) traverses the NPC and is therefore a major channel between the nuclear and cytoplasmic compartments. The central aperture appears to open and close which offers a possible control mechanism for transport (Kiseleva et al., 1998; Akey, 1990; Feldherr et al., 2001). However, the transporter remains a controversial component, believed to be an artifact by some, and there is some evidence that it is not always present (Stoffler et al., 2003).
On the nucleoplasmic side there is a peripheral ring structure similar to the star ring. Attached to this is a fishtrap- (Ris, 1991) or basket-like structure (Goldberg and Allen, 1992), consisting of eight filaments which join together at the distal basket ring.

The 3D structure of unfixed isolated yeast NPCs (Yang et al., 1998), rapidly frozen and embedded in amorphous ice showed that the core of the yeast structure (the inner spoke ring, transmembrane domain and transporter) was very similar to the vertebrate structure. However, the cytoplasmic ring and filaments and the nucleoplasmic ring and basket were not evident, although weak densities of some structures were detected in these regions. Thin section TEM has been used to demonstrate that filaments are attached to both sides of the NPC in yeast, and that there is probably a nucleoplasmic basket (Fahrenkrog et al., 1998; Rout and Blobel, 1992). However, the extent of conservation of NPC structure between yeast and higher eukaryotes remains uncertain. The 2D projection maps and 3D structures have been calculated from transmitted images of isolated yeast NPCs (Yang et al., 1998). Although this strategy has provided a wealth of information concerning the 3D architecture particularly of the core of the structure, the structures were calculated from many individual NPCs to give a reasonable signal to noise ratio. Therefore, the methods could be biased against structural components that are variable between NPCs or which are flexible and hence do not occupy a clearly defined position. Such structural components would be predicted from the protein composition of the NPC, which includes several proteins which appear to be “mobile” in vertebrates (Nakielny et al., 1999; Griffis et al., 2002) and in yeast (Dilworth et al., 2001; Denning et al., 2001). It is also possible that isolation of NPCs, necessary for these structural studies could disrupt particularly peripheral components.

Because of the experimental advantages of yeast, it is important to determine in finer detail the yeast NPC architecture. FeSEM can be used to image individual NPCs at a resolution comparable to 3D TEM reconstructions (Allen et al., 1997). FeSEM therefore complements the 3D structure of the core obtained by TEM and provides information on the more variable flexible components found at the periphery of the NPC. Although cryo-TEM showed weak peripheral densities, that were consistent with peripheral structures (Yang et al., 1998), which are also seen in thin sections, here we directly demonstrate that these peripheral structures, such as the basket and internal filaments, appear to be very similar to those in vertebrates and other organisms.

The yeast NPC is formed from a series of distinct nucleoporin complexes, each of which may contribute substructure to the NPC. Several nucleoporin subcomplexes have been identified (Grandi et al., 1993; Grandi et al., 1995a; Grandi et al., 1995b; Nehrbass et al., 1996; Zabel et al., 1996; Siniossoglou et al., 1996; Ivone et al., 1997; Bailer et al., 1998; Belgareh et al., 1998; Marelli et al., 1998; Lutzmann et al., 2002). Different nucleoporins have been localised to distinct substructures of both the vertebrate and yeast NPCs (reviewed in Stoffler et al., 1999; Doye and Hurt, 1997; Rout et al., 2000).

In this report we further resolve the structure of NPC substructures in the evolutionarily diverse yeasts Saccharomyces cerevisiae and Schizosaccharomyces pombe by combining new methods for isolation of nuclei with FeSEM. We demonstrate that in both yeast types, cytoplasmic and nucleoplasmic NPC
compartments are comprised of similar individual components that are inherent to vertebrate NPCs. Immuno-feSEM suggests that Nup159p maybe part of the cytoplasmic ring whereas Nup116p could be partly associated with the cytoplasmic filaments. Finally we found that the loss of Nup57p in a temperature sensitive mutant correlates with striking perturbations of the NPC structure at the cytoplasmic face of the NE. This further refinement of both wild type and mutant yeast NPC structure is necessary to couple yeast genetic and biochemical analysis of NPC function with the structural architecture.

RESULTS

NPC numbers

Nuclei were isolated from *Saccharomyces cerevisiae* cells by one of two protocols (Fig. 1): from spheroplasts (Fig. 1b,e,g) or by a freeze-grinding technique (Fig. 1 c,d,f) (Materials and Methods). These two methods gave essentially the same results, and thus *Schizosaccharomyces pombe* nuclei were only isolated from spheroplasts (Fig. 2). NPCs were difficult to make out amongst the densely packed ribosomes, especially in *S. cerevisiae* (Fig. 1a). Therefore ribosomes were partially dissociated with 0.3M NaCl, which resulted in clearly visible NPCs (Figs. 1b-g). The NPCs often appeared in clusters of 3-20 (circled in Fig. 1b), consistent with a previous TEM thin section study (Winey et al., 1997).

The NPCs in nuclei prepared from *S. pombe* were similar to those from *S. cerevisiae* (Fig. 2) but were more easily detected amongst the ribosomes (circled in Fig. 2b). The number of NPCs per nucleus ranged from about 20 to about 40 on the region of the nucleus accessed by feSEM possibly reflecting that cells were at different stages of the cell cycle. These results are in agreement with studies analyzing NPC numbers and distribution in reconstructions of serially sectioned yeast cells (Winey et al., 1997).

Yeast NPCs have a cytoplasmic ring

The structural organization of the NPC was similar in samples prepared either from spheroplasts (Figs.1b,e,g) or by the freeze-grinding technique (Figs. 1c,d,f). The structure was also similar in budding and fission yeasts (Figs 1 and 2, respectively). NPCs have a diameter of ~ 95nm, in the feSEM, which is consistent with 3D reconstructions of frozen-hydrated yeast NPCs (Yang et al., 1998) and smaller than *Xenopus* oocytes NPCs (110-120nm) visualized by the feSEM (Goldberg and Allen, 1996).

In feSEM images there was a bulky cytoplasmic ring structure (Fig. 1b-g), which projected above the level of the outer nuclear membrane. The ring appeared to consist of at least two components, which are shown and illustrated in the insets in Fig. 1c.

Firstly, and most clearly, are eight bulky subunits (Fig. 1c, numbered in the bottom inset and gray in the cartoon) which appear to be equivalent to the cytoplasmic ring subunits in *Xenopus* oocytes NPCs (see Fig.7 for comparison). These subunits are above the level of the membrane, which can be seen most clearly when the NPCs are tilted with respect to the axis of view (e.g. arrow Fig.1b). In addition, extra brightness
around the edge of these particles shows that they have topographical contrast, and therefore are not at the level on the membrane. This is important because it shows that this structure cannot be part of the spoke complex which only extends to the level of the outer nuclear membrane in yeast (Yang et al., 1998) and in *Xenopus* (Akey and Radermacher, 1993; Goldberg and Allen, 1993). Therefore we believe that they are subunits of the cytoplasmic ring. The finding that yeast NPCs have a cytoplasmic ring is significant because although the previous 3D reconstruction data (Yang et al., 1998) showed some density in this region, it did not show a definitive ring structure consisting of subunits. The peripheral density could therefore have been interpreted as other peripheral structures such as filaments being attached directly to the spoke ring complex (accounting for the extra peripheral density). FeSEM data, in contrast, suggest that the cytoplasmic filaments or particles are, like in *Xenopus*, attached to bulky cytoplasmic ring subunits. Additional cytoplasmic filaments/particles are observed attached to these ring subunits (Fig.1c and d, small arrows and Fig.7).

The reason for this discrepancy is not clear but is likely to be related to one or more of the following, which vary between the current and the previous (Yang et al., 1998) studies:
1) isolation of the NPCs (here NPCs are left within the intact isolated nucleus, whereas previously NPCs were isolated);
2) processing for imaging (here they are fixed and critical point dried, whereas previously they were viewed unfixed, frozen hydrated);
3) the imaging itself (surface imaging by feSEM compared to projection imaging).

Because cytoplasmic rings were previously observed in TEM reconstructions in *Xenopus* (Hinshaw et al., 1992; Akey and Radermacher, 1993) it is unlikely that the imaging method could cause this discrepancy. It is possible that the fixation and drying used here could cause collapse and aggregation of peripheral structures to form subunit like structures similar to the *Xenopus* cytoplasmic ring. We do not favor this interpretation because of the apparent similarity in the structure observed here by feSEM compared to that observed in *Xenopus* by both feSEM and cryo-TEM. We favor the interpretation that when NPCs are isolated (Yang et al., 1998) there may be some disruption or disordering of the ring structure, which may preclude the detection of a defined ring and the resolution of individual subunits. It may be that the density detected in this region in 3D reconstructions (Yang et al., 1998) can be attributed to disordered rings and/or cytoplasmic filaments that have collapsed onto them to differing degrees. However, confirmation of the cytoplasmic ring in yeast will require its observation by an alternative method, particularly of sample preparation. To this end we are currently developing methods to examine frozen samples by feSEM.

Thin section TEM studies also suggest a structural similarity between yeast and vertebrates in this region, particularly the presence of cytoplasmic filaments, but could not demonstrate the presence of rings (Fahrenkrog et al., 1998; Rout and Blobel, 1993; Allen and Douglas, 1989).

The size of these putative yeast cytoplasmic ring subunits is also not strikingly different from *Xenopus* oocyte cytoplasmic ring subunits at about 25nm across (Fig.7 and Goldberg and Allen, 1993).
The second component of the cytoplasmic ring that was observed is the “thin ring”. The thin ring underlies the cytoplasmic ring subunits and is observed in *Xenopus* in 3D reconstructions (Akey and Radermacher, 1993), by feSEM after partial proteolysis (Goldberg and Allen, 1996) and as a putative assembly intermediate in *Xenopus* egg extracts and *Drosophila* early embryos (Goldberg and Wiese et al., 1997; Kiseleva et al., 2001). As with other specimens, the thin ring in yeast is difficult to detect in intact NPCs. However, sometimes glimpses of a structure consistent with the thin ring can be observed between two cytoplasmic ring subunits (arrows in the bottom inset in Fig. 1c and colored black in the cartoon in the upper inset). A similar structure is also seen in *S. pombe* (Fig. 2d, arrow). We have also observed a consistent structure in putative incomplete NPCs (see below), which could be the thin ring without the cytoplasmic ring subunits (Fig. 3f and 5b), although it remains to be shown for certain that these are NPC-related substructures. We suggest therefore that, like *Xenopus*, yeasts too have a thin ring, which is an integral part of the cytoplasmic ring.

**Yeast NPCs appear to have cytoplasmic particles or filaments**

In *S. cerevisiae* a variable number of globular particles were found attached to the cytoplasmic ring subunits (Fig. 1d and e, small arrows). It was possible that these were the cytoplasmic filaments. In nuclei isolated from *S. pombe* without 0.3M NaCl, these putative cytoplasmic filaments were more consistently present and had a more extended conformation, appearing more like a rod or filament than a particle (Fig. 2f). The filaments could be quite extended (Fig. 2f and g, marked “1”) or more compact (marked “2” and “3”), which was reminiscent of the apparent conformational flexibility observed in vertebrates (Goldberg et al., 2000; Pante and Aebi, 1996; Rutherford et al., 1997).

Such filaments emanating from the cytoplasmic face of the yeast NPC have previously been suggested from structures observed in TEM thin sections (Fahrenkrog et al., 1998; Rout and Blobel, 1993). The thin section data together with our observation of globular or extended particles attached to the cytoplasmic ring suggests that both yeasts have equivalent cytoplasmic filaments to vertebrates, although it remains to be shown directly whether there are extensive filaments that attach to the cytoplasmic ring and extend some distance into the cytoplasm.

**Central NPC structures**

As feSEM is a surface imaging method it is most suited for visualizing peripheral structures. However, more internal components of the NPC can be detected when NPCs are incomplete, either during assembly (Goldberg and Wiese et al., 1997; Kiseleva et al., 2001) or when they are fractured (Kiseleva et al., 1998; Goldberg and Allen, 1996). These structures include the putative transporter, the inner spoke ring and the internal filaments. We have found evidence for these central core structures in yeast (Fig. 3).

The evidence for these structures comes from images of nuclei that have structures on their cytoplasmic membrane surface, which are consistent with previously identified vertebrate putative assembly intermediates and fractured NPCs, as well as structures that are consistent with components identified in thin ice reconstructions (Yang et al., 1998). These structures could be assembly intermediates or they could be fractured.
NPCs, that have lost certain peripheral components to reveal underlying structure, and we shall refer to them as “intermediates”. Because the intermediates are significantly different in appearance to NPCs it is possible that they are not NPC-related structures. The reason we think they are is because they are about the right diameter and have the characteristic eight fold rotational symmetry of NPCs, and their components are consistent with previously identified NPC substructures. We know of no other similar structure that might be found on the nuclear envelope.

We observed, in regions of some nuclei, a striking structure (Fig. 3a, circles) which, consisted of eight filaments radiating out from the center to give a wagon wheel like structure. Such structures were also observed in Chironomus tentans salivary gland NEs after NPCs were fractured by “rolling” the nucleus (Fig 3e). The filaments are consistent with the internal filaments identified in Xenopus and other organisms (Goldberg and Allen, 1996: Kiseleva et al., 1998; Goldberg et al., 1997; Kiseleva et al., 2001). Again, this may not be an NPC-related structure, but the diameter and the fact that there are eight radiating filaments is suggestive of a NPC-related structure and not suggestive of any other known different structure.

Internal filaments were difficult to identify in complete NPCs, although we did find evidence of them (Fig. 3b smaller arrows). It may be that they cannot be detected easily in complete NPCs because they are obscured by the overlying cytoplasmic ring subunits. The internal filaments are also clear in putative “intermediates” where the thin ring appears to be present but the cytoplasmic ring subunits do not (Fig. 3f). It should be noted that internal filaments have not been observed by any method other than feSEM and it could be argued that they may be artifacts of fixation or critical point drying. It is difficult to avoid chemical fixation in feSEM preparation, but we have observed internal filaments, by feSEM, in frozen hydrated specimens, showing that this potential artifact is not caused by critical point drying (unpublished result).

In Xenopus the internal filaments appear to attach to the “top” of the transporter, which is a controversial cylindrical structure found in the central channel of the NPC (Akey and Radermacher, 1993; Kiseleva et al., 1993; Goldberg and Allen, 1996; see also Stoffler at al., 2003). In yeast a central mass is also observed (Fig. 1g large arrows) and could represents the top part of the central cylinder or transporter. In some NPCs the central globule protrudes from the central channel (Fig. 1f and g small arrows). Such a structure was also reported in Chironomus and was suggested to be a peripheral component of the transporter (Kiseleva et al., 1993), but alternatively it could be material, such as mRNP, in transit.

The nucleoplasmic face of the NPC

To reveal the nucleoplasmic face of the yeast NPC, nuclei were fractured by centrifugation at 10,000 X g to disrupt the NE. This revealed NEs where the nucleoplasmic face of NPCs were either mostly obscured by intranuclear material (Fig. 4a) or NEs where some NPCs were visible and some were damaged (Fig. 4b). However some NPCs retained a morphology that was similar to that in Xenopus and other previously studied organisms. Yeast NPC baskets were however mostly at least partially damaged compared to Xenopus. This is mostly likely because manual isolation of Xenopus NEs is very mild compared to exposure of the nucleoplasmic face of the yeast NE.
As with all other organisms studied so far, we found filaments attached to the outer periphery of the NPC and extending into the nucleus (Fig. 4c white arrows). Sometimes these filaments joined together to form the distinctive basket or fish-trap structure (Goldberg and Allen, 1992; 1996; Ris, 1991). Yeast baskets have previously been suggested from analysis of TEM thin sections (Stoffler et al., 1999; Fahrenkrog et al., 1998; Rout and Blobel, 1993), but we can show here that the yeast basket appears to be very similar to that of vertebrates and other organisms. There are up to eight basket filaments, which are about 10nm in diameter and up to 40nm long, which is similar to Xenopus when viewed by feSEM (Goldberg and Allen, 1996).

Evidence for the nucleoplasmic ring could also be observed (Fig. 4c black arrows). The nucleoplasmic ring is less massive than the cytoplasmic ring (Akey and Radermacher, 1993; Reichelt et al., 1990) and is not always so clearly observed, even in Xenopus oocyte NEs. In S. cerevisiae we can detect an electron emission from some topographical structure between the filaments of the basket, where they appose the membrane (black arrows in Fig.4c), which is consistent with the nucleoplasmic ring. The ring is not always clearly visible and requires careful inspection but can sometimes be seen, even at low magnification (Fig. 4b, small arrows). As the structure is not clearly visualized we cannot say whether its morphology is similar to other organisms. What we can see, suggests that it may be a simpler smooth ring, rather than having the eight clear subunits observed in Xenopus. The putative nucleoplasmic ring diameter is 90-100nm which is larger than that previously reported for the nuclear membrane pore (Yang et al., 1998). This suggests that, in our view, it is unlikely to be part of the spoke ring complex, which lies within the pore.

Occasionally, thick filaments of about 20nm in diameter were attached to the top of some baskets (Fig. 4b large arrows). These could correspond to filamentous structures stretching from the nucleoplasmic face of the NE into the nucleoplasm as described previously in budding yeast (Strambio-de-Castillia et al., 1999) and could represent a structure similar to the NE lattice and “cables” observed in Xenopus (Goldberg and Allen, 1996;Ris and Malecki, 1993). Alternatively they could be remnants of chromatin.

**Role of specific nucleoporins in the NPC**

Having shown that yeast NPCs do have cytoplasmic rings and filaments we wanted to determine what proteins were involved in these structures. Immuno-gold labeling can help to determine the location of a protein, whereas the structural analysis of NPCs containing mutant proteins gives clues as to the structural role of a given protein or subcomplex.

**Nup159p may be associated with the cytoplasmic ring**

Anti-Nup159p labeling was observed at a low level on most nuclei (not shown), and was increased slightly by a 0.3M NaCl wash. The highest level of labeling was found on NPCs that appeared incomplete which was about 10 fold more gold particles per NPC (1-4 particles/NPC or 30-35 particles/half nucleus compared to 0-3 particles/nucleus in the control where the primary antibody was omitted) than in complete NPCs. We speculate that incomplete NPCs could be assembly intermediates
(Goldberg and Wiese et al., 1997; Kiseleva et al., 2001) or damaged NPCs. Our interpretation of this is that the Nup159p epitope is not accessible in complete NPCs but would be made partially accessible by the salt wash and is most accessible when certain components are missing, such as during assembly or disassembly, or if they are damaged during isolation.

The position of the labeling was at the periphery of the cytoplasmic face. Labeling was not in the central channel (Fig. 5a and b), so we can eliminate a location at the central transporter. Such a position is consistent with the location at the cytoplasmic ring or the cytoplasmic filaments. However the fact that the level of labeling is greater when more peripheral structures, such as cytoplasmic filaments, are missing suggests that it is not part of the filaments. The labeling is therefore consistent with a location on the cytoplasmic ring.

**Nup116p may be associated with the cytoplasmic filaments**

An anti-Nup116p antibody (which is specific on Western blots and immunoprecipitation) did label intact NPCs at the periphery of the cytoplasmic face (Fig. 5c-f and see Ho et al., 2000), showing that the epitope is more accessible. In general the labeling appears to be particularly peripheral and possibly associated with the cytoplasmic filaments (see particularly Figs 5e and f). However, this only represents one position of Nup116p, as thin section TEM shows that it is also in the central domain and on the nucleoplasmic side (Ho et al., 2000), although it is biased to the cytoplasmic side (Rout et al., 2000).

**FeSEM analysis of a mutant with perturbations in NPC function**

Analysis of nucleoporin mutants will provide essential information for understanding the mechanism of both the nuclear transport and NPC structure. Here we looked at the temperature sensitive nup57- E17 mutant (Bucci and Wente, 1998). We have focused on this mutant for several reasons. First, protein-protein interactions between Nup57p and two other essential S. cerevisiae nucleoporins (Nup49p and Nsp1p) have been defined (Grandi et al., 1995b; Shlaich et al., 1997). Second, the nup57- E17 mutant strain has been extensively characterized for perturbations in nuclear transport and NPC composition (Bucci and Wente, 1998). Temperature sensitive growth arrest of nup57- E17 cells is coincident with specific defects in nuclear import. The NE in temperature arrested nup57- E17 cells also results in a small number of “herniations” previously observed by TEM as membranous seals over the NPC cytoplasmic face (Bucci and Wente, 1998).

We isolated nuclei from nup57- E17 cells that were grown at:
1. 23°C (Fig. 6a);
2. 37°C for 2 hours (Fig. 6b);
3. 37°C for 5 hours (Fig. 6c).

At 23°C the NPCs appeared quite normal (Fig. 6a and d) compared to NPCs with wild type nucleoporins (Fig. 1). There may however be some subtle differences. In particular, the NPCs do not appear to be as consistent or symmetric, suggesting they may be more fragile. A temperature shift to 37°C for 2 hours caused dramatic changes (Fig. 6b and e), the most obvious of which were the loss of internal structures (transporter and internal filaments) and removal of the cytoplasmic filaments. The cytoplasmic ring appeared to be present in most NPC-like structures (Fig. 6b arrows)
but sometimes was partially fragmented. The ring appeared less massive as it had less topography and emitted less signal, possibly because the cytoplasmic filaments were missing. The channel through the pore appeared to be stabilized at a fairly consistent size, suggesting that components of the spoke ring complex were still intact. However, after 5 hours at 37°C (Fig. 6c and f) most of the remaining recognizable structures were gone and the “hole” apparently closed up (Fig 6c arrows), leaving structures that resembled the “dimples” previously suggested to be early assembly intermediates (Goldberg and Wiese et al., 1997). NPCs of nuclei isolated from wild type S. cerevisiae, that were incubated at 37°C were identical to those incubated at 23°C, showing that the structural alterations were due to the mutation.

These results show that disruption of Nup57p function initially causes a disruption of the internal structures, as well as parts of the cytoplasmic ring (after 2 hours), but ultimately the whole cytoplasmic side of the NPC is destabilized (after 5 hours). This final step could represent an intermediate in the herniation of the NE observed in TEM thin sections (Wente and Blobel, 1993). As all NPCs are affected in this way these effects are most likely due to disruption of pre-existing NPCs rather than mis-assembly of new ones, where only a proportion would be affected.

Discussion

Yeast has proved an invaluable tool for identifying nucleoporins and obtaining information about their functional roles. However, until recently little was known about the structure of the yeast NPC and how it related to the well defined structure of the vertebrate NPC. TEM reconstructions provided the first 3D information for the yeast NPC (Yang et al., 1998) and it appeared to be significantly smaller and simpler than in vertebrates (Akey and Radermacher, 1993). This provided a puzzle because recent proteomic analysis of both yeast (Rout et al., 2000) and rat (Cronshaw et al., 2002) NPCs showed that they both contained a similar complexity of constituent proteins. One explanation for this is that the 3D reconstructions (Yang et al., 1998) contained data for part of the structure (the spoke ring complex and transporter) but not for the more peripheral components (cytoplasmic ring and filaments and nucleoplasmic ring and basket). This could be because they do not occupy a defined position and are therefore not clearly resolved, or that they are disrupted, disordered or lost during isolation or distorted upon interaction with carbon grid during freezing. On the other hand, biochemical data (Rout et al., 2000) does suggest that there is not an overall loss of nucleoporins upon isolation. It is difficult, however, to be sure that there is no loss of unknown nucleoporins or partial loss of others, which could result in the loss of a structural component without loosing all its constituent proteins. Thin section data (Fahrenkrog et al., 1998) showed the attachment of filaments to both sides of the yeast NPC, but components such as the cytoplasmic and nucleoplasmic rings, the transporter and internal filaments are not readily visible in thin sections.

Yeast and Vertebrate NPCs are structurally similar

Here we have used feSEM to visualize the cytoplasmic face of individual NPCs in situ in intact nuclei from fission and budding yeast. This has allowed us to look at the surface of relatively intact NPCs. We believe, from this data, that the yeast NPC, like that of other organisms so far studied, does have a cytoplasmic ring (consisting of a thin ring and subunits) as well as cytoplasmic filaments and internal filaments. By
disrupting the nucleus and exposing the nucleoplasmic face we also found evidence that the yeast NPC has a basket and possibly a nucleoplasmic ring, but these were difficult to preserve and image. A comparison of yeast and *Xenopus* oocyte NPCs is shown in Fig. 7.

The data presented here show, by feSEM, that the peripheral components of yeast NPCs are morphologically similar to other organisms so far studied. This view is supported by thin section data (Fahrenkrog et al., 1998; Rout and Blobel, 1993). However 3D reconstructions (Yang et al., 1998) did not detect cytoplasmic or nucleoplasmic rings or baskets, but rather a weak density in these regions, which were attributed to collapsed peripheral filaments. Here we show evidence, at least viewed by feSEM, that there are cytoplasmic rings and filaments, which are very similar to those observed in *Xenopus*. We favour the view that these structures are lost or disordered when the yeast NPCs are isolated from the nucleus, which is avoided in work presented here. Alternatively, it is possible that the ring, which we observe, is an artifactual structure formed by chemical fixation and drying. We do not favour this view because the structure is so similar to the *Xenopus* cytoplasmic ring seen by feSEM. This in turn is the same as the ring clearly shown in unfixed frozen hydrated *Xenopus* oocyte NEs (Akey and Radermacher, 1993) and negatively stained NEs (Hinshaw et al., 1992) in TEM reconstructions.

The nucleoplasmic face of the NPC appears to be more difficult to preserve, essentially because we have to use harsher conditions to disrupt the NE and remove the associated nuclear contents in order to expose the nucleoplasmic face of the NPC. For instance it was recently shown that Mlp1p and Mlp2p, which are the yeast Tpr homologues, organise telomeres to the NPC (Galy et al., 2000). Therefore removal of the NE from the chromosomes is likely to disrupt components of the NPC that contain these proteins, such as the basket. Despite this, by looking at NEs that have been differently disrupted, we have found structures that appear to be baskets. This was no great surprise as fairly convincing thin section data showing possible baskets has previously been published (Fahrenkrog et al., 1998; Rout and Blobel, 1993). We also show evidence for nucleoplasmic rings. These appear to be thin and faintly contrasted in the feSEM. This could be because they, too, are disrupted or extracted in the isolation of the NE. Alternatively they could in fact be significantly different to *Xenopus* nucleoplasmic rings. In particular they could be much less massive.

**FeSEM Immunolocalisation**

Having established the structural organisation of the yeast NPC, our long-term objective is to determine which nucleoporins were involved in which structural components. This is important because although immuno-TEM has been used to locate yeast nucleoporins, the structural components of the NPC cannot be directly visualized in thin sections. The immuno-feSEM presented here also has its limitations, chiefly that the nuclear surface has to be exposed, providing the opportunity for loss or rearrangement of epitopes. With further work it should also be possible to increase the resolution of the labeling, by direct labeling and the use of smaller probes.

First we looked at Nup159p, which is part of the Nup82p-Nup159p-Nsp1p complex (Belgareh et al., 1998) and is thought to be the yeast homologue of vertebrate Nup214/CAN. In yeast, Nup159p localises to the cytoplasmic face (Rout et al., 2000),
as does Nup214 in vertebrates (Kraemer et al., 1994). Nup214 is localised quite centrally (Walther and Pickersgill et al., 2002), possibly on the inner rim of the cytoplasmic ring. Here we confirm that in budding yeast Nup159p may also be on the cytoplasmic ring. Removal of Nup214 from Xenopus nuclear assembly extracts resulted in the assembly of NPCs with apparently normal cytoplasmic rings and filaments (Walther and Pickersgill et al., 2002). Likewise TEM thin sections (Gorsch et al., 1995) could not detect any structural effect of a Nup159p mutation. The mutation did however inhibit mRNA export, but not import, which is also consistent with Nup214 (Walther and Pickersgill et al., 2002). The results presented here, therefore corroborate the view that yeast Nup159p is the functional homologue of vertebrate Nup214/CAN.

**Temperate sensitive mutant of Nup57**

Like Nup159p, Nup57p also forms a complex with Nsp1, which is distinct from the Nup159p complex above and contains Nup49p. A temperature sensitive mutation in Nup57p resulted in NPCs that lack internal structures after 2 hours at the non-permissive temperature, but retained elements of the cytoplasmic ring. It was previously shown that this mutation perturbed the incorporation of not only the Nup57-Nsp1-Nup49 complex but also of Nup116p (Bucci and Wente, 1998). The lack of Nup116p is consistent with the lack of cytoplasmic filaments in this nup57-E17 mutant at the non-permissive temperature. The mutation however permitted the incorporation of the Nup159-Nup82-Nsp1 complex, which is consistent with our finding that the cytoplasmic ring is at least partially present after 2 hours at 37°C.

Nup62 is the proposed vertebrate orthologue of Nsp1. In two similar studies, immunodepletion of Nup62 from Xenopus egg nuclear assembly extracts resulted either in an apparent inhibition of NPC assembly (Dabauvalle et al., 1990) or assembly of non-functional NPCs that were missing central structures (Finlay et al., 1991). Although apparently inconsistent with each other, these results are both similar to certain aspects to our findings. After 2 hours at the non-permissive temperature our mutant had lost its central structures, presumably as the Nup57-Nsp1-Nup49 complex disassociated. Then after 5 hours the cytoplasmic ring and at least parts of the spoke ring complex dismantled, presumably because, although these structures are unlikely to contain the Nsp1 complex, they must rely on it for their continued stability. The position of Nup57p labelling (Fahrenkrog et al., 1998; Rout et al., 2000) is consistent with the internal filaments that link the transporter to the cytoplasmic ring. Likewise Nup62 is in a similar position (Guan et al., 1995) and certainly in the region of the internal filaments. Therefore it is possible that the internal filaments consist of the Nup57-Nsp1-Nup82 complex. So when this linkage is broken, in the nup57-E17 temperature sensitive mutant at 37°C (because Nup57p holds together the Nsp1 complex - Schlaich et al., 1997), the structures that it may link together (the transporter and the cytoplasmic ring) are no longer stably associated. Indeed the transporter and the internal filaments, are the first structures to disappear at 37°C. We therefore speculate that the Nup57-Nsp1-Nup49 complex could be a component of the internal filaments. The predicted coiled-coil nature of all the components of this complex would be consistent with them forming a filament.

**Cytoplasmic filaments**

One surprising result is that yeast NPCs contain cytoplasmic filaments, which appear to be very similar to those of vertebrates, although such filaments have been indicated
in thin sections (Fahrenkrog et al., 1998). Firstly, they are not detected in 3D reconstructions (Yang et al., 1998). Secondly, it has been shown that the Ran binding nucleoporin, Nup358, is a major constituent of the filaments (Wu et al., 1995; Delphin et al., 1997; Walther and Pickersgill et al., 2002), possibly even the filament itself, and there is apparently no similar protein in yeast. Despite the fact that Nup358 appears to be the major component of cytoplasmic filaments in the vertebrate NPC (Walther and Pickersgill et al., 2002) in yeast a similar filament structure could be constructed from one or more different proteins. As Nup358 is a multidomain, multifunctional giant protein, it is entirely possible that several smaller proteins could execute the same function.

In summary, we have shown that NPC structure, particularly with respect to the peripheral components, is very similar in both budding and fission yeast compared to vertebrates and invertebrates. We have shown evidence for yeast cytoplasmic and possibly nucleoplasmic rings and internal filaments. We have confirmed evidence that the basket exists in yeast and probably has the same structure as in other organisms. Likewise we have confirmed that there are cytoplasmic filaments and shown that they have a similar conformation as in other organisms. Nup159p, like its putative vertebrate homologue Nup214, appears to be associated with the cytoplasmic ring. Nup116p appears to be associated with the cytoplasmic filaments, although it also occupies other positions. We have also shown that without Nup57p, which is required to stabilise the Nup57-Nsp1-Nup49 complex, the internal filaments and the transporter are absent. With the tools available to yeast cell biologists, such as antibodies, tagged nucleoporins, temperature sensitive, deletion, and other nucleoporin mutants, the approach shown here could be a useful way to determine the roles of nucleoporins in the structural organisation and function of the NPC.

MATERIALS AND METHODS

Strain growth, media and reagents

Saccharomyces cerevisiae wild type strains AB1380 (Mata ade2-1 ura3-1 trp1 lys2 his5 can1-100) and J 1003-10 (Mata ade2-1 ura3-1 trp1-1 leu2-3,112 can1-100) and Schizosaccharomyces pombe wild type strain HM123 h leu1-32 (lab stock) were analyzed. In addition, the yeast strain SWY1587 (Mata ura3-1 his3-11,15 trp1-1 leu2-3,112 can1-100 ade2-1:ADE2 nup57-E17 ) was analyzed at the permissive (23°C) and restrictive (37°C) growth temperatures as described (Bucci and Wente, 1998). The cells were grown in liquid YPD medium (1% yeast extract, 2% bactopeptone, 2% glucose). A stationary yeast culture was diluted at 1:2000 in YPD medium and grown for ~10 hours at 30° C with moderate shaking (130 rpm) to logarithmic phase and cells were used when about 80% of cells were dividing. Reagents were obtained as follows: Zymolase 20T (Sigma), dithiothreitol (Sigma), Novozyme (Sigma); Sucrose ultrapure (Agar), sorbitol (Serva), Poly-L-lysine and Hoechst 33258 (Sigma).

Isolation of nuclei

Two methods of nuclear isolation were used: from spheroplasts (method A) and from whole frozen cells (method B). The spheroplast methods for budding and fission yeast were modified from previously described studies (Rozijn and Tonino, 1964; Hurt, 1988). About 3 g weight of wet yeast cells were harvested by centrifugation in a
Sorvall GS3-rotor, washed once in 100 ml water and incubated in 30 ml 0.1M Tris-HCl, pH 9.4, 10 mM dithiothreitol for 25 min at 30°C with moderate shaking. Cells were pelleted, washed in 50 ml spheroplast buffer (1.2M sorbitol, 20 mM potassium phosphate, pH 7.4) and resuspended in 25 ml spheroplast buffer containing 1 mg zymolase 20T per gram of wet cells for budding yeast and 10mg/ml Novozym for fission yeast, then incubated for 20-40 min at 30°C with gentle shaking. When about 70% of the cells were converted to spheroplasts, they were pelleted for 3 min at 3500 rpm in a Sorvall SS34 rotor and washed twice in spheroplast buffer. The final pellet was resuspended at room temperature in 10 ml spheroplast buffer with 0.5 mM MgCl₂ pH 6.5. All subsequent steps were at 4°C. 200μl of spheroplast homogenate were transferred into a glass tube and diluted with 200μl distilled water. This was incubated for 2 min to allow the spheroplasts to swell and was then vortexed for 2 seconds. The specimens were then checked using light and fluorescence microscopy by staining with 5 μg/ml Hoechst 33258. Before or after additional incubation of the samples for 10 min with 0.3 M NaCl in 20 mM potassium phosphate pH 6.5 to remove ribosomes from the NE surface, they were immediately prepared for the feSEM (below).

Nuclear isolation from whole cells involved manual grinding of frozen yeast cells, and was modified from a previously described method (den Hollander et al., 1986). Yeast cells were suspended in 1.2 M sorbitol, 2 M sucrose, 20 mM potassium phosphate, 0.5 mM MgCl₂ pH 6.5 and frozen as small drops in liquid nitrogen. Frozen drops covered by liquid nitrogen were ground into powder with a pestle and fixed immediately (see fixative solution below) after thawing to 4°C.

**Preparation of samples for feSEM**

For feSEM, 5mm x 5mm silicon chips were treated with 1 mg/ml poly-L-lysine in distilled water for 30 min, then rinsed in water and transferred to buffer containing 20 mM potassium phosphate, 0.5 mM MgCl₂, pH 6.5. 10μl of the samples prepared by method A or by method B were layered on top of 40μl fixative (20 mM potassium phosphate, 0.5 mM MgCl₂, 0.2M sucrose, 4% paraformaldehyde, pH 6.5) and spun onto a silicon chip (Agar) at 4000 X g for 3 minutes in a swing-out rotor. During this centrifugation, spheroplast membranes in samples from methods A were broken due to impact with the chip surface, and their contents were released and fixed immediately on the chips. To visualize the nucleoplasmic side of the NPC, the spheroplast samples were pelleted for 3 min at 10,000 X g to disrupt the NE. After centrifugation, the chips were further fixed in 2% glutaraldehyde, 0.2% tannic acid, 20 mM potassium phosphate, 0.5 mM MgCl₂, pH 7 for 10 minutes at room temperature, postfixed with 1% (w/v) osmium tetroxide for 10 minutes, then stained with 1% (w/v) uranyl acetate for 10 minutes (both solutions in distilled water), dehydrated in ethanol and critical-point dried from CO₂ via Ark lone (ICI, Runcorn, Cheshire, UK). Specimens were sputter coated with 2-4 nm of tantalum in an Edwards 306 cryo-pumped vacuum system with a magnetron head (Edwards High Vacuum International, Crawley, West Sussex, UK). Samples were examined at 30kV in the top stage of a Topcon (ABT) ISI DS-130F field emission scanning electron microscope (Topcon Corporation, Tokyo, Japan). Measurements of individual NPC components were done on micrographs of 20 different samples prepared by spheroplast or freeze/grind methods.
Preparation of samples for immuno-feSEM
Yeast nuclei before or after removal of the ribosomes were isolated from spheroplasts by method A and centrifuged onto silicon chips for 3 min at 4000 g through buffer A (20mM Tris-HCl pH 7.5, 0.5mM MgCl₂, 0.2 M Sucrose). Samples were then transferred to buffer A without sucrose containing 3.7% formaldehyde and fixed for 15 min, and then given two washes in buffer A without sucrose for 5 min. Samples were blocked for 20 min in 1% BSA, 20 mM Tris-HCl, then incubated (with shaking) for 60 min with either the rabbit anti-Nup116-C antibody (1:10 dilution) (raised against the C-terminal 60kD domain and specific for Nup116p by Western blots and immunoprecipitation and affinity purified against the cognate epitope) or mouse Mab against Nup159p (Mab 165C10; 1:50 - Kraemer et al., 1995), or with buffer as a control, and washed twice for 5 min with 0.001% (wt/vol) Tween 20, 20 mM Tris-HCl. The samples were incubated for 30 min with secondary gold-conjugated goat anti-rabbit or anti-mouse antibody (AuroProb TM EM GAR IgG G10; 10 nm in diameter; Amersham) in 0.2% BSA-20 mM Tris-HCl (1:20). The samples were washed twice with 20mM Tris-HCl and fixed for 10 min in 2% (wt/vol) glutaraldehyde, 0.2% (wt/vol) tannic acid in 20 mM Tris-HCl, rinsed with buffer, and incubated for 10 min in 1% OsO₄ in water. Further processing and feSEM analysis were performed as described above, except that after critical point drying, the samples were coated with 4 nm of chromium. The backscattered electron imaging mode was used to resolve the gold particles by feSEM, using a solid-state retractable backscatter electron detector in the top stage of the TOPCON (ISI) DS 130F SEM. To determine the position of the gold particles, backscatter electron images were precisely overlaid onto the secondary image in Adobe Photoshop, the position of the colloidal gold signal marked with a black dot, then the backscatter image layer (which obscures some of the secondary electron detail) removed.

Preparation of Chironomus nuclei
Isolated salivary gland cells were kept in cold TKM buffer (100mM KCl, 1mM MgCl₂, and 10mM trietahanolamine-HCl, pH 7.0), supplemented with 2% NP-40 for 10-30 seconds and then transferred into 0.025% NP-40. The nuclei were isolated from glands by pipetting. To remove the cytoplasmic rings from the NPC, the nuclei were allowed to adhere to the silicon chip and then gently rolled over the surface by pushing with a glass needle as described previously (Kiseleva et al., 1998). The samples were fixed and processed as described for the yeast nuclei.

Acknowledgements
We thank Dr. M. Rout for advice on the use of the freeze/grind method for nuclear isolation from yeast cells. We also thank our colleagues in the Wente laboratory (A. Ho, E. Ives, Y. Lee, K. Ryan, and L. Strawn) for critical discussions and comments on the manuscript. The research was supported by the Wellcome Trust Foundation (E.K., M.W.G. and T.D.A.), Cancer Research UK (T. D. A. and M.W.G.), the Russian Foundation for Basic Research (E.K.), a Beckman Young Investigator Award from the Arnold and Mabel Beckman Foundation (S.R.W.), and a grant from the National Institutes of Health (S.R.W.). Artwork (Fig. 7) was by P. Chantry.


Figure legends

Figure 1. *S. cerevisiae* nuclei isolated with ribosomes still attached (a) where the NPCs (arrows and inset) are difficult to make out and (b-g) with the ribosomes removed with 0.3M NaCl. Nuclei were isolated either from spheroplasts (a,b,e,g) or by a freeze-grinding technique (c,d,f) with similar results. NPCs were sometimes clustered (b – circles). Previously recognised peripheral NPC components are observed (c-g). Components include the eight subunit cytoplasmic ring (c - numbered in bottom inset and grey in top inset, which is a cartoon of the image in the bottom inset); the thin ring (c – white arrows in bottom inset and black in top inset); cytoplasmic particles/filaments (smaller arrows in d and e); a central mass (large arrows in e and g) and a central globule (small arrows f and g). Bars = 100nm

Figure 2. *S. pombe* nuclei isolated from spheroplasts. (a) Nucleus protruding from broken cell. (b) NPCs (circled) are seen amongst the ribosomes (arrows). (c-e) Examples of NPCs showing the eight subunits (numbered) and the thin ring (arrow). Sometimes the filamentous nature of the cytoplasmic particles is evident (f) and can be compact (illustrated in g “3”) or more extended (illustrated in g “1”). Bar in (a) = 500nm; Other Bars = 100nm.

Figure 3. Evidence for internal filaments can be seen in intact NPCs (b small arrows). Structures were also found that could be internal filaments of incomplete NPCs (a - circled) where the cytoplasmic ring has not yet formed or detached. These structures consist of eight radiating filaments and have a diameter of about 90nm, which is suggestive of an NPC structure. Examples of these structures are shown at high magnification (c and d) and an equivalent structure that was found in *Chironomus tentans* salivary glands is shown in (e). Structures were also observed that were similar but with an apparent peripheral ring, which could be the thin ring, as illustrated (f). Bars = 100nm. Bar in (f) refers to (b-f).

Figure 4. Nucleoplasmic face of *S. cerevisiae* NE from ruptured nuclei. Often NPCs are obscured by overlying material (a), but baskets (b - circled) and the nucleoplasmic ring (b - small arrows) can be observed as well as filaments that interconnect the distal parts of the baskets (b - large arrows). At high magnification (c) the basket filaments (white arrows) and nucleoplasmic ring (black arrows) were evident. Bars = 100nm

Figure 5. Immuno-fESEM of Nup159p (a and b) and Nup116p (c -f). (a) Some NPCs (circled) are labelled by indirect immuno-gold labelling (black dots). The position of the label appears peripheral and NPCs are most often labelled when they appear incomplete (b). Nup116p labelling however could be more peripheral and labels complete NPCs more frequently (c-f). Bars = 100nm

Figure 6. The *nup57-E17* temperature sensitive mutant maintained at 23°C (a and d), 37°C for 2 hours (b and e) and 37°C for 5 hours (c and f). At 23°C, the NPCs appeared quite normal. After 2 hours of cell growth at non-permissive temperature the NPCs lost central structures (transporter and internal filaments). Only small holes or dimples of about 20nm in diameter instead of NPCs were observed in the nuclear envelope after 5 hours of cell growth at 37°C. Bar = 100nm
Figure 7. Model (adapted from Yang et al., 1998) and images of the cytoplasmic face of the NPC comparing yeast and *Xenopus*. CF = cytoplasmic filaments; CRS = cytoplasmic ring subunits; TR = thin ring; ONM = outer nuclear membrane; IF = internal filaments; MR = membrane ring; T = transporter; ISD = inner spoke domain; INM = inner nuclear membrane; NR = nucleoplasmic ring; B = basket. Coloured components are those observed by feSEM, although the basket had previously been indicated and the nucleoplasmic internal filaments remain a speculation. Bar = 50nm
Figure 1
Figure 2

(a) Nucleus

(b) Detailed view of cellular structures

(c) Magnified view of specific areas

(d) Enhanced focus on certain regions

(e) Further magnification of interest

(f) Additional details

(g) Illustrative model

Figure 2
Figure 5
Figure 7