

## Identification of Karyopherins involved in the nuclear import of RNA exosome subunit Rrp6 in *Saccharomyces cerevisiae*

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### ABSTRACT

The exosome is a conserved multiprotein complex essential for RNA processing and degradation. The nuclear exosome is a key factor for pre-rRNA processing through the activity of its catalytic subunits, Rrp6 and Rrp44. In *Saccharomyces cerevisiae*, Rrp6 is exclusively nuclear and has been shown to interact with exosome cofactors. With the aim of analyzing proteins associated with the nuclear exosome, in this work, we purified the complex with Rrp6-TAP, identified the co-purified proteins by mass spectrometry, and found karyopherins to be one of the major groups of proteins enriched in the samples. By investigating the biological importance of these protein interactions, we identified Srp1, Kap95 and Sxm1 as the most important karyopherins for Rrp6 nuclear import and the nuclear localization signals recognized by them. Based on the results shown here, we propose a model of multiple pathways for the transport of Rrp6 to the nucleus.

### INTRODUCTION

The RNA exosome is a protein complex involved in processing and degradation of different classes of RNA in the cell. This complex was first identified in the yeast *Saccharomyces cerevisiae* (1) and later identified in other eukaryotes. The exosome is not present in bacteria, but has been identified in archaea, being structurally conserved throughout evolution (2).

In eukaryotes, the exosome is present in both nucleus and cytoplasm and its nuclease activity is provided by two catalytic subunits, Rrp44/Dis3 and Rrp6 (3). In yeast, the difference between the nuclear and cytoplasmic exosomes is the presence of the subunit Rrp6 in the nucleus. The exosome core is composed of nine subunits: six different subunits each containing an inactive *RNase PH*

domain (Rrp41, Rrp42, Rrp43, Rrp45, Rrp46, Mtr3), and three RNA binding subunits (Rrp4, Rrp40, Csl4). The catalytically active subunits bind to opposite sides of the core, Rrp44 is an RNase R-like with both endonucleolytic and processive 3'-5' exonucleolytic activities (3-6), whereas Rrp6 shows a distributive 3'-5' exonucleolytic activity (7).

Nuclear exosome function comprises processing of ribosomal RNAs (rRNAs), small nuclear RNAs (snRNAs) and small nucleolar RNAs (snoRNAs), as well as surveillance and degradation of incorrectly processed RNAs (8). In the pre-rRNA processing pathway, the exosome is directly responsible for the degradation of the 5' external transcribed spacer sequence, 5'-ETS, after cleavage at site A<sub>0</sub>, and for trimming of the internal transcribed sequence ITS2 segment present in the intermediate 7S for the generation of the mature

5.8S rRNA (9). In addition to being important for the RNA quality control in the nucleus, the exosome has also been described to be involved in cytoplasmic mRNA degradation (10,11).

Since the exosome does not show substrate specificity *in vitro*, the recruitment of the complex *in vivo* might be performed by its cofactors (12). Interestingly, most of the proteins identified as nuclear exosome cofactors have been shown to interact with Rrp6 (13-15). Due to the low concentration of Rrp6 in the cell, or to its restriction to the cell nucleus, however, attempts to isolate nuclear exosome cofactors copurifying with the exosome when using a core exosome subunit as bait has been shown to be inefficient (16,17).

A significant amount of information is now available on the structure and function of the exosome, but relatively little is known about the transport of this complex to the nucleus. Protein transport from cytoplasm to nucleus is primarily mediated by specific interactions between karyopherins and signal sequences (NLS; Nuclear Localization Signal) present in the cargo proteins (18,19). The first identified sequence was the simian virus 40 large tumor antigen (SV40TAg)-like nuclear localization signal (classical NLS), which is recognized by karyopherin  $\alpha$  (Srp1 in yeast) and transported to the nucleus as a trimeric complex with the karyopherin  $\beta$  (Kap95 in yeast) and the cargo protein (19,20). Another nuclear import route occurs through the recognition of a different NLS in the cargo by a karyopherin  $\beta$  and the transport of a heterodimer to the nucleus (21). In *Saccharomyces cerevisiae*, there are 14 karyopherins  $\beta$ , ten of which are involved in transport to the nucleus (importins - Kap95, Kap104, Sxm1/Kap108, Mtr10, Kap114, Nmd5, Kap120/Lph2, Pse1/Kap121, Kap122, Kap123), three are involved in transport to the cytoplasm (exportins - Cse1, Crm1/Xpo1, Los1), and one karyopherin is involved in transport in both directions, Msn5 (22).

Despite the association described previously of Rrp6 with Srp1 and Kap95 (23-26), there are no conclusive studies describing the nuclear import pathway of yeast Rrp6. In this work, we purified the *S. cerevisiae* nuclear exosome with Rrp6-TAP for the identification of proteins interacting with this complex. One of the major groups of proteins co-purified with the exosome was that of the karyopherins/importins. The presence of different karyopherins associated with

Rrp6/exosome in our purifications raised the possibility of multiple pathways for nuclear import of Rrp6. Here we show the participation of different karyopherins in the transport of the exosome subunit Rrp6 to the nucleus and the sequences that these karyopherins might recognize in Rrp6. The results shown here provide evidence for alternative pathways of Rrp6 transport to the cell nucleus.

## RESULTS

### Purification of the nuclear exosome with Rrp6-TAP

To identify proteins interacting with the yeast nuclear exosome (Exo11), we took advantage of the TAP-tag method (27) using an Rrp6-TAP fusion. Since Rrp6 expression in yeast cells is lower than that of the core subunits (16,17; Fig. 1), twenty liters of culture were used here for the purification of the exosome with Rrp6-TAP. Despite the difference in expression levels of the bait proteins, the bands profile of the proteins co-immunoprecipitated with Rrp6-TAP was similar to that of the exosome co-immunoprecipitated with Rrp43-TAP (16), and all the exosome subunits were identified in the elution fraction (Fig. 1A). Samples from the same Rrp6-TAP elution fraction were analyzed by western blot with antibody against the CBP portion of the TAP-tag, confirming that Rrp6-TAP bound efficiently to the IgG-sepharose resin and was eluted after the cleavage reaction with TEV protease (Fig. 1B).

For the identification of the proteins co-immunoprecipitated with Rrp6-TAP, the eluted fraction was analyzed by mass spectrometry (Suppl. Table S1). All the exosome subunits were identified, those that are part of the RNA binding "cap", Rrp4, Rrp40 and Csl4, the subunits that form the RNase PH ring, Rrp41, Rrp45, Rrp46, Rrp43, Mtr3 and Rrp42, as well as the catalytic subunits, Rrp44 and Rrp6 (4,28-30). Based on the exosome structure previously described, Rrp6 and Rrp44 do not interact directly, but rather bind to opposite sides of the exosome core (4,28). The presence of Rrp44 in all the purifications, therefore, indicates that the exosome obtained here was stable during the co-immunoprecipitation with Rrp6-TAP.

To confirm that the exosome subunits eluted from the column were assembled in the Exo11 complex, proteins co-purified with Rrp6 were subjected to gel filtration for the separation of

complexes from free subunits. The collected fractions were subsequently analyzed by SDS-PAGE, showing that the main protein bands were concentrated between fractions 31 and 39 (Fig. 2B). Protein identification of the fractions by mass spectrometry showed that the exosome was eluted mainly in fractions 29 through 32 (Suppl. Table S2), corresponding to complexes of 440 to 660 kDa. These results confirm that the exosome complex was stable under the conditions used here (Fig. 2C; Suppl. Fig. S1).

Many fractions in which the exosome was identified corresponded to complexes with masses larger than expected (478 kDa), indicating that the exosome could be associated with additional proteins. Accordingly, some of the previously characterized exosome cofactors that had been shown to bind Rrp6, such as Mpp6 and Mtr4 (26), were identified in the same fractions in which the exosome was concentrated (Fig. 2). Additionally, other proteins were copurified with the exosome, many of them, chaperones (Suppl. Table S2), which may interact with the exosome to facilitate the assembly of this large protein complex (31). Interestingly, karyopherins were also identified among the proteins copurified with the exosome (Fig. 2, Suppl. Table S2).

### Karyopherins copurify with Rrp6-TAP

In nine independent experiments of Rrp6-TAP co-immunoprecipitations, we detected a total of 298 proteins associated with Rrp6 (Suppl. Table S1). Gene-ontology (GO) analysis of the proteins co-precipitated with Rrp6-TAP showed that the term “Importins/Karyopherins” was significantly enriched (Fig. 2D). The Karyopherins identified copurifying with Rrp6-TAP were Kap95, Srp1, Kap114, Kap123, Sxm1, and Cse1 (Suppl. Table S1). The interaction of Srp1 and Kap95 with Rrp6 had previously been observed in protein purification experiments (23-26). Interestingly, however, the interaction of the exosome with the other karyopherins has not been previously reported. Kap114 has been shown to be responsible for nuclear transport of TBP (TATA binding protein), the transcription factor TFIIB, histones H2A and H2B, and the cofactor associated with histone, NAP1 (32-35). Kap123 is involved in nuclear transport of ribosomal proteins before their association with the ribosome subunits, and also in

transport of other proteins associated with ribosomes, of histones H2A, H2B, H3 and H4, the histone acetyltransferase complex, SRP protein (signal recognition particle), and the endonuclease HO (32,33,36-41). Sxm1 is involved in nuclear transport of Lhp1 chaperone (La protein in humans), and complements the absence of Kap123 (42). Finally, Cse1 is an exportin responsible for transporting SRP1 to the cytoplasm (43).

Since various nuclear import pathways can be responsible for the transport of proteins to the nucleus, including the recognition of the cargo nuclear localization signals by many karyopherins (32,44,45), the identification of different karyopherins complexed with the exosome could suggest multiple nuclear import pathways for the exosome (46).

### Depletion of various karyopherins affect Rrp6 localization

Earlier studies showed the presence of a classical NLS at the C-terminal region of Rrp6 that could be recognized by importin  $\alpha$  Srp1 (47), suggesting that the transport of Rrp6 to the nucleus would be performed by the heterodimer Kap95/Srp1. In order to confirm this hypothesis, we first tested the effect of inhibition of Srp1 expression on Rrp6 subcellular localization.

A conditional mutant of *SRP1* ( $\Delta$ *srp1/GAL1::A-SRP1*) was transformed with plasmids coding for either GFP or GFP-Rrp6 for analysis of Rrp6 localization upon inhibition of expression of Srp1 for 14 hours in glucose. The results show that GFP-Rrp6 was localized to the nucleus when cells were incubated in galactose medium, but after incubation for 14 hours in glucose medium, a weak protein signal was visualized in the cytoplasm, although GFP-Rrp6 remained concentrated in the nucleus, most probably due to the low levels of Srp1 still present in the cells (Fig. 3A). Control experiment shows GFP in the cytoplasm of these cells (Suppl. Fig. S2A). Control western blot showed that A-Srp1 expression was inhibited after 12 hours in glucose, but its levels decreased significantly only after 15 hours in glucose (Fig. 4A). Interestingly, the results also show that the levels of Rrp6 decreased upon inhibition of A-Srp1 expression, whereas the levels of GFP alone did not change (Fig. 4A, left panel). These results confirm the involvement of Srp1 in the Rrp6 nuclear import, and could suggest that

when not efficiently transported to the nucleus, Rrp6 may be destabilized.

Kap95 is the yeast ortholog of human importin  $\beta$ , and has been shown to be important for the transport of some transcription factors (48). To analyze the involvement of Kap95 in the nuclear import of Rrp6, a conditional mutant expressing A-Kap95 under control of the *GAL1* promoter ( $\Delta kap95/GAL1::A-KAP95$ ) was transformed with plasmids expressing either GFP or GFP-Rrp6. Inhibition of Kap95 expression strongly affected GFP-Rrp6 localization, although this latter protein remained concentrated in the nucleus (Fig. 3B). It is noteworthy that some of the  $\Delta kap95/GAL1::A-KAP95$  cells showed an elongated form when grown in glucose medium (Suppl. Fig. S2B), probably due to the role of Kap95 in the transport of proteins involved in cell cycle regulation (48). The analysis of GFP-Rrp6 upon inhibition of Kap95 expression shows that the levels of Rrp6 also decreased (Fig. 4B), as observed after lowering Srp1 levels. The molecular masses of GFP-Rrp6 and ProtA-Kap95 are similar, and therefore, to better visualize GFP-Rrp6 band, the gels were also run longer to separate the bands (Fig. 4B, right panel).

The results showing that lower levels of Kap95 resulted in a stronger mislocalization of GFP-Rrp6 than lower levels of Srp1 suggest that the complex Srp1/Kap95 is not the only responsible for Rrp6 transport, but rather, Kap95 may be involved in the import of Rrp6 to the nucleus, either on its own, or associated with other adaptor proteins in addition to Srp1, as shown for other proteins (45).

To test the hypothesis that more than one transport pathway might be involved in the nuclear import of Rrp6, we investigated the subcellular localization of Rrp6 in mutants of other karyopherins found to co-purify with Rrp6-TAP: Kap114, Kap123, and Sxm1. Upon testing deletion of *KAP114* or *KAP123*, however, no effect on Rrp6 localization in the cell was detected (data not shown). These results are interesting because although the karyopherins Kap95, Srp1, Kap114 and Kap123 were co-immunoprecipitated in the same fraction as the exosome in the gel filtration chromatography, the individual depletion of Kap95 and Srp1 affected partially Rrp6 localization, whereas deletion of Kap114 and Kap123 did not have any effect, which might suggest that these latter karyopherins interact with other proteins co-purifying with Rrp6-TAP.

Strengthening our hypothesis of alternative pathways for Rrp6 nuclear import, deletion of the  $\beta$ -karyopherin Sxm1/Kap108 gene affected partially the localization of Rrp6. Sxm1 is not essential for growth and despite not being described as temperature sensitive (42), the localization of GFP-Rrp6 was tested in the deletion strain  $\Delta sxm1$  under different temperatures, 25°C and 37°C. The results show that in this strain, although still concentrated in the nucleus, Rrp6 was detected in the cytoplasm, and its mislocalization was stronger at 37°C (Fig. 3C). Interestingly, when the cells were shifted to 37°C, the levels of GFP-Rrp6 decreased, what could be visualized by western blot (Fig. 4C). Because GFP-Rrp6 expression was lower in this strain growing at 37°C, gain of the confocal microscope had to be increased in order to Rrp6 signal be detected. Importantly, this signal was above background levels. Control experiment shows that GFP-Rrp6 remains nuclear in WT cells incubated at 37°C (Suppl. Fig. S2C). The results of Srp1, Kap95 and Sxm1, an  $\alpha$ - and two  $\beta$ -karyopherins, affecting the nuclear localization of Rrp6 further suggest multiple transport pathways for this protein. Quantification of karyopherin mutant cells clearly shows the stronger effect of the deletion of *SXM1* on GFP-Rrp6 localization (Fig. 4D).

To determine whether Rrp6 could directly interact with these karyopherins in the absence of other yeast factors, protein pull-down experiments were performed using recombinant epitope-tagged proteins expressed in *E. coli*. GST or GST-Rrp6 were immobilized in glutathione-sepharose beads and incubated with extracts of cells expressing either His-Srp1 or His-Kap95. After extensive washing, proteins were eluted and analyzed by western blot with antibodies against the tags. The results show that GST-Rrp6 interacts directly with His-Srp1 and with His-Kap95 (Fig. 4E). Rrp6 interacts more strongly with Srp1 than with Kap95, as deduced by the relative amounts of proteins recovered in the elution fractions. Importantly, however, the results shown here confirm that Kap95 can interact with Rrp6 independently of Srp1, strongly indicating alternative pathways for Rrp6 nuclear import.

Because the effect of Sxm1 on Rrp6 localization was stronger than that of Kap95, we also attempted to test Rrp6-Sxm1 interaction. However, despite obtaining satisfactory His-Sxm1 expression levels in *E. coli*, this protein was mainly

present in inclusion bodies, and the little soluble protein was very labile (data not shown), making the performance of pull-down experiments unviable.

We next analyzed the participation of additional karyopherins in Rrp6 nuclear import. Mutants of karyopherins were therefore transformed with the plasmid coding for the GFP-Rrp6 fusion. Deletion of Kap120 had very small effect on Rrp6 nuclear accumulation when the cells were incubated at 25°C (Suppl. Fig. S3). Deletion of *KAP120* did not affect significantly GFP-Rrp6 levels, but when the cells were incubated at 37°C, GFP-Rrp6 was visualized in an area apparently larger than the nucleus, which could correspond to a perinuclear localization of this protein. This phenotype may be due to a defect in pre-60S maturation caused by the absence of Rpf1 in the nucleus, which has been shown to be transported by Kap120 (49). Additionally, deletion of *KAP120* has been shown to cause accumulation of 60S in the nucleus, with stronger defects when cells were incubated at 37°C (50).

Deletion of the gene of karyopherin Msn5 resulted in partial mislocalization of Rrp6 (Suppl. Fig. 3B). When analyzing GFP-Rrp6 expression in this strain, however, full-length GFP-Rrp6 band was not visualized on western blots (Suppl. Fig. 3D), even though the GFP-Rrp6 signal was visible by fluorescence microscopy. The GFP-Rrp6 degradation product visualized on western blots cannot correspond to GFP alone because of its mass and its concentration in the nucleus, as visualized by fluorescence microscopy (Suppl. Fig. 3B). Full-length GFP-Rrp6 might be present in these cells below the levels of detection by western blot, but sufficient for the detection by fluorescence microscopy. These results of GFP-Rrp6 localization in *Δmsn5* cells show that despite being concentrated in the nucleus, Rrp6 is not stable, suggesting that nuclear accumulation is not the only important factor for maintenance of Rrp6 levels.

Kap104 (homolog of human karyopherin  $\beta$ 2/Transportin) has been shown to mediate nuclear import of Nab2 (a nuclear polyadenylated RNA-binding protein), Hrp1 (subunit of cleavage complex required for maturation of pre-mRNA 3' ends), and the transcription factor Tfg2 (51-54). Kap104 recognizes a multipartite proline-tyrosine nuclear localization signals (PY-NLS) present in its cargos. Multipartite PY-NLS share a common C-terminal (R/H/K)-X(2-5)-P-Y motif within a

positively charged region of approximately 30 amino acids. The central region can contain a basic enriched motif or a hydrophobic motif (55). Interestingly, Rrp6 contains a similar PY-NLS motif in its sequence (Fig. 6A – NLS2). To test the involvement of Kap104 in the transport of Rrp6, *Δkap104* strain was transformed with plasmids coding for GFP, GFP-Rrp6, or GFP-Nab2, and the subcellular localization of these proteins was determined by fluorescence microscopy. Absence of Kap104 in cells growing at 37°C caused some mislocalization of Rrp6, but it remained concentrated in the nucleus (Fig. 5A). Interestingly, from a total of 119 cells observed, 16% showed granules close to the nucleus as well (data not shown). As expected, absence of Kap104 affected very strongly the transport of Nab2, used here as a control (Fig. 5A). Interestingly, upon assessing the levels of expression of GFP-Rrp6 and GFP-Nab2 in *Δkap104* cells, western blots results show that although GFP-Nab2 levels do not vary when incubating cells at different temperatures, levels of GFP-Rrp6 decreased dramatically at 37°C (Fig. 5B).

Combined, the results shown here confirm the hypothesis that Rrp6 can associate with different  $\beta$ -importins for its transport to the nucleus. Accordingly, further analysis of Rrp6 primary sequence using NLS prediction softwares revealed a third putative nuclear localization signal in its N-terminal region (Fig. 6A – NLS1), in addition to the classical NLS in the C-terminal portion of the protein, and the PY-NLS pointed out above.

### Deletion mutants of Rrp6 show different subcellular localization profiles

To determine whether the putative NLS at the N-terminal portion of Rrp6 could be involved in its nuclear import, we next constructed GFP-fused Rrp6 deletion mutants lacking one or more putative NLS (Fig. 6). To analyze the expression levels of the mutants in *Δrrp6* strain, the proteins were first visualized by western blot. Although the expression levels of the mutants vary, bands corresponding to the GFP-fused proteins were visualized for all of them, with the exception of Rrp6<sup>532-733</sup> (Fig. 6B). Levels of mutants Rrp6<sup>1-106</sup>, Rrp6<sup>1-398</sup>, and Rrp6<sup>153-398</sup> were similar to that of full-length Rrp6, whereas mutants Rrp6<sup>1-186</sup>, Rrp6<sup>1-619</sup>, Rrp6<sup>1-693</sup>, Rrp6<sup>153-186</sup>, Rrp6<sup>620-733</sup>, and Rrp6<sup>532-619</sup> showed higher levels than the full-length protein. To determine whether

any of the deletion mutants could complement growth of the temperature-sensitive *Arrp6* strain (56), *Arrp6* cells expressing the deletion mutants were incubated at either 30°C or 37°C. Interestingly, mutants Rrp6<sub>1-619</sub> and Rrp6<sub>1-693</sub>, which lack the C-terminal region of Rrp6, and therefore the classical NLS (NLS3), did complement growth at 37°C (Fig. 6C). Remarkably, mutant Rrp6<sub>532-619</sub>, containing only the exosome interacting domain, partially complemented growth of *Arrp6* at 37°C (Fig. 6C). These findings suggest that the C-terminal NLS is not essential for Rrp6 function, and, consequently, for its subcellular localization. Surprisingly, mutants Rrp6<sub>1-106</sub>, Rrp6<sub>1-186</sub>, and Rrp6<sub>1-398</sub>, bearing the Rrp47 interacting region, and mutants Rrp6<sub>153-398</sub> and Rrp6<sub>532-733</sub> negatively affected growth of *Arrp6* (Fig. 6C and data not shown). Rrp6<sub>153-398</sub> might interact with exosome subunits or exosome cofactors, sequestering these proteins and causing the negative effects. As indicated above, Rrp6<sub>532-733</sub> could not be detected by western blot (Fig. 6B).

The analysis of the subcellular localization of the GFP-Rrp6 deletion mutants in strain *Arrp6* by fluorescence microscopy showed that the full length GFP-Rrp6 was concentrated in the cell nucleus (Fig. 7), as expected, confirming that the GFP-fused protein was functional, as also seen by growth complementation at 37°C (Fig. 6C). To test the role of the classical NLS in Rrp6 localization, mutants Rrp6<sub>1-106</sub>, Rrp6<sub>1-186</sub>, Rrp6<sub>1-398</sub>, Rrp6<sub>1-619</sub>, Rrp6<sub>1-693</sub>, which lack NLS3, were analyzed. Rrp6<sub>1-106</sub>, Rrp6<sub>1-186</sub>, Rrp6<sub>1-398</sub>, Rrp6<sub>1-619</sub>, were concentrated in the nucleus, although significant amounts of the proteins were present in the cytoplasm. Rrp6<sub>1-693</sub>, on the other hand, was present throughout the cells (Fig. 7). This observation suggests that the putative NLS elements found in the N-terminal portion of Rrp6 could be recognized by karyopherins, allowing the transport of the mutants to the nucleus (Fig. 7). Alternatively, these mutants could interact with proteins containing NLS and be transported to the nucleus as subcomplexes. This might be the case especially for mutant Rrp6<sub>1-106</sub>, which contains only the three first amino acid residues of putative NLS1 (residues 104-132). Because these five mutants contain the Rrp47-interacting domain, these results also suggest that Rrp47 is not sufficient for the nuclear retention of Rrp6.

Mutants Rrp6<sub>153-398</sub> and Rrp6<sub>153-186</sub>, in spite of containing NLS2 at the N-terminus of the

exoribonuclease domain, were present throughout the cells (Fig. 7). The signal difference between these mutants are due to their different levels of expression. These results indicate that conformation of the mutant proteins, as well as protein interactions play a significant role on NLS recognition. Mutant Rrp6<sub>620-733</sub>, containing only the C-terminal portion of Rrp6 encompassing the classical NLS, was transported to the nucleus, albeit with very low efficiency, since the protein was mainly visualized in the cytoplasm (Fig. 7). Strikingly, Rrp6<sub>532-619</sub>, a mutant that does not contain any NLS, but only the Rrp6 portion involved in interaction with the exosome core, although present in the cytoplasm, concentrated in the nucleus (Fig. 7), further confirming that alternative pathways might be used for the transport of Rrp6 through the nuclear pore. These results also suggest that Rrp6 might interact with other proteins containing an NLS sequence and be imported to the nucleus as part of a complex. Based on these results, the NLS in the C-terminal portion of Rrp6 is undoubtedly not the only region of the protein responsible for its nuclear localization. Additionally, there seems to be a correlation between Rrp6 function and cell form. In the case of Rrp6 mutants that did not complement growth and did not localize to the nucleus, the cells seemed more elongated than normal, suggesting some impairment of cell division, as has been described for *Drosophila* (57). Confirming our observations, changes in Rrp6 expression have been shown to cause filamentous growth (58). *Arrp6* expressing mutants Rrp6<sub>1-106</sub>, Rrp6<sub>1-186</sub>, Rrp6<sub>1-398</sub>, Rrp6<sub>153-186</sub>, and Rrp6<sub>532-733</sub> showed the same elongated cell phenotype as *Arrp6* strain, which correlates with the lack of growth complementation at 37°C (Fig. 6C and data not shown). Mutants Rrp6<sub>1-619</sub>, Rrp6<sub>1-693</sub>, and Rrp6<sub>532-619</sub>, on the other hand, did not show the elongated phenotype, were expressed at high levels, complemented growth, and localized to the nucleus, despite lacking the canonical NLS. The results shown here strongly indicate that alternative import pathways are responsible for Rrp6 nuclear localization.

### Sxm1 is involved in Rrp6 nuclear import

Since Rrp6 mutants lacking its canonical C-terminal NLS were transported to the nucleus and deletion of *SXM1* affected Rrp6 localization (Figs. 7 and 3C, respectively), we tested the localization of

mutants Rrp6<sub>1-106</sub> and Rrp6<sub>532-619</sub> in  $\Delta$ *sxm1* strain. Rrp6<sub>1-106</sub>, containing Rrp47 interaction domain, was concentrated in the nucleus in  $\Delta$ *rrp6*, but not in  $\Delta$ *sxm1* (Fig. 8A). Rrp6<sub>532-619</sub>, containing only the exosome core-interacting domain, localized throughout the cell but was more concentrated in the nucleus in  $\Delta$ *sxm1* cells (Fig. 8A). These results strongly suggest that Sxm1 participates in Rrp6 nuclear import, either by recognizing the Rrp6 noncanonical NLS, or the NLS of a protein complexed with Rrp6. Importantly, as pointed out above, full-length Rrp6 is not stable in  $\Delta$ *sxm1* cells at 37°C, whereas the mutants Rrp6<sub>1-106</sub> and Rrp6<sub>532-619</sub> were expressed at high levels in  $\Delta$ *sxm1* cells (Fig. 8B). As in  $\Delta$ *rrp6* cells, Rrp6<sub>532-733</sub> was not detected by western blot in  $\Delta$ *sxm1* (Fig. 8B).

### Rrp47 is not the only factor affecting Rrp6 nuclear retention

Rrp6 has been shown to interact with Rrp47 through its N-terminal domain (59). To determine whether Rrp47 could influence Rrp6 mutants localization, full-length Rrp6 and deletion mutants containing Rrp47-interaction region, Rrp6<sub>1-106</sub>, Rrp6<sub>1-398</sub>, Rrp6<sub>1-619</sub>, and Rrp6<sub>1-693</sub>, were transformed into  $\Delta$ *rrp47* cells for analysis. The results show that localization of full-length GFP-Rrp6 was not affected by the absence of Rrp47, remaining exclusively nuclear (Fig. 9). Rrp6 deletion mutants, on the other hand, were mainly visualized in the cytoplasm (Fig. 9). These results are very interesting because although mutants Rrp6<sub>1-106</sub> and Rrp6<sub>1-398</sub> do not contain the canonical C-terminal NLS, they were present in the nucleus in strain  $\Delta$ *rrp6*, but remarkably, significantly lower amounts of these proteins were visible in the nucleus in  $\Delta$ *rrp47* cells (Figs. 7 and 9A). Mutants Rrp6<sub>1-619</sub> and Rrp6<sub>1-693</sub> showed similar localization in  $\Delta$ *rrp6* and  $\Delta$ *rrp47* cells (Figs. 7 and 9).

Similar to what was seen in  $\Delta$ *rrp6* strain, Rrp6 mutants showed different levels of expression in  $\Delta$ *rrp47* cells. Most of the mutants were expressed at higher levels than full-length Rrp6 (Fig. 9B). Interestingly, overexpression of Rrp6 and the mutants Rrp6<sub>1-619</sub> and Rrp6<sub>1-693</sub> complemented growth of  $\Delta$ *rrp47* cells at 37°C (Fig. 9C). Combined, these results suggest that Rrp47 plays a role, but is not the only factor regulating the nuclear retention of Rrp6.

## DISCUSSION

Previous attempts to purify the exosome from yeast cells with tagged core subunits resulted in very little recovery of Rrp6 (16). By using Rrp6 as bait, however, Exo11 complex was purified in a stable form that could be separated by gel filtration. In addition, exosome cofactors and karyopherins were identified that were not present in the complex isolated with TAP-Rrp43 (16). These results show that different proteins remain associated with the exosome depending on the bait used for the purification, probably because the core exosome subunits are present both in nucleus and cytoplasm, whereas Rrp6 is exclusively nuclear.

Among the karyopherins identified with Exo11 were the yeast  $\alpha$ -importin Srp1/Kap60 and the  $\beta$ -importins Kap95, Sxm1, Kap114, and Kap123. As shown here, lower levels of Srp1 or of Kap95 affect partially the localization of Rrp6, because in spite of being concentrated in the nucleus, Rrp6 can also be visualized in the cytoplasm upon inhibition of expression of these karyopherins. Kap114 and Kap123, on the other hand, do not affect Rrp6 nuclear localization. Copurification of Kap114 and Kap123 with Rrp6-TAP may therefore be due to the interaction of these karyopherins with other proteins co-purifying with Rrp6.

It has been proposed that the complex Srp1/Kap95 is responsible for the nuclear import of Rrp6 based on pull-down of Srp1 with ProtA-Rrp6 (59), however, as shown here, despite the copurification of these proteins from yeast cells, inhibition of expression of Srp1 or Kap95 has different effects on the localization of Rrp6. Srp1 possibly recognizes the classical NLS sequence at Rrp6 C-terminus, but in its absence, alternative NLS sequences in the N-terminal portion of Rrp6 may be recognized by other karyopherins, since Rrp6 partially localizes to the nucleus in  $\Delta$ *srp1*/*GAL::SRP1* and  $\Delta$ *kap95*/*GAL::KAP95* strains growing in glucose medium. Importantly, Rrp6 mutants lacking the canonical C-terminal NLS, Rrp6<sub>1-619</sub> and Rrp6<sub>1-693</sub> are not concentrated, but are transported to the nucleus and complement growth of  $\Delta$ *rrp6* cells at the nonpermissive temperature. Overlapping and redundant import pathways have been reported for other proteins and may also occur in the case of Rrp6 (60).

Confirming this hypothesis, by performing protein pull-down experiments, we show here that not only Rrp6 interacts directly with Srp1, but also with Kap95 in the absence of any other yeast proteins. To the best of our knowledge, this is the first time such Rrp6 direct interactions are shown. Importantly, by binding directly to the  $\beta$ -karyopherin Kap95, Rrp6 may be transported to the nucleus independently of the  $\alpha$ -karyopherin Srp1.

Upon testing other karyopherins, we identified the involvement of Sxm1 in the transport of Rrp6. The deletion *Sxm1* gene strongly affects Rrp6 localization in the cells, suggesting that alternative pathways might be used for the transport of Rrp6 to the nucleus. Confirming that hypothesis, Rrp6<sub>1-106</sub> mislocalized in  $\Delta$ *sxm1* cells, a very different phenotype from that observed in  $\Delta$ *rrp6* strain. Sxm1 might recognize alternative NLS in Rrp6 sequence, or in proteins interacting with Rrp6. This hypothesis is supported by the observation that Rrp6 mutants lacking the canonical NLS still localize to the nucleus, but not in the absence of Sxm1. In all karyopherin mutants tested here, despite Rrp6 presence in the cytoplasm in some cases, the full-length protein was concentrated in the nucleus. Srp1, Kap95, and Sxm1 were the karyopherins that affected Rrp6 more strongly. Deletion of Msn5 and Kap104 affected mildly Rrp6 localization, whereas Kap120, Kap114, Kap122 and Kap123 had little or no effect (Fig. 10). These results strongly support the idea of overlapping mechanisms for Rrp6 nuclear import.

Rrp6 structure has been determined in the context of the exosome complex (4,28,61), but the cell compartment in which its association with the exosome occurs has not been described. Considering the structure of Rrp6 when bound to the exosome core, its N-terminal portion is free to interact with Rrp47 (61), and possibly, with karyopherins that might recognize the putative N-terminal NLS (Fig. 6A). The classical NLS is positioned in the C-terminal portion of Rrp6, also exposed in the exosome structure (61). Alternatively, these NLS could be recognized in Rrp6 molecules not complexed with the exosome core.

One possible nuclear import pathway for Rrp6 is through the recognition of its classical C-terminal NLS by Srp1/Kap95 heterodimer. However, as shown here, another pathway involves the recognition of additional N-terminal NLS in

Rrp6. We have identified one additional  $\beta$ -karyopherin involved in Rrp6 nuclear import, Sxm1. In the absence of this protein, transport of full-length Rrp6 is less efficient, whereas mutant Rrp6<sub>1-106</sub> is no longer transported to the nucleus. Because of Rrp6 interactions with other exosome subunits, and with exosome co-factors, it may also be transported in the form of protein subcomplexes.

As shown here, inhibition of Kap95 expression leads to the appearance of abnormal elongated cells in the culture, similar to  $\Delta$ *rrp6* strain, or  $\Delta$ *rrp6* expressing deletion mutants of Rrp6. Rrp6 has been shown to be involved in processing of histone mRNAs in yeast, indirectly affecting cell cycle regulation (62,63). Taken together, these results suggest that there may be a cell division impairment in the absence of functional Rrp6 in the cells, or when Rrp6 is mislocalized.

Another important observation described here was that Rrp6 levels decreased in most of the karyopherin mutants, suggesting that when not efficiently transported to the nucleus, Rrp6 may be directed for degradation. Incubation of the deletion strains  $\Delta$ *msn5* and  $\Delta$ *kap104* at 37°C led to very strong decrease in Rrp6 levels, so that it could no longer be detected by western blot. These results suggest that not only the subcellular localization of Rrp6 affects its levels, but additional factors are important as well. It is tempting to hypothesize that Msn5 may be involved in the transport of other proteins that are important for maintaining Rrp6 levels.

Based on the data shown here, on the direct interactions between Rrp6 and karyopherins, and on the sequences of the putative NLS sequences present in Rrp6, we propose a model according to which Rrp6 transport to the nucleus can be facilitated by the  $\alpha/\beta$  dimer Srp1/Kap95 recognizing the canonical NLS3 at the C-terminal portion of Rrp6, or by the  $\beta$  karyopherins Kap95 and Sxm1 recognizing one of the nuclear localization signals. In addition to these pathways, Rrp6 may be transported to the nucleus complexed with other exosome subunits which could be recognized by Sxm1 and other  $\beta$  karyopherin (Fig. 10E). Importantly, interaction of Rrp6 with the exosome core or with Rrp47 may be necessary for nuclear retention and maintenance of Rrp6 stability.

## EXPERIMENTAL PROCEDURES

### DNA manipulation and plasmid construction

Plasmids used in this study, described in Table 1, were constructed according to the cloning techniques described by Sambrook and Russell (2001) and sequenced by the Big Dye method (Perkin-Elmer). Cloning strategies are briefly described below.

Plasmids expressing *Srp1* and *Kap95* fused to Protein A were constructed by inserting the PCR-amplified open reading frames into YCplac33-GAL-A-RRP43 (67), which was previously digested to remove the *RRP43* coding sequence. *SRP1* fragment was cloned using *Bam*HI and *Sal*I restriction sites and *KAP95* fragment with *Bam*HI and *Pst*I restriction sites. In both constructs, the expression of the fusion proteins was regulated by the *GAL1* promoter.

Plasmids expressing the GFP fusions in yeast were constructed by inserting DNA fragments into pUG34 plasmid (64) using oligonucleotides with specific restriction sites. An RRP6 fragment was extracted from pGADC2-RRP6 (11) and inserted into pUG34 using *Eco*RI and *Sal*I restriction sites. Constructs expressing Rrp6 truncated mutants 1–106, 1–186, 1–398, 1–619, 1–693, 153–398, 153–186, 532–733, 620–733, and 532–619 (numbers denote amino acid positions in full-length Rrp6) were amplified by PCR using specific oligonucleotides and inserted into pUG34 with the same restriction enzymes. *NAB2* gene was inserted into pUG34 vector using *Spe*I and *Sal*I restriction enzymes. Expression of these GFP fusions was regulated by *MET25* promoter. MET-RFP-NOP1 fragment was extracted from pUG36-DsRed-NOP1 (65) and inserted into YCplac111 *Sac*I and *Hind*III restriction sites. To construct the plasmid YCplac111-GFP-RRP6, fragment MET25-GFP-RRP6 from pUG34-RRP6 was extracted after digestion with *Sac*I and *Sal*I restrictions enzymes and inserted into YCplac111 plasmid digested with the same enzymes.

### Yeast maintenance, transformation and sporulation

Yeast genetic techniques were conducted as described by Guthrie and Fink, 1991. Yeast strains (Table 2) were maintained in synthetic dropout (SD) medium complemented with the appropriate amino acids or nitrogenous bases mix, or synthetic

complete (SC) medium. Glucose or galactose was added as carbon source to a final concentration of 2% (w/v), as indicated. Yeast cells were transformed using the lithium acetate method (68).

### Tandem Affinity Purification

RRP6-TAP and empty-TAP purifications were performed as described previously (16). Briefly, 20 liters of yeast cells expressing Rrp6-TAP-tagged were grown to an OD<sub>600</sub> of 1.0–1.2 in synthetic complete medium containing glucose as a carbon source. Cells were harvested by centrifugation, resuspended in 50 mM Tris-HCl pH8.0, 150 mM NaCl, 10% glycerol, and 1 mM phenylmethylsulfonyl fluoride (PMSF), immediately frozen in liquid N<sub>2</sub>, and stored at -80°C. Cells were lysed by grinding using a ball mill device (Retsch, Mixer Mill MM 200 or Planetary Ball Mill PM 100) and centrifuged at 40,000 rpm for 1 h at 4°C in a rotor P50AT2-716 (Hitachi). The supernatant was incubated for 2 h at 4°C with IgG-Sepharose beads (GE Healthcare), followed by extensive washing with the same buffer. Proteins were eluted from beads by incubating the resin with 20 U of Tobacco Etch Virus protease (Invitrogen) for 16 h at 4°C, in the presence of 1mM DTT and 0.5 mM EDTA pH 8.0. Elution fraction from IgG-sepharose chromatography was subjected to gel filtration using a pre-packed glass column Superose™ 6 10/300 GL (GE Healthcare, Cat No 17-5172-01) connected to an AKTA-FPLC (GE Healthcare, Cat No 18-1900-26) at 0.5 ml/min flow rate.

### Protein digestion and mass spectrometric identification by LC-MS/MS

For identification of proteins obtained from TAP purification, elution fraction was resolved by SDS-PAGE, and bands were removed by cutting gels in slices, reduced, alkylated, and digested with trypsin (69). The digested samples were desalted using Sep-Pak C18 Plus Short Cartridge (WATERS) or EMD Millipore ZipTip™ Pipet Tips (MILLIPORE) according to manufacturer instructions. Tryptic peptides were resuspended in 20 µL formic acid (0.1%) and an aliquot (4.5 µL) was injected onto a QTOF Ultima mass spectrometer (Waters, Milford, USA) through a coupled nanoUPLC system (Acquity, Waters, Milford, USA). The peptide mixture was first desalted into a C18 trap column (180 µm i.d.×20 mm -Waters, Milford, USA) with

100% solvent A (0.1% formic acid) at 5  $\mu$ L/min for 3 min. Peptides were fractionated onto an analytical C18 column (75  $\mu$ m i.d.  $\times$  100 mm) (Waters) in a 20 min gradient (5–40% acetonitrile in 0.1% formic acid) at a flow rate of 600 nL/min. Spray voltage was set at 3.2 kV and the instrument was operated in data dependent mode in which one full MS scan was acquired in the  $m/z$  range of 200–2000 followed by MS/MS acquisition using collision induced dissociation of the three most intense ions from the MS scan. The phosphoric acid (0.05% in acetonitrile) was used as a lock mass and therefore was continuously sprayed into the ESI source and detected every 15 seconds. Alternatively, peptides originated from in-gel digestion were also analyzed by LC-MS/MS using a Q-ToF PREMIER mass spectrometer (Waters) as described (70). The raw data were processed and transformed into pkl format using ProteinLynx Global Server (Waters) after lock mass (at  $m/z$  784,823) correction. In solution digestion was also performed for proteins both directly eluted from TAP constructs (including empty vector) and from those eluted from TAP constructs followed by gel filtration chromatography (71). Glycerol was previously removed from protein mixtures through acetone precipitation. Peptide mixture originated from total TAP eluates were analyzed by LC-MS/MS using a Q-ToF PREMIER mass spectrometer (Waters) as described (16). Tryptic peptides from gel filtration chromatography eluates were analysed by LC-MS/MS using a LTQ-Velos Orbitrap (Thermo) as described (16). Proteins were identified by searching against a database sequence of *Saccharomyces cerevisiae* (S288c strain, downloaded at UNIPROT). Carbamidomethylation (C) was set as fixed modification while oxidation (M) as variable modifications. For Q-TOF and LTQ-Velos Orbitrap mass spectrometry, MS1 tolerance was set of 0.1–0.5 Da and 10 ppm, respectively, while, MS2 of 0.1–0.5 Da and 0.5 Da, respectively. High resolution data was also analyzed at Proteome Discover 1.4 (Thermo) where a FDR 1% was set. Proteins present in the empty-TAP purification (negative control) were excluded from the final list of Rrp6-TAP (interactors detected in all nine purifications). Functional annotation and GO enrichment analysis was performed by DAVID (72,73) with the parameter Ease=0.01.

### Protein pull-down and immunoblot analysis

In the pull-down assay, cellular extracts generated in 20mM Tris/150mM NaCl/1mM EDTA/0.8% Nonidet/1mM DTT of *E. coli* cells expressing either GST or GST-Rrp6 were incubated for 1 hour at 4°C with 250 $\mu$ L glutathione-sepharose beads (GE Healthcare) and the unbound material was washed. Beads were then incubated with cellular extract containing His-Srp1 or His-Kap95, flow through was collected, and beads were washed with the same buffer, followed by washing with buffer containing 250mM NaCl. Bound proteins were eluted with 50 mM Tris pH 8.0/ 10 mM reduced glutathione.

### Immunoblot Experiments

Protein samples were resolved by SDS-PAGE and transferred to polyvinylidene difluoride (PVDF) membranes (GE Healthcare). Membranes were incubated with primary antibodies against CBP, GFP (Sigma Aldrich), or Arp2 (Santa Cruz Biotech) in PBST/nonfat milk. Secondary antibodies used were anti-rabbit or anti-goat IgG conjugated to peroxidase (Sigma-Aldrich). Western blots were developed using Immobilon Western Chemiluminescent HRP Substrate (Millipore Corporation).

### Fluorescence microscopy

Cells were fixed in 70% methanol for 15 minutes, rinsed with cold phosphate-buffered saline (PBS), and then treated with 1 mg/mL RNase for 30 min. Nuclei were counterstained in a dye solution containing 3 mg/ml propidium iodide (PI) for 15 minutes. Cells were observed using a Nikon Eclipse Ti microscope equipped with filters for green fluorescence (GFP-3035B-000-ZERO, Semrock) and red fluorescence (Texas Red BrightLine set; TXRED4040-B, Semrock). The exposure times varied from 1 to 3 seconds. Images were processed and analyzed using the programs Nis Elements (version 3.07; Nikon) and ImageJ (<http://rsbweb.nih.gov/ij/>). Confocal images were captured in a 1024 $\times$ 1024 pixel format using a Zeiss LSM 780 confocal laser scanning inverted microscope (Carl Zeiss, Germany) at Centro de Facilidades para a Pesquisa (CEFAP-USP). Image stacks comprised 8 images captured with an alpha Plan-Apochromat 100x /1.46 Oil DIC M27 objective (Zeiss), applying a zoom factor of 1.5. Step intervals along the Z-axis ranged from 200 to

250 nm. Image processing was performed using the Zen 2011 software (Zeiss, version 11.00.190).

#### Identification of putative NLS elements

PSORT II Prediction software (<http://www.genscript.com/psort/psort2.html>), NLS Mapper (74) and NLStradamus (75) were used for the identification of putative NLS elements.

Examples of monopartite, bipartite and basic enriched NLS can be found in (19,52,53). NLS1: 104-NSKSRGSDLQYLGEFSGKNFSPTKRVEKP-132; NLS2: 153-KEKPNALKPLSESLRLVDDDENNP SHYPHY-183; NLS3: 697-RQQKRRFDPSSSDSNGPRAAKRRPA-723.

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#### Author Contributions

CCO conceived and coordinated the study. FAGZ designed, performed and analyzed the experiments. EKO contributed to the confocal analysis, protein pull-down experiments and preparation of the figures. FAGZ and JPCDC analyzed the mass spectrometry results. FAGZ, JPCDC and CCO wrote the manuscript. All authors reviewed the results and approved the final version of the manuscript.

#### Conflict of interest

The authors declare that they have no conflicts of interest with the contents of this article.

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## TABLES

**Table 1.** Plasmids used in this work.

Name	Description	Reference
pGADC - RRP6	AD::RRP6; LEU2; 2mm	11
pUG34	MET25::GFP, HIS3, CEN6	64
pUG36	URA3, CEN6	64
pUG34-ASR1	MET25::GFP-ASR1, HIS3, CEN6	This study
pUG34-H2A	MET25::GFP-H2A, HIS3, CEN6	This study
pUG34-GCN4	MET25::GFP-GCN4, HIS3, CEN6	This study
pUG34-NAB2	MET25::GFP-NAB2, HIS3, CEN6	This study
pUG34-RRP6	MET25::GFP-RRP6, HIS3, CEN6	This study
pUG34-rrp6 (1-106)	MET25::GFP-rrp6 1-106, HIS3, CEN6	This study
pUG34-rrp6 (1-398)	MET25::GFP-rrp6 1-398, HIS3, CEN6	This study
pUG34-rrp6 (1-619)	MET25::GFP-rrp6 1-619, HIS3, CEN6	This study
pUG34-rrp6 (1-693)	MET25::GFP-rrp6 1-693, HIS3, CEN6	This study
pUG34-rrp6 (153-398)	MET25::GFP-rrp6 153-398, HIS3, CEN6	This study
pUG34-rrp6 (532-733)	MET25::GFP-rrp6 532-733, HIS3, CEN6	This study
pUG34-rrp6 (620-733)	MET25::GFP-rrp6 620-733, HIS3, CEN6	This study
pUG34-rrp6 (532-619)	MET25::GFP-rrp6 532-619, HIS3, CEN6	This study
pUG36-DsRed-NOP1	MET25::DsRED-NOP1, URA3, CEN6	65
YCplac33	URA3; CEN4	66
YCplac111	LEU2; CEN4	66
YCplac33-GAL-A-RRP43	GAL1::ProtA-RRP43, URA3, CEN4	67
YCplac33-GAL-A-SRP1	GAL1::ProtA-SRP1, URA3, CEN4	This study
YCplac33-GAL-A-KAP95	GAL1::ProtA-KAP95, URA3, CEN4	This study
YCplac111-RFP-NOP1	MET25::DsRED-NOP1, LEU2, CEN4	This study
YCplac111-GFP-RRP6	MET25::GFP-RRP6, LEU2, CEN4	This study
pGEX-RRP6	<i>GST::RRP6</i> , Amp <sup>R</sup>	This study
pET28-KAP95	<i>His::KAP95</i> , Kan <sup>R</sup>	This study
pET29-SRP1	<i>His::SRP1</i> , Kan <sup>R</sup>	This study
pET28-SXM1	<i>His::SXM1</i> , Kan <sup>R</sup>	This study

**Table 2.** Strains used in this work.

Strain	Name	Genotype	Reference
FGY-25	KAP95/ $\Delta$ kap95	BY4743; Mat a/ $\alpha$ ; his3 $\Delta$ 1/his3 $\Delta$ 1; leu2 $\Delta$ 0/leu2 $\Delta$ 0; lys2 $\Delta$ 0/LYS2; MET15/met15 $\Delta$ 0; ura3 $\Delta$ 0/ura3 $\Delta$ 0; YLR347c::kanMX4/YLR347c	EUROSCARF
FGY-53	$\Delta$ kap95	YLR347c::kanMX4; his3 $\Delta$ 1; leu2 $\Delta$ 0; lys2 $\Delta$ 0; ura3 $\Delta$ 0; Ycplac33-GAL-A-KAP95	This study
FGY-97	KAP104/ $\Delta$ kap104	BY4743; Mat a/a; his3 $\Delta$ 1/his3 $\Delta$ 1; leu2 $\Delta$ 0/leu2 $\Delta$ 0; lys2 $\Delta$ 0/LYS2; MET15/met15 $\Delta$ 0; ura3 $\Delta$ 0/ura3 $\Delta$ 0; YBR017c::kanMX4/YBR017c	EUROSCARF
FGY-112	$\Delta$ kap104	CEN.ZI5-3B; CEN.PK; Mat a; ura3-52; his3 $\Delta$ 1; leu2-3_112; trp1-289; YBR017c::URA3	EUROSCARF
FGY-60	$\Delta$ kap114	BY4742; Mat a; his3 $\Delta$ 1; leu2 $\Delta$ 0; lys2 $\Delta$ 0; ura3 $\Delta$ 0; YGL241w::kanMX4	EUROSCARF
FGY-100	$\Delta$ kap120	BY4742; Mat alpha; his3 $\Delta$ 1; leu2 $\Delta$ 0; lys2 $\Delta$ 0; ura3 $\Delta$ 0; YPL125w::kanMX4	EUROSCARF
FGY-101	$\Delta$ kap122	BY4742; Mat a; his3 $\Delta$ 1; leu2 $\Delta$ 0; lys2 $\Delta$ 0; ura3 $\Delta$ 0; YGL016w::kanMX4	EUROSCARF
FGY-61	$\Delta$ kap123	BY4742; Mat a; his3 $\Delta$ 1; leu2 $\Delta$ 0; lys2 $\Delta$ 0; ura3 $\Delta$ 0; YER110C::kanMX4	EUROSCARF
FGY-105	$\Delta$ msn5	BY4742; Mat a; his3 $\Delta$ 1; leu2 $\Delta$ 0; lys2 $\Delta$ 0; ura3 $\Delta$ 0; YDR335w::kanMX4	EUROSCARF
FGY-87	$\Delta$ nmd5	CEN.RO22-4B; CEN.PK; Mat a; ura3-52; his3 $\Delta$ 1; leu2-3_112; trp1-289; YJR132w::HIS3	EUROSCARF
FGY-5	RRP6-TAP	MATa; ade2; arg4; leu2-3,112; trp1-289; ura3-52; YOR001w::TAP	EUROSCARF
FGY-88	$\Delta$ rrp6	BY4742; Mat a; his3 $\Delta$ 1; leu2 $\Delta$ 0; lys2 $\Delta$ 0; ura3 $\Delta$ 0; YOR001w::kanMX4	EUROSCARF
FGY-209	$\Delta$ rrp47	BY4742; Mat alpha; his3 $\Delta$ 1; leu2 $\Delta$ 0; lys2 $\Delta$ 0; ura3 $\Delta$ 0; YOR001w::kanMX4	EUROSCARF
FGY-26	SRP1/ $\Delta$ srp1	BY4743; Mat a/ $\alpha$ ; his3 $\Delta$ 1/his3 $\Delta$ 1; leu2 $\Delta$ 0/leu2 $\Delta$ 0; lys2 $\Delta$ 0/LYS2; MET15/met15 $\Delta$ 0; ura3 $\Delta$ 0/ura3 $\Delta$ 0; YNL189w::kanMX4/YNL189w	EUROSCARF
FGY-41	$\Delta$ srp1	YNL189w::kanMX4; his3 $\Delta$ 1; leu2 $\Delta$ 0; lys2 $\Delta$ 0; ura3 $\Delta$ 0; Ycplac33-GAL-A-SRP1	This study
FGY-86	$\Delta$ sxm1	BY4742; Mat a; his3 $\Delta$ 1; leu2 $\Delta$ 0; lys2 $\Delta$ 0; ura3 $\Delta$ 0; YDR395w::kanMX4	EUROSCARF

## FIGURES LEGENDS

**Figure 1.** Coimmunoprecipitation of the exosome with Rrp6-TAP. Total extract was incubated with IgG-sepharose beads, and proteins were eluted with TEV protease. Samples from total extract (input - In), flow through (FT), and elution (Elu) were subjected to SDS-PAGE and silver stained (**A**). The exosome subunits were identified by mass spectrometry. (**B**) Western blot of the same samples with anti-CBP. Rrp6-TAP bound efficiently to the resin and was eluted after cleavage with TEV protease.

**Figure 2.** Proteins co-purified with Rrp6-TAP were subjected to gel-filtration on superose 6 column for the separation of the exosome complex. (**A**) Chromatographic profile of the proteins. Arrows indicate the elution volumes of the molecular mass controls run through the same column. (**B**) SDS-PAGE of the fractions obtained. Fractions were also analyzed by mass spectrometry for the identification of the proteins. (**C**) Fractions 29-32 contained, in addition to exosome subunits, some exosome cofactors, karyopherins and chaperones, grouped in different colors. (**D**) Proteins copurifying with Rrp6-TAP were classified by Gene Ontology (GO) using DAVID software. Karyopherins was the second most predominant category of proteins.

**Figure 3.** Inhibition of karyopherins expression affects the subcellular localization of GFP-Rrp6. (**A**) Laser scanning confocal microscope images show the subcellular localization of GFP-Rrp6 after inhibition of Srp1 expression for 14h in glucose medium in *Δsrp1/GAL::SRP1* cells. Analysis of GFP-Rrp6 relative to PI by using ImageJ is shown on the right. Green lines represent GFP and red lines represent PI. Localization of GFP in *Δsrp1/GAL::SRP1* cells is shown in Suppl. Fig. S2. (**B**) Analysis of the subcellular localization of GFP-Rrp6 after inhibition of Kap95 expression for 14h in glucose medium in *Δkap95/GAL::KAP95* cells. Analysis of GFP-Rrp6 relative to PI by using ImageJ is shown on the right. (**C**) Analysis of the subcellular localization of GFP-Rrp6 in *Δsxm1* cells growing at 25°C or 37°C. Because GFP-Rrp6 expression is lower in this strain growing at 37°C, gain of the confocal microscope had to be increased in order for the Rrp6 signal to be detected. Importantly, this signal was above background levels.

**Figure 4.** Inhibition of karyopherins expression affects the levels of GFP-Rrp6. Western blot of total cell extract from karyopherins mutants expressing either GFP or GFP-Rrp6, growing in galactose- or glucose-containing media was performed with antibody against GFP, which also allowed the detection of ProtA-Srp1 and ProtA-Kap95. (**A**) *Δsrp1/GAL::A-SRP1* growing in glucose shows the lower levels of ProtA-Srp1 after 12h or 15h in glucose. GFP-Rrp6 also decrease upon inhibition of Srp1 expression. (**B**) Western blot of total cell extract from *Δkap95/GAL::A-KAP95* expressing either GFP or GFP-Rrp6 shows the lower levels of ProtA-Kap95 in glucose. GFP-Rrp6 and ProtA-Kap95 have very similar molecular masses. Longer run and exposure for separation and visualization of the bands is shown on the right-hand side. (**C**) Western blot of total cell extract from *Δsxm1* expressing either GFP or GFP-Rrp6, growing at 25°C or 37°C show that the expression levels of GFP-Rrp6 are lower at 37°C. Western blot with antibody against Arp2 was used as an internal control. (**D**) Quantitative analysis of cells expressing GFP-Rrp6. Approximately 100 of each strain was analyzed by fluorescence microscopy, cells were counted that showed protein localized to the nucleus (N), present both in nucleus and cytoplasm (N+C), or visible mainly in the cytoplasm (C). Numbers correspond to the percentage of cells showing each phenotype. (**E**) Pull-down of Srp1 and Kap95 with Rrp6. GST or GST-Rrp6 bound to glutathione-sepharose beads were incubated with His-Srp1, or His-Kap95-containing extracts. Elution fractions are shown. The same membrane was incubated with antibody against His-tag, and subsequently with antibody against GST-tag. Figure shown is representative of three independent experiments.

**Figure 5.** Deletion of *KAP104* does not affect localization of GFP-Rrp6 at 37°C. (**A**) Fluorescence microscopy for the analysis of the subcellular localization of GFP-Rrp6 in *Δkap104* cells. GFP-Nab2 was used as a control of a protein affected by the absence of Kap104. Analysis of GFP-Rrp6 and GFP-Nab2

relative to PI in *Δkap104* cells growing at 37°C by is shown on the right. **(B)** Western blot of total cell extract from *Δkap104* cells expressing GFP-Rrp6 or GFP-Nab2, incubated at 25°C or 37°C was performed with antibody against GFP. Expression levels of GFP-Rrp6 decreased at 37°C in *Δkap120* cells, but not those of GFP-Nab2. Antibody against Arp2 was used as an internal control.

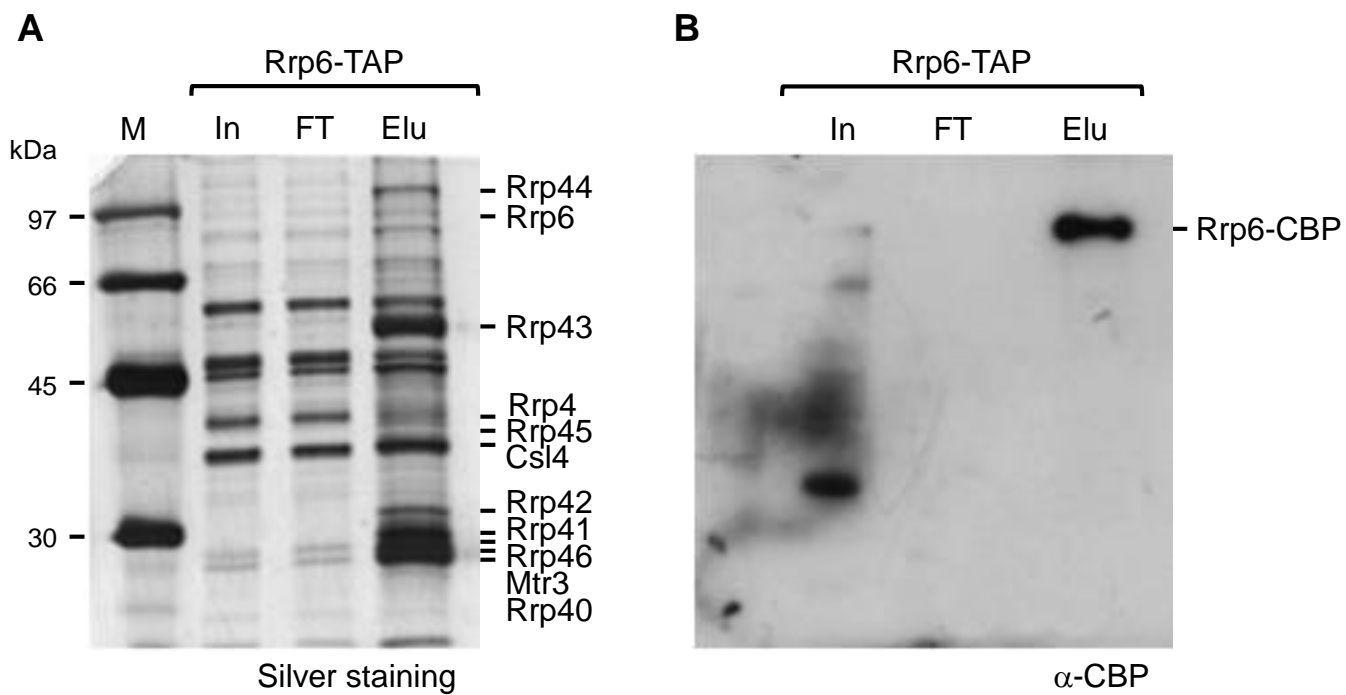
**Figure 6.** Expression of Rrp6 deletion mutants in *Δrrp6*. **(A)** Schematic representation of the deletion mutants of Rrp6. Rrp47 and core exosome interaction regions, as well as active domains of Rrp6 are highlighted. Positions of putative NLS are in shown blue. **(B)** Western blot for the determination of the expression of the GFP-Rrp6 mutants in *Δrrp6* strain. Arp2 was used as an internal control. **(C)** Analysis of growth of *Δrrp6* cells expressing either GFP or the different GFP-Rrp6 constructs at 30°C or 37°C. Mutants lacking canonical NLS, Rrp6<sub>1-619</sub> and Rrp6<sub>1-693</sub>, complement growth of *Δrrp6* at 37°C.

**Figure 7.** Presence of canonical NLS is not the only determinant of Rrp6 nuclear import. Laser scanning confocal microscope images show the subcellular localization of the Rrp6 deletion mutants expressed in *Δrrp6* cells growing at 25°C. **(A)** GFP-Rrp6, Rrp6<sub>1-106</sub>, Rrp6<sub>1-186</sub>, Rrp6<sub>1-398</sub>, Rrp6<sub>1-619</sub>, Rrp6<sub>1-693</sub>. **(B)** Rrp6<sub>153-398</sub>, Rrp6<sub>153-186</sub>, Rrp6<sub>620-733</sub>, Rrp6<sub>532-619</sub>, GFP. Various mutants accumulate in the nucleus, although to different levels. Quantification is shown on the right. **(C)** Quantitative analysis of cells expressing GFP-fused Rrp6 mutants. (N), protein concentrated in the nucleus; (N+C) protein present both in nucleus and cytoplasm; (C), protein visible mainly in the cytoplasm. Numbers correspond to the percentage of cells showing each phenotype.

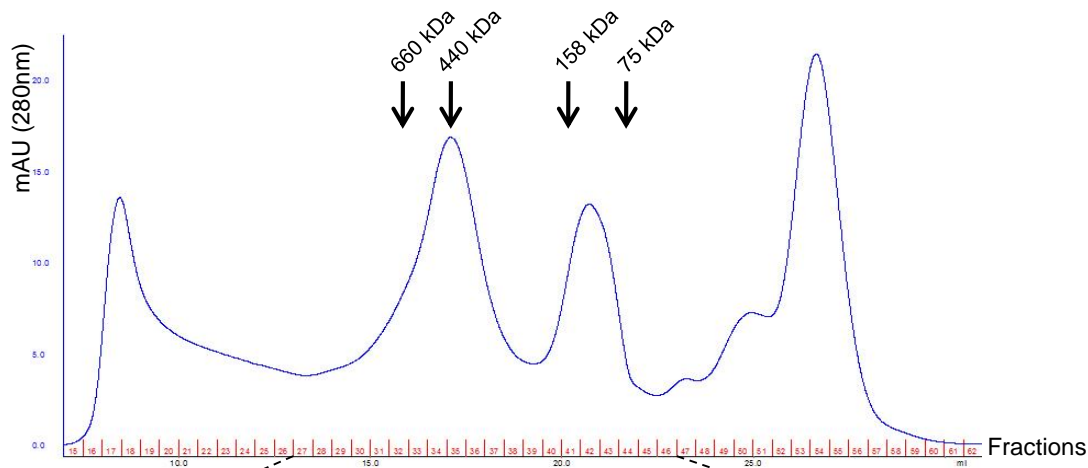
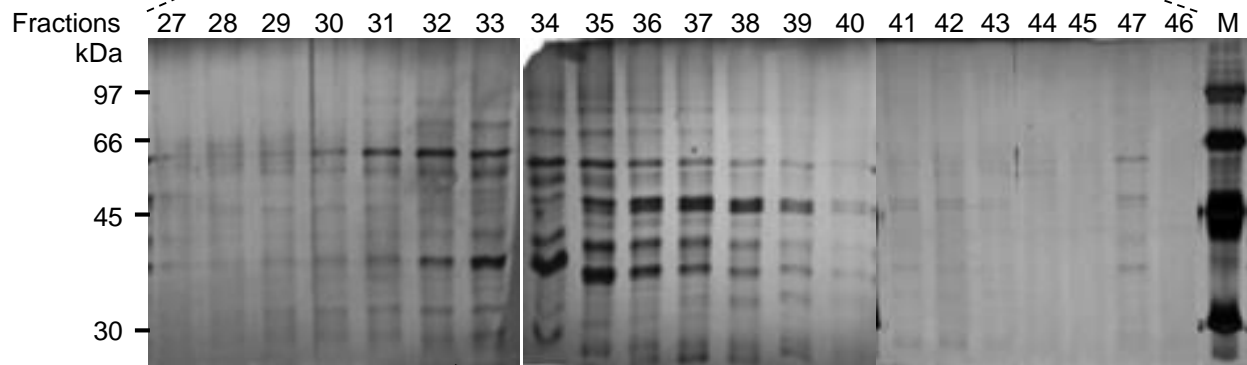
**Figure 8.** Deletion of *SXM1* strongly affects the subcellular localization of Rrp6 mutants. **(A)** Fluorescence microscopy for the analysis of the subcellular localization of GFP-Rrp6 mutants in *Δsxm1* cells growing at 25°C. Mutants Rrp6<sub>1-106</sub> and Rrp6<sub>532-619</sub> were analyzed in this strain. Quantification is shown on the right. **(B)** Western blot for the determination of the expression of the GFP-Rrp6 mutants in *Δsxm1* strain. Arp2 was used as an internal control. Mutants are expressed in higher levels than full-length Rrp6, and their levels do not decrease at 37°C. **(C)** Quantitative analysis of *Δsxm1* cells expressing Rrp6 mutants. Quantitative analysis of cells expressing GFP-fused Rrp6 mutants. (N), protein concentrated in the nucleus; (N+C) protein present both in nucleus and cytoplasm; (C), protein visible mainly in the cytoplasm. Numbers correspond to the percentage of cells showing each phenotype.

**Figure 9.** Analysis of the subcellular localization of Rrp6 or Rrp6 mutants in *Δrrp47* cells. Fluorescence microscopy for the analysis of the subcellular localization of GFP-Rrp6, or the mutants containing the Rrp47-interacting domain Rrp6<sub>1-106</sub>, Rrp6<sub>1-398</sub>, Rrp6<sub>1-619</sub>, and Rrp6<sub>1-693</sub> in *Δrrp47* cells. Absence of Rrp47 affects localization of these Rrp6<sub>1-106</sub> and Rrp6<sub>1-398</sub> mutants. **(B)** Analysis of the expression levels of Rrp6 deletion mutants in *Δrrp47* cells by western blot. Mutants are expressed in higher levels than full-length Rrp6. Arp2 was used as an internal control. **(C)** Analysis of growth of *Δrrp47* cells expressing either GFP or the different GFP-Rrp6 constructs at 25°C or 37°C. Full-length Rrp6 and mutants Rrp6<sub>1-619</sub> and Rrp6<sub>1-693</sub> complement growth of *Δrrp47* at 37°C.

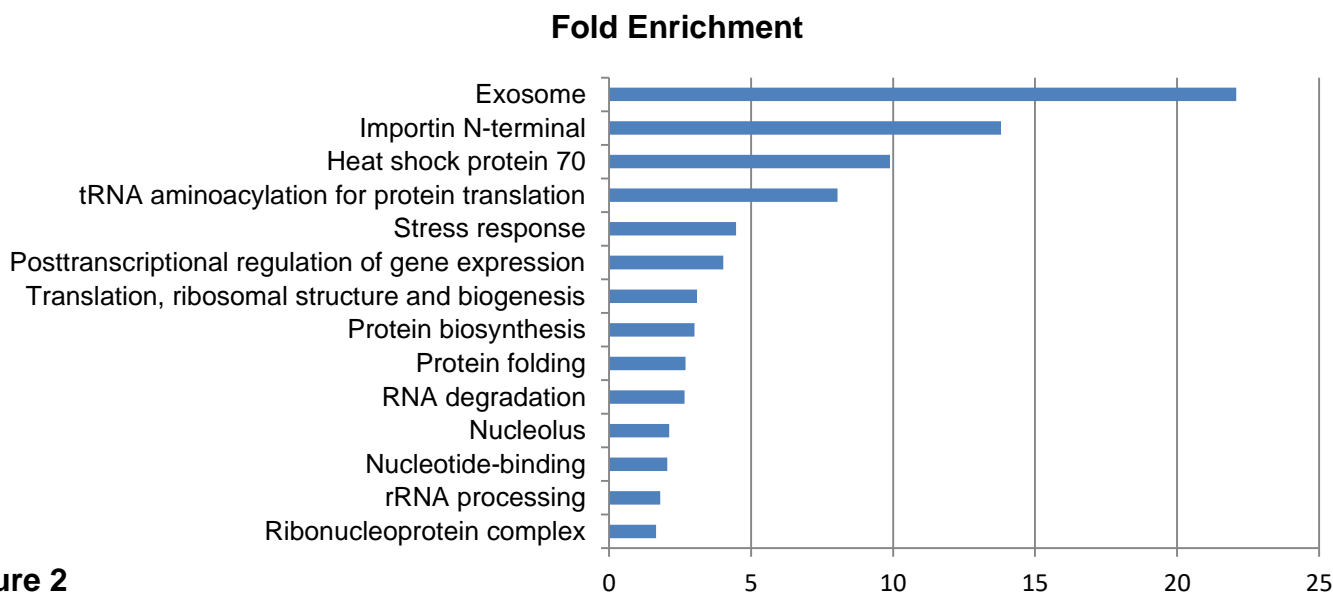
**Figure 10.** Score chart to summarize the effects of karyopherins depletions (Srp1 and Kap95; Gal or Glu) or deletions (Sxm1, Msn5, Kap104, Nmd5, Kap120, Kap114, Kap122, Kap123; 25°C or 37°C) on Rrp6 localization **(A)**. **(B)** Localization of Rrp6 deletion mutants in *Δrrp6* strain. **(C)** Localization of Rrp6 deletion mutants in *Δsxm1* strain. **(D)** Localization of Rrp6 deletion mutants in *Δrrp47* strain. N, nuclear localization; N>C, mainly nuclear, but also present in cytoplasm; N=C, protein visualized both in nucleus and cytoplasm; N<C, protein mainly present in cytoplasm. **(E)** Model of the alternative and overlapping pathways for the nuclear import of Rrp6. Rrp6 transport to the nucleus can be facilitated by the α/β dimer Srp1/Kap95 recognizing the canonical NLS3 at the C-terminal portion of Rrp6, or by the β importins Kap95 and Sxm1 recognizing one of its nuclear localization signals. Alternatively, Rrp6 could be transported in a subcomplex with the exosome complex or other exosome-interacting proteins.

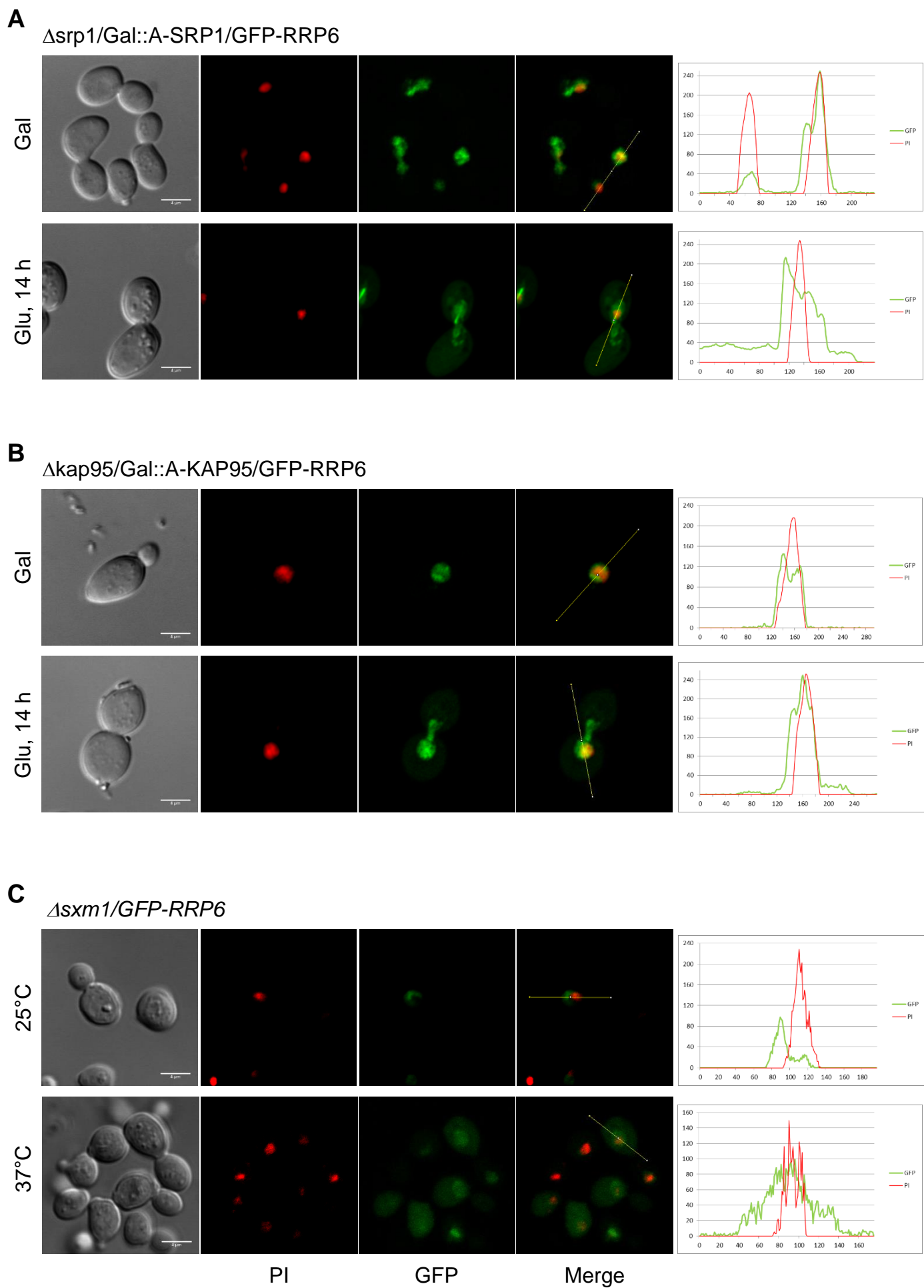


**Figure 1**

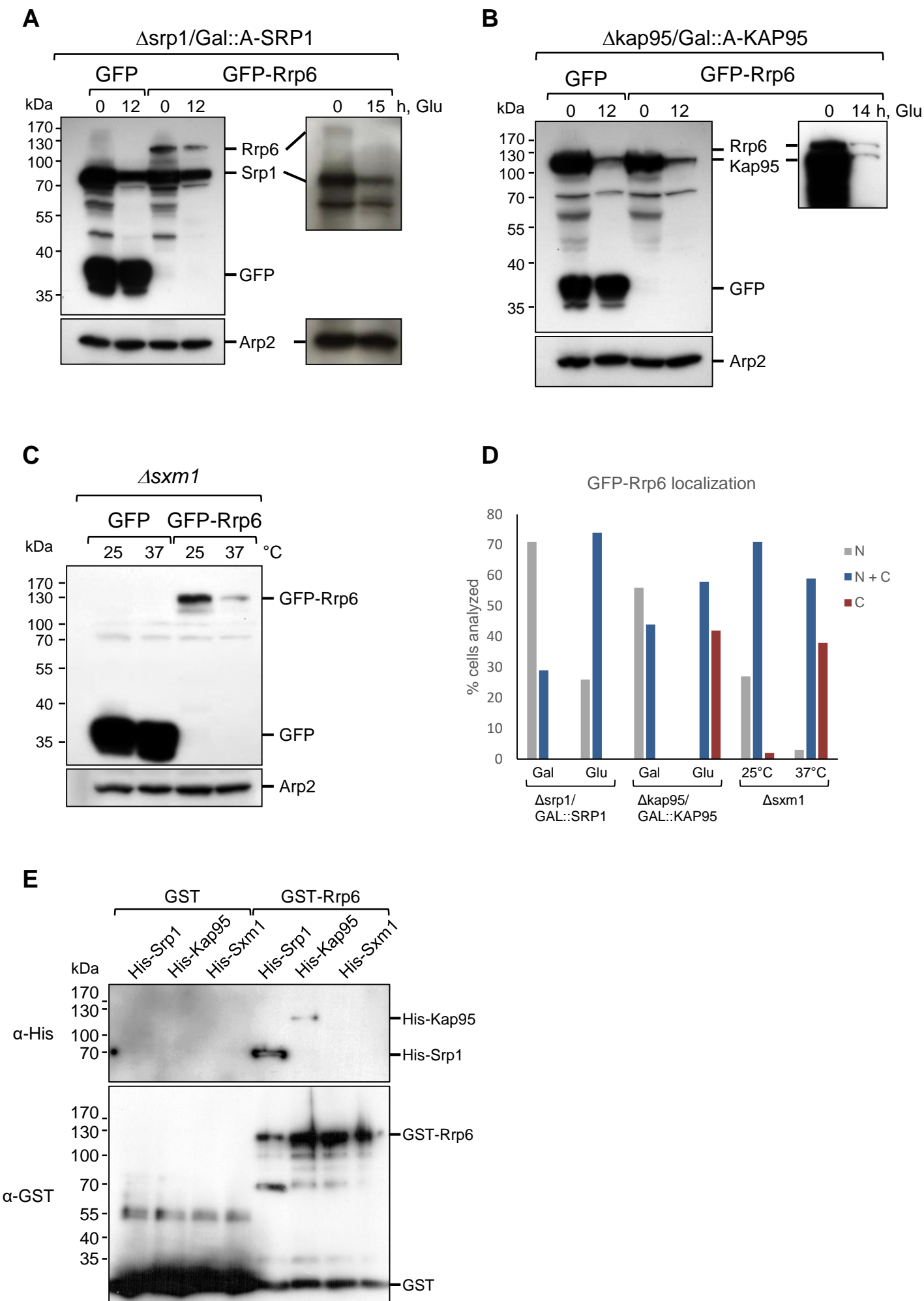
**A****B****C**

Exosome subunits		Exosome cofactors	Karyopherins	Chaperones	
Rrp6	Rrp44	Mpp6	Kap95	Ssa1	Cdc37
Rrp4	Rrp45	Mtr4	Srp1	Ssa2	Hsp82
Rrp40	Rrp46		Cse1	Ssb1	Hsc82
Rrp41	Mtr3		Kap123	Sse1	Hsp104
Rrp42	Csl4		Kap114	Sse2	Sti1
Rrp43			Sxm1		

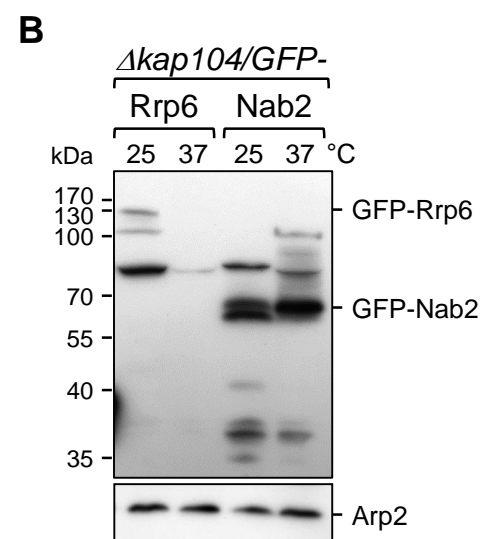
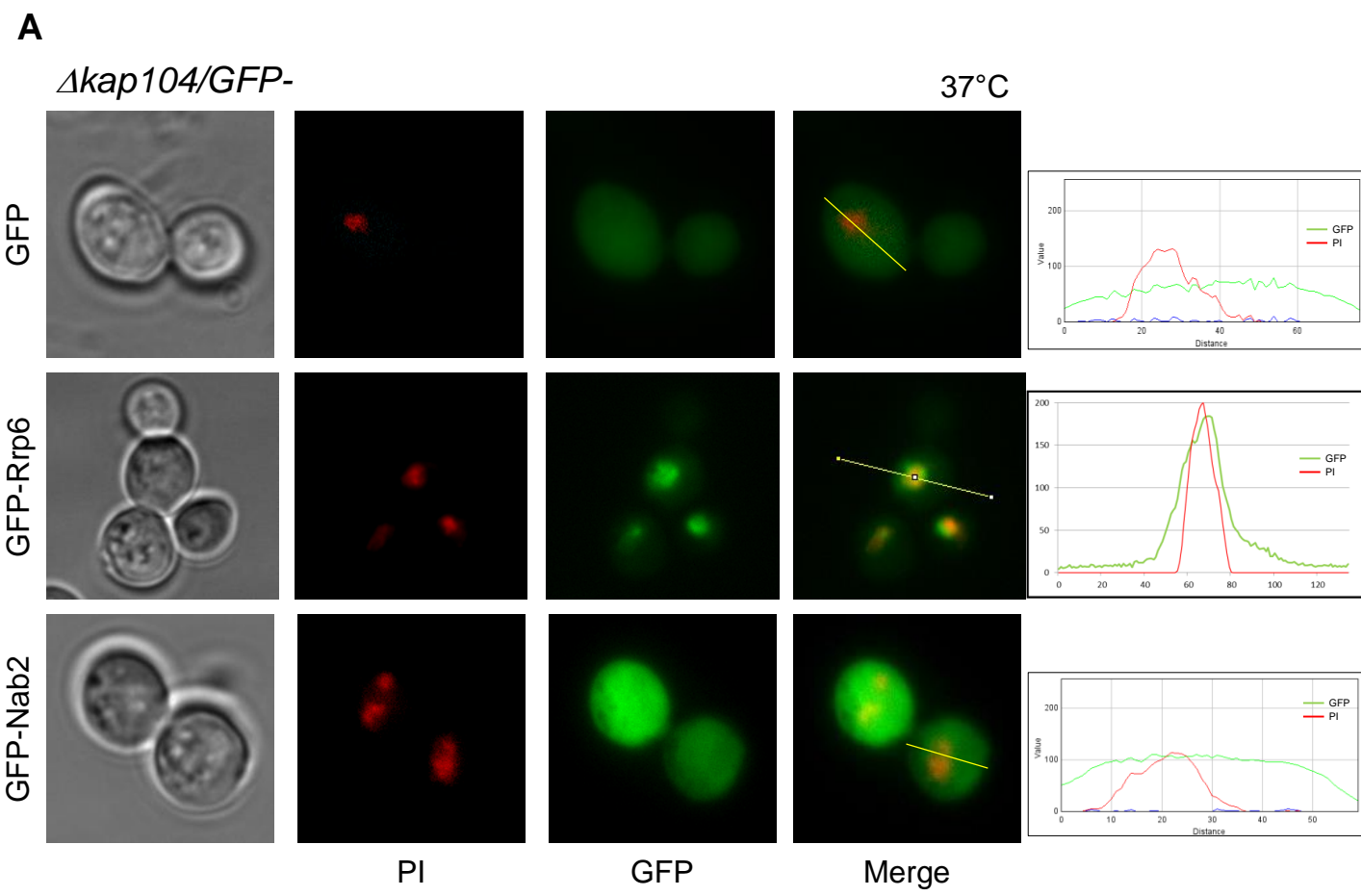
**D****Figure 2**



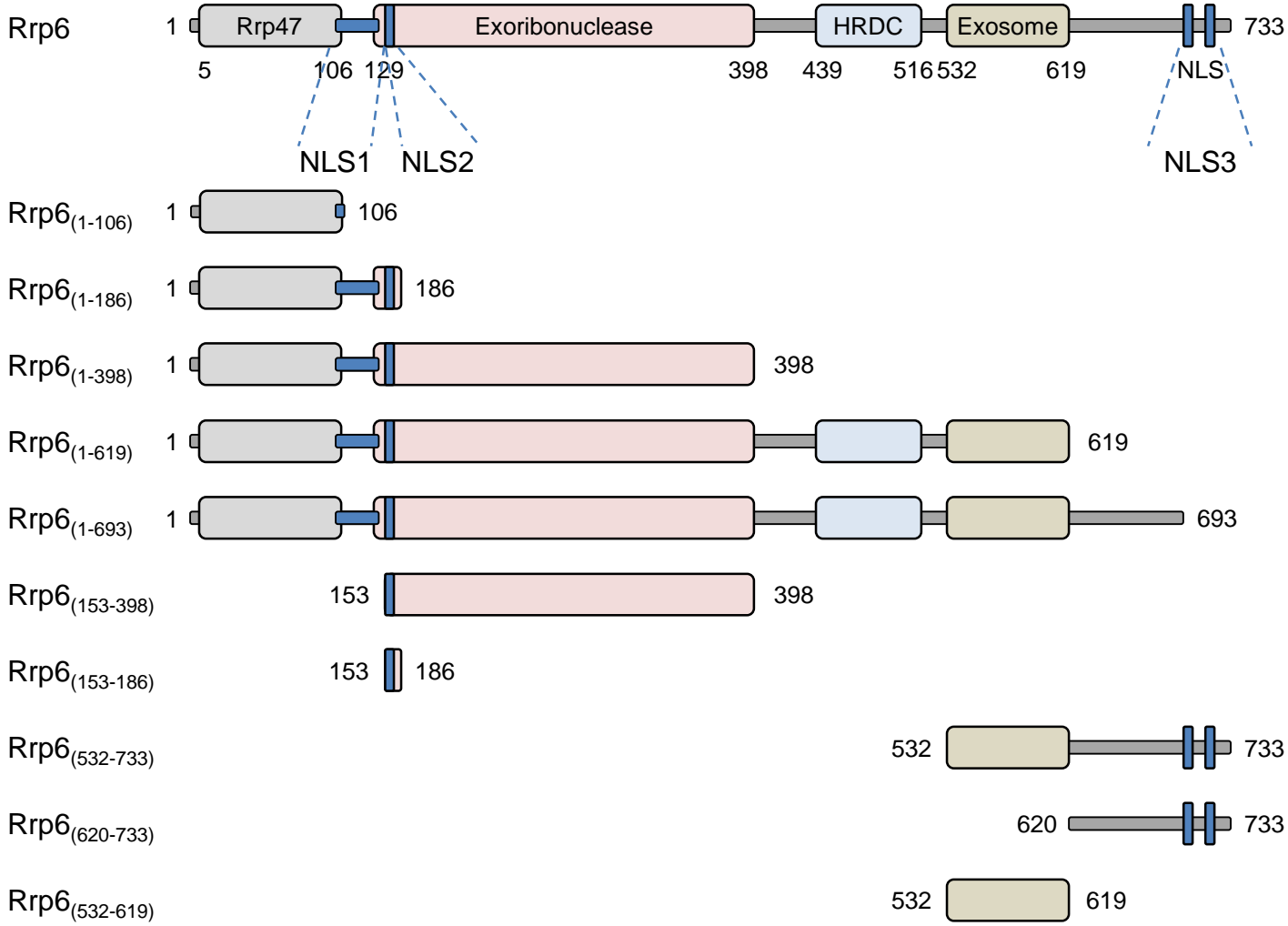
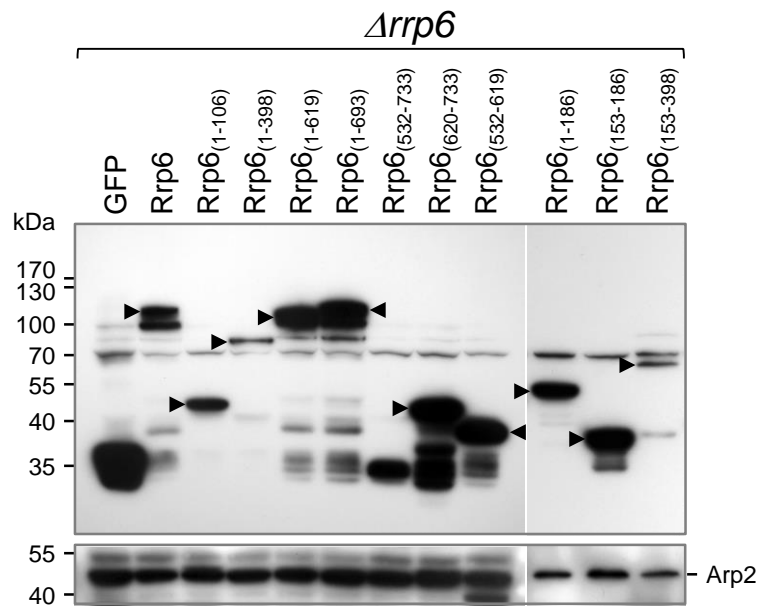
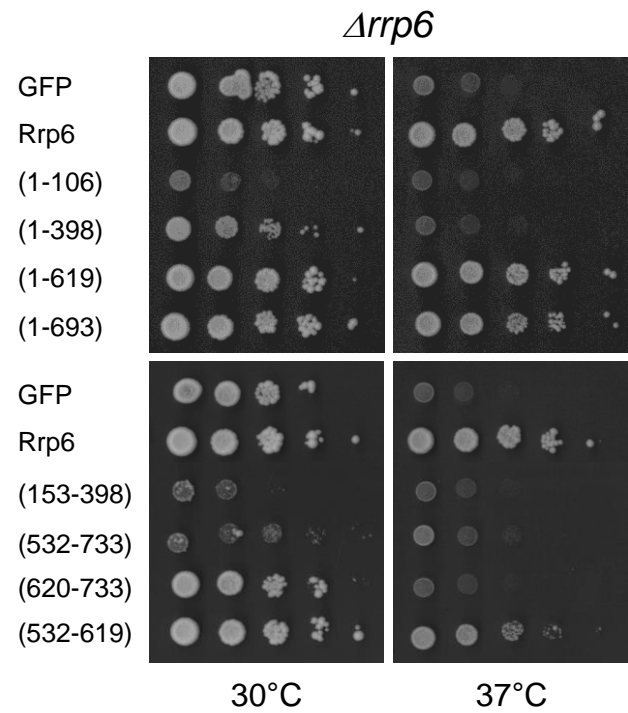
**Figure 3**

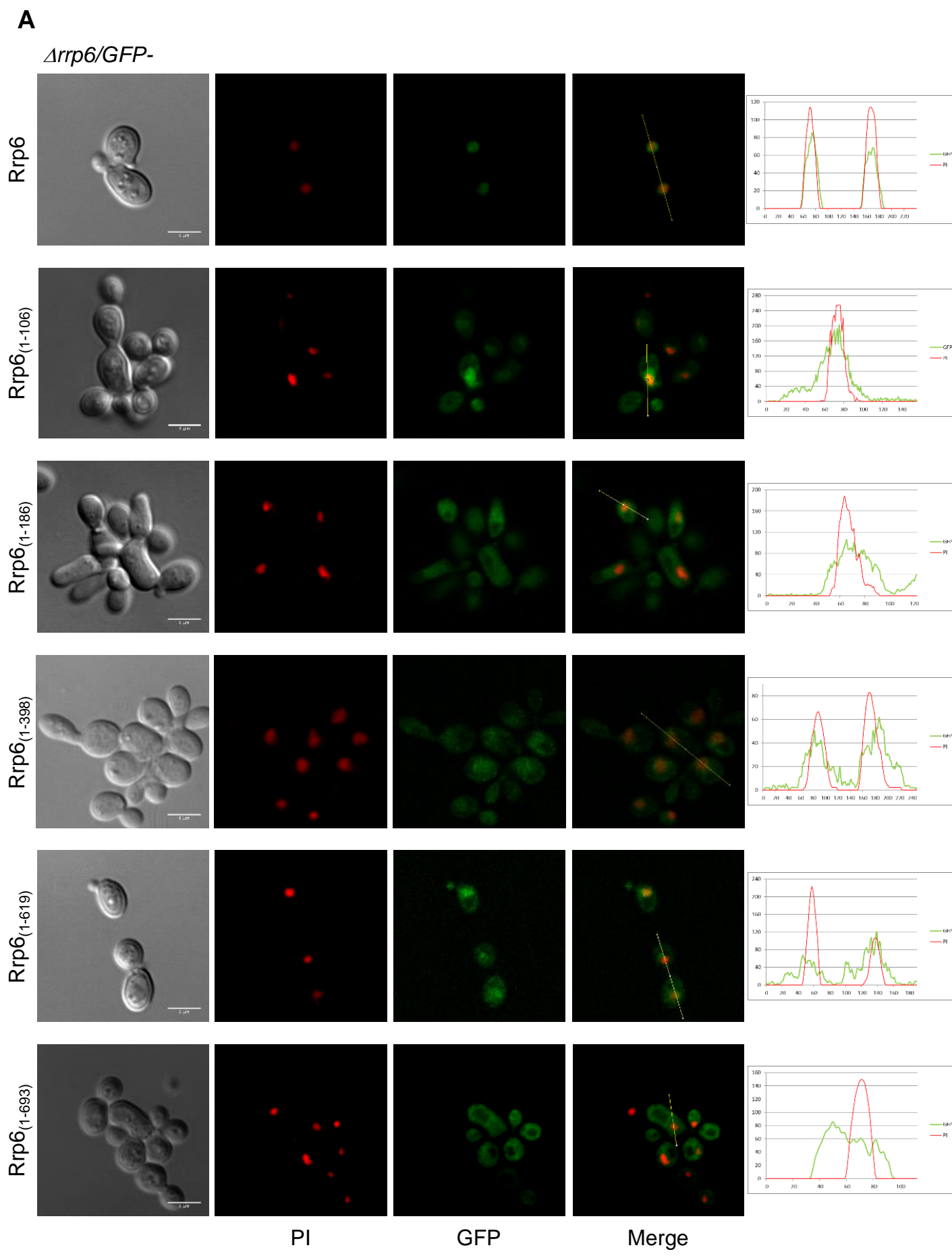


**Figure 4**

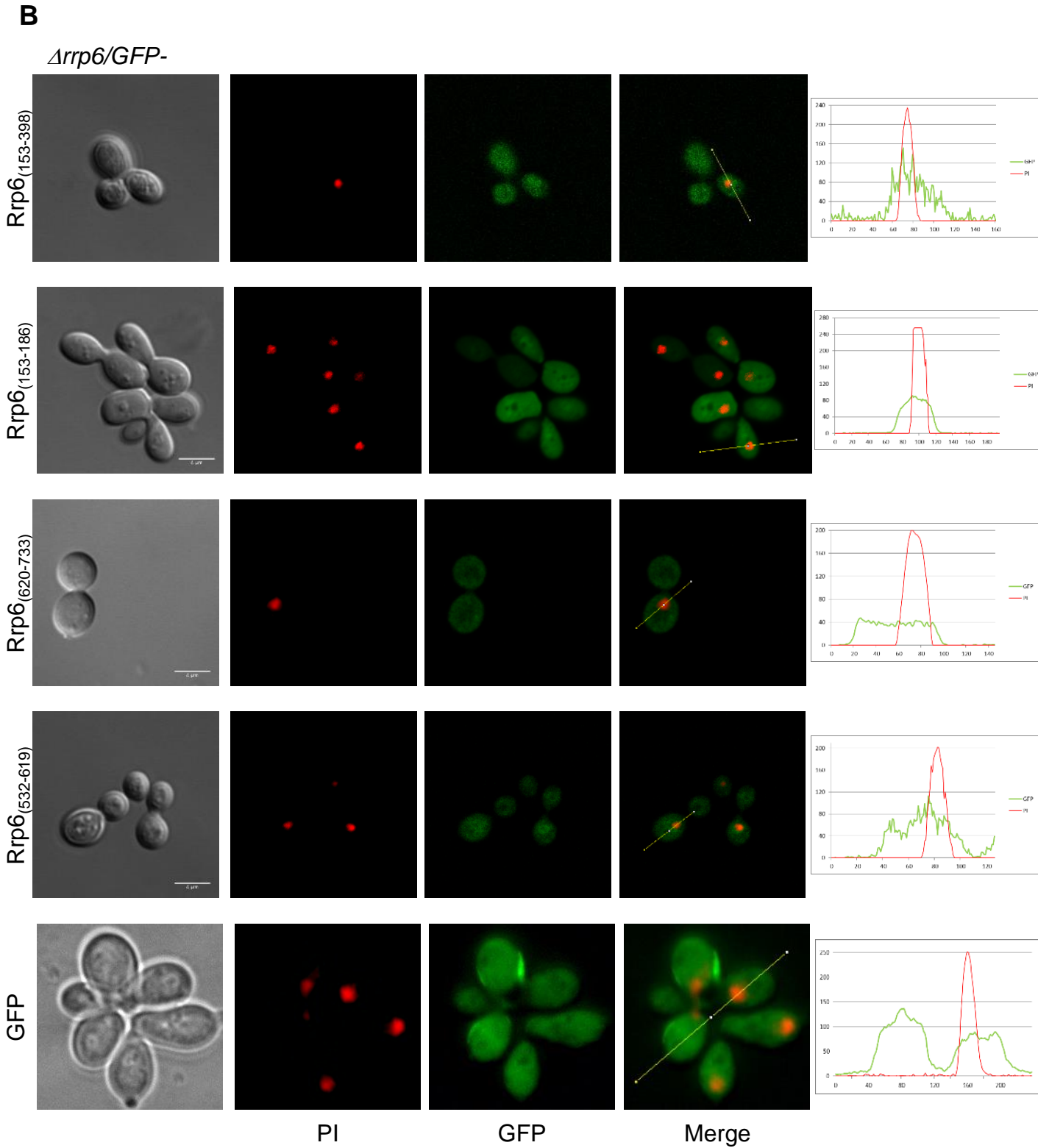


**Figure 5**

**A****B****C****Figure 6**



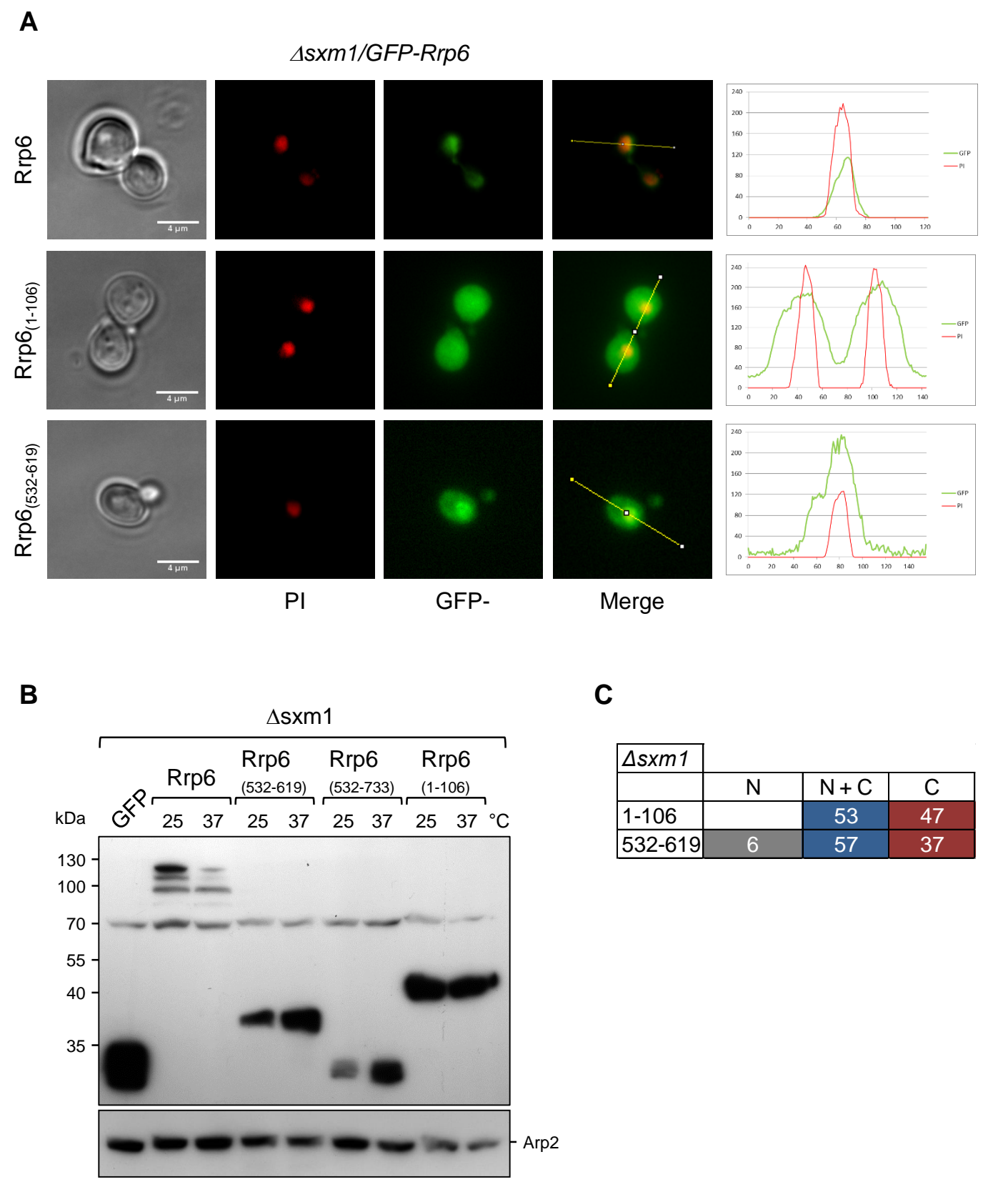
**Figure 7**



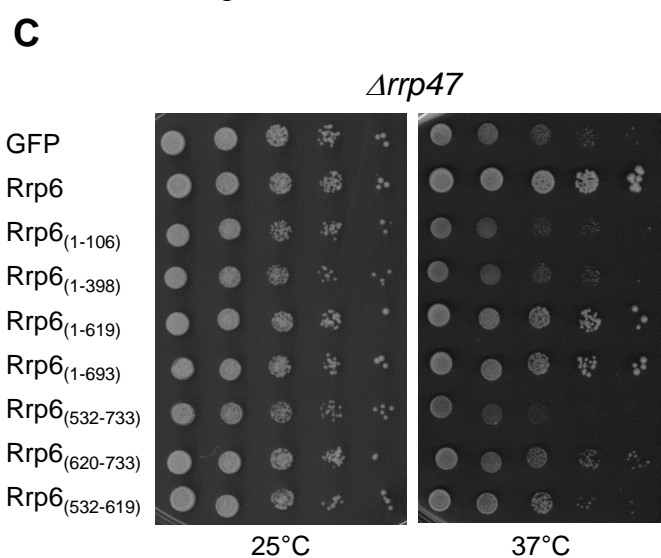
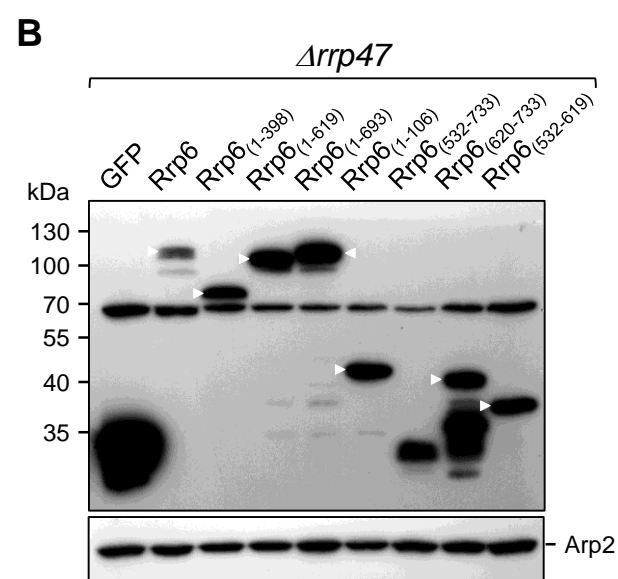
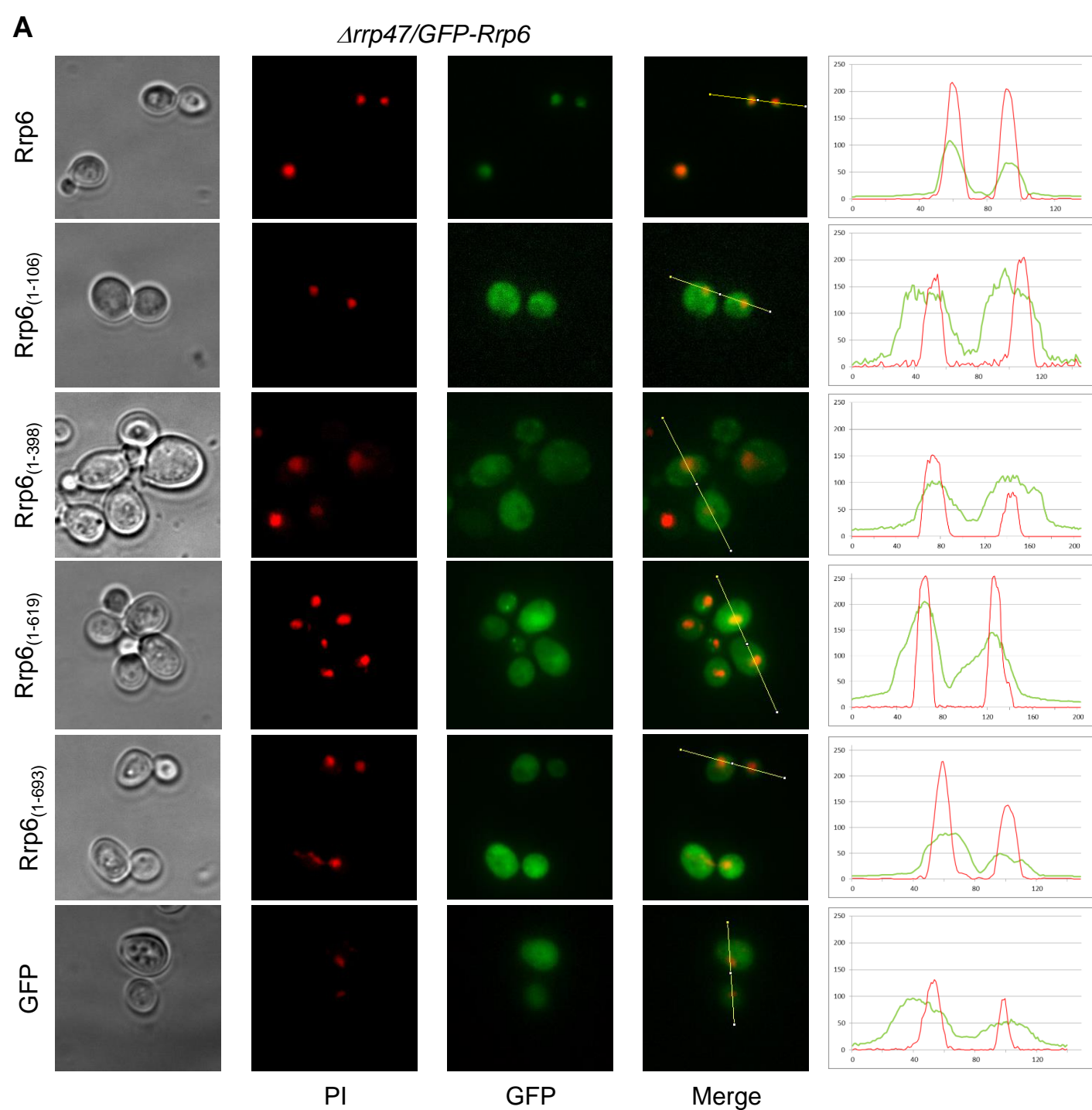
**C**

	N	N + C	C
Rrp6	100		
1-106		100	
1-186		100	
1-398		100	
1-619		100	
1-693			100
153-398			100
153-186			100
620-733		63	37
532-619		100	

**Figure 7**  
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**Figure 8**



**Figure 9**

**A**

		N + C		
	N	N > C	N = C	N < C
Gal::Srp1	Gal	Glu		
Gal::Kap95	Gal	Glu		
$\Delta sxm1$		25°C	37°C	
$\Delta msn5$	25°C	37°C		
$\Delta kap120$	25°C	37°C		
$\Delta kap104$	25°C	37°C		
$\Delta kap122$	25°C			
	37°C			
$\Delta kap123$	25°C			
	37°C			

**B**

$\Delta rrp6$

		N + C		
	N	N > C	N = C	N < C
Rrp6				
Rrp6(1-106)				
Rrp6(1-186)				
Rrp6(1-398)				
Rrp6(1-619)				
Rrp6(1-693)				
Rrp6(153-398)				
Rrp6(153-186)				
Rrp6(620-733)				
Rrp6(532-619)				

**C**

$\Delta sxm1$

		N + C		
	N	N > C	N = C	N < C
Rrp6				
Rrp6(1-106)				
Rrp6(532-619)				

**D**

$\Delta rrp47$

		N + C		
	N	N > C	N = C	N < C
Rrp6				
Rrp6(1-106)				
Rrp6(1-398)				
Rrp6(1-619)				
Rrp6(1-693)				

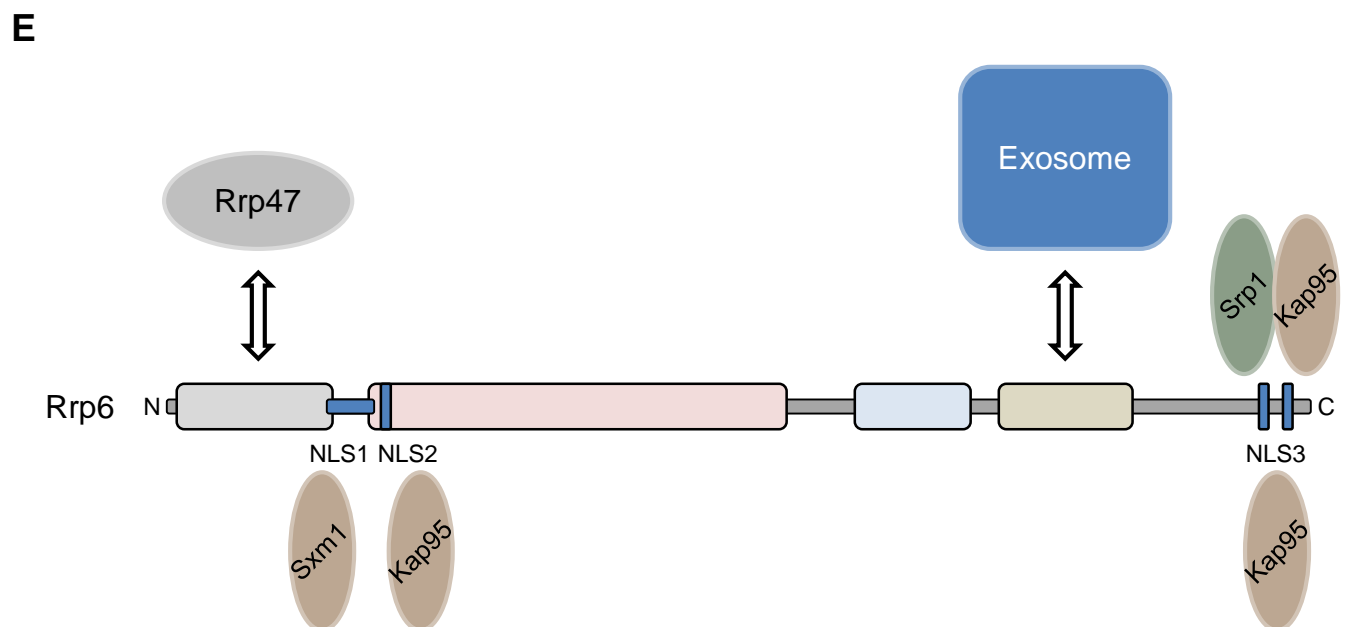


Figure 10

**Identification of karyopherins involved in the nuclear import of RNA exosome subunit Rrp6 in *Saccharomyces cerevisiae***

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