Karyopherin Msn5 is involved in a novel mechanism controlling the cellular level of cell cycle regulators Cln2 and Swi5

Inma Quilis*abc, Francisco J. Taberner*†abc, Carlos A. Martínez-Garay‡abc, Paula Alepuzabc, and J. Carlos Igualabc

*Departament de Bioquímica i Biologia Molecular, Universitat de València, Valencia, Spain; †Estructura de Recerca Interdisciplinar en Biotecnologia i Biomedicina (ERI BIOTECMED), Universitat de València, Valencia, Spain

ABSTRACT

The yeast β-karyopherin Msn5 controls the SBF cell-cycle transcription factor, responsible for the periodic expression of CLN2 cyclin gene at G1/S, and the nuclear export of Cln2 protein. Here we show that Msn5 regulates Cln2 by an additional mechanism. Inactivation of Msn5 causes a severe reduction in the cellular content of Cln2. This occurs by a post-transcriptional mechanism, since CLN2 mRNA level is not importantly affected in asynchronous cultures. Cln2 stability is not significantly altered in msn5 cells and inactivation of Msn5 causes a reduction in protein level even when Cln2 is stabilized. Therefore, the reduced amount of Cln2 in msn5 cells is mainly due not to a higher rate of protein degradation but to a defect in Cln2 synthesis. In fact, analysis of polysome profiles indicated that Msn5 inactivation causes a shift of CLN2 and SWI5 mRNAs from heavy-polysomal to light-polysomal and non-polysomal fractions, supporting a defect in Cln2 and Swi5 protein synthesis in the msn5 mutant. The analysis of truncated versions of Cln2 and of chimeric cyclins combining distinct domains from Cln2 and the related Cln1 cyclin identified an internal region in Cln2 from 181 to 225 residues that when fused to GFP is able to confer Msn5-dependent regulation of protein cellular content. Finally, we showed that a high level of Cln2 is toxic in the absence of Msn5. In summary, we described that Msn5 is required for the proper protein synthesis of specific proteins, introducing a new level of control of cell cycle regulators.

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Introduction

Nucleocytoplasmic trafficking is a relevant mechanism for the regulation of different cellular processes. The β-karyopherins are soluble receptors that mediate the transport of proteins into and out of the nucleus through the nuclear pore complex (NPC), which is composed in S. cerevisiae of approximately 30 different nucleoporins (Nups)
β-karyopherins act as importins or exportins by recognizing specific features in their cargoes: nuclear localization signals (NLS) or nuclear export signals (NES), respectively [4,5]. The translocation of cargo-bound β-karyopherins through the NPC is facilitated by specific interaction with a class of nucleoporins rich in phenylalanine-glycine repeat motifs (FG-Nups) [6,7]. The binding and release of cargo proteins by karyopherins is controlled by the Ran GTPase cycle [8].

Msn5 is a member of the β-karyopherin family with many pleotropic functions [9,10]. It controls the nuclear export of the transcription factors Pho4 [11], Crz1 [12], Mig1 [13], Msn2/4 [14], Rtg1/3 [15], Aft1 [16], Maf1 [17], and Haa1 [18], which are involved in phosphate metabolism, calcium signaling, glucose repression, stress response, nitrogen regulation, iron response, RNApol III transcription and lactic acid tolerance, respectively. It also participates in different aspects of cell cycle control by mediating nuclear export of the transcription factors Swi5 [19], Swi6 [20] and Whi5 [21], the CKI inhibitor Far1 [22], the APC activator Cdh1 [23], and the Ste5 scaffold protein [24] and the HO endonuclease [25] involved in mating. In addition to its role in protein export, Msn5 may play a secondary role in the trafficking of tRNA between the nucleus and the cytoplasm [26,27]. In particular, Msn5 is involved in the re-export of mature tRNAs to the cytoplasm in the tRNA retrograde pathway [28]. As a consequence, tRNAs accumulate in the nucleus in msn5 mutant cells. In spite of this, no general translational defects have been described for the msn5 mutant; rather, translation of only twelve mRNAs related to methionine and arginine biosynthetic pathways are apparently affected by Msn5 inactivation [29].

Cell cycle progression is governed by the sequential activation of different cyclin-CDK complexes. For the yeast S. cerevisiae, nine different cyclins (Cln1-3 and Clb1-6) activate the Cdc28 protein, the only yeast CDK that performs an essential function in cell cycle progression [30]. Kinase activities existing at any time are determined by the levels of cyclins and CDK inhibitors (CKI). Two molecular mechanisms, gene transcription and protein degradation, are fundamental to control these key cell cycle regulators. The coordinated expression of different sets of genes, organized in transcriptional waves along the cell cycle, is a very common strategy in cell cycle control in all eukaryotes [31]. All the cyclins associated with Cdc28 CDK are periodically expressed. In the case of the cyclin studied in this work, CLN2 gene is expressed during the G1/S transition by the transcription factor SBF, a heterodimer composed by the Swi4 and Swi6 proteins [32]. The second major mechanism involved in the control of the cellular levels of cell cycle regulators is proteolysis by means of ubiquitination and degradation in the proteasome [33,34]. Two ubiquitin ligases play a central role in cell cycle control, APC, which regulate mitosis, and SCF, involved mainly in the control of the G1/S transition. SCF regulates the G1/S transition through the degradation of G1 cyclins and CKI [35]. The SCF complex consists of four subunits: Skp1, Cdc53, Rbx1, and an adapter protein with an F-box, which is responsible for substrate recognition. SCF associated to the F-box protein Grr1 (SCF\(^{Grr1}\)) is the main ubiquitin ligase involved in Cln2 degradation [36,37].

In addition to transcription and protein degradation, other cellular processes contribute to the fine-tuning of cell cycle machinery. Spatial regulation is a common strategy to control protein function and this is also the case for cell cycle regulators. Indeed, as commented above distinct cell-cycle transcription factors shuttle between the nucleus and the cytoplasm and functional specificity between cyclins can be determined, at least in part, by the differences in their subcellular localization which targets their associated CDK activity to specific locations. This is well illustrated by the case of Cln cyclins [38–40]. In the case of Cln2, it is present in both the nucleus and the cytoplasm. It contains a NLS sequence in its N-terminal region that is responsible for nuclear import via the Kap95-Kap60 classical nuclear import pathway and a NES region between amino acids 225–299 that mediates its nuclear export by the Msn5 karyopherin [41]. This export mechanism helps to confer specific functionality to Cln2. On the other hand, examples of post-transcriptional regulation of cyclin mRNAs have been described. Thus, the translation of CLN3 mRNA is repressed under nutrient-deprivation conditions, providing a mechanism coupling cell growth and division [42,43], and the RNA binding
protein Whi3 binds to CLN3 mRNA to promote turnover and to inhibit translation [44,45]. As regards the other G1 cyclins, it has been reported that Cln1 and Cln2 are down-regulated by iron and that this occurs by translational repression of their mRNA [46]. Also, the degradation of CLB2 mRNA by the MRP RNA endonuclease is important to promote mitotic exit [47].

The combination of different strategies is, therefore, a common trait in the control of cell cycle regulators in order to ensure the correct order and timing of the successive processes. We have previously described that Msn5 controls SBF, the transcription factor that regulates the periodic expression of Cln2, and the subcellular localization of Cln2 [20,41]. In this work we investigate in more depth the connection between Msn5 and Cln2 cyclin regulation describing a new mechanism by which Msn5 karyopherin regulates Cln2 protein synthesis.

Materials and methods

Strains, plasmids and growth conditions

Yeast strains used in this work are shown in Supplemental material Table S1. The msn5 mutant strains were obtained by integrating a DNA fragment amplified from the pFA6a series plasmids (from J.R. Pringle) or from strains containing the msn5A3::HIS3 or msn5Δ2::HIS3 disruption cassette [20]. Strain GAL1:HA-MSN5 was constructed by integrating at the appropriate position a DNA fragments amplified from the pFA6a series plasmids. The swi4::LEU2, swi6::TRP1 and grr1::LEU2 strains derived by using disruption-cassette plasmids BdB194 and BdB197 (from L. Breeden) and pBM1829 (from M. Johnston), respectively. The cln2::TRP1 mutant strain was obtained by integrating a DNA fragment amplified from the pFA6a series plasmids. Strains expressing the Cln2 protein or truncated versions of it tagged at C-terminal with 3xHA, were constructed by integrating at the appropriate position DNA fragments amplified from the pFA6a series plasmids. Strains expressing the Cln2 protein tagged at C-terminal with 6xHA, was constructed by integrating at the appropriate position a DNA fragment amplified from pGA2256 plasmid (from Dr. G. Ammerer). Strains JCY1775 and JCY1845 were constructed by integrating at the URA3 locus EcoRV-digested Yip-CLN2p::GFP or Yip-CLN2p::CLN2181–225-GFP plasmids, respectively.

Centromeric plasmids pCLN1 and pCLN2, containing HA-tagged versions of Cln1 or Cln2, and plasmids ptetO-CLN2 overexpressing the CLN2 gene under the control of the tetO2 promoter, were a gift from Dr. M. Aldea. Centromeric plasmids expressing the chimeric cyclin Cln21–476–Cln1477–546, Cln21–370–Cln1371–546, Cln21–299–Cln1313–546, Cln21–225–Cln1227–356 and Cln21–200–Cln1201–546, were constructed as previously described [41]. To obtain pCLN24T3S, a XhoI-KpnI fragment from pCLN2 containing the CLN2 promoter and the N-terminal part of the protein, was cloned in XhoI-KpnI digested pGAL1:CLN2−4T3S-HA plasmid (from C. Wittenberg). Now, Cln24T3S is expressed under the control of the CLN2 promoter. Yip-CLN2p::GFP contains, in order, the CLN2 promoter, the GFP coding region and the ADH1 terminator in YIpplac211. This plasmid was constructed in a three-step process. First, the ADH1 terminator including the stop codon (+772, +1011), amplified from pFA6a-GFP(S65T) with a forward oligo containing a SalI site and a reverse oligo with a PstI site, was cloned in YIp-CLN2p::GFP or Yip-CLN2p:GFP.

Yeast cells were grown on standard yeast extract-peptone-dextrose (YPD) media or synthetic dextrose (SD) minimal media supplemented as required. To repress the tetO7 promoter, doxycycline was added to a concentration of 10 µg/mL. To evaluate protein stability, 100 µg/mL cycloheximide was added to exponentially growing cells.
Western blot analysis

Approximately 10^8 cells were collected, resuspended in 100 μL water and, after adding 100 μL 0.2 M NaOH, they were incubated for 5 min at room temperature. Cells were collected by centrifugation, resuspended in 50 μL sample buffer, and incubated for 5 min at 95°C. Extracts were clarified by centrifugation, and equivalent amounts of protein were resolved in an SDS-PAGE gel and transferred onto a nitrocellulose membrane. The primary antibodies used were monoclonal anti-HA 12CA5 antibody (Roche Diagnostics), monoclonal anti-HA peroxidase 3F10 antibody (Roche Diagnostics), monoclonal anti-GFP (Roche Diagnostics), monoclonal anti Cdc2 p34 (PSTAIRE) (Santa Cruz Biotechnology Inc.), and anti α-tubulin antibody (Serotec Ltd.). Blots were developed with anti-mouse IgG and anti-rabbit IgG Horseradish Peroxidase conjugate (Thermo Fisher Scientific,) or anti-rat IgG Horseradish Peroxidase conjugate (GE Healthcare) using the Supersignal West Femto Maximum Sensitivity Substrate (Thermo Scientific). Bands were quantified with an ImageQuant™ LAS 4000mini biomolecular imager (GE Healthcare).

Gene expression analysis

RNA was purified from exponentially growing cells using RNeasy®Mini Kit (Qiagen) following the manufacturer instructions. Previously, yeast cells were broken in a FastPrep Precellys24 (Bertin technologies) with glass beads in the recommended kit buffer. The sample was incubated with Turbo DNase (Ambion) and after DNase inactivation and incubation with oligo dT, cDNA was obtained with Improm-II® Reverse Transcriptase and Recombinant RNasin® (Promega) following the manufacturer instructions. The cDNA was analyzed by quantitative RT-PCR in a DNA Engine Peltier Thermal Cycler (Bio Rad) using the SYBR Premix Ex Taq Tli RNase H Plus Green with ROX (Takara).

Northern analysis were carried out as previously described [19]. CLN2 and ACT1 mRNA were detected using 32P-labeled probes obtained with HighPrime (Roche) according to the manufacturer instructions.

Polyribosome profile analysis

Cells were grown to the exponential phase in YPD. Preparation of cells and polysome gradients were performed as previously described [48], with some modifications. Approximately 10^9 cells were chilled for 5 min on ice in the presence of 0.1 mg/mL cycloheximide. Cells were harvested by centrifugation at 6000 × g for 4 min at 4°C and washed twice with 1 mL of lysis buffer (20 mM Tris-HCl, pH 8, 140 mM KCl, 5 mM MgCl2, 0.5 mM dithiothreitol, 1% Triton X-100, 0.1 mg/mL CHX, and 0.5 mg/mL heparin). Cells were resuspended in 700 µl of lysis buffer, a 0.5-mL volume of glass beads was added, and cells were disrupted by vortexing 8 times for 30 s with 30 s of incubation on ice in between. Lysates were cleared by centrifugation at 5000 rpm for 5 min at 4°C, after which the supernatant was recovered and centrifuged at 8000 rpm for 5 min at 4°C. Finally, glycerol was added to the supernatant at a final concentration of 5% and extracts were frozen in liquid nitrogen and stored at −70°C. Samples of 8.5 A260 units were loaded onto 5–50% sucrose gradients and were separated by ultracentrifugation for 2 h 40 min at 35,000 rpm in a Beckman SW41 rotor at 4°C. Gradients were then fractionated by isotonic pumping of 60% sucrose from the bottom and either eleven 1 mL-samples or twenty-two 0.5 mL-samples were recovered. This was followed by a recording of the polysomal profiles by online UV detection at 240 nm (Density Gradient Fractionation System; Teledyne Isco, Lincoln, NE). For the RNA analyzes of the polysomal fractions, 6 µL of mixed lys and spo mRNAs at 3 ng/µl from Bacillus subtilis were added to each fraction before extraction in order to normalize all the values with spiked-in mRNA levels of B. subtilis lys and spo.

The fractions were mixed with a 2-fold volume of 8 M guanidine-HCl; the RNAs were then precipitated in 3 volumes of 100% ethanol and stored at 20°C overnight. Nucleic acids were collected by centrifugation (13,000 rpm for 20 min at 4°C), followed by a wash step with 70% ethanol. The pellets were then dissolved in 500 μl of water, followed by phenol-chloroform-isoamyllic alcohol extraction. After centrifugation (10,000 rpm for
5 min at 4°C), the supernatant was precipitated with LiCl (final concentration, 2.5 M) at −80°C. RNA samples were collected at 12,000 rpm for 15 min at 4°C and washed with 70% ethanol. The RNA pellets were resuspended in nuclease-free H₂O. Equal volumes of RNA from each fraction were retrotranscribed using SuperScript™ IV VILO™ Master Mix with ezDNase (invitrogen, Thermo Fisher Scientific) following the manufacturer instructions. The cDNA was analyzed by quantitative RT-PCR as described above.

Results

Msn5 regulates Cln2 protein level through a post-transcriptional mechanism

Our group had previously described that Msn5 mediates nuclear export of Cln2 [41] and that Msn5 regulates the SBF transcription factor responsible for periodic expression of the CLN2 gene [20]. During the course of these works, we envisaged the possibility that Msn5 could affect Cln2 by an additional mechanism. As shown in Figure 1(a), the amount of Cln2 in msn5 mutant cells is severely reduced compared to the wild-type cells and Cln2 levels are recovered after introduction of a plasmid containing the MSN5 gene. A similar effect was observed in four additional msn5 mutant strains (Figure 1(b)), confirming that Msn5 is required for proper levels of Cln2 protein. CLN2 is a periodically expressed gene; therefore, the change in Cln2 protein level in msn5 mutant cells could be due to an alteration in the distribution of cells along the different phases of the cell cycle. However, this was not the case, because there are not differences in the cell-cycle distribution of cells between the wild type and msn5 mutant strains (Supplemental material Figure S1). Alternatively, the reduced cellular level of Cln2 protein could be attributed to a defect in CLN2 gene expression since Msn5 controls the SBF transcription factor [20]. Strikingly, the inactivation of the SBF components Swi4 or Swi6 in asynchronous cultures has very mild or no effect in Cln2 protein level compared to what is observed in msn5 mutant cells (Figure 1(b)). This suggests that other mechanisms besides transcriptional regulation must affect the control of cellular content of Cln2 by Msn5. Supporting this, Cln2 protein level also diminished when CLN2 was ectopically expressed under the control of the tetO₂ promoter (Figure 1(b)).

In order to confirm that the severe reduction in Cln2 protein level in the absence of Msn5 is caused by a post-transcriptional mechanism, we carried out, in parallel, an analysis of CLN2 mRNA and Cln2 protein in the wild type and msn5 mutant strains. Remarkably, CLN2 mRNA level in the msn5 mutant strain in asynchronous cultures was only slightly affected in comparison with the severe effect observed in Cln2 protein (Figure 1(c,d)). These results rule out the possibility of a reduced expression or stability of the CLN2 mRNA as the origin for the reduced Cln2 protein level in msn5 mutant cells, and clearly demonstrate that Msn5 is involved in a post-transcriptional mechanism important for a proper protein synthesis from the CLN2 mRNA.

Stabilization of Cln2 does not abolish the reduction of protein level in msn5 mutant cells

Msn5 acts as an exportin of Cln2 and it has been reported in several cases that there are changes in protein stability depending on subcellular localization [49,50]. Taking this into account, it is possible that the reduced amount of Cln2 in msn5 mutant cells was caused by a higher rate of protein degradation due to Cln2 accumulation in the nucleus. Nevertheless, other results raise doubts on this hypothesis. The analysis of Cln2 protein decay in a transcriptional shut-off assay indicated that the stability of Cln2 is not significantly affected in the msn5 mutant strain when compared to that observed in the wild-type strain (Figure 2(a)). Degradation of Cln2 is mainly mediated by the SCF₆Grr1 ubiquitin ligase [36,37]. Importantly, the inactivation of Msn5 in mutant strains in Cdc53 or Grr1, components of the SCF ubiquitin ligase, still caused a reduction in Cln2 protein levels (Figure 2(b,c)). Finally, a variant protein mutated in phosphorylation sites, Cln2₄T₃S, which is stable and accumulates in the nucleus [38,51], also displayed a reduced level upon Msn5 inactivation (Figure 2(d)).
These results strongly support that Msn5 inactivation affects the Cln2 protein level by a mechanism distinct to changes in protein stability. The steady-state level of a protein is the result of the balance between the synthesis and degradation rates. Therefore, our results strongly support that Cln2 protein synthesis takes place at a reduced rate in msn5 mutant cells.

**Msn5 is required for efficient translation of CLN2 and SWI5 mRNAs**

The results described above recalls the situation for another cell-cycle regulator protein, the transcription factor Swi5, whose protein synthesis rate is impaired in the absence of Msn5 [19]. It is necessary to clarify that this is a specific response
of Cln2 and Swi5, since Msn5 inactivation did not affect the protein content of other cell cycle transcription factors such as Swi6, Whi5 or Ace2 [19].

In order to investigate how Msn5 could control Cln2 and Swi5 synthesis, we analyzed the association of their mRNA to ribosomes in polysome profiles in sucrose gradient. The polysome profile of the \( msn5 \) mutant was very similar to the wild type profile (Figure 3(a)), in agreement with previously reported results in other experimental conditions [29]. Thus, global translation capacity does not seem to be altered in \( msn5 \) mutant cells, which is consistent with the fact that reduction of protein level is only observed in some proteins like Cln2 and Swi5 and not in other proteins. Interestingly, the cumulative amount of both \( CLN2 \) and \( SWI5 \) mRNAs relative to the control \( ACT1 \) mRNA in the monosomal and polysomal fractions is similar to that observed in total crude extracts (Figure 3(b); crude extracts in Figure 1(d) for \( CLN2 \) and [19] for \( SWI5 \)). This is an important observation that rules out any defect in \( CLN2 \) and \( SWI5 \) mRNAs export or mRNA access to the translational machinery as the cause of the reduced levels of Cln2 and Swi5 proteins; rather, it points to a defect in the translation process itself.

Although global translation was not altered in \( msn5 \) cells, it is possible that translation of a subset of mRNAs was affected. To compare the translational states of particular target mRNAs, the distribution of mRNA between the different fractions was examined. In the wild type cells, \( CLN2 \) and \( SWI5 \) mRNAs manifested a distribution very similar to that of \( ACT1 \) and \( HXX2 \) mRNA, used as positive controls of optimal translation activity. However, in the case of \( msn5 \) mutant cells there is a specific shift of \( CLN2 \) and \( SWI5 \) mRNAs from the heaviest polysome region to the non-polysomal (in particular the 80S monosomes) and light polysomal fractions (Figure 3(c) and Supplemental material Figure S2). This strongly suggests that translational efficiency of \( CLN2 \) and \( SWI5 \) mRNAs is specifically diminished in the absence of Msn5. This could certainly help to

\begin{figure}
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\includegraphics[width=\textwidth]{figure2}
\caption{Analysis of Cln2 protein levels in SCF ubiquitin-ligase mutant strains. (a) Doxycycline to a final concentration of 10 \( \mu \)g/mL was added to exponentially growing cultures of the wild type (W303-1a) and the \( msn5 \) mutant (JCY1018) strains transformed with a centromeric plasmid expressing a HA-epitope tagged Cln2 protein under the control of the tetO promoter (ptetO:CLN2). The Cln2 protein level in crude cell extracts was determined at the indicated time by western analysis with a specific anti-HA antibody. A non-specific band (n.s.) that cross-reacts with the antibody is shown as loading control. The shown result is representative of three independent assays. (b) Cells of the thermosensitive \( cdc53 \) and \( cdc53 msn5 \) (JCY684) mutants transformed with a centromeric plasmid expressing a HA-epitope tagged Cln2 protein under the control of the tetO promoter were grown at 25\(^\circ\) and transferred for 3 hours at 37\(^\circ\). Cln2 protein level was analyzed by western blot. Ponceau staining of the membrane is shown as loading control. The shown result is representative of five replicates. (c) Cln2 protein level in the wild type (JCY1357), \( msn5 \) (JCY1359), \( grr1 \) (JCY2082) and \( grr1 msn5 \) (JCY2083) strains expressing a HA-tagged version of Cln2. The shown result is representative of three replicates. (d) Cln2 protein level in the wild type (W303-1a) and the \( msn5 \) mutant (JCY1018) strains transformed with a plasmid expressing the stabilized HA-tagged Cln2 \(^{4T3S} \) variant. The shown result is representative of five replicates.}
\end{figure}
explain the reduced amount of Cln2 and Swi5 proteins observed in msn5 mutant cells.

**An internal region of Cln2 mediates the Msn5-dependent control of protein level**

After establishing that Msn5 controls Cln2 synthesis, we decided to analyze whether a specific region of CLN2 mRNA/Cln2 protein might be responsive to Msn5. It is important to point out that the decrease in Cln2 levels in the msn5 mutant strain was observed in constructs lacking the 3′-UTR (C-terminal epitope-tagged proteins) or 5′-UTR (tetO constructs), so the effect of Msn5 on Cln2 synthesis must be mediated by sequences in the coding region. In a first approach, the levels of various C-terminal truncated forms of Cln2 were analyzed in both wild type and msn5 mutant strains (Figure 4(a)). Truncated Cln2 forms encompassing fragments 1–370, 1–299 and 1–225 showed diminished protein level in msn5 compared to wild type cells. Thus, the C-terminal part of Cln2 was not required for its regulation by Msn5 and, reciprocally, region 1–225 was sufficient to mediate control by Msn5. Interestingly, this control was lost in the Cln21–181 truncated protein, since the protein concentration was the same in both msn5 and wild-type cells. This result
supports that the region from 181 to 225 is necessary for this Msn5-dependent regulatory mechanism.

With the aim of addressing this point in the context of a functional cyclin protein instead of truncated variants, we carried out a second
approach consisting in the construction of chimeric cyclins by interchanging equivalent domains between the Cln2 and Cln1 cyclins. It has to be noted that in contrast to what happen in Cln2, Cln1 protein level is not severely affected by Msn5 inactivation (Figure 4(b)). In agreement with the previous results, different chimeric cyclins containing the N-terminal region of Cln2 as far as amino acid 225 manifested a severe reduction in their protein level after inactivation of Msn5 (Figure 4(c)). Importantly, this Msn5-dependent effect was absent in the chimera Cln2^{1-200}-Cln1^{201-546}. This indicates that region from amino acid 200 to 225 of Cln2 is important for the Msn5-dependent regulation of protein level.

Once we have identified an internal region in Cln2 protein necessary for its regulation by Msn5, we wonder whether this region would be sufficient to confer Msn5-regulated synthesis to a protein. To test this possibility, the Cln2^{181-225} fragment was fused to GFP. The cellular content of control GFP was similar in wild type and msn5 mutant cells. However, the fusion to GFP of the Cln2^{181-225} fragment caused a reduction in its protein level in msn5 compared to wild type cells (Figure 5(a)). This is not due to alterations in transcription or mRNA stability since levels of mRNAs are not affected in msn5 cells (Figure 5(b)). Moreover, it is not related to changes in protein stability since Cln2^{181-225}-GFP shows the same degradation rate as GFP (Figure 5(c)). Thus, the synthesis of the Cln2^{181-225}-GFP protein is defective in the absence of Msn5.

In short, all our results indicate that a small fragment of Cln2 from residue 181 to 225, or its corresponding mRNA region, imposes a requirement of Msn5 karyopherin for the efficient synthesis of Cln2 protein.

**msn5 mutant cells are sensitive to overexpression of Cln2**

During the described work, we envisaged the possibility that msn5 cells were sensitive to high levels of Cln2. To explore this possibility in detail, we analyzed growth of wild type and msn5 mutant cells containing the ptetO:CLN2 plasmid. As it can be seen in Figure 6, while overexpression of CLN2 from the tetO promoter has no effect in wild type cells, the three msn5 mutant strains tested manifested an impaired cell growth. This result indicates that the msn5 mutant was sensitive.

![Figure 5](image)

Figure 5. Analysis of Cln2^{181-225}-GFP protein level and stability in the msn5 mutant strain. (a) The level of Cln2^{181-225}-GFP and the GFP control protein in the wild type (JCY1775 and JCY1845 respectively) and the msn5 mutant (JCY1778 and JCY1847 respectively) strains, was determined by western analysis with a specific anti-GFP antibody. Cdc28 protein level is shown as loading control. (b) Graph represents the amount of Cln2 protein (relative to Cdc28) determined by western analysis and CLN2 mRNA (relative to CDC28 mRNA) determined by qPCR, as well as the protein/mRNA ratio, in the msn5 mutant strain relative to the wild type strain. Values are mean and standard deviation derived from 11 replicates. Student’s t test was performed from Cln2^{181-225}-GFP to the GFP control. *: P < 0.0005. (c) Cycloheximide to a final concentration of 100 μg/mL was added to exponentially growing cultures of the wild type (JCY1845) and the msn5 mutant (JCY1847) strains expressing the Cln2^{181-225}-GFP protein. Protein level in crude cell extracts was determined at the indicated time by western analysis with a specific anti-GFP antibody. Cdc28 protein level is shown as loading control.
to Cln2 in a dose-dependent manner. Considering the nuclear accumulation of Cln2 protein in msn5 mutant cells [41], this observation strongly suggests that abnormal accumulation of high levels of Cln2 inside the nucleus could prove toxic for cells.

**Discussion**

Previous work in our group connected Msn5 with Cln2 [20,41]. Msn5 controls subcellular localization of Swi6, a component of the cell cycle transcription factor SBF responsible for the periodic expression of many genes in G1/S, among them the CLN2 gene. On the other hand, Msn5 mediates the nuclear export of Cln2 protein through a NES sequence located between amino acids 225 and 299. In this report, we described that Msn5 also affects Cln2 by an additional mechanism since Msn5 participates in regulating Cln2 protein levels. The msn5 mutant cells had a severe reduction in Cln2 levels compared to wild-type cells. Initially, the drop in Cln2 protein could be thought to be a consequence of Msn5 controlling CLN2 gene expression through the regulation of SBF. However, different observations raise this idea into question. Analysis in asynchronous cultures revealed a very small reduction of CLN2 mRNA compared to the drastic drop of Cln2 protein in msn5 cells. The slight effect in CLN2 mRNAs in asynchronous cultures is not a surprise since many other genes regulated by SBF are affected in their cell-cycle regulation but not in their global level of expression [52]. Consistent with this, mutants in the two components of SBF, Swi4 and Swi6, have no significant effect in Cln2 protein level in contrast to what is observed in the msn5 mutant. Furthermore, protein level of some chimeric cyclins (Cln21−200-Cln1201−546) and truncated version of Cln2 (Cln21−181) are not affected by Msn5 inactivation despite being expressed from the CLN2 promoter. Finally, when CLN2 is ectopically expressed, although the effect is softened, Msn5 absence still caused a reduction in Cln2 protein level. All these observations strongly indicate that the drastic reduction in Cln2 protein level is not caused by alterations in CLN2 gene transcription (although it cannot be ruled out that the slight effect in transcription could contribute to some extent); rather, Msn5 control of Cln2 protein level occurs mainly at posttranscriptional level.

Protein stability could be affected by subcellular localization [49,50]. Msn5 mediates nuclear export of Cln2 [41]. Therefore, it is conceivable, in principle, that nuclear accumulation of Cln2 in msn5 mutant cells could lead to greater protein instability and, consequently, to lower Cln2 cellular levels. However, several results contradict this possibility. The Cln1 and Cln2 cyclins contain a NLS activity in the N-terminal 1–224 region, and Cln2, but not Cln1, contains a Msn5-dependent NES between 225–299 amino acids. However, both cyclins are distributed through the whole cell and the presence of the Cln2-NES only alters the nuclear/cytoplasmic ratio modestly [41]. Because of this,

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**Figure 6.** Effect of Cln2 missregulation on cell growth. (a) 10-fold serial dilutions from exponentially growing cultures of the wild type (W303-1a), msn5 (JCY1018), GAL1:MSN5 (JCY313) and msn5::kanMX (JCY705) strains transformed with a centromeric plasmid expressing CLN2 under the control of the tetO2 promoter (ptetO:CLN2), were spotted onto YPD medium with or without 10 μg/mL doxycycline and incubated at 25º for 3 days.
we would not expect drastic differences in the subcellular localization of chimeric cyclins (all containing Cln2-NLS) based on the presence of the Cln2-NES sequence. Moreover, we observed that a variant of Cln2 (Cln2\textsuperscript{1\textsuperscript{7}T\textsuperscript{3}S}) that is stable and accumulates in the nucleus \cite{38,51} also had a reduced protein level in \textit{msn5} mutant cells. Importantly, the protein level of the truncated version Cln2\textsuperscript{1\textsuperscript{2}2\textsuperscript{5}}-Cln1\textsuperscript{2\textsuperscript{2}7\textsuperscript{-}546} or the fusion protein Cln2\textsuperscript{1\textsuperscript{8}1\textsuperscript{-}22\textsuperscript{5}}-GFP are affected by the inactivation of Msn5 despite lacking the NES region. All of these observations indicate that the control of Cln2 protein level by Msn5 is a new mechanism not related with the regulation of Cln2 subcellular localization. On the other hand, the drop in Cln2 protein levels caused by Msn5 inactivation also occurred in mutant strains in the SCF\textsuperscript{Grr1}, which is the main activity involved in Cln2 degradation. Finally, the shut-off experiments showed that the stability of Cln2 and Cln2\textsuperscript{1\textsuperscript{8}1\textsuperscript{-}22\textsuperscript{5}}-GFP was not significantly affected in the \textit{msn5} strain. All of these observations led us to conclude that the reduced amount of Cln2 in \textit{msn5} cells is not caused by an increased rate of degradation as a result of an altered localization. Instead, the results indicate that in the absence of Msn5 there are problems in Cln2 protein synthesis.

The analysis of polysome profiles indicated that inactivation of Msn5 apparently does not have a significant impact on global mRNA translation efficiency. This is consistent with the fact that Msn5 only affects protein level of specific proteins and it is in agreement with previously reported results in other growth conditions \cite{29}. Importantly, a clear shift in the distribution of specific mRNA to lighter polysomal and non-polysomal fractions is observed. This is the case for \textit{CLN2} and \textit{SWI5} mRNAs. It has to be noted that the distribution of \textit{ACT1} and \textit{HXK2} mRNA along the polysomal profile is also affected, although at a minor level, in \textit{msn5} cells compared to wild type cells. This could indicate that Msn5 affects a general aspect of mRNA translation but that specific properties of particular mRNAs (like \textit{CLN2} and \textit{SWI5}) impose a more restrictive need for Msn5 for their efficient translation. Importantly, the altered polysomal association confirms a role for Msn5 in proper translation of \textit{CLN2} and \textit{SWI5} mRNA, and could explain the reduced protein synthesis rate of Cln2 and Swi5 in the \textit{msn5} mutant cells.

We have tried to elucidate the molecular basis of Cln2 protein synthesis control by Msn5. Typically, mRNA is regulated by sequences present in the 5′- or 3′-untranslated regions. However, the drop in Cln2 levels in the \textit{msn5} mutant strain was observed in constructs lacking the 3′-UTR (C-terminal epitope-tagged proteins) or 5′-UTR (tetO constructs) and, some constructs containing the 5′-UTR are not affected by \textit{msn5} mutation. Thus, the effect of Msn5 on the Cln2 synthesis rate must be mediated by sequences in the coding region. In support of this, we show that the region coding for amino acids 181–225 was able to confer Msn5-regulated synthesis when fused to GFP. This is a very similar case to that of Swi5 \cite{19} and suggests that Msn5 could affect synthesis of specific proteins by a common mechanism. Susceptibility to Msn5 regulation could rely on specific properties of mRNAs like sequence, codon usage or secondary structures that could affect mRNA translation. However, the analysis of mRNA sequences coding the Cln2\textsuperscript{1\textsuperscript{8}1\textsuperscript{-}22\textsuperscript{5}} and Swi5\textsuperscript{1\textsuperscript{9}2\textsuperscript{5}-42\textsuperscript{2}} fragments that mediate the Msn5-dependent regulation were not conclusive.

Different scenarios can be envisaged on how Msn5 could control the synthesis of Cln2, Swi5 and other putative proteins. Msn5 has a well-defined role in exporting phosphorylated protein cargos. Therefore, it can be argued that the reduced levels of target proteins resulted from defects in Msn5-mediated nuclear export of specific factors required for proper \textit{CLN2} and \textit{SWI5} mRNA translation in the cytoplasm. These factors may act directly, increasing the translation efficiency of these mRNAs, or indirectly, inactivating or removing a specific RNA-binding translational repressor. In any case, the effect on translation would occur in a stage after mRNA export and its association to the translation machinery since the amount of \textit{CLN2} and \textit{SWI5} mRNAs associated to ribosomes is very similar in wild type and \textit{msn5} cells. On the other hand, the tRNAs perform an essential function to decode information in mRNAs into proteins, and Msn5 may affect tRNAs in different ways. Msn5 mediates the nuclear export of Maf1, a repressor of RNApol III transcription \cite{17}. However, \textit{msn5} mutation affected Cln2 protein levels even in the absence of Maf1 (Supplemental material Figure S3). In addition to a role in protein trafficking, Msn5 is
also involved in tRNA export to the cytoplasm, the tRNAs accumulating inside the nucleus in its absence [26,27,53]. It is possible that an altered tRNA nuclear-cytoplasmic trafficking could affect the cytosolic pools of tRNAs in such a way that compromises the translation efficiency of some specific messengers like CLN2 and SWI5 mRNAs. It has been suggested that there may be substrate specificity among the different tRNA exporters, so they could influence the cytoplasmic pools of particular tRNAs and hence possibly influence the proteome in different ways [28]. The tRNA<sup>Met</sup> and tRNA<sup>Ser</sup> are the most affected tRNAs by msn5 mutation among the analyzed cases [53]. It is interesting to point out that the Cln2<sup>181-225</sup> fragment mediating the Msn5-dependent regulation of Cln2 protein level contains a significant higher proportion of Met residues (6.8%) than the average proportion found in proteins (1.7%) and that the Swi5<sup>325-422</sup> fragment mediating the Msn5-dependent regulation of Swi5 protein level contains a higher proportion of Ser residues (18.8%) than the average proportion found in proteins (7.1%). Moreover, Msn5-mediated tRNA export involves the formation of a quaternary nuclear export complex consisting of Msn5, aminoacylated tRNA, RanGTP and the Tef1/2 protein [54], the yeast ortholog of translation elongation factor eEF1A that functions in delivering aa-tRNAs to the ribosome. In fact, msn5 mutant cells partially accumulate not only tRNA but also Tef1/2 in the nucleus [26]. Together with the reduced pools of tRNA, the reduced amount of cytosolic Tef1/2 could also contribute to defects in translation.

It is interesting to comment that we have observed a toxic effect of high Cln2 activity in the absence of Msn5, regardless of the reduced amount of the Cln2 protein. This could be because the toxicity of Cln2 was due to its nuclear accumulation. Consistent with this, the stable variant Cln2<sup>G1TS</sup> that accumulates inside the nucleus affects proper cell growth of a wild-type strain [36]. We can speculate that unscheduled Cln2 nuclear function is toxic and that Cln2 export by Msn5 helps to restrict the period during which this activity takes place. For example, Cln2 export by Msn5 could contribute, together with transcriptional shut-off and degradation of the protein, to the inactivation of Cln2 after the G1/S transition. If excess nuclear Cln2 is indeed toxic for cells, the concomitant protein level reduction accompanying its nuclear accumulation when exportin Msn5 is absent may be an evolutionary advantage.

How extended could the case of CLN2 and SWI5 be? The protein level of cyclin Cln1 and the transcription factors Swi6, Whi5 and Ace2 [19] are not affected by Msn5 inactivation, indicating that the effect of Msn5 is restricted to specific proteins. In fact, in a genomic approach only 12 genes were identified as having an altered translation profile in a msn5 mutant strain, all of them being translationally down-regulated compared to wild type cells [29]. It has to be pointed out that in this analysis the translation status was calculated as a ratio between the polysomal and non-polysomal fractions. In the light of our results with CLN2 and SWI5, this is a binary extreme situation in which genes with a shift in mRNA distribution inside the polysomal fractions (i.e. a shift to lighter polysomal fractions) will escape the analysis. Future work will help to identify more targets for Msn5-dependent regulation of protein synthesis and to clarify the molecular basis of the regulation.

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**Disclosure statement**

No potential conflict of interest was reported by the authors.

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**ORCID**

Paula Alepuz [http://orcid.org/0000-0003-1472-2373](http://orcid.org/0000-0003-1472-2373)

J. Carlos Igual [http://orcid.org/0000-0001-6910-5845](http://orcid.org/0000-0001-6910-5845)
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