Nuclear Import of Ho Endonuclease Utilizes Two Nuclear Localization Signals and Four Importins of the Ribosomal Import System*

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Activity of Ho, the yeast mating switch endonuclease, is restricted to a narrow time window of the cell cycle. Ho is unstable and despite being a nuclear protein is exported to the cytoplasm for proteasomal degradation. We report here the molecular basis for the highly efficient nuclear import of Ho and the relation between its short half-life and passage through the nucleus. The Ho nuclear import machinery is functionally redundant, being based on two bipartite nuclear localization signals, recognized by four importins of the ribosomal import system. Ho degradation is regulated by the DNA damage response (DDR), specifically the MEC1, RAD9, and CHK1 pathway (5). Despite being a nuclear protein, Ho must exit the nucleus to be degraded in the proteasomes. The DDR functions are important for Ho phosphorylation: phosphorylation of threonine 225 is crucial for Ho nuclear export and additional phosphorylations are required for recruitment of Ho for ubiquitination. Ho is ubiquitylated by the SCF (Skp1-Cdc53-F-box protein) E3 ubiquitin ligase complex, to which it is recruited by the F-box protein Ufo1 (6). In mec1 mutants Ho is stabilized and accumulates in the nucleus; conversely trapping Ho in the nucleus by deletion of its nuclear importin, Msn5, leads to stabilization of the protein (4). Ddi1 binds ubiquitylated Ho and is required for interaction of Ho with the proteasome; in its absence Ho is stabilized. The finding that Ho is not degraded within the nucleus, but in the cytoplasm, is further strengthened by the direct demonstration of accumulation of ubiquitylated Ho in the cytoplasm of Δddi1 mutants (7).

Ho nuclear import is very rapid and efficient. Ectopic expression of HO leads to rapid cleavage of MAT (8), and to a mating type switch at any phase of the cell cycle in both mother and daughter cells. This indicates that there is no impediment to its nuclear import (9). Macromolecules are conveyed through nuclear pore complexes in the nuclear envelope by soluble karyopherins. Karyopherins comprise two structurally related families, α- and β-karyopherins. These recognize specific nuclear localization sequence (NLS) peptide motifs in the cargo molecule: NLSs may comprise a short stretch of basic residues (classical NLSs) or two basic clusters 10–12 residues apart (bipartite NLS) (10). Cargos may be recognized by an adaptor protein, α-karyopherin/Srp1, which mediates their binding to the transport receptor, β-karyopherin/Kap95 (11). Additionally, a family of about 14 β-karyopherins bind an array of cargoes directly and also makes contacts with the nucleoporin subunits of the nuclear pore complex. Directionality of transport is determined by interaction with the GTPase Ran/Rayest Gsp1. RanGTP is at a high concentration in the nucleus due to the asymmetric distribution of the Ran regulators. The nuclear guanine nucleotide exchange factor, RanGef/Rayest Prp20, converts RanGDP to RanGTP, whereas the GTPase activating protein, RanGAP/Rayest Rna1, is localized in the cytoplasm and catalyzes the hydrolysis of RanGTP. Importin-cargo complexes assemble in the cytoplasm and after translocation into the nucleus they dissociate upon binding of RanGTP to the importin (12).

To investigate how the efficient nuclear import that supports the unique biological function of Ho is achieved we located and analyzed its nuclear localization signal: GFP, green fluorescent protein; IP, immunoprecipitation; PI, protein introns.
import signals, and identified the nuclear importins that mediate its import.

Proteasomes and other components of the SCF are found throughout the cell and it is not clear why Ho cannot be degraded within the nucleus. One possibility is that it is not fully marked as a degradation substrate in the nucleus and needs to acquire additional phosphorylations by a cytoplasmic protein kinase. In this respect Ho may resemble the kinase inhibitor, Sic1, that requires multiple phosphorylations to reach a critical threshold for stable binding to its F-box receptor, Cdc4 (13). We therefore examined whether Ho acquires all the post-translational modifications that target it for degradation within the nucleus, or whether it can be marked as a proteasome substrate by a cytoplasmic protein kinase(s).

Homothallism in budding yeast arose by stepwise replications of the MAT locus leading to the establishment of three cassettes of mating type information. This was followed by acquisition of Ho by “domestication” of a fungal VMA1 protein-splicing intein element (14–17). Intein protein domains are encoded by selfish genetic elements present within the protein coding regions of diverse genes. Inteins include a protein-splicing domain that catalyzes their extraction from the host polypeptide (18, 19). They typically also have an endonuclease domain that mediates copying of the intein gene into an orthologous unoccupied intein-integration point (20). Homology modeling of Ho based on the crystal structure of S. cerevisiae PI-SceI, a typical VMA1 intein intein (21), indicates that Ho residues from the N-terminal domain (1–185) together with residues downstream of the endonuclease domain (428–465) form a protein splicing domain (with a small DNA recognition region between residues 93 and 157). The intervening region forms a LAGLIDADG-type homing endonuclease domain including all expected catalytic active site residues (16). In addition, Ho has a unique C-terminal 120-residue zinc finger domain (22) that is critical for mating type switching and may contribute to specificity of cognate site recognition (16, 23).

Despite their common ancestry Ho and PI-SceI have very different life strategies. PI-SceI is produced as part of the vacuolar ATPase protein precursor at all stages of the cell cycle, but is active only in meiotic cells (20). In contrast to Ho nuclear import that can occur in all cell types at all stages of the cell cycle (above), activity of the PI-SceI intein is regulated by its nuclear import. The spliced out intein is retained in the cytoplasm during vegetative growth and imported into the nucleus in premeiotic cells when the TOR (target of rapamycin) pathway is inactivated in response to nutrient depletion. Nuclear import of PI-SceI is mediated by the importin Srp1 (24). In diploids heterozygous for the intein, cleavage of the PI-SceI cognate site of the inteinless allele is followed by double strand break repair during which a copy of the endonuclease open reading frame is inserted at the cleaved site (homing). Homing disrupts the cognate site preventing further cleavage by the endonuclease (25). This is different from the repair of the MAT allele that regenerates the Ho cognate site during the mating type switch. To determine when the Ho nuclear import mechanism arose we therefore compared the protein sequences of Ho and PI-SceI from different Saccharomyces yeasts.

**MATERIALS AND METHODS**

**Yeast Strains** are described in Table 1.

**Plasmid Construction**

For two-hybrid clones, Gal4 activating domain fusions of Ho and subclones were constructed by R. Buchnik using restriction fragments (56). The PseI-DNA binding domain fusion in pOBW was obtained from Peter Uetz (26). Construction of pYES2-HO and HO and HO CS08A, CS11A in pH315-GFP are described in Ref. 16. GFP-NLS1 was made by amplifying a fragment centered around the NLS1 of HO with primers NLS1Not1F (GAGAAATGCGGCCGC-CTGGGGTCTCTACCTTACG) and NLS1Sac1R (GCAACTCTTTATGAGCTCGCCGTACATAATTTCGG). The PCR product was digested with NotI and SacI and introduced in-frame to GFP in pH315-GFP. HO-NLS2 was constructed by overlap primer extension using primer pairs: HoSpe1F and ΔNLS1aR (CGTATAGATCTACCCAGACAGT) and HoSac1R (GCAATCGATGAATTGTAGCTACACAGCCGG). Deletion of NLS2 was employed primer pair ΔNLS2F (GAAAGAATTGAGTGTAGAAGAC) and ΔNLS2R (CCAATCCTGTCTCCCCACACACG); mutation of NLS2 residues, HoR491A, K492A employed primer pair NLS2RF (CCCGCGGCCGCAATCAAGTATTITTTAAGAGTGG) and NLS2RK (CCCGCGGCCGGAACAGGATTGGGAGAGAG) and each karyopherin mutant we analyzed cells according to the following procedure: three regions of interest corresponding to nucleus, cytoplasm, and adjacent cell background were chosen and the pixel value for each region (N, nucleus; C, cytoplasm; B, background) was used to determine the nuclear/cytoplasmic ratio for each cell. A ratio for each cell sampled was calculated using (N-B)/(C-B). Based on this we calculated the median ratio for each cell population. Table 2 shows the median values of the nuclear/cytoplasmic ratio for Ho, NLS1, and NLS2 in wild type and each karyopherin mutant we analyzed cells according to the following procedure: three regions of interest corresponding to nucleus, cytoplasm, and adjacent cell background were chosen and the pixel value for each region (N, nucleus; C, cytoplasm; B, background) was used to determine the nuclear/cytoplasmic ratio for each cell. A ratio for each cell sampled was calculated using (N-B)/(C-B). Based on this we calculated the median ratio for each cell population. Table 2 shows the median values of the nuclear/cytoplasmic ratio for Ho, NLS1, and NLS2 in wild type and mutant cells. For temperature-shift experiments, cultures after overnight induction were shifted to 37 °C with continuous induction for a further 2 h. Co-immunoprecipitation-Immunoblotting Yeast cells were grown overnight to late logarithmic phase (A460 = 0.8) in 50 ml of the appropriate inductive synthetic minimal medium with 2% galactose for expression of GAL–HO. The cells were harvested by centrifugation, washed in 50 ml of TE, and resuspended in 400 μl of co-IP buffer (0.1% Nonidet P-40, 250 mM NaCl, 5 mM EDTA, 50 mM Tris–Cl, pH 7.5, 1:25 of Boehringer Protease Inhibitor mixture). 0.5–0.6-g Glass beads were added and cells were broken with a glass beater (Biospec Products) using five 1-min cycles at 4 °C. The lysate was clarified by centrifugation at 13,000 × g for 15 min at 4 °C and protein concentration was measured with the Bio-Rad protein reagent. 5–15 mg of protein lysate were used for IP with the appropriate antibodies in co-IP buffer at 4 °C for 1–2 h with mild shaking. 30 μl of 50% Protein A-Sepharose (Amersham Biosciences) were added to each sample and incubation was continued under the same conditions for 0.5–1 h. The samples were washed 6 times with co-IP buffer with 1% Triton. The pellet was resuspended in 30 μl of 2× Laemmli sample buffer, boiled for 5 min, and the samples were loaded onto SDS–polyacrylamide gels. The proteins were transferred to a nitrocellulose membrane and immunoblotted.
Ho Endonuclease Nuclear Import System

**TABLE 1**

Yeast strains with karyopherin mutations

<table>
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<th>Strain</th>
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<td>Y00000-BY4714</td>
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</table>

10 min, and electrophoresed on a 10% polyacrylamide SDS gel (PAGE) with protein size standards. The gel was transferred to nitrocellulose membrane (Protran BA 85, Schleicher & Schuell) and Western blotting was performed with the appropriate antisera. 1 mg of crude protein was used to determine input by IP-Western blot and 30 μg was taken for direct Western blotting.

For determination of Ho half-life cells were induced as above. For promoter shutoff 3% glucose was added at the zero time point, together with cycloheximide to a final concentration of 10 μM to inhibit translation. Aliquots were collected at the times indicated for anti-GFP Western blotting; goat anti-rabbit and goat anti-mouse at 1:1000 for Western blotting; goat anti-rabbit and goat anti-mouse at 1:1000 for immunoprecipitation and 1:25 of BioRad Western blotting kit. Western blotting was performed with the appropriate antisera. 1 mg of crude protein was used to determine input by IP-Western blot and 30 μg was taken for direct Western blotting.

For Pulldown Experiments—GST fusion proteins were produced in logarithmic bacterial cultures by overnight incubation with 0.1 mM isopropyl-1-thio-β-D-galactopyranoside at 20 °C. The cells were pelleted by centrifugation at 3,500 × g for 10 min, washed in 0.1× phosphate-buffered saline, and resuspended in 10% GTP-bound Gsp1, with 0.5% Nonidet P-40, and 1:25 of BioRad Western blotting kit. Western blotting was performed with the appropriate antisera. 1 mg of crude protein was used to determine input by IP-Western blot and 30 μg was taken for direct Western blotting.

For Western blotting—GST fusion proteins were produced in logarithmic bacterial cultures by overnight incubation with 0.1 mM isopropyl-1-thio-β-D-galactopyranoside at 20 °C. The cells were pelleted by centrifugation at 3,500 × g for 10 min, washed in 0.1× phosphate-buffered saline, and resuspended in 10% GTP-bound Gsp1, with 0.5% Nonidet P-40, and 1:25 of BioRad Western blotting kit. Western blotting was performed with the appropriate antisera. 1 mg of crude protein was used to determine input by IP-Western blot and 30 μg was taken for direct Western blotting.

6His-Pse1p and 6His-Gsp1p—6His-Pse1p and 6His-Gsp1p were purified with nickel-nitrotriacetic-acid agarose (Qiagen). Pse1p was further purified by Mono Q (Amersham Biosciences) chromatography. The GTP-bound form of 6His-Gsp1p was separated on a Mono Q column (Amersham Biosciences) and the GTP/GDP contents were determined by high pressure liquid chromatography analysis. The Gsp1p-GTP preparation contained 50% GTP-bound protein.

Antibodies were purchased from Santa Cruz Biotechnology and used at the following dilutions: anti-GFP, 1:200 for immunoprecipitation and 1:1000 for Western blotting; anti-LacZ and anti-His, 1:6000 for immunoprecipitation and 1:1000 for Western blotting; anti-GST, 1:5000 for Western blotting; goat anti-rabbit and goat anti-mouse at 1:1000 for Western blotting. Detection was by ECL using an Amersham Biosciences ECL Western blotting kit.

**Phylogenetic Analysis and Sequence Data**

Multiple sequence alignments were calculated by the BlockMaker (28) and dialign2 (29) programs. Sequence logos, showing the multiple alignment conservation, were calculated according to Ref. 28. Phylogenetic trees were calculated with the PHYLP10 program (30). A consensus tree was made from a neighbor joining trees calculated from protein distance matrices of 1000 bootstrapped sequence versions, using default program parameters. NCBI database accession of the Ho sequences analyzed are Saccharomyces cerevisiae 1431383, Saccharomyces pastorianus HO-Sc 7549713, S. pastorianus HO-Lg 17756436, Saccharomyces bayanus HO-17756438, S. bayanus HO-22953602, Saccharomyces mikatae 9364028, Saccharomyces paradoxus 22830763, Saccharomyces castellii 30988068, Saccharomyces krudiatzevi 30995262, Klyveromyces delphensis 4255759, Candida glabrata 42557533, and S. castellii HO-like 30988030. The full sequence of Zygosaccharomyces rouxii Ho was kindly provided by Ken Wolfe, University of Dublin. Sequences of the fungal VMA1 inteins are available on the NEB intein database (31) including those we identified and deposited there.

**RESULTS**

Identification of Karyopherins and NLSs Involved in Nuclear Import of Ho—The subcellular location of Ho was directly visualized by fusing it to GFP. We screened mutants in the karyopherins Srp1, Ran1/Kap95, Kap104, Kap114, Kap120, Pse1/Kap121, Pdr6/Kap122, Yrb4/Kap123, Nmd5, Sxm1, Cse1, Xpo1, and Msn5; in Prp20 (RanGEF) and Rna1 (RanGAP); and in a mutant of the Ran-binding protein Yrb1 (Table 1). Conditional mutants were used to test essential genes. We found Ho in the nucleus of wild type cells (Fig. 1A) and in all the karyopherin mutants screened above. In ma1 mutants nuclear import of GFP-Ho was severely impaired at the restrictive temperature indicating that Ho
import depends on Ran (Fig. 1B). In parallel we assayed a series of isogenic wild type and yeast mutants for their ability to support a mating type switch. **HO** was induced from the **GAL** promoter in *pYES2-HO* and the cells were mated with a tester of the opposite mating type to test for their mating efficiency, and with a tester of their original mating type to test for a mating type switch. *rna1* mutants did not switch, however, all the karyopherin mutants switched mating type, indicating that nuclear import of Ho was not abrogated by deletion of any single karyopherin. These results imply functional redundancy between the karyopherins that mediate Ho nuclear import and/or the existence of more than one NLS on Ho imported by different karyopherins.

The Ho sequence shows a putative bipartite NLS between residues 280 and 302. This region is highly conserved in Ho from different species, in particular the crucial basic residues (Fig. 2A). In the past we reported that mutation of Arg-286 abrogates mating type switching in yeast (16). Arg-286 aligns with PI-Scel Asn-281, a residue that contacts the DNA in the cocrystal (32). We fused Ho residues 248–311, centered around this putative NLS (NLS1), to GFP and found that this Ho fragment led to nuclear accumulation of GFP. However, when we deleted NLS1 from Ho this did not affect Ho nuclear import. Further inspection of the primary sequence indicated another very well conserved putative bipartite NLS in the zinc finger domain between residues 480 and 492 (Fig. 2). Indeed fusion of residues 441–596 (NLS2) to GFP led to accumulation of GFP in the nucleus (Fig. 2B). This supports an earlier observation that mutation of cysteine residues in the zinc finger domain of Ho besides leading to loss of Ho activity greatly reduces nuclear accumulation of GFP-Ho (16). We deleted both NLS1 and NLS2 regions (residues 286–302 and 480–496) and found that the resulting Ho protein was not imported at all into the nucleus in wild type cells. Similarly the above NLS1 deletion combined with NLS2 point mutations of R491A,K492A also abrogated nuclear import, indicating that it is the basic residues of the NLS that are important for binding the karyopherin (Fig. 2B). These experiments indicate that we have correctly delineated the functional NLSs of Ho.

Having identified two NLSs in Ho we rescreened the karyopherin mutants for their ability to import GFP-Ho, and each GFP-NLS1 and GFP-NLS2, separately. GFP-NLS1 nuclear import was reduced in *pse1* mutants at the restrictive temperature as evidenced by the ratio of nuclear to cytoplasmic GFP signal; GFP-NLS1 was also predominantly cytoplasmic in *kap123* mutants. In addition in *sxm1* mutants GFP-NLS1 showed a relatively high cytoplasmic signal compared with GFP-NLS2. Nuclear import of GFP-Ho and GFP-NLS2 was not affected in *pse1, kap123, or sxm1* mutants; the only mutant in which GFP-NLS2 was predominantly cytoplasmic was *kap120* (Fig. 3 and Table 2). Although they could support a mating type switch, microscopic observation of *nmn5* mutants revealed a general import defect that severely reduced nuclear import of GFP-Ho, GFP-NLS1, and GFP-NLS2 (not shown) and were not analyzed further.

**Interaction of Karyopherins with Ho**—To further support the GFP-Ho nuclear import data we tested for an interaction between Ho and Pse1 in the two-hybrid protein interaction trap. Full-length Ho, and a number of subclones (Fig. 4A) were fused to the Gal4 activation domain in plasmid pGAD424 (Clontech) and co-transformed into reporter strains (33) together with plasmids expressing *PSE1* fused to the Gal4 DNA binding domain (pOBD) (26). Transformants in which Ho interacted with Pse1 activated the **GAL1-HIS3** and **GAL2-ADE2** reporter genes and grew on the selective plates. Ho, residues 1–566, and Ho subclone 186–441 interacted with Pse1. However, subclones encompassing residues 1–297, 299–565, and 441–566 did not interact. This analysis confirmed
the bipartite NLS1 sequence we identified between residues 280 and 302. This NLS would be intact in the interacting subclones, but split in the bipartite NLS1 sequence we identified between residues 280 and 302. This NLS would be intact in the interacting subclones, but split in

In contrast to \( \kappa \)-karyopherins that form a heterodimer with a nucleoporin-binding \( \beta \)-karyopherin, Kap95, subunit, most \( \beta \)-karyopherins bind both their cargo and the nucleoporin subunits. However, it was recently reported that the nucleolar protein Sof1, a cargo of Psel, does not bind Psel directly. Instead Sof1 employs a Nop1 bridge that mediates its nuclear import via Psel by a “piggyback” mechanism (34). We therefore tested whether Ho interacts directly with Psel using recombinant proteins expressed in \textit{Escherichia coli}. Full-length Ho (residues 1–586) and a C-terminal truncation of Ho that extends from residues 1 to 451 were fused to GST and isolated on glutathione-Sepha-

<table>
<thead>
<tr>
<th>Strain</th>
<th>NLS1</th>
<th>NLS2</th>
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<tbody>
<tr>
<td>WT 37 °C</td>
<td>2.9 ± 0.7 (23)</td>
<td>2.2 ± 0.4 (36)</td>
</tr>
<tr>
<td>pse1 37 °C</td>
<td>2.4 ± 0.5 (38)</td>
<td>1.8 ± 0.3 (21)</td>
</tr>
<tr>
<td>kap123</td>
<td>2.0 ± 0.4 (38)</td>
<td>1.0 ± 0.1 (20)</td>
</tr>
<tr>
<td>sxm1</td>
<td>1.9 ± 0.3 (24)</td>
<td>1.4 ± 0.2 (15)</td>
</tr>
<tr>
<td>kap120</td>
<td>1.6 ± 0.1 (20)</td>
<td>1.8 ± 0.3 (18)</td>
</tr>
</tbody>
</table>

TABLE 2
Nuclear/cytoplasmic ratio (\( R \equiv E \)) for Ho, NLS1, and NLS2 in wild type and karyopherin mutant cells

\[ R = \text{median} \{ R_i \} \]
\[ E = \text{median} \{ R_i - R_i^* \} \]

\( R \) is the median of multiple measurements \( R_i \) on the same type of cells; \( E = \text{median} \{ R_i - R_i^* \} \). The number of cells analyzed appears in parentheses.

the bipartite NLS1 sequence we identified between residues 280 and 302. This NLS would be intact in the interacting subclones, but split in clones 1–297 and 299–565. NLS2 is between residues 480 and 496 and our observations on nuclear import of GFP-NLS2 indicated that its import is indeed not affected in \( pse1 \) mutants (Fig. 3).
The Ho NLSs Arise Early in Its Domestication—Ho arose by domestication of a fungal VMA1-type intein, exemplified by the well studied PI-SceI (14–17). The Ho protein in all species of Saccharomyces is extremely highly conserved. This Ho protein has retained the intein protein splicing, homing endonuclease, and DNA binding domains, and acquired a unique zinc finger region. NLS1 is in the endonuclease domain, whereas NLS2 is in the zinc finger domain. PI-SceI is imported into the nucleus by the α-importin, Srp1 (24). Here we found that at the restrictive temperature for Srp1–31, Ho is nuclear, indicating that Srp1 is not required for nuclear import of Ho. A control bipartite NLS derived from the cNLS of SV40 (35) remains in the cytoplasm indicating that Srp1 activity is indeed lost in the mutant (Fig. 5). We compared the Ho and VMA1-type inteins to examine the origin of the two Ho NLSs. The region corresponding to Ho NLS1 (PI-SceI–275–297) does not have any NLS features and is not conserved in different yeast VMA1 inteins (results not shown). In structures of PI-SceI this region forms a surface patch and its N-terminal half becomes ordered upon co-crystallization with the endonuclease DNA target (32). Inspection of other fungal inteins (such as those of the PRP8 proteins (36)) did not identify any conserved NLS-like regions in their endonuclease domains. The additional NLSs of Ho, NLS2, is in the zinc finger domain of Ho, the domain unique to Ho and not present in inteins. We compared the common sequence regions of Ho and the VMA1-type inteins (all domains apart from the zinc fingers) and found that the Ho proteins and fungal VMA1-type inteins form two separate groups (Fig. 6). The Ho protein closest to the divergence point is that of Z. rouxii, a species that did not undergo genome duplication but is closer to species that have acquired Ho than to those that have not (37). This phylogenetic analysis shows that the efficient nuclear import of Ho is based on two NLSs acquired after divergence from its intein ancestor.

Disruption of Ho Nuclear Import Stabilizes the Protein—The sensitivity of Ho activity to mutation in the regions of NLS1 and NLS2 (16) precluded the use of mating type switching as a functional bioassay for mutational analysis of the putative NLSs. We therefore devised an alternative functional assay based on our findings that Ho is marked as a proteasome substrate within the nucleus by functions of the DDR and is then exported to the cytoplasm to be degraded (3, 4, 7). These experiments while showing that Ho nuclear export is essential for its degradation, did not address the question of whether Ho acquires additional phosphorylations in the cytoplasm. Lack of nuclear import of Ho could perhaps direct it for rapid degradation due to its mislocalization, or alternatively if the crucial modifications that mark Ho as a proteasome substrate occur exclusively within the nucleus this could extend its half-life. Here delineation of the Ho NLSs and the karyopherins that mediate its nuclear import allow us to address this question. We previously reported that mutation of two upstream cysteine residues of each of the putative zinc fingers of Ho abolished mating type switching. These Ho mutants showed reduced nuclear import, although in Δmsn5 mutants it
Ho Endonuclease Nuclear Import System

FIGURE 7. A, half-life of Ho (GFP-Ho) and Ho(C508A,C511A), GFP-Ho(CACA), in a kap123 mutant determined by microscopy after translation shutoff at the zero time point. The GFP-Ho signal is nuclear at the zero time point, but 10 min after addition of cycloheximide there is no longer a nuclear GFP-Ho signal and by 20 min there is no GFP signal in these cells. Cells expressing Ho(C508A,C511A) have a strong GFP cytoplasmic signal and during the course of the experiment, a large part of the cytoplasmic signal accumulates in discrete spots, probably aggresomes. B, top panel: half-life of Ho and Ho(C508A,C511A) in a kap123 mutant determined by translation shutoff and immunoprecipitation of aliquots at the designated time points followed by Western blotting with anti-GFP antisera. Lower panel: half-life of GFP-Ho/ho, GFP-Ho/ΔNL51, ΔNL52 (Δ1,Δ2), and GFP-Ho/ΔNL51/ΔNL52(R491A,K492A) (Δ1,Δ2) in wild type cells determined as described above.

was possible to accumulate some GFP-Ho within the nucleus (16). Based on our results above this nuclear import can now be attributed to the activity of NLS1 as the zinc finger mutations probably affect NLS2 (residues 480–492)-mediated nuclear import. We therefore expressed the zinc finger mutant, Ho(C508A,C511A), in a Δkap123 mutant that recognizes NLS1, to investigate the half-life of Ho misdirected to the cytoplasm. We produced GFP-Ho and GFP-Ho(C508A,C511A) by growing the cells in galactose for 18 h and added 10 mM cycloheximide to inhibit further translation of Ho and Ho(C508A,C511A) in a kap123 mutant determined by microscopy after translation shutoff and immunoprecipitation of aliquots at the designated time points followed by Western blotting with anti-GFP antisera. Lower panel: half-life of GFP-Ho/ho, GFP-Ho/ΔNL51, ΔNL52 (Δ1,Δ2), and GFP-Ho/ΔNL51/ΔNL52(R491A,K492A) (Δ1,Δ2) in wild type cells determined as described above.

DISCUSSION

Our experiments demonstrate that efficient nuclear import of Ho is achieved through two separate bipartite NLSs and involves at least four different importins. Functional redundancy is a hallmark of yeast karyopherins as only four β-karyopherins are encoded by essential genes (11). The nuclear import observations of GFP fusion proteins suggest that NLS1 in the endonuclease domain of Ho is recognized chiefly by Kap123. However, the two-hybrid and in vitro binding experiments show a strong interaction with PseI too. In fact the two-hybrid Ho subclones that interacted with PseI enabled precise delineation of NLS1. Kap123 and Psel are the major importins of ribosomal proteins (38, 39). Kap123 is not essential for growth due to its functional redundancy with Psel and Sxm1, each of which complements growth of kap123 mutants (38–40). Kap123 and Psel are also crucial for export of mRNA; here too the synthetic lethality of kap123, pse1 mutants can be suppressed by overexpression of SXM1 (41). We find the same functional redundancy between Kap123, Psel, and Sxm1 in Ho import via NLS1. Ribosomal biosynthesis imposes a considerable metabolic burden on actively growing yeast cells (42) and use of the functionally redundant highly abundant ribosomal importins by Ho has doubtless contributed to the evolution of Ho activity in yeast. Proteins that function in the same pathway are often imported by the same karyopherin whose abundance may fluctuate during the cell cycle. In this context it is interesting to find that a major importin of Ho, Psel, also mediates import of Ste12 required for yeast mating (43). Identification of the Ho bipartite NLS1 as the signal recognized by Psel is the first characterization of a NLS recognized by this karyopherin. In a previous report (16) we showed that mutation of Arg-286 to alanine abrogates a mating type switch. This residue was chosen for mutagenesis as our structural model of Ho indicated that the parallel residue, Asn-281, is essential for P-Scel activity and indeed contacts the DNA. Arg-286 is part of NLS1 of Ho and our present analysis in which we show that in the absence of NLS1, Ho nuclear import is supported by NLS2, confirms that it is the effect of the point mutation R286A on Ho activity and not on its nuclear import that leads to abrogation of mating type switching. Given the sensitivity of Ho activity to mutation we used native Ho in all our mating type tests of karyopherin mutants. Psel and Kap123 also import the histone acetyltransferase complex, SAS-1, however, the consensus NLS signal derived from these experiments and a bioinformatics analysis of other putative cargoes of this importin (44) does not resemble NLS1 of Ho.

The zinc finger region of Ho, unique among LAGLIDADG endonucleases, provides a second bipartite NLS recognized by Kap120. Interestingly Kap120 too has been identified as a karyopherin of ribosomal proteins (45). Here our mutational analysis shows that Kap120 binds a bipartite NLS. We previously showed that mutations in the zinc finger domain of Ho lead to loss of mating type switching. This is due to the effect of the mutations on Ho structure as using a Δmsn5 mutant in which Ho nuclear export is inhibited, we showed that Ho with mutations in this region of the protein does enter the nucleus (16). Based on the results presented here we conclude that this nuclear import is mediated by NLS1, as in kap120 mutants it is only nuclear import of GFP-NLS2 that is affected, both GFP-Ho and GFP-NLS1 are imported into the nucleus. Therefore acquisition of a unique zinc finger domain with NLS2 further increased the efficiency of Ho import. Use of multiple importins each present in copious amounts (46) can explain the high efficiency of Ho import.

The identification of the Ho nuclear import machinery enabled us to accumulates in the cytoplasm it is protected from the ubiquitylation machinery that mediates its rapid degradation.

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revisit the role of the DDR phosphorylations in targeting Ho for degradation in the proteasome. We find that when its nuclear import is abrogated Ho is stabilized. We hypothesize that in the cytoplasm Ho is not recognized and marked as a degradation substrate. In this respect Ho turnover may resemble the GC17–GC26-specific degradation of the cyclin-dependent kinase inhibitor, p27Kip1, which occurs in the cytoplasm. At this stage of the cell cycle p27Kip1 is ubiquitylated by the KPC (Kip1 ubiquitylation-promoting complex) complex. This E3 complex does not recognize p27Kip1 as a ubiquitylation substrate unless it is first imported into the nucleus (47).

Phylogenetic distribution and sequence conservation of Ho and yeast VMA1 inteins coincide with their biological functions and evolutionary history. Ho mediation of mating type switching is a relatively recent innovation of S. sensu stricto and close species, most likely occurring 50–70 million years ago (17, 48). Ho proteins are known only in this group and cluster together in phylogenetic analyses (Fig. 6). The closest proteins to Ho are the fungal VMA1 inteins (49). Besides S. sensu stricto these are found in other Saccharomycete yeasts such as Kluyveromyces lactis (50) and Candida tropicalis (51), which split from the S. sensu stricto group before the latter acquired the HO gene (17, 48). This is further indication regarding the origin of Ho from an intestine (15). VMA1 yeast inteins are also more diverse then Ho proteins (Ref. 48 and Fig. 6) indicating their earlier origin, and probably also reflecting the reduced evolutionary selection they face. Intein genes must efficiently protein-splice to survive (18, 19); their homing capability (mediated by their endonuclease domains) is only necessary for long-term survival and can be inactivated or lost (5, 52–54).

S. castellii includes in addition to its typical HO gene a gene coding for a Ho-like protein. This gene has all the Ho protein sequence features: intein protein splicing, LAGLIDADG homing endonuclease, DNA recognition region, and zinc finger domains. It is most similar to the K. lactis HO pseudogene (5). Despite its Ho-specific features that include the unique zinc finger domain (whose sequence cannot be used in the phylogenetic analysis with inteins), the S. castellii Ho-like protein does not cluster with the typical Ho proteins, nor with the VMA1 inteins (Fig. 6). This protein and the K. lactis HO-like pseudogene include sequence regions corresponding to the two Ho NLSs, but lacks the basic residues that define these NLSs (not shown). These regions are also not very similar between these two published Ho-like genes. An HO-like gene was recently reported in K. thermotolerans and based on this gene and the K. lactis HO-like pseudogene an earlier date for appearance of HO prior to the branching of K. lactis was proposed (5). Further examination of this hypothesis awaits public release of the sequence of the K. thermotolerans HO-like gene. Our phylogenetic analysis supports the acquisition of HO closer to the appearance of Z. rouxii (17, 48). This species branched off before the Saccharomyces genome duplication (37) has a typical HO gene with both NLS1 and NLS2, and confidently appears as the earliest branch of the known Ho sequences (Fig. 6).

The totally different strategies adopted by Ho and PI-SceI to regulate their activity since diverging from their common ancestor can now be explained at the molecular level. Ho nuclear import is very efficient and involves two NLSs and at least four abundant importins. Both Ho NLSs are conserved in HO of all Saccharomyces species, but have no apparent parallel in yeast VMA1 inteins. This suggests that adoption of a new nuclear import strategy occurred very early in the evolution of Ho. The divergence in NLS is supported by the different karyopherins employed by each endonuclease. Whereas PI-SceI nuclear import is mediated by Srpl, Ho is nuclear in srpl mutants and needs Pse1, Kap123, and Sxm1 for its nuclear import via NLS1, and Kap120 for import via NLS2. The highly efficient Ho nuclear import is thus a successful strategy for achieving a high rate of mating type switching. Diploidization of yeast cells that have switched mating type enables sporulation in conditions of nutrient deprivation thus increasing fitness of HO yeast. HO cells that accumulate stable protein within the nucleus suffer from perturbation of the cell cycle (4). We find that the highly efficient nuclear import of Ho has evolved in parallel with equally efficient DDR-mediated Ho nuclear export and ubiquitylation by SCF(Skp1) that occurs exclusively in the cytoplasm (3, 4, 6, 7). This promotes degradation of Ho in the proteasome and provides a mechanism for maintaining genome stability. Thus the combination of highly efficient nuclear import together with fortuitous recruitment of functions of the DDR that target Ho to the proteasome for rapid destruction has enabled the establishment of homothallic mating type interconversion.

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REFERENCES

Ho Endonuclease Nuclear Import System