The inner nuclear membrane protein Src1 associates with subtelomeric genes and alters their regulated gene expression

Stefanie E. Grund,1 Tamás Fischer,1 Ghislain G. Cabal,2 Oreto Antúnez,3,4 José E. Pérez-Ortín,3,4 and Ed Hurt1

1Biochemie-Zentrum der Universität Heidelberg, D-69120 Heidelberg, Germany
2Unite de Biologie Cellulaire du Noyau, Institut Pasteur, 75724 Paris, Cedex 15, France
3Departamento de Bioquímica y Biologia Molecular and 4Sección de Chips de DNA Servei de Soutport a la Investigació Experimental, Universitat de Valéncia, E-46100 Valéncia, Spain

Introduction

Among the numerous steps of gene expression, formation, and maturation of messenger RNP particles (mRNP) are crucial steps before transcripts can be exported from the nucleus and translated in the cytoplasm. Studies over the past years have revealed that these various steps, including chromatin organization, gene expression, and nuclear envelope biogenesis. In this study, we identify a robust genetic interaction between transcription export (TREX) factors and yeast Src1, an integral inner nuclear membrane protein that is homologous to vertebrate LEM2. DNA macroarray analysis revealed that the expression of the phosphate-regulated genes PHO11, PHO12, and PHO84 is up-regulated in src1Δ cells. Notably, these PHO genes are located in subtelomeric regions of chromatin and exhibit a perinuclear location in vivo. Src1 spans the nuclear membrane twice and exposes its N and C domains with putative DNA-binding motifs to the nucleoplasm. Genome-wide chromatin immunoprecipitation-on-chip analyses indicated that Src1 is highly enriched at telomeres and subtelomeric regions of the yeast chromosomes. Our data show that the inner nuclear membrane protein Src1 functions at the interface between subtelomeric gene expression and TREX-dependent messenger RNA export through the nuclear pore complexes.
and genes located in the subtelomeric regions (e.g., nuclear periphery. Moreover, the regulation of expression of chromatin immunoprecipitation (ChIP) – on-chip analysis, Src1 binding N and C domains to the nucleoplasm. As shown by nuclear membrane twice, thereby exposing the putative DNA-

that Src1 is an integral membrane protein that spans the inner

2005 ; Wagner and Krohne, 2007 ). Our analyses further revealed for chromatin at the nuclear periphery and modulating higher expression and mRNA export (for reviews see Akhtar and Gasser, 2007 ; Köhler and Hurt, 2007 ).

Historically, the nuclear periphery was known as a zone that harbors silenced regions of the genome and thus was believed to be an area of transcription repression. However, recent studies have revealed that genes can be recruited to the nuclear periphery upon their transcriptional activation (for reviews see Akhtar and Gasser, 2007 ; Köhler and Hurt, 2007 ). Thus, activating and repressing chromatin environments coexist but appear to be spatially partitioned. Whereas the heterochromatin, like telomeres and the mating-type locus, line the nuclear envelope, the active chromatin domains are in the vicinity of the NPCs (Taddei et al., 2004 ; for review see Akhtar and Gasser, 2007 ). Gene recruitment to the nuclear periphery involves components of the nuclear basket and associated factors implicated in transcription and mRNA export, including SAGA and TREX-2 factors as well as Mex67. Regarding the mechanism, gene gating requires the nascent transcript or posttranscriptional events. However, other studies suggest that gene gating can also be independent of transcription and is then mediated by direct interaction of the gene with components of the nuclear periphery (for reviews see Akhtar and Gasser, 2007 ; Köhler and Hurt, 2007 ). In general, gene recruitment to the periphery could allow access to a favorable environment, including chromatin remodeling, transcription, and export machineries, thereby optimizing gene expression and mRNA export (for reviews see Akhtar and Gasser, 2007 ; Köhler and Hurt, 2007 ).

In this study, we found Src1 (also called Heh1; King et al., 2006 ) in genetic screens using mutants of the THO–TREX and TREX-2 complexes. Src1 exhibits a domain organization similar to higher eukaryotic LEM2 and MAN1, which are integral inner nuclear membrane proteins that consist of an N-terminal LEM domain, two transmembrane-spanning sequences, and a Man1-Src1 C-terminal (MSC) domain (Mans et al., 2004 ; Brachner et al., 2005 ). LEM proteins (named after LAP2, emerin, and MAN1) can interact with the nuclear lamina and/or chromatin-binding factors, thereby providing anchoring sites for chromatin at the nuclear periphery and modulating higher order chromatin structure (for reviews see Gruenbaum et al., 2005 ; Wagner and Krohne, 2007 ). Our analyses further revealed that Src1 is an integral membrane protein that spans the inner nuclear membrane twice, thereby exposing the putative DNA-binding N and C domains to the nucleoplasm. As shown by chromatin immunoprecipitation (ChIP)–on-chip analysis, Src1 is associated with subtelomeric regions that are located at the nuclear periphery. Moreover, the regulation of expression of genes located in the subtelomeric regions (e.g., PHO11, PHO12, and PHO84) is altered in cells lacking Src1. These data suggest that Src1 functions at the interface between chromatin organization of subtelomeric genes at the inner nuclear membrane and TREX-dependent mRNA export.

**Results**

**TREX factors interact genetically with SRC1**

To unravel the genetic network in which TREX factors carry out their function in vivo, we performed synthetic lethal screens with the mutant strains hpr1Δ (a THO–TREX factor), which is thermo-sensitive for growth at 37°C, and thp1Δ (a TREX-2 factor). Each screen yielded a synthetic lethal mutant that was complemented by SRC1 (unpublished data). DNA sequencing of the SRC1 allele recovered from the thp1Δ synthetic lethal mutant indeed revealed that Src1 was inactivated in the synthetic lethal strain by a premature stop codon at amino acid 455 (unpublished data). SRC1 was previously identified as an intron-containing gene involved in sister chromatid segregation (Rodríguez-Navarro et al., 2002 ). In another study, Src1/Heh1 was shown to be located at the inner side of the nuclear membrane (King et al., 2006 ). To directly verify the identified genetic interactions, we combined the nonessential src1Δ (null) allele with the nonessential thp1Δ or hpr1Δ disruption alleles. This analysis confirmed that the generated double mutants, src1Δhpr1Δ and src1Δthp1Δ, were not viable (i.e., synthetic lethal; Fig. 1 ).

To gain further insight in the genetic network in which SRC1 is active, we tested additional factors with known roles in transcription-coupled mRNA export for a functional overlap with SRC1. This analysis indicated that SRC1 is genetically linked to another TREX-2 factor SAC3 (Fig. 1 ) but not to SUS1 or CDC31 (Fig. 1 and not depicted). This correlates with the fact that src1Δ is not synthetic lethal with sac3ΔCId, where the binding site for Sus1 and Cdc31 is deleted (Fig. 1 ). Unexpectedly, the other THO–TREX mutants thox2Δ, mft1Δ, and thp2Δ were not found to be genetically linked to src1Δ (Fig. 1 and not depicted). Moreover, factors acting more downstream in transcription-coupled mRNA export (e.g., mutants of YRA1 or MEX67 ) and mlp1Δmp2Δ were not synthetic lethal with src1Δ (unpublished data). In contrast, a synthetically enhanced growth inhibition was observed between src1Δ and the sub2-85 allele (Fig. 1 ). Collectively, these genetic studies indicated that SRC1 is functionally linked to factors of the THO–TREX and TREX-2 complex, and thus, Src1 might functionally overlap with an upstream step in the formation of an export-competent mRNP.

**Two forms of Src1 protein generated by alternative splicing are functionally not equivalent**

To study the in vivo role of Src1 with respect to its genetic linkage to TREX factors, we sought to tag chromosomal SRC1 at the C terminus with the tandem affinity purification (TAP) and GFP tag to perform affinity purification and subcellular location experiments, respectively. However, C-terminal tagging was not straightforward because SRC1 contains an intron that can be alternatively spliced (Davis et al., 2000 ; Rodríguez-Navarro et al., 2002 ).
cal functions, as the long form of Src1 complements the synthetic lethal phenotype of THO–TREX and TREX-2 mutants significantly better than the short form (Fig. 2C).

To find out whether both Src1s have a similar subcellular location, we performed fluorescence microscopy. Both Src1-L and Src1-S tagged at the N terminus with GFP showed a distinct concentration at the nuclear envelope with no apparent staining of other cellular membranes (Fig. S1, left, available at http://www.jcb.org/cgi/content/full/jcb.200803098/DC1). Moreover, Src1-L and Src1-S did not cluster with NPCs in the NPC-clustering nup133/H9004 mutant (Fig. S1, right). Thus, Src1-L and Src1-S are located at the inner nuclear membrane and appear not to have stable physical contact to the NPCs, although a transient interaction cannot be excluded.

Specifically, the SRC1 intron has two alternative 5′ splice sites, which could potentially encode two different Src1 proteins: a long form with 834 (Src1-L) and a shorter form with 687 amino acids (Src1-S). Importantly, Src1-L and Src1-S would differ in their amino acid sequences at the C-terminal end because the alternative 5′ splice sites shift the reading frame in the 3′ exon (Fig. 2A).

To demonstrate that both Src1 splice variants are produced in vivo, we inserted the TAP tag at the two alternative stop codons by homologous recombination (Fig. 2A). Both Src1-L and Src1-S were detected in about equimolar amounts in whole cell lysates (Fig. 2B). Notably, Src1-L and Src1-S do not have identical functions, as the long form of Src1 complements the synthetic lethal phenotype of THO–TREX and TREX-2 mutants significantly better than the short form (Fig. 2C).

To find out whether both Src1s have a similar subcellular location, we performed fluorescence microscopy. Both Src1-L and Src1-S tagged at the N terminus with GFP exhibited a distinct concentration at the nuclear envelope with no apparent staining of other cellular membranes (Fig. S1, left, available at http://www.jcb.org/cgi/content/full/jcb.200803098/DC1). Moreover, Src1-L and Src1-S did not cluster with NPCs in the NPC-clustering nup133Δ mutant (Fig. S1, right). Thus, Src1-L and Src1-S are located at the inner nuclear membrane and appear not to have stable physical contact to the NPCs, although a transient interaction cannot be excluded.
Figure 2. Alternative splicing of SRC1 results in two different spliced protein forms. (A) Schematic overview of pre-mRNA, mRNA, and protein products upon alternative splicing. Either a 126- or a 130-nt intron can be excised by using two alternative 5′ splice sites. In the latter case, a frame shift results in an earlier stop codon and, therefore, in a shorter protein with a different amino acid sequence at the C terminus compared with Src1-L. Conserved domains (HEH/LEM and MSC) and transmembrane domains (M) are indicated. Numbers represent amino acid residues. (B) Whole cell lysates of N- (TAP-Src1) and C-terminal TAP-tagged Src1-L or Src1-S were analyzed by SDS-PAGE followed by Western blotting using anti-ProtA antibodies. (C) Genetic relationship of SRC1 splice variants with TREX–THO and TREX-2 components. The double-disruption strains were transformed with empty vector, GFP-Src1 splice variants, and the respective TREX component. Transformants were spotted in 10-fold serial dilutions on 5-FOA–containing plates for 5 d at the indicated temperatures.

**Src1-L and Src1-S are integral inner nuclear membrane proteins with different topology**

Biocomputational analysis suggested that Src1-L could be a membrane protein with two membrane-spanning sequences, but only the first transmembrane span has a strong prediction (Fig. 3 A). To experimentally verify that Src1-L is an integral membrane protein, whole cell lysates containing Src1-TAP were extracted under different conditions, and the partitioning of Src1-TAP into soluble and insoluble (membrane) fractions was tested and compared with...
the behavior of a peripheral membrane protein (Nsp1) and an integral membrane protein (vacuolar Vam3). Whereas high salt (1 M NaCl) or pH 11.5 treatment did not release Src1-L and Src1-S, detergent (Triton X-100) efficiently solubilized both Src1-TAP splice variants, as with Vam3 (Fig. 3B). Thus, the biochemical behavior of Src1 is typical of an integral membrane protein.

Because the putative transmembrane span M2 (residues 708–725) of Src1 has a lower hydrophobicity than transmembrane span M1 (residues 450–474) and M2 is absent from the Src1-S form because of alternative splicing (Fig. 3A), we wanted to analyze the topology of both Src1 forms within the inner nuclear membrane in living cells (see models in Fig. 4A). Thus, we developed a novel in vivo assay to probe for the topology of Src1-L and Src1-S within the inner nuclear membrane. Src1 was genomically TAP tagged at the N terminus (protein A [ProtA]–tobacco etch virus [TEV]–calmodulin-binding peptide–Src1) or C terminus (Src1–calmodulin-binding peptide–TEV-ProtA). The accessible TEV cleavage site in these constructs could be exploited to test whether it is cleaved by an inducible TEV protease carrying an NLS and a myc tag. We anticipated that the TEV protease should remove the ProtA moiety only from those Src1 constructs in which the epitope is exposed to the nucleoplasm but not when hidden in the perinuclear space (Fig. 4B, left). When cells were grown in glucose (no TEV expression), all Src1 constructs retained the ProtA epitope, as shown by Western blotting with anti-ProtA antibodies (Fig. 4B, lanes 1, 3, and 5).
To determine the contribution of the transmembrane spans M1 and M2 for membrane insertion of Src1, mutants lacking M1 or M2 were expressed in the src1/H9004 mutant and subsequently analyzed in vitro and in vivo. Biochemical studies revealed that Src1/H9004 M2 behaved like an integral membrane protein, whereas Src1/H9004 M1 was no longer inserted into the membrane and became soluble (Fig. 3B). Thus, M1 is necessary and sufficient for membrane insertion of Src1. However, M2 alone could not confer a stable membrane insertion and only integrated into the nuclear membrane in the context of full-length Src1 (Fig. 4B). When cells were shifted to galactose to induce the TEV protease, both short and long N-terminally tagged Src1 as well as C-terminally tagged Src1-L lost their ProtA tags, suggesting that the N terminus of Src1-L and Src1-S and the C terminus of Src1-L are facing the nucleoplasm (Fig. 4B, lanes 2 and 4). However, the TEV protease could not remove the ProtA epitope from the C-terminally tagged Src1-S (Fig. 4B, lane 6). Altogether, these data suggest that Src1-L spans the inner nuclear membrane twice with the N and C domain exposed to the nucleoplasm (Fig. 4A, boxed in red). In contrast, Src1-S spans the membrane once with the N domain located in the nucleoplasm and a short C-terminal domain hidden in the perinuclear space (Fig. 4A, boxed in red).

**Role of the various Src1 domains for membrane insertion and nuclear envelope targeting**

To determine the contribution of the transmembrane spans M1 and M2 for membrane insertion of Src1, mutants lacking M1 or M2 were expressed in the src1Δ mutant and subsequently analyzed in vitro and in vivo. Biochemical studies revealed that Src1ΔM2 behaved like an integral membrane protein, whereas Src1ΔM1 was no longer inserted into the membrane and became soluble (Fig. 3B). Thus, M1 is necessary and sufficient for membrane insertion of Src1. However, M2 alone could not confer a stable membrane insertion and only integrated into the nuclear membrane in the context of full-length Src1 (Fig. 4B).
Figure 5. **Src1 domain analysis.** (A) Schematic representation of Src1 deletion constructs. (B) Fluorescence microscopy of src1Δ cells harboring the indicated GFP-tagged Src1 full-length and truncation constructs. (C) The double-disruption strain src1Δthp1Δ carrying plasmid-borne THP1 was transformed with the indicated constructs, and cells were spotted in 10-fold serial dilutions on 5-FOA-containing plates and incubated for 5 d at 30°C. Only cDNA-based constructs of Src1-L are shown.
Thus, the first transmembrane domain in Src1 is necessary for membrane insertion, whereas the second membrane span alone cannot confer integral insertion into the membrane but develops insertion activity when M1 is present.

To study the role of the transmembrane sequences M1 and M2 and of other domains for nuclear envelope targeting of Src1 in vivo, GFP-tagged Src1 mutant constructs (Fig. 5 A) were analyzed for their subcellular location by fluorescence microscopy (Fig. 5 B). This analysis showed that GFP-Src1/H9004M2 was still located at the nuclear envelope, in contrast to GFP-Src1ΔM1, which was detached from the nuclear periphery and mislocalized to the nucleoplasm.

Consistent with these findings, the N-terminal domain of Src1 devoid of any membrane span showed a strong nuclear accumulation. The reciprocal construct, GFP-Src1ΔN, was still targeted to the nuclear envelope, albeit with lower efficiency, and was partly found to be associated with other membranes (cortical ER or plasma membrane). Deletion of only the N-terminal
oligo(dT) nucleotide probes did not reveal a nuclear mRNA export defect in src1Δ/H9004 cells (unpublished data), consistent with the finding that Src1 is not genetically linked to mRNA export factors such as Mex67 (see the first Results section). To investigate whether Src1 plays a role in transcription, we analyzed the expression profile of the 6,000 yeast genes in the src1Δ strain using DNA macroarrays. This genome-wide analysis indicated that only a small number of genes (60) is affected in the src1Δ/H9004 mutant (increased or decreased expression), which also included genes that have a subtelomeric location (Table S1, available at http://www.jcb.org/cgi/content/full/jcb.200803098/DC1; and Gene Expression Omnibus database). To analyze whether subtelomeric genes have a higher probability to be affected in their expression than other genes when SRC1 is deleted, the relative increases (src1Δ/wild type [wt] when src1Δ > wt) or decreases (wt/src1Δ when src1Δ < wt) of gene expression were plotted versus their distance to the closest telomere or centromere using a sliding window of 100 genes. This analysis suggested a statistically significant misregulation of subtelomeric genes in src1Δ/H9004 cells (subtelomeric regions are defined to be within 25 kb from each telomere; Fig. 6A; Louis, 1995). This effect is specific to telomeres because it does not occur (e.g., in regions close to the centromeres; Fig. 6B). In particular, several PHO genes (PHO84, PHO11, PHO12, and, to a lesser extent, also PHO4) were markedly up-regulated in cells devoid of Src1 (Table S1). Quantitative real-time PCR independently showed that PHO84, PHO11, and PHO12 transcript levels increased approximately fivefold in src1Δ cells when grown in high phosphate (HP) medium. However, further up-regulation of these PHO genes was no longer observed in src1Δ cells when shifted from high to low

Src1 is involved in expression of subtelomeric PHO genes that exhibit a perinuclear location

As Src1 was found in this study to be linked to TREX factors, we determined whether Src1 participates in transcription and/or mRNA export. In situ hybridization using fluorescently labeled oligo(dT) nucleotide probes did not reveal a nuclear mRNA export defect in src1Δ cells (unpublished data), consistent with the finding that Src1 is not genetically linked to mRNA export factors such as Mex67 (see the first Results section). To investigate whether Src1 plays a role in transcription, we analyzed the expression profile of the 6,000 yeast genes in the src1Δ strain using DNA macroarrays. This genome-wide analysis indicated that only a small number of genes (60) is affected in the src1Δ/H9004 mutant (increased or decreased expression), which also included genes that have a subtelomeric location (Table S1, available at http://www.jcb.org/cgi/content/full/jcb.200803098/DC1; and Gene Expression Omnibus database). To analyze whether subtelomeric genes have a higher probability to be affected in their expression than other genes when SRC1 is deleted, the relative increases (src1Δ/wild type [wt] when src1Δ > wt) or decreases (wt/src1Δ when src1Δ < wt) of gene expression were plotted versus their distance to the closest telomere or centromere using a sliding window of 100 genes. This analysis suggested a statistically significant misregulation of subtelomeric genes in src1Δ/H9004 cells (subtelomeric regions are defined to be within 25 kb from each telomere; Fig. 6A; Louis, 1995). This effect is specific to telomeres because it does not occur (e.g., in regions close to the centromeres; Fig. 6B). In particular, several PHO genes (PHO84, PHO11, PHO12, and, to a lesser extent, also PHO4) were markedly up-regulated in cells devoid of Src1 (Table S1). Quantitative real-time PCR independently showed that PHO84, PHO11, and PHO12 transcript levels increased approximately fivefold in src1Δ cells when grown in high phosphate (HP) medium. However, further up-regulation of these PHO genes was no longer observed in src1Δ cells when shifted from high to low

LEM domain (ΔLEM) did not affect the nuclear envelope location of Src1. Notably, the Src1ΔNΔM1 construct was also located in the nucleoplasm with occasional perinuclear spots. Altogether, the data suggest that the N domain and, to a lesser extent, also the C domain of Src1-L contribute to nuclear targeting, whereas the first transmembrane domain is necessary and sufficient to insert Src1 into the inner nuclear envelope.

To assess which Src1 domains are functionally important, we tested the ability of the different truncated Src1 forms to complement the synthetic lethal phenotype of the src1Δ/hplΔ double mutant (Fig. 5C). None of the constructs, which lacked either M1 or M2, complemented the src1Δ/hplΔ strain. Thus, although Src1ΔM2 apparently is targeted to and inserted into the nuclear membrane, it is not functional in our complementation assay. Notably, the C-terminal domain of Src1-L is exposed to the nucleoplasm (see previous section), but in the Src1ΔM2 strain it is hidden in the lumen of the perinuclear space (Fig. 4B). Thus, the C-terminal domain of Src1 performs a crucial role with respect to the Hpl1 function and has to be both targeted to the membrane and exposed to the nucleoplasm. In contrast, Src1 devoid of the LEM domain or the entire N terminus is able to significantly complement the src1Δ/hplΔ strain and thus may perform additional functions.

Src1 is involved in expression of subtelomeric PHO genes that exhibit a perinuclear location

As Src1 was found in this study to be linked to TREX factors, we determined whether Src1 participates in transcription and/or mRNA export. In situ hybridization using fluorescently labeled oligo(dT) nucleotide probes did not reveal a nuclear mRNA export defect in src1Δ cells (unpublished data), consistent with the finding that Src1 is not genetically linked to mRNA export factors such as Mex67 (see the first Results section). To investigate whether Src1 plays a role in transcription, we analyzed the expression profile of the 6,000 yeast genes in the src1Δ strain using DNA macroarrays. This genome-wide analysis indicated that only a small number of genes (60) is affected in the src1Δ/H9004 mutant (increased or decreased expression), which also included genes that have a subtelomeric location (Table S1, available at http://www.jcb.org/cgi/content/full/jcb.200803098/DC1; and Gene Expression Omnibus database). To analyze whether subtelomeric genes have a higher probability to be affected in their expression than other genes when SRC1 is deleted, the relative increases (src1Δ/wild type [wt] when src1Δ > wt) or decreases (wt/src1Δ when src1Δ < wt) of gene expression were plotted versus their distance to the closest telomere or centromere using a sliding window of 100 genes. This analysis suggested a statistically significant misregulation of subtelomeric genes in src1Δ/H9004 cells (subtelomeric regions are defined to be within 25 kb from each telomere; Fig. 6A; Louis, 1995). This effect is specific to telomeres because it does not occur (e.g., in regions close to the centromeres; Fig. 6B). In particular, several PHO genes (PHO84, PHO11, PHO12, and, to a lesser extent, also PHO4) were markedly up-regulated in cells devoid of Src1 (Table S1). Quantitative real-time PCR independently showed that PHO84, PHO11, and PHO12 transcript levels increased approximately fivefold in src1Δ cells when grown in high phosphate (HP) medium. However, further up-regulation of these PHO genes was no longer observed in src1Δ cells when shifted from high to low
or – Src1-S could be significantly enriched using a protocol for isolation of membrane proteins, coenrichment of stoichiometrically interacting proteins was not observed (Fig. S2, available at http://www.jcb.org/cgi/content/full/jcb.200803098/DC1).

Next, we tested whether ProtA-Src1 is associated with chromatin. We performed genome-wide ChIP-on-chip analyses to determine the chromosomal distribution profile of Src1-L. Overall, three independent ChIP-on-chip experiments were performed with ProtA-Src1, and the data were highly similar. As shown in Fig. 8 and Fig. S3 (available at http://www.jcb.org/cgi/content/full/jcb.200803098/DC1) for one set of these experiments, a significant enrichment of Src1 at subtelomeric regions of all 16 yeast chromosomes was observed (note that the genomic chips used in this study did not contain the repetitive telomeric DNA). Moreover, Src1 localizes also to other heterochromatin-like regions such as the ribosomal DNA (rDNA) locus, the mating-type loci (within subtelomeric regions of chromosome III), and, albeit only to a minor degree, to centromeric regions (Fig. 8). Most of the other locations to which Src1 maps to a minor extent are likely to be unspecific, as those loci are also enriched in the ProtA control. Altogether, these data indicate that Src1 is preferentially associated with subtelomeric chromatin, and this interaction can affect expression of subtelomeric-located genes.

To test whether Src1 plays a role in gene silencing, we monitored the expression of a silenced URA3 reporter inserted into the telomeric region of chromosome 7 in the presence and absence of SRC1. Deletion of SRC1 did not affect telomeric silencing as revealed by a normal growth of cells on 5-fluoroorotic acid (5-FOA)–containing plates in comparison with a wt strain. In contrast, when yku70Δ cells known to have a telomeric silencing defect (Laroche et al., 1998) were plated on 5-FOA, a significant growth inhibition was observed. Moreover, the combination of src1Δ and yku70Δ in haploid cells caused a

Src1 is associated with subtelomeric chromatin

To identify molecules, which potentially link Src1 to chromatin, we sought to affinity purify ProtA-Src1. Although ProtA–Src1-L or –Src1-S could be significantly enriched using a protocol for isolation of membrane proteins, coenrichment of stoichiometrically interacting proteins was not observed (Fig. S2, available at http://www.jcb.org/cgi/content/full/jcb.200803098/DC1).

Next, we tested whether ProtA-Src1 is associated with chromatin. We performed genome-wide ChIP-on-chip analyses to determine the chromosomal distribution profile of Src1-L. Overall, three independent ChIP-on-chip experiments were performed with ProtA-Src1, and the data were highly similar. As shown in Fig. 8 and Fig. S3 (available at http://www.jcb.org/cgi/content/full/jcb.200803098/DC1) for one set of these experiments, a significant enrichment of Src1 at subtelomeric regions of all 16 yeast chromosomes was observed (note that the genomic chips used in this study did not contain the repetitive telomeric DNA). Moreover, Src1 localizes also to other heterochromatin-like regions such as the ribosomal DNA (rDNA) locus, the mating-type loci (within subtelomeric regions of chromosome III), and, albeit only to a minor degree, to centromeric regions (Fig. 8). Most of the other locations to which Src1 maps to a minor extent are likely to be unspecific, as those loci are also enriched in the ProtA control. Altogether, these data indicate that Src1 is preferentially associated with subtelomeric chromatin, and this interaction can affect expression of subtelomeric-located genes.

To test whether Src1 plays a role in gene silencing, we monitored the expression of a silenced URA3 reporter inserted into the telomeric region of chromosome 7 in the presence and absence of SRC1. Deletion of SRC1 did not affect telomeric silencing as revealed by a normal growth of cells on 5-fluoroorotic acid (5-FOA)–containing plates in comparison with a wt strain. In contrast, when yku70Δ cells known to have a telomeric silencing defect (Laroche et al., 1998) were plated on 5-FOA, a significant growth inhibition was observed. Moreover, the combination of src1Δ and yku70Δ in haploid cells caused a...
synergistic inhibition of growth on 5-FOA plates (Fig. 9). These data suggest that Src1 in combination with another silencing factor can synergistically affect telomeric silencing.

Discussion

This study has revealed that the integral inner nuclear membrane protein Src1 functions in gene regulation of subtelomeric genes and is embedded functionally in a network of factors, which participate in transcription-coupled mRNA export. Importantly, Src1 is associated with subtelomeric chromatin and thus can help to organize this region of the chromosomes. In previous studies, Src1 was shown to contain an intron with the possibility of alternative splicing (Davis et al., 2000; Rodríguez-Navarro et al., 2002). Our study revealed that both splice forms localize to the nuclear periphery and are integral inner nuclear membrane proteins (King et al., 2006). However, both proteins are not functionally equivalent. To the best of our knowledge, this is the first demonstration that two forms of a protein generated by alternative splicing can function within the N domain of Src1.

Src1 mediates nuclear targeting, and insertion into the nuclear membrane requires the first transmembrane domain. These findings are consistent with studies of vertebrate LEM2 and MAN1 in which nuclear envelope targeting and retention are also dependent on the first transmembrane domain and the N terminus (Wu et al., 2002; Brachner et al., 2005). Previously, it has been shown that Src1 is imported to the inner nuclear membrane by the Kap60–Kap95 pathway in a RanGTP-dependent manner (King et al., 2006). Replacing the entire N terminus of Src1 with the classical SV40 NLS, however, was not sufficient to restore its normal localization at the inner nuclear membrane (unpublished data). This suggests that besides an NLS, additional targeting/retention information is retained within the N domain of Src1.

Src1 is preferentially associated with heterochromatin such as subtelomeric and telomeric chromatin. As both N- and C-terminal domains of Src1-L face the nucleoplasm and each harbors a potential DNA-interacting domain (LEM and MSC), it is possible that the interaction with DNA is direct, but adaptor proteins could also be involved in chromatin binding. LEM domain proteins in higher eukaryotes are able to associate with chromatin either directly (Cai et al., 2001; Caputo et al., 2006) or via the chromatin-associated barrier to autointegration factor (for review see Wagner and Krohne, 2007). However, a homologue of barrier to autointegration factor does not exist in yeast, and we were unable to find stoichiometric protein interaction partners of Src1.

Telomeres are anchored to the nuclear periphery in yeast by redundant mechanisms involving the heterodimeric yKu complex and Sir4, which associates with the membrane-associated Esc1 (Hediger et al., 2002; Taddei et al., 2004). Moreover, the SUN domain protein Mps3 assists in Sir4-dependent telomere tethering and silencing (Bupp et al., 2007). Whether Src1 functions directly in silencing remains open. SRC1 is not genetically linked to ESC1 or YKU70 (unpublished data), and telomere silencing was not affected in src1Δ cells. However, disruption of SRC1 affected telomere length (Askree et al., 2004), and a synergistically reduced silencing of subtelomeric genes was observed when src1Δ was combined with the yku70Δ allele.

In addition to telomeres, Src1 is also associated with the rDNA locus. Both consist of repetitive sequences, which have to be protected from uncontrolled homologous recombination to maintain genomic integrity. Notably, Src1 was copurified with Lrs4, a factor located at the rDNA locus with a role in rDNA silencing and suppression of rDNA recombination (Huang et al., 2006). Lrs4 together with cohesins are thought to prevent the formation of chromatid junctions, and Src1 could aid in this step locally at the inner nuclear membrane. In line with a function together with cohesins, Src1 has been reported to be involved in sister chromatid separation and is genetically linked to SCC1 and ESP1 (Rodríguez-Navarro et al., 2002).

Our study further reveals that Src1 is required for proper gene expression of subtelomeric genes, such as several PHO genes, that are associated with the inner nuclear membrane. Transcription of PHO genes in response to the phosphate concentration is regulated by the phosphorylation status of the transcription factor Pho4, which in turn affects its nucleocytoplasmic shuttling (O’Neill et al., 1996). Our data suggest that the subtelomeric PHO genes may exhibit, besides specific control by Pho4, a global regulation as a result of their association with the inner nuclear membrane, which is repressive for gene expression. By which mechanism Src1 could affect subtelomeric gene expression remains unclear. It is known that in contrast to telomere silencing mediated by the Sir complex, silencing of genes within subtelomeric/HAST (Hda1-affected subtelomeric) domains depends on the local nucleosome structure (Wyrick et al., 1999). This is characterized by histone H3 hypomethylation and hypoacetylation (Bernstein et al., 2002; Roby et al., 2002). Whether Src1 specifically affects histone modifications in subtelomeric regions remains to be determined, but its role in gene expression could be envisaged in a broader sense (e.g., by interfering with different types of mechanisms that activate or repress gene expression). The effect of both up- and down-regulation of genes observed in the src1Δ mutant could be caused by transcription factors concentrated at the nuclear periphery that act as both activators and repressors. Lack of Src1 might restrict the access of these factors to subtelomeric loci as a result of an altered chromatin structure. The macroarray data also show that expression of some genes that are not located at subtelomeric regions, but randomly across the chromosomes, were affected...
by loss of \textit{SRC1}. Because Src1 did not interact significantly with these genes according to ChIP-on-chip, the effect could be indirect (e.g., caused by misregulation of subtelomeric genes, which in turn alter expression of genes located elsewhere in the genome).

The finding that Src1 is genetically linked to factors of the THO–TREX and TREX-2 complexes suggested an involvement in transcription and/or export. However, the disruption of \textit{SRC1} did not cause an mRNA export defect. Thus, Src1 could be connected to the transcription-assisted (upstream) functions of these TREX complexes. It is well established that mutations in these TREX factors (e.g., in \textit{Thp1}, \textit{Sac3}, and \textit{Hpr1}) not only inhibit transcription elongation but also induce transcription-dependent hyperrecombination and genome instability (for review see Reed and Cheng, 2005). Src1 is not genetically linked to all of the subunits within the TREX complexes may have different roles with respect to Src1 function. For example, Sub2, a subunit of THO–TREX, not only functions in transcription, splicing, and mRNA export but was also shown to be localized to telomeres and to affect heterochromatin gene expression (Lahue et al., 2005).

Collectively, the simplest model of how Src1 functions as an integral inner nuclear membrane protein is to help in recruiting and organizing the peripheral (telomeric and subtelomeric) chromatin, perhaps by directly binding to specific DNA sequences or nucleosomes (see above). By organizing subtelomeric and telomeric DNA in a distinct nuclear compartment, the nuclear periphery, Src1 could help to cluster a group of genes in a zone in which silencing factors, transcription activators, and repressors cooperate for regulated gene expression and which also has access to an efficient mRNA export pathway via the TREX machineries associated with NPCs. Subtelomeric and telomeric chromatin and the rDNA locus are repetitive, and their preservation is crucial for maintaining genome integrity. Thus, localization of this chromatin to the nuclear membrane could also play a role in preventing unwanted recombination. Many of these described functions attributed to Src1 can now be tested by exploiting \textit{src1} mutants in different functional assays.

Last but not least, Src1 is related to higher eukaryotic LEM domain–containing inner nuclear membrane proteins that were shown to interact with transcription regulators (e.g., R-Smads and HDAC3) and thus affect gene expression and signal transduction by recruitment of transcription regulators to the nuclear periphery (for reviews see Gruenbaum et al., 2005; Wagner and Krohne, 2007). Our study showed that the LEM domain protein Src1 is associated with subtelomeric chromatin and affects gene expression in this region. Thus, Src1 could serve as a model protein to gain further insight into the complex role of inner nuclear membrane proteins, including their involvement in diseases such as laminopathies.

### Materials and methods

#### Yeast strains and plasmid constructs

\textit{S. cerevisiae} strains and plasmids are listed in Table S2 (available at \url{http://www.jcb.org/cgi/content/full/jcb.200803098/DC1}). The synthetic lethal screens were performed as described previously (Segref et al., 1997). In the \textit{thp1Δ} screen, 54 synthetic lethal mutants were obtained from \(70,000\) colonies. 24 of these mutants were complemented by plasmid-borne \textit{THP1} and used for further analysis.

#### Fluorescence microscopy

Cells were grown in selective medium at 30°C to logarithmic phase. Fluorescence microscopy was performed using an Imager Z1 (Carl Zeiss, Inc.) equipped with a 63x NA 1.4 Plan-Apochromat oil immersion lens (Carl Zeiss, Inc.) and using DICII and H6E GFP filters. Pictures were acquired with a camera (AxioCamMRm; Carl Zeiss, Inc.) and AxioVision 4.3 software (Carl Zeiss, Inc.). Pictures were exported as jpg files and processed in Photoshop 7.0 (Adobe) for levels.

#### Src1 membrane extraction, topology analysis, and affinity purification

500 mg of spheroplasts prepared from early log-phase cultures were lysed with a Dounce homogenizer in 3 ml of buffer [150 mM NaCl and 20 mM Tris-HCl, pH 8.0] containing protease inhibitors and was pelleted at 2,000 rpm for 2 min. The supernatant was centrifuged at 13,000 rpm for 20 min at 4°C, resulting in pellet and supernatant. One fourth of the pellet was re-suspended in 400 pl of buffer (\(1\) Triton X-100 and 1 M NaCl or in 0.1 M sodium carbonate, pH 11.5), incubated 30 min on ice, and separated into a soluble and pellet fraction by ultracentrifugation at 100,000 g for 1 h at 4°C. Equal amounts were analyzed by Western blotting using anti-ProA, anti-Vam3, and anti-Nsp1 antibodies.

The galactose promoter–driven TEV protease fused to NLS and a myc tag was integrated into strains expressing N- or C-terminal–tagged Src1 forms. An equivalent of 3 OD_{600} of exponentially growing cells in YPD or YPG at 30°C was harvested, TCA lysis was performed, and equal amounts were analyzed by Western blotting using anti-ProA, anti-myc, and anti-Arc1 antibodies. Affinity purification of ProtA-tagged bait Src1 was performed as described previously (Gavin et al., 2002).

#### DNA macroarray measurement of RNA levels

50 ml of wt and \textit{src1Δ} cells (OD_{600} 0.5) were pelleted and immediately frozen. After thawing the samples on ice, total RNA was isolated and reverse transcribed using [32P]-labeled deoxyctydine triphosphate and oligo (d[5’T1VN-3’]) (García-Martínez et al., 2004). Hybridization was performed on nylon filters using PCR-amplified whole ORF sequences as probes (Alberola et al., 2004; García-Martínez et al., 2004) except that hybridizations were 48–48 h. Different controls were used for each strain. Scanning and analysis of the macroarrays were performed essentially as described previously using ArrayStat software (García-Martínez et al., 2004; Rodríguez-Navarro et al., 2004). Apart from the genes detected by ArrayStat as significantly up- or down-regulated, we obtained an additional list of 16 genes that were expressed over background in the three replicates in \textit{src1Δ} and below background in all three replicates in wt cells. Array data have been submitted to the Gene Expression Omnibus data repository (accession no. GSE6570).

#### RNA isolation and real-time PCR

Cells were grown in selective HP medium [11 mM KH_{2}PO_{4}] at 30°C to an OD_{600} of 0.5. Half of the cells was washed and grown for 2 h in selective LP medium [220 μM KH_{2}PO_{4}] at 30°C. Total RNA was isolated using the RNeasy Mini kit (QIAGEN). cDNA was synthesized from 1 μg of total RNA using the Quantitect Reverse Transcription kit (QIAGEN). PHO11, PHO12, and PHO84 cDNAs were detected by quantitative real-time PCR (ABI-Prism 7000; Applied Biosystems) using specific primers and TaqMan probes (Table S3, available at \url{http://www.jcb.org/cgi/content/full/jcb.200803098/DC1}). Triplicate reactions containing cDNA equivalent to 0.08 μg RNA were analyzed per experiment.

#### ChIP-on-chip analysis

ChIP-on-chip analyses were performed as described previously (Cam et al., 2003) with minor modifications. Cells were grown in selective HP medium at 30°C to an OD_{600} of 0.5. After washing, the cells were grown for 2 h in selective LP medium at 30°C. The PFA-dimethyl adipimidate cross-linked chromatin was sheared by sonication and without preclearing was incubated with IgG Sepharose 6 Fast Flow (GE Healthcare) for 2–3 h at 4°C. After extensive washing and amplification, we combined 500 ng Cy5-labeled and Cy3-labeled whole cell extract DNA and hybridized it onto a 4 × 44 K \textit{S. cerevisiae} Whole Genome Chip-on-chip Microarray (Agilent Technologies). Hybridization, washes, and processing slides were performed in accordance with the yeast ChIP-on-chip protocol (version 9.1; Agilent Technologies).

For statistical analysis, we have taken a 10-kb region from the end of the chromosomes. The 599 independent probes representing subtelomeric regions show an average of 4.45 × Src1 enrichment over the base-line. The remaining 40,881 probes give a 1.06 × average enrichment. In the negative control (cells expressing only ProtA), we did not detect any enrichment of immunoprecipitated ProtA at telomeres, centromeres, HML...
and HMR loci, or on rDNA (1.08x average enrichment at subtelomeric region vs. 1.03x).}

**Localization of single chromosomal loci**
Growing and mounting yeast cells for in vivo microscopy were performed as described previously (Cabal et al., 2006) using a Revolution Nikolov disk confocal system (Andor Technology). The system was controlled using Revolution IQ software (version 1.5; Andor Technology). Z stacks of 41 images with a 250-nm Z step were acquired using a 100×/1.4 NA Plan Apochromat oil immersion objective (Carl Zeiss, Inc.). Pixel size was 77.3 nm. Quantitative analysis of microscopy data was performed using Matlab software as described previously (Cabal et al., 2006). In brief, each nucleus was processed to extract 3D coordinates of the locus, the nuclear center, and the nuclear envelope. In the following step, loci positions relative to the nuclear center obtained from different nuclei were plotted on the same 3D graph. Finally, and for reading facility, this 3D graph was projected on a 2D graph using radial projection (i.e., the loci—nuclear center distance is conserved; Cabal et al., 2006).

**Online supplemental material**
Fig. S1 contains the localization of Src1 splice forms in wt and nup133Δ cells. Fig. S2 shows that Src1-L and Src1-S does not purify stoichiometric binding partners. Fig. S3 shows the genomic-wide distribution maps of Src1-L Table S1 contains the up- and down-regulated genes in src1Δ cells. Table S2 indicates the yeast strains and plasmids used, and Table S3 presents the real-time PCR primers used in this study. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.200803098/DC1.

We thank Dr. Christian Ungermann for providing the anti-Vam3 antibody and Dr. Francisco Jiménez-Díaz for providing the FUS1-NES(NAl)TEV2V-NII construct. We thank Dr. Katja Sträßer for performing the hprt IA assay and Sheila Lutz, Dr. Dieter Kessler, and Jonas Binding for discussions. We thank Axel B. Berger, Dr. Christophe Zimmer, and Dr. Olivier Gadal for sharing unpublished data, providing yeast strains, and providing access to bioinformatics tools. We wish to thank Vicent Pelechano for his advice with the macroarray data analysis.

The work in J.E. Pérez-Ortín’s laboratory was funded by Ministerio de Educación y Ciencia (Spain) with grants BFU2006-15446-C03-02/BMC and BFU2007-67575-C03-01/BMC and EMBIO J. 21:5843–5852.


Segref, A., K. Sharma, V. Doye, A. Hellwig, J. Huber, R. Luhrmann, and E. Hurt.
1997. Mex67p, a novel factor for nuclear mRNA export, binds to both
poly(A)+ RNA and nuclear pores. EMBO J. 16:3256–3271.

Sommer, P., and U. Nehrbass. 2005. Quality control of messenger ribonucleo-
17:294–301.

Separation of silencing from perinuclear anchoring functions in yeast

Wagner, N., and G. Krohne. 2007. LEM-domain proteins: new insights into

Wu, W., F. Lin, and H.J. Worman. 2002. Intracellular trafficking of MAN1, an
115:1361–1371.

Wyrick, J.J., F.C. Holstege, E.G. Jennings, H.C. Causton, D. Shore, M.
Grunstein, E.S. Lander, and R.A. Young. 1999. Chromosomal landscape
402:418–421.
<table>
<thead>
<tr>
<th>ORF name</th>
<th>Gene name</th>
<th>Chromosomal location</th>
<th>Biological process</th>
<th>src1/ADpb wt</th>
<th>wt</th>
<th>src1 or src1/ADpb</th>
</tr>
</thead>
<tbody>
<tr>
<td>Up-regulated</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>YML123C</td>
<td>PHO84</td>
<td>Subtelomeric</td>
<td>Phosphate transport</td>
<td>7.06</td>
<td>23.1</td>
<td>163.0</td>
</tr>
<tr>
<td>YHR145C</td>
<td>Unknown</td>
<td></td>
<td></td>
<td>4.76</td>
<td>33.3</td>
<td>158.2</td>
</tr>
<tr>
<td>YAR071W</td>
<td>PHO11</td>
<td>Subtelomeric</td>
<td>Phosphate metabolism</td>
<td>4.05</td>
<td>31.1</td>
<td>126.0</td>
</tr>
<tr>
<td>YHR215W</td>
<td>PHO12</td>
<td>Subtelomeric</td>
<td></td>
<td>3.15</td>
<td>21.4</td>
<td>67.4</td>
</tr>
<tr>
<td>YPL019C</td>
<td>VTC3</td>
<td>Subtelomeric</td>
<td>Vacuole fusion (nonautophagic)</td>
<td>2.86</td>
<td>28.8</td>
<td>82.4</td>
</tr>
<tr>
<td>YER181C</td>
<td>Unknown</td>
<td></td>
<td></td>
<td>2.75</td>
<td>27.1</td>
<td>74.5</td>
</tr>
<tr>
<td>YJR106W</td>
<td>IME2</td>
<td></td>
<td>Protein amino acid phosphorylation, regulation of meiosis</td>
<td>2.47</td>
<td>34.2</td>
<td>84.4</td>
</tr>
<tr>
<td>YMR135W</td>
<td>Unknown</td>
<td></td>
<td></td>
<td>2.46</td>
<td>38.6</td>
<td>94.8</td>
</tr>
<tr>
<td>YDR266C</td>
<td>PHO8</td>
<td></td>
<td></td>
<td>2.45</td>
<td>91.0</td>
<td>222.9</td>
</tr>
<tr>
<td>YLR289W</td>
<td>GUF1</td>
<td></td>
<td></td>
<td>2.37</td>
<td>36.8</td>
<td>87.2</td>
</tr>
<tr>
<td>YLR234W</td>
<td>TOP3</td>
<td></td>
<td>Meiotic recombination, regulation of DNA recombination,</td>
<td>2.37</td>
<td>32.7</td>
<td>77.5</td>
</tr>
<tr>
<td>YMR106C</td>
<td>YKU80</td>
<td></td>
<td>telomerase-dependent telomere maintenance</td>
<td>2.37</td>
<td>32.7</td>
<td>77.5</td>
</tr>
<tr>
<td>YLR403W</td>
<td>SFP1</td>
<td></td>
<td>Chromatin assembly/disassembly, double-strand break repair via homologous</td>
<td>2.33</td>
<td>33.3</td>
<td>77.5</td>
</tr>
<tr>
<td>YNL047W</td>
<td>Unknown</td>
<td></td>
<td>recombination and nonhomologous end joining, telomere maintenance</td>
<td>2.33</td>
<td>33.3</td>
<td>77.5</td>
</tr>
<tr>
<td>YKIO29C</td>
<td>MAE1</td>
<td></td>
<td></td>
<td>2.32</td>
<td>68.7</td>
<td>159.5</td>
</tr>
<tr>
<td>YMR075W</td>
<td>RCO1</td>
<td></td>
<td></td>
<td>2.31</td>
<td>40.5</td>
<td>93.5</td>
</tr>
<tr>
<td>YKR019C</td>
<td>IRS4</td>
<td></td>
<td></td>
<td>2.31</td>
<td>60.4</td>
<td>139.3</td>
</tr>
<tr>
<td>YMR232W</td>
<td>FUS2</td>
<td></td>
<td></td>
<td>2.30</td>
<td>66.7</td>
<td>153.4</td>
</tr>
<tr>
<td>YML082W</td>
<td>Unknown</td>
<td></td>
<td></td>
<td>2.29</td>
<td>59.8</td>
<td>137.2</td>
</tr>
<tr>
<td>YMR278W</td>
<td>Unknown</td>
<td></td>
<td></td>
<td>2.29</td>
<td>31.0</td>
<td>71.2</td>
</tr>
<tr>
<td>YJL057C</td>
<td>IKS1</td>
<td></td>
<td></td>
<td>2.29</td>
<td>48.8</td>
<td>111.7</td>
</tr>
<tr>
<td>YJL045W</td>
<td>Unknown</td>
<td></td>
<td></td>
<td>2.27</td>
<td>91.6</td>
<td>207.6</td>
</tr>
<tr>
<td>YIL060W</td>
<td>Unknown</td>
<td></td>
<td></td>
<td>2.24</td>
<td>43.5</td>
<td>95.0</td>
</tr>
<tr>
<td>YMR008C</td>
<td>PIB1</td>
<td></td>
<td></td>
<td>2.24</td>
<td>48.6</td>
<td>108.7</td>
</tr>
<tr>
<td>YMR092C</td>
<td>AIP1</td>
<td></td>
<td></td>
<td>2.22</td>
<td>69.8</td>
<td>154.7</td>
</tr>
<tr>
<td>YLR273C</td>
<td>PIG1</td>
<td></td>
<td></td>
<td>2.22</td>
<td>88.9</td>
<td>196.5</td>
</tr>
<tr>
<td>YLR019W</td>
<td>MPS3</td>
<td></td>
<td></td>
<td>2.18</td>
<td>54.3</td>
<td>118.0</td>
</tr>
<tr>
<td>YPR135W</td>
<td>Unknown</td>
<td></td>
<td></td>
<td>2.17</td>
<td>34.3</td>
<td>74.2</td>
</tr>
<tr>
<td>YCL075W</td>
<td>Unknown</td>
<td></td>
<td></td>
<td>2.17</td>
<td>47.3</td>
<td>102.2</td>
</tr>
<tr>
<td>YML042W</td>
<td>CAT2</td>
<td></td>
<td></td>
<td>2.14</td>
<td>40.8</td>
<td>87.5</td>
</tr>
<tr>
<td>YIL160C</td>
<td>POT1</td>
<td></td>
<td></td>
<td>2.14</td>
<td>47.6</td>
<td>102.0</td>
</tr>
<tr>
<td>YIR027C</td>
<td>DAL1</td>
<td></td>
<td></td>
<td>2.11</td>
<td>44.4</td>
<td>93.6</td>
</tr>
<tr>
<td>YHR136C</td>
<td>SPL2</td>
<td></td>
<td></td>
<td>2.11</td>
<td>44.5</td>
<td>93.6</td>
</tr>
<tr>
<td>YPR192W</td>
<td>AQY1</td>
<td>Subtelomeric</td>
<td></td>
<td>2.11</td>
<td>44.5</td>
<td>93.6</td>
</tr>
<tr>
<td>YAL010C</td>
<td>MDM10</td>
<td></td>
<td>Mitochondrial genome maintenance, mitochondrial inheritance,</td>
<td>2.11</td>
<td>44.5</td>
<td>93.6</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>mitochodrial organization and biogenesis</td>
<td>2.11</td>
<td>44.5</td>
<td>93.6</td>
</tr>
<tr>
<td>Gene</td>
<td>Protein</td>
<td>Function</td>
<td>Ratio</td>
<td>Background</td>
<td>Expression</td>
<td></td>
</tr>
<tr>
<td>-----------</td>
<td>---------</td>
<td>--------------------------------------------------------------------------</td>
<td>-------</td>
<td>------------</td>
<td>------------</td>
<td></td>
</tr>
<tr>
<td>YFR034C</td>
<td>PHO4</td>
<td>Cellular response to phosphate starvation, phosphate metabolism</td>
<td>NA</td>
<td>BB 6.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>YPR150W</td>
<td>Unknown</td>
<td>Ion transport</td>
<td>NA</td>
<td>BB 8.7</td>
<td></td>
<td></td>
</tr>
<tr>
<td>YLR114C</td>
<td>POR2</td>
<td>Cytogamy, cellular morphogenesis during conjugation with cellular fusion</td>
<td>NA</td>
<td>BB 6.4</td>
<td></td>
<td></td>
</tr>
<tr>
<td>YCR089W</td>
<td>FIG2</td>
<td>Mitochondrion organization and biogenesis</td>
<td>NA</td>
<td>BB 9.3</td>
<td></td>
<td></td>
</tr>
<tr>
<td>YOR147W</td>
<td>MDM32</td>
<td>Cell transport</td>
<td>NA</td>
<td>BB 10.3</td>
<td></td>
<td></td>
</tr>
<tr>
<td>YER187W-A</td>
<td>KHS1</td>
<td>Unknown</td>
<td>NA</td>
<td>BB 8.6</td>
<td></td>
<td></td>
</tr>
<tr>
<td>YPL076W</td>
<td>GRP2</td>
<td>Nucleolus organization and biogenesis</td>
<td>NA</td>
<td>BB 28.9</td>
<td></td>
<td></td>
</tr>
<tr>
<td>YOR018W</td>
<td>ROD1</td>
<td>Response to drug</td>
<td>NA</td>
<td>BB 17.9</td>
<td></td>
<td></td>
</tr>
<tr>
<td>YOR376W</td>
<td>Unknown</td>
<td>Unknown</td>
<td>NA</td>
<td>BB 16.0</td>
<td></td>
<td></td>
</tr>
<tr>
<td>YFL051C</td>
<td>FIG2</td>
<td>Subtelomeric</td>
<td>NA</td>
<td>BB 12.2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>YDL034W</td>
<td>Unknown</td>
<td>Subtelomeric</td>
<td>NA</td>
<td>BB 10.4</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Down-regulated

<table>
<thead>
<tr>
<th>Gene</th>
<th>Protein</th>
<th>Function</th>
<th>Ratio</th>
<th>Background</th>
<th>Expression</th>
</tr>
</thead>
<tbody>
<tr>
<td>YOR383C</td>
<td>FIT3</td>
<td>Siderochrome transport</td>
<td>0.10</td>
<td>515.9</td>
<td>51.4</td>
</tr>
<tr>
<td>YLR114W</td>
<td>TIR3</td>
<td>Unknown</td>
<td>0.28</td>
<td>48.6</td>
<td>13.5</td>
</tr>
<tr>
<td>YGL055W</td>
<td>OLE1</td>
<td>Fatty acid desaturation, mitochondrion inheritance</td>
<td>0.31</td>
<td>72.5</td>
<td>22.6</td>
</tr>
<tr>
<td>YOL016C</td>
<td>CMK2</td>
<td>Protein amino acid phosphorylation, signal transduction</td>
<td>0.32</td>
<td>70.5</td>
<td>22.2</td>
</tr>
<tr>
<td>YBR268W</td>
<td>MRPL37</td>
<td>Protein biosynthesis</td>
<td>0.33</td>
<td>150.6</td>
<td>50.2</td>
</tr>
<tr>
<td>YGL136W</td>
<td>AMS1</td>
<td>Carbohydrate metabolism</td>
<td>0.35</td>
<td>30.6</td>
<td>10.8</td>
</tr>
<tr>
<td>YJR071W</td>
<td>Unknown</td>
<td>Subtelomeric</td>
<td>0.36</td>
<td>62.0</td>
<td>22.1</td>
</tr>
<tr>
<td>YGL261C</td>
<td>Subtelomeric</td>
<td>Unknown</td>
<td>0.38</td>
<td>20.0</td>
<td>7.5</td>
</tr>
<tr>
<td>YHL046C</td>
<td>Subtelomeric</td>
<td>Unknown</td>
<td>0.39</td>
<td>18.4</td>
<td>7.1</td>
</tr>
<tr>
<td>YBR019C</td>
<td>GAL10</td>
<td>Galactose metabolism</td>
<td>0.39</td>
<td>22.6</td>
<td>8.8</td>
</tr>
<tr>
<td>YMR317W</td>
<td>Subtelomeric</td>
<td>Unknown</td>
<td>0.39</td>
<td>31.6</td>
<td>12.4</td>
</tr>
<tr>
<td>YBL021C</td>
<td>HAP3</td>
<td>Transcription, regulation of carbohydrate metabolism</td>
<td>0.40</td>
<td>44.7</td>
<td>17.8</td>
</tr>
<tr>
<td>YOR152C</td>
<td>Unknown</td>
<td>Subtelomeric</td>
<td>0.42</td>
<td>29.5</td>
<td>12.2</td>
</tr>
<tr>
<td>YFL020C</td>
<td>PAU5</td>
<td>Unknown</td>
<td>0.42</td>
<td>22.3</td>
<td>9.5</td>
</tr>
<tr>
<td>YOL161C</td>
<td>Subtelomeric</td>
<td>Unknown</td>
<td>0.43</td>
<td>18.2</td>
<td>7.8</td>
</tr>
<tr>
<td>YFR024C</td>
<td>Unknown</td>
<td>Subtelomeric</td>
<td>0.43</td>
<td>28.4</td>
<td>12.3</td>
</tr>
<tr>
<td>YHL040C</td>
<td>ARN1</td>
<td>Subtelomeric</td>
<td>0.44</td>
<td>37.3</td>
<td>16.5</td>
</tr>
<tr>
<td>YCR104W</td>
<td>PAU3</td>
<td>Subtelomeric</td>
<td>0.45</td>
<td>18.7</td>
<td>8.4</td>
</tr>
</tbody>
</table>

**BB:** below background. Ratios of expression levels between src1 and an isogenic wt strain of 5,937 genes were obtained by dividing arbitrary expression units derived from three independent sets of experiments. Ratios >1 represent increased transcript levels, and ratios <1 represent decreased transcript levels. Only genes that exhibit a statistically significant altered expression in src1 cells (threshold level: twofold increase/decrease) are shown. Subtelomeric means <27 kb away from the respective telomere. Genes in bold were obtained apart from ArrayStat analysis because they had no significant expression (below the background) in wt cells.
<table>
<thead>
<tr>
<th>No.</th>
<th>Strain/plasmid</th>
<th>Relevant genotype/information</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Y1476</td>
<td>BY4741</td>
<td>MATa his3Δ1 leu2Δ0 met15Δ0 ura3Δ0</td>
<td>Euroscarf</td>
</tr>
<tr>
<td>Y2948</td>
<td>src1Δ</td>
<td>MATa ymr034w::kanMX4 his3Δ1 leu2Δ0 met15Δ0 ura3Δ0</td>
<td>Euroscarf</td>
</tr>
<tr>
<td>Y3060</td>
<td>src1Δhsp1Δ</td>
<td>MATA src1::kanMX4 hsp1::kanMX4 trp1 leu2 ura3</td>
<td>This study</td>
</tr>
<tr>
<td>Y3030</td>
<td>src1Δsac3Δ</td>
<td>MATA src1::kanMX sac3::kanMX4 his3Δ1 leu2Δ0 ura3Δ0</td>
<td>This study</td>
</tr>
<tr>
<td>Y3748</td>
<td>src1Δsus1Δ</td>
<td>MATA src1::kanMX sus1::kanMX4 trp1 leu2 ura3</td>
<td>This study</td>
</tr>
<tr>
<td>Y3045</td>
<td>src1Δhpr1Δ</td>
<td>MATA src1::kanMX hpr1::His3 his3 leu2 ura3 trp1 ade2</td>
<td>This study</td>
</tr>
<tr>
<td>Y3758</td>
<td>src1Δhsp2Δ</td>
<td>MATA src1::kanMX hsp2::kanMX4 trp1 leu2 ura3</td>
<td>This study</td>
</tr>
<tr>
<td>Y4277</td>
<td>src1Δmlp1Δ</td>
<td>MATA src1::natNT2 sub2::kanMX4 ura3Δ2</td>
<td>This study</td>
</tr>
<tr>
<td>Y3245</td>
<td>src1Δmlp2Δ</td>
<td>MATA src1::natMX mlp1::His3 mlp2::His3 ade2 ura3</td>
<td>This study</td>
</tr>
<tr>
<td>Y523</td>
<td>nup133Δ</td>
<td>MATA nup133::His3 ade2-1 his3-11,15 leu2-3,112 trp1-1, ura3-32</td>
<td>Doye et al., 1994</td>
</tr>
<tr>
<td>Y3049</td>
<td>TAP-Src1</td>
<td>MATA TAP-Src1 ade2-1 his3-11,15 leu2-3,112 trp1-1, ura3-32</td>
<td>This study</td>
</tr>
<tr>
<td>Y3752</td>
<td>Src1-L-TAP</td>
<td>MATA SRC1 long-TAP::URA3 trp1 his3 leu2 ura3</td>
<td>This study</td>
</tr>
<tr>
<td>Y3658</td>
<td>Src1-S-TAP</td>
<td>MATA SRC1 short-TAP::URA3 leu2Δ1 trp1Δ63</td>
<td>This study</td>
</tr>
<tr>
<td>Y3735</td>
<td>TAP-Src1 TEV</td>
<td>MATA TAP-SRC1 PGALSV40NLS9mycTEV-2xSV40NLS::TRP1 leu2 ura3 his3</td>
<td>This study</td>
</tr>
<tr>
<td>Y3736</td>
<td>Src1-L-TAP TEV</td>
<td>MATA SRC1 long-TAP::URA3 GAL:SV40NLS9mycTEV-2xSV40NLS::TRP1 ura3-1 his3-11,15 leu2-3,112 ade2-1</td>
<td>This study</td>
</tr>
<tr>
<td>Y3737</td>
<td>Src1-S-TAP TEV</td>
<td>MATA SRC1 short-TAP::URA3 GAL:SV40NLS9mycTEV-2xSV40NLS::TRP1 ura3-1 his3-11,15 leu2-3,112 ade2-1</td>
<td>This study</td>
</tr>
<tr>
<td>Y3955</td>
<td>src1Δ TEV</td>
<td>MATA src1::natNT2 GAL:SV40NLS9mycTEV-2xSV40NLS::TRP1 ura3-1 his3-11,15 leu2-3,112 ade2-1</td>
<td>This study</td>
</tr>
<tr>
<td>Y4001</td>
<td>Nucloc2</td>
<td>MATA leu2Δ1 his3Δ200 ura3Δ851 nup49Δ1 hphNT1 tetR-GFP::LYS2 (pASZ11-mCherry-NOP1-GFP::NUP49ADE2)</td>
<td>Laboratory library; provided by A.B. Berger and O. Gadali</td>
</tr>
<tr>
<td>Y4124</td>
<td>PHO1-TelI</td>
<td>Nucloc2, YHR210::URA3::Δtel::Wlk (TelI*112)</td>
<td>This study</td>
</tr>
<tr>
<td>Strain</td>
<td>Plasmid</td>
<td>Description</td>
<td></td>
</tr>
<tr>
<td>--------</td>
<td>---------</td>
<td>-------------</td>
<td></td>
</tr>
<tr>
<td>Y4125</td>
<td>pRS314-SRC1</td>
<td>CEN, TRP1, PSRC1, TSRC1</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>SRC1 sequence containing the intron</td>
<td></td>
</tr>
<tr>
<td>Y4126</td>
<td>pRS314-SUB2</td>
<td>CEN, TRP1, PSUB2, TSUB2</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Laboratory library, provided by V. Zakian</td>
<td></td>
</tr>
<tr>
<td>Y4122</td>
<td>pRS314-sub2-85</td>
<td>CEN, TRP1, PSUB2, TSUB2</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Strässer and Hurt, 2001</td>
<td></td>
</tr>
<tr>
<td>Y4130</td>
<td>pRS315-THO2</td>
<td>CEN, LEU2, PTHO2, THO2</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Pirvat and Aguilera, 1998</td>
<td></td>
</tr>
<tr>
<td>Y4131</td>
<td>pRS315-SUS1</td>
<td>CEN, LEU2, PSUS1, TSUS1</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Rodriguez-Navarro et al., 2004</td>
<td></td>
</tr>
<tr>
<td>Y4132</td>
<td>pRS315-THP1</td>
<td>CEN, LEU2, PTHP1, THP1</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Laboratory library</td>
<td></td>
</tr>
<tr>
<td>Y4137</td>
<td>pRS315-SAC3</td>
<td>CEN, LEU2, PSAC3, TSAC3</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Fischer et al., 2002</td>
<td></td>
</tr>
<tr>
<td>Y3139</td>
<td>pRS315-MLP1</td>
<td>CEN, LEU2, PMLP1, TMLP1</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Laboratory library</td>
<td></td>
</tr>
<tr>
<td>Y4165</td>
<td>pRS315-GFP-SRC1L+</td>
<td>CEN, LEU2, PSRC1, TSRC1, N-terminal GFP tag</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>SRC1 sequence containing the intron</td>
<td></td>
</tr>
<tr>
<td>Y4166</td>
<td>pRS315-GFP-SRC1L-</td>
<td>CEN, LEU2, PSRC1, TSRC1, N-terminal GFP tag</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>aa 1–834, cDNA of SRC1Δ1 126bp intron</td>
<td></td>
</tr>
<tr>
<td>Y4165</td>
<td>pRS315-GFP-SRC1S</td>
<td>CEN, LEU2, PSRC1, TSRC1, N-terminal GFP tag</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>aa 1–688, cDNA of SRC1Δ1 130bp intron</td>
<td></td>
</tr>
<tr>
<td>Y4166</td>
<td>pRS315-GFP-SRC1ΔM1</td>
<td>CEN, LEU2, PSRC1, TSRC1, N-terminal GFP tag</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>aa 1–834 Δ454475; cDNA of SRC1Δ1 126bp intron</td>
<td></td>
</tr>
<tr>
<td>Y4166</td>
<td>pRS315-GFP-SRC1ΔM2</td>
<td>CEN, LEU2, PSRC1, TSRC1, N-terminal GFP tag</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>aa 1–834 Δ704–725; cDNA of SRC1Δ1 126bp intron</td>
<td></td>
</tr>
<tr>
<td>Y4166</td>
<td>pRS315-GFP-SRC1ΔN</td>
<td>CEN, LEU2, PSRC1, TSRC1, N-terminal GFP tag</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>aa 447–834; cDNA of SRC1Δ1 126bp intron</td>
<td></td>
</tr>
<tr>
<td>Y4166</td>
<td>pRS315-GFP-SRC1N</td>
<td>CEN, LEU2, PSRC1, TSRC1, N-terminal GFP tag</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>aa 1–453; cDNA of SRC1Δ1 126bp intron</td>
<td></td>
</tr>
</tbody>
</table>

This study
<table>
<thead>
<tr>
<th>Line</th>
<th>Description</th>
<th>Details</th>
<th>Source</th>
</tr>
</thead>
<tbody>
<tr>
<td>3604</td>
<td>pRS315-GFP-SRC1ΔC</td>
<td>CEN, LEU2, PSRC1, TSRC1, N-terminal GFP tag aa 1–737; cDNA of SRC1ΔI 126bp intron</td>
<td>This study</td>
</tr>
<tr>
<td>3605</td>
<td>pRS315-GFP-SRC1ΔNΔM1</td>
<td>CEN, LEU2, PSRC1, TSRC1, N-terminal GFP tag aa 481–834; cDNA of SRC1ΔI 126bp intron</td>
<td>This study</td>
</tr>
<tr>
<td>3606</td>
<td>pRS315-GFP-SRC1ΔNΔM</td>
<td>CEN, LEU2, PSRC1, TSRC1, N-terminal GFP tag aa 1–480; cDNA of SRC1ΔI 126bp intron</td>
<td>This study</td>
</tr>
<tr>
<td>3607</td>
<td>pRS315-GFP-SRC1ΔLEM</td>
<td>CEN, LEU2, PSRC1, TSRC1, N-terminal GFP tag aa 60–834; cDNA of SRC1ΔI 126bp intron</td>
<td>This study</td>
</tr>
<tr>
<td>3608</td>
<td>pRS315-SRC1ΔM2TAP</td>
<td>CEN, LEU2, PSRC1, TADH1, C-terminal TAP tag aa 1–834 Δ704–725; cDNA of SRC1ΔI 126bp intron</td>
<td>This study</td>
</tr>
<tr>
<td>3609</td>
<td>pNOPPATA1L-SRC1L + S</td>
<td>CEN, LEU2, PNOP1, TADH1, N-terminal ptA tag, SRC1 sequence containing the intron</td>
<td>This study</td>
</tr>
<tr>
<td>3610</td>
<td>pNOPPATA1L-SRC1ΔI</td>
<td>CEN, LEU2, PNOP1, TADH1, N-terminal ptA tag, SRC1 sequence containing the intron, Δaa 454–475</td>
<td>This study</td>
</tr>
<tr>
<td>3611</td>
<td>pNOPPATA1L-SRC1ΔM2</td>
<td>CEN, LEU2, PNOP1, TADH1, N-terminal ptA tag, SRC1 sequence containing the intron, Δaa 704–725</td>
<td>This study</td>
</tr>
<tr>
<td>3612</td>
<td>pNOPPATA1L-SRC1L + S</td>
<td>CEN, LEU2, PNOP1, TADH1, N-terminal ptA tag aa 1–834, cDNA of SRC1ΔI 126bp intron</td>
<td>This study</td>
</tr>
<tr>
<td>1732</td>
<td>pUN100-HPR1</td>
<td>CEN, LEU2, PHPR1, THR1</td>
<td>Laboratory library</td>
</tr>
<tr>
<td>2302</td>
<td>pRS316-MIp2</td>
<td>CEN, URA3, PMIp2, TMIp2</td>
<td>Laboratory library</td>
</tr>
<tr>
<td>220</td>
<td>pRS316-NUP133</td>
<td>CEN, URA3, PNU133, TNUP133</td>
<td>Laboratory library</td>
</tr>
<tr>
<td>3746</td>
<td>pAS211-SRC1</td>
<td>CEN, ADE2, PSRC1, TSRC1</td>
<td>This study</td>
</tr>
<tr>
<td>1107</td>
<td>pASZ11-GFP-NUP49</td>
<td>CEN, ADE2, PNU49, N-terminal GFP tag</td>
<td>Siniosoglu et al., 1998</td>
</tr>
<tr>
<td>2846</td>
<td>pH4467ΔThp1</td>
<td>CEN6 (instable), URA3, ADE3, PTHP1, TTHP1</td>
<td>This study</td>
</tr>
<tr>
<td>3615</td>
<td>pGAL::NLSmyc9TEV2xNLS</td>
<td>TEV protease; PGAL; N-terminal fused to SV40NLS and 9myc; C-terminal fused to 2xSV40NLS; integrative (TRP1)</td>
<td>Uhlmann et al., 2000</td>
</tr>
<tr>
<td>3647</td>
<td>p15816</td>
<td>112 TeLO repeats; natNT2</td>
<td>Gottschling et al., 1990</td>
</tr>
<tr>
<td>3647</td>
<td>pGC389</td>
<td>p15816-TeLO-interYAR046-YAR047; for deletion of telVv</td>
<td>Cabal et al., 2006</td>
</tr>
<tr>
<td>3647</td>
<td>pGC390</td>
<td>p15816-TeLO-interYHR210-FLO5; for deletion of telVv</td>
<td>This study</td>
</tr>
<tr>
<td>3647</td>
<td>pGC405</td>
<td>p15816-YAR069c-YHR214c-D; for [TeLO*112]-natNT2 insertion close to PHO11/PHO12</td>
<td>This study</td>
</tr>
<tr>
<td>3647</td>
<td>pGC20</td>
<td>p15816-interPHO84-GTR1; for [TeLO*112]-natNT2 insertion close to PHO84</td>
<td>Cabal et al., 2006</td>
</tr>
</tbody>
</table>
This study close to GIS1

References


Table S3. Real-time PCR primers used in this study

<table>
<thead>
<tr>
<th>Primer</th>
<th>Sequence</th>
</tr>
</thead>
<tbody>
<tr>
<td>ACT1_for</td>
<td>5'-CGCCTGGACTCAACAAG3'</td>
</tr>
<tr>
<td>ACT1_rev</td>
<td>5'-TCTGGAAGTCGTAGGATTITCAA3'</td>
</tr>
<tr>
<td>ACT1_probe</td>
<td>5'-AATGCAAAACGCTGCTCAATCTTCTCA3'</td>
</tr>
<tr>
<td>PHO11+PHO12_for</td>
<td>5'-GTTCCTGCTGACCTAATTGGAATGACTC3'</td>
</tr>
<tr>
<td>PHO11_rev</td>
<td>5'-GCATAGTCGTAGAAGACATTITTCACAG3'</td>
</tr>
<tr>
<td>PHO12_rev</td>
<td>5'-ATAAGAGCTGCTAATTTTTCAC3'</td>
</tr>
<tr>
<td>PHO11+PHO12_probe</td>
<td>5'-CGATGCTGCTGACCTAATTGGAATGACTC3'</td>
</tr>
<tr>
<td>PHO84_for</td>
<td>5'-CTCTGCTGCTGATGTGACTC3'</td>
</tr>
<tr>
<td>PHO84_rev</td>
<td>5'-AGAACGCGTACCAACCTAATAAG3'</td>
</tr>
</tbody>
</table>

The probes are labeled 5’ FAM and 3’ TAMRA. To distinguish between PHO11 and PHO12, which share 99% similarity, we used primers complementary to a sequence that differs in two base pairs between the two genes. Differences in Pho11_rev and Pho12_rev primers are underlined.