Phosphorylation of Elp1 by Hrr25 Is Required for Elongator-Dependent tRNA Modification in Yeast

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Abstract

Elongator is a conserved protein complex comprising six different polypeptides that has been ascribed a wide range of functions, but which is now known to be required for modification of uridine residues in the wobble position of a subset of tRNAs in yeast, plants, worms and mammals. In previous work, we showed that Elongator’s largest subunit (Elp1; also known as Iki3) was phosphorylated and implicated the yeast casein kinase I Hrr25 in Elongator function. Here we report identification of nine in vivo phosphorylation sites within Elp1 and show that four of these, clustered close to the Elp1 C-terminus and adjacent to a region that binds tRNA, are important for Elongator’s tRNA modification function. Hrr25 protein kinase directly modifies Elp1 on two sites (Ser-1198 and Ser-1202) and through analyzing non-phosphorylatable (alanine) and acidic, phosphomimic substitutions at Ser-1198, Ser-1202 and Ser-1209, we provide evidence that phosphorylation plays a positive role in the tRNA modification function of Elongator and may regulate the interaction of Elongator both with its accessory protein Kti12 and with Hrr25 kinase.

Introduction

Elongator is a conserved, multi-subunit protein complex containing six different polypeptides (Elp1-Elp6), first discovered in yeast in association with the elongating form of RNA polymerase II and initially proposed to play a role in transcriptional elongation [1,2]. Although Elongator is non-essential in yeast, knockout of the mouse IKKBAP gene encoding Elongator’s largest subunit leads to embryonic lethality and the protein is crucial for vascular and neural development [3]. The hereditary neuropathy Familial Dysautonomia results from human IKKBAP mutations, while mutations in other Elongator subunits have been associated with Amyotrophic Lateral Sclerosis [4] and Rolandic Epilepsy [3]. Elongator in Caenorhabditis elegans is also involved in neuronal function and development [6,7], while in plants it plays a role in proliferation during organ growth [8].

While it is therefore clear that Elongator is important for neuronal function in higher organisms [9,10], it has been proposed to have a bewildering range of seemingly unrelated functions. The Elp3 subunit of Elongator has a histone acetyltransferase (HAT) domain that can acetylate histones in vitro [11] and yeast Elongator mutants show changes in histone acetylation in vivo [12]. Elongator has also been proposed to acetylate α-tubulin, and the neuronal protein Bruchpilot [7,13,14] and has been implicated in paternal DNA demethylation in mouse zygotes [15]. In yeast, Elongator mutants adapt slowly to changing growth conditions, show sensitivity to high temperature, rapamycin, caffeine, hydroxyurea and various other chemical stressors, and are resistant to zymocin [16,17], a protein toxin secreted by the yeast Kluyveromyces lactis that kills other yeasts including Saccharomyces cerevisiae [18]. Yeast Elongator has also been implicated in transcriptional silencing, replication-coupled nucleosome assembly [17] and polarized secretion [19].

However, while these apparently diverse roles could imply that Elongator is multifunctional, work in yeast [20], C. elegans [6], plants [21] and mammals [22] has demonstrated that many if not all of these proposed functions reflect a primary role for Elongator in tRNA modification, specifically in the addition of mcm2 (5-methoxycarbonylmethyl) and mcm5 (5-carbamoylmethyl) groups to uridine when present at the ‘wobble’ position (U34) in tRNA anticodons. Eleven out of 13 such tRNAs in yeast contain either mcm2U, mcm5U or 5-methoxycarbonylmethyl-2-thiouridine.

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Data Availability: The authors confirm that all data underlying the findings are fully available without restriction. All relevant data are within the paper and its Supporting information files.

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tRNAs molecules function as adapters in protein synthesis, bringing amino acids to the ribosome and reading the genetic code through codon-anticodon base pairing. When the tRNA contains a uridine residue in the “wobble position” of its anticodon, which base-pairs with purine residues in the third position of a cognate codon, it is almost always chemically modified and modification is required for efficient decoding. In eukaryotic cells, these wobble uridine modifications require a conserved protein complex called Elongator. Our work shows that Elp1, Elongator’s largest subunit, is phosphorylated on several sites. By blocking phosphorylation at these positions using mutations, we identified four phosphorylation sites that are important for Elongator’s role in tRNA modification. We have also shown that Hr25 protein kinase, a member of the casein kinase 1 (CK1) family, is responsible for modification of two of the sites that are important for Elongator function. Phosphorylation appears to affect interaction of the Elongator complex both with its kinase (Hr25) and with Kt12, an accessory protein previously implicated in Elongator function. Our studies imply that Elp1 phosphorylation plays a positive role in Elongator-mediated tRNA modification and raise the possibility that wobble uridine modification may be regulated, representing a potential translational control mechanism.

Author Summary

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Identification of in vivo phosphorylation sites in Elp1 and other Elongator subunits

To identify sites of phosphorylation in Elongator we affinity isolated the complex from yeast cells expressing a TAP-tagged version of Elp1. Tryptic digests of the affinity-purified material were subjected to phosphopeptide enrichment involving a two-step procedure using Hypersep SCX and TiO₂, followed by tandem mass spectrometry to locate sites of phosphorylation. To maximize the chance of detecting phosphopeptides, in addition to isolating Elongator from wild-type yeast cells we also prepared and analyzed the complex from a sit4Δ strain, in which lack of Sit4 phosphatase leads to Elp1 hyperphosphorylation [37,39]. We also analyzed Elongator prepared from a Kit12Δ mutant in which Elongator is hypophosphorylated [37,39] in case additional phosphorylation sites could also be detected under these circumstances. In this way we identified eight phosphorylation sites in Elp1, one each in Elp2 and Elp4 and two in Elp5 (Table 1). With the exception of Ser-222 in Elp4 [40], all of these sites are novel and were not identified in any of the recent proteome-wide phosphoproteomics studies. Since we previously showed that Elp1 phosphorylation state changes are associated with altered Elongator function [37,39], we focused on the phosphorylation sites we identified in Elp1. SI Fig. shows representative MS/MS spectra providing evidence for the eight sites identified in Elp1, which with one exception were identified within monophosphorylated peptides. Apart from Ser-1205/Thr-1206, where phosphorylation of the two sites cannot be unambiguously distinguished from the mass spectra, all phosphorylation sites can be identified with high confidence. All Elp1 sites identified in Elongator isolated from either the sit4Δ mutant or the kit12Δ mutant were also found in Elp1 from the wild-type strain (Table 1).

Since analysis of our mass spectrometry data also provided additional weak evidence for phosphorylation on Elp1 Ser-1209, we raised a phosphospecific antibody against a synthetic peptide carrying phosphate on this residue to examine whether it was a genuine phosphorylation site. As shown in Fig. 1B, when used to probe Elp1 by Western blotting, this phosphospecific antibody gave a strong signal that was lost when Ser-1209 was mutated to

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alanine, thereby demonstrating its phosphospecificity. Thus Elp1 Ser-1209 represents a ninth Elp1 phosphorylation site (Table 1).

Table 1. Summary of phosphorylation sites identified in Elp1 and other Elongator subunits.

| Elp1          | Elp2     | Elp4     | Elp5     |
|---------------|----------|----------|----------|
| Ser-529 (wt)  | Ser-1198 (wt, k) | Ser-492 (s) | Ser-3 (w, s) |
| Ser-539 (wt)  | Ser-1202 (wt) | Ser-222A (s) | Ser-4 (w, s) |
| Ser-551 (wt)  | Ser-1205/Thr-1206 (wt, s) | Ser-1209² | Ser-4 (w, s) |
| Ser-636 (wt, k) | Ser-1209² | Ser-828 (wt, s) | Ser-3 (w, s) |

¹For sites identified by mass spectrometry: wt, identified in Elongator from wild-type yeast cells; k, identified in Elongator from a kbt124 strain; s, identified in Elongator from a sit4D mutant.
²Phosphorylation on these two adjacent sites cannot be unambiguously distinguished.
³Demonstrated using a phosphospecific antibody; all other sites identified by mass spectrometry.
⁴Previously identified by Soulard et al.[40].

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Phenotype of Elp1 C-terminal domain phosphorylation site mutants

To investigate the Elp1 C-terminal phosphorylation region, additional combinations of alanine, aspartate or glutamate substitutions at the phosphorylation sites were generated. In addition to using the eclipse assay, we monitored tolerance to intracellular expression of the zymocin tRNase (γ) subunit from the galactose-inducible GAL1 promoter as a more quantitative measure of zymocin resistance [43]. Since zymocin sensitivity provides a readout largely for the mcm5 modification state of just tRNA\(^{\text{3OS}}\)(UUC), we also monitored Elongator function via efficiency of ochre suppression mediated by SUP4, a tRNA\(^{\text{1OS}}\)(UUA) that requires Elongator-dependent wobble uridine modification for efficient ochre (UAA) codon readthrough [20,24]. This involved single copy integration of a plasmid that carried both SUP4 and a ura3 ochre allele [32], such that suppression (and hence Elongator’s tRNA modification function) could be monitored by growth in the absence of uracil.

Fig. 2 shows that the S1209A substitution conferred complete resistance to intracellular expression of the zymocin γ subunit and greatly reduced SUP4-dependent ura3\(^{\text{3OS}}\) suppression, in each case conferring a phenotype comparable to that observed upon complete loss of Elongator function (elp1D). In contrast, an S1209D substitution showed considerable restoration of function, while the equivalent glutamate substitution (potentially a poorer mimic of phosphoserine due to its longer side chain) was less effective. The regain of functionality caused by the phosphomimic aspartate substitution therefore provides evidence that phosphorylation of Ser-1209 acts positively for Elongator function. Although the double S1198A S1202A substitution mutant conferred strong zymocin resistance it showed considerable residual Elongator function in the SUP4 assay, allowing for growth in the absence of uracil similar to that shown by the ELP1 wild-type control. However, concurrent glutamate substitutions at both positions largely restored zymocin sensitivity, supporting the notion that phosphorylation of these two sites acts positively for Elongator function. S1198A S1202A in combination with T1204A S1205A T1206A, a triple alanine substitution that on its own had essentially no effect on Elongator function in any of the assays, conferred stronger zymocin resistance than S1198A S1202A alone and dramatically reduced SUP4-dependent suppression efficiency, indicative of an additive Elongator defect in the quintuple mutant. The triple T1204A S1205A T1206A mutation was used here because phosphorylation at Ser-1205/Thr-1206 could not be unambiguously distinguished (see above). Mutants where acidic residues substituted combinations of these five positions, either...
alone or in combination with alanine substitutions, indicated that Elongator functionality was not greatly affected by acidic substitutions, although the T1204E S1205D T1206E triple substitution improved Elongator function when combined with the S1198A S1202A double mutant (seen most clearly by the eclipse assay). While the difference in the severity of phenotype observed between the K. lactis colony placed at the edge of each patch of cells being tested indicates zymocin resistance and hence loss of Elongator function. (D) Conservation of the phosphorylation sites identified in the Elp1 C-terminal domain. The full sequences were first aligned using T-Coffee [83] but only the relevant detail from the alignment is shown. NCBI database accessions for the aligned sequences are *Saccharomyces cerevisiae* (NP_013488), *Ashbya gossypii* (NP_984907), *Candida albicans* (XP_710040), *Mus musculus* (NP_080355), *Homo sapiens* (NP_003631), *Schizosaccharomyces pombe* (NP_595335), *Arabidopsis thaliana* (NP_196872). (E) C-terminal phosphorylated region of *S. cerevisiae* Elp1 showing the downstream region with a repeating pattern of serine and threonine residues. Mapped phosphorylation sites are underlined and shown in bold, potential phosphorylation sites downstream of the mapped sites are underlined.

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S1198 S1202A T1204A S1205A T1206A strains showed almost no mcm 5U or ncm 5U, consistent with the major defect in Elongator-dependent wobble uridine modification indicated by the phenotypic assays (Fig. 2). The S1198A S1202A double mutant showed reduced levels of mcm 5U and ncm 5U consistent with a less severe Elongator defect, while strains with the S1198E S1202E double or S1209D single phosphomimic alleles, or with the triple T1204A S1205A T1206A allele, had levels of mcm 5U and ncm 5U intermediate between those of the S1198A S1202A mutant and ELP1 wild-type. Taken together, the profiles for the ncm5 and mcm5 modification nucleosides are consistent with our phenotypic analyses (Fig. 2) and our data therefore support the notion that phosphorylation of Ser-1209 plays a major, positive role in Elongator-dependent tRNA modification and that phosphorylation at Ser-1198, Ser-1202 and Ser-1205/Thr-1206 makes a similar but partly redundant contribution to Elongator activity.

Phosphorylation site mutations in Elp1 do not prevent Elongator assembly but lead to changes in association with Kti12 and with Hrr25 kinase

Phosphorylation site mutations that alter Elongator functionality could in principle do so by affecting assembly of the holo-Elongator complex, either because they interfere with assembly or because phosphorylation of these sites could regulate assembly of the Elongator complex. When myc-tagged Elp2 was used to immunoprecipitate Elongator from cell extracts, elp1 mutant derivatives carrying mutations in ELP1 as shown. (A) Key to substitutions present in the different alleles tested (wild-type sequence at the top). (B) Zymocin sensitivity measured by eclipse assay (see Fig. 1 legend), with strains being tested at either 1.0 or 0.1 OD600/ml following growth in SCD-Leu medium. (C) Zymocin sensitivity monitored by intracellular expression of the zymocin γ subunit. Cells were grown on SCD-Leu-Ura to select for both plasmids, diluted to 1.0 OD600/ml and then equivalent 10-fold dilutions plated on YPD agar (to control for cell viability) and YP agar containing 2% raffinose and 2% galactose to induce expression of zymocin γ-toxin from pAE1. (D) Cells were grown in SCD-Leu (to select for YCplac111 and its derivatives), diluted to 1.0 OD600/ml and then equivalent 10-fold dilutions plated on SCD-Leu (to control for cell viability) and SCD-Leu-Ura (to assess efficiency of SUP4-dependent ura3+ suppression).

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Fig. 2. Effect on Elongator function of phosphorylation site mutations in the C-terminal domain of Elp1. All strains were based on WAY034 (elp1Δ; panel B), YRDS119 (elp1Δ [pAE1]; panel C) or SBY138 (elp1Δ his3Δ1::pSB3; panel D) transformed with YCplac111, YCplac111-ELP1-6HA (wild-type) or mutant derivatives carrying mutations in ELP1 as shown. (A) Key to substitutions present in the different alleles tested (wild-type sequence at the top). (B) Zymocin sensitivity measured by eclipse assay (see Fig. 1 legend), with strains being tested at either 1.0 or 0.1 OD600/ml following growth in SCD-Leu medium. (C) Zymocin sensitivity monitored by intracellular expression of the zymocin γ subunit. Cells were grown on SCD-Leu-Ura to select for both plasmids, diluted to 1.0 OD600/ml and then equivalent 10-fold dilutions plated on YPD agar (to control for cell viability) and YP agar containing 2% raffinose and 2% galactose to induce expression of zymocin γ-toxin from pAE1. (D) Cells were grown in SCD-Leu (to select for YCplac111 and its derivatives), diluted to 1.0 OD600/ml and then equivalent 10-fold dilutions plated on SCD-Leu (to control for cell viability) and SCD-Leu-Ura (to assess efficiency of SUP4-dependent ura3+ suppression).

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Since Elongator interacts with both its accessory factor Kti12 and with Hrr25 kinase and because Elongator function requires both Kti12 and Hrr25 kinase activity [16,37,45,46], we next used co-immunoprecipitation to examine the effect of selected phosphorylation site mutations on Elongator’s association with Kti12 and Hrr25. Elongator complex in which Elp1 carries the double S1198A S1202A mutation showed reduced interaction with Hrr25 that was not seen with the corresponding phosphomimic allele (S1198E S1202E), but combining S1198A S1202A with T1204A S1205A T1206A did not further reduce interaction with Hrr25 (Fig. 5) despite the greater loss of Elongator function in the quintuple mutant. Conversely, mutation of Elp1 Ser-1209 to...
alanine led to enhanced interaction between Elongator and Hrr25 (Fig. 5). In all ELP1 mutants tested, Elongator retained its ability to interact with Kti12, but the elp1-S1209A allele also led to enhanced Kti12 interaction (Fig. 6A). We examined the effect of the Elp1 S1209A substitution on Kti12 association in more detail by tagging the genomic copy of ELP1 with GFP and carefully quantitating the recovery of HA-tagged Kti12 in strains expressing either the wild-type or mutant Elp1-GFP fusions. This confirmed that the S1209A mutation leads to enhanced Kti12 association: in comparison with wild-type Elp1, Elp1-S1209A was reproducibly associated with approximately twice as much Kti12 (Fig. 6B, C). Thus the S1198A S1202A and S1209A mutations have opposite effects on association with Hrr25 and the 1209A mutation enhances association with Kti12, suggesting that phosphorylation at these sites affects the interaction between Elongator and key proteins required for its functionality.

Hrr25 directly phosphorylates Elp1 on functionally important sites

We next wished to identify the protein kinase(s) that mediate modification of the phosphorylation sites in the Elp1 carboxy-terminal domain that are important for Elongator function. To take an unbiased approach, we screened a library of 119 GST-protein kinase fusions [47] for their ability to phosphorylate Elp1 in vitro using TAP-purified Elongator as a substrate. Purified Elongator showed significant background phosphorylation specifically on Elp1 when incubated with [γ-32P]ATP in the absence of added kinase. However, five protein kinases were identified that could clearly phosphorylate Elp1 in vitro (Hrr25, Yck1, Yck2, Yck3 and Hal5; S2A Fig.). Given that Hrr25 interacts with Elongator [37,49] as seen in Fig. 5 and because Elp1 is already known to become hypophosphorylated in a hrr25 mutant [37], it is an excellent candidate for an in vivo Elp1 kinase. Hrr25, Yck1, Yck2 and Yck3 all belong to the casein kinase I (CKI) family of protein kinases [50], but while hrr25 mutants show clear defects in Elongator function [37,38], Yck1, Yck2 and Yck3 are membrane associated via lipid modification [51,52] and do not confer detectable zymocin resistance when the corresponding genes are deleted [38]. Thus we consider it unlikely that Yck1-3 mediate functionally important phosphorylation events on Elp1 in vitro and that they were identified in our in vitro screen due to shared substrate specificity with the related Hrr25 CKI isoform. Preliminary analysis also failed to generate data supporting a role for Hal5 in Elongator function (S2B-S2C Fig.).

In further support of a role for Hrr25 as a direct Elp1 kinase, we next showed that the Elp1 kinase activity present in affinity-isolated Elongator preparations was due to Hrr25. This made use of a yeast strain dependent on an ‘analog-sensitive’ HRR25 allele (hrr25-I82G) in which the mutant Hrr25 kinase has acquired the capacity to be inhibited specifically by addition of the ATP analogs 1NM-PP1 or 3MB-PP1 [53]. When Elongator was isolated from the hrr25-I82G strain, the Elp1 phosphorylation observed upon incubation of the isolated complex with [γ-32P]ATP was blocked by addition of either 1NM-PP1 or 3MB-PP1 (Fig. 7A). This was in contrast to Elongator isolated from a control strain expressing wild-type HRR25, phosphorylation of which was refractory to these inhibitors. In fact the hrr25-I82G strain became zymocin resistant when grown in the presence of INM-PP1, emphasizing the positive role of Hrr25 kinase in Elongator’s tRNA modification in vivo (Fig. 7B) and consistent with Elongator-minus phenotypes of hrr25 mutants [37,38].

To determine whether Hrr25 phosphorylation of Elp1 occurred at any of the functionally relevant sites that we had identified, Elongator containing wild-type or mutant forms of Elp1 was purified and tested for incorporation of phosphate from [γ-32P]ATP both in the absence and presence of recombinant Hrr25. Fig. 7C shows that even when reactions were supplemented with Hrr25, Elp1 in which the C-terminal region had been deleted was not phosphorylated. Elongator in which both Elp1 Ser-1198 and Ser-1202 had been replaced by either alanine or glutamate showed negligible incorporation of radiolabel in comparison with the Elp1 in wild-type Elongator. In contrast, Elongator in which Elp1 Ser-1209 was substituted with either alanine or aspartate showed high levels of 32P incorporation into Elp1 in the presence of recombinant Hrr25. Since HRR25 is an essential gene we were unable to examine Ser-1209 phosphorylation with the anti-phosphoserine-1209 antibody in the complete absence of Hrr25. However, in the kinase-dead hrr25-3 mutant, which shows loss of Elongator function and absence of the phosphorylated isoform detected in wild-type cells by gel shift assay [37], a similar level of Ser-1209 phosphorylation to that seen in a wild-type strain was observed (S3 Fig.). Although more complex interpretations are possible, these observations are therefore most simply explained if Ser-1198 and Ser-1202 are downstream targets of Hrr25.

Fig. 3. Quantitation of modified U34 nucleosides in tRNA from selected elp1 mutants. Content of ncm5U and mcm5U in small RNA samples from the indicated yeast strains. In each case, A (modified U)/A (U) indicates that the modified nucleoside signal was normalized using the total uridine content to allow comparison of the different samples. Inset: chemical structures of uridine and its Elongator-dependent ncm5U and mcm5U derivatives.

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major sites of Hrr25 phosphorylation and Ser-1209 is not a major site for modification by the kinase. In the absence of added Hrr25, Elongator complex containing Elp1-S1209A still showed high levels of Elp1 phosphorylation that were not seen with the corresponding aspartate substitution or when Elp1 was wild-type, although after addition of recombinant Hrr25 the level of Elp1 phosphorylation of the Elp1-S1209A and Elp1-S1209D complexes was similar as noted above (Fig. 7C). These data are consistent with the enhanced interaction of Elp1-S1209A-containing Elongator with Hrr25 that was seen by co-immune precipitation (Fig. 5).

To complement these experiments, the ability of recombinant Hrr25 in vitro and analyzed by mass spectrometry, three types of phosphopeptide were detected: monophosphorylated peptide modified on either Ser-1198 (S4A Fig.) or Ser-1202 (S4B Fig.), and diphosphorylated peptide modified on both these residues (S4C Fig.). These data therefore confirm the identity of Hrr25 as a protein kinase that can directly phosphorylate Elp1 on serine residues that are important for Elongator function, and are consistent with a model in which phosphorylation at one of these two sites may then favor phosphorylation of the second site. When all phosphorylatable residues apart from Ser-1198 and Ser-1202 were changed to alanines, dual phosphorylation on Ser-1198 and Ser-1202 was still seen, indicating that phosphorylation of these two positions was not dependent on phosphorylation of Ser-1205 or any other downstream residues.

When the 1193-1213 peptide was incubated with Hrr25 together with γ-[32P]ATP, it readily incorporated radiolabelled phosphate (Fig. 8B and C; peptide 89). Although the related peptide in which all serine and threonine residues apart from Ser-1198 and Ser-1202 had been substituted by alanines was still an excellent Hrr25 substrate (Fig. 8C; peptide 92), incorporation of 32P at later times was reduced in comparison (Fig. 8B), consistent with the possibility of additional phosphorylation to the right of Ser-1202. Additional single alanine substitutions at either Ser-1198 or Ser-1202 in peptide 92 greatly reduced but did not completely prevent phosphorylation (Fig. 8C, peptides 1001, 1002). However, replacement of both Ser-1198 and Ser-1202 by either alanines or glutamates completely blocked phosphorylation of the peptide despite the presence of the five serine and threonine

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**Fig. 4.** Elongator assembly is unaffected by a range of phosphorylation site mutations in Elp1. *elp1Δ* yeast strains expressing myc-tagged ELP2 and either HA-tagged ELP3 (A) or ELP5 (B) were transformed with either empty vector or plasmids expressing the indicated ELP1 alleles (Table S2). Elp2-myc was immunoprecipitated from extracts and immunoprecipitates were examined by Western blotting with anti-myc and anti-HA antibodies to detect co-immunoprecipitated Elp1, Elp3 and Elp5 as indicated. Note that Elp1 runs as a doublet as observed previously [84].

**Fig. 5.** Phosphorylation site mutations in Elp1 lead to changes in Hrr25 association with Elongator. *elp1Δ ELP2-myc, KT12-HA* yeast strains were transformed with either empty vector or plasmids expressing the indicated HAα-tagged ELP1 alleles (Table S2). Elp2-myc was immunoprecipitated from extracts and immunoprecipitates were examined by Western blotting with anti-myc to detect Elp2, anti-HA antibodies to detect co-immunoprecipitated Elp1 and anti-Hrr25 (to detect co-immunoprecipitated Hrr25) as indicated. Western blots of the extracts before immunoprecipitation (lower panels) confirm similar levels of each protein in the different strains.

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residues downstream (peptides 90 and 91). Another peptide encompassing Elp1 residues 1207-1229 was not phosphorylated at all following incubation with Hrr25 (Fig. 8D, peptide 95), supporting the notion that neither Ser-1209 nor any of the repeating threonine and serine shown in Fig. 1E can be directly phosphorylated by Hrr25 kinase. These data are therefore consistent with interdependence of Ser-1198 and Ser-1202 phosphorylation by Hrr25 and absence of Hrr25 phosphorylation on the downstream serines and threonines.

Since phosphorylation of specific Ser or Thr residues by CKIs such as Hrr25 is often primed by phosphorylation at another Ser or Thr residue 2-4 positions upstream [54], it was possible that Hrr25 might phosphorylate Ser-1205 once Ser-1202 had been modified, and that phosphorylation at Ser-1205 might then promote modification of Ser-1209. However, phosphomimetic glutamate substitutions at 1198 and 1202, which supported essentially normal Elongator functionality in vivo (Fig. 2), also prevented phosphorylation of the 1193-1213 peptide on any other downstream site. Furthermore, phosphorylation of a related peptide (Elp1 1192-1212; Fig. 8E), which was also a good in vitro substrate for Hrr25, was also completely blocked for Hrr25 phosphorylation when synthesized with phosphoserine at the two positions corresponding to Elp1 Ser-1198 and Ser-1202. This again supports the model that Ser-1198 and Ser-1202 are the only major sites of Hrr25 phosphorylation and also indicates that it is unlikely that efficient priming of phosphorylation by Hrr25 on downstream residues, such as Ser-1205 and Ser-1209, occurs as a result of the upstream phosphorylation events at Ser-1198 and Ser-1202. Thus although these experiments do not as yet fully explain all the intricacies of Hrr25 phosphorylation in this region, taken together they nonetheless demonstrate that Hrr25 phosphorylates Elp1 directly on Ser-1198 and Ser-1202, two serine residues that we identified as functionally relevant in vivo phosphorylation sites.

Discussion

Identification of functionally relevant phosphorylation sites in Elp1

We have identified nine phosphorylation sites on the Elp1 subunit of yeast Elongator complex and based on the phenotypes...
Fig. 7. Purified Elongator shows Hrr25-dependent phosphorylation that requires Ser-1198 and Ser-1202. (A) Elongator was purified using TAP-tagged Elp3 from either a HRR25 wild-type or a hrr25-I82G analog-sensitive mutant strain and supplemented with [γ-32P]ATP to examine endogenous Elp1 phosphorylation in the presence of DMSO (drug vehicle), 1NM-PP1 or 3MB-PP1. Phosphorylated Elp1 was visualized by autoradiography following SDS-PAGE. (B) Equivalent 10-fold serial dilutions of HRR25 ELP1, hrr25-I82G ELP1 and HRR25 elp1Δ yeast strains were plated out on YPAD agar with or without 1% (v/v) zymocin in the presence or absence of 10 μM 1NM-PP1 or an equivalent volume of DMSO (drug vehicle) as indicated and photographed after 2 days’ growth at 30°C. (C) Elongator was prepared as in (A) from elp1Δ ELP3-TAP strains carrying plasmids encoding the indicated ELP1 alleles and incubated with [γ-32P]ATP in the presence or absence of added recombinant Hrr25 (100 ng). Phosphorylated Elp1 was visualized by autoradiography following SDS-PAGE. Western blotting to detect Elp3-TAP and Elp1-6HA demonstrates equivalent recovery of Elongator from each strain.

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Fig. 8. In vitro phosphorylation of peptides corresponding to the Elp1 C-terminal region by Hrr25. (A) Sequences of synthetic peptides used as substrates shown in alignment with Elp1 residues 1191-1230. Lower-case bold letters represent alanine and/or glutamate substitutions relative to the native Elp1 sequence. (B) Time course of Hrr25-dependent incorporation of 32P from [γ-32P]ATP into Peptides 89 and 92. (C-E) Incorporation of 32P from [γ-32P]ATP into peptides as assayed by SDS-PAGE and autoradiography. Note that in (D) and (E) two panels are shown in each case because some unnecessary intervening lanes were removed when preparing the figure. * indicates autophosphorylated recombinant Hrr25.

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of non-phosphorylatable and phosphomimic mutations, provide evidence that phosphorylation on four sites near the Elp1 C-terminus (Ser-1198, Ser-1202, Ser-1205/Thr-1206 and Ser-1209) plays a positive role in Elongator function. Previously, we showed that a slid phosphate mutant trapped Elp1 in a slower-migrating, hyperphosphorylated form whereas hrr25 kinase mutations led to presence of just a faster-migrating, hypophosphorylated Elp1 isoform [37,39]. Both types of mutation cause loss of Elongator function [37,39], suggesting that dynamic phosphorylation and dephosphorylation of Elp1 is needed in functional Elongator and predicting that mimicking constitutive phosphorylation on at least some of the sites we have identified should be inhibitory. It is therefore surprising that all of the phosphomimic alleles we tested conferred significant Elongator function. Possibly the acidic substitutions do not fully mimic constitutive phosphorylation and thereby allow for substantial residual function, even though constitutive phosphorylation might inhibit Elongator. Thus while our alanine substitution mutants lend support to the idea that phosphorylation at the mapped sites functions positively for Elongator activity, we cannot rule out a requirement for dynamic phosphorylation/dephosphorylation at the sites we have identified. It is also possible that additional, inhibitory phosphorylation sites exist that were not found in our study. For example, although we could not detect phosphorylation of Thr-1212 in vivo or in vitro, both alanine and glutamate substitutions at this site caused severe loss of Elongator function (S5 Fig.), consistent with the notion that dynamic phosphorylation and dephosphorylation of this site could be important if it is phosphorylated in vivo.

**Hrr25 directly phosphorylates Elp1 on sites important for Elongator function**

Despite conducting an unbiased, ‘kinome-wide’ screen to identify kinases responsible for Elp1 C-terminal domain phosphorylation, Hrr25, the yeast CKI that associates with Elongator and has already been implicated in Elongator function [37,48,49] was the sole convincing candidate to be identified. Through several lines of evidence, we have now established that Hrr25 directly phosphorylates Elp1 on two in vivo phosphorylation sites that we have mapped and shown to be relevant for Elongator function: Ser-1198 and Ser-1202. Thus purified Elongator shows Hrr25-dependent phosphorylation that requires the presence of these two residues, while peptides derived from the Elp1 C-terminal region show direct, interdependent phosphorylation by recombinant Hrr25 on Ser-1198 and Ser-1202. Conversely, we found no evidence for direct phosphorylation of Ser-1209 by Hrr25 using three different synthetic peptides containing Ser-1209, or using purified Elongator complex in which Ser-1198 and Ser-1202 were mutated but Ser-1209 was intact.

Two types of consensus phosphorylation site have been proposed for CKIs: pS [X]_{1-3} [ST] and [DE]_{2-4} [X]_{0-2} [ST], where pS indicates an upstream phosphoserine residue that is needed to prime phosphorylation at the downstream Ser or Thr (shown in bold and underlined) but which can be substituted by an acidic patch in the second class of motif [34]. Ser-1198 conforms to the latter type of motif and is closely related to other mapped Hrr25 phosphorylation sites [55-57] in yeast proteins (S6 Fig.). In contrast, Ser-1202 matches the former motif, suggesting that priming-independent phosphorylation at Ser-1198 by Hrr25 might then prime phosphorylation by Hrr25 at Ser-1202 and predicting that Ser-1198 should still be efficiently phosphorylated when Ser-1202 is replaced by alanine. However, the latter mutation largely blocked phosphorylation at Ser-1198 in our in vitro peptide kinase assays and instead we observed interdependent phosphorylation at these two residues. Furthermore, replacement of one of the two upstream aspartate residues with alanine did not obviously interfere with phosphorylation of Ser-1198 and Ser-1202. Thus although the sites are direct targets of Hrr25 and match the accepted consensus for CKI phosphorylation, dependency of Ser-1202 phosphorylation on prior Ser-1198 modification remains to be demonstrated. Similarly, Ser-1205 and Ser-1209 both fit the pS [X]_{1-3} [ST] consensus and might be modified once Ser-1202 phosphorylation has occurred, and yet we could not demonstrate Hrr25-dependent incorporation of phosphate at these sites even using a peptide where the upstream sites were already phosphorylated. Given the known substrate specificity of CKIs, it is therefore surprising that we can find no evidence for such priming of phosphorylation at Ser-1205 and Ser-1209 following modification of Ser1202. Furthermore, since the repeating pattern of phosphorylatable residues shown in Fig. 1E also fits the pS [X]_{1-3} [ST] motif we are even more surprised that we failed to detect phosphorylation of these sites either in vivo or in vitro.

In spite of the strong evidence we have provided that Hrr25 is a direct Elp1 kinase, inability to detect direct phosphorylation of either the key residue Ser-1209 or of Ser-1205/Thr-1206 by Hrr25 implies that at least one additional Elp1 kinase is involved in Elp1 phosphorylation. While these sites might become better substrates for Hrr25 in the context of fully assembled Elongator complex rather than within synthetic peptides, the fact that we detect normal levels of Ser-1209 phosphorylation in a hrr25 mutant that is defective for Elongator function strongly supports the involvement of a different kinase at Ser-1209.

**Elp1 phosphorylation may regulate Elongator’s interaction with Kti12 and Hrr25**

Although all phosphorylation site mutants examined showed essentially normal assembly of the Elongator complex and retained the ability to interact with the accessory protein Kti12, the S1209A mutation enhanced association of Kti12 with Elongator in comparison with cells expressing either the wild-type protein or the S1198A S1202A mutant. Kti12 stoichiometry is important – either too much or too little interferes with Elongator function [39,43,45] and thus Ser-1209 phosphorylation may regulate Elongator’s interaction with its accessory protein. Furthermore, the S1209A mutation caused increased Hrr25 association with Elongator whereas the double S1198A 1202A mutation led to reduced Hrr25 association. Thus Hrr25 not only directly phosphorylates Elp1 but may also regulate its own interaction with Elongator through doing so. Although the S1209A single and S1198A S1202A T1204A S1205S T1206A quintuple mutants both conferred strong loss of Elongator function, their differing effects on Hrr25 association implies that they are defective for different reasons – phosphorylation at S1198A S1202A may stabilize Hrr25 binding whereas phosphorylation at Ser-1209 may be required to displace the kinase. Since the interaction between Hrr25 and Elongator is dependent on Kti12 [37] it is also possible that the enhanced interaction of both proteins with Elongator seen when Elp1 cannot be phosphorylated on Ser-1209 are directly related.

Intriguingly, enhanced interaction of both Kti12 and Hrr25 with the Elongator complex is also seen in hrr25 mutants that cause Elp1 phosphorylation defects and zymocin resistance [37 and S7 Fig.]. However, these properties of hrr25 mutants are more similar to those of the elp1-S1209A mutant, which is mutated at a site apparently not directly phosphorylated by Hrr25, rather than mirroring the properties of the elp1-S1198A, S1202A mutant that removes the only sites in Elp1 that we have shown to
be direct Hrr25 targets. This suggests an as yet undiscovered connection between phosphorylation at Ser-1209 and Hrr25 kinase. Regardless of this, our data nonetheless indicate that Elp1 phosphorylation by Hrr25 and other kinases could modulate the interaction between Elongator and Kti12.

What type of role does Elp1 phosphorylation play in Elongator function?

Although there is clearly more to learn about the role of phosphorylation in Elongator function, two types of model can be proposed. On the one hand, phosphorylation might regulate Elongator, turning its wobble uridine modification activity up or down in response to growth conditions and the demand for protein synthesis, or perhaps in response to cellular stresses. Given that the translation of some mRNAs is particularly dependent on wobble base modification and Elongator function [58-60] and that tRNA modifications (including Elongator-dependent ones) oscillate during the cell cycle and in response to stress signals [61,62], this raises the interesting possibility that Elongator may be part of a translational control mechanism functioning through tRNA modification. Our proposal that Elp1 phosphorylation operates in a largely positive sense for Elongator, based on the properties of the phosphorylation sites that we have examined, is consistent with such a regulatory role. The Elp1 kinase Hrr25 is needed for a wide range of cellular functions [49,56,57,63-65] that do not appear to provide a clear insight into the signals that might regulate Elp1 phosphorylation. However, the Hrr25 requirement for full functionality of two different components of the translation machinery - wobble uridine-containing tRNAs [37, this work] and ribosomes [49,57] - might reflect a role in regulation of the cell’s capacity for protein synthesis. In addition, both hrr25 mutants [66] and Elongator-deficient yeast [17] are sensitive to hydroxyurea and methyl methanesulfonate. Since efficient translation of the ribonucleotide reductase gene RNR1 requires mcm2-modified tRNAs [62,67], Elp1 phosphorylation may also be linked to the known role of Hrr25 in expression of genes such as RNR2 and RNR3 in response to DNA damage [66].

Alternatively, Elp1 phosphorylation might be dynamic, with sequential phosphorylation and dephosphorylation of specific residues driving the biochemical mechanism through which Elongator carries out the tRNA modification reactions. Such a dynamic role might operate through modulation of Elongator’s interaction with factors such as Kti12 or with tRNA. It is intriguing in this context that the C-terminal phosphorylated region in Elp1 is immediately adjacent to a basic region that binds specifically to tRNA and that mutation of the tRNA binding domain leads to reduced interaction with Kti12 [Fig. 6C and ref. 32]. Thus phosphorylation of Elp1 could potentially influence how Elp1 interacts with tRNA, perhaps through interaction between the acidic phosphate groups and the basic residues present in the tRNA binding domain.

In conclusion, while there is still much to learn about Elp1 phosphorylation and its involvement in Elongator function, our work clearly demonstrates the importance of at least four in vivo phosphorylation sites in the C-terminal domain of Elp1 for Elongator-dependent tRNA wobble uridine modification, shows that Hrr25 kinase directly modifies two of these sites and provides evidence that phosphorylation regulates the association between Elongator and both its accessory protein Kti12 and its kinase Hrr25.

Materials and Methods

Yeast strains and plasmids

Basic yeast methods, growth media, and routine recombinant DNA methodology were performed as previously described [68,69]. All plasmids used in this study are listed in Table S2 and yeast strains are listed in Table S3. To generate yeast strains dependent on wild-type or mutant forms of Elp1, the Elp1 gene was deleted from BY4741 and WAY008 (ELP3-TAP) using pFA6a-KanMX6 and pCORE-UH deletion cassettes to obtain the elp1Δ knockout mutant strains WAP034 and WAY037, respectively. Wild-type or mutant ELP1 variants carried on the low-copy plasmid YCplac111-ELP1-6HA (Table S2) were then introduced into these elp1Δ strains as the sole source of Elp1. The mutant versions were made by either site directed mutagenesis (Qiagen QuickChange) using YCplac111-ELP1-6HA as template, or by replacing the relevant region using standard cloning procedures and synthetic DNA carrying the desired mutations. The majority of plasmids made using the latter approach were produced by DNA2.0, Inc (Menlo Park, CA, USA). All elp1Δ mutants made by site-directed mutagenesis were verified by DNA sequencing of the entire ELP1 ORF plus approximately 200 base pairs upstream and downstream. When mutations were introduced by cloning, the region that had been replaced was sequenced to exclude the possibility of gene synthesis errors. The wild-type and mutant versions of YCplac111-ELP1-6HA were transformed into WAY034, WAY037 or elp1Δ::KanMX6 ELP2- myc3 strains carrying additional epitope-tagged components (Table S3) for phenotypic screening, Elongator-complex purification or Western blot analysis, respectively. pFA6a-CTAP1- HIS3MX6 was made by ligating the 1322 bp BglII-Pmel fragment from pFA6a-3HA-HIS3MX6 to the 3603 bp BglII-Pmel fragment from pFA6a-CTAP4-NatMX6 [70] such that the NatMX marker was replaced by HIS3MX. Using this template, a C-terminal TAP tag was added to ELP1 in RL-343-F0 and RL-343-F1 (Table S3) using standard one-step tagging methodology [71]. stI4Δ::LEU2 and kit12A::LEU2 knockouts were made as previously described [39,43], as was introduction of elp1Δ::KanMX6 in strains used for co-immune precipitation with myc-tagged Elp2 [16]. Construction of ELP1- HA6::KTRIP1 utilized the pYM3 tagging plasmid and S2/S3 primer set described by Knop et al. [72].

Purification of TAP-tagged Elongator

For phosphorylation site mapping, Elongator complexes were purified from WAY009, WAY010, WAY011, WAY-HPIT and WAY-Has-PIT (Table S3) by two-step tandem affinity purification of Elp1-TAP from 2-12 liter cultures in YPAD medium grown to OD600 1.0-1.5, as described previously [73]. All purifications were done in the presence of PhosSTOP and Complete protease inhibitor cocktail (Roche). Elongator complexes in which Elp1 contained phosphorylation site mutations were similarly purified from WAY037 (elp1Δ::pCORE-UH ELP3-TAP::HIS3MX) or elp1Δ::KanMX6 or its mutant derivatives (Table S2), growing cells as above but in SCD-Leu to select for retention of the plasmid.

Phosphorylation site mapping

Elongator protein preparations were digested in solution with Trypsin (Trypsin Gold, mass spectrometry grade; Promega V5280). The resulting peptides were cleaned over Hypersep C18 columns (Thermo Scientific) to remove buffer contaminants, eluting the peptides in 70% acetonitrile, 0.1% trifluoroacetic acid (TFA). Phosphopeptide enrichment was done in a two-step procedure using a Hypersep SCX column (Thermo Scientific) followed by TiO2 enrichment of mono-phosphorylated peptides from the SCX flow-through. SCX binding and washing buffer contained 10 mM KH2PO4, 25% acetonitrile (pH 3.0) and the elution buffer contained 10 mM KH2PO4, 25% acetonitrile, 350 mM KCl (pH 3.0). The flow-through was reduced in volume.
to ~100 μl, supplemented with 100 μl 80% acetonitrile/2% TFA containing 200 mg/ml 2,3-dihydroxybenzoic acid (DHB), pH 2.0, then the sample applied to 5 micron TitanSphere TiO₂ beads (GL Sciences) and rotated gently at room temperature for 1 h. The beads were washed twice with 80% acetonitrile/2% TFA, 200 mg/ml DHB (pH 2.0), then 3 times with 80% acetonitrile/2% TFA (pH 2.0) before eluting with 60 μl 100 mM NH₄OH (pH 11.0) and then supplementing with 2 μl of 100% formic acid. All phosphopeptide fractions were cleaned over C18 and submitted for mass spectrometry in 0.1% TFA.

Phosphopeptide fractions were analyzed by LC-MS/MS using a Dionex U3000 system (Dionex California) with a PepMap C18 column coupled to either an Orbitrap XL or Orbitrap Velos (Thermo Fisher Scientific). Peptides were eluted using a 45 min 5%-90% acetonitrile gradient, sequencing the top 5 most intense ions with the following settings: CID, FTMS 335-1800 Da, 60,000 resolution, MS/MS charge state 1+ accepted. Peak picking, recalibration and peptide mass fingerprinting was done using MaxQuant software [74,75], searching masses against the Saccharomyces Genome Database orf_trans_all database [January 3rd, 2010 release] with 10 ppm MS error, ≤ 2 missed cleavages, Trypsin/P enzyme, variable modifications set to Acetyl (protein N-term), Oxidation (M), Phospho (ST) and Phospho (Y) and a fixed modification of Carbomidomethyl (C). The MS/MS tolerance was set to 0.5 Da and the false discovery rate for site, protein and modification of Carbomidomethyl (C). The MS/MS tolerance was set to 0.5 Da and the false discovery rate for site, protein and modification of Carbomidomethyl (C). The MS/MS tolerance was set to 0.5 Da and the false discovery rate for site, protein and modification of Carbomidomethyl (C). The MS/MS tolerance was set to 0.5 Da and the false discovery rate for site, protein and modification of Carbomidomethyl (C).

Relative quantitation of tRNA wobble uridine modifications by HPLC-DAD-MS/MS analysis

elp1A yeast strains containing YCPlac111-ELP1-HA6 or its mutant derivatives were grown in SCD-Leu to select for the plasmid and then total tRNA was prepared by RNA extraction and LiCl precipitation as described previously [32]. Prior to LC-MS/MS/MS analysis, 3 μg of each tRNA sample were digested into nucleosides by incubation at 37°C for 2 h in the presence of 1/10 volume of 10× nucleases P1 buffer (0.2 M NH₄OAc pH 5.0, ZnCl₂ 0.2 mM, 0.3 U nuclease P1 (Sigma Aldrich, Munich, Germany) and 0.1 U snake venom phosphodiesterase (Worthington, Lakewood, USA). Next, 1/10 volume of 10× fast alkaline phosphatase buffer and 1 U fast alkaline phosphatase (Fermentas, St. Leon-Roth, Germany) were added, and samples were incubated for additional 60 min at 37°C. The digested tRNA samples were analyzed on an Agilent 1260 HPLC series equipped with a diode array detector (DAD) and a triple quadrupole mass spectrometer (Agilent 6460). A Synergy Fusion RP column (4 μm particle size, 80 Å pore size, 250 mm length, 2 mm inner diameter) from Phenomenex (Aschaffenburg, Germany) was used at 35°C column temperature. The solvents consisted of 5 mM ammonium acetate buffer adjusted to pH 5.3 using acetic acid (solvent A) and pure acetonitrile (solvent B). The elution was performed at a flow rate of 0.35 ml/min using a linear gradient from 0% to 8% solvent B at 10 min, 40% solvent B at 20 min and 0% solvent B at 23 min. For additional 7 min, the column was rinsed with 100% solvent A to restore the initial conditions. Prior to entering the mass spectrometer, the effluent from the column was measured photometrically at 254 nm by the DAD for detection of the 4 canonical nucleosides. The triple quadruple mass spectrometer, equipped with an electrospray ion source (Agilent Jet Stream), was run at the following ESI parameters: gas temperature 350°C, gas (N₂) flow 8 L/min, nebulizer pressure 50 psi, sheath gas (N₂) temperature 350°C, sheath gas (N₂) flow 12 L/min and capillary voltage 3000 V. The MS was operated in positive ion mode using Agilent MassHunter software and modified nucleosides were monitored by multiple reaction monitoring (dynamic MRM mode). Identification of mcG-U and mcGmU peaks was performed as described previously [77]. Peak areas were determined employing Agilent MassHunter Qualitative Analysis Software. In the case of the major nucleosides, peak areas were extracted from the recorded UV chromatograms in order to avoid saturation of the mass signals. For inter-sample comparability of the detected modifications, the peak areas of the modified nucleosides were normalized to the UV peak area of uridine.

Phenotypic assays

To test the effect of extracellular zymocin on wild-type and mutant strains, killer-eclipse assays were performed as described previously [76] using the K. lactis zymocin producer strain NCYC1368 and YPD plates prepared using Kobe I agar (Roth 5210). To test the effect of intracellular expression of zymocin’s γ subunit on growth, wild-type and mutant elp1 strains were transformed with pAE1, which expresses the γ subunit from the galactose-inducible GAL1 promoter [43]. The response to zymocin γ-toxin induction was monitored on galactose plates after 3-4 days at 30°C. SUP4 suppression efficiency was measured following integration of pSB3 in single copy at the his3Δ10 locus and then monitoring growth on SCD-Leu-Ura after 3 days at 30°C as described previously [32]. To test the effect of Hrr25 inhibition on Elongator function, strains dependent on w.t. or analog-sensitive Hrr25 were grown overnight in YPAD medium, diluted to 1.0 A₆₀₀/ml and then 10-fold dilutions were spotted onto YPAD agar with or without 1%/v/v zymocin and containing 10 μM 1NM-PP1 or an equivalent volume of DMSO as drug vehicle control. Growth was documented after 2 days growth at 30°C. Zymocin was prepared using cell-free culture medium from K. lactis NCYC1368 which had been grown at 30°C for 2 days in YPAD medium, by 50-fold concentration using Amicon Ultra-15 Centrifugal Filter Units (Millipore) followed by sterilization by filtration.

Western blotting and co-immune precipitation

Detection of tagged proteins used anti-TAP (Thermo Scientific, CAB1001), anti-myc and anti-HA antibodies (Roche) and was performed as previously described [16,45]. Protein concentrations were determined using Quick StartTM Bradford Protein Assay (BioRAD) [78] and checked with anti-Pik1 antibodies recognizing yeast Pik1 (1:50,000, kindly provided by Dr. J. Heinisch) or anti-Cdc19 serum (1:10,000, kindly provided by Dr. J. Thorner) so as...
to ensure equivalent loadings. For detection of the Hrr25 kinase in total yeast extracts and in immune precipitates, a generic ant-Hrr25 antibody [64] was used (1:10,000 dilution). Antibody cross-linking to Dynabeads M-270 Epoxy (Invitrogen), preparation of protein extracts and immune precipitation were performed according to the manufacturer’s instruction and as described previously [16,79]. In general, 1 μg of antibody coupled to Dynabeads was used per 1 mg total cell extract in B60 buffer.

Identification of protein kinases that phosphorylate Elp1

All 119 GST-protein kinases constructs in the library described by Zhu et al. [47] were expressed in yeast and purified as described previously [80]. Yeast Elongator complex was isolated as described in the main paper and used as a substrate for the GST-kinases, which were initially assayed in 23 pools of 5 kinases under similar conditions to those described below. Reactions were separated by SDS-PAGE and radiolabelled Elp1 detected by autoradiography. Eight out of the 23 pools showed phosphorylation occurred above the background level observed due to co-purification of Hrr25 with Elongator (see Results). From these eight pools, 40 individual kinases were individually screened for their ability to phosphorylate Elp1. Reactions in which Elongator was omitted was controlled for any radiolabeled bands that co-migrated with Elp1 and were therefore due to kinase autophosphorylation or co-purified kinase substrates.

In vitro protein kinase assays

E. coli BL21 (DE3) pLysS (Novagen) was transformed with pTrcHis-HRR25 for overexpression and purification of His6-Hrr25 using HisPur cobalt resin (Thermo Scientific) according to the manufacturer’s instructions. To analyze Elp1 phosphorylation in vitro the HisPur cobalt resin (Thermo Scientific) according to the manufacturer’s instructions. To analyze Elp1 phosphorylation using HisPur cobalt resin (Thermo Scientific) according to the manufacturer’s instructions.

Phosphorylation reactions were stopped by the addition of NuPAGE 4× LDS Sample Buffer (Life Technologies) and heated for 5 min at 95°C. Samples were run on a NuPAGE (4-12%) polyacrylamide/Bis-Tris SDS gel (Life Technologies) at 200 V for 1 h. Gels were stained with BioSafe Coomassie (BioRad), dried and subjected to autoradiography. To examine the effect of chemical inactivation of yeast Hrr25 on Elp1 phosphorylation in purified Elongator complex, Elongator was purified as above from strains WAY-H-PIT (wild-type HRR25) and WAY-Has-P1T (allele-sensitive hrr25). Elongator was omitted from the reaction, and phosphorylated peptides were visualized by autoradiography. For mass spectrometric analysis of phosphorylated peptides, 150 μl reactions were conducted as above but omitting the radiolabelled ATP.

Supporting Information

S1 Fig. Mapping in vivo phosphorylation sites in yeast Elp1. Representative MS/MS spectra are shown in which diagnostic B (green) and Y (blue) ions are mapped onto the corresponding peptide sequence. S(Ph), phosphoserine; M(Ox), oxidized methionine. (A) Phosphorylation on Ser-529 within the Elp1 peptide 521-531, parent MH+ ion m/z 675.29. (B) Dual phosphorylation on Ser-559 and Ser-551 within the Elp1 peptide 552-552, parent MH+ ion m/z 817.41. (C) Phosphorylation on Ser-636 within the Elp1 peptide 624-652, parent MH+ ion m/z 1063.46. (D) Phosphorylation on Ser-828 within the Elp1 peptide 828-844, parent MH+ ion m/z 1051.44. (E) Phosphorylation on Ser-1198 within the Elp1 peptide 1180-1213, parent MH+ ion m/z 1314.56. (F) Phosphorylation on Ser-1202 within the Elp1 peptide 1180-1213, parent MH+ ion m/z 1314.56. This spectrum represents a mixture of monophosphorylated isoforms of the peptide: * denotes peaks consistent with Ser-1198 phosphorylation, # denotes peaks consistent with Ser-1205 phosphorylation and ■ indicates an ion consistent with Thr-1206 phosphorylation. (G) Phosphorylation on Ser-1205 or Thr-1206 within the Elp1 peptide 1180-1213, parent MH+ ion m/z 1314.56. (TIF)

S2 Fig. Identification of Elp1 kinases. (A) Phosphorylation of yeast Elongator by yeast protein kinases. Elongator was purified from yeast using TAP-tagged Elp3 and incorporation of 32P from [γ-32P]ATP to GST-kinase fusions identified from the GST-kinase library of Zhu et al. [47]. Background incorporation of 32P in the absence of added kinase is due to co-purification of Hrr25 with affinity-purified Elongator complex as shown in the main paper. Elp1 runs as a doublet, the faster-migrating component of which is truncated at the amino-terminus [84]. (B) Relatedness of Hal5 to Kqs8 and Sat4. A standard BLAST search (http://blast.ncbi.nlm.nih.gov/Blast.cgi) of the yeast proteome with Hal5 using the default parameters indicates that Kqs8 and Sat4 are its closest relatives within the yeast kinome. (C) Deletion of HAL5, either alone or in combination with KKS8 and/or SAT4, fails to confer zymocin sensitivity by eclipse assay. (TIF)

S3 Fig. Elp1 shows similar levels of Ser-1209 phosphorylation in wild-type and hrr25-3 mutant strains. Elp1 phosphorylation in samples of extract containing equivalent amounts of protein was monitored by Western blot analysis using the phosphoSer-1209 phosphospecific antibody. (TIF)

S4 Fig. Mapping of phosphorylation sites following in vitro phosphorylation of a peptide corresponding to Elp1 residues 1193–1213. MS/MS spectra are shown in which diagnostic B (green) and Y (blue) ions are mapped onto the corresponding peptide sequence. S(Ph), phosphoserine. (A) Phosphorylation on Ser-1198, parent MH+ ion m/z 889.08 corresponding to monophosphorylated peptide. * denotes peaks consistent with phosphorylation on Ser-1198, suggesting that this is a mixed spectrum. (B) Phosphorylation on Ser-1202, parent MH+ ion m/z 889.08 corresponding to monophosphorylated peptide. # indicates peaks consistent with phosphorylation of Ser-1198, suggesting that this is a mixed spectrum. (C) Phosphorylation on Ser-1198 and Ser-1202, parent MH+ ion m/z 925.73 corresponding to diphosphorylated peptide. * denotes peaks consistent with one
phosphate on Ser-1202 and a second phosphate to the right, most likely on Ser-1205.

(THF)

**S3 Fig.** Neither alanine nor glutamate substitutions at Elp1 Thr-1212 support Elongator function. All strains were based on WAY034 transformed with YCplac111 (empty vector; *elp1Δ*), YCplac111-ELP1-6HA (*ELP1* wild-type) or its Thr-1212 mutant derivatives as indicated. Zymocin sensitivity was measured by eclipse assay [see Fig. 1], testing samples of cells at either 1.0 or 0.1 OD₅₅₀/ml following growth in SCD-yeast medium to select for the plasmids.

(THF)

**S6 Fig.** Alignment of Hrr25 phosphorylation sites. Alignment of the Elp1 Ser-1198 phosphorylation site with mapped Hrr25 phosphorylation sites in Tifs [57], Ash1 [56] and Man1 [55]. Sites known to be phosphorylated directly by Hrr25 are shown in red. Residues surrounding the mapped phosphorylation sites are highlighted in yellow (acidic), grey (small hydrophobic) or green (polar). In each case the phosphorylated residue is preceded by two acidic and one small hydrophobic residue and followed by two small hydrophobic residues.

(THF)

**S7 Fig.** *hrr25-2* leads to increased association of Hrr25 and Kti12 with Elongator. Elp1-HA was immunoprecipitated from extracts of the indicated *HRR25* and *hrr25-2* strains and immunoprecipitates were examined by Western blotting with anti-HA, anti-myc and anti-Hrr25 antibodies to detect immunoprecipitated Elp1 and co-immunoprecipitated Kti12 and Hrr25 respectively. The ELP1 untagged control confirms that recovery of Kti12 and Hrr25 is dependent on Elp1.

(THF)

**S1 Table** Summary of Zymocin phenotype monitored by Eclipse assay associated with different mutant *elp1* alleles. WAY034 (*elp1Δ*) and WAY031 (*elp1Δ sit4Δ*) were transformed with YCplac111, YCplac111-ELP1-6HA or with mutant derivatives of the latter plasmid and monitored by Eclipse assay for Zymocin sensitivity/resistance. R, Zymocin resistant; S, Zymocin sensitive; R/S, intermediate.

(PDF)

**S2 Table** Plasmids used or generated in this study.

(PDF)

**S3 Table** Yeast strains used or generated in this study.

(PDF)

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**Author Contributions**

Conceived and designed the experiments: WAF MH RS MJRS. Performed the experiments: WAF DJ RDS KL7 VS AH SH MH. Analyzed the data: WAF DJ RDS KL7 SH MH RS MJRS. Contributed reagents/materials/analysis tools: WAF DJ RDS KL7 SH MH RS MJRS. Wrote the paper: WA RS MJRS.

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