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To cite this version:
Marie-Claire Daugeron, Tineke L. Lenstra, Martina Frizzarin, Basma El Yacoubi, Xipeng Liu, et al.. Gcn4 misregulation reveals a direct role for the evolutionary conserved EKC/KEOPS in the t6A modification of tRNAs. Nucleic Acids Research, Oxford University Press, 2011, 39 (14), pp.6148–6160. 10.1093/nar/gkr178. pasteur-01404019

HAL Id: pasteur-01404019
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Submitted on 28 Nov 2016

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Gcn4 misregulation reveals a direct role for the evolutionary conserved EKC/KEOPS in the t6A modification of tRNAs

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Received November 16, 2010; Revised March 10, 2011; Accepted March 11, 2011

ABSTRACT

The EKC/KEOPS complex is universally conserved in Archaea and Eukarya and has been implicated in several cellular processes, including transcription, telomere homeostasis and genomic instability. However, the molecular function of the complex has remained elusive so far. We analyzed the transcriptome of EKC/KEOPS mutants and observed a specific profile that is highly enriched in targets of the Gcn4p transcriptional activator. GCN4 expression was found to be activated at the translational level in mutants via the defective recognition of the inhibitory upstream ORFs (uORFs) present in its leader. We show that EKC/KEOPS mutants are defective for the N6-threonylcarbamoyl adenosine modification at position 37 (t6A37) of tRNAs decoding ANN codons, which affects initiation at the inhibitory uORFs and provokes Gcn4 derepression. Structural modeling reveals similarities between Kae1 and bacterial enzymes involved in carbamoylation reactions analogous to t6A37 formation, supporting a direct role for the EKC in tRNA modification. These findings are further supported by strong genetic interactions of EKC mutants with a translation initiation factor and with threonine biosynthesis genes. Overall, our data provide a novel twist to understanding the primary function of the EKC/KEOPS and its impact on several essential cellular functions like transcription and telomere homeostasis.

INTRODUCTION

Subunits of the Endopeptidase-like Kinase Chromatin (EKC)-associated complex [also called Kinase putative Endopeptidase and Other Proteins of Small size (KEOPS)] were first identified by genetic analyses (1,2). We isolated Pcc1p, a protein of small size with no predicted functional domains in a search for suppressors of a splicing defect and later demonstrated that its function is not related to splicing (2). Additional genetic, biochemical and cell biology experiments led to the definition of a five subunits complex (Pcc1p, Pcc2p/Gon7p, Cgi121p, Bud32p and Kae1p) the integrity of which is required for normal induction of pheromone- and galactose-responsive genes.
Components of the EKC/KEOPS have been found to associate with transcribed chromatin and the biochemical defect in transcription activation of GAL genes was mapped to the recruitment of transcriptional co-activators (2). A parallel genome-wide screen led to the identification of the cgl121 null mutant as a suppressor of the telomere-capping defect of ade13-1 cells. Deletion of CGL121 or other EKC/KEOPS subunits suppresses telomeres resection and single strand DNA production due to defective protection by the capping complex (1). All the tested EKC/KEOPS mutants have short telomeres in an otherwise wild-type context, an observation seemingly at odds with the telomere protective effect in the context of telomeres capping deficiencies. The molecular mechanism linking the involvement of the complex in transcription and telomere maintenance is unclear but it has been suggested to be related to effects on chromatin structure (1,2). Finally, DNA instability has also been associated with mutation of EKC/KEOPS components and the bacterial counterparts of Kae1p [our unpublished data; (3)].

Of the five subunits, the only ones containing predicted functional domains are Kae1p and Bud32p. Kae1p was originally proposed to be a candidate endopeptidase, based on sequence similarities with bacterial zinc-dependent metalloproteases of the M22 family (4). However, such a function could never be demonstrated and it does not appear to be compatible with the structure of Kae1p homologues from Archae and Eukaryotes (5–7). Structural studies (5,7) have revealed the existence of an actin-like ATPase domain and a metal (iron) binding site. It has been shown that archaeal PaKae1 has DNA binding capabilities and apurinic site endonuclease activities (7).

Bud32p is an atypical kinase that lacks some of the canonical domains required for substrate recognition (5,6). In vitro phosphorylation assays have demonstrated that this protein is a functional kinase that is also responsible for its autophosphorylation (8). It has been suggested that the association of Kae1p with Bud32p inhibits the phosphorylation activity of the latter (6).

One of the most striking features of EKC/KEOPS is its conservation and Kae1p belongs to the category of universal protein families (9). With the exception of Pcc2p/Gon7p that is restricted to Fungi, the whole complex is conserved in Archaea and Eukaryas (2). With roughly 60% identity from Saccharomyces cerevisiae to man, Kae1p is the most conserved subunit and is also the most ancient member of the complex, with clear homologues (YgiD) present in Bacillus (3). Widespread conservation is however at odd with a primary function that would be limited to features restricted to eukaryotes, such as telomere maintenance or the establishment of specific chromatin structures. Therefore, in spite of the large number of phenotypes described for the mutants of the EKC/KEOPS complex, the molecular function of this set of proteins remains elusive. To address this important question we undertook genetic and transcriptome analyses of EKC/KEOPS mutants. We describe unexpected links to transcription and tRNA metabolism as well as a specific transcriptome profile characterized by the inappropriate expression of amino acid biosynthesis genes. These phenotypes are causally linked by the translational misregulation of GCN4, encoding the transcriptional activator responsible for regulation of most amino acid biosynthesis genes. Phylogenetic and biochemical analyses strongly suggest a direct role of the EKC/KEOPS complex in N6-threonylcarbamoyl adenosine modification at position 37 (t6A37) of the anticodon stem-loop of all tRNAs decoding ANN codons. We show that the molecular mechanism underlying the translational defects relates to defective t6A37 formation, which affects the codon-anticodon interaction at the first AUG codon. The universality of this modification fits the evolutionary conservation of Kae1p, providing a novel twist to understanding the impact of this important set of proteins on several essential cellular functions.

**MATERIAL AND METHODS**

**Manipulation of yeast strains and genetic screens**

All the strains used in this study are listed in Supplementary Table S1. Genetic crossing and analysis were performed with standard methods. The synthetic lethal screen with the pcc1-4 mutant was based on a colorimetric assay to detect the ability of double mutant cells to grow in the absence of a wild-type copy of PCC1. Briefly, an ade8Δ, ade2-1 W303 strain containing an integrated pcc1-4 allele was transformed with a centromeric plasmid containing a wild-type copy of PCC1, ADE8 and the URA3 markers (pCM188-PCC1-ADE8). This strain is red because of the accumulation of a pigment in ade2-1/ADE8 cells. Random loss of the plasmid leads to the appearance of white sectors (the phenotype of ade2-1/ade8Δ cells). Cells were UV-mutagenized to 10% viability and mutants that failed to lose the plasmid (i.e. that were unable to live without a wt copy of PCC1) were retained for further analysis. The tfi5-K55E mutation was isolated with standard genetic methods. Genomewide genetic interaction mapping (GIM) was performed as described (10).

**Microarray experiments**

Expression-profiling of mutant strains was performed using two-channel, long oligomer arrays as previously described (11). In brief, each strain was cultured twice and amplified, labeled cRNA was hybridized twice in dye-swap replicate against a single common reference wt cRNA, yielding a total of four estimates of changes in gene expression for each strain versus wt. Normalization was performed using print-tip Loess followed by dye-bias correction as described (11). Statistical analysis was performed by LIMMA (12) versus a collection of wt profiles hybridized versus the same common reference. A gene was regarded as significantly changed if the P-value was <0.01 and the fold-change >1.5. Microarray data have been submitted to the ArrayExpress database (http://www.ebi.ac.uk/microarray) under accession number E-TABM-1070 (username Reviewer_E-TABM-1070, password 1283942451014)
Polysome profile analysis

Polysomes extracts were prepared and analyzed by sedimentation in 10–50% sucrose gradients as previously described (13). Gradients were fractionated with an ISCO gradient fractionator and the absorbance profiles at 254 nm were monitored within an ISCO UA-5 absorbance monitor. Cellular extract for ribosomal subunit quantification were prepared and analyzed on low-Mg++ sucrose gradients as described (14).

β-Galactosidase assays with Gcn4-LacZ reporters

Wild-type or mutant strains containing the reporter plasmids p180, p227 or p4164 (kind gift of A. G. Hinnebusch) were grown at 30°C in CSM-Ura to OD600 0.3–0.6. Cells pellets were resuspended in 500 μl of buffer Z (100 mM HNaPO4, pH 7.2; 10 mM KCl; 1 mM MgSO4; 50 mM β-mercaptoethanol) and permeabilized by the addition of 200 μl of ether. After ether evaporation, 100 μl of ONPG (4 mg/ml in buffer Z) was added and the reaction incubated at 30°C until the development of a yellow color. The reaction was stopped by the addition of 250 μl of 1M Na2CO3 and the OD420 of the supernatants was measured. β-Galactosidase activity was expressed in arbitrary units (similar to Miller units) as follow: activity = 1000 × OD420 / [OD600 of cultured sample × volume of cultured sample (ml) × reaction time (s)].

t6A analysis

Bulk tRNA was prepared, hydrolyzed, and analyzed by liquid chromatography-tandem mass spectrometry (LC–MS–MS) as described in (15). Levels of t6A were measured by integrating the peak area from the extraction ion chromatograms. The ratios of Ψ-modified base/m1G was used to normalize for tRNA concentration across samples. Levels for mutant strains were expressed relatively to wild-type levels. The MS–MS fragmentation data were also used to confirm the presence of t6A.

RESULTS

Genetic interactions of EKC/KEOPS mutants

To elucidate the molecular mechanism of action of the complex we analyzed the genetic interaction profile of kae1 and pcc1 EKC/KEOPS mutants. A genetic screen for additional mutations preventing growth of thermosensitive pcc1-4 cells at permissive temperature resulted, as expected, in the isolation of alleles of two other subunits of the complex, Kae1p and Pcc2/Gon7. We also isolated a mutant allele of the translation initiation factor Tif5p/eIF5, an essential GTpase activating protein (GAP) (Figure 1A). This factor is required for stimulating GTP hydrolysis by eIF2 in the 43S pre-initiation complex (PIC) upon AUG codon–anticodon base-pairing, thereby allowing the transition from initiation to translation elongation. Virtue of its physical interactions with other translation factors, eIF5 is also required for the formation of the PIC and the selection of the starting AUG codon [for recent reviews see (16,17)]. The sequence of the mutated allele revealed that a single A to G change is present in the coding sequence, leading to mutation of K55 to E. Accordingly, we named this allele if5-K55E.

To extend these results we performed a genomewide genetic interaction mapping (GIM) (10). For better sensitivity and specificity we used a hypomorphic PCC1 allele that contains a substitution of its terminator with a CYC1 terminator (unpublished data). To validate the results obtained and to filter out false positives, the deletion strains identified in the screen were individually crossed to the EKC/KEOPS mutant kae1-18 that is more severely affected than PCC1-CYC1ter cells. Strong negative genetic interaction leading to inviability at 30°C was observed for thr1 and thr4 deletion mutants (Figure 1B). Thr1p and Thr4p are the enzymes required for the conversion of homoserine to threonine in the two last steps of the biosynthesis of this amino acid. Genetic interactions (although not leading to lethality) were also confirmed with PUS1, required for pseudouridylation of tRNAs and U2 snRNA, and, to a lesser extent, HOM2, involved in an earlier step of threonine biosynthesis (Figure 1C) as well as TRM1 encoding the enzyme responsible for N2,N2-dimethylguanosine modification in both cytoplasmic and mitochondrial tRNAs (data not shown).

Together, these results suggested that the EKC/KEOPS might be involved in translation and/or tRNA metabolism. The stronger genetic interaction with genes involved in threonine biosynthesis also suggests a specific relationship between this pathway and the function of the EKC/KEOPS (see below).

A peculiar transcriptome profile in EKC complex mutants

The unexpected results of the genetic screens prompted us to perform transcriptome analysis of EKC/KEOPS mutants. We expected that a transcriptome profile similar to that of mutants altering known cellular functions would reveal important aspects of the role of the complex. Since most EKC null mutants are extremely sick (2), we integrated less affected thermosensitive mutants of Kae1p (kae1-18), Pcc1p (pcc1-4) and Pcc2/Gon7 (pcc2-ts1) in the BY4742 strain and analyzed the transcriptome of these cells using DNA microarrays at the semi-permissive temperature of 30°C (‘Methods’ section). Isogenic bud32 and cgl121 null mutants from the yeast deletion strain collection were also profiled. In several of the independent isolates analyzed we noticed the presence of whole chromosomal duplications, suggesting the occurrence of genomic instability in these mutants. Only euploid isolates were considered for further analyses. Since different isolates of bud32 null mutants, obtained from different sources, all contained chromosomal duplications, we did not further pursue analysis of this mutant.

The EKC/KEOPS mutants show significant changes in gene expression, with a total of 579 genes upregulated and 282 genes downregulated collectively in all single mutants compared with wt (P < 0.01, fold-change >1.5, Figure 2 and Supplementary Figure S1). As is often observed for mutants derived from subunits of the same protein complex (18), the profiles of the EKC/KEOPS mutants are related to each other (Figure 2A). The kae1-18 and pcc1-4 mutants show the strongest expression signature,
consisting of essentially the same set of mRNA expression changes, with the pcc2 and cgil21 mutants affecting a subset of the same genes, in most cases to a lower degree (Figure 2 and Supplementary Figure S1). Strikingly, the expression-profile signature of the EKC/KEOPS mutants is highly enriched for upregulated genes involved in amino acid biosynthesis (Gene Ontology biological process, cellular amino acid biosynthetic process, enrichment \( P = 4 \times 10^{-34} \)), as well as related metabolic pathways (Supplementary Table S2). Genes encoding enzymes involved in most cellular amino acid biosynthesis pathways are upregulated in the pcc1-4 mutant (Supplementary Figures S1 and S2). These results were independently confirmed by quantitative RT-PCR analysis using a subset of the upregulated genes. Moreover, we showed that upregulation was suppressed by expression of the corresponding wt gene in mutant cells, demonstrating that the expression profile is indeed a consequence of the EKC/KEOPS mutations (data not shown). Surprisingly, such a massive upregulation of amino acid biosynthesis genes was not observed in the collection of over 1500 mutants for which the transcriptome was analyzed during growth in identical conditions (F.C.P.H., unpublished results), indicating that the EKC/KEOPS mutant profiles are highly specific, at least among the mutants analyzed.

To investigate how the genetic interactions revealed by the genome-wide screen relate to the expression-signature of EKC/KEOPS mutants, the thr1, thr4, trm1 and pus1 null mutants were also expression-profiled. Interestingly, the gene expression changes observed in thr1 and thr4 mutants overlap highly significantly with pcc1 and kael mutants (\( P \) ranges from \( 10^{-21} \) to \( 7 \times 10^{-41} \)), and includes genes involved in amino acid biosynthesis (Figure 2 and Supplementary Figure S1). The expression-profile of trm1 and pus1 cells do not show a significant overlap with EKC/KEOPS or thr1/4 mutants (data not shown), suggesting the existence of different mechanisms underlying the observed genetic interactions.
Translation of the transcription activator Gcn4 is de-repressed in EKC mutants

Amino acid biosynthesis genes are generally activated in conditions of amino acids starvation. We observed upregulation of these genes in EKC mutants despite growth in the presence of normal levels of amino acids, suggesting that integrity of the EKC is required for their regulation. The transcriptional activator Gcn4p is involved in the activation of amino acid biosynthesis genes [for a review see (19)] and its binding site targets (20) are highly enriched in the EKC/KEOPS mutant signature genes ($P = 3.25 \times 10^{-11}$ and $2 \times 10^{-10}$, respectively, for kae1 and pcc1 mutants, Figure 2B) as well as in the signature genes shared by EKC/KEOPS mutants and thr1/4 mutants ($P$ ranging from $4 \times 10^{-11}$ to $2 \times 10^{-13}$).
Defective recognition of initiator AUG codons in EKC and tif5-K55E mutants

Activation of GCN4 translation in pcel and kael mutant cells could be due to defective recognition of the inhibitory uORFs 2–4 after post-termination re-initiation (see above), to activation of the Gcn2 kinase pathway, for instance as a consequence of some molecular event that creates or mimics amino acid shortage or both. To assess whether defective recognition of AUG codons occurs in EKC/KEOPS mutants, we employed a construct containing an out of frame AUG upstream of the LacZ ORF (Figure 3A, p4164). In this construct, β-galactosidase can only be produced by leaky scanning through the upstream AUG allowing the occurrence of translation initiation at the downstream start codon (note that leaky scanning at the second AUG would still allow a positive read out of the reporter) (22). Interestingly, a strong increase in β-galactosidase production was observed with this construct in pcel and kael mutants relative to the wild-type, indicating the occurrence of leaky AUG recognition during scanning. To extend these results we analyzed the implication of another EKC/KEOPS component, the Bud32 kinase, in translational repression of Gcn4p. Bud32p is closely related to the RIO family of protein kinases and the Bud32-D161A mutant has been shown to be catalytically inactive in vitro (8). De-repression of Gcn4p translation and leaky AUG scanning were observed both in bud32 cells and the catalytically inactive mutant (Figure 3C, constructs p180 and p4164), although the effect was less prominent in the latter.

These results indicate that de-repression of Gcn4 translation occurs as a result of defective recognition of the upstream inhibitory ORFs. However, we cannot formally rule out that the amino acids shortage response is also somehow activated in mutants. As outlined earlier, the latter depends critically on the phosphorylation of eIF2α by the Gcn2 kinase and gen2A cells have been shown to be defective for Gcn4 activation under amino acids shortage (19). A direct activation of Gcn4 translation by defective AUG recognition in EKC/KEOPS mutants would imply shortcutting the Gcn2p pathway. Consistent with this notion, mutation of Kae1p and Pcc1p activated β-galactosidase expression from the construct containing the wild-type Gcn4 leader even in a gen2Δ context (Figure 3D, construct p180 and Supplementary Figure S3). As expected, expression of LacZ from the construct containing the out of frame upstream AUG was not affected by the absence of Gcn2p (Figure 3D, construct p4164). Note that in a gen2Δ background the overall levels of β-galactosidase expressed from the p180 construct are generally lower, possibly because the Gcn2 pathway is activated to low levels in the growth conditions employed. However the relative effect (fold activation) of EKC mutations is similar in gen2Δ and GCN2 background.

Overall, these results support the notion that integrity of the EKC/KEOPS (and to some extent of the kinase activity of Bud32p) is required to maintain Gcn4 translational repression. They strongly suggest that de-repression of GCN4 in EKC/KEOPS mutants results from skipping non-productive translation of uORFs in the GCN4 leader via the defective recognition of the respective upstream AUGs.
Figure 3. Translational regulation of Gnc4 expression is altered in EKC and tif5 mutants. (A) Schematic of the GCN4-lacZ fusions reporters. The 5' region of GCN4 RNA contains four upstream reading frames (uORFs 1-4, empty boxes) upstream of the main GCN4-lacZ ORF (light gray box, the arrow indicates its start codon). Construct P180 contains the unmodified region. P227 contains point mutations (X) in the four upstream AUG codons. p4164 contains an elongated version of uORF1 (thin gray box, the arrow indicates its start codon) that is not in frame with the GCN4 ORF. (B) EKC/KEOPS mutants kae1, pcc1 and pcc2 de-repress GCN4 translation and increase AUG leaky scanning. β-Galactosidase activity (arbitrary units) was measured from wild-type or mutant cells transformed with the GCN4-LacZ reporters indicated above each panel. Average of four independent experiments each one performed in triplicate. Errors bars represent standard deviations. (C) As in (B) GCN4-LacZ analysis of bud32 and the catalytic mutant bud32D161A. Since the latter is also HA tagged, analysis was performed in parallel with a tagged version of wild-type BUD32. (D) Gcn4 activation by EKC mutants is independent of the Gcn2-dependent amino acids shortage pathway. Double gcn2Δ/kae1-18 mutants transformed with the Gcn4-LacZ reporters were analyzed as in (B). Note that in the absence of Gcn2p, the overall levels of β-galactosidase obtained from the p180 reporter are decreased suggesting that the Gcn2 pathway is partially activated in these growth conditions. The relative levels of activation of the reporter are however similar in the presence and absence of Gcn2 (E). Leaky scanning in tif5-K55E mutant cells. β-Galactosidase activities were measured as in (B) from wild-type or tif5-3 cells containing the GCN4-LacZ reporters grown at the permissive (30°C) temperature. Average of four independent experiments each one performed in triplicate. Errors bars represent standard deviations.
initiation at different steps (16,23). Therefore, we transformed the isolated tif5-K55E strain with the Gcn4-LacZ constructs. In contrast to EKC mutants, a global effect on translation was observed in tif5-K55E cells (Figure 3E, p227), possibly reflecting a global decrease of translation initiation efficiency.

Importantly, β-galactosidase colorimetric assays revealed the occurrence of leaky scanning in tif5-K55E cells (Figure 3E, construct p4164), which is even more prominent considering the global effect observed with the construct lacking the out of frame AUGs. Interestingly, we did not observe activation of Gcn4p translation in the construct containing all four uORFs (Figure 3C, construct p180) in tif5-K55E. This result might either imply that tif5-K55E cells are also defective for initiation at uORF1 [which is required for Gcn4 regulation, (19)], or that TC recruitment to the post-termination scanning ribosome is not affected in this mutant (see ‘Discussion’ section).

Sucrose gradient analysis of polysomes in tif5-K55E cells (Supplementary Figure S4A, left panels) revealed a profile very similar to the ones previously observed for elf5-depleted or mutant cells that are defective in translation initiation (24,25). The marked decrease in polyribosome content relative to the free ribosomal particles (that most likely constitute the bulk of the 80S peak) is characteristic of a defect in translation initiation. A similar profile was also observed in kae1A cells (Supplementary Figure S4). The translational phenotype of tif5 and kae1 is not due to a defect in the biogenesis of the 40S and 60S ribosomal particles as a parallel analysis of total dissociated ribosomes (Supplementary Figure S4A, right panels) revealed very similar levels of the two particles in the mutant and the wild-type strains (Supplementary Figure S4). Finally, no apparent differences were observed in the levels of ribosomal 25S and 18S mature ribosomal RNAs (Supplementary Figure S4B).

Overall, these results strongly suggest that the stringent requirement for elf5 function in EKC/KEOPS mutant cells relates to the efficient selection of the initiator AUG. Overall, they strongly support the involvement of EKC/KEOPS in AUG start site selection and translation initiation.

Phylogenetic and biochemical analyses implicate EKC/KEOPS in t6A37 synthesis

Deregulation of Gcn4p translation was recently observed in sua5 mutants (26). Sua5p was originally isolated because its inactivation allowed expression of a cyc1 allele containing an out-of-frame upstream AUG codon (27). The molecular mechanism of this suppression was recently elucidated by showing that Sua5p is required for the N6-threonylcarbamoyl modification present at adenine 37 (t6A37) in the anticodon stem-loop of all tRNAs decoding ANN codons (15). Defective modification of the initiator tRNA\textsuperscript{Met} in sua5 cells leads to a Gcd‘ phenotype, i.e. constitutive expression of the GCN4 gene (26). The Gcd‘ phenotype and the genetic interaction of EKC mutants with threonine biosynthesis (threonine is part of the t6A37 modification) and tRNA modification genes are suggestive of similarities with Sua5p.

While this work was in progress, a connection between Sua5p and Kae1p was revealed. Sua5 contains an N-terminal YrdC domain (residues 1–250) that is also present in a number of bacterial enzymes, such as NodU, HypF, NovN and CmcH. Remarkably these enzymes are all involved in the carbamoylation of different substrates (28).

Three-dimensional (3D) atomic models from multiple threading alignments and iterative structural assembly simulations indicates that NodU, NovN, HypF and CmcH all have a similar domain architecture that also contains an additional homologous module in addition to the YrdC module. In spite of low sequence similarity (16% identity on average), this module can be convincingly predicted to have the same fold as Pyrococcus abyssii Kae1 (PaKae1). Using the I-tasser meta server, we constructed a 3D model for this module and analyzed conserved amino acid positions between the carbamoyl transferases and Kae1. The structure of PaKae1 was obtained in the presence of the non-hydrolysable AMPPNP nucleotide, which is bound in a tunnel between two domains (7). The gamma phosphate group of AMPPNP interacts with the side chains of His107, Tyr127, Ser129 and Asp285. The phosphate oxygens of the nucleotide are further liganded to a Fe ion that is bound to Kae1 through the side chains of His107, Tyr127, His111 and Asp285 in a configuration that is very similar to that of the Fe ions found in the active site of acid phosphatases. All these residues are absolutely conserved in eukaryotic and archaeal Kae1 sequences. The 3D model of the Kae1 modules of NodU (Figure 4) shows that both histidines are conserved in the structure and that the groove could easily accommodate a nucleotide as is the case for Kae1. These data indicate that the carbamoylating enzymes contain distinct modules that have
active site configurations respectively similar to Kae1p and Sua5p, suggesting that they catalyze similar chemical reactions.

Inspired by these considerations, we performed LC–MS analysis of tRNAs extracted from wt, kael-18 and pcc1-4 strains. To minimize indirect effects the experiment was performed at the permissive temperature (30°C) for the thermosensitive alleles. The characteristic t\(^6\)A\(_{37}\) 25.5 min peak was strongly reduced in kael-18 and pcc1-4 cells to, respectively, 25 and 30% of wild-type levels, which is highly significant considering that the analysis was performed with an allele that is only partially defective at this temperature. The absence of t\(^6\)A\(_{37}\) in EKC mutants was independently reported by two other groups while this work was under revision (28,29).

These experiments together with the data reported by El Yacoubi et al. (28,29) indicate that EKC mutants are defective for t\(^6\)A\(_{37}\) modification of ANN decoding tRNAs. Since this modification affects the efficiency of codon–anticodon recognition and tRNA\(_{\text{Met}}\) is also t\(^6\)A\(_{37}\) modified, these findings strongly suggest that translational de-repression of Gcn4p in EKC mutants is mechanistically linked to defective recognition of uORF AUG start codons by unmodified tRNA\(_{\text{Met}}\). Overall, these data set the basis for understanding the molecular events associated with alteration of EKC/KEOPS integrity.

**DISCUSSION**

In spite of several phenotypes associated to mutants of the EKC/KEOPS, the primary function of this complex that is conserved in Eukarya and Archaea has remained elusive so far. The conservation of one subunit of the complex, Kae1p, in Bacteria adds a universal twist to this important question. Several theoretical possibilities exist to relate universal conservation and function. First, it is possible, that an ancient function, which was conserved in Bacteria, has been diverted in Eukarya and Archaea to serve evolutionary novel and distinct purposes and that this requires the other EKC/KEOPS components. In this view, the ancestral function of Kae1-like proteins might be lost in Eukarya and Archaea. Another possibility is that the ancestral function is universally present, but requires the additional complexity of the whole EKC/KEOPS in Eukarya and Archaea. A corollary of this possibility is that the phenotypes observed in EKC/KEOPS mutants are indirect effects of the perturbation of this ancestral function. Finally, both the ancestral and the archaeeal/eukaryotic specific functions might co-exist, and depend on the same or different biochemical forms of the complex. The results presented here allow ruling out the first possibility by providing evidence for a biochemical function of the complex that is universally conserved and impacts the efficiency and specificity of translation.

We have shown that EKC/KEOPS mutants are defective in the translational regulation of Gcn4p, which mechanistically relates to defective t\(^6\)A\(_{37}\) modification of ANN-decoding tRNAs. These findings are strengthened by the genetic interactions of pcc1 and kael mutants with genes involved in the synthesis of threonine, an amino acid that is part of the t\(^6\)A\(_{37}\) modification (30,31), and by the upregulation of Gcn4p targets in thr1 and thr4 mutants. One additional strong connection to translation is provided by the synthetic lethality of EKC mutants with a mutant in the translation factor eIF5, a protein that is at the heart of the translation initiation process and the selection of the initiator AUG, which is also impacted by the t\(^6\)A\(_{37}\) modification. Finally, structural modeling underscores the similarities between Kae1p and bacterial enzymes catalyzing similar chemical reactions as t\(^6\)A\(_{37}\) synthesis (Figure 4). While this work was in progress El Yacoubi et al. revealed a phylogenetic connection between Sua5p and Kae1p and showed that kaelA cells are defective for t\(^6\)A\(_{37}\) modification (28,29). A defect in t\(^6\)A\(_{37}\) synthesis was also reported in pcc1Δ, bud32A and the catalytic mutant allele of Bud32, bud32-D161A (29), which is fully consistent with our genetic assays on Gcn4 activation and our LC–MS data.

**Deregulation of Gcn4p expression in EKC/KEOPS mutants is linked to defective t\(^6\)A synthesis**

In the quest for a molecular function of the EKC/KEOPS that could relate the different phenotypes associated to its alterations, we first performed genetic screens and transcriptome analyses. The specific transcriptome profile of EKC/KEOPS mutants was particularly striking as a large majority of genes involved in the synthesis of amino acids were found to be upregulated (Supplementary Figures S1 and S2). We show that this is due to increased expression of the Gcn4p transcription factor that controls a large majority of the upregulated genes. Regulation of the GCN4 locus occurs at the translational level (19). Translation starts in the GCN4 leader at an upstream ORF (uORF1) but most ribosomes resume scanning after termination. Regulation occurs by altering the efficiency of recognition of three additional inhibitory upstream ORFs (uORFs2-4): whenever these are skipped, translation can occur at the downstream in frame Gcn4 AUG (Figure 3). Under amino acids shortage, phosphorylation of eIF2\(\alpha\) by the Gcn2p kinase decreases the levels of the ternary complex (containing the charged tRNA\(_{\text{Met}}\) and eIF2-GTP), which impairs recognition of the inhibitory uORFs and allows expression of Gcn4p. Mutations altering formation of the PIC, the scanning process or the recognition of the AUG by tRNA\(_{\text{Met}}\), allows reading through ORFs2-4 AUGs and expression of Gcn4p, i.e. a Gcd- phenotype [for a review see (19)].

The Gcd- phenotype we observed for EKC/KEOPS mutants could in principle be due to the activation of the Gcn2p pathway by some events causing or mimicking amino acid shortage. For instance, it has been shown that t\(^6\)A\(_{37}\) is an important determinant of the recognition of tRNA\(_{\text{Leu}}\) by the cognate isoleucyl-tRNA synthetase (32) and the presence of uncharged tRNAs is known to activate the Gen2 pathway. We show, however, that Gcn4p translation is activated in EKC mutants even in the absence of Gen2p (Figure 3D), which does not support this hypothesis. The Gcd- phenotype of EKC mutants is unlikely due to a defect in ribosomal biogenesis.
as previously described for an rpl16BA mutant (33) as the levels of 40S and the 60S particles and their ratio was very similar to the wild-type even in the severely affected kae1A mutant (Supplementary Figure S4 and data not shown).

The most likely mechanisms underlying Gcn4p activation in EKC mutants is the occurrence of scanning through the inhibitory uORFs2-4 in the GCN4 leader after translation of uORF1 (which is required for regulation). We have shown that leaky scanning indeed occurs in EKC mutants by using a reporter gene that allows expression of β-galactosidase only upon skipping of an upstream AUG (Figure 3B, construct p4164).

The most likely mechanism underlying leaky scanning in EKC mutants is the faulty recognition of the starting AUG codon, as a consequence of the strong reduction in t6A37 modified initiator tRNA\(^{\text{Met}}\) that we observed in kae1-18 cells (Figure 5). Indeed, the presence of t6A37 has been shown to confer a particular structure to the anticodon loop that stabilizes the codon–anticodon interaction and increases the binding to the ribosome (34–37).

Similar findings have been reported for the Sua5p protein and to be defective for the t6A37 modification (15,26). Consistent with this notion, several recessive tif5 alleles have been isolated in the CTD inducing a Gcd\(^{-}\) phenotype that can be mechanistically associated with defective recruitment of the TC to the re-initiating ribosome on the GCN4 RNA (23). We show that the recessive tif⁵ allele that we isolated based on its synthetic lethality with pcc1-4 (tif⁵-K55E) has a poly-somatic profile similar to the one of previously characterized tif⁵ alleles and consistent with a defect in translation initiation (24,25).

Consistent with this notion, tif⁵-K55E have a strong leaky scanning phenotype, which parallels the one observed with pcc1 and kae1 cells. In contrast to EKC mutants, however, this particular allele does not induce a Gcd\(^{-}\) phenotype (i.e. it does not constitutively activate translation of the wild-type Gcn4-LacZ reporter, Figure 3E, p180). This might be due to the fact that efficient Gcn4 activation requires translation of uORF1 and that tif⁵-K55E cells might severely affect recognition of the uORF1 start codon. Alternatively, this mutation of eIF5 might not affect TC recruitment to the post-termination 40S particle, which would allow translation of the inhibitory uORFs2-4. This would be consistent with the finding that the mutation isolated is not in the CTD but in the N-terminal GAP domain. Dominant mutations in this region have been isolated that have a relaxed AUG recognition and enhanced GAP activity (SUI phenotype) (42). Alleles with reduced GAP activity have also been described that have a reduced rate of translation initiation (43), but, generally, mutations in the N-terminal domain do not lead to a leaky scanning phenotype (23). Therefore, the K55E allele behaves differently from described mutations in the GAP domain, possibly implicating GTP hydrolysis in scanning and AUG selection. We suggest that, while EKC mutants affect the efficiency of AUG recognition via the unmodified initiator tRNA\(^{\text{Met}}\), the tif⁵-K55E allele slows down GTP hydrolysis after AUG codon–anticodon interaction, delaying (or preventing) the structural rearrangements required for the interruption of scanning and the transition to elongation. The future biochemical characterization of this allele might reveal novel interesting features of the mechanism of translation initiation and eIF5 function.

**Relationship to other EKC phenotypes**

Several phenotypes have been described for mutants of the EKC/KEOPS complex, ranging from defective

The genetic interactions of EKC mutants provide strong support for the proposed function of the complex in t6A synthesis.

The finding that integrity of the threonine synthesis pathway is essential in a context of defective EKC/KEOPS substantiates the implication of EKC/KEOPS in t6A37 synthesis. Threonine is part of the t6A37 and has been shown to be required, together with carbonate and ATP, for in vitro synthesis of this modification (40,41).

Deletion of **THR1** or **THR4**, involved in the two last steps of threonine synthesis, leads to lethality in a kae1-18 mutant. This finding is particularly striking considering that all genetic analyses have been performed in rich, threonine-containing medium, which allows normal growth of *thr1* or *thr4* threonine auxotrophs. This suggests that impairment of endogenous synthesis in *thr* mutants leads to a decrease in threonine availability to levels that do not significantly affect normal cell metabolism and protein synthesis still strongly enhance the effect of partial alterations in t6A37 synthesis due to EKC/KEOPS mutations. The stronger effect of *thr1* and *thr4* deletion on kae1-18 mutants compared to upstream mutations in the threonine pathway (e.g. hom2) might suggest the alternative, provocative, hypothesis that Thr1p and Thr4p directly take part in t6A synthesis, which might for instance imply the existence of a homoserine intermediate that would be converted to threonine by the sequential action of the two enzymes. This homoserine pathway might be a preferred but not an obligatory pathway, which would be consistent with early *in vivo* and *in vitro* studies reporting the incorporation of threonine in t6A (30,31,40,41).

An additional piece of evidence that the EKC/KEOPS impacts translation via the t6A modification, is the strong genetic interaction with the translation initiation GTPase-activating protein (GAP) Tif5p/eIF5. This protein is involved in several steps of translation initiation (16,17). After the stable interaction of initiator tRNA\(^{\text{Met}}\) with the starting AUG eIF5 enhances eIF2 GTPase activity, which prompts structural rearrangements in the scanning ribosome and allows the transition from initiation to elongation. By virtue of the multiple interactions of its carboxy-terminal domain (CTD) with other translation initiation factors, eIF5 also functions as an assembly platform that favors the interaction of the TC with the scanning ribosome and the formation of the translation PIC (16). Consistent with this function, several recessive tif5 alleles have been isolated in the CTD inducing a Gcd\(^{-}\) phenotype that can be mechanistically associated with defective recruitment of the TC to the re-initiating ribosome on the GCN4 RNA (23). We show that the recessive tif5 allele that we isolated based on its synthetic lethality with pcc1-4 (tif5-K55E) has a poly-somatic profile similar to the one of previously characterized tif5 alleles and consistent with a defect in translation initiation (24,25). Consistent with this notion, tif5-K55E have a strong leaky scanning phenotype, which parallels the one observed with pcc1 and kae1 cells. In contrast to EKC mutants, however, this particular allele does not induce a Gcd\(^{-}\) phenotype (i.e. it does not constitutively activate translation of the wild-type Gcn4-LacZ reporter, Figure 3E, p180). This might be due to the fact that efficient Gcn4 activation requires translation of uORF1 and that tif5-K55E cells might severely affect recognition of the uORF1 start codon. Alternatively, this mutation of eIF5 might not affect TC recruitment to the post-termination 40S particle, which would allow translation of the inhibitory uORFs2-4. This would be consistent with the finding that the mutation isolated is not in the CTD but in the N-terminal GAP domain. Dominant mutations in this region have been isolated that have a relaxed AUG recognition and enhanced GAP activity (SUI phenotype) (42). Alleles with reduced GAP activity have also been described that have a reduced rate of translation initiation (43), but, generally, mutations in the N-terminal domain do not lead to a leaky scanning phenotype (23). Therefore, the K55E allele behaves differently from described mutations in the GAP domain, possibly implicating GTP hydrolysis in scanning and AUG selection. We suggest that, while EKC mutants affect the efficiency of AUG recognition via the unmodified initiator tRNA\(^{\text{Met}}\), the tif5-K55E allele slows down GTP hydrolysis after AUG codon–anticodon interaction, delaying (or preventing) the structural rearrangements required for the interruption of scanning and the transition to elongation. The future biochemical characterization of this allele might reveal novel interesting features of the mechanism of translation initiation and eIF5 function.
transcription (2) to telomere homeostasis (1) and genomic instability (our unpublished data; 3). It is obviously possible that these phenotypes result from indirect effects due to defective t6A modification in several tRNAs that affect the translation of transcription or telomere maintenance factors. One obvious candidate would be the Est3p component of telomerase that requires a frameshift for its expression (44). However, defective codon recognition due to lack of t6A modification rather favors Est3p production by enhancing frameshifting [as was shown for sua5Δ cells, (26)] and it has been shown that a frameshift independent EST3-fsc allele has normal growth and telomere lengths (44). Another possibility is that the EKC/KEOPS has additional functions that might or might not be related to carbamoylation. This might correlate to the increased complexity of the archaeal/eukaryotic enzymes relative to the prokaryotic counterparts and to the physical separation of the Sua5 and Kael modules. It is important to point out, however, that both sua5 and EKC mutants have telomeric defects (1,45), suggesting that, if a eukaryotic specific effect in telomere homeostasis exists, this might involve a similar chemistry as t6A37 formation. One interesting possibility is the existence of an additional target for a carbamoylation reaction analogous to t6A37 formation, which might be a protein factor or an RNA with similar or different characteristics from the tRNA anticodon stem loop. In the context of the whole EKC the carbamoylation reaction might acquire additional substrate specificity and even be regulated based on cellular physiology or cell cycle stage. In the absence of reported physical interactions between Sua5p and EKC components, the molecular mechanism ensuring the coordination of their activities remains, however, elusive.

The long sought elucidation of at least one of the EKC/KEOPS primary functions is an important step for understanding the impact of these conserved factors on several cellular processes. Future challenges include the identification of additional targets or the discovery of the mechanistic pathways linking tRNA modification, transcription and telomere homeostasis.

**SUPPLEMENTARY DATA**

Supplementary Data are available at NAR Online.

**ACKNOWLEDGEMENTS**

The authors would like to thank A. Hinnebusch for the kind gift of Gcn4-LacZ constructs, J. Boulay M. G. Koerkamp, D. van Leenen, C. Ko, P. Kemmeren, L. Bakker and S. van Hooff for expert technical assistance, F. Lacroute for help with yeast genetics and P. Forterre, C. Mann and H. Grosjean for fruitful discussions.

**FUNDING**

Centre National pour la Recherche Scientifique (CNRS), the Agence Nationale pour la Recherche (ANR) (grant ANR-09-BLAN-0349-03 to D.L. and H.v.T. and ANR-08-JCJC-0019-01 to C.S.) Netherlands Organization of Scientific Research (NWO) (grants 021.002.035, 817.02.015, 050.71.057, 911.06.009, 016.108.607 to T.L.L.) Netherlands Bioinformatics
Centre (NBIC), the U.S. Department of Energy (grant no. DEFG0207ER64498 to V.deC.-L.) and by the National Institutes of Health (grant no. R01 GM70641-01 to V.deC.-L.). Fondation pour la Recherche Médicale (FRM) to X.L. Funding for open access charge: CNRS.

Conflict of interest statement. None declared.

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