

Hydroxylation of the eukaryotic ribosomal decoding center affects translational accuracy

Christoph Loenarz^{a,1}, Rok Sekirnik^{a,2}, Armin Thalhammer^{a,2}, Wei Ge^a, Ekaterina Spivakovskiy^a, Mukram M. Mackeen^{a,b,3}, Michael A. McDonough^a, Matthew E. Cockman^c, Benedikt M. Kessler^b, Peter J. Ratcliffe^c, Alexander Wolf^{a,4}, and Christopher J. Schofield^{a,1}

^aChemistry Research Laboratory and Oxford Centre for Integrative Systems Biology, University of Oxford, Oxford OX1 3TA, United Kingdom; ^bTarget Discovery Institute, University of Oxford, Oxford OX3 7FZ, United Kingdom; and ^cCentre for Cellular and Molecular Physiology, University of Oxford, Oxford OX3 7BN, United Kingdom

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The mechanisms by which gene expression is regulated by oxygen are of considerable interest from basic science and therapeutic perspectives. Using mass spectrometric analyses of *Saccharomyces cerevisiae* ribosomes, we found that the amino acid residue in closest proximity to the decoding center, Pro-64 of the 40S subunit ribosomal protein Rps23p (RPS23 Pro-62 in humans) undergoes posttranslational hydroxylation. We identify RPS23 hydroxylases as a highly conserved eukaryotic subfamily of Fe(II) and 2-oxoglutarate dependent oxygenases; their catalytic domain is closely related to transcription factor prolyl *trans*-4-hydroxylases that act as oxygen sensors in the hypoxic response in animals. The RPS23 hydroxylases in *S. cerevisiae* (Tpa1p), *Schizosaccharomyces pombe* and green algae catalyze an unprecedented dihydroxylation modification. This observation contrasts with higher eukaryotes, where RPS23 is monohydroxylated; the human Tpa1p homolog OGFOD1 catalyzes prolyl *trans*-3-hydroxylation. *TPA1* deletion modulates termination efficiency up to ~10-fold, including of pathophysiologically relevant sequences; we reveal Rps23p hydroxylation as its molecular basis. In contrast to most previously characterized accuracy modulators, including antibiotics and the prion state of the *S. cerevisiae* translation termination factor eRF3, Rps23p hydroxylation can either increase or decrease translational accuracy in a stop codon context-dependent manner. We identify conditions where Rps23p hydroxylation status determines viability as a consequence of nonsense codon suppression. The results reveal a direct link between oxygenase catalysis and the regulation of gene expression at the translational level. They will also aid in the development of small molecules altering translational accuracy for the treatment of genetic diseases linked to nonsense mutations.

translation | hypoxia | ribosomal hydroxylation | 2-oxoglutarate oxygenase | nonsense readthrough

The rises in atmospheric oxygen levels provided life with a new energy source, but necessitated the evolution of regulatory mechanisms (1). Defining how cells regulate protein biosynthesis by the direct addition of oxygen to cellular molecules is of current basic and medicinal interest. Recent work suggests the presence of multiple regulatory levels and interfaces between O₂ and gene expression, many of which are catalyzed by Fe(II)- and 2-oxoglutarate (2OG)-dependent oxygenases, some of which are therapeutic targets (2). The hypoxic response in animals, but not lower organisms, is substantially mediated by prolyl *trans*-4-hydroxylation of the hypoxia-inducible transcription factor (HIF) (3–6). In addition to the gene-specific regulation of transcription via HIF, oxygen-dependent modifications at other interfaces have been discovered: hydroxylation of splicing-related proteins (7), histone lysyl demethylation (8), and 5-methylcytosine hydroxylation (9). In addition to signaling, posttranslational hydroxylations can also have structural roles: prolyl *trans*-4-hydroxylation stabilizes the collagen triple helix, whereas *trans*-3-hydroxylation appears to be destabilizing in collagen (10).

Enzyme-catalyzed hydroxylation of intracellularly localized proteins was once thought to be rare, but accumulating recent evidence suggests it is widespread (11). Motivated by these findings, we investigated whether the translation of mRNA to protein is affected by oxygen-dependent modifications. A rapidly growing eukaryotic cell devotes most of its resources to the transcription, splicing, and transport of ribosomal proteins and rRNA (12). We therefore reasoned that ribosomal modification is a candidate mechanism for the regulation of protein expression.

Here we provide evidence that posttranslational prolyl hydroxylation of ribosomal protein S23 (RPS23) is an evolutionarily conserved modification that regulates translation termination. Mutations to RPS23 and its bacterial homolog (S12) have long been associated with translational accuracy, following from seminal work characterizing *Escherichia coli* growth phenotypes in the presence of the antibiotic streptomycin (13, 14). The finding that S12 mutations that increased or decreased the accuracy of stop codon recognition in *E. coli* had similar effects in yeast suggested

Significance

The processing of DNA sequences into proteins is fine-tuned to meet the conflicting demands of accuracy and speed. DNA mutations can introduce premature stop codons, leading to inactive proteins. We report that oxygen-dependent post-translational modification of the ribosomal decoding center affects stop codon readthrough in an mRNA sequence-dependent manner. Our work demonstrates that oxygenases catalyzing RPS23 hydroxylation are conserved in eukaryotes, including yeasts, flies, and humans. In basal eukaryotes, RPS23 undergoes two hydroxylations, whereas in animals we only observe one hydroxylation. Yeast ribosomes lacking hydroxylation manifest altered stop codon readthrough up to ~10-fold. The results reveal oxygen-dependent modifications that regulate translational accuracy and suggest unprecedented approaches to modulating ribosomal accuracy for medicinal application.

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¹To whom correspondence may be addressed. E-mail: christopher.schofield@chem.ox.ac.uk or christoph.loenarz@chem.ox.ac.uk.

²R.S. and A.T. contributed equally to this work.

³Present address: School of Chemical Sciences, Faculty of Science and Technology and Institute of Systems Biology, Universiti Kebangsaan Malaysia, Bangi 43600, Malaysia.

⁴Present address: Institute of Molecular Toxicology and Pharmacology, Helmholtz Zentrum München—German Research Center for Environmental Health, 85764 Neuherberg, Germany.

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RPS23 serves as a conserved “accuracy center” (15), a proposal later supported by crystallographic analyses (16, 17).

We report that in basal eukaryotes, RPS23 contains an unprecedented dihydroxyprolyl modification, whereas human RPS23 only undergoes a single prolyl *trans*-3-hydroxylation. RPS23 prolyl-3-hydroxylation affects stop codon readthrough in a sequence-specific manner, consistent with the location of the hydroxylation site within the ribosomal decoding center. Small-molecule inhibition of the RPS23 hydroxylases increases production of full-length proteins from sequences containing clinically relevant nonsense mutations, suggesting that ribosome hydroxylation is of therapeutic interest.

Results

Eukaryotic Ribosomes Undergo Posttranslational Hydroxylation. In *Saccharomyces cerevisiae*, hypoxic stress [3% (vol/vol) atmospheric O₂] decreases polysome levels and increases free 40S ribosomal subunits (Fig. S1A and B). This effect is exacerbated in severe hypoxia (0.1% O₂; Fig. S1C). The only known oxidative modification to the yeast translational machinery is the essential hypusine residue in the eukaryotic translation initiation factor 5A (18); however, this modification does not account for the majority of differences observed in hypoxia (Fig. S1E). To investigate whether these differences reflect O₂-dependent modifications, we analyzed ribosomal proteins by intact protein mass spectrometry (MS). We found that a chromatographic peak corresponding to 15,938 Da (± 1 Da) is absent in *S. cerevisiae* grown at lower O₂ levels (Fig. 1A). In severe hypoxia (0.1% O₂) and near anoxia (1 ppm O₂) peaks with masses 15,922 and 15,906 Da are

observed (Fig. 1B). Together with immunoreactivity against an RPS23 antibody, these observations identify the modified protein as Rps23p (unmodified calculated 15,906 Da), and imply Rps23p undergoes posttranslational mono- and dihydroxylation modifications (+16 and +32 Da).

Anoxia Affects Translational Accuracy via Tpa1p. Because RPS23 mutations are associated with the modulation of translational accuracy (14, 15), we investigated whether the identified oxidative modifications to Rps23p act similarly, using a dual-luciferase assay measuring readthrough of the yeast bypass of stop codon 4 (*BSC4*) (19) stop codon sequence (Table S1). *Renilla* and Firefly luciferase (Rluc and Fluc) cDNAs are separated by an in-frame stop codon or a sense codon control; their relative ratio allows quantification of stop codon readthrough as a measure of translational accuracy (Fig. S2A). Whereas hypoxic (0.1% O₂) *S. cerevisiae* gave similar results to normoxia, translational accuracy significantly increased in an anoxic environment (Fig. 1C). Although we were unable to analyze the Rps23p hydroxylation level during reoxygenation due to the low levels of cell mass obtained under anoxic conditions, luciferase assays imply that the effect may be reversible: ~7 h after a shift from anoxia to normoxia, the extent of readthrough returned to that in normoxia.

We investigated whether the Rps23p hydroxylations are enzyme mediated, focusing on 2OG oxygenases because of their roles in animal oxygen sensing (5, 20). Bioinformatic analyses imply there are ~10 *S. cerevisiae* candidate 2OG oxygenases (Fig. S1F). Tpa1p is a component of mRNA–protein complexes binding to the mRNA/poly(A) 3'-end, with *TPA1* deletion reported to

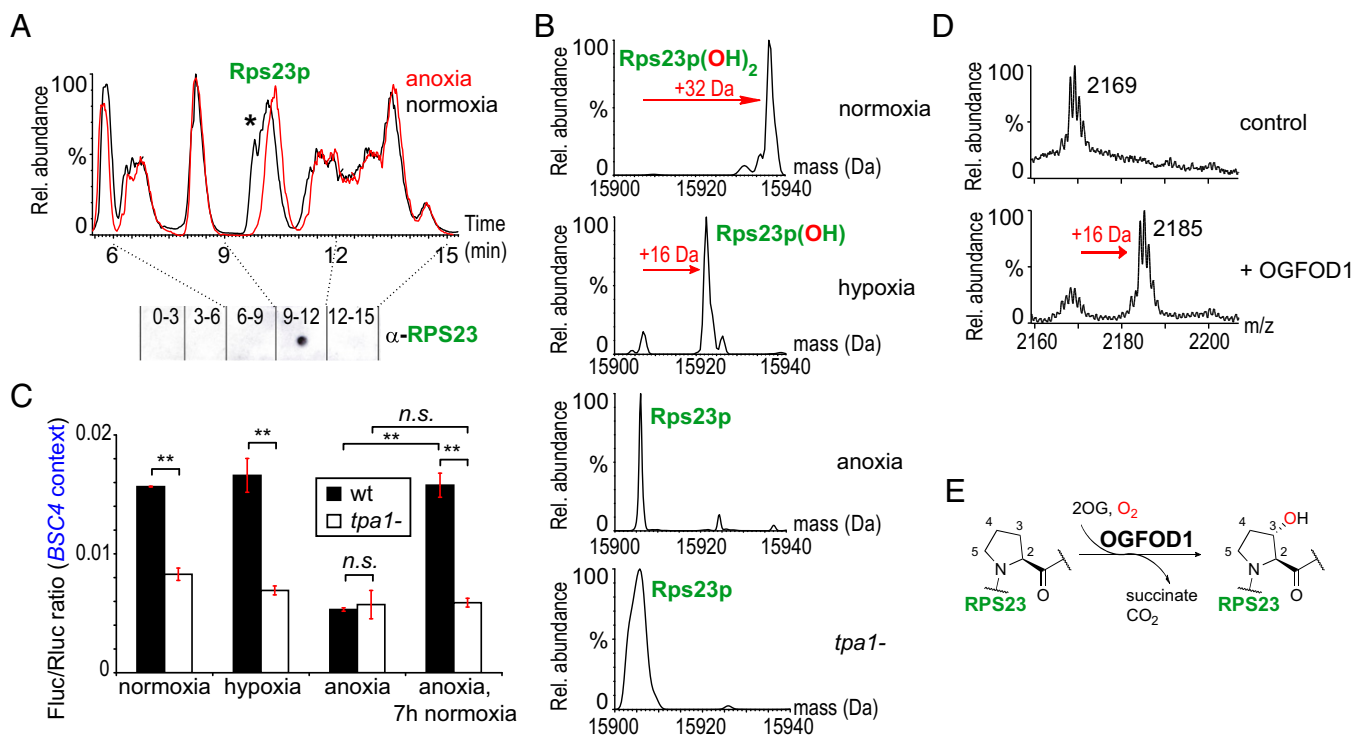


Fig. 1. Tpa1p and its human homolog OGFOD1 catalyze RPS23 prolyl hydroxylation, which affects translational accuracy in yeast. (A) UPLC chromatogram of ribosomal proteins from *S. cerevisiae* BY4742 grown in normoxia [21% (vol/vol) O₂] and anoxia (1 ppm O₂). Note the difference in the elution profile: a peak [retention time (R_t) ~9.8 min; highlighted by asterisk] present in normoxia is absent in anoxia. Dot-blot analysis of normoxic UPLC fractions (pooled every 3 min) reveals that fraction 9–12 min cross-reacts with human anti-RPS23 antibody. In anoxia, Rps23p coelutes (R_t ~10.5 min) with Rpl21ap and Rpl21bp (Fig. S7A). (B) Deconvoluted ribosome ESI-MS spectrum of Rps23p (R_t ~10 min) from *S. cerevisiae* grown in normoxia, hypoxia, and anoxia [21% (vol/vol), 0.1%, and 1 ppm O₂, respectively] and a *tpa1*[−] strain. Masses correspond to Rps23p unmodified (15,906 Da, calculated 15,906 Da; N-terminal methionine cleaved) and with addition of one and two oxygen atoms (15,922 Da and 15,938 Da, respectively). (C) Effect of changes in O₂ atmosphere [normoxia, 21% (vol/vol) O₂; hypoxia, 0.1% O₂; anoxia, 0.1 ppm O₂] on translational accuracy (Fig. S2A) in *S. cerevisiae*. Reoxygenation for 7 h after growth in near-anoxia affects accuracy of WT ($P = 0.003$) but not *tpa1*[−] strains ($P = 0.84$). $n = 3$; mean \pm SD; ** $P < 0.01$; n.s., not significant. (D) MALDI MS analyses of RPS23_{51–70} (calculated 2,168.5 Da) after incubation with OGFOD1 and cofactors. (E) Reaction scheme showing *trans*-3-hydroxyproline as the catalytic product of OGFOD1-catalyzed hydroxylation of RPS23 Pro-62, as identified by amino acid analysis (Fig. S5).

increase stop codon readthrough (21, 22); however, its substrate was hitherto unknown. Strikingly, the *tpa1⁻* strain displays reduced readthrough that was apparently unaffected by O₂ levels and similar to wild-type (WT) levels in anoxia (Fig. 1C), suggesting Tpa1p mediates the O₂-dependent pathway that affects translational accuracy. Ribosomal Rps23p from a *tpa1⁻* strain lacks oxidative modifications, implying Tpa1p as the Rps23p oxygenase (Fig. 1B). Other aspects of translation, including tested frameshifting sites (+1 and -1) and processivity through consecutive rare codons, were unaffected by *TPA1* deletion within limits of detection (Fig. S2 B–D). Polysome profiles were unaffected by *TPA1* deletion, suggesting Tpa1p does not directly influence ribosomal biogenesis (Fig. S1D).

RPS23 Hydroxylation Conserved Across Eukaryotes. Tpa1p homologs are present in most eukaryotes; they contain an N-terminal 2OG oxygenase domain and a C-terminal double-stranded beta-helix fold (Fig. S3E) (22, 23). Consistent with an oxygen-dependent role for Tpa1p in translation, its human homolog 2OG and iron-dependent oxygenase domain containing 1 (OGFOD1) is linked to ischemic signaling/stress responses (24, 25). The Tpa1p homolog in *Schizosaccharomyces pombe* (Ofd1) regulates transcription in a hypoxia-dependent manner via the sterol-responsive transcription factor Sre1 (not present in *S. cerevisiae*) (26, 27); although proposed as a prolyl-4-hydroxylase, no Ofd1 catalytic activity has been described.

As described in the companion articles (28, 29), human OGFOD1 and fruit fly *Sudestada1* also catalyze RPS23 oxidation, but result in monohydroxylation of Pro-62. To investigate the evolution of oxidative RPS23 modifications, we prepared Tpa1p/OGFOD1/Ofd1 homologs including from the green algae *Ostreococcus tauri* (otOGFOD1) and fission yeast (Ofd1) as purified proteins. All Tpa1p homologs display catalytic activity with purified GST-RPS23_{44–143} protein, as demonstrated by 2OG turnover (Fig. S4 D and E). However, to date we have observed that only purified human OGFOD1 catalyzes hydroxylation of RPS23_{51–70} peptide fragments in a 2OG oxygenase characteristic manner; alanine-scanning analyses are consistent with monohydroxylation at Pro-62 (Fig. 1D and Fig. S4 A–C). Unexpectedly, and in contrast to HIF α prolyl *trans*-4-hydroxylation (5), amino acid analyses using synthetic standards identify *trans*-3-hydroxyproline as the OGFOD1 product (Fig. 1E and Fig. S5).

Using coexpression studies of Tpa1p homologs and RPS23 in *E. coli* coupled with proteomic tandem MS, we identify RPS23 Pro-62 (Pro-64 in yeast Rps23p; Fig. S3D) as the conserved site of oxidative modifications (Fig. 2A and Fig. S6). Human OGFOD1 and fruit fly *Sudestada1* were found to catalyze Pro-62 monohydroxylation [companion articles (28, 29)], whereas Tpa1p, Ofd1, and otOGFOD1 produce a mass increment corresponding to a dihydroxyproline residue (Fig. S6). Dihydroxyproline is an unusual amino acid, previously identified as a component of diatom cell walls and *Amanita* mushroom toxin peptides, but

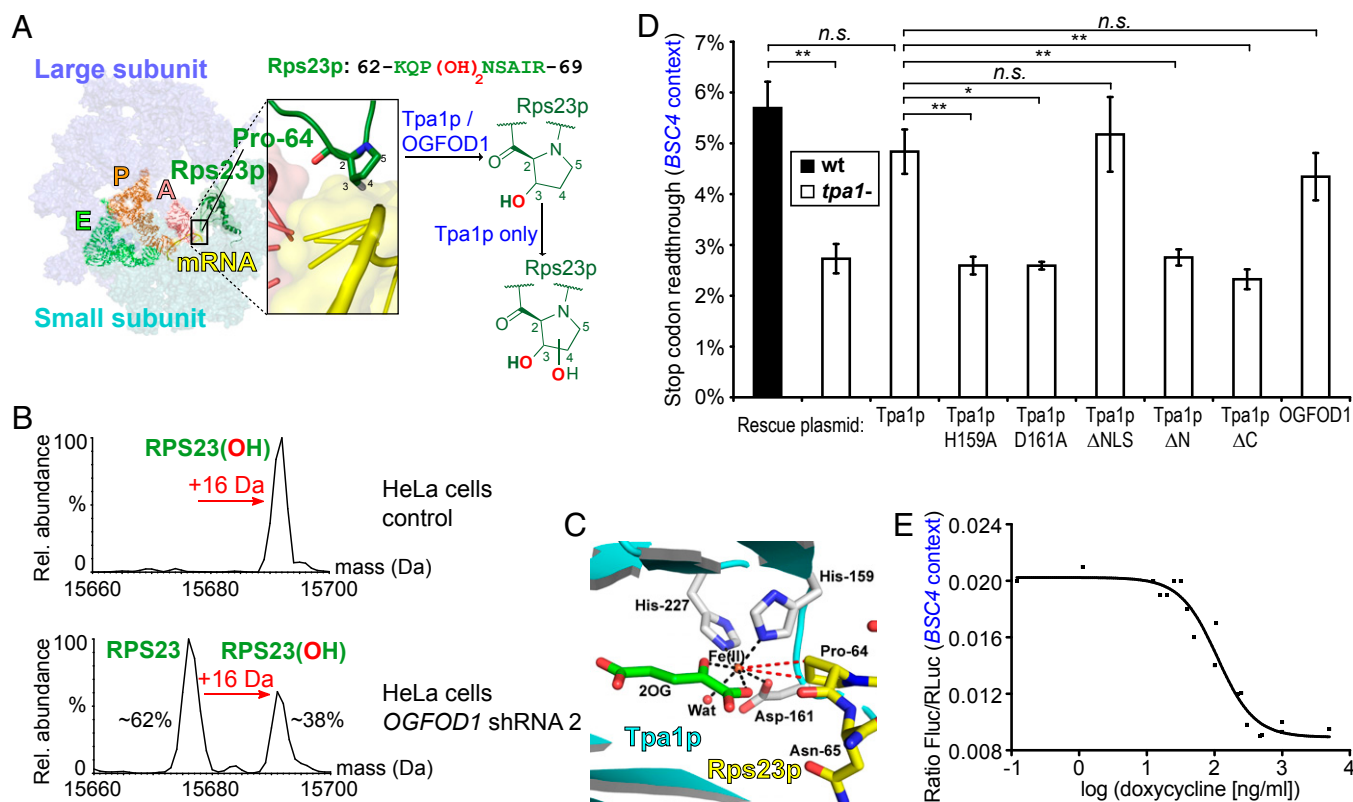


Fig. 2. OGFOD1 and Tpa1p catalyze RPS23 *trans*-3-hydroxyprolyl and dihydroxyprolyl formation, respectively. (A) Schematic depicting Tpa1p/OGFOD1 hydroxylation near the ribosomal decoding center; Rps23p Pro-64 is located at the apex of a loop protruding from a β -strand. View of a structure of the bacterial 70S ribosome including tRNAs and mRNA (yellow), with Rps23p (forest green) modeled onto bacterial S12 [Protein Data Bank (PDB) ID codes 2J00, 2J01, and 3U5C] (17, 30). (B) Deconvoluted ESI-MS spectrum of RPS23 after UPLC-separation of ribosomes from HeLa cells stably transfected with doxycycline-inducible OGFOD1 shRNA 2. Observed masses in control (15,692 Da) and induced (15,676 Da) cells correspond to monohydroxylated and unmodified RPS23 (calculated 15,676 Da; N-terminal methionine cleaved), respectively. (C) View of the Tpa1p active site (cyan/gray; PDB ID code 3KT7) (22) with ribosomal Rps23p (yellow; PDB ID code 3U5C) (30) modeled into the active site (Fig. S3C). Wat, water. (D) Reduced BSC4 stop codon readthrough from *TPA1* deletion ($P = 0.003$) is rescued by a Tpa1p plasmid ($P = 0.095$). Tpa1p Fe(II) binding (H159A, D161A) and domain deletion (Δ N, Δ C) variants are inactive ($P = 0.006$, 0.011, 0.008, and 0.004, respectively), whereas Tpa1p Δ NLS ($P = 0.54$) and OGFOD1 ($P = 0.25$) are similar to WT Tpa1p. $n = 3$; mean \pm SD; * $P < 0.05$; ** $P < 0.01$; n.s., not significant. (E) A doxycycline-repressible *TPA1* plasmid affects BSC4 stop codon readthrough in a *tpa1⁻* strain ($IC_{50} = 110.1 \pm 1.2$ ng/mL doxycycline; 95% confidence interval 76.1–159.2 ng/mL; $R^2 = 0.9573$; values are pooled from $n = 2$).

which has not previously been identified in intracellular proteins (30, 31). Given the chemically stable nature of the dihydroxylated RPS23 product, the identification of the OGFOD1 product as a *trans*-3-hydroxyproline residue, and a model of ribosomal (32) Rps23p complexed with Tpa1p (22) (Fig. 2C and Fig. S3C), we propose a 3,4-dihydroxyproline residue as the likely product of two Tpa1p, Ofd1, or otOGFOD1 catalytic cycles.

Consistent with the *in vitro* studies, ribosomal RPS23 from *S. pombe* (Fig. S7C) and *Drosophila melanogaster* S2 cells [companion article (29)] undergoes di- and monohydroxylation, respectively. MS analysis of intact ribosomal proteins in HeLa cells reveals monohydroxylation without evidence of dihydroxylation (Fig. 2B). These results suggest that the second RPS23 hydroxylation identified in simpler eukaryotes is not evolutionarily conserved in the tested animals. To investigate whether OGFOD1 catalyzes RPS23 monohydroxylation in cells, we reduced OGFOD1 levels to ~5% of control using shRNA (Fig. S3A and B); ribosomal RPS23 monohydroxylation decreased substantially (to ~40% of normal levels; Fig. 2B).

Rps23p Prolyl *Trans*-3-Hydroxylation Affects Translation Termination in Yeast. To investigate differences between mono-/dihydroxylation, we rescued *S. cerevisiae tpa1⁻* strains with *OGFOD1* or *TPA1*; ribosomal Rps23p was monohydroxylated (+16 Da) by human OGFOD1 demonstrating that mono- or dihydroxylation product selectivity is not a consequence of different cellular contexts (Fig. S7B). Whereas the Tpa1p nuclear localization sequence (NLS) is not required for Rps23p dihydroxylation, inactivating variants of its iron-binding site (D161A) and domain deletion (Δ N, Δ C) variants ablate Rps23p hydroxylation. Readthrough assays reveal that whereas *tpa1⁻* rescue with Tpa1p or OGFOD1 restores WT accuracy levels, variants (H159A and D161A) that disrupt Fe(II) binding have no effect (Fig. 2D). Tpa1p Δ NLS, but not its Δ N or Δ C variants, restore WT accuracy levels. A dose–response relationship of Tpa1p levels on accuracy was identified using doxycycline-repressible *TPA1* rescue; at intermediate doxycycline levels, ribosomal Rps23p was present in unmodified, mono- and dihydroxylated forms (Fig. 2E and Fig. S7B). The combined results imply that prolyl *trans*-3-hydroxylation exerts the major effect on accuracy whereas the second hydroxylation, observed in some eukaryotes, does not play a substantial role, at least for the tested stop codon context.

Biological Relevance of Rps23p Hydroxylations. To investigate the molecular consequences of Rps23p hydroxylation, we undertook comparisons with known modulators of translational accuracy. The prion state of the *S. cerevisiae* translation termination factor 3 (eRF3), termed “[PSI⁺]” cells (33), reduces termination efficiency with benefits for survival (34, 35), but does not affect ribosomal Rps23p hydroxylation (Fig. S7B). *TPA1* deletion in [PSI⁺] cells also decreases readthrough of the *BSC4* stop codon, suggesting the role of Tpa1p is independent of eRF3 levels (Fig. 3A). Consistent with a critical role for the structure of the Rps23p Pro-64 loop in translation (Fig. 3B), N65P and all tested Pro-64 variants are lethal; some substitutions at nearby residues affect growth (Fig. S8A–C). As observed in yeast and prokaryotes (13–15), Rps23p Lys-62 variants have opposing effects: K62T is hyperaccurate (approximately fivefold), whereas K62R substitution decreases accuracy approximately ninefold (Fig. S8D).

In some yeast strains, a nonsense stop codon in an adenine biosynthesis gene (ADenine-requiring; *ADE1*) causes red pigmentation; this pigmentation is reduced under conditions that decrease accuracy via nonsense suppression, providing an alternative assay for termination efficiency (36). Notably, *TPA1* deletion ablates red pigmentation, which is rescued by WT but not inactive Tpa1p (Fig. 3C). Further, *tpa1⁻* but not WT yeast grows on medium lacking adenine (Fig. 3D). Together, these results suggest that Rps23p hydroxylation decreases *ADE1* readthrough, but increases readthrough of *BSC4* (the initial context of our luciferase reporters). Consistent with the pigmentation observations, luciferase reporter-measured *ADE1* readthrough significantly

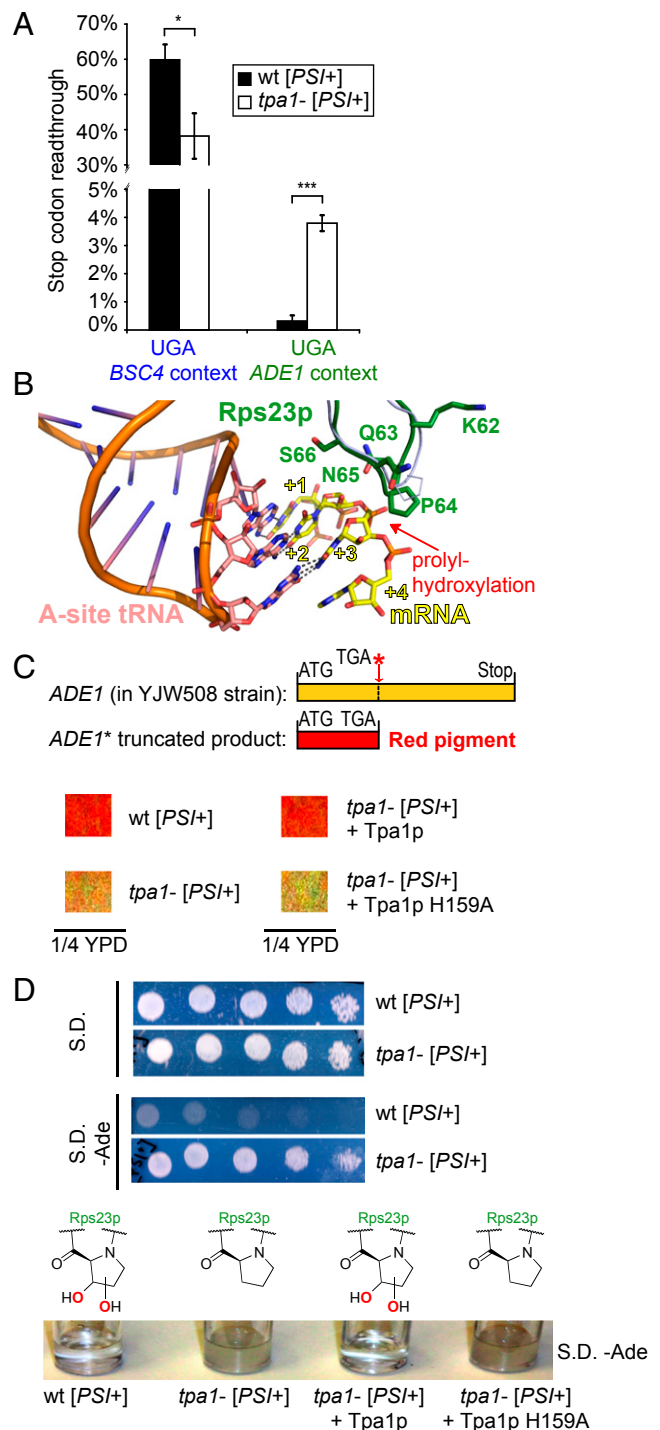


Fig. 3. Physiological relevance of altered stop codon readthrough via Rps23p hydroxylation. (A) *TPA1* deletion affects readthrough of the *BSC4* and *ADE1* stop codons ($P = 0.012$ and 0.00013 , respectively) ($n = 3$; mean \pm SD; * $P < 0.05$; *** $P < 0.001$; y axis interrupted for clarity). (B) View of the decoding site of the bacterial ribosome (17) with Rps23p (30) (green) modeled onto prokaryotic S12 (light-blue ribbon) (PDB ID codes 2J00, 2J01, and 3U5C). Integers indicate mRNA (yellow) codon positions. (C) *TPA1* deletion leads to loss of red pigmentation as a consequence of increased readthrough of an *ADE1* nonsense codon present in the YJW508 strain; this effect is reversed by plasmid rescue with WT Tpa1p but not an inactive variant (H159A). Views from suspension cultures in diluted yeast extract peptone dextrose (YPD) medium (Fig. S8E). (D) *tpa1⁻* but not WT YJW508 strains grow in medium lacking adenine both on agar (Upper; fivefold serial dilutions) and in suspension (Lower); an effect that is reversed by plasmid rescue with WT but not inactive (H159A) Tpa1p.

increases ~10-fold upon *TPA1* deletion (Fig. 3A). The directional effect of Rps23p hydroxylation on accuracy is independent of stop codon identity (UGA, UAA, and UAG) for both *BSC4* and *ADE1* contexts (Fig. S2E and F). These results imply that Tpa1p regulates accuracy in a sequence-/context-dependent and biologically relevant manner.

Relevance to Clinical Nonsense Mutations. In contrast to the finding that Rps23p hydroxylation increases *BSC4* but decreases *ADE1* readthrough, other modulating factors consistently either increase (Rps23p K62T) or decrease (Rps23p K62R, reduced eRF3 levels, paromomycin) accuracy independent of context (Fig. 4A). We therefore analyzed additional stop codon contexts, including from Tobacco mosaic virus (TMV) and nonsense codons of medicinal interest (Fig. S2G). Nonsense suppression of *Mdx* from the Duchenne muscular dystrophy mouse model increases upon *TPA1* deletion, an effect that was mirrored by WT cells grown with a cell-penetrating Tpa1p inhibitor (Fig. 4B), suggesting that inhibition of RPS23 hydroxylation may be of therapeutic interest (37).

Sequence alignments suggest that the identity of the first nucleotide after the stop codon (+4) may contribute to the directional effect of Rps23p hydroxylation (Fig. 4C); statistically, the +4 nucleotide is the least random within mammalian termination contexts (38), and can significantly affect translation termination (39, 40). Structural analyses of Rps23p predict that the Pro-64 *trans*-3-hydroxyl group introduced by Tpa1p/OGFOD1 catalysis is positioned to interact in the ribosomal A site with the phosphate backbone between the third base of the mRNA triplet codon and that in the +4 position (Fig. 3B). Importantly, substituting the +4 nucleotide from C to A (or A to C) inverts the directionality of accuracy modulation by Tpa1p for several tested contexts (Fig. 4D), i.e., *TPA1* deletion decreases readthrough of the *ADE1* UGAC and TMV UAGC stop codon but increases *ADE1* UGAA and TMV UAGA readthrough. Thus, Rps23p hydroxylation modulates the mRNA-ribosome chemical microenvironment, in a manner relevant to translational accuracy.

Discussion

Overall, given the roles of 2OG oxygenases in transcription factor, chromatin, and DNA hydroxylation/demethylation (3, 4, 7–9, 20), there is accumulating evidence that 2OG oxygenases contribute to an integrated and context-dependent regulation of gene expression. The finding that OGFOD1/Tpa1p catalysis can regulate the accuracy of translation via termination efficiency in an mRNA sequence-dependent manner opens up a unique interface between oxygenase catalysis and the regulation of protein biosynthesis. One possibility is that OGFOD1/Tpa1p catalysis is involved in regulating the translation of specific sets of mRNA sequences, including those involved in the hypoxic response, although at present it is unclear whether ribosome hydroxylation plays a direct role in cellular oxygen sensing. It is possible that ribosomal oxygenase-catalyzed modifications in animals, including those to RPL27A and RPL8 that we recently identified (41), as well as to tRNAs (42–44), help to regulate the sets of genes up-regulated by hypoxia in a context-dependent manner. In the two companion articles (28, 29), we provide evidence that Tpa1p catalyzed Rps23p hydroxylation is conserved in humans and fruit flies, and that it is of physiological importance.

The observation that a small posttranslational modification, i.e., addition of a single oxygen atom, can either increase or decrease translational accuracy, is unprecedented. We also note that RPS23 hydroxylation occurs in close proximity to the binding site of known antibiotics, including streptomycin and paromomycin, offering prospects for the targeting of hypoxic tissue via selective inhibition of unhydroxylated RPS23.

Materials and Methods

Luciferase Assay Measurements. Cells from liquid cultures of *S. cerevisiae* strains grown to OD₆₀₀ 0.4–0.6 were lysed for 25 s using Passive Lysis Buffer (all reagents from Promega Dual-Luciferase Reporter Assay System). Fluc

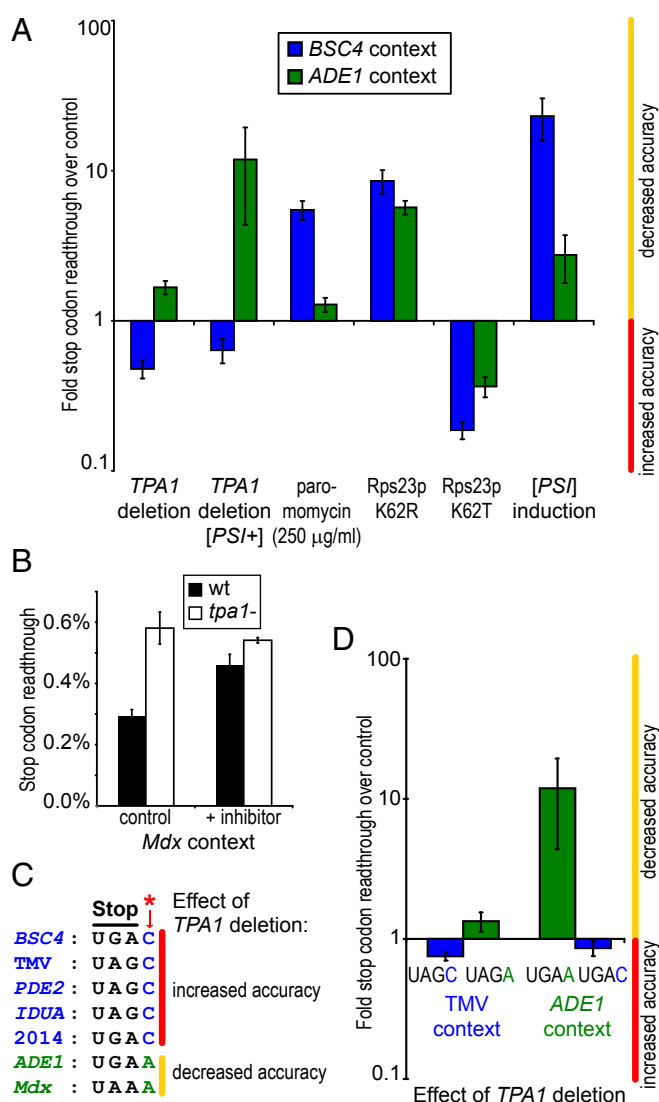


Fig. 4. Effect of Rps23p hydroxylation on translation termination is sensitive to mRNA sequence context. (A) Whereas the directional effect of known accuracy modulators is sequence independent, *TPA1* deletion reduces *BSC4* but increases *ADE1* readthrough ($n = 3$; mean \pm SD; logarithmic y axis). Relative ratios refer to controls [WT BY4742, except for *TPA1* deletion [PSI⁺] (WT [PSI⁺] YJW508) and PSI induction (WT [psi⁻] YJW508)]. (B) The generic 2OG oxygenase inhibitor dimethylallylglycine (10 mM) increases readthrough of the *Mdx* stop codon in WT but not *tpa1*⁻ BY4742 strains ($n = 2$; mean \pm SD). (C) Sequence alignment of tested stop codon contexts depicting the directional effect of Rps23p hydroxylation on readthrough (Fig. S2G). (D) Swapping of the +4 nucleotide within TMV and *ADE1* termination sequences inverts the directional effect of *TPA1* deletion on readthrough in BY4742 and YJW508 [PSI⁺] strains, respectively. Relative ratios refer to WT controls ($n = 3$; mean \pm SD; logarithmic y axis).

activity was measured (10-s integration time) using reconstituted Luciferase Assay Buffer in a GloMax 20/20 luminometer (Promega). Rluc activity (10-s integration time) was determined subsequent to quenching of Fluc activity using Stop & Glo Buffer. Signals were >10-fold above background. The Fluc:Rluc activity ratio is reported. To account for potential differences in translational processivity, the percentage stop codon readthrough of a strain was determined as the ratio of Fluc:Rluc signals of a stop codon bearing insert to that of a sense codon control.

Hypoxic and Anaerobic Cell Growth. Hypoxic culture growth was performed in an Invivo₂ Hypoxic Workstation (Ruskin Life Sciences) at 30 °C, attached to a Ruskin gas mixer module. Near-anoxic growth was performed in an anaerobic glove box (Belle Technology UK Ltd.) at 18 °C. All media, starter

cultures, growth flasks, and cell-harvesting equipment were preequilibrated at the desired O₂ concentration for 2–4 d. Cell lysis was performed in the presence of 10 mM pyridine-2,4-dicarboxylic acid (Sigma) to inhibit oxygenase activity.

Ribosomal Ultrapformance Liquid Chromatography–Electrospray Ionization Time-of-Flight MS. Samples of purified ribosomal proteins were analyzed by reversed-phase-ultrapformance liquid chromatography (RP-UPLC) and electrospray ionization time-of-flight MS (ESI-TOF MS). The method uses an Acquity UPLC BEH, C4 RP column (2.1 × 50 mm, 1.7-μm particle size, 300-Å pore size; Waters Ltd.). A flow rate of 0.3 mL/min was used with the column held at 50 °C using a Waters Acquity UPLC system connected directly to an LCT ESI-TOF MS (Waters Ltd.). The column was equilibrated with solvent A [0.2% formic acid (HCOOH) in H₂O]. Typically, 5 μL ribosomal protein sample was injected onto the column and proteins eluted using a stepped gradient from solvent A to solvent B (0.1% HCOOH in acetonitrile). The following MS parameters were used: polarity, ES⁺; capillary voltage, 3,000 V; sample cone voltage, 35 V; extraction cone voltage, 2.5 V; desolvation temperature, 250 °C; cone gas flow, 10 L/h; and desolvation gas flow (N₂), 500 L/h. The mass spectra were acquired from 400 to 2,400 *m/z* using MassLynx 4.1 software (Waters Ltd.) and protein spectra deconvoluted using Maxent 1 with a range 5–30 kDa (1-Da resolution). Masses were confirmed using manual component analysis. Sodium formate was used as the external calibrant for the instrument. Leucine enkephalin was used as the lock-mass calibrant for internal calibration of samples.

Cell Culture for OGFOD1 shRNA Knockdown Experiments. For lentiviral transduction, subconfluent cells were incubated with lentiviral stock for 24 h before puromycin selection (1–5 μg/mL). Clonal selection was performed by the dilution series of puromycin-resistant cell populations, until

formation of distinct colonies could be observed. All work involving doxycycline-inducible lentivirus was conducted in medium containing high-quality Tet system-approved FBS (Clontech) to avoid fortuitous induction of shRNA expression. Induction of shRNA used 1.5 μg/mL doxycycline every 2 d. Primary antibodies were polyclonal rabbit anti-OGFOD1 (1:500; HPA003215; Sigma-Aldrich), rat HRP-anti-β-actin (1:25,000; ab49900 [AC-15]; Abcam), and mouse anti-RPS23 (1:1,000; MCA34332; AbD Serotec). Secondary antibodies were goat HRP-anti-rabbit (1:500; 170-6515; Bio-Rad Immun-star) and goat HRP-anti-mouse (1:100,000; A9917; Sigma-Aldrich).

Statistical Analyses. For dual-luciferase assays, the probability (*P* values) that two populations are the same was tested using Student *t* test (two-tailed; assuming groups have unequal variance).

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