Control of translation efficiency in yeast by codon–anticodon interactions

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ABSTRACT

The choice of synonymous codons used to encode a polypeptide contributes to substantial differences in translation efficiency between genes. However, both the magnitude and the mechanisms of codon-mediated effects are unknown, as neither the effects of individual codons nor the parameters that modulate codon-mediated regulation are understood, particularly in eukaryotes. To explore this problem in *Saccharomyces cerevisiae*, we performed the first systematic analysis of codon effects on expression. We find that the arginine codon CGA is strongly inhibitory, resulting in progressively and sharply reduced expression with increased CGA codon dosage. CGA-mediated inhibition of expression is primarily due to wobble decoding of CGA, since it is nearly completely suppressed by coexpression of an exact match anticodon-mutated tRNA^{Arg(UCG)}, and is associated with generation of a smaller RNA fragment, likely due to endonucleolytic cleavage at a stalled ribosome. Moreover, CGA codon pairs are more effective inhibitors of expression than individual CGA codons. These results directly implicate decoding by the ribosome and interactions at neighboring sites within the ribosome as mediators of codon-specific translation efficiency.

Keywords: yeast; translation; genetic code; ribosome; codons; tRNA

INTRODUCTION

Differences in translation efficiency contribute substantially to the large variation in protein expression, with translation of individual mRNAs ranging over at least three orders of magnitude in the yeast *Saccharomyces cerevisiae* (Arava et al. 2003; de Godoy et al. 2008; Ingolia et al. 2009). While differential translation efficiency is the result of many factors, substantial evidence from both bacteria and eukaryotes indicates that the amount of polypeptide produced per mRNA is influenced by the particular choice of synonymous codons, codons that specify insertion of the same amino acid (Ghaemmamghami et al. 2003; Brockmann et al. 2007).

In organisms such as *Escherichia coli*, *S. cerevisiae*, *Caenorhabditis elegans*, and *Drosophila melanogaster*, a subset of synonymous codons, generally those decoded by the most abundant tRNAs without wobble decoding (Ikemura 1982), are “preferred codons” for efficient translation and are used more frequently than their synonyms (dos Reis et al. 2004; Parmley and Hurst 2007). Preferred codons are used nearly exclusively in many highly expressed genes,

sharply differentiating these genes from the majority of other genes (Grantham et al. 1980; Bennetzen and Hall 1982; Duret and Mouchiroud 1999). Indeed, codon usage and protein abundance correlate genome-wide in *S. cerevisiae* and *E. coli* (Ghaemmamghami et al. 2003; Tuller et al. 2007; Ishihama et al. 2008; Tuller et al. 2010b). Remarkably, despite changes in the actual identity of the preferred codons, the selection for preferred codons in many gene families is maintained across species (Man and Pilpel 2007). In addition, expression of genes in *E. coli* and in *S. cerevisiae* is frequently improved by recoding genes with preferred codons (Gustafsson et al. 2004; Burgess-Brown et al. 2008; Keppler-Ross et al. 2008; Quartley et al. 2009; Welch et al. 2009), or by overproducing rare tRNAs in *E. coli* (Burgess-Brown et al. 2008).

Nonetheless, the degree to which codons affect expression in *E. coli* has been the topic of some debate, with two recent papers arriving at very different conclusions about the quantitative importance of codon effects. Kudla et al. (2009) concluded that codon bias had little significant effect on protein levels of GFP variants, while Welch et al. (2009) found that variation in expression of two genes was strongly correlated with codon biases for 10 amino acids. However, it is likely that the coupling between transcription and translation complicates the analysis of codon effects in bacteria, since some codon changes are known to cause premature rho-mediated transcription termination (Deana

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the effect of repeats of 10 identical codons in both positions, for 59 codons (except [CCC]₁₀ and [GGG]₁₀ for technical reasons). We also examined the effect of repeats of four identical codons in the N-terminal position, for 23 codons, specifying nine amino acids that were found to be inhibitory with all specifying codons in the 10-mers. Relative luciferase activity of yeast expressing each of these constructs was assayed and reported, both with respect to expression from a vector with no insert (%V), and with respect to the identical polypeptide encoded by the synonymous codon that gave maximal activity (%C) (Table 1).

We draw three conclusions from the analysis of the 57 codons with one or more synonymous codons. First, most synonymous codons have little effect on expression in these reporters. Most constructs yielded >66%C activity, including 36 and 51 of 57 codon repeats with 10 amino acids in X₄F and in RX₃₁₄F, respectively, and 20 of 23 codon repeats with four amino acids in X₄F. As expected, most of the codons identified as preferred codons in *S. cerevisiae* (Jansen et al. 2003) yielded high expression (19 of 22 in X₄F and 22 of 22 in RX₃₁₄F) (see Table 1).

Second, the inhibitory effects of many codon repeats in X₄F are likely due to formation of secondary structures in the mRNA near the AUG, and thus may not reflect codon effects. For 15 codon repeats, we find that poor expression of X₄F correlates with predicted thermodynamic stability in the first 50 bases (folding free energy less than -16 kcal/mol) (Fig. 1B). Since strong secondary structures near the 5' end are inhibitory in *E. coli* (Kudla et al. 2009) and are selected against in both *E. coli* and *S. cerevisiae* (Tuller et al. 2010b), we infer that secondary structure may explain some or all of the inhibitory effects of these codon repeats in the N-terminal location, and, thus, these codons were not further analyzed. Similarly, the effects of Lys AAA (24%C) are likely due to the polyA sequence, which is known to inhibit expression (Dimitrova et al. 2009).

Third, Arg CGA was strongly inhibitory both in the N-terminal and internal

TABLE 1. Systematic analysis of codon effects on firefly luciferase activity

Codon	Amino acid	X ₄ F-10 mer		RX ₃₁₄ F-10mer		X ₄ F-4mer	
		% V	%C	% V	%C	%V	%C
UUU	Phe	0.2	92.8	1.5	57.7	44.4	85.0
UUC	Phe	0.3	100.0	2.6	100.0	52.2	100.0
UUA	Leu	1.6	94.1	5.9	92.2	103.3	100.0
UUG	Leu	1.6	94.1	5.7	89.1	87.9	85.1
CUU	Leu	1.5	88.2	6.4	100.0	85.2	82.5
CUC	Leu	1.6	94.1	5.6	87.5	80.0	77.4
CUA	Leu	1.7	100.0	5.8	90.6	91.9	88.9
CUG	Leu	1.4	82.4	6.4	100.0	86.0	83.3
AUU	Ile	4.7	91.9	5.5	100.0	14.5	68.6
AUC	Ile	5.0	98.1	5.5	100.0	18.0	85.3
AUA	Ile	5.1	100.0	5.0	90.4	21.1	100.0
AUG	Met	11.4	—	7.8	—	127.4	—
GUU	Val	1.3	100.0	2.9	81.4	51.5	75.9
GUC	Val	0.2	15.4	3.1	88.6	5.1	7.5
GUA	Val	0.7	53.8	3.1	88.6	67.9	100.0
GUG	Val	0.7	53.8	3.5	100.0	42.0	61.9
UCU	Ser	233.9	100.0	103.7	68.7	nd	nd
UCC	Ser	176.0	75.2	109.1	72.3	nd	nd
UCA	Ser	227.3	97.2	151.0	100.0	nd	nd
UCG	Ser	6.4	2.7	150.2	99.5	nd	nd
CCU	Pro	132.8	99.0	156.3	100.0	nd	nd
CCC	Pro	nd	nd	nd	nd	nd	nd
CCA	Pro	134.2	100.0	136.8	87.5	nd	nd
CCG	Pro	57.6	42.9	149.0	95.3	nd	nd
ACU	Thr	171.2	100.0	123.5	89.1	nd	nd
ACC	Thr	156.5	94.5	114.4	82.6	nd	nd
ACA	Thr	160.3	93.6	138.5	100.0	nd	nd
ACG	Thr	59.8	34.9	100.7	72.7	nd	nd
GCU	Ala	52.3	37.7	113.2	89.7	nd	nd
GCC	Ala	6.4	4.6	104.0	82.4	nd	nd
GCA	Ala	138.7	100.0	116.4	92.2	nd	nd
GCG	Ala	10.5	7.6	126.2	100.0	nd	nd
UAU	Tyr	2.5	73.2	3.9	100.0	59.4	100.0
UAC	Tyr	3.4	100.0	3.7	93.6	57.7	97.2
UAA	STOP	—	—	—	—	nd	nd
UAG	STOP	—	—	—	—	nd	nd
CAU	His	90.9	81.4	100.2	100.0	nd	nd
CAC	His	111.7	100.0	73.3	73.2	nd	nd
CAA	Gln	212.7	100.0	90.7	71.2	nd	nd
CAG	Gln	113.8	53.5	127.4	100.0	nd	nd
AAU	Asn	177.5	74.5	107.0	94.8	nd	nd
AAC	Asn	238.4	100.0	112.8	100.0	nd	nd
AAA	Lys	2.4	23.9	21.7	92.0	85.6	100.0
AAG	Lys	9.9	100.0	23.6	100.0	84.7	98.9
GAU	Asp	12.6	100.0	72.1	100.0	65.4	96.4
GAC	Asp	11.6	92.0	70.8	98.2	67.8	100.0
GAA	Glu	109.9	100.0	110.8	84.6	nd	nd
GAG	Glu	34.0	30.9	130.9	100.0	nd	nd
UGU	Cys	17.8	95.2	27.3	100.0	nd	nd
UGC	Cys	18.7	100.0	23.2	85.0	nd	nd
UGA	STOP	—	—	—	—	—	—
UGG	Trp	1.3	—	0.6	—	8.1	—
CGU	Arg	0.2	0.6	9.4	37.1	nd	nd
CGC	Arg	1.2	3.8	15.6	61.9	nd	nd
CGA	Arg	0.2	0.5	0.3	1.0	2.9	5.2
CGG	Arg	0.8	2.6	11.1	44.0	nd	nd
AGU	Ser	78.2	33.4	145.2	96.2	nd	nd
AGC	Ser	141.0	60.3	136.1	90.1	nd	nd

(continued)

TABLE 1. Continued

Codon	Amino acid	X ₄ F-10 mer		RX ₃₁₄ F-10mer		X ₄ F-4mer	
		% V	%C	% V	%C	%V	%C
AGA	Arg	32.1	100.0	25.3	100.0	55.4	100
AGG	Arg	5.8	17.9	16.4	65.1	nd	nd
GGU	Gly	21.5	19.6	100.8	89.7	nd	nd
GGC	Gly	29.2	26.6	112.4	100.0	nd	nd
GGA	Gly	109.8	100.0	97.4	86.7	nd	nd
GGG	Gly	nd	nd	nd	nd	nd	nd

Firefly luciferase activity of X₄F and RX₃₁₄F constructs with insertion of either 10 or four copies of each codon, as indicated. Two values are reported for each construct: (%V) is the percent activity with respect to the wild-type vector without an insertion, and (%C) is the percent activity with respect to the maximum activity obtained with each amino acid insertion. Optimal codons, as reported by Jansen et al. (2003), are shaded and bolded. (nd = not determined.)

positions (Fig. 2A; Table 1). In the N-terminal position, four other sequences likely encode inhibitory codons (Fig. 1B; Table 1), including Thr ACG, Val GUA, and possibly Ser UCG codons, which are decoded by essential single (or two copy) tRNAs (Johansson et al. 2008) and Ser AGU, which, like Arg CGA, is decoded by wobble decoding. In the internal position, four other Arg codons, as well as Phe UUU, yielded <66% C (Fig. 2A; Table 1).

CGA codon repeats severely inhibit expression in a dose-dependent manner

The Arg CGA codon stood out as a candidate for a strongly inhibitory codon, because of both the magnitude and generality of its effects on expression. The CGA codon repeats were strong inhibitors of expression in both positions, with 10 codon repeats resulting in 0.5% C and 1% C firefly luciferase activity, respectively (Fig. 2A). CGA repeats also exerted strong inhibitory effects upstream of *Renilla* luciferase (Fig. 2A). Although all of the unpreferred Arg codon repeats were inhibitory, none affected all three reporters to nearly the same extent as CGA.

The effects of CGA codons on expression are unlikely to be due to RNA secondary structure for two reasons. First, the predicted free energy of the CGA repeat (−10.2 kcal/mol) is inconsistent with an inhibitory RNA secondary structure (Fig. 1B). Second, CGA repeats in the other reading frames did not cause similar effects. The Asp GAC and Thr ACG codons, which are permutations of CGA, form repeats of the same sequence, and thus, likely form structures similar to (CGA)₁₀, but neither codon repeat inhibits expression nearly as much as CGA, either in terms of absolute, or polypeptide-adjusted, levels of luciferase (Table 1).

We also found that very few CGA codons are required for severe inhibition of expression. First, to determine the extent to which CGA codons affect expression, we examined the expression of firefly luciferase as a function of CGA codon dosage. In each case, we made identical Arg₁₀–

containing luciferase proteins, with the Arg residues coded by increasing numbers of CGA codons (Fig. 2B). In this context, two CGA codons in the N-terminal position caused a decrease in firefly luciferase activity to 46% C, while four CGA codons caused a decrease to 12% C (Fig. 2C). Within the first 14 amino acids, the position of the codons did not matter, as the constructs (CGA)₅–(AGA)₅ and (AGA)₅–(CGA)₅ yielded similar levels of luciferase activity (see Fig. 7B below). Furthermore, CGA codons were also strongly inhibitory in the internal location (RX₃₁₄F), although slightly more CGA codons were required to exert similar levels of inhibition in this position compared to the N-terminal location (Fig. 2C).

Second, the inhibitory effects of CGA were apparent with only three or four Arg residues. Inhibition by CGA codons (5.2% C) is observed with only four Arg residues upstream of firefly luciferase (Fig. 2D; Table 1), and three arginines upstream of GFP (6.7% C) (Fig. 2E). Thus, inhibition by CGA is effective with very few codons and does not require an extended arginine repeat.

Inhibition by CGA is due to I•A wobble decoding

To determine explicitly if the CGA codon inhibits expression, we examined the effect of overproducing the corresponding tRNA species on expression of (CGA)₁₀ constructs. CGA codons are decoded by tRNA^{Arg(ICG)}, which is encoded by six tRNA^{Arg} genes (Fig. 3A) bearing an ACG anticodon that is subsequently converted to ICG (Grosjean et al. 2010). Because the number of gene copies of tRNAs vary from one to 17, and the amounts of various tRNA species correspond closely to the number of genes that specify each variant (Tuller et al. 2010a), these six copies should yield sufficient amounts of tRNA to decode CGA. Nonetheless, overproduction of tRNA^{Arg(ICG)}, but not any other Arg isoacceptor, resulted in an ~10-fold increase in X₄F–(CGA)₁₀ luciferase activity compared to the vector control (Fig. 3B), and similarly suppressed the inhibitory effect of the internal CGA codon repeat (Fig. 3C). This increase in expression to ~7% of X₄F–(AGA)₁₀ luciferase activity (Fig. 3B) was as much as would be expected from the likely increase in the concentration of tRNA^{Arg(ICG)} due to the tRNA gene on the multicopy plasmid, since the multicopy plasmid added ~25 copies of the tRNA^{Arg(ICG)} gene (Schneider and Guarente 1991) to the six genomic copies, resulting in an approximately fivefold increase in gene copy number. Thus, we conclude that some of the CGA-mediated reduced expression is due to a defect in translation of CGA codons.

Since suppression by the overproduced tRNA^{Arg(ICG)} was incomplete, we inferred that either a portion of the CGA

FIGURE 2. CGA codon repeats inhibit expression in a dose-dependent manner. (A) Luciferase activity of three different reporter constructs (X₄F, RX₃₁₄F, and X₄R), containing (Arg)₁₀ insertions specified by each of the six Arg codons. (B) The relationship between CGA codon dose and luciferase activity was examined with (Arg)₁₀ inserts containing increasing CGA content, as illustrated in the schematic. (C) Firefly luciferase activity as a function of CGA codon dose in X₄F (Δ) and RX₃₁₄F (■)—reporter constructs. (D) Firefly luciferase activity of X₄F-reporter constructs containing either no insertion, or insertion of four Arg residues, specified with the indicated CGA or AGA codons. Values were normalized to the X₄F-(AGA)₄ reporter. (E) Expression of integrated GFP constructs with three Arg residues inserted at amino acid 4 specified by AGA or CGA codons. GFP expression was assessed based on the distribution of yeast cells exhibiting different intensities of fluorescence emission at 530 nm.

effects is not codon-dependent, or that decoding of the CGA codon by its cognate tRNA^{Arg(ICG)} is inherently inefficient. Decoding of the CGA codon might be inefficient either due to inherently poor tRNA function of tRNA^{Arg(ICG)} or due to the I•A wobble decoding necessary to decode the CGA codon. To address these issues, we constructed two tRNAs that should exactly base pair (an anticodon-match) to the CGA codon, by mutating the anticodon of the major tRNA^{Arg(UCU)} species from UCU to UCG, and of tRNA^{Arg(ICG)} from ACG to UCG.

The introduction of an anticodon-matched isoaccepting tRNA almost completely suppressed the expression defect caused by the 10 CGA codons. Overproduction of either the mutant tRNA^{Arg(UCU to UCG)} or the mutant tRNA^{Arg(ICG to UCG)} resulted in a 60-fold increase in luciferase activity, to nearly the levels of X₄F-(AGA)₁₀ (Fig. 4A). Similar suppression was observed for the reporter with CGA codons at amino acid 314 (Fig. 4B). This result demonstrates that almost all effects of the (CGA)₁₀ sequence are due to the use of this codon in translation, suggests that CGA decoding is severely deficient, and implies that I•A wobble decoding is a prominent factor in CGA effects on translation efficiency.

To evaluate the importance of tRNA abundance in decoding CGA with an exact match tRNA, we examined suppression of CGA-mediated inhibition of expression by a single copy of the anticodon mutated tRNA genes integrated into the yeast chromosome (Fig. 4C,D). Remarkably, we found that an integrated copy of either tRNA^{Arg(ICG to UCG)} or tRNA^{Arg(UCU to UCG)} strongly suppressed the expression defect caused by CGA codons, resulting in a 40- to 50-fold improvement in expression of the (CGA)₁₀ construct (Fig. 4D). Similarly, expression of a (CGA)₅(AGA)₅ luciferase construct improved ~20-fold in strains expressing these exact match single copy tRNAs (Fig. 4C). The simplest interpretation of this strong suppression is that the exact match tRNA^{Arg(UCG)} decodes a disproportionate fraction of CGA codons relative to that decoded by the more

FIGURE 3. Suppression of CGA-mediated expression defects by its cognate tRNA. (A) The Arg codons are compared with respect to use in the genome, gene copy number of the decoding tRNA^{Arg} species and the requirement for wobble decoding. (B) Effect of overproduction of the indicated tRNA^{Arg} species from high copy plasmids on firefly luciferase activity of X₄F-reporter constructs containing (CGA)₁₀ insertions. Values were compared to activity of X₄F-(AGA)₁₀ luciferase in a strain with an empty vector. (C) Effect of overproduction of tRNA^{Arg(ICG)} from a high copy plasmid on firefly luciferase activity of RX₃₁₄F-reporter constructs containing (CGA)₁₀ insertions at amino acid 314. Values were normalized to (AGA)₁₀ overexpressing a vector control.

abundant tRNA^{Arg(ICG)}. Thus, the decoding properties of the exact match tRNA^{Arg(UCG)}, not its abundance, are responsible for efficient expression.

CGA codons do not result in reduced mRNA amounts, but do yield a shortened RNA

Defects in translation are often closely coupled to mRNA degradation (Frischmeyer et al. 2002; van Hoof et al. 2002; Inada and Aiba 2005; Doma and Parker 2006; Isken and Maquat 2007) and sometimes also affect protein folding and activity (Kimchi-Sarfaty et al. 2007). To directly assess mRNA and protein levels, we introduced an RGS-(His)₆ epitope tag upstream of an (Arg)₈ insertion, coded by (AGA)₈ or increasing increments of CGA up to (CGA)₈. The introduction of progressively more CGA codons resulted in reduced firefly luciferase activity (Fig. 5) and in a parallel loss of protein (Fig. 5). Firefly luciferase protein was weakly detectable from strains bearing the (CGA)₃(AGA)₅ and (CGA)₄(AGA)₄ constructs, which produced 8%–12% C luciferase activity, and was undetectable from strains bearing the (CGA)₈ constructs. However, we did not observe concurrent loss of firefly luciferase mRNA from any of the (CGA)-containing constructs, relative to that from constructs with (AGA)₈, using an RT-PCR assay with the indicated probes (Fig. 5).

To ascertain whether mRNA stability of transcripts bearing CGA codons is a general phenomenon and to examine the RNA species more carefully, we also examined *Renilla* and firefly luciferase mRNAs by Northern analysis of RNA from strains bearing the RGS-(His)₆-(Arg)₈ insertions at either amino acid 4 of *Renilla* luciferase (X₄R) or at amino acid 314 in the *Renilla*-firefly luciferase fusion (RX₃₁₄F). In both cases, we observed that the amount of full-length luciferase RNA from the (CGA)₈ construct is similar to (or slightly greater than) that from the corresponding (AGA)₈ construct (Fig. 6B,F). Neither the amount nor the approximate size of *Renilla* luciferase

mRNA from the X₄R reporter was reduced in RNA from strains bearing either the (CGA)₄(AGA)₄ or the (CGA)₈ constructs, relative to that of the (AGA)₈ or no-insert constructs (Fig. 6A,B). In fact, the CGA-containing mRNAs were approximately twofold more abundant (Fig. 6B), a phenomenon that was not observed if tRNA^{Arg(ICG to UCG)} was coexpressed (Fig. 6C). As expected, the amounts of *Renilla* luciferase protein were reduced by CGA codons, and restored by coexpression of the exact match tRNA^{Arg(UCG)} (data not shown). Likewise, the large RNA from the RX₃₁₄F fusion that hybridized to both the *Renilla* luciferase and firefly luciferase probes was approximately twofold more abundant in strains bearing constructs with (CGA)₈, compared to strains bearing constructs with (AGA)₈, [(CGA)₄(AGA)₄], or no insert (Fig. 6F). Furthermore, we note that *Renilla* luciferase mRNA is itself suitable for RNA degradation, since introduction of either a UAA or a UGA stop codon at amino acid 4, which would be expected to elicit nonsense-mediated decay in susceptible genes (Isken and Maquat 2007), results in reduction of *Renilla* luciferase mRNA to ~32% of the no-insert control (Fig. 6D). Thus, we conclude that CGA-mediated inhibition of expression is not generally accompanied by mRNA degradation.

In addition to the full-length *RLuc*-FLuc fusion RNA, we also detected a smaller RNA species from strains expressing either the (CGA)₈ or the [(CGA)₄(AGA)₄] RX₃₁₄F reporters (Fig. 6E,F). If the ribosome stops or stalls at CGA codons, the mRNA might undergo cleavage, since many stalls during translation trigger endonucleolytic cleavage near the site at which the ribosome stalls (Doma and Parker 2006). The smaller RNA (labeled FLuc mRNA 2 in Fig. 6F), had the properties expected for an RNA cleaved at or near the CGA codons. This smaller RNA was detected with the firefly luciferase probe, but not the *Renilla* luciferase probe, and was found only in strains bearing the (CGA)₈ or [(CGA)₄(AGA)₄] constructs (Fig. 6F). Moreover, there is significantly more small RNA from strains with the (CGA)₈ reporters than in the [(CGA)₄(AGA)₄] reporters. We did

FIGURE 4. Suppression of CGA-mediated expression defects by an exact base-pairing anticodon mutated tRNA^{Arg(UCG)}. (A) Effect of indicated native and anticodon-mutated tRNA^{Arg} species, expressed on high copy plasmids, on firefly luciferase activity of X_4F -reporter constructs containing (CGA)₁₀ insertions. The parent tRNA of each anticodon-mutated tRNA^{Arg} is indicated by the first anticodon shown below the figure. Values were compared to activity of X_4F -(AGA)₁₀ luciferase in a strain with an empty vector. (B) Effect of an anticodon-mutated tRNA^{Arg} species, expressed on a high copy plasmid, on firefly luciferase activity of a $RX_{314}F$ -reporter construct containing a (CGA)₁₀ insertion. (C) Suppression of the inhibitory effects of CGA codons by integrated copies of the mutant tRNA^{Arg(UCG)} species that decode CGA without wobble. Luciferase activity of X_4F -reporter constructs containing (Arg)₁₀ insertions, specified by (AGA)₁₀, [(CGA)₅(AGA)₅], or (CGA)₁₀, in yeast strains bearing an integrated copy of either of two native tRNA^{Arg} species, two nonnatural exact match tRNA^{Arg} species, or a vector control as indicated. Two individual integrant strains were analyzed in each case. Values were normalized to the average of the activity of X_4F -(AGA)₁₀ luciferase in the two integrant strains expressing a vector control. Error bars represent the standard deviation from three independent transformants, each assayed in triplicate. (D) A closer examination of expression of X_4F -reporter constructs containing (CGA)₁₀ insertions in indicated strains with an expanded scale of the y-axis. Values are normalized as in C.

not observe a unique band corresponding to the RNA fragment from strains with constructs in which the CGA codons are near the N-terminus, most likely because the expected difference in size between the full-length and truncated RNAs is small (39–60 bases out of >800 bases) (Fig. 6B). Chen et al. (2010) have reported similar results: they observed an RNA cleavage product from the *PGK1* gene bearing four CGA codons. Thus, we consider it highly likely that this shorter mRNA with homology to firefly luciferase is the result of a ribosome stall at or near the CGA codons.

Adjacent CGA codons are more effective inhibitors than separated CGA codons

Although the existence of codon pair bias is well documented, its significance in gene expression has not been explicitly examined (Smith and Yarus 1989; Irwin et al. 1995; Kane 1995; Moura et al. 2005; Buchan et al. 2006; Coleman et al. 2008). To elucidate the importance of codon interactions in modulating expression, we compared the inhibitory effects of different arrangements of five CGA and five AGA codons (Fig. 7A,B). Since there was ~15-fold

more luciferase from strains in which the construct lacks any adjacent CGA codons [(CGA–AGA)₅] compared to those with five adjacent CGA codons [(AGA)₅(CGA)₅] or (CGA)₅(AGA)₅, codon arrangement plays an important role in CGA-mediated inhibition of expression.

To quantitatively assess the contribution of individual CGA codons, adjacent codon pairs and alternating codons, we increased the Arg repeat to 16. This number allowed us to insert spacers of (AGA)₂ between various CGA-containing test sequences, such that CGA codons in adjacent test sequences could not reside within a ribosome simultaneously, and to insert a sufficient number of test sequences to establish the relationship between expression and CGA dosage by systematic variation (Fig. 7C).

For all three arrangements of CGA codon test sequences, there was an exponential dependence of the inhibition of expression on the number of CGA codons (Fig. 7D–F), indicating that each test sequence acts independently. For each series, we derived the exponential coefficient of translation inhibition, called k_{TI} . In each case, this coefficient had a very similar value for CGA codons upstream of firefly or *Renilla* luciferase (Fig. 7D–F; Table 2).

FIGURE 5. CGA-mediated inhibition affects firefly luciferase activity and protein levels, but does not result in reduced luciferase mRNA. Analysis of firefly luciferase protein, mRNA, and activity in X₄F-reporter constructs containing RGS-(His)₆-(Arg)₈ insertions, from yeast strains bearing the indicated constructs (lanes 1–10). Firefly luciferase protein was detected with antibody to RGS-(His)₆, with antibody to enolase serving as an internal control. In lanes 11–14, the top two panels contain dilutions of crude extracts expressing the X₄F-RGS-(His)₆ (AGA)₈ reporter construct (10, 7.5, 5.0, 2.5 μg). Firefly luciferase mRNA was measured by RT-PCR analysis using primers homologous to bases 21–46 and 148–173, as indicated on the diagram below the panel; sequences are given in Supplemental Table S3; measurements of Actin mRNA served as an internal control. In the third panel, lanes 11–14 contain RT-PCR reactions in which dilutions of DNase-treated RNA from X₄F-RGS-(His)₆ (CGA)₈-containing sample (0.6, 0.3, 0.15, 0.075 μg) were used in the RT reactions, with equal volumes of the RT reaction in each PCR reaction. Reverse transcriptase reactions in lanes 1–10 contain 0.3 μg total RNA input. Firefly luciferase mRNA amounts were first normalized to the levels of Actin mRNA and then quantified relative to the wild-type X₄F-reporter construct with no insertion; the FLuc mRNA levels in the wild-type strain were arbitrarily set to 100. The presence of RGS-(His)₆ is indicated by an asterisk (*). Firefly luciferase activity of X₄F-reporter constructs containing RGS-(His)₆-(Arg)₈ insertions was normalized to the wild-type X₄F-reporter construct with no insertion.

Strikingly, CGA–CGA codon pairs were far more effective inhibitors of expression than isolated CGA codons or alternating CGA–AGA codon combinations. Whereas the k_{TI}^S (for single isolated codons) was -0.21 and -0.25 for the two reporters, the k_{TI}^P (for codon pairs) was -0.53 and -0.48 for CGA codon pairs, for the two reporters, respectively (Table 2). Thus, there appears to be a synergistic interaction between codons that occupy the ribosome simultaneously in adjacent sites. This synergy does not occur with CGA codons that might occupy the separated A and E sites, since the k_{TI}^A for alternating (CGA–AGA) codons was -0.22 , nearly identical to that of isolated CGA codons (Fig. 7F). We note, however, that even individual CGA codons were themselves strongly inhibitory, since 10 isolated CGA codons would be expected to yield $\sim 14\%C$ ($y = 114e^{-0.21x}$, where $x = 10$) (Fig. 7D).

DISCUSSION

The major result that emerges from this study is that decoding of the CGA codon in yeast interferes with an intrinsic aspect of the translation elongation reaction. Three features of the CGA translational inhibitory signal are notable. First, the critical defect in decoding CGA appears to be I•A wobble decoding, and not tRNA abundance, since a single copy of an exact base-pairing mutant tRNA^{Arg(UCG)} efficiently suppressed the expression defects caused by the CGA codon, whereas an approximate sixfold excess of tRNA^{Arg(ICG)} only mildly suppressed the expression defect. This means that wobble decoding modulates the efficiency of elongation, a primary catalytic function of the ribosome. The inhibition of expression reported here is clearly caused by translation of the CGA codon, and not by poly-arginine, since the expression effect is codon-dependent, is suppressible by the anticodon-mutated tRNA^{Arg(UCG)}, and does not require long stretches of arginine. Second, although the loss of expression caused by defects in decoding CGA codons is not mediated by mRNA degradation, CGA codons do result in formation of a novel RNA, most likely due to endonucleolytic cleavage near a stalled ribosome. This means that translation of sense codons can induce a strong and efficient halt to translation. Third, adjacent CGA codons are far more potent inhibitors of expression than separated CGA codons, a phenomenon most easily ascribed to interactions within the ribosome. Thus, profound differences in translation efficiency can stem directly from decoding interactions in the ribosome.

Observations from our work and others provide two clues to the mechanism by which CGA decoding impairs translation efficiency. First, the rate of acceptance of the tRNA^{Arg(ICG)} into an A site occupied with a CGA codon may be slow, since overproduction of the native tRNA^{Arg(ICG)} partially suppressed the CGA-mediated expression defect, indicating that charged tRNA^{Arg(ICG)} is limiting for this codon. Acceptance into the A site may be slow because of the altered geometry of the ICG anticodon base-paired with a CGA codon in the ribosome (Murphy and Ramakrishnan 2004). Second, CGA codons are likely to exert their effects in two successive steps during translation, since adjacent CGA codons have a synergistic effect on expression. This might mean that ribosomes have a memory of the last decoding reaction in the A site, an idea for which there is no precedent, or that the I•A wobble interaction is also recognized as defective in the P site. Zaher and Green (2009) recently identified a quality control pathway in *E. coli* in which a mismatch in the P site provokes an increased rate of misincorporation into the A site, and ultimately results in premature release. A similar pathway in yeast that probes the codon–anticodon interaction in the P site would account for the increased potency of adjacent CGA codons.

Many translation defects are closely coupled to mRNA degradation. For example, the presence of a premature

FIGURE 6. The effects of CGA codons at amino acids 4 and 314 on full-length mRNA, examined by Northern analysis. (A) Diagram of the X₄R *Renilla* luciferase reporter indicating the position of the oligonucleotide probe used in panels B–D. Probe c is located at base pairs 682–705 in the 935 base-pair *Renilla* luciferase coding sequence. (B) Analysis of the effects of insertion of CGA codons at amino acid 4 upstream of *Renilla* luciferase on *Renilla* luciferase mRNA and activity. *Renilla* luciferase mRNA from X₄R reporter constructs containing RGS-(His)₆-(Arg)₈ was examined by Northern blots with the probe c indicated in panel A; U2 snRNA served as an internal control. *Renilla* luciferase mRNA levels were first normalized to levels of U2 snRNA and then compared to the wild-type *Renilla* luciferase mRNA containing RGS-(His)₆. In the lower panel, luciferase activity of *Renilla* luciferase reporter constructs containing RGS-(His)₆-(Arg)₈ insertions are shown. (Off, expression under repressing conditions.) Lanes 1–4 contain different amounts of input total RNA from the X₄R- RGS-(His)₆ (CGA)₈ bearing strain (20, 10, 5, 2.5 μg), while lanes 5–12 contain RNA (20 μg) from yeast strains bearing the indicated constructs. (C) Decoding of CGA codons with an anticodon-mutated tRNA^{Arg} species suppresses the CGA-mediated increase in *Renilla* luciferase mRNA. Northern blot analysis of *Renilla* luciferase mRNA from *Renilla* luciferase reporter constructs containing RGS-(His)₆-(Arg)₈, using Actin as an internal control. Strains expressed either a nonnatural exact match tRNA or a vector control. Luciferase mRNA amounts were normalized to the levels of Actin mRNA and then to RGS-(His)₆ (AGA)₈; In the titration indicated with the triangle, mRNA input from the strain X₄R- RGS-(His)₆ (CGA)₈ in a vector control was varied (20, 10, 5 μg). (D) Insertion of stop codons at amino acid 4 of *Renilla* luciferase resulted in a reduction in stable *Renilla* luciferase mRNA. Northern blot analysis of *Renilla* luciferase mRNA from indicated *Renilla* luciferase reporter constructs. *Renilla* luciferase mRNA was first normalized to levels of U1 snRNA and then to RLuc (wt) mRNA. (E) A diagram of the RX₃₁₄F reporter indicating the relative position of the probes used in panel F. Probe c is located at base pairs 682–705 in *Renilla* luciferase (935 base pairs), while probe d is located at base pairs 1564–1593 in the RX₃₁₄F reporter (2631 base pairs); this corresponds to base pairs 601–630 in firefly luciferase (1668 base pairs). Probe sequences are shown in Supplemental Table S3. (F) Analysis of the effects of insertion of CGA codons at amino acid 314 in the RX₃₁₄F reporter on *Renilla*-firefly luciferase mRNA. RNA from strains bearing RX₃₁₄F constructs with RGS-(His)₆, or RGS-(His)₆-(Arg)₈ encoded with either (AGA)₈, [(CGA)₄(AGA)₄], or (CGA)₈ as indicated was examined by Northern blots with the probes indicated in panel E. U2 snRNA served as an internal control.

stop codon results in nonsense-mediated decay (Isken and Maquat 2007), an RNA structure that blocks ribosome progress results in no-go decay (Doma and Parker 2006), and the absence of a stop codon or translation into the 3' UTR results in nonstop decay (Frischmeyer et al. 2002; van Hoof et al. 2002; Inada and Aiba 2005). It is therefore not surprising that CGA-mediated inhibition of expression is associated with cleavage of the mRNA at or near the CGA codons, although it is surprising that the resulting 3' fragment appears to be stable, since there is a large amount of it. We infer that ribosomes arrest at CGA codons, since

CGA codons in two contexts result in a new RNA with properties expected of an endonucleolytic cleavage product near the CGA codons (our data; as well as Chen et al. 2010). The ribosome may remain bound to the 5' end of the fragment either because the arginine residues in the exit tunnel hold it there (Dimitrova et al. 2009), or because the ribosome with both the A and P sites occupied with tRNA^{Arg(ICG)} is arrested but left in a translation competent mode. Tuller et al. (2010a) noted that clusters of poorly adapted codons are enriched within the first 50 codons, and proposed that such clusters function to stall ribosomes

FIGURE 7. CGA codon pairs are more inhibitory than single CGA codons. (A,B) The effects of the arrangement of five CGA codons within an (Arg)₁₀ insert on *Renilla* luciferase activity were compared to each other and to *Renilla* luciferase activity from constructs containing (Arg)₁₀ insertions specified by (AGA)₁₀ or (CGA)₁₀. (A) The arrangements of Arg CGA codons with the number of CGA codons and codon pairs indicated and (B) the resulting luciferase activity with lanes corresponding to constructs in A. Reported values were corrected to an independently transcribed firefly luciferase gene, and then normalized to (AGA)₁₀. (C) Schematic of the arrangement of CGA codons in test constructs designed to examine the effects of single CGA codons, pairs of CGA codons, and alternating CGA codons. (D) Relative luciferase activity of firefly luciferase reporter constructs containing (Arg)₁₆ insertions with increasing CGA content arranged either as single codons or codon pairs. Reported values were corrected to an independently transcribed *Renilla* luciferase gene, and then normalized to (AGA)₁₆. (E) Relative luciferase activity of *Renilla* luciferase reporter constructs containing (Arg)₁₆ insertions with increasing CGA content arranged either as single codons or codon pairs. Reported values were corrected to an independently transcribed firefly luciferase gene, and then normalized to (AGA)₁₆. (F) Relative luciferase activity of *Renilla* luciferase reporter constructs containing (Arg)₁₆ insertions with increasing CGA content. CGA codons are clustered either as single codons separated by two AGA codons or as alternating codons (CGA-AGA). Reported values were corrected to an independently transcribed firefly luciferase gene, and then normalized to (AGA)₁₆.

early in translation to prevent collisions between elongating ribosomes. If rare codons do play a role in mediating ribosome density, then mechanisms may have evolved to prevent mRNA decay in response to codon-mediated ribosome arrest.

Although codon choice has been linked for many years to the large disparities in translation efficiency (Grantham et al. 1980; Bennetzen and Hall 1982; Ikemura 1982; Duret and Mouchiroud 1999; Ghaemmaghami et al. 2003; Brockmann et al. 2007), we still do not know the extent to which codon choice directly influences translation efficiency, or the identity and properties of all of the codon-initiated signals, particularly in eukaryotes. With this systematic analysis, we found that most codon repeats have only small effects on

expression, making the CGA codon conspicuous in its effects. Given that CGA codon pairs are substantially more effective inhibitors of expression than isolated CGA codons, it is likely that other codon combinations, that are more potent inhibitors of expression than the individual codon repeats, remain to be discovered.

Three lines of reasoning support the idea that codon-mediated regulation of expression is functionally important and nearly universal. First, I•A wobble decoding of CGA is conserved in the hemiascomycetes fungi, as well as in many bacteria (Curran 1995; Grosjean et al. 2010), even though it is presumed to be inefficient, as described here. Although CGA is the only codon in yeast decoded with an I•A wobble interaction (Johansson et al. 2008), it is highly unlikely that

TABLE 2. Exponential coefficient of translation inhibition, k_{TI} values, for CGA in different contexts

Test arrangement	Layout	Reporter	k_{TI}
CGA singles	(CGA·AGA·AGA) _x AGA _(16-3x) *AGA AGA (CGA·AGA·AGA) _x AGA _(14-3x)	FLuc	−0.21
		RLuc*	−0.24
		RLuc	−0.25
CGA pairs	(CGA·CGA·AGA·AGA) _x AGA _(16-4x)	FLuc	−0.53
		RLuc	−0.48
CGA alternating	AGA AGA (CGA·AGA) _x AGA _(14-2x)	RLuc	−0.22

widespread retention of this decoding strategy is accidental. Since expression of the exact match tRNA^{Arg(UCG)} does not result in obvious growth defects on either minimal or rich plates at temperatures from 18°C to 37°C (data not shown), such mutations have undoubtedly occurred in nature and been discarded. Presumably the retention of this inefficient decoding has an important, but as yet undiscovered, role. Second, organisms go to great lengths to vary the decoding efficiency at different codons. All organisms use 49 (or fewer) tRNAs to decode the 61 sense codons, making wobble decoding of some codons obligatory. In many organisms, there is tremendous variation in the amounts of tRNAs that decode particular codons, as exemplified by the one to 17 copies of different tRNA genes in yeast (Grosjean et al. 2010). In addition, the sequences of tRNAs that all decode the same codon (isodecoders) vary. In humans, the 49 isoaccepting tRNA families are encoded by 450 tRNA genes, which include ~270 isodecoders, some of which appear to be functionally distinct (Geslain and Pan 2010). Third, reduced translation efficiency of some genes may be under selective pressure. Even in human genes, clusters of codons decoded by rare codons are found more frequently than expected, are frequently conserved between human and chimp orthologs, and sometimes cluster into distinct functional groups (Neafsey and Galagan 2007; Parmley and Huynen 2009). A resolution to these issues of codon usage requires further identification of the precise codon combinations that reduce translation efficiency, and an analysis of their distribution in genomes and functional importance.

MATERIALS AND METHODS

Plasmids and strains

Both X₄F-firefly luciferase (pDL306) and RX₃₁₄F dual luciferase (pDL202) constructs were constructed from dual luciferase vector pDB688 (Keeling et al. 2004) and contain unique SalI and BamHI sites for codon insertions. To construct X₄F (pDL306), the *Renilla* luciferase gene was removed by BamHI digestion followed by ligation of oligonucleotides DLO188 and DLO189 containing an ATG start codon (Supplemental Table S3). To construct RX₃₁₄F (pDL202), the BamHI site upstream of the *Renilla* luciferase gene was mutated by PCR amplification of pDB688 with DLO99 and

DLO100 (Supplemental Table S3). Vectors (JP0054 and JP0056), in which *Renilla* and firefly luciferase genes were expressed under P_{GAL1,10} control, were constructed by individual PCR amplification of the firefly and *Renilla* luciferase genes with additional sequences required for ligation-independent cloning into a 2μ *URA3* vector BG2596, which is derived from BG2483 by insertion of sequences containing P_{GAL10} and the CYC1 transcription terminator (details available upon request) (Malkowski et al. 2007).

Unique NheI and BamHI sites were inserted in firefly luciferase (JP0054) or *Renilla* luciferase (JP0056). Oligonucleotides containing codon insertions (Supplemental Table S3) were annealed and ligated into SalI-BamHI or NheI-BamHI digested vectors, which were transformed into *E. coli* (XL1-Blue) and confirmed by sequence analysis. Vector pEKD1024 carries the superfolder GFP (Pedelacq et al. 2006) and yeast optimized RFP (Keppler-Ross et al. 2008) under control of the bidirectional *GAL1,10* promoter inserted into integrating vector JW132 (J Whipple and E Phizicky, unpubl.).

To construct 2μ plasmids containing tRNA^{Arg} species with their own promoters, tRNAs *tR(CCG)L*, *tR(CCU)J*, *tR(UCU)K*, and *tR(ACG)D*, with ~300 base pairs of upstream and downstream sequences, were amplified by PCR from yeast genomic DNA with oligonucleotides listed in Supplemental Table S3. Purified PCR products were cloned into 2μ vector AVA0577 (Alexandrov et al. 2006) by standard ligation-independent cloning (LIC) (Aslanidis and de Jong 1990; Alexandrov et al. 2004). To construct the anticodon mutated exact base-pairing tRNA^{Arg} variants, pDL866 (*tR(UCU)K*) and pDL867 (*tR(ACG)D*) were subjected to site-directed mutagenesis with a Quickchange kit (Stratagene 200518) according to the manufacturer's instructions using oligonucleotides DLO444, DLO445, DLO508, and DLO509 (Supplemental Table S3). To integrate tRNA^{Arg} variants, tRNA genes from pDL866 (*tR(UCU)K*), pDL867 (*tR(ACG)D*), pDL869 (*tR(UCU-U35G)*), and pDL870 (*tR(ACG-A34U)*) were amplified by PCR with oligonucleotides DLO575 and DLO576, ligated into integrating vector JW132 by standard LIC cloning procedures (Alexandrov et al. 2004). A linear fragment containing the tRNA^{Arg} variants was produced by restriction digestion and transformed into yeast strain BY4741. Yeast strains, derived from BY4741 (*MATa his3Δ1, leu2Δ0, met15Δ0, ura3Δ0*), are listed in Supplemental Table S1 and plasmids used in these studies are listed in Supplemental Table S2.

Luciferase assays

Transformants were grown in SD-uracil or SC-uracil with 2% raffinose, 2% galactose media (Sherman et al. 1986) to OD₆₀₀ between 0.8 and 1.0. Cells (0.8 OD·mL) were harvested by centrifugation at 4000 rpm, washed with cold ddH₂O, and resuspended in 1X Passive Lysis Buffer (100 μL) (Promega), nutated for 15 min at 25°C, followed by incubation at 25°C for 20 min., then 2 μL lysate was added to 20 μL Luciferase Assay Reagent II (Dual Luciferase Reporter Assay System, Promega PAE1910). Firefly luciferase activity was measured in Relative Luminescence Units (RLUs), with a 2-sec delay and a 10-sec measurement using a Berthold FB12 luminometer. To assay *Renilla* luciferase activity in the same

reaction, 20 μ L 1X Stop and Glo (Promega) was added, and activity was measured as described above. In the systematic analysis of codon effects, activity was measured from three independent transformants, and three readings were performed on each lysate. Error bars reflect the standard deviation between three independent transformants.

Protein expression and RNA analysis

Yeast transformants, in which *Renilla* and firefly luciferase genes were under control of the *GAL1,10* promoter, were grown as described (Quartley et al. 2009) except that strains were grown overnight in raffinose to an OD₆₀₀ between 1.8 and 2.4, and expression was induced by addition of 3X YP media with 6% galactose (1/3 final volume), followed by growth for 6 h. Crude extract preparation and Western blot analysis were performed as described (Quartley et al. 2009). Membranes were incubated either with mouse anti-RGS-(His)₆ (1:2500 dilution) or with rabbit anti-enolase (1:25,000 dilution), followed by washing as described previously, and incubation with either HRP-conjugated Goat IgG anti-mouse (1:10,000 dilution, BioRad) for anti-RGS-(His)₆ or HRP-conjugated Goat IgG anti-rabbit (1:10,000 dilution, Biorad) for anti-enolase, and development with ECL Plus Western blotting detection system (GE Healthcare RPN2132).

RNA for RT-PCR was prepared from 50 OD-mL cells as described (Schmitt et al. 1990). RNA (5 μ g) was treated with 1.25 U RNase free RQ DNase I (Promega M6101), followed by cDNA synthesis from DNase-treated RNA (0.3 μ g) using Moloney murine leukemia virus reverse transcriptase (Invitrogen); the resulting cDNA was amplified by PCR using primers DLO527, DLO528, DLO531, and DLO532 (Supplemental Table S3).

Expression of green fluorescent protein in yeast strains bearing an integrated copy of pEKD1163, pEKD1170, or pEKD1167 was assessed after growth in YP + adenine + 2% galactose + 2% raffinose. Cells were washed in cold ddH₂O, collected by centrifugation, resuspended in 1X PBS, and subjected to analysis on the FACSCantoII with excitation at 488 nm and detection of fluorescence emission using a 530/30 band pass filter.

For analysis by Northern blots, yeast transformants (150 OD-mL), grown as described above, were resuspended in 3 mL RNA isobuffer (0.5 M NaCl, 0.2 M Tris-Cl, pH 7.5, 0.01 M EDTA, 1% SDS) followed by addition of 3 mL phenol-chloroform-isoamyl alcohol (PCA) (25:24:1), equilibrated at pH 7.5, and glass beads, and lysed by vortexing 10 times for 20 sec with 1 min on ice between cycles. Additional RNA isobuffer (4.5 mL) and PCA (4.5 mL) were added, and lysates were centrifuged; the aqueous phase was re-extracted with 4.5 mL PCA, nucleic acids were precipitated with ethanol, and the pellet was resuspended in TE, pH 7.5, and stored at -80°C . Total RNA (20 μ g) was separated by electrophoresis either in a 1.5% formaldehyde agarose gel (Sambrook et al. 1989) or in a 6% polyacrylamide gel (19:1) containing 7 M urea, transferred to Genescreen plus membrane, and UV crosslinked to the membrane at 254 nm for 2 min. mRNA was visualized by synthetic oligonucleotide probes DLO511, DLO531, and SHO21 (Supplemental Table S3) labeled at the 5' end with ³²P using T4 polynucleotide kinase (Roche).

SUPPLEMENTAL MATERIAL

Supplemental material can be found at <http://www.rnajournal.org>.

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