Identification of novel Arabidopsis thaliana upstream open reading frames that control expression of the main coding sequences in a peptide sequence-dependent manner

Isao Ebina1,†, Mariko Takemoto-Tsutsumi2,†, Shun Watanabe1, Hiroaki Koyama2, Yayoi Endo3, Kaori Kimata2, Takuya Igarashi2, Karin Murakami2, Rin Kudo2, Arisa Ohsumi2, Abdul Latif Noh2, Hiro Takahashi4, Satoshi Naito1,2 and Hitoshi Onouchi2,*

1Graduate School of Life Science, Hokkaido University, Sapporo 060-0810, Japan, 2Graduate School of Agriculture, Hokkaido University, Sapporo 060-8589, Japan, 3Faculty of Agriculture, Hokkaido University, Sapporo 060-8589, Japan and 4Graduate School of Horticulture, Chiba University, Matsudo 271-8510, Japan

Received August 5, 2014; Revised December 30, 2014; Accepted January 8, 2015

ABSTRACT

Upstream open reading frames (uORFs) are often found in the 5'-leader regions of eukaryotic mRNAs and can negatively modulate the translational efficiency of the downstream main ORF. Although the effects of most uORFs are thought to be independent of their encoded peptide sequences, certain uORFs control translation of the main ORF in a peptide sequence-dependent manner. For genome-wide identification of such peptide sequence-dependent regulatory uORFs, exhaustive searches for uORFs with conserved amino acid sequences have been conducted using bioinformatic analyses. However, whether the conserved uORFs identified by these bioinformatic approaches encode regulatory peptides has not been experimentally determined. Here we analyzed 16 recently identified Arabidopsis thaliana conserved uORFs for the effects of their amino acid sequences on the expression of the main ORF using a transient expression assay. We identified five novel uORFs that repress main ORF expression in a peptide sequence-dependent manner. Mutational analysis revealed that, in four of them, the C-terminal region of the uORF-encoded peptide is critical for the repression of main ORF expression. Intriguingly, we also identified one exceptional sequence-dependent regulatory uORF, in which the stop codon position is not conserved and the C-terminal region is not important for the repression of main ORF expression.

INTRODUCTION

Nascent peptides with certain specific sequences cause ribosome stalling during mRNA translation and thereby regulate gene expression (1–5). In prokaryotes, small open reading frames (ORFs) located in the 5'-leader regions of several genes encode regulatory nascent peptides that cause ribosome stalling in the middle of or at the stop codon of the small ORFs under specific conditions. Ribosome stalling in these small ORFs induces expression of the downstream cistron by destabilizing the secondary structure to make the Shine–Dalgarno sequence accessible (1,6–10) or by inhibiting transcriptional termination before the downstream cistron (11). In eukaryotes, most documented regulatory nascent peptides are encoded by upstream open reading frames (uORFs), which are located in the 5' untranslated regions (5'-UTRs). Although uORFs often negatively modulate the translational efficiency of the downstream main ORF, most uORFs exert their effects in a sequence-independent manner (12). By contrast, certain uORFs control translation of the main ORF in a peptide sequence-dependent manner (1–5,13). In the previously
characterized sequence-dependent regulatory uORFs, such as those in the cytomegalovirus gpUL4, fungal arg-2 and CPA1, mammalian AdoMetDC, and Arabidopsis thaliana AdoMetDC1 genes, the uORF-encoded peptides cause ribosome stalling at the stop codon of the uORF (14–18). The stalled ribosome prevents other scanning ribosomes from reaching the start codon of the main ORF, resulting in translational repression of the main ORF. In many of these uORF peptide-mediated translational regulations, a metabolite acts as an effector molecule. For example, arginine induces ribosome stalling in the arg-2 and CPA1 uORFs (16,17), and polyamine induces ribosome stalling in the mammalian AdoMetDC and A. thaliana AdoMetDC1 uORFs (15,18). In both cases, the effector molecule is a metabolite produced by the pathway involving the main ORF-encoded enzyme (15–18). Therefore, the translational repression mediated by these uORF-encoded peptides acts as feedback regulation.

To date, only a limited number of regulatory nascent peptides have been found in prokaryotes and eukaryotes, and the prevalence of nascent peptide-mediated gene regulation is yet to be determined. In attempts to identify uORF-encoded regulatory peptides on a genome-wide scale, exhaustive searches for uORFs encoding conserved amino acid sequences, which are referred to as ‘conserved peptide uORFs (CPuORFs)’ (19), have been conducted in various organisms, such as mammals (20), plants (19,21) and insects (22), using comparative genomic analyses. In plants, Hayden and Jorgensen identified 26 homology groups of CPuORFs by comparing uORF sequences between A. thaliana and rice homologous genes or among A. thaliana paralogous genes (19). Recently, Vaughan et al. additionally identified four homology groups of CPuORFs by comparing uORF sequences between A. thaliana, cotton, orange, soybean, grape and tobacco (21). For a more comprehensive identification of CPuORFs, we developed a BLAST-based program, BAIUCAS, which permits comparisons of uORF sequences of certain species to those of any other species in the expressed sequence tag databases. Using BAIUCAS, we identified 13 additional homology groups of A. thaliana CPuORFs that are conserved beyond Brassicales (23). These bioinformatic analyses revealed that a plant genome contains more than 40 homology groups of CPuORFs. However, it has not been assessed how many of the identified CPuORFs encode regulatory peptides that control main ORF translation.

In our previous report, we classified the recently identified plant CPuORFs into two classes based on the conservation pattern of their encoded amino acid sequences (23). In class I CPuORFs, the C-terminal amino acid sequence and the stop codon position are evolutionarily conserved. In class II CPuORFs, the amino acid sequence is conserved entirely or the N-terminal and/or middle region is conserved, but the stop codon position is not conserved. Cryo-electron microscopy studies by Bhushan et al. revealed that the C-terminal regions of the gpUL4 and arg-2 uORF-encoded nascent peptides interact with components of the ribosomal exit tunnel when ribosome stalling occurs at the uORF stop codon (24). Additionally, genetic and biochemical studies have revealed that interaction between a regulatory nascent peptide and the exit tunnel components is important for ribosome stalling (7,9,25–31). Considering these observations, class I CPuORFs may be more likely to encode a regulatory nascent peptide. However, the relationship between the conservation patterns of the CPuORF-encoded peptide sequences and the abilities as regulatory nascent peptides has not been addressed.

In this study, to address what percentage and type of CPuORFs encode regulatory peptides, we selected 16 recently identified A. thaliana CPuORFs, all of which belong to distinct homology groups, and analyzed the effects of their peptide sequences on the expression of the main ORF. From this analysis, we identified five novel peptide sequence-dependent regulatory uORFs that repress main ORF expression. Additionally, we found that CPuORFs belonging to class I have an increased tendency to encode regulatory peptides compared with class II CPuORFs. However, we identified one sequence-dependent regulatory uORF from class II CPuORFs, and found that it has an exceptional feature in that the C-terminal region of the encoded peptide is not important for the repression of the main ORF expression.

MATERIALS AND METHODS

Plant material and growth condition

Arabidopsis thaliana MM2d suspension cells (32) were cultured in modified Linsmaier and Skoog (LS) medium (33) at 26°C in the dark with orbital shaking at 130 rpm. Cells were transferred to fresh medium every week.

Plasmid construction

For cloning of the 5′-UTRs of At1g67480 and At3g55050, poly(A)+ RNA was prepared from A. thaliana (Col-0 ecotype) flower buds and opened flowers using a Qiagen Plant RNAeasy Mini Kit (Qiagen) and a GenElute mRNA Miniprep Kit (Sigma-Aldrich). For cloning of the 5′-UTRs of the other genes, poly(A)+ RNA was prepared from A. thaliana (Col-0 ecotype) seedlings using the same kits. cDNA of the 5′-UTRs was amplified from poly(A)+ RNA using the OneStep RT-PCR Kit (Qiagen) and the primers listed in Supplementary Table S1. The sense and antisense primers contained XbaI and SalI restriction endonuclease sites, respectively.

Plasmid pIE0, which harbors the cauliflower mosaic virus 35S RNA (35S) promoter, the 5′-UTR of the A. thaliana AdoMetDC1 gene, the Renilla luciferase (RLUC) coding sequence and the polyadenylation signal of the Agrobacterium tumefaciens nopaline synthase (NOS) gene in pUC19, was used as the cloning vector for the amplified 5′-UTRs. To construct this vector, we first digested plasmid pSY209 (18), which contains the AdoMetDC1 5′-UTR and the RLUC coding sequence in the pSP64 Poly(A) vector (Promega), with XbaI and Smal at sites downstream of RLUC, treated the digested DNA with a T4 DNA polymerase to fill in the XbaI end, and then ligated the blunt-ended XbaI site to the Smal site to remove the XbaI and Smal sites. The modified pSY209 was then digested with HindIII, treated with the Klenow fragment to fill in the digested ends, and digested with SacI. The HindIII-SacI fragment of the modified pSY209 containing the AdoMetDC1 5′-UTR and
the RLUC coding sequence was inserted between the 35S promoter and the NOS polyadenylation signal of plasmid pMI4(WT) (34,35) to generate pIE0, after pMI4(WT) was digested with XbaI, treated with the Klenow fragment, and digested with SacI.

The amplified 5′-UTR cDNA fragments containing the CPuORFs analyzed in this study were digested with XbaI and SacI and ligated between the XbaI and SacI sites of pIE0 to generate 35S::UTR:RLUC reporter plasmids. The deletion and insertion mutations and the codon changes were introduced into the CPuORFs using the overlap extension polymerase chain reaction (PCR) method (36) with primers listed in Supplementary Tables S2–S8. In all of the constructs, sequence analysis confirmed the integrity of the PCR-amplified regions.

The 35S::UTR:FLUC reporter plasmids carry the 5′-UTR of the ANAC082, CIPK6, At3g15430, At5g27920 or OTLD1 genes between the 35S promoter and a firefly luciferase (FLUC) coding sequence. To construct these plasmids, oligonucleotides LUCSalIF (5′-CTCCTCTAGATATCAATCTCTTCTCTCAAAAGATG GCGTCGACCATGGAAGC-3′) and LUCSalIR (5′-GCTTCCATGGTCGACGCCATCTTTTGAGAAGA GATTGATATCTAGAGGA-3′) were annealed, digested with XbaI and SacI, and the SalI-SacI fragment containing the FLUC coding sequence was ligated into the SalI and SacI sites of the constructs plasmids to yield the 35S::UTR:RLUC reporter plasmids.

**Transient expression assay**

In the transient expression experiments, plasmid DNAs were introduced into MM2d protoplasts by electroporation or polyethylene glycol (PEG) treatment. To prepare protoplasts, MM2d cells were collected by centrifugation on the third day after transfer to fresh media and suspended in modified LS medium containing 1% (w/v) cellulase Onozuka RS (Yakult Pharmaceutical) and 5% (w/v) pectolyase Y23 (Seishin Pharmaceutical) and 0.4 M mannitol, and incubated at 26 °C with gentle shaking until the suspension became turbid with protoplasts (~3 h). The protoplasts were then washed five times with wash buffer (0.4 M mannitol, 5 mM CaCl₂ and 12.5 mM NaOAc, pH 5.8). For electroporation, protoplasts were suspended in electroporation buffer (5 mM morpholinooethanesulfonic acid, 70 mM KCl and 0.3 M mannitol, pH 5.8). Ten micrograms each of a 35S::UTR:RLUC reporter plasmid and the 35S::FLUC internal control plasmid, pBI221 (Clontech), which carries an Escherichia coli β-glucuronidase (GUS) coding sequence under control of the 35S promoter, were mixed with 3 × 10⁶ protoplasts in 100 μl of ManMg solution and 110 μl of PEG solution (40% PEG4000, 0.5 M CaCl₂, 0.4 M mannitol). This mixture was incubated for 15 min at room temperature, and diluted by adding 800 μl of wash buffer. The protoplasts were centrifuged (60 × g, 2 min at room temperature) and resuspended in 1 ml of the modified LS medium containing 0.4 M mannitol.

For PEG-mediated transfection, protoplasts were suspended in ManMg solution (5 mM morpholinooethanesulfonic acid, 15 mM MgCl₂ and 0.4 M mannitol, pH 5.8). Five micrometers each of a 35S::UTR:RLUC reporter plasmid, a 35S::UTR:FLUC reporter plasmid and the 35S::GUS internal control plasmid, pBI221 (Clontech), which carries an Escherichia coli β-glucuronidase (GUS) coding sequence under control of the 35S promoter, were mixed with 3 × 10⁶ protoplasts in 100 μl of ManMg solution and 110 μl of PEG solution (40% PEG4000, 0.5 M CaCl₂, 0.4 M mannitol). This mixture was incubated for 15 min at room temperature, and diluted by adding 800 μl of wash buffer. The protoplasts were centrifuged (60 × g, 2 min at room temperature) and resuspended in 1 ml of the modified LS medium containing 0.4 M mannitol.

After 48 h of incubation at 23 °C in the dark, cells were harvested and suspended in 200 μl of extraction buffer [100 mM(NaH₂/Na₂H)PO₄, 5 mM dithiothreitol, pH 7] by sonication on ice with a Branson Sonifier 250. A Dual-LUC Reporter Assay Kit (Promega) was used to measure the RLUC and FLUC activities. GUS activities were determined as described by Jefferson (39) with 4-methylumbelliferyl-β-D-glucuronide as the substrate, using a spectrofluorimeter (Hitachi, Fluorescence Spectrophotometer F-2500).

**RESULTS**

**Identifying sequence-dependent regulatory uORFs**

Among the recently identified CPuORFs, we selected 16 CPuORFs for analysis of their sequence-dependent effects on main ORF expression. Ten of the CPuORFs (in the ANAC082, ANAC096, ARF4, ATMPK20, CIPK6, At1g67480, At3g55050, At5g27920 and At5g27920 genes) belong to class I, whereas six (in the AT3G55050, ATMPK20, AT4G12790 and AT5G63640 genes) belong to class II (23).

To investigate the regulatory functions of these CPuORFs, the 5′-UTR of each gene was amplified by reverse transcription PCR (RT-PCR) using primers designed based on full-length cDNA sequence information available at the Arabidopsis Information Resource (TAIR) website (http://arabidopsis.org/) (Figure 1. Supplementary Figure S1, Table S1). According to the current TAIR gene models (TAIR10), splice variant forms of the 5′-UTRs exist for some of the selected genes. In fact, multiple bands were detected when the 5′-UTRs of some genes were amplified by RT-PCR. In all such cases, the most abundant RT-PCR product was cloned and sequencing confirmed the presences of the CPuORFs in the cloned 5′-UTRs (Supplementary Figure S1). Each cloned 5′-UTR was fused to the RLUC coding sequence and placed under the control of the 35S promoter (Figure 2A).

To assess the sequence-dependence of the effect of each CPuORF on main ORF expression, frameshift (fs) mutations were introduced to alter the amino acid sequence of each CPuORF. A +1 or −1 fs mutation was introduced upstream or in the conserved region of each CPuORF, and another fs mutation was introduced before the stop codon to shift the reading frame back to the original frame (Supplementary Figure S1). In the AT3G55050 CPuORF, two sets of fs mutations were introduced to avoid changing
Figure 1. Schematic representation of the 5′-UTRs containing CPuORFs analyzed in this study. Shaded, open and closed boxes represent the CPuORFs, the other uORFs and the main ORFs, respectively. Arrows indicate the positions of primers used to clone the 5′-UTRs. The translation initiation contexts of the first and internal in-frame AUG codons of each CPuORF are shown.

Peptides translated from the five CPuORFs repress main ORF expression

In the fs mutants, only a few nucleotide changes were introduced into each CPuORF, whereas most amino acid residues in the frameshifted region were altered. However, it is possible that those nucleotide changes affected main ORF expression independently of CPuORF-encoded peptide function; for example, by altering the secondary structure of the mRNA. To address this possibility, we eliminated the start codon of the *ANAC082*, *CIPK6*, *At3g15430*, *At5g27920* and *OTLD1* CPuORFs by changing them to an

the length of ORF beginning at the internal Met codon, Met-40 (Supplementary Figure S1K). In the *ANAC096*, *OTLD1* and *AT4G12790* CPuORFs, introduction of the fs mutations generated an in-frame premature stop codon. Therefore, an additional nucleotide change was made to replace the premature stop codon by an amino acid-encoding codon (Supplementary Figure S1B, H and M). In the *ATMPK20*, *CIPK6*, *CIPK23*, *AT1G67480*, *AT3G55050* and *AT4G12790* genes, another uORF overlaps with the CPuORF (Figure 1, Supplementary Figure S1D, E, F, I, K and M). In these cases, fs mutations were introduced leaving the presence and length of the overlapping uORF unaltered, because alteration in the presence or length of other uORFs may affect the main ORF expression.

The 35S::UTR:RLUC reporter plasmid containing the wild-type (WT) or fs mutant version of each CPuORF was introduced into protoplasts prepared from *A. thaliana* MM2d suspension cultured cells. After 48 h of incubation, cells were harvested and disrupted for analysis of RLUC activity. As shown in Figure 2, among the class I CPuORFs, the fs mutations of the *ANAC082*, *CIPK6*, *At3g15430* and *At5g27920* CPuORFs exhibited a more than two-fold increase in RLUC activity level compared with the corresponding WT (Figure 2B). In five of the remaining class I CPuORFs, the fs mutations had weaker effects, with 1.2- to 1.5-fold increases. By contrast, the *At1g67480* uORF mutant showed no significant effect (Figure 2B). Among the class II CPuORFs, the fs mutations of the *OTLD1* and *At4g10170* CPuORFs significantly enhanced the RLUC activity by 2.1- and 1.4-fold, respectively (Figure 2C). The fs mutations of the other class II CPuORFs had no significant effect (Figure 2C). These results suggest that nine class I CPuORFs and two class II CPuORFs may have a sequence-dependent inhibitory effect on main ORF expression. We further analyzed the five CPuORFs whose fs mutation caused a more than two-fold increase in reporter expression level.
AAG codon, and tested if the fs mutations of these uORFs affected main ORF expression even in the absence of the uORF start codon by the transient expression assay. In the ANAC082 CPuORF, the internal Met codon, Met-20, was also replaced by an AAG codon. In all of the CPuORFs tested, eliminating the start codon elevated RLUC activity, indicating that translation of these CPuORFs have an inhibitory effect on main ORF expression (Figure 3). However, the fs mutations caused no further increase in RLUC activity in the absence of the CPuORF start codon (Figure 3). These results indicate that the effects of the fs mutations of the five CPuORFs depend on translation of the CPuORFs, and suggest that the peptides encoded by these CPuORFs are responsible for the sequence-dependent inhibitory effect on main ORF expression.

The newly identified regulatory uORF peptides act in cis

If the peptides encoded by the five CPuORFs function as regulatory nascent peptides to repress main ORF expression, as do the previously characterized regulatory uORF peptides, they should act in cis to exert their effects only on the downstream main ORF on the same mRNA. To address this hypothesis, we tested whether the five CPuORFs acted in cis or in trans to repress main ORF expression. For this analysis, the 5'-UTR containing the WT or fs version of each CPuORF was fused to the FLUC reporter gene and placed under control of the 35S promoter to yield 35S:UTR:FLUC reporter plasmids (Figure 4A). The 35S:UTR:FLUC reporter plasmid harboring the WT or fs version of each CPuORF was co-transfected into MM2d protoplasts with the 35S:UTR:RLUC reporter plasmid carrying the WT or fs version of the corresponding CPuORF. As shown in Figure 4, for all the CPuORFs analyzed, the reporter activities of each reporter plasmid were not significantly different, regardless of whether it was co-transfected with the WT or fs version of the other reporter plasmid. These results indicate that neither the WT nor the fs mutant CPuORF affected the reporter activity of the other reporter plasmid, and suggest that the peptides encoded by these five CPuORFs act in cis to repress main ORF expression.

Identification of critical residues of the uORF-encoded regulatory peptides

To identify critical amino acid residues of the five CPuORF-encoded peptides responsible for the inhibitory function, we next performed Ala scanning mutagenesis. Several amino acids of each CPuORF peptide were individually changed...
to Ala, and their effects on expression of the downstream RLUC reporter gene were examined by the transient expression assay. To compare the functional importance of the amino acid residues in the CPuORF peptides and their evolutionary conservation levels, conservation scores (40) were calculated based on the alignments shown in Supplementary Figure S2 and indicated below the graphs showing data of the transient expression assays in Figure 5.

In the ANAC082 CPuORF peptide, the C-terminal region comprising the 24th to the 37th amino acid residues is highly conserved (Figure 5A, Supplementary Figure S2A). As shown in Figure 5A, all the Ala substitutions tested in this region increased RLUC activity compared with the WT, suggesting that this region is crucial for the repression of main ORF expression. Even the Ala substitution of Q29, which shows a low conservation score, had a strong effect (Figure 5A). By contrast, the Ala substitutions introduced outside of the highly conserved region exhibited no significant effect. Although Y19 and M20 are highly conserved, we did not test Ala substitutions of these residues because they may also affect translation initiation from M20 and it would be difficult to assess their effects solely on the peptide function.

In the CIPK6 CPuORF peptide, the C-terminal region comprising the 20th to the 32nd amino acid residues is highly conserved (Figure 5B, Supplementary Figure S2B). As shown in Figure 5B, many of the Ala substitutions in this region affected RLUC activity. Of these, D31A had a stronger repressive effect than the WT. Among the Ala substitutions introduced outside of the highly conserved region, only R17A caused an increase in RLUC activity; however, its effect was much weaker than the Ala substitutions of Arg residues in the highly conserved region (R22 and R26). These results suggest that the C-terminal region comprising 12 amino acid residues is critical for the repression of main ORF expression and that R17 may have an accessory role.

The At3g15430 CPuORF peptide has a long highly conserved C-terminal region comprising the 19th to the 48th amino acid residues (Figure 5C, Supplementary Figure S2C). Of the four first amino acid residues in the highly conserved region, three residues showing relatively high conservation scores (P19, F20 and Y22) were individually replaced by Ala, and their effects were tested. As shown in Figure 5C, none of them exhibited a significant effect. By contrast, in the remaining C-terminal highly conserved region, many of the Ala substitutions did affect RLUC activity level. These results suggest that the C-terminal region comprising 26 amino acid residues is important for the repression of main ORF expression.

In the At5g27920 CPuORF peptide, the C-terminal region comprising the 19th to the 34th amino acid residues is relatively highly conserved, and the region comprising the 10th to 18th amino acid residues is weakly conserved (Figure 5D, Supplementary Figure S2D). In the highly conserved region, all the Ala substitutions tested increased RLUC activity except for S21A. By contrast, none of the Ala substitutions in the weakly conserved region showed a significant effect, except that R12A slightly upregulated RLUC activity (Figure 5D). These results suggest that the C-terminal highly conserved region is crucial for the repression of main ORF expression, and that R12 in the weakly conserved region may have an accessory role.

In the OTLD1 CPuORF peptide, the region comprising the 20th to 33rd amino acid residues is highly conserved (Figure 5E, Supplementary Figure S2E). In addition, the region comprising the 16th to 18th amino acid residues is weakly conserved. In the highly conserved region, many of the Ala substitutions tested enhanced RLUC activity level. In the weakly conserved region, S16A slightly increased...
The peptides encoded by the five sequence-dependent CPuORFs act in cis. (A) Schematic representation of the 35S::UTR:FLUC and 35S::UTR:RLUC reporter constructs. (B-F) Transient expression studies of the co-transfected 35S::UTR:FLUC and 35S::UTR:RLUC reporter plasmids. MM2d protoplasts were co-transfected with three plasmids, the 35S::UTR:FLUC and 35S::UTR:RLUC reporter plasmids and the 35S::GUS internal control plasmid, by PEG treatment. The 35S::UTR:FLUC and 35S::UTR:RLUC reporter plasmids contained the WT or fs version of the ANAC082 (B), CIPK6 (C), At3g15430 (D), At5g27920 (E) or OTLD1 (F) CPuORFs. Co-transfection was carried out for all four combinations for each CPuORF, as indicated. After 48 h of incubation, the transfected cells were harvested and disrupted for luciferase and GUS assays. FLUC and RLUC activities were normalized to GUS activity, and the FLUC and RLUC activities relative to those in the experiment where both reporter plasmids had the WT CPuORF were calculated. Means ± S.D. of at least three biological replicates are shown. Each graph is representative of two or more separate experiments using independently prepared protoplasts. In each graph, bars with the same colors are not significantly different, whereas bars with different colors differ significantly (P < 0.05 by t-test).

RLUC activity (Figure 5E). These results suggest that the highly conserved region is important for the repression of main ORF expression, and that S16 in the weakly conserved region may have an accessory role.

Overall, the Ala scanning mutagenesis revealed that, in the newly identified regulatory uORF peptides, the regions comprising 12 to 26 amino acid residues in the highly conserved regions have a pivotal role in the repression of main ORF expression, and that, in some of the uORF peptides, the weakly conserved region located upstream of the highly conserved region may have an accessory role. In the ANAC082 CPuORF, Q29A exhibited a strong effect despite its low conservation, whereas G20A and H28A in CIPK6, F34A in At3g15430 and P26A in OTLD1 showed no significant effect despite their high conservation. These results suggest that changes of these amino acid residues to certain specific amino acids are tolerated, but changes to the other amino acids are not.

Synonymous codon changes in the critical region

Although the analysis in Figure 3 suggested that the peptides encoded by the five CPuORFs are responsible for main ORF repression, to further confirm the peptide sequence-dependence of the effects of the five CPuORFs, we investigated the effects of synonymous codon changes in the crucial region of the five CPuORFs. For this analysis, we introduced synonymous changes in the Arg codons in the crucial region of each CPuORF, because the Ala substitutions of the Arg residues showed relatively strong effects in many cases (Figure 5) and two of three nucleotides in an Arg codon can be altered synonymously. We examined the effects of the synonymous codon changes in the five CPuORFs on expression of the downstream RLUC reporter gene by the transient expression assay. As shown in Figure 6, in all five CPuORFs, none of the synonymous changes tested affected RLUC activity, in contrast to the effects of the Ala substitutions of the same codons. These results confirmed that the amino acid sequences of the five
Figure 5. Alanine scanning of the five sequence-dependent CPuORFs (A–E) Effects of Ala substitutions and conservation scores of amino acid residues in the ANAC082 (A), CIPK6 (B), At3g15430 (C), At5g27920 (D) and OTLD1 (E) CPuORFs. The 35S: UTR: RLUC reporter plasmid harboring a WT CPuORF or its mutant with an Ala substitution was co-transfected with the 35S: FLUC internal control plasmid into MM2d protoplasts by electroporation, and the reporter activities were analyzed as in Figure 2. Means ± S.D. of at least three biological replicates are shown. Each graph is representative of at least two separate experiments using independently prepared protoplasts. Single and double asterisks indicate significant differences from the corresponding WT at $P < 0.05$ and $P < 0.01$ by t-test, respectively. The amino acid sequence of each CPuORF is indicated below the graph. Amino acid residues analyzed in the Ala scanning mutagenesis are shown in bold. The numbers below the amino acid residues indicate the positions of the residues in each CPuORF peptide. Conservation scores of the amino acid residues of each CPuORF were calculated based on the alignments presented in Supplementary Figure S2, using the Scorecons server (40), and shown in the graph below the numbers indicating the positions of the amino acid residues.
Figure 6. Effects of synonymous codon changes in the five CPuORFs. (A–F) Transient expression studies to compare the effects of Ala substitutions and synonymous codon changes in the ANAC082 (A), CIPK6 (B), At3g15430 (C), At5g27920 (D) and OTLD1 (E and F) CPuORFs. The 35S: UTR:RLUC reporter plasmid harboring a WT CPuORF or its mutant with an Ala substitution or a synonymous codon change was co-transfected with the 35S: FLUC internal control plasmid into MM2d protoplasts by electroporation, and the reporter activities were analyzed as in Figure 2. Means ± S.D. of at least three biological replicates are shown. Each graph is representative of three separate experiments using independently prepared protoplasts. Single and double asterisks indicate significant differences between two constructs at $P < 0.05$ and $P < 0.01$ by t-test, respectively, whereas ‘n/s’ indicate a non-significant difference. In (A) and (D), the data for the R24A, R34A and R26A mutants are the same as those presented in Figure 5A and D.

CPuORFs are responsible for the sequence-dependent inhibitory effects on main ORF expression.

The 3' non-conserved region and the stop codon position of the OTLD1 uORF are not important

The C-terminal amino acid sequence and the stop codon position of the OTLD1 CPuORF are not evolutionarily conserved (Supplementary Figure S2E), implying that the C-terminal region of the OTLD1 CPuORF-encoded peptide is not essential for the repression of main ORF expression. To address this possibility, we generated a deletion series of the 3' non-conserved region of the OTLD1 CPuORF (Figure 7A). The changes in the uORF length by these deletions may affect gene expression, regardless of the function of the uORF peptide, because the efficiencies of translational reinitiation following uORF translation and uORF-induced nonsense-mediated mRNA decay depend on uORF size (41,42). Therefore, to assess the effects of the deletions solely on the uORF peptide function, we also constructed an fs version (fs2) of each deletion mutant (Figure 7A) and compared the effects of the deletion mutant OTLD1 CPuORFs on expression of the downstream RLUC reporter gene with those of their corresponding fs versions, using the transient expression assay. As shown in Figure 7B, even in the presence of any of these deletions, the fs2 mutation showed a similar effect to that in their absence. These results suggest that the C-terminal non-conserved region of the OTLD1 CPuORF-encoded peptide is not essential for the repression of main ORF expression. Additionally, these results also suggest that the stop codon position of the OTLD1 CPuORF is not important for the repression, because the position of the stop codon was moved four or five codons upstream in the deletion mutants.

To further confirm the unimportance of the stop codon position of the OTLD1 CPuORF, we next substituted an Ala codon for the stop codon (Stop39A) (Figure 7C). This
Figure 7. The 3′-non-conserved region and the stop codon of the OTLD1 CPuORF are not important for the peptide sequence-dependent repressive effect. (A) Amino acid sequences of the WT OTLD1 CPuORF and its mutants with a deletion (ΔPWDI, ΔWDIL or ΔPWDIL) and/or a fs (fs2) mutation. The frameshifted region in the fs2 mutant is underlined. Hyphens indicate the deleted amino acid residues. (B) Transient expression assay to test the effect of the C-terminal deletion series. (C) Amino acid sequences of the WT OTLD1 CPuORF and its mutants with the Ala substitution of the stop codon (Stop39A) and/or a fs mutation. Asterisks represent stop codons. The frameshifted region in the fs mutant is underlined. (D) Transient expression assay to examine the effect of uORF stop codon elimination. In (B) and (D), the 35S::UTR:RLUC reporter plasmids containing the WT or mutant OTLD1 CPuORF whose sequence is presented in (A) and (C), respectively, was co-transfected with the 35S::FLUC internal control plasmid into MM2d protoplasts by electroporation, and the reporter activities were analyzed as in Figure 2. Means ± S.D. of four and three biological replicates are shown in (B) and (D), respectively. Each graph is representative of two or more separate experiments using independently prepared protoplasts. Double asterisks indicate a significant difference between two constructs (P < 0.01 by t-test).

Ala substitution moved the stop codon position six codons downstream, because there is another in-frame stop codon six codons downstream of the original stop codon (Supplementary Figure S1H). We compared the effect of the CPuORF carrying the Stop39A mutation on the RLUC gene expression with that of its fs version using the transient expression assay. As shown in Figure 7D, even in the presence of the Stop39A mutation, the fs mutation exhibited a similar effect to that in its absence. This result suggests that the stop codon position of the OTLD1 CPuORF is not critical for the repression of main ORF expression.

DISCUSSION

Previously, ~10 peptide sequence-dependent regulatory uORFs have been reported in eukaryotes (2–3,43–46). In plants, although five CPuORFs have been reported to be involved in the regulation of main ORF expression (44,46–51), to date, their peptide sequence-dependence has only been shown in two of them (44,46). In the present study, we analyzed 16 A. thaliana CPuORFs for their effects on main ORF expression, and identified five novel regulatory uORFs that control main ORF expression in a peptide sequence-dependent manner.

Identification of peptide sequence-dependent regulatory uORF

Of the CPuORFs analyzed in this study, the fs mutations of the ANAC082, CIPK6, At3g15340, At5g27920 and OTLD1 CPuORFs conferred a more than two-fold increase in main ORF expression compared with their corresponding WT constructs (Figure 2). The effects of the fs mutations were abolished in the absence of the uORF start codon (Figure 3). These results indicated that translation of the CPuORFs
is required for the fs mutations to exert their effects, and suggested that the effects of the fs mutations are caused by amino acid sequence alterations of the CPuORFs rather than by nucleotide sequence changes. Another possibility to explain the dependence of the fs mutations' effects on uORF translation is that the fs mutations affected the translation initiation efficiencies of the CPuORFs, and that the reduced translation efficiencies of the CPuORFs resulted in increased main ORF translation. However, this possibility is unlikely for the following reasons: firstly, in the five CPuORFs, the fs mutations were introduced more than eight nucleotides downstream of the uORF start codon (Supplementary Figure S1A, E, H, J and O), and, therefore, the fs mutations did not change the translation initiation context sequence (52,53). Secondly, in all five CPuORFs, some Ala substitutions introduced at positions further away from the uORF initiation codon and in different positions from the fs mutations showed similar effects to the corresponding fs mutation (Figure 5). Furthermore, synonymous codon changes introduced at codons where an Ala substitution elevated main ORF expression showed no significant repressive effect (Figure 6). These observations strongly suggest that the peptides encoded by these five CPuORFs function to repress main ORF expression.

Of these five CPuORFs, only the OTLD1 CPuORF belongs to class II, whereas the other four CPuORFs belong to class I. Among the remaining class I CPuORFs analyzed, the fs mutations of the ATPMK20, At3g55050 and At1g02480 CPuORFs conferred ∼1.5-fold increases in the main ORF expression level compared with their corresponding WT (Figure 2B), suggesting that these CPuORFs may have a modest sequence-dependent repressive effect, although further mutational analyses are necessary to establish the peptide sequence-dependence of these uORFs. The remaining class I CPuORFs (At1g67480, ANAC096 and ARF4 uORFs) exhibited little or no significant sequence-dependent effect on main ORF expression (Figure 2B). The amino acid sequence conservations of these three CPuORFs are relatively low in Arabidopsis compared with those of other plant orthologs (23). Therefore, little or no significant sequence-dependent effects of these CPuORFs are likely because of the low sequence conservation. By contrast, in only two of the six class II CPuORFs, the fs mutations exhibited a significant effect on main ORF expression. In particular, the CIPK23 and DIC1 CPuORFs showed no sequence-dependent effect, despite their highly conserved amino acid sequences (Figure 2C) (23). These observations suggest that class I CPuORF peptides tend to possess a regulatory function that controls main ORF expression, compared with class II CPuORF peptides. However, in all the transient assays performed in this study, the protoplasts were cultured under normal culture conditions; therefore, we cannot rule out the possibility that the CPuORFs that showed little or no sequence-dependent effect may exert their effects under certain specific conditions. In fact, many of the previously reported sequence-dependent regulatory uORFs repress main ORF translation in response to metabolites, such as polyamine, arginine and sucrose (15–18, 46). Therefore, it is possible that the CPuORFs that showed little or no sequence-dependent effect in this study may require a metabolite as an effector molecule to exert their effects, and that the cellular concentration of the metabolite was not sufficient under our conditions.

Possible mechanisms of gene expression control by the newly identified regulatory uORF peptides

The co-transfection assays in Figure 4 revealed that the peptides encoded by the five newly identified sequence-dependent uORFs act in cis to repress main ORF expression, suggesting that these uORF peptides function as regulatory nascent peptides. The most likely underlying mechanism for the repression is that the nascent peptides encoded by these uORFs cause ribosome stalling. Genetic and biochemical studies have revealed that some regulatory nascent peptides cause ribosome stalling by interacting with components of the ribosomal exit tunnel (7, 9, 25–31). Cryo-electron microscopy studies have observed how regulatory nascent peptides interact with exit tunnel components. In prokaryotes, several of the C-terminal 16 and 11 amino acid residues in the TnaC and SecM nascent peptides interact with exit tunnel components when ribosome stalling occurs (54, 55). In eukaryotes, several of the C-terminal 16 and 19 amino acid residues in the gpUL4 and arg-2 uORF-encoded nascent peptides interact with exit tunnel components (24). Ala scanning mutagenesis in this study revealed that, in the five newly identified regulatory uORF peptides, the regions comprising 12 to 26 amino acid residues in the highly conserved regions have a pivotal role for the repression of main ORF expression (Figure 5). Thus, the lengths of the crucial regions in these uORF peptides are roughly consistent with those of the regions interacting with the exit tunnel components in the previously characterized regulatory nascent peptides. In some of the newly identified uORF peptides, an Ala substitution in the weakly conserved region located upstream of the highly conserved region showed a weak effect on reporter expression (Figure 5). This suggested that the weakly conserved regions might have an accessory role in ribosome stalling. Alternatively, because a eukaryotic ribosomal exit tunnel holds 30–40 amino acid residues (56, 57) and, therefore, the weakly conserved regions in the nascent uORF peptides should be inside the exit tunnel when ribosome stalling occurs, the Ala substitutions in the weakly conserved regions may have affected the structure or position of the crucial region in the nascent peptide to some extent, resulting in a slight impairment of the uORF peptide function.

The Ala scanning mutagenesis also indicated that, in the peptides encoded by ANAC082, CIPK6, At3g15430 and At1g27920 CPuORFs, all of which belong to class I, the crucial regions are located at the C-terminus. By contrast, the deletion analysis of the OTLD1 CPuORF, which belongs to class II, revealed that the C-terminal five amino acid residues are essentially dispensable for the repression of main ORF expression (Figure 7B). If the OTLD1 CPuORF-encoded peptide causes ribosome stalling by interacting with exit tunnel components, it is unlikely that the ribosome is stalled at the uORF stop codon as seen in the gpUL4 and arg-2 uORFs, because deletion of the C-terminal five amino acid residues, which would change the position of the crucial region of the uORF-encoded pep-
tide in the exit tunnel if ribosome stalling occurs at the stop codon, had little effect (Figure 7B). Additionally, elimination of the stop codon, which changed the stop codon position to one six codons downstream from the original position, did not affect the sequence-dependent inhibitory effect of the OTLD1 CPuORF (Figure 7D). Therefore, it is more likely that ribosome stalling occurs at the translation elongation step before the 3′-terminal non-conserved region of the OTLD1 CPuORF. In prokaryotes, there are examples of nascent peptide-mediated translation elongation arrest, in which ribosomal stalling occurs in the middle of small ORFs (1, 4). In eukaryotes, the regulatory nascent peptide encoded by the A. thaliana CGS1 gene causes ribosomal stalling in the middle of the main ORF (58, 59). However, in all of the previously characterized regulatory uORFs in which the stall position has been determined (i.e. the gpUL4, arg-2, CPA1, mammalian AdoMetDC and A. thaliana AdoMetDC1 uORFs), ribosome stalling mainly occurs at the uORF stop codon (14–18, 60), although the arg-2 uORF-encoded peptide can cause ribosome stalling at the translation elongation step if the uORF start codon is removed (61). Therefore, the OTLD1 CPuORF-mediated regulation may involve, at least in part, a distinct mechanism from the regulation mediated by the previously characterized native uORF peptides.

In the analysis of Figure 3, eliminating the start codon of the At5g27920 and OTLD1 CPuORFs caused stronger derepression than the corresponding fs mutation; whereas, in the ANAC082, CIPK6 and At3g15430 CPuORFs, start codon elimination and the fs mutation showed a similar derepressing effect. The presence of a uORF can have an inhibitory effect on main ORF translation, regardless of its peptide function, if the uORF is translated and ribosomes dissociate after translation. It is likely that the fs mutant version of the At5g27920 and OTLD1 CPuORFs had a repressive effect to some extent by this mechanism. By contrast, the results shown in Figure 3A–C suggest that, for ANAC082, CIPK6 and At3g15430, the presence of the fs mutant CPuORF had little or no effect on main ORF translation. One possibility to explain this observation is that translational initiation efficiencies of these CPuORFs are low and therefore most scanning ribosomes bypass these CPuORFs. Another possibility is that ribosomes that had translated these CPuORFs efficiently reinitiate translation at the main ORF. However, this latter possibility is unlikely, because reinitiation efficiency depends on the length of the uORF, and the ANAC082, CIPK6 and At3g15430 CPuORFs, whose sizes are 114, 99 and 147 nt, respectively, are too long for efficient reinitiation to occur (41). By contrast, the former possibility is more likely for the following reason. In A. thaliana, a purine (A or G) at position −3 and a guanine at position +4, where the A of AUG is defined as +1, are the optimal context for efficient translation initiation (62, 63), as established in mammals by Kozak (52, 53). The initiation contexts of the At5g27920 and OTLD1 CPuORFs are partially consistent with the optimal context, whereas those of the ANAC082 and At3g15430 CPuORFs are completely inconsistent with the optimal context (Figure 1). Although the initiation context of the CIPK6 CPuORF is consistent with the optimal context, another uORF overlaps with the CPuORF and has its start codon upstream of the CPuORF (Figure 1, Supplementary Figure S1E); therefore, it is likely that the presence of the overlapping uORF reduces translation initiation efficiency of the CIPK6 CPuORF. In addition, the former possibility is also supported by the observation that, in the Saccharomyces cerevisiae CPA1 uORF, whose translation initiation efficiency is low (64), a missense mutation impairing the regulatory function of the uORF peptide showed a similar derepressing effect to removal of the uORF start codon in yeast cells (65). It has been shown in vitro that scanning ribosomes frequently bypass the CPA1 uORF and reach the main ORF start codon, and that they are blocked when ribosome stalling occurs at the uORF in response to arginine (64).

The main ORFs regulated by the five newly identified sequence-dependent uORFs encode proteins involved in the control of gene expression or protein activity. ANAC082 encodes a NAC (NAM, ATAF1,2 and CUC2) domain-containing transcription factor. CIPK6 encodes a serine/threonine protein kinase that modulates the activity of a potassium channel, AKT2 (66), and is involved in the response to salt and osmotic stresses (67). At3g15430 encodes a protein related to the mammalian regulator of chromosome condensation, RCC1, which is the Ran guanine-exchange factor that regulates nuclear transport and mitotic spindle formation (68). At5g27920 encodes an F-box family protein. OTLD1 encodes an otubain-like histone deubiquitinase, which is involved in transcriptional repression via histone deubiquitination (69). Among the previously reported eukaryotic regulatory nascent peptides whose regulatory roles have been elucidated, many of them control the expression of metabolic enzyme genes and act in feedback regulation of the metabolic pathways (15–17, 34, 44). By contrast, the genes regulated by the uORF peptides identified in this study do not include any metabolic enzyme genes. The five newly identified regulatory uORF peptides had an inhibitory effect under normal culture conditions; therefore, these uORF peptides are likely to repress main ORF expression constitutively or in response to a metabolite that is present at a sufficient level in MM2d protoplast cells cultured under normal conditions. Even in the case where the uORF-encoded peptides always repress main ORF expression when the uORFs are translated, these uORFs can be involved in conditional regulation of gene expression if the translational initiation efficiencies of these uORFs are conditionally modulated. Such conditional modulation of translational initiation efficiencies of sequence-dependent regulatory uORFs has been reported in the human CHOP and A. thaliana AdoMetDC1 genes. In the CHOP gene, scanning ribosomes frequently bypass the sequence-dependent inhibitory uORF under endoplasmic reticulum stress condition, and thereby translation of the main ORF is enhanced (70). In the A. thaliana AdoMetDC1 gene, an overlapping uORF, whose start codon is located upstream of the sequence-dependent inhibitory uORF, was suggested to be involved in polyamine-responsive modulation of the translational initiation efficiency of the inhibitory uORF (44). As mentioned above, there is also an overlapping uORF upstream of the CIPK6 CPuORF (Figure 1, Supplementary Figure S1E). The size of the overlapping uORF and its position
relative to the CPuORF are evolutionarily conserved. Additionally, it has been suggested that expression of the chick-
pea CIPK6 ortholog is regulated in response to salt stress at both the transcriptional and post-transcriptional level (71). Therefore, the CIPK6 uORFs might be involved in the post-
transcriptional regulation in response to salt stress.

CONCLUSIONS

The present study showed that at least five of the 16 CPuORFs tested encode regulatory peptides. In \textit{A. thaliana}, 43 homology groups of CPuORFs have been identified (19,21,23) and two of them have been reported to exert a peptide sequence-dependent effect on main ORF expression (44,46). If the remaining CPuORFs contain peptide sequence-dependent regulatory uORFs at the same ratio as seen in this study, there would be at least 15 homology groups of sequence-dependent regulatory uORFs in the \textit{A. thaliana} genome. As mentioned above, it is possible that the CPuORFs that showed only weak or no significant effect in this study may exert a strong effect under certain conditions. If that is the case, even higher numbers of sequence-dependent regulatory uORFs would exist in the \textit{A. thaliana} genome. In addition, the genes regulated by the peptide sequence-dependent uORFs identified in this study comprise a variety of regulatory genes. Thus, this study suggests that uORF peptide-mediated gene regulation is more prevalent than previously thought and is involved in the control of a wide variety of genes. Further studies on the regulatory roles of the identified uORFs will reveal novel roles of the uORF-encoded regulatory peptides.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

ACKNOWLEDGEMENTS

We thank Ms Hitomi Sekihara, Ms Eriko Tanaka and Ms Saeko Yasokawa for skillful technical assistance and Ms Maki Mori for general assistance. We used the DNA sequencing facility of the Graduate School of Agriculture, Hokkaido University.

FUNDING

Japan Society for the Promotion of Science [Grant-in-aid for Scientific Research (C) 25440122 to H.O.]; Ministry of Education, Culture, Sports, Science and Technology of Japan [Grant-in-aid for Scientific Research on Innovative Areas 21112501 to H.O., 22119006 to S.N.]. Funding for open access charge: Japan Society for the Promotion of Science [Grant-in-aid for Scientific Research (C) 25440122 to H.O.].

Conflict of interest statement. None declared.

REFERENCES


