

Upstream open reading frames cause widespread reduction of protein expression and are polymorphic among humans

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Upstream ORFs (uORFs) are mRNA elements defined by a start codon in the 5' UTR that is out-of-frame with the main coding sequence. Although uORFs are present in approximately half of human and mouse transcripts, no study has investigated their global impact on protein expression. Here, we report that uORFs correlate with significantly reduced protein expression of the downstream ORF, based on analysis of 11,649 matched mRNA and protein measurements from 4 published mammalian studies. Using reporter constructs to test 25 selected uORFs, we estimate that uORFs typically reduce protein expression by 30–80%, with a modest impact on mRNA levels. We additionally identify polymorphisms that alter uORF presence in 509 human genes. Finally, we report that 5 uORF-altering mutations, detected within genes previously linked to human diseases, dramatically silence expression of the downstream protein. Together, our results suggest that uORFs influence the protein expression of thousands of mammalian genes and that variation in these elements can influence human phenotype and disease.

polymorphism | post-transcriptional control | proteomics | translation | uORF

The regulation of gene expression is controlled at many levels, including transcription, mRNA processing, protein translation, and protein turnover. Posttranscriptional regulation is often controlled by short sequence elements in the UTRs of mRNA. One such 5' UTR element is the upstream ORF (uORF) depicted in Fig. 1A. Because eukaryotic ribosomes usually load on the 5' cap of mRNA transcripts and scan for the presence of the first AUG start codon, uORFs can disrupt the efficient translation of the downstream coding sequence (1, 2). Previous reports have shown that ribosomes encountering a uORF can (i) translate the uORF and stall, triggering mRNA decay, (ii) translate the uORF and then, with some probability, reinitiate to translate the downstream ORF, or (iii) simply scan through the uORF (2). uORFs have been shown to reduce protein levels in ≈ 100 eukaryotic genes [supporting information (SI) Table S1]. Additionally, mutations that introduce or disrupt a uORF have found to cause 3 human diseases (3–5). In several interesting cases, the uORF-derived protein is functional; however, in most cases, the mere presence of the uORF is sufficient to reduce expression of the downstream ORF (1, 2, 6–8). Previous genomic analyses suggest that uORFs may be widely functional for several reasons: They correlate with lower mRNA expression levels (9), they are less common in 5' UTRs than would be expected by chance (6, 10), they are more conserved than expected when present (6), and several hundred have evidence of translation in yeast (11). However, no study has demonstrated that these elements have a widespread impact on cellular protein levels. Moreover, no study has investigated whether uORF presence varies in the human population. Here, we take advantage of recently available datasets of protein abundance (12–17) and genetic variation (18, 19) to assess the impact and natural variation of mammalian uORFs.

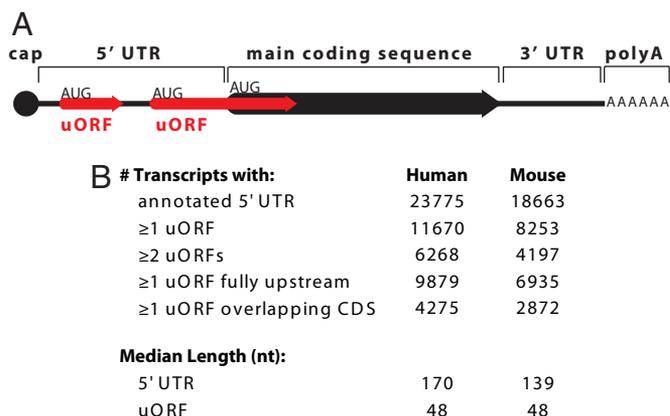


Fig. 1. uORF definition and prevalence. (A) Schematic representation of mRNA transcript with 2 uORFs (red arrows), 1 fully upstream and 1 overlapping the main coding sequence (black arrow). uORFs are defined by a start codon (AUG) in the 5' UTR, an in-frame stop codon (arrowhead) preceding the end of the main coding sequence, and length ≥ 9 nt. (B) Number and length of uORFs in human and mouse RefSeq transcripts.

Results

uORF Prevalence Within Mammalian Transcripts. We define a uORF as formed by a start codon within a 5' UTR, an in-frame stop codon preceding the end of the main coding sequence (CDS), and length at least 9 nt including the stop codon. As shown in Fig. 1A, this definition includes uORFs both fully upstream and overlapping the CDS, because both types are predicted to be functional (20). We searched for uORFs within all human and mouse RefSeq transcripts with annotated 5' UTRs >10 nt. Consistent with previous estimates (9, 10), we find that 49% of human and 44% of mouse transcripts contain at least 1 uORF (Fig. 1B). Interestingly, human and mouse uORF start codons (uAUGs) are the most conserved 5' UTR trinucleotide across vertebrate species (Fig. S1), consistent with a widespread functional role.

uORF Impact on Cellular Protein Levels. If uORFs cause widespread reduction in protein expression, as predicted by ribosome scanning

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models, we would expect uORF-containing transcripts to correlate with lower protein levels when compared with uORF-less transcripts. To test this hypothesis, we analyzed a total of 11,649 matched mRNA and protein abundance measurements from 4 published studies across a variety of mouse tissues and developmental stages. These included: 2,484 genes expressed in liver (12), 722 genes expressed in 6 stages of lung development (13), 487 mitochondria-localized gene products expressed in 14 tissues (14), and 925 genes expressed in 6 tissues (15) (see *SI Text* for details). Proteins were detected via tandem mass spectrometry (MS/MS), and abundance was estimated by standard methods using the normalized number (12, 13, 15) or total peak area (14) of matching MS spectra. mRNA abundance in these conditions was measured by microarrays (21, 22). Although neither technology provides absolute quantitation, these large-scale datasets can reveal trends across thousands of genes. Because MS/MS technology cannot reliably distinguish splice variants, we analyzed expression at the gene level and considered only those genes whose collective splice variants either all contain, or all lack, uORFs. Consistent with previous reports (23), we observed that the 10% most highly expressed transcripts based on microarray tissue atlases (21) tend to lack uORFs (Fig. S2 and *SI Text*), and therefore, we conservatively excluded these genes to avoid overestimating uORF effects.

Despite differences in experimental methodology, all 4 independent datasets showed a reduced distribution of protein levels for genes containing versus lacking uORFs (Fig. 2 A–D). Median protein levels were reduced, respectively, by 39% ($P = 1e-5$), 29% ($P = 0.007$), 34% ($P = 0.008$), and 13% ($P = 0.36$), where significance was determined by empirical permutation testing. mRNA levels were reduced to a lesser extent with only the liver dataset (12) showing a statistically significant median reduction (Fig. 2E and Fig. S3). Importantly, the ratio of protein to mRNA was significantly reduced for uORF-containing genes in 3 of 4 datasets (Fig. 2E and Fig. S3), suggesting that uORF presence likely inhibits translation of the main coding sequence. We observed the same trends when we modified the definition of a uORF by altering length and overlap criteria, and when we included the 10% most highly expressed genes (Fig. S4). Analysis of 2 additional MS/MS studies of mouse adipocyte cells (16) and differentiating embryonic stem cells (17) also showed reduced protein levels for uORF-containing genes, although matched mRNA data were not available (Fig. S3). Collectively, these analyses across 3,297 mouse genes demonstrated the first large-scale correlation of uORF presence with reduced protein levels.

To determine whether uORFs play a causal role in reducing protein levels, and to more accurately quantify their effect size, we performed a series of experiments on 15 uORF-containing genes using dual-luciferase reporter constructs (see *Materials and Methods*). Five genes were chosen randomly from the set of all mouse transcripts containing single uORFs and where, for technical ease, 5' UTR length exceeded 100 nt (Fig. 3 B and F). An additional 10 were selected from our mitochondrial study (14) where MS/MS and conservation data suggested functionality (Fig. 3 C and G). We cloned the 5' UTR of each selected gene upstream of a luciferase reporter (Fig. 3A). HEK 293A cells were then transfected with uORF-containing luciferase constructs or control constructs where the uORF's start codon (ATG) was mutated to TTG. After 48 h, cells were assayed for luciferase transcript levels by quantitative PCR and for luciferase activity by luminometry. These experiments showed that, on average, uORFs cause a 58% decrease in protein levels (Fig. 3 B and C) and a 5% decrease in transcript levels (Fig. 3 F and G). All individual protein differences and 4 mRNA differences were statistically significant (Fig. 3), and all protein/mRNA ratio differences were statistically significant except for gene *Hsd12* (Table S2). The constructs with randomly selected uORFs showed higher protein levels compared with the uORFs selected with evidence of functionality ($P = 1e-5$ based on *t* test). Similar results were obtained using HEK 293T cells. Together, the

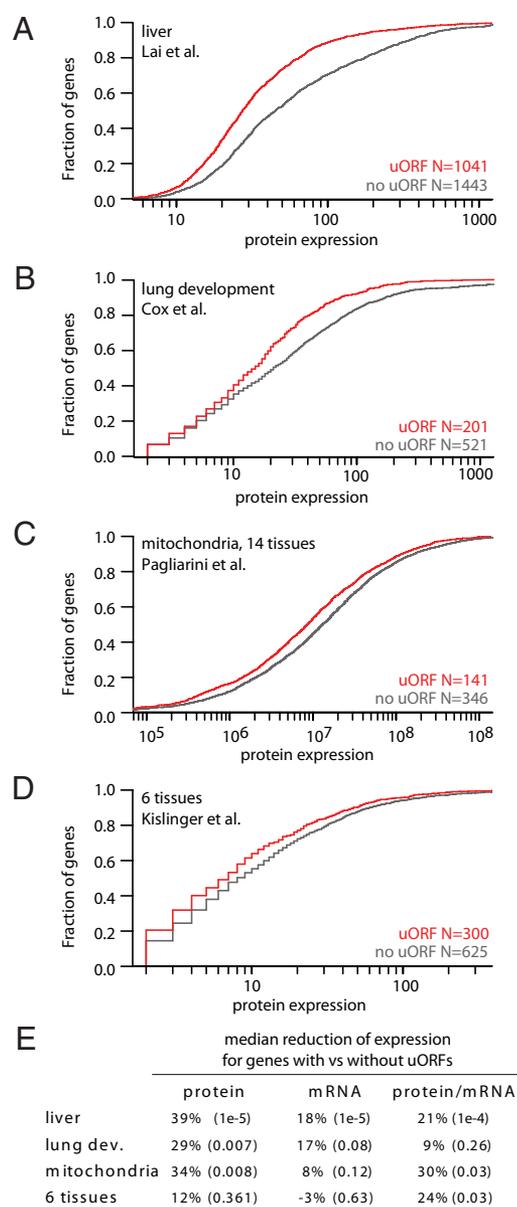


Fig. 2. Protein expression of uORF-containing genes. (A–D) Cumulative distribution of protein expression for mouse genes containing uORFs (red curve) or lacking uORFs (gray curve) in each of 4 independent MS/MS studies (12–15). N indicates the number of unique genes in each set. (E) Median reduction of protein and mRNA expression for genes containing uORFs compared with genes lacking uORFs, with *P* values (in parentheses) computed by empirical permutation testing.

large-scale correlations and validation experiments demonstrate that uORFs cause blunted protein expression of downstream coding sequences.

Influence of uORF Context, Position, and Conservation. We next investigated whether specific uORF properties were associated with stronger translational inhibition. We analyzed uORF length, number, conservation, position relative to the cap, position relative to the CDS, and uAUG context (also called “Kozak sequence”) (see *Materials and Methods*). We quantified uORF effects using the Kolmogorov–Smirnov (KS) *D* statistic within the largest dataset (liver), which offered statistical power for these analyses. All tested subsets of uORFs showed reduced protein levels compared with uORF-less genes ($P < 0.05$), although certain properties modified

Table 1. Notable human variants that create polymorphic uORFs

#	SNP	AvHet, %	Gene	Gene description
1	rs1801020	50	<i>F12</i>	Coagulation factor XII (Hageman factor)
2	rs12272467	50	<i>TRIM6</i>	Tripartite motif-containing 6
3	rs1108842	50	<i>GNL3</i>	Guanine nucleotide-binding protein-like 3 (nucleolar)
4	rs6460054	50	<i>CLDN3</i>	Claudin 3
5	rs1046188	50	<i>SCAMP3</i>	Secretory carrier membrane protein 3
6	rs13104310	49	<i>C4orf21</i>	Chromosome 4 open reading frame 21
7	rs7667298	49	<i>KDR</i>	Kinase insert domain receptor
8	rs7331765	49	<i>RASL11A</i>	RAS-like, family 11, member A
9	rs2001216	49	<i>RCCD1</i>	RCC1 domain containing 1
10	rs12975585	48	<i>HNRNPUL1</i>	Heterogeneous nuclear ribonucleoprotein U-like 1
11	rs2838343	46	<i>HSF2BP</i>	Heat shock transcription factor 2-binding protein
12	rs765007	46	<i>TAS2R3</i>	Taste receptor, type 2, member 3
13	rs17499247	45	<i>CREM</i>	cAMP responsive element modulator
14	rs1048371	42	<i>MUCL1</i>	Mucin-like 1
15	rs1800070	*	<i>CFTR</i>	Cystic fibrosis transmembrane conductance regulator
16	rs34704828	*	<i>HBB</i>	Hemoglobin, β
17	rs28926176	0.2	<i>MC2R</i>	Melanocortin 2 (ACTH hormone) receptor
18	rs41409645	4	<i>CCL3</i>	Chemokine (C-C motif) ligand 3
19	rs2856759	*	<i>CCR5</i>	Chemokine (C-C motif) receptor 5
20	rs34819868	*	<i>HAVCR1</i>	Hepatitis A virus cellular receptor 1
21	rs41275166	*	<i>CD59</i>	CD59 molecule, complement regulatory protein
22	rs6057688	*	<i>DEFB119</i>	Defensin, β 119
23	rs2234011	*	<i>TAS2R5</i>	Taste receptor, type 2, member 5
24	rs1091826	*	<i>OXTR</i>	Oxytocin receptor
25	rs6781226	*	<i>HTR1F</i>	5-hydroxytryptamine (serotonin) receptor 1F

List contains common SNP variants (#1–14) and genes associated with monogenic disease (#15–17), immune response (#18–22), and receptor activity (#23–25 and 7, 12, 17, 19, 20). Table S3 contains a complete list. AvHet indicates SNP's average heterozygosity.

*Data not available.

in the remaining case (36). To our knowledge, the mechanistic link between the gene mutation and uORFs had not been previously proposed for *SRY* (32), *IRF6* (33), or *GCH1* (34).

To assess whether the uORF-altering mutations influenced protein expression, we used luciferase reporter constructs to test patient mutations in 5 genes (*HBB*, *PRKARIA*, *IRF6*, *SRY*, and *SPINK1*). The uORF-altering mutations in these genes reduced luciferase mRNA levels by <20% and luciferase activity levels by 70–100% (Fig. 3 E and I). These effects on protein levels were highly significant ($P < 2e-12$) and were larger than in the other uORFs experimentally tested ($P = 4e-4$). Thus, these uORF-altering mutations cause dramatically reduced protein levels in our reporter assays, suggesting that they may indeed be responsible for the observed disease phenotypes.

Discussion

Our analyses provide an assessment of the widespread impact of uORFs on mammalian protein expression. Many previous studies

of individual genes demonstrated that the presence of uORFs can lead to reduced mRNA stability and protein translation. Here, we show that approximately half of human and mouse protein-encoding genes contain uORFs and that uORF presence correlates with reduced protein expression across thousands of mammalian genes in a variety of tissues and conditions (Fig. 2). We quantify uORF effects using mutation experiments on 25 selected 5' UTRs (Fig. 3), which have typical length, context, position, and conservation features (Fig. S6). These experiments indicate that uORFs typically affect mRNA levels by <30% and reduce protein levels by 30–80%, although complete protein suppression is possible (Fig. 3). Although our mutation experiments focused chiefly on 5' UTRs containing single uORFs, our MS/MS data suggest that multiple uORFs lead to greater reduction of protein expression (Fig. S5E). Collectively, these data suggest that uORFs cause reduced protein levels of thousands of mammalian genes.

Our data provide insight into the mechanism by which uORFs influence protein expression. Without exception, uORF-containing

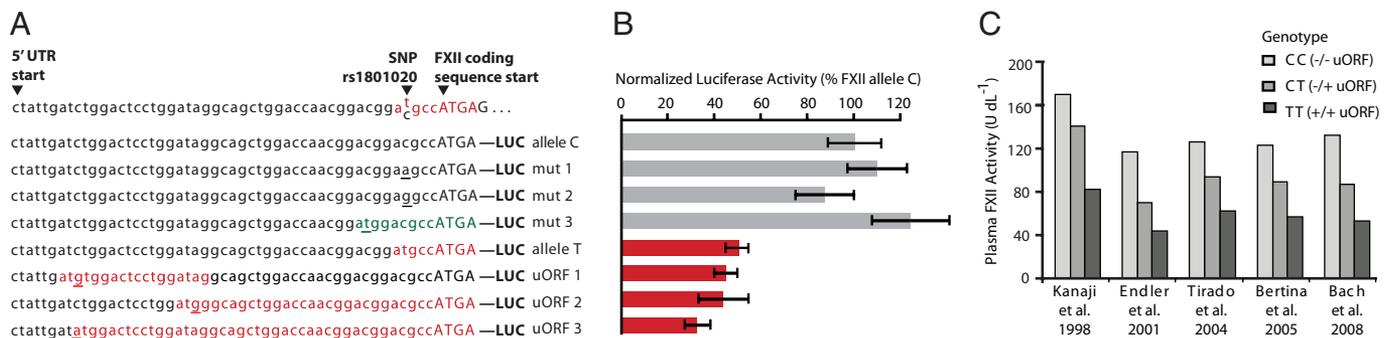


Fig. 4. Polymorphic uORF alters FXII protein expression. (A) 5' UTR sequence of FXII shown with 2 SNP variants, where the T allele creates a uORF (red text). Below are 8 constructs with introduced mutations (underlined text), where colored text indicates a uORF (red) or in-frame alternative start (green). (B) Luciferase activity from reporter constructs listed in A. Error bars represent \pm SD of ≥ 6 biological replicates. (C) Metaanalysis of plasma FXII activity levels measured by 5 independent studies, stratified by genotype of SNP rs1801020.

Table 2. uORF-altering mutations linked to disease

#	Gene	Disease	Mutation	uORF link
1	<i>THPO</i>	Thrombocythemia	splice site (3)	Known
2	<i>CDKN2A</i>	Melanoma	G-34T (4)	Known
3	<i>HR</i>	Marie Unna hereditary hypotrichosis	A-321G (5)	Known
4	<i>SRY</i>	Gonadal dysgenesis	G-75A (32)	Novel*
5	<i>IRF6</i>	Van der Woude syndrome	A-48T (33)	Novel*
6	<i>GCH1</i>	DOPA-responsive dystonia	C-22T (34)	Novel
7	<i>HAMP</i>	Juvenile hemochromatosis	G-25A (35)	Predicted
8	<i>KCNJ11</i>	Hyperinsulinemic hypoglycemia, 2	C-54T (36) [†]	Predicted
9	<i>LDLR</i>	Familial hypercholesterolemia	delC-22 (37) [†]	Predicted
10	<i>PEX7</i>	Rhizomelic chondrodysplasia punctata	C-45T (38) [†]	Predicted
11	<i>POMC</i>	Proopiomelanocortin deficiency	C-11A (39)	Predicted
12	<i>PRKAR1A</i>	Carney complex type 1	G-97A (40)	Predicted*
13	<i>SPINK1</i>	Hereditary pancreatitis	C-53T (41)	Predicted*
14	<i>HBB</i>	Thalassaemia β	G-29A (42) [†]	Predicted*

uORF-altering mutations detected in patients but not population controls. Mutation column includes 5' UTR position relative to translation start and literature reference (in parentheses). The links between mutations and uORFs were previously known, previously predicted, or not previously known (novel).

*Mutations tested experimentally in this study.

[†]Compound heterozygous mutations.

reporter constructs exhibit more pronounced reduction of protein compared with mRNA levels (Fig. 3), in agreement with the trend observed in large-scale datasets (Fig. 2E). This suggests that uORFs act primarily by reducing translational efficiency, and more modestly by affecting mRNA levels. Additionally, because uORF effects do not correlate with the distance between the uORF and CDS (Fig. S5D), it is likely that CDS translation generally proceeds from ribosomes that scan through the uORF rather than from ribosomes that reinitiate after uORF translation—at least in genes that contain only a single uORF.

Given that uORFs reduce translation, variants that create or delete uORFs are likely to alter cellular protein levels and in some cases may influence phenotype. uORF-altering variants are likely to be widespread, because each human transcript contains, on average, 28 nt that could be mutated to introduce a uORF. We identified 509 human genes with polymorphic uORFs (puORFs), although more are likely to be identified as genome variation databases expand. Our data suggest that puORFs will typically alter cellular protein levels by 30–80% in cases where the 5' UTR contains a single uORF. When these puORFs cause physiologically relevant changes in protein levels, as we showed for factor XII, they may cause phenotypic variation. Indeed, the factor XII puORF has been associated with several thromboembolic conditions, although the associations are in contention due to small sample sizes (44). We speculate that other puORFs in our collection (Table S3) may also affect phenotype. For instance, the puORF in chemokine receptor CCR5 might mediate susceptibility to HIV-1 infection, because previous studies showed that variants affecting CCR5 expression alter susceptibility to HIV-1 infection and progression of AIDS (45). Similarly, the puORFs in bitter-taste receptors TAS2R5 and TAS2R3 might lead to common variation in taste perception, and puORFs in receptors for ACTH, serotonin, and oxytocin may modulate neurohormonal response (Table 1).

In addition to common polymorphisms, rare uORF-altering mutations that alter levels of essential proteins can cause human disease. To date, 3 such mutations have been reported. First, a hereditary form of thrombocythemia is caused by a mutation in *THPO* mRNA that eliminates a uORF through a splicing defect, and thus causes increased translation of thrombopoietin (3). Second, a mutation introducing a uORF into *CDKN2A* causes a familial predisposition to melanoma (4). Third, disruption of uORF presence and coding sequence in gene *HR* causes Marie Unna hereditary hypotrichosis (5). Additional uORF-altering mutations detected in patients with 11 diseases have been reported in the

literature, although they were not followed up experimentally (Table 2). In each case, the patient mutation was present within a gene known to underlie the disease when disrupted and was the sole mutation detected or was compound heterozygous with a nonsynonymous variant. Using reporter assays, we tested 5 patient mutations in genes associated with disease: Gonadal dysgenesis (*SRY*), Van der Woude syndrome (*IRF6*), Carney complex type 1 (*PRKAR1A*), Hereditary pancreatitis (*SPINK1*), and Thalassaemia- β (*HBB*). We found that the uORF-altering patient mutation caused severely reduced protein levels, and in 2 cases almost no reporter protein was detected (*SRY* and *SPINK1*, Fig. 3E). In these 2 cases, the patient mutation created a second uORF within the gene 5' UTR, rather than creating a single uORF. The strong suppression of protein expression by these 5 patient mutations offers a simple mechanistic basis for their pathogenicity. These cases add to the growing list of uORF-altering mutations linked to disease and highlight the importance of searching for uORF changes in addition to coding changes underlying disease.

In summary, our analyses demonstrate that uORFs have a widespread impact on the expression of human and mouse genes and that the human genome contains hundreds of polymorphic uORFs. With the routine application of newer generation sequencing technologies, an important challenge will be to link variation in genome sequences to physiology and disease—and puORFs may represent an important class of functional variants that can be readily linked to phenotype. Although the current analyses focused on the constitutive effects of uORFs on steady-state protein levels, an important next step is to determine whether the influence of uORFs is widely regulated by environmental conditions or signaling pathways, as has been shown for a handful of examples (2).

Materials and Methods

Human and Mouse uORFs. RefSeq transcripts for human (hg18) and mouse (mm9) were obtained from the University of California, Santa Cruz (UCSC) Genome Browser Database (46) (accessed May 20, 2008), along with 28-vertebrate species alignments (47). Custom Perl scripts annotated uORFs and computed features: uORF context ("strong" indicates a -3 purine and $+4$ guanine relative to uAUG, otherwise "weak"), cap-to-uORF distance (length between mRNA cap and uAUG), uORF length (including start and stop codon), uORF-to-CDS distance (length between uORF stop codon and CDS start codon), uORF number (number of distinct uORFs in a transcript, where uORFs may overlap but not in the same frame), and conservation (number of species with aligned start codons within 28 species alignments). The first 4 features were analyzed on transcripts containing single uORFs. uORF properties were compared using a Bonferroni-corrected,

1-sided KS test. 5' UTR trinucleotide conservation was scored by number of identities in 28 species alignments.

Matched mRNA and Protein Datasets. MS/MS protein abundance measurements were obtained from published studies (12–17). Matched mRNA data were available in 3 studies (13–15). For the liver study (12), we used mean mRNA expression from GNF1M liver replicates (21). All data were mapped to Entrez Gene identifiers with the gene inheriting the highest score from any splice form. We excluded genes with poorly quantified mRNA values (expression values <40) and the top 10% most highly expressed genes, based on mean mRNA expression values from the GNF1M atlas. We analyzed mouse genes with annotated 5' UTRs (>10nt), where all splice forms contained a uORF (6,933 genes) or lacked a uORF (9,343 genes). Differences in median protein expression were measured as percentage reduction from uORF-less genes, using 10,000 permutations of gene uORF labels to assess significance. See *SI Text* for details.

Luciferase Assays. UTR sequences, up to and including the primary ATG initiation codon, were synthesized (IDT), cloned and ligated into the NheI site directly preceding the *Renilla* luciferase gene in the dual-luciferase vector psiCHECK-2 (Promega) (Table S4). Before cloning, the ATG of the *Renilla* luciferase was mutated to TTG so that the *Renilla* luciferase expression would be driven by the primary ATG initiation codon of the gene under investigation. HEK 293A cells were seeded at 6,000 cells per well in 96-well opaque white cell culture plates (Nunc). After overnight incubation, cells were transfected with 20–100 ng of each construct by using Fugene 6 (Roche). Forty-eight hours after transfection, cells were washed with PBS and lysed with Passive Lysis Buffer (Promega). *Renilla* and Firefly luciferase signals were generated by using Promega's Dual-Luciferase Assay System according to the manufacturer's protocol. For each construct, *Renilla* luciferase signal was normalized to the Firefly luciferase internal control signal. Plates were read by using a Victor³ plate reader (PerkinElmer) and the data

analyzed by using Wallac 1420 Workstation software. Two-sided, homoscedastic *t* tests assessed significance.

Real-Time PCR. HEK 293A cells were seeded at 2×10^5 cells per well in 6-well cell culture plates 24 h before transfection. One microgram of each construct was transfected per well by using Fugene 6 as above. Forty-eight hours after transfection, cells were washed with PBS, and RNA was harvested by using a Qiagen RNeasy kit. First-strand cDNA synthesis was performed by using SuperScript III (Invitrogen) using 1 μ g of RNA from each transfection as starting material. Real-time PCR was performed by using custom TaqMan Assays (ABI) designed against *Renilla* luciferase (target) and Firefly luciferase (endogenous control). Two-sided, homoscedastic *t* tests assessed significance.

uORF-Altering Variants. Human dbSNP version 128 (18) was obtained from UCSC (46) and filtered for SNPs (class "single") that mapped to single locations within hg18 and overlapped annotated RefSeq 5' UTRs, excluding those that overlapped RefSeq CDSs. Custom perl scripts mapped SNPs onto mRNA transcripts and determined those that altered uORF presence. The Human Gene Mutation Database professional release 2008.2 (19), was searched for all noncoding substitutions or microlesions that altered presence of ATG codons and that overlapped 5' UTRs based on manual inspection of BLAT alignments to hg18.

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