Structural insights into translational recoding by frameshift suppressor tRNA^{SufJ}

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Footnotes:

Running title: Structure of ASL^{SufJ} bound to the ribosomal A site

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Figure S2. Electron density maps for ASL^{SufJ} **bound to the A site**. Unbiased F_o - F_c electron density map for ASL^{SufJ} (blue) bound to the +1 suppressible codons (A) ACC-<u>A</u> (B) ACC-<u>U</u> and (C) ACC-<u>C</u> shows strong density for nucleotides 29-42 (contoured at 3 σ). The 70S structures containing the (B) ACC-<u>U</u> and (C) ACC-<u>C</u> codons contain the antibiotic paromomycin while the 70S structure containing the (A) ACC-<u>A</u> codon is without.



Figure S3. The mRNA path of each +1 suppressible codon is similar. The mRNA path is unaffected by the fourth nucleotide in the ACC-<u>A</u> (brown), ACC-<u>U</u> (dark blue) and ACC-<u>C</u> (green) codons. The P-site tRNA^{fMet} is in magenta.



Figure S4. ASL^{Thr} **interactions with mRNA in the A site.** ASL^{Thr} (gray) forms three Watson-Crick base pairs with the first three nucleotides of the +1 suppressible codons **(A)** ACC-<u>A</u> (brown) and **(B)** ACC-<u>C</u> (green) in the zero frame. These structures were solved with the antibiotic paromomycin. The ASL is stabilized by a hydrogen bonding with 23S rRNA nucleotide A1913 (tan).