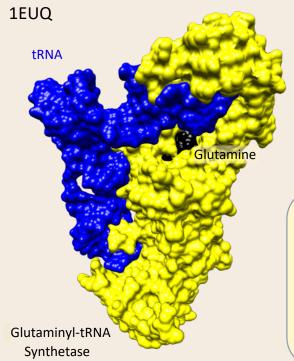
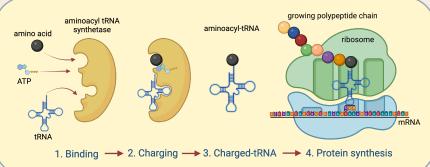
tRNA CHARGER



Aminoacyl-tRNA synthetase (AARS)

- an enzyme that attaches a particular amino acid onto its corresponding tRNA ("charging")
- charged tRNAs bring amino acids to ribosomes for production of proteins



STRUCTURE¹

- Class I: mostly monomeric; binds the minor groove of tRNA and attaches aminoacyl to the 2'-OH of tRNA; charge hydrophobic amino acids
- Class II: dimeric or multimeric; binds to major groove of tRNA and prefers 3'-OH; charge small and polar amino acids
- tRNA binding involves alpha helix shared by both classes
- consist of catalytic domain, anticodon recognition and editing domain

LOCATION

- ubiquitously expressed enzymes by all forms of life
- eukaryotic cells have AARSs in the cytoplasm and inside the mitochondria
- each of the 20 amino acids has its own specific AARS
- in bacteria mostly as freestanding proteins
- in archaea and eukaryotes
 AARSs assemble with
 accessory proteins into
 mutisynthetase complexes

FUNCTION

- link amino acids to their cognate tRNAs forming aminoacyl-tRNAs (aa-tRNAs)
- in the ribosome, the aa-tRNAs decode information from mRNA into a polypeptide
- also involved in regulation of gene expression, splicing of introns, and biosynthesis
- plays a critical role in the development and activation of immune system
- act as signaling molecules in autoimmune and infectious diseases

Disorders and Application to Medicine

- Autoimmune Antisynthetase Syndrome² autoantibodies present in the cytoplasm begin targeting tRNA synthetases, causing myositis, polyarthritis, and lung disease among other symptoms.
- Charcot-Marie-Tooth Disease³ a neurodegenerative disease; results from dominant, monoallelic mutations; described for 6 different AARSs; causes nerve damage in arms and legs and muscle weakness.
- Pharmacological targets for development of antibiotic therapies agains pathogenic bacteria and antifungal treatments⁴.

- Cavarelli, J., & Moras, D. (1993). Recognition of tRNAs by aminoacyl-tRNA synthetases. FASEB journal: official publication of the Federation of American Societies for Experimental Biology, 7(1), 79–86. https://doi.org/10.1096/fasebj.7.1.8422978
- Galani, K., Grosshans, H., Deinert, K., Hurt, E. C., & December 3). The intracellular location of two aminoacyl-trna synthetases depends on complex formation with arc1p. The EMBO journal. Retrieved February 6, 2022, from https://www.ncbi.nlm.nih.gov/pmc/articles/PMC125769/
- Jiang, L., Jones, J., & Diang, X.-L. (2020, September 28). Human diseases linked to cytoplasmic aminoacyl-trna synthetases. The Enzymes. Retrieved January 20, 2022, from https://www.sciencedirect.com/science/article/abs/pii/S1874604720300251
- McClain, W. H. (2002, May 25). Rules that govern trna identity in protein synthesis. Journal of Molecular Biology. Retrieved February 6, 2022, from https://www.sciencedirect.com/science/article/pii/S0022283683715822?via%3Dihub
- Figure 1 created with BioRender.com