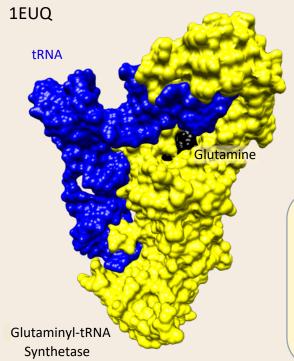
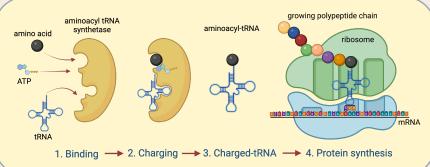
# **tRNA CHARGER**



# 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



## STRUCTURE<sup>1</sup>

- 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 Syndrome<sup>2</sup> autoantibodies present in the cytoplasm begin targeting tRNA synthetases, causing myositis, polyarthritis, and lung disease among other symptoms.
- Charcot-Marie-Tooth Disease<sup>3</sup> 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 treatments<sup>4</sup>.

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