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博士学位論文

Mechanistic insights into tRNA thiolation catalyzed by iron-sulfur enzymes

(鉄硫黄クラスター含有酵素が触媒する tRNA 硫黄修飾の反応機構解析)

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Major abbreviations

AA	Amino acid
AdetRNA	Activated tRNA: Adenylated tRNA
AF2	AlphaFold version 2
AMPPNP	ATP analog: Adenylyl-imidodiphosphate
APM	[(<i>N</i> -Acryloylamino) phenyl]mercuric chloride
APM-PAGE	APM-polyacrylamide gel electrophoresis
ATP	Adenosine triphosphate
<i>B. subtilis</i>	<i>Bacillus subtilis</i>
<i>Bsu</i> MnmA	MnmA from <i>Bacillus subtilis</i> strain 168
C α RMSD	Root-mean-square deviation of the alpha carbon
CBB	Coomassie brilliant blue
cmnm ⁵ s ² U	5-carboxymethylaminomethyl-2-thiouridine
CV	Column volume
DT	Reductant: Dithionite
DTT	Reductant: Dithiothreitol
<i>E. coli</i>	<i>Escherichia coli</i>
<i>Eco</i> MnmA	MnmA from <i>Escherichia coli</i> strain K12
<i>Eco</i> ThiI	ThiI from <i>Escherichia coli</i> strain K12
<i>Eco</i> TtcA	TtcA from <i>Escherichia coli</i> strain K12
EPR	Electron paramagnetic resonance
Fe-S	Iron-sulfur
Gm	2'- <i>O</i> -methylguanosine
<i>H. sapiens</i>	Human: <i>Homo sapiens</i>
HPLC	High-performance liquid chromatography
<i>Hsa</i> Ncs6	Ncs6 from <i>Homo sapiens</i>
<i>Hsa</i> Urm1	Urm1 from <i>Homo sapiens</i>
I34	Inosine at position 34 of tRNA (the wobble position)
IPTG	Isopropyl β -D-1-thiogalactopyranoside
KO	Knockout
LB	Lysogeny broth

m ⁵ s ² U	5-methyl-2-thiouridine
m ⁵ s ² U54	5-methyl-2-thiouridine at position 54 of tRNA
m ⁵ U	5-methyluridine
m ⁵ U54	5-methyluridine at position 54 of tRNA
<i>M. maripaludis</i>	<i>Methanococcus maripaludis</i> strain S2
MALDI-TOF MS	Matrix-assisted laser desorption/ionization-time-of-flight mass spectrometry
mcm ⁵ s ² U	5-methoxycarbonylmethyl-2-thiouridine
mcm ⁵ s ² U34	5-methoxycarbonylmethyl-2-thiouridine at position 34 of tRNA
MERRF	Mitochondrial disease: Myoclonic epilepsy with ragged-red fibers
mnm ⁵ s ² U	5-methylaminomethyl-2-thiouridine
<i>MmaNcs6</i>	Ncs6 from <i>Methanococcus maripaludis</i> strain S2
<i>MmaThiI</i>	ThiI from <i>Methanococcus maripaludis</i> strain S2
mRNA	Messenger ribonucleic acid
ms ² A	2-methylthioadenosine
ms ² ct ⁶ A	2-methylthio-cyclic- <i>N</i> ⁶ -threonylcarbamoyladenosine
ms ² i ⁶ A	2-methylthio- <i>N</i> ⁶ -isopentenyladenosine
ms ² t ⁶ A	2-methylthio- <i>N</i> ⁶ -threonylcarbamoyladenosine
ms ² x ⁶ A37	Modified 2-methylthioadenosine at position 37 of tRNA
MT	Mutant
mt-tRNA	Mitochondrial tRNA
Mw	Molecular weight
m/z	Mass-to-charge ratio
NiAC	Ni-affinity chromatography
OD ₆₀₀	Absorption at 600 nm
<i>P. horikoshii</i>	<i>Pyrococcus horikoshii</i>
PDB	Protein Data Bank
<i>PhoTtuA</i>	TtuA from <i>Pyrococcus horikoshii</i> strain OT-3
P _i	Inorganic phosphate: PO ₄ ³⁻
pLDDT	Predicted local-distance difference test
PPase	Pyrophosphatase
PP _i	Pyrophosphate: P ₂ O ₇ ⁴⁻

R ²	Coefficient of determination
RT	Room temperature: 25°C
s ² C	2-thiocytidine
s ² C32	2-thiocytidine at position 32 of tRNA
s ² U	2-thiouridine
s ² U34	2-thiouridine at position 34 of tRNA (the wobble position)
s ⁴ U	4-thiouridine
s ⁴ U8	4-thiouridine at position 8 of tRNA
<i>S. cerevisiae</i>	Budding yeast: <i>Saccharomyces cerevisiae</i>
SAM	S-adenosyl methionine
SD	Standard deviation
SDS	Sodium dodecyl sulfate
SDS-PAGE	Sodium dodecyl sulfate-polyacrylamide gel electrophoresis
SEC	Size exclusion chromatography
<i>T. thermophilus</i>	<i>Thermus thermophilus</i>
T _m	Melting temperature
TmaThiI	ThiI from <i>Thermotoga maritima</i> strain MSB8
tRNA	Transfer ribonucleic acid
tRNA ^{AA(NNN)}	tRNA that carry AA with anticodon of NNN
TthMnmA	MnmA from <i>Thermus thermophilus</i> strain HB27
TthTtuA	TtuA from <i>Thermus thermophilus</i> strain HB27
TtuB-COOH	Non-sulfur donor TtuB with carboxyl group at the C-terminus
TtuB-COSH	Sulfur donor TtuB with thiocarboxyl group at the C-terminus
UV-Vis	Ultra violet and visible light
UV280	Ultra violet at wavelength of 280 nm
v/v	Volume/Volume
WT	Wild type
xm ⁵ s ² U34	Modified 2-thiouridine at position 34 of tRNA (the wobble position)
τm ⁵ s ² U	5-taurinomethyl-2-thiouridine
Ψ	Pseudouridine

*Abbreviations of standard amino acids and bases comply with international rules.

Abstract

tRNA transfers amino acids to ribosomes to translate genetic information into proteins. However, immature tRNA does not function immediately after transcription. Hence, tRNA undergoes post-transcriptional processes such as base/ribose modifications for maturation. To date, more than 110 base/ribose modifications have been discovered in tRNA. Particularly, thiolation (sulfur modification) is a universal and essential enzymatic reaction that improves the thermal stability and translational accuracy of tRNA.

My target is thiolation at position 54 of tRNA (5-methyl-2-thiouridine, m^5s^2U54), which is essential for thermophiles to survive above 70°C. m^5s^2U54 modification is catalyzed by 2-thiouridine synthetase TtuA with sulfur donor protein TtuB in *Thermus thermophilus*. Our recent structural analysis of the TtuA-TtuB complex showed that an oxygen-sensitive [4Fe-4S] cluster is required for the enzymatic activity of TtuA. Interestingly, a non-cysteine coordinated Fe (the unique Fe) of the [4Fe-4S] bound to the C-terminus of TtuB. This structure suggested that the unique Fe in TtuA relates to the sulfur transition from TtuB to tRNA, which is a novel reaction mechanism of m^5s^2U54 biosynthesis involving the unique Fe. On the other hand, TtuA homolog enzyme Ncs6 catalyzes thiolation at position 34 of tRNA (mcm^5s^2U34) with sulfur donor Urm1, which is similar to TtuA. However, a spectroscopic study indicated that Ncs6 contains [3Fe-4S], whereas crystallography supported [4Fe-4S]. Therefore, it was unclear whether the active form of tRNA-thiolation enzymes only require [4Fe-4S] or both [4Fe-4S] and [3Fe-4S].

In this study, I analyzed the structural change of Fe-S clusters in TtuA in time-course and evaluate their enzymatic activity. As a result, [3Fe-4S] spontaneously transformed into [4Fe-4S] even without the additional iron source, and the activity of TtuA gradually recovered corresponding to an increase in [4Fe-4S]. I also revealed that [3Fe-4S]-TtuA cannot bind to the C-terminus of TtuB, indicating that only [4Fe-4S]-TtuA is an active form. Furthermore, I found that TtuB does not release sulfur until tRNA is activated by ATP, and identified the critical residues of TtuA. Considering the similarity, I proposed that [4Fe-4S] is generally an active form in tRNA-thiolation enzymes and the detailed tRNA-thiolation mechanism catalyzed by TtuA. These findings showed that the time-course analysis of the structure and activity under strictly anaerobic conditions is necessary to elucidate the reaction mechanism of enzymes containing Fe-S clusters.

Chapter 1. Introduction

1-1. Post-transcriptional modifications of RNA

In protein biosynthesis, various ribonucleic acids (RNA) are involved such as transfer RNA (tRNA), messenger RNA (mRNA), and ribosomal RNA (rRNA). These RNAs do not function immediately after transcription *in vivo*. Thus, immature RNA undergoes enzymatic reactions called post-transcriptional processes including splicing and modifications for maturation. Among these processes, more than 150 kinds of base/ribose modifications have been discovered in RNA, and these modifications are the most diverse in tRNA^[1]. Recently, the relationship between abnormalities in tRNA modifications and diseases has been reported^[2,3]. Therefore, it is significant to understand the molecular mechanisms of tRNA maturation.

1-2. tRNA modifications

During translation, tRNA transfers amino acids to ribosomes according to their anticodon for protein biosynthesis. Immature tRNA undergoes post-transcriptional processes such as splicing, CCA addition, ribose/base modification, and aminoacylation (Fig. 1-1)^[4]. For the ribose/base modification, more than 110 kinds of tRNA modifications have been identified such as methylation, acetylation, and thiolation^[1]. In particular, diverse base modifications are found at position 1 of tRNA anticodon (position 3 of the codon), and loss of these modifications reduces the translation rate, causing ribosomal arrest and protein misfolding/aggregation^[5].

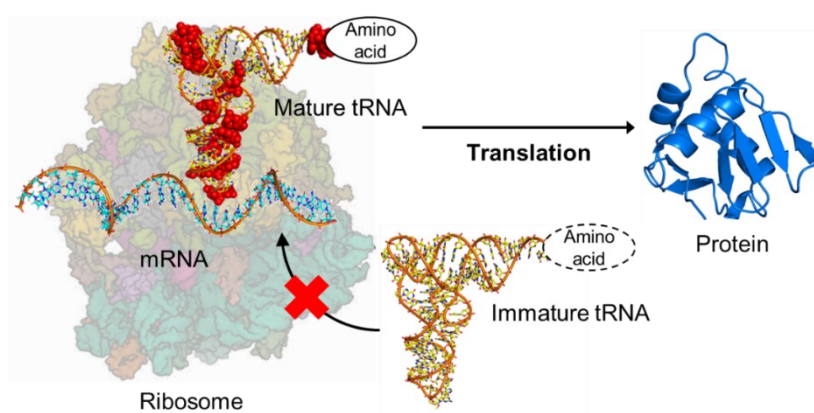


Fig. 1-1 The role of tRNA modifications in protein synthesis.

Red balls show modification sites on tRNA.

1-3. tRNA thiolation (sulfur modification)

tRNA thiolations are enzymatic modifications that replace the oxygen atom with a sulfur atom in the specific site in tRNA, which have been roughly classified into four types: 2-thiouridine (s^2U), 4-thiouridine (s^4U), 2-thiocytidine (s^2C), and 2-methylthioadenosine (ms^2A)^[6]. These modifications are found in a specific position of tRNA and play universal roles in bacteria, archaea, and eukaryotes (Fig. 1-2, 1-3). tRNA thiolation improves the thermal stability of tRNA^[7,8], translational speed and accuracy^[9–11], and is responsible for near-UV sensing^[12,13] depending on its modification site.

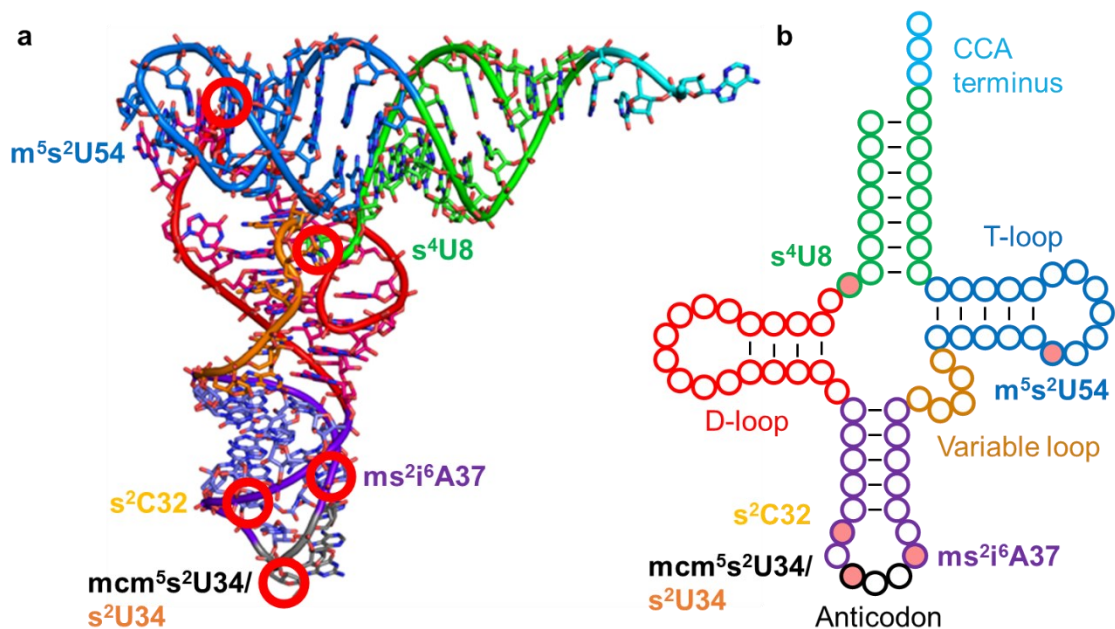


Fig. 1-2 Structures of tRNA.

(a) L-shaped 3D structure of tRNA (PDB ID: 1EHZ). The thiolation sites are indicated with red circles and the names of typical tRNA thiolation; 5-methoxycarbonylmethyl-2-thiouridine (mcm^5s^2U), 2-methylthio- N^6 -isopentenyladenosine (ms^2i^6A), and 5-methyl-2-thiouridine (m^5s^2U). The numbers following the name of the thiolated base are the base number counted from the 5' end of tRNA. (b) The clover-leaf model of the 2D structure of tRNA. The length of the variable loop varies depending on the species and type of tRNA. The color code is the same as the 3D structure.

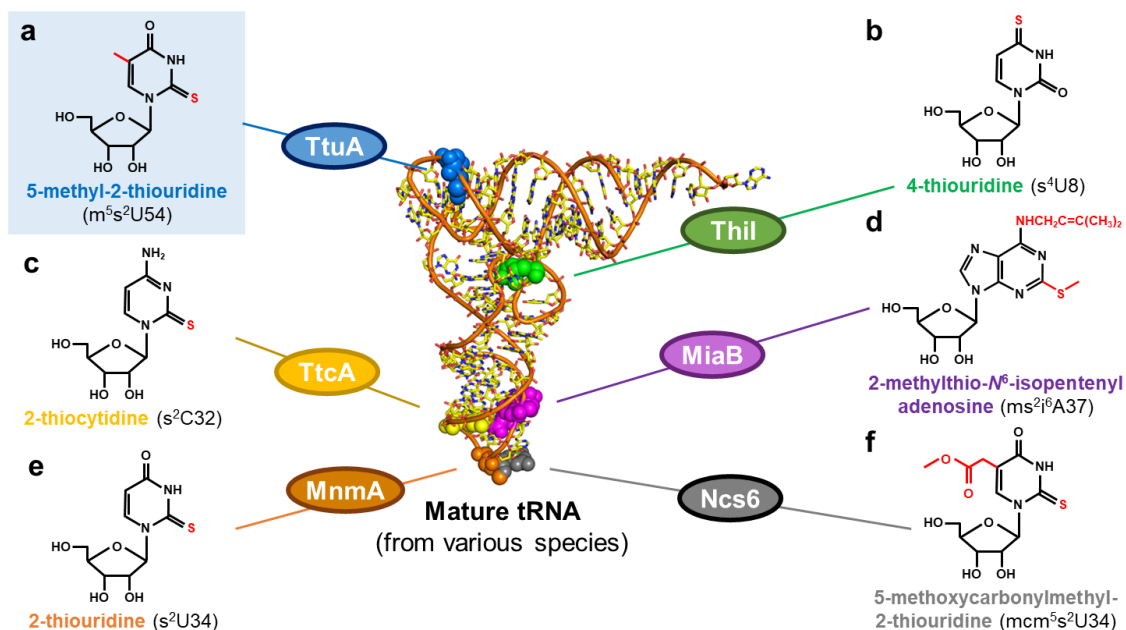


Fig. 1-3 Typical tRNA thiolations at the specific sites of tRNA, and their enzymes. TtuA, ThiI, TtcA, MiaB, MnmA, and Ncs6 catalyze m^5s^2U54 , s^4U8 , s^2C32 , ms^2i^6A37 , s^2U34 , and mcm^5s^2U34 synthesis, respectively. My target, m^5s^2U54 is highlighted in blue.

1-3-1. Thermal stability of tRNA

Thermophilic prokaryotes have 5-methyl-2-thiouridine modifications at position 54 of tRNA (m^5s^2U54 , s^2T54). Thermal adaption using m^5s^2U54 is found in a wide variety of substrate tRNAs^[14], unlike other tRNA thiolations. Since the longer van der Waals radius of sulfur (1.80 Å) than that of oxygen (1.52 Å)^[15], steric repulsion between a 2-thiocarbonyl group and a 2'-hydroxyl group of m^5s^2U54 induces the conformational change of the ribose from the unstable C2'-endo to the stable C3'-endo (Fig. 1-4a)^[16]. This conformational change triggers the formation of base pairing between m^5s^2U54 and 1-methyladenine at position 58 of tRNA (m^1A58), which stacks between base pairs G53-C61 and pseudouridine (Ψ55)-2'-O-methylguanosine (Gm18) (Fig. 1-4b)^[16,17]. The local stabilization around m^5s^2U54 contributes interloop base pairs across T-loop (Ψ55) and D-loop (Gm18), resulting in high thermostability of tRNA. The melting temperature (T_m) of tRNA^{Met} with m^5s^2U54 modification is 89°C^[14], allowing thermophiles to survive in high-temperature environments of more than 80°C^[8]. Note that m^5s^2U54 synthesis is catalyzed by 2-thiouridine synthetase (TtuA)^[18].

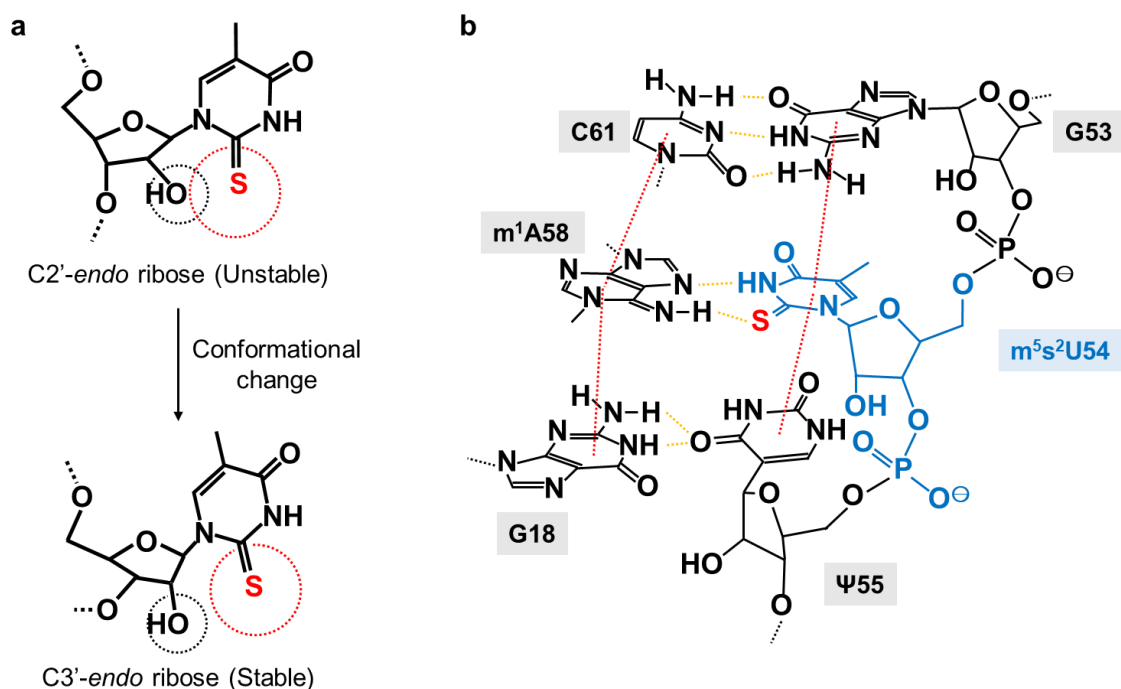


Fig. 1-4 Molecular mechanism of thermal adaption regulated by 5-methyl-2-thiouridine at position 54 of tRNA (m^5s^2U54)^[16].

(a) Conformational change of the ribose in m^5s^2U54 . The red and black dotted lines represent the van der Waals surface of the sulfur and oxygen atoms, respectively. (b) Base pairing across the T and D-loops of thermophilic tRNA. The dashed red and yellow lines represent π - π stacking interactions and hydrogen bonds, respectively. m^5s^2U54 is indicated in blue.

1-3-2. Translational fidelity regulated by thiolation at the wobble position

Accurate translation from genetic information into amino acids is critical for all organisms. To ensure this validation, modified 2-thiouridines (xm^5s^2U34) are universally found at position 34 (the wobble position) of tRNA^{Glu}, tRNA^{Gln}, and tRNA^{Lys} such as 5-methylaminomethyl-2-thiouridine (mm^5s^2U) or 5-carboxymethylaminomethyl-2-thiouridine ($cmnm^5s^2U$) in bacterial tRNA^[19], 5-methoxycarbonylmethyl-2-thiouridine (mcm^5s^2U) in eukaryotic cytosolic tRNA^[2], and 5-taurinomethyl-2-thiouridine (τm^5s^2U) or 5-carboxymethylaminomethyl-2-thiouridine ($cmnm^5s^2U$) in mitochondrial tRNA (Fig. 1-5)^[20]. xm^5U34 alone facilitates base pairing between the oxygen atom at position 2 of the wobble uridine and mRNA guanosine as well as mRNA adenosine (Fig. 1-6a)^[21].

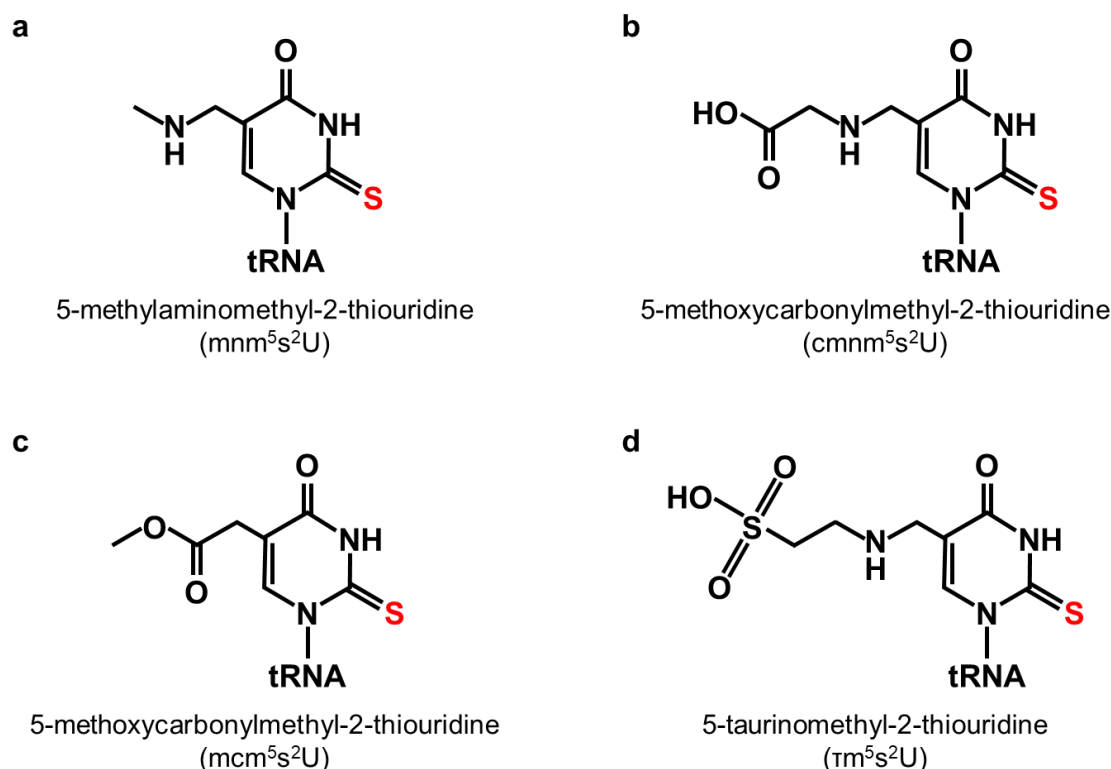


Fig. 1-5 Hypermodified uridines at position 34 of tRNA

(a) 5-methylaminomethyl-2-thiouridine. (b) 5-methoxycarbonylmethyl-2-thiouridine. (c) 5-methoxycarbonylmethyl-2-thiouridine. (d) 5-taurinomethyl-2-thiouridine.

On the other hand, xm⁵s²U34 promotes the formation of C3'-*endo* ribose due to steric effects similar to m⁵s²U54, which stabilizes base pairing with purine bases in mRNA^[21]. Furthermore, the electronegativity of sulfur (2.58) is lower than that of oxygen (3.44)^[22], making hydrogen bonds between sulfur and hydrogen weaker than those between oxygen and hydrogen^[23]. Thus, xm⁵s²U34 interacts more weakly with guanosine (Fig. 1-6b) than with adenosine in the third letter of the codon (Fig. 1-6c)^[21]. As a result, tRNA^{Glu(UUC)}, tRNA^{Gln(UUG)}, and tRNA^{Lys(UUU)} with xm⁵s²U34 modification preferentially translate Glu (GAA), Gln (CCA), and Lys (AAA), respectively.

Notably, it has been reported that the abnormalities in human tRNA thiolations are related to various diseases. For example, defects in the tm⁵s²U34 modification in mitochondrial tRNA^{Lys} (mt-tRNA^{Lys}) are associated with an incurable mitochondrial diseases (MERRF) with muscle weakness and stroke-like symptoms^[24,25]. On the other

hand, excessive mcm⁵s²U34 modification of cytoplasmic tRNAs also promotes the translation of hypoxia-induced factor 1 α (HIF1 α) mRNA and glycolysis in melanoma cells^[2,26]. The mcm⁵s²U34 synthetase complex Ncs6-Ncs2 (also known as Ctu1 and Ctu2 in eukaryotes) are overexpressed in breast cancer, which sustains metastasis^[27,28]. Therefore, understanding the molecular mechanisms of tRNA thiolation is important for biology and medicine.

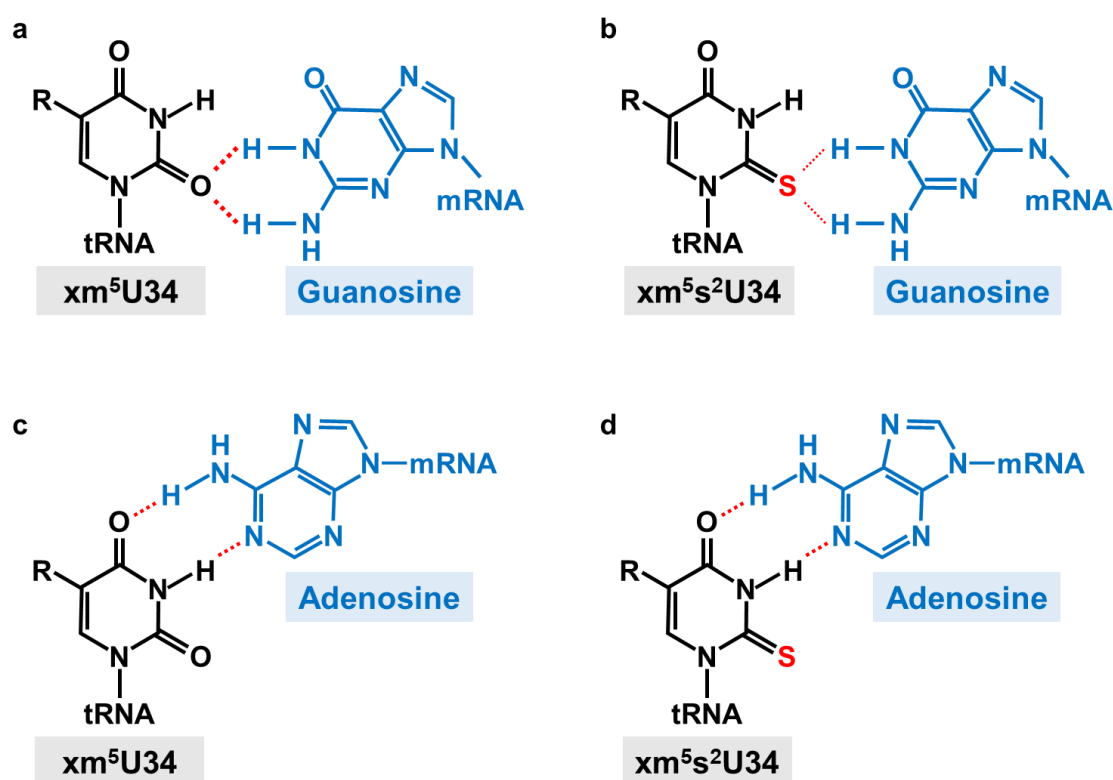


Fig. 1-6 Molecular mechanism of translational fidelity of tRNA with thiolated wobble uridine (xm⁵s²U34).

Wobble base pairing between (a) xm⁵U34 or (b) xm⁵s²U34 of tRNA and mRNA guanosine. The red dotted lines in the non-thiolated state indicate stronger hydrogen bonds than those in the thiolated state. Base pairing between (c) xm⁵U34 or (d) xm⁵s²U34 of tRNA and mRNA adenosine. The binding force does not change regardless of the sulfurization state of tRNA.

1-3-3. Translational fidelity regulated by other thiolation

tRNA thiolation responsible for the accuracy of translation is not only wobble uridines, but also 2-thiocytidine at position 32 of tRNA (s^2C32) and modified 2-methylthioadenosine at position 37 of tRNA (ms^2x^6A37). s^2C32 is found in bacterial and archaeal tRNA^{Arg1(ICG)} with inosine at the wobble position (I34)^[29]. tRNA^{Arg1(ICG)} can decode CGC, CGU, and CGA codons using I34 (Fig. 1-7)^[30]. However, the frequency of CGA codon usage in *E. coli* is only 0.43%^[31], meaning the decoding of adenosine in the third letter of the codon by I34 is inefficient^[32]. Although s^2C32 weakens the hydrogen bond with A38, s^2C32 stabilizes π - π stacking interactions at positions 31-33 of tRNA because sulfur has higher polarizability than oxygen^[33]. Furthermore, s^2C32 destabilizes the structure of the tRNA anticodon, inhibiting the base pairing between I34 at the wobble position and adenosine at position three of the codon^[29]. Thus, tRNA^{Arg1(ICG)} can efficiently translate only CGC and CGU codons. Note that s^2C32 synthesis is catalyzed by 2-thiocytidine synthetase (TtcA)^[33].

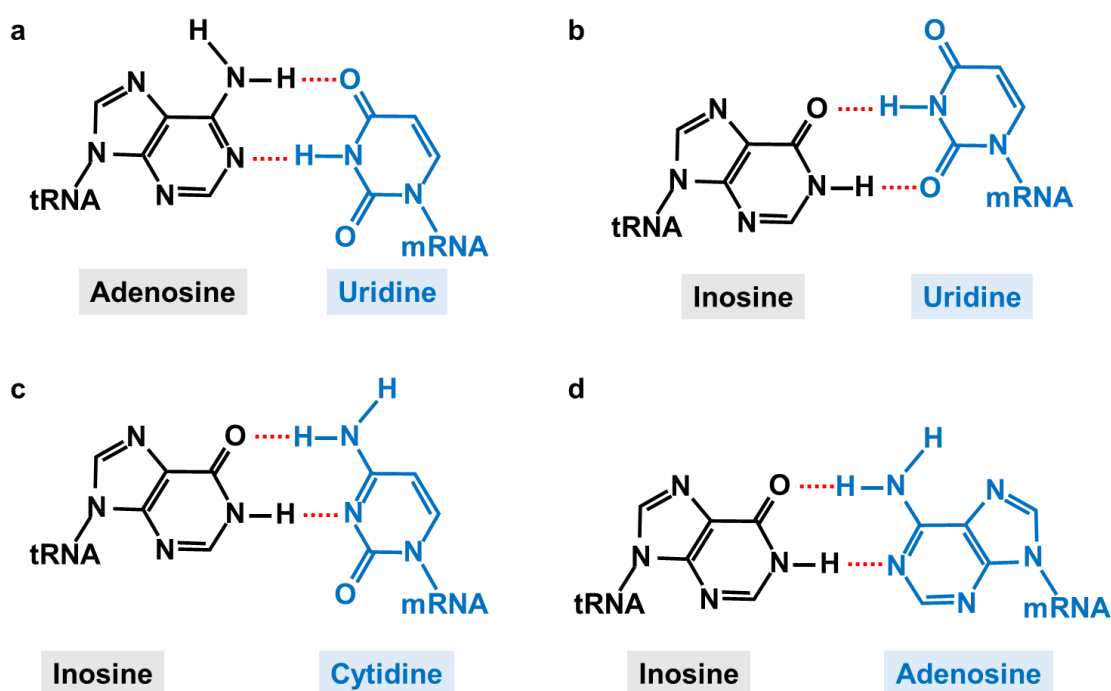


Fig. 1-7 Base pairs expanded by inosine at position 34 of tRNA (I34).

(a) Canonical Watson-Crick base pairing between A34 and mRNA uridine. (b) Base pairing between I34 and mRNA uridine. (c) Major wobble base pairing between I34 and mRNA cytidine. (d) Minor wobble base pairing between I34 and mRNA adenosine.

The ms^2x^6A37 modification is found at tRNA^(NNA) decoding the UNN codon. The sulfur atom of the methylthio group of ms^2x^6A37 stacks with the uridine at position 1 of the codon (Fig. 1-8)^[34]. This stacking strengthens the base pairing between U of mRNA (position 3 of the codon) and A of tRNA (position 1 of the anticodon), preventing slippage of the peptidyl-tRNA in the ribosomal P site^[35]. As a result, ms^2x^6A37 contributes to maintaining the reading frame. Position 6 of the ms^2x^6A37 modification is diverse, with 2-methylthio-*N*⁶-isopentenyladenosine (ms^2i^6A37 , Fig. 1-9a) in tRNA^{Lys(UUU)} from *E. coli*^[36], 2-methylthio-*N*⁶-threonylcarbamoyladenine (ms^2t^6A37 , Fig. 1-9b) in tRNA^{Lys(UUU)} from *Bacillus subtilis*^[37]. Recently, 2-methylthio-cyclic-*N*⁶-threonylcarbamoyladenine (ms^2ct^6A37 , Fig. 1-9c) has also been discovered at position 37 of tRNA^{Lys(UUU)} in *B. subtilis*, plants, and *Trypanosoma brucei*^[38]. In mammals, ms^2t^6A37 is found at position 37 of cytoplasmic tRNA^{Lys(UUU)}^[39], and ms^2i^6A37 at position 37 of mitochondrial tRNA^{Phe}, tRNA^{Tyr}, tRNA^{Trp}, and tRNA^{Ser(UCN)}^[40].

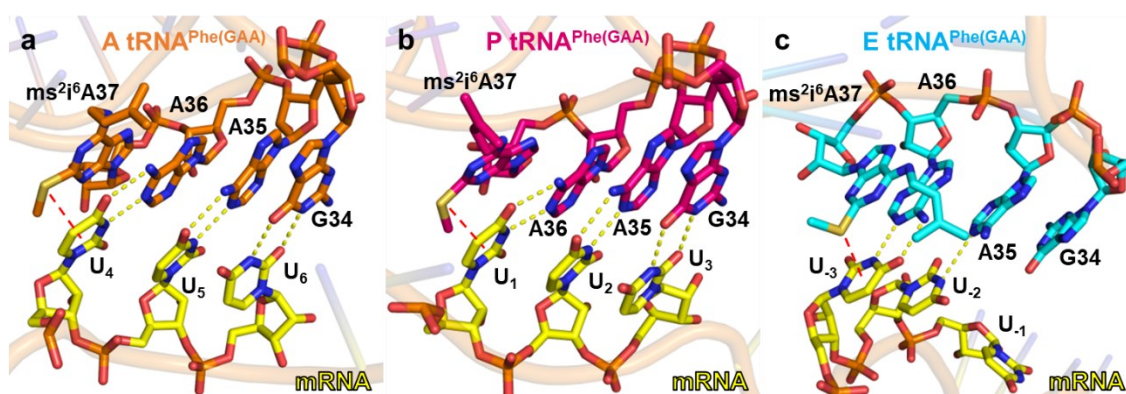


Fig. 1-8 Codon-anticodon interactions stabilized by 2-methylthio-*N*⁶-isopentenyl adenosine at position 37 of tRNA (ms^2i^6A37)^[34].

Base pairing between mRNA codon and tRNA^{Phe} anticodon (a) at the ribosomal A site, (b) P site, and (c) E site (PDB ID: 4V6F). The red and yellow dashed lines indicate stacking and hydrogen bond, respectively.

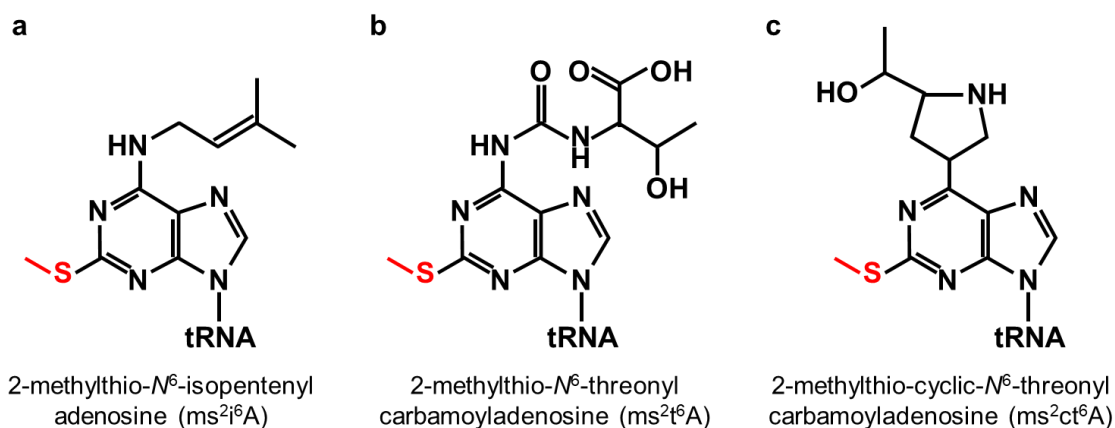


Fig. 1-9 Hypermodified adenosines at position 37 of tRNA

(a) 2-methylthio-*N*⁶-isopentenyladenosine. (b) 2-methylthio-*N*⁶-threonylcarbamoyl adenosine. (c) 2-methylthio-cyclic-*N*⁶-threonylcarbamoyladenosine.

Bacterial *ms*²*i*⁶A37 and *ms*²*t*⁶A37 synthesis is catalyzed by MiaB^[36] and MtaB^[37], respectively. *ms*²*ct*⁶A37 synthesis is catalyzed by TcdA^[38], but the detailed mechanism of biosynthesis is not clear. Mammalian cytoplasmic *ms*²*t*⁶A37 synthesis is catalyzed by MtaB homolog CDKAL1^[37], and mitochondrial *ms*²*i*⁶A37 is synthesized by Cdk5rap1^[41]. Notably, CDKAL1 KO mice showed a deficit of *ms*²*t*⁶A37, a decrease in insulin secretion, and lost blood glucose control^[39]. Thus, *ms*²*t*⁶A37 is considered to be involved in insulin maturation and the development of type II diabetes^[42]. It is also reported that *ms*²*t*⁶A37 is potentially linked to the development of neurodegenerative diseases^[40,43].

1-3-4. Near-UV sensing

*s*⁴U8 (or *s*⁴U9 in some tRNAs) is found in many kinds of prokaryotic tRNA such as tRNA^{Ala}, tRNA^{Met}, and tRNA^{Phe}, which contain C13 (Fig. 1-10)^[44,45]. When *s*⁴U8 is exposed to near UV light (wavelengths: 310 nm-400 nm), it undergoes intramolecular cross-linking with C13 because sulfur is more reactive than oxygen (Fig. 1-11)^[46]. As a result, the cross-linked tRNA changes its conformation and reduces its affinity for aminoacyl-tRNA synthetases, inhibiting aminoacylation, translation^[45], and cellular growth^[47,48]. In addition, *s*⁴U8 stabilizes the structure of tRNA and prevents its degradation by RNase *in vivo*^[49]. Interestingly, biosynthesis of *s*⁴U8 and thiamine (vitamin B1) are catalyzed by the same enzyme, 4-thiouridine synthetase (ThiI)^[50].

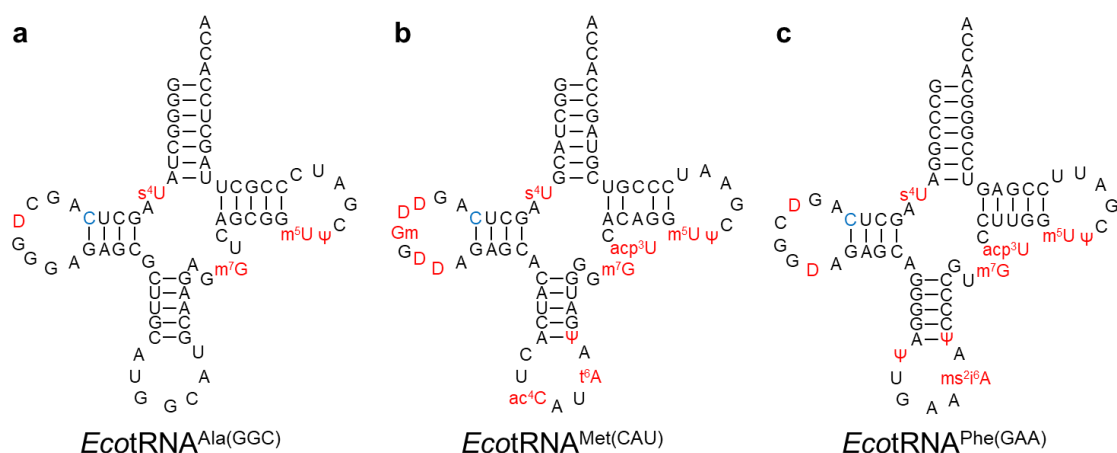


Fig. 1-10 Example of tRNA from *E. coli* which contain s⁴U8 modification and C13.

Secondary structures of (a) tRNA^{Ala(GGC)}[51]. (b) tRNA^{Met(CAU)}[52]. (c) tRNA^{Phe(GAA)}[53].

Modified nucleotides and C13 are colored in red and blue, respectively. The full name of modified nucleotides are as follows: Dihydrouridine (D), 7-methylguanosine (m⁷G), 5-methyluridine (m⁵U), pseudouridine (Ψ), 2'-O-methylguanosine (Gm), N⁴-acetylcytidine (ac⁴C), N⁶-threonylcarbamoyladenine (t⁶A), 3-(3-amino-3-carboxypropyl)uridine (acp³U), and 2-methylthio-N⁶-isopentenyladenine (ms²i⁶A).

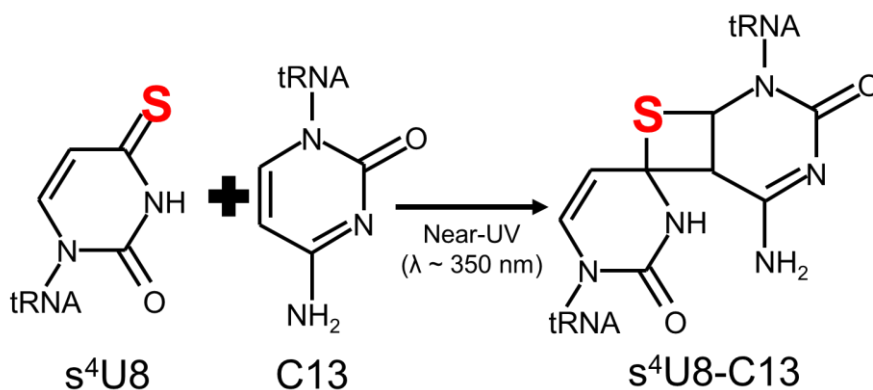


Fig. 1-11 Molecular mechanism of UV sensing regulated by 4-thiouridine at position 8 of tRNA (s⁴U8).

1-4. Iron-sulfur clusters

To date, various thiolated nucleotides are discovered in tRNA, and their biosynthesis is catalyzed by tRNA-thiolation enzymes with an oxygen-sensitive cofactor, iron-sulfur (Fe-S) cluster^[54]. Fe-S clusters are readily decayed in the air^[55], but they are one of the most widely used as cofactors in proteins (Fe-S proteins) from ancient organisms to humans^[56,57]. The fundamental functions of Fe-S clusters are electron transition, protein stabilization, and ligand binding, which are essential for cellular processes such as photosynthesis, respiration, nitrogen fixation, and oxygen sensing^[56,58]. In addition, recent studies show that some Fe-S proteins regulate DNA replication and repair^[59,60], RNA replication and function^[61,62], and their dysfunction is linked to diseases^[63–65].

These broad functions of Fe-S clusters are derived from the wide variety of the type of Fe-S clusters such as [2Fe-2S], [3Fe-4S], [4Fe-4S], and [8Fe-7S] (Fig. 1-12). Notably, [4Fe-4S] clusters can be classified into two groups: ferredoxin-type which is coordinated by four amino acids (typically by cysteine residues, Fig. 1-12c) and aconitase-type which is coordinated by three residues (Fig. 1-12d). Exceptionally, there is a unique [4Fe-4S] coordinated by five cysteine residues (Fig. 1-12f)^[66], but not adopted in tRNA-thiolation enzymes^[67].

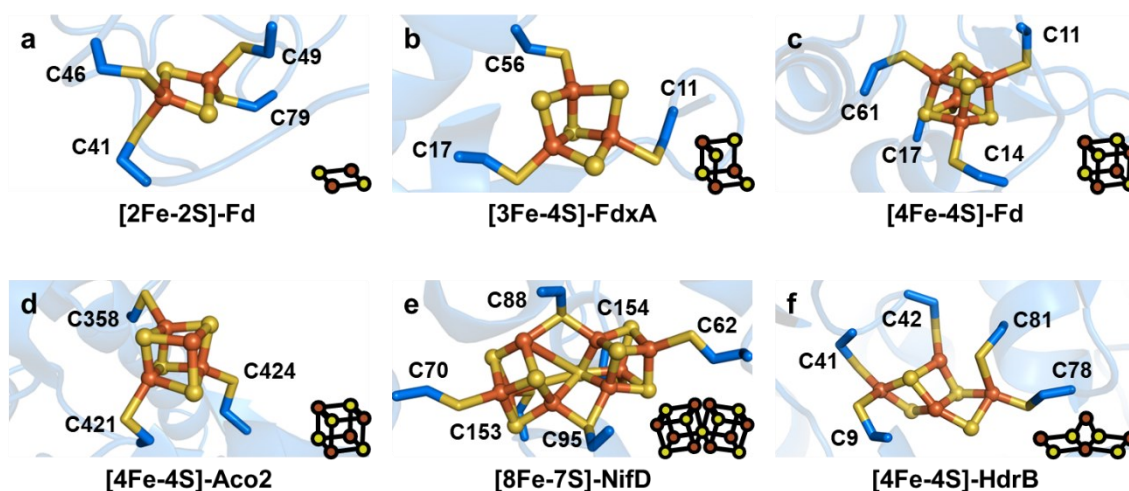


Fig. 1-12 Biological iron-sulfur clusters.

(a) [2Fe-2S] in ferredoxin (PDB ID: 4FXC). (b) [3Fe-4S] in ferredoxin (PDB ID: 1SIZ). (c) [4Fe-4S] in ferredoxin (PDB ID: 1IQZ). (d) [4Fe-4S] in aconitase (PDB ID: 1C96). (e) [8Fe-7S] in nitrogenase (PDB ID: 2MIN). (g) Non-cubane [4Fe-4S] in heterodisulfide reductase (PDB ID: 5ODC). The brown and yellow balls represent Fe and S, respectively.

1-5. tRNA-thiolation enzymes

Studies have identified several tRNA-thiolation enzymes, such as TtuA, TtcA, and Ncs6 of the TtuA/Ncs6 family; MnmA; ThiI; and MiaB. These enzymes are sulfurtransferases that share a PP-loop (SGGXD[S/T]) motif in the pyrophosphatase (PPase) domain and a CXXC...C motif in the binding site of the Fe-S cluster except for the MiaB (methylthiotransferases) (Fig. 1-13)^[67]. As discussed below, I will skip the detailed explanations on MiaB-type enzymes in this paper because their properties are significantly different from those of other tRNA-thiolation enzymes.

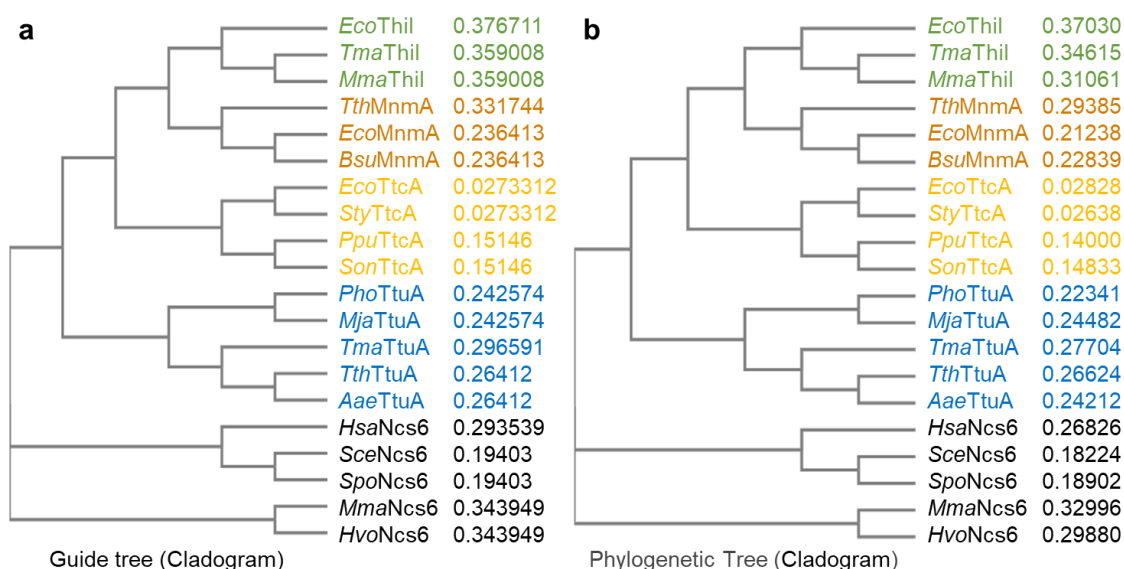


Fig. 1-13 Relevance of tRNA-thiolation enzymes discussed in this paper.

(a) Guide tree. (b) Phylogenetic tree. These Neighbour-joining trees without distance corrections are built by Clustal Omega (<https://www.ebi.ac.uk/Tools/msa/clustalo/>)^[68]. ThiI from *Escherichia coli* K12 (*EcoThiI*), *Thermotoga maritima* MSB8 (*TmaThiI*), and *Methanococcus maripaludis* S2 (*MmaThiI*)^[69,67]. MnmA from *Thermus thermophilus* HB27 (*TthMnmA*), *E. coli* K12 (*EcoMnmA*), and *Bacillus subtilis* 168 (*BsuMnmA*)^[70]. TtcA from *E. coli* K12 (*EcoTtcA*), *Salmonella typhimurium* LT2 (*StyTtcA*), *Pseudomonas putida* W619 (*PpuTtcA*), and *Shewanella oneidensis* MR-1 (*SonTtcA*)^[71]. TtuA from *Pyrococcus horikoshii* OT-3 (*PhoTtuA*), *Methanocaldococcus jannaschii* ATCC 43067 (*MjaTtuA*), *T. maritima* MSB8 (*TmaTtuA*), *T. thermophilus* HB27 (*TthTtuA*), and *Aquifex aeolicus* VF5 (*AaeTtuA*)^[18,72]. Ncs6 from *Homo sapiens* (*HsaNcs6*), *Saccharomyces cerevisiae* ATCC 204508 (*SceNcs6*), *Schizosaccharomyces pombe* 972 strain (*SpoNcs6*), *M. maripaludis* S2 (*MmaNcs6*), and *Haloferax volcanii* ATCC 29605 (*HvoNcs6*)^[67,73].

From the early 2000s until around 2014, tRNA-thiolation enzymes were roughly classified into two groups: Fe-S-dependent enzymes (for s^2C32 and ms^2x^6A synthesis) and Fe-S-independent enzymes (for s^4U8 and x^5s^2U synthesis) (Fig. 1-14a)^[19,74,75]. Then, it was reported that s^2C32 synthetase TtcA depends on a [4Fe-4S] cluster^[71], which triggered the discoveries of the presence of [3Fe-4S] cluster in archaeal s^4U8 synthetases ThiI (also known as TtuI) and mcm s^5s^2U synthase Ncs6 (also known as NcsA) and the eukaryotic mcm s^5s^2U synthetase Ncs6 (also known as Ctu1)^[69]. Furthermore, structural and functional analyses of the m s^5s^2U54 synthetase TtuA showed that TtuA has a [4Fe-4S] cluster^[72,76]. Around 2018, these results enabled us to suggest a new classification that tRNA-thiolation enzymes can be classified into three types: the MnmA/ThiI type which is independent of Fe-S clusters; the TtuA/Ncs6/TtcA type which is dependent on Fe-S clusters and adenosine triphosphate (ATP); and the MiaB type which is dependent on Fe-S clusters and *S*-adenosyl methionine (SAM) (Fig. 1-14b)^[6].

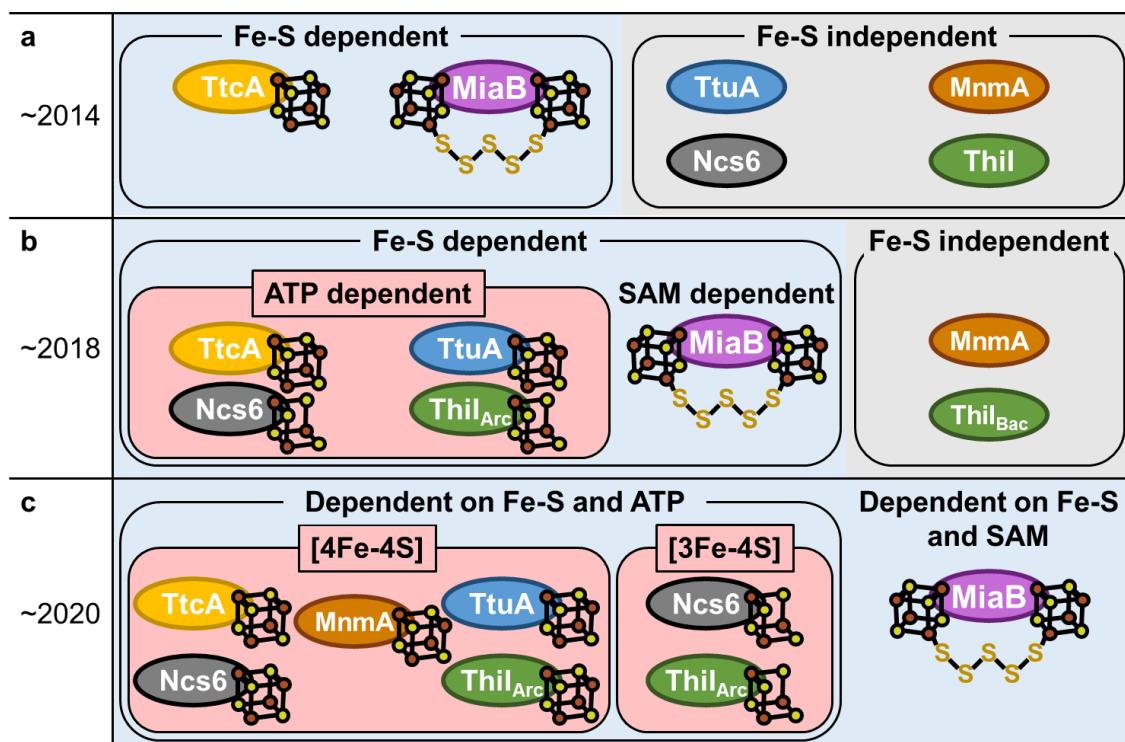


Fig. 1-14 Changes in the classification of typical tRNA-thiolation enzymes.

(a) Two major groups depending on the Fe-S clusters until 2014. (b) Three major groups depending on the Fe-S clusters and other cofactors. (c) Three major groups depending on the type of Fe-S clusters. Note that bacterial ThiI (ThiI_{Bac}) is independent of Fe-S clusters whereas archaeal ThiI (ThiI_{Arc}) is dependent on Fe-S clusters.

In the 2020s, spectroscopic and biochemical experiments revealed that the enzymatic active form of MnmA is [4Fe-4S] cluster^[70,77]. Furthermore, [4Fe-4S] cluster is required for the enzymatic activity of bacterial ThiI under strict anaerobic conditions^[67]. Interestingly, some research groups proposed that Ncs6 and ThiI have [4Fe-4S], but others proposed [3Fe-4S] despite the sequence similarity of these enzymes (Table 1-1). It is now proposed that tRNA-thiolation enzymes are generally dependent on Fe-S clusters and can be classified into two types: TtuA/Ncs6 and MiaB (Fig. 1-14c)^[54,67].

	Tth	Aae	Tma	Pho	Mja	Eco	Ppu	Son	Sty	Has	Sce	Spo	Mma	Hvo	Eco	Bsu	Tth	Eco	Tma	Mma
	TtuA	TtuA	TtuA	TtuA	TtuA	TtcA	TtcA	TtcA	TtcA	Ncs6	Ncs6	Ncs6	Ncs6	Ncs6	MnmA	MnmA	MnmA	Thil	Thil	Thil
Tth		45.5	37.0	39.6	33.5	17.4	18.5	17.4	16.6	27.1	23.6	22.6	23.4	23.8	13.2	11.3	15.6	13.7	13.9	12.8
TtuA		(74.0)	(71.6)	(72.9)	(67.7)	(52.6)	(48.0)	(50.0)	(52.6)	(52.9)	(56.1)	(59.8)	(57.8)	(54.7)	(44.6)	(40.2)	(40.9)	(37.9)	(46.2)	(47.3)
Aae			47.3	44.3	39.1	19.2	20.4	17.2	18.3	22.5	23.4	22.6	26.9	29.0	12.1	11.3	13.1	13.1	14.8	12.8
TtuA			(76.1)	(75.5)	(71.7)	(52.9)	(54.9)	(50.6)	(52.3)	(49.3)	(51.3)	(54.8)	(60.9)	(63.3)	(41.2)	(42.1)	(39.1)	(37.8)	(45.7)	(44.7)
Tma				41.1	41.3	18.0	17.7	16.8	18.1	20.7	23.4	22.7	26.0	30.3	10.5	10.2	11.8	11.0	14.3	14.8
TtuA				(76.1)	(74.8)	(50.2)	(51.5)	(51.1)	(51.5)	(50.3)	(53.8)	(59.1)	(60.2)	(59.6)	(30.9)	(39.1)	(40.6)	(36.5)	(42.5)	(46.2)
Pho					50.6	17.5	18.0	16.7	18.3	25.1	27.3	25.7	30.2	27.5	12.4	13.2	12.5	13.0	13.4	12.8
TtuA					(80.6)	(51.4)	(49.8)	(51.1)	(51.9)	(52.3)	(55.2)	(58.8)	(62.2)	(59.7)	(41.2)	(44.0)	(40.9)	(38.5)	(43.8)	(43.6)
Mja						18.9	19.9	19.2	18.4	21.7	23.4	24.1	27.1	26.7	12.8	12.3	11.7	11.8	12.6	15.6
TtuA						(50.9)	(51.3)	(51.4)	(50.5)	(50.6)	(55.4)	(60.7)	(62.5)	(61.5)	(42.9)	(43.2)	(42.0)	(39)	(45.9)	(45.1)
Eco						59.5	60.8	94.5	17.7	16.9	17.7	19.8	20.4	10.8	13.6	10.5	11.0	13.6	12.5	
TtcA						(78.5)	(85.7)	(99.7)	(47.6)	(54.7)	(52.7)	(53.0)	(51.4)	(41.5)	(44.1)	(38.1)	(38.0)	(44.2)	(47.4)	
Ppu							63.8	59.2	19.9	17.2	17.8	21.1	21.6	12.0	14.0	14.5	10.6	12.5	12.7	
TtcA							(80.7)	(78.8)	(49.0)	(49.2)	(49.9)	(53.8)	(54.6)	(39.7)	(39.4)	(41.1)	(36.7)	(42.5)	(40.9)	
Son							61.5	19.1	18.7	18.2	18.5	20.9	11.7	12.7	14.0	13.2	12.3	12.5		
TtcA							(85.0)	(49.0)	(52.4)	(52.4)	(50.5)	(51.6)	(42.6)	(42.6)	(44.7)	(37.8)	(40.8)	(45.7)		
Sty									18.8	18.9	17.7	20.1	20.4	12.7	13.6	10.8	10.4	15.9	12.5	
TtcA									(48.7)	(53.9)	(51.5)	(52.1)	(49.0)	(45.4)	(43.3)	(38.8)	(38.6)	(44.3)	(44.8)	
Has									43.6	42.9	26.9	29.5	11.8	13.3	15.0	11.0	11.4	11.6		
Ncs6									(72.2)	(74.2)	(58.2)	(56.8)	(44.0)	(44.7)	(41.0)	(38.4)	(43.2)	(39.3)		
Sce											58.3	25.6	29.3	11.6	12.6	12.8	10.6	12.9	11.3	
Ncs6											(80.0)	(59.2)	(57.8)	(44.2)	(42.4)	(43.0)	(50.1)	(47.7)		
Spo												28.0	28.3	10.2	8.3	10.3	13.0	12.2	13.0	
Ncs6												(62.8)	(59.1)	(44.7)	(44.8)	(44.1)	(38.9)	(50.1)	(44.4)	
Mma													33.7	11.7	10.7	12.7	9.1	12.1	14.3	
Ncs6													(68.1)	(39.6)	(42.9)	(44.1)	(38.6)	(45.5)	(43.9)	
Hvo														13.3	13.5	12.8	11.6	11.0	12.0	
Ncs6														(46.2)	(44.3)	(42.7)	(41.0)	(44.4)	(42.1)	
Eco															54.3	37.7	12.0	12.4	10.2	
MnmA															(79.5)	(65.4)	(41.7)	(44.7)	(43.0)	
Bsu																36.6	12.9	12.3	11.3	
MnmA																(66.5)	(41.5)	(46.9)	(45.5)	
Tth																	11.6	12.2	9.3	
MnmA																	(42.5)	(42.5)	(43.8)	
Eco																	21.8	23.1		
Thil																	(52.1)	(51.2)		
Tma																			31.9	
Thil																			(66.1)	

Table 1-1 Sequence identity of tRNA-thiolation enzymes. The upper value indicates identity (%) and the lower value indicates similarity (%) of the amino acid sequence evaluated using ClustalW available on the NPS@ server (https://npsa-prabi.ibcp.fr/cgi-bin/npsa_automat.pl?page=/NPSA/npsa_clustalw.html)^[78]. Identity values $\geq 40\%$; $\geq 30\%$ but $< 40\%$; and $\geq 20\%$ but $< 30\%$ are highlighted in red, yellow, and green, respectively.

1-5-1. 2-thiouridine synthetase TtuA

TtuA catalyzes m^5s^2U54 biosynthesis with ATP and sulfur donor protein TtuB (Fig. 1-15), and these proteins were identified in the thermophilic bacterium *T. thermophilus* in 2006^[18]. TtuB is a ubiquitin-like protein with a conserved GG motif at the thiocarboxyl C-terminus (R-COSH), like other sulfur donors (Fig. 1-16)^[79]. The substrate tRNA of TtuA is modified with 5-methyluridine (m^5U54) by the methyltransferase TrmFO^[80], and it is suggested that m^1A58 modification also promotes m^5s^2U54 formation^[81].

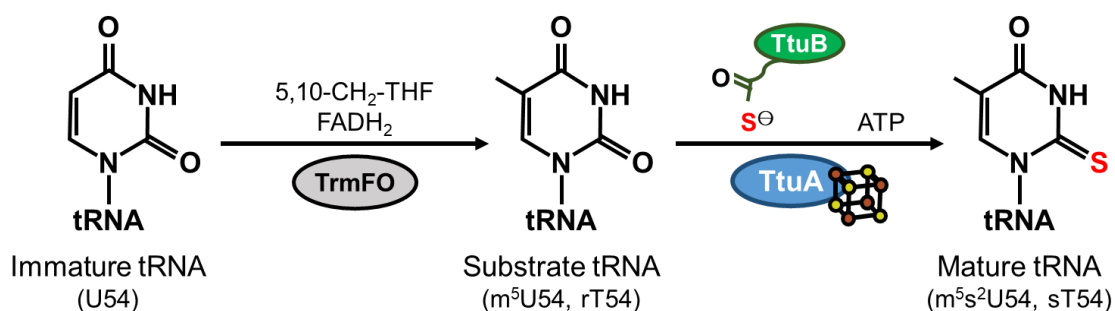


Fig. 1-15 Scheme of 5-methyl-2-thiouridine biosynthesis.

TrmFO catalyzes m^5U54 biosynthesis with N^5,N^{10} -methylenetetrahydroforate (CH₂THF) and reduced flavin adenine dinucleotide (FADH₂). TtuA transfers sulfur from the C-terminus of TtuB to substrate tRNA with ATP. Note that some organisms without *ttub* gene catalyze m^5s^2U54 biosynthesis using inorganic sulfur from the environment.

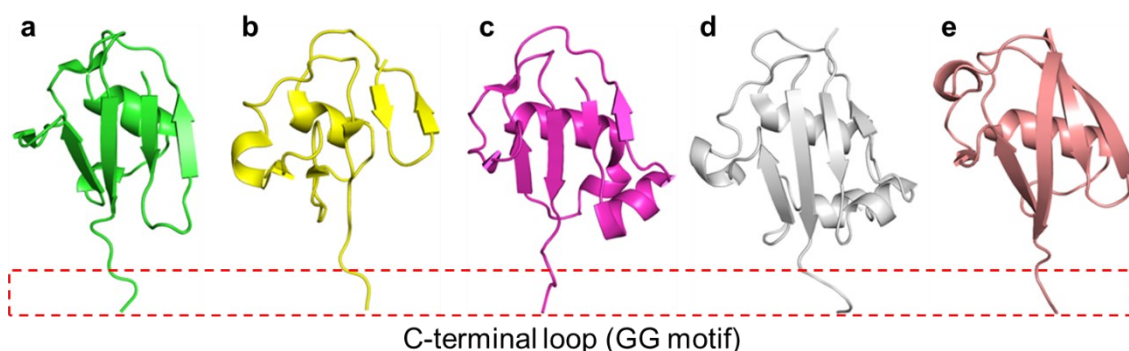


Fig. 1-16 Ubiquitin-like sulfur donor proteins.

(a) TtuB is a sulfur donor of m^5s^2U54 (PDB ID: 5ZTB). (b) ThiS is a sulfur donor of thiamine (PDB ID: 1TYG). (c) MoaD is a sulfur donor of molybdopterin (PDB ID: 2QIE). (d) Urm1 is a sulfur donor of mcm^5s^2U34 (PDB ID: 2AX5). (e) Ubiquitin is not a sulfur donor, but responsible for the degradation of proteins in eukaryotes (PDB ID: 1UBQ).

The crystal structure of apo-TtuA from *Pyrococcus horikoshii* (*Pho*TtuA) was determined and the critical residues were also identified in 2013^[82], then it was revealed that *Pho*TtuA and TtuA from *T. thermophilus* (*Tth*TtuA) contain a [4Fe-4S] cluster in their catalytic site in 2017^[72,76]. Interestingly, whereas three of the four Fe atoms in the [4Fe-4S] were coordinated to the CXXC...C motif, the remaining one bare Fe (the unique Fe) was exposed to the solution^[72]. Furthermore, we have recently determined the crystal structure of the [4Fe-4S]-TtuA-TtuB-ATP complex, showing that the unique Fe binds to the C-terminus of TtuB (Fig. 1-17)^[83]. Based on these structures, we proposed a novel reaction mechanism of tRNA thiolation in which the unique Fe receives sulfur from TtuB, then transfers sulfur to the adenylated tRNA (AdetRNA)^[83]. However, the reaction mechanism of tRNA activation potentially involved in [4Fe-4S] has not been elucidated because the TtuA-tRNA complex structure is unknown.

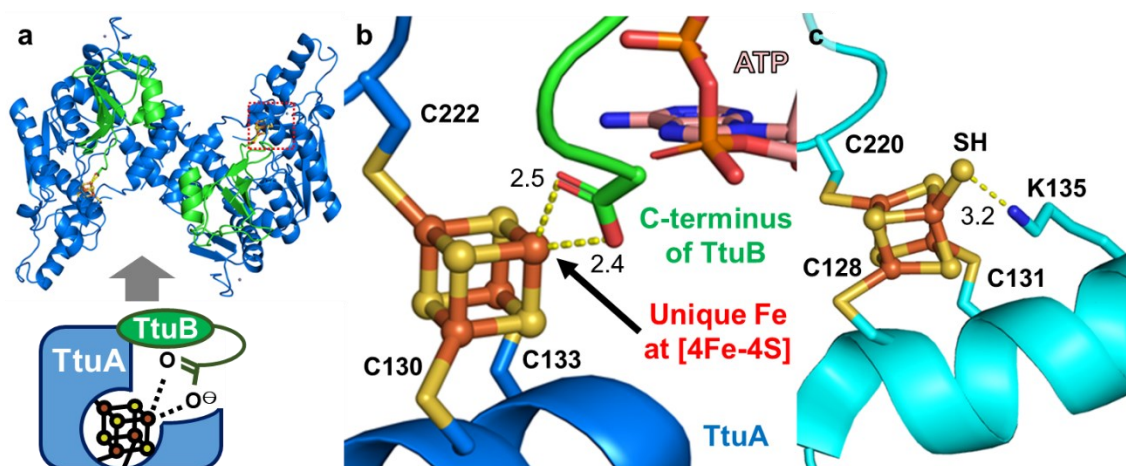


Fig. 1-17 Crystal structure of the [4Fe-4S]-TtuA complex.

(a) Overall structure of the [4Fe-4S]-*Tth*TtuA-*Tth*TtuB-ATP complex dimer (PDB ID: 5ZTB). TtuA and TtuB are colored blue and green, respectively as indicated in the illustration. The catalytic site is marked by the red dotted frame. (b) Close-up view of the catalytic center of *Tth*TtuA. The brown and yellow balls represent Fe and S, respectively. [4Fe-4S] is coordinated by Cys130, Cys133, and Cys222. The yellow dotted lines indicate coordination bonds between the unique Fe and O atoms at the C-terminus of TtuB with distance in angstrom units. (c) Close-up view of the catalytic site of *Pho*TtuA (PDB ID: 5MKP). [4Fe-4S] is coordinated by Cys128, Cys131, and Cys220. The unique Fe captures inorganic sulfur and forms the [4Fe-5S]-TtuA intermediate stabilized by Lys135.

1-5-2. 2-thiouridine synthetase Ncs6

Ncs6 catalyzes mcm⁵s²U34 biosynthesis in eukaryotic tRNA^{Glu}, tRNA^{Gln}, and tRNA^{Lys} with ATP, Ncs2, and sulfur donor protein Urm1 (Fig. 1-18), and Ncs6/Ncs2 and Urm1 were identified in *S. cerevisiae* in 2003 and 2000, respectively^[84,85]. Urm1 is a ubiquitin-like sulfur donor similar to TtuB (Fig. 1-16d)^[86]. The substrate tRNA of Ncs6 has mcm⁵U34 modification involving more than 10 genes including the methyltransferase Trm9^[87]. The previous reports that Ncs6 recruits Ncs2, indicating the formation of the Ncs6-Ncs2 complex, but the detailed role of Ncs2 is still unclear^[73,87].

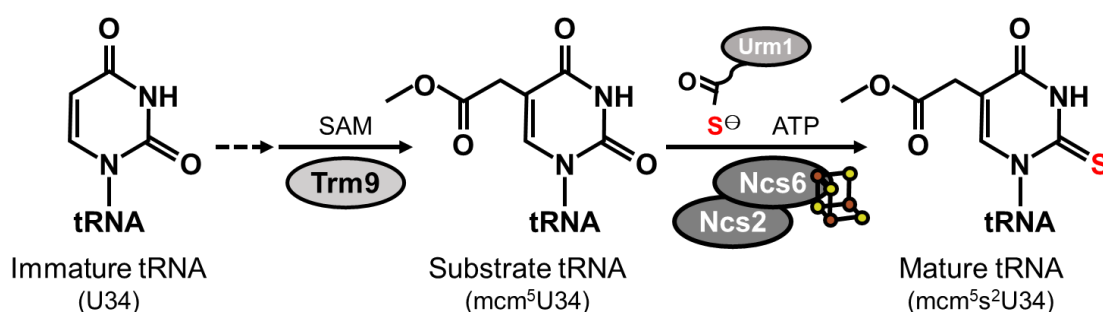


Fig. 1-18 Scheme of 5-methoxycarbonylmethyl-2-thiouridine biosynthesis.

Trm9 catalyzes mcm⁵U34 synthesis using a co-factor SAM after multiple modifications of U34. Ncs6 transfers sulfur from the C-terminus of Urm1 to substrate tRNA with ATP and Ncs2. The type of Fe-S cluster is under discussion, but is shown as [3Fe-4S] in this figure. Note that some organisms without *urm1* gene catalyze mcm⁵s²U34 biosynthesis using inorganic sulfur from the environment.

In 2016, electron paramagnetic resonance (EPR) and Mössbauer spectroscopic research showed that Ncs6 from *M. maripaludis* (*Mma*Ncs6) and from *S. cerevisiae* (*Sce*Ncs6) contains [3Fe-4S] cluster, respectively (Fig. 1-19a, b)^[69]. On the other hand, more recently, the crystal structure of [4Fe-4S]-*Mma*Ncs6 has been determined under strictly anaerobic conditions. This structure showed that the CXXC...C motif of Ncs6 coordinates the [4Fe-4S] and the presence of the unique Fe similar to TtuA (Fig. 1-19c, d)^[67]. The inconsistent identification of the Fe-S cluster in Ncs6 comes from the oxidative sensitivity of [4Fe-4S] similar to [4Fe-4S] in aconitase (Aco2)^[88], pyruvate formate-lyase activating enzyme (PFL-AE)^[89], and isopentenyl-diphosphate (IspH)^[90] (Fig. 1-20).

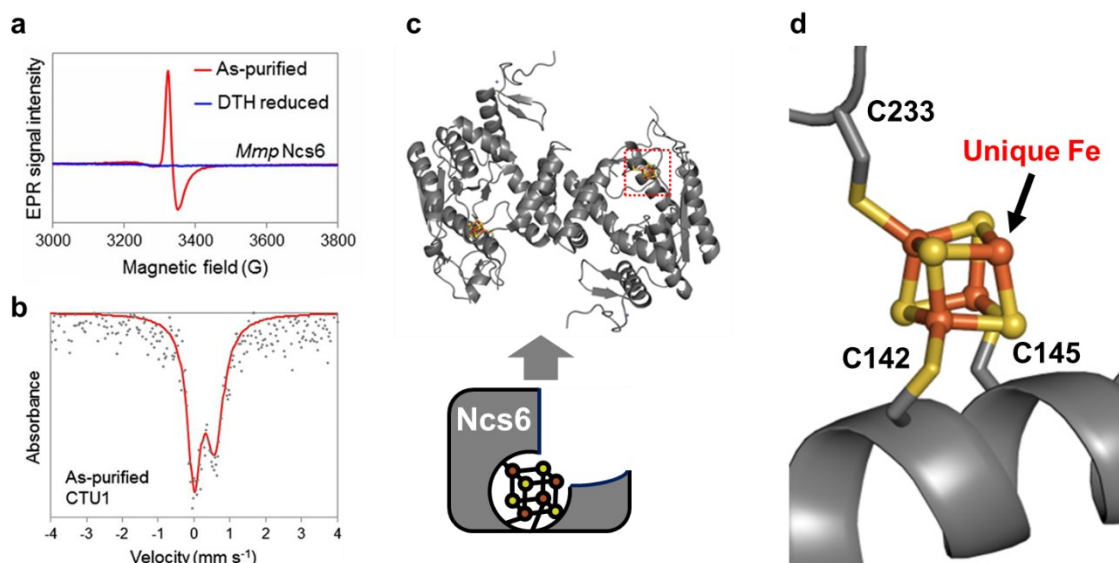


Fig. 1-19 Experimental evidences for the type of Fe-S clusters in Ncs6.

(a) X-band EPR spectra of [3Fe-4S]-*MmaNcs6* in the as-purified (red) and sodium dithionite (DTH)-reduced (blue) states^[69]. (b) The zero-field ⁵⁷Fe Mössbauer spectrum of the as-purified *SceNcs6*^[69]. (c) Overall structure of the [4Fe-4S]-*MmaNcs6* complex dimer (PDB ID: 6SCY)^[67]. The catalytic site is marked by the red dotted frame. (d) Close-up view of the catalytic center of *MmaNcs6*. The brown and yellow balls represent Fe and S, respectively. Three of the four Fe atoms in [4Fe-4S] are coordinated by Cys142, Cys145, and Cys233.

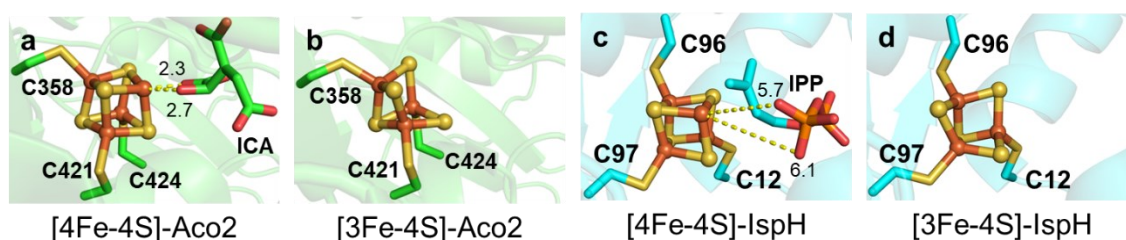


Fig. 1-20 Misidentification of Fe-S clusters.

(a) The unique Fe of [4Fe-4S]-Aco2 (PDB ID: 1B0J) coordinates with isocitric acid (ICA). (b) Inactive [3Fe-4S]-Aco2 (PDB ID: 5ACN) produced by oxidation of [4Fe-4S]-Aco2. (c) The unique Fe of [4Fe-4S]-IspH (PDB ID: 3KE9) coordinates with isopentenyl diphosphate (IPP). (d) Inactive [3Fe-4S]-IspH (PDB ID: 3F7T) produced by oxidation of [4Fe-4S]-IspH. The dashed lines indicate distances between unique Fe atoms and each ligand with distance in angstrom units.

1-5-3. 2-thiocytidine synthetase TtcA

TtcA catalyzes s^2C32 biosynthesis in bacterial tRNA^{Arg1(ICG)} with ATP and inorganic sulfur (Fig. 1-21), which was identified in *E. coli* (*EcoTtcA*) and *Salmonella typhimurium* (*StyTtcA*) in 2004^[33]. The sulfur source for s^2C32 is free L-cysteine, which is desulfurized by desulfurase (IscS)^[91]. In 2014, spectroscopic and biochemical research showed that [4Fe-4S] is essential for the enzymatic activity of *EcoTtcA*^[71]. TtcA contains the PPase domain and two CXXC motifs, and three of the four cysteines (Cys122, Cys125, and Cys213) are essential for the catalytic activity of *EcoTtcA*, whereas the C210A mutant showed 50% of enzymatic activity^[71]. These biochemical results indicated that [4Fe-4S]-TtcA contains the unique Fe.

In addition to the biochemical characterization, physiological insight on TtcA has recently been provided, which showed that TtcA from *Pseudomonas aeruginosa* (*PaeTtcA*) protects against oxidative stress via catalase activity^[92]. Although the crystal structure of TtcA has not been determined, 3D structural models of *EcoTtcA* have been reported using AlphaFold and RoseTTAFold^[93]. The predicted structures also support the presence of the unique Fe, but the possibility that C210 could be a ligand of [4Fe-4S] still remains. Therefore, the structural determination of TtcA by X-ray crystallography is highly desired.

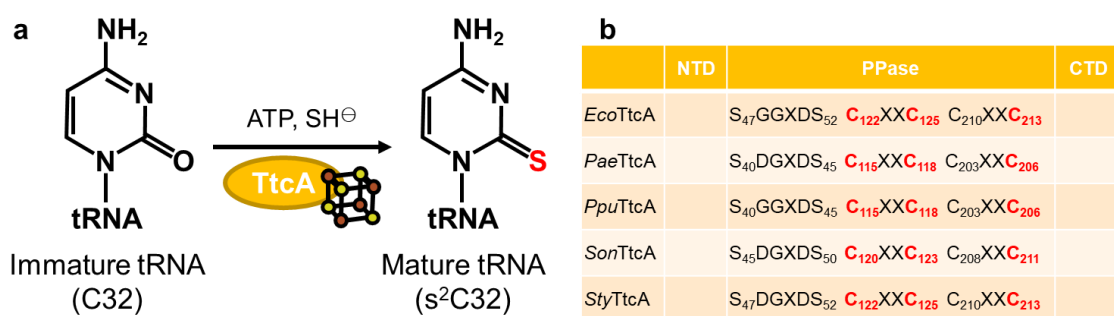


Fig. 1-21 Scheme of 2-thiocytidine biosynthesis and characterization of TtcA.

(a) TtcA catalyzes s^2C32 synthesis using ATP and inorganic sulfur source. TtcA transfers sulfur provided by IscS system. (b) Domain construct and the location of critical residues in typical TtcA. TtcA contains the N-terminal domain (NTD), catalytic PPase domain, and C-terminal domain (CTD). The cysteine residues binding to [4Fe-4S] are indicated in red based on the previous study on *EcoTtcA*^[71]. The abbreviations of the protein source are the same as in Fig. 1-13.

1-5-4. 2-thiouridine synthetase MnmA

MnmA (formerly AsuE or TrmU) catalyzes s²U34 biosynthesis in bacterial tRNA^{Glu}, tRNA^{Gln}, and tRNA^{Lys} with ATP and an inorganic sulfur (Fig. 1-22). MnmA was identified in *E. coli* (*EcoMnmA*) in 2003^[94]. In 2006, the structures of the *EcoMnmA*-tRNA^{Glu} complex have been determined (Fig. 1-23), showing that the flexible loop of MnmA prevents the inactivation of the AdetRNA by the influx of solvent^[95]. Based on this structure, the two possible catalytic mechanisms were proposed that MnmA catalyzes s²U34 synthesis via persulfide (R-SSH).

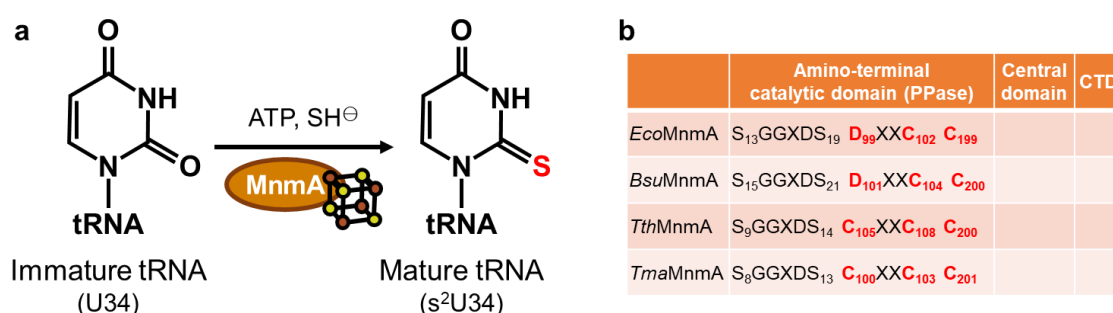


Fig. 1-22 Scheme of 2-thiouridine biosynthesis and characterization of MnmA.

(a) MnmA catalyzes s²U34 synthesis using ATP and inorganic sulfur source. (b) Domain construct and the location of critical residues in typical MnmA. MnmA contains the amino-terminal catalytic domain (PPase), central domain, and carboxy-terminal β-barrel domain (CTD)^[95]. The [4Fe-4S]-binding residues are indicated in red.

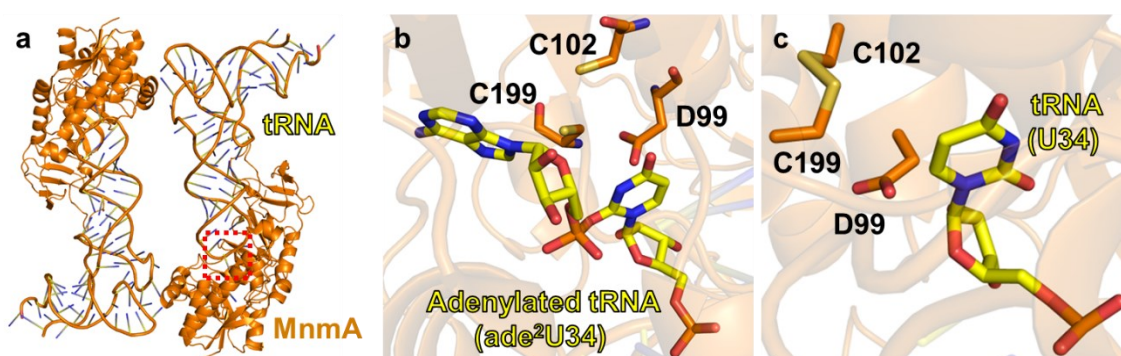


Fig. 1-23 Crystal structure of the *EcoMnmA*-tRNA^{Glu} complexes^[95].

(a) Overall structure of the MnmA-adenylated tRNA complex (PDB ID: 2DEU). The catalytic site is marked by the red dotted frame. Close-up view of the catalytic center of MnmA (b) in the adenylated intermediate state and (c) in the initial tRNA binding state (PDB ID: 2DER). Cys102 and Cys199 can form a disulfide bond.

Although the structural understanding of s^2U34 biosynthesis was advanced, the enzymatic activity of apo-MnmA was very weak^[94–97]. Recently, spectroscopic and biochemical research showed that MnmA from *T. thermophilus* (*Tth*MnmA) contains [4Fe-4S] cluster coordinated by Cys105, Cys108, and C200^[70]. Furthermore, it was reported that the enzymatically active form of *Eco*MnmA contains a [4Fe-4S] coordinated by Asp99, Cys102, and Cys199^[77]. Although the experimental structure of the [4Fe-4S]-MnmA complex has not been determined, the [4Fe-4S]-MnmA model has been reported^[93]. These recent studies updated the sulfur relay system. Previously, the sulfur source for s^2U34 is considered to be free L-cysteine transferred via the sulfur relay system involving persulfide proteins IscS^[98,99] and TusABCDE in typical bacteria (Fig. 1-24a)^[96,100] or YrvO in *B. subtilis* (Fig. 1-24b)^[97]. On the other hand, owing to the structural insights of [4Fe-5S]-*Pho*TtuA and [4Fe-5S]-TudS^[76,101], we can assume a new sulfur relay system involving [4Fe-5S]-MnmA (Fig. 1-24c).

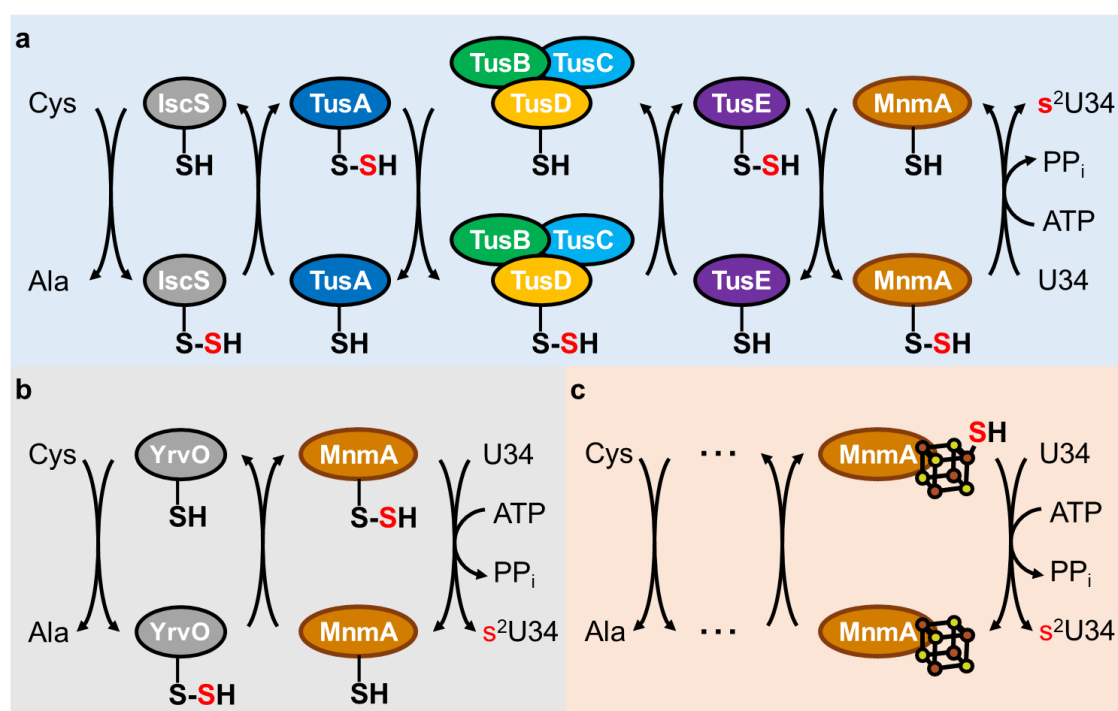


Fig. 1-24 Sulfur relay system for the biosynthesis of s^2U34 catalyzed by MnmA.

Previously proposed pathway mediated via (a) the IscS-TusABCDE system in typical bacteria and (b) YrvO in some bacteria. (c) New putative pathway mediated via [4Fe-5S]-MnmA. The upstream of MnmA involves persulfide proteins that differ among species, as previous studies have shown.

1-5-5. 4-thiouridine synthetase ThiI

ThiI catalyzes the s^4 U8 biosynthesis in prokaryotic tRNA^{Trp}, tRNA^{Phe}, tRNA^{Arg}, tRNA^{Val}, tRNA^{Asp}, tRNA^{Gly}, tRNA^{Met}, and tRNA^{His} with ATP and inorganic sulfur (Fig. 1-25a). ThiI was identified in *E. coli* (*EcoThiI*) in 1998^[13], and it was proposed the catalytic mechanism of s^4 U8 biosynthesis involves the disulfide bond between Cys344 at the PP-loop domain and Cys456 at the rhodanese-like domain (RLD) (Fig. 1-26)^[98]. However, the structures of ThiI from *B. anthracis* (*BanThiI*) with the THUMP domain responsible for tRNA-binding showed that RLD is absent in some organisms (Fig. 1-25b)^[102]. These bacteria also have specific cysteine desulfurase NifZ, which enables s^4 U8 synthesis without RLD in ThiI^[103]. In most archaea in the absence of NifZ, their ThiI contains catalytic cysteines in the PP-loop domain instead of RLD^[104].

Furthermore, the crystal structure of the ThiI from *Thermotoga maritima* (*TmaThiI*) complex with truncated tRNA^{Phe} has been determined (Fig. 1-27), which revealed the interaction manner between tRNA and the THUMP domain of *TmaThiI*, and demonstrated that dynamic structural change of the substrate tRNA with ACCA 3'-end is required for s^4 U8 formation^[105]. However, the catalytic mechanism of ThiI is still unclear because the substrate base U8 did not enter the active site.

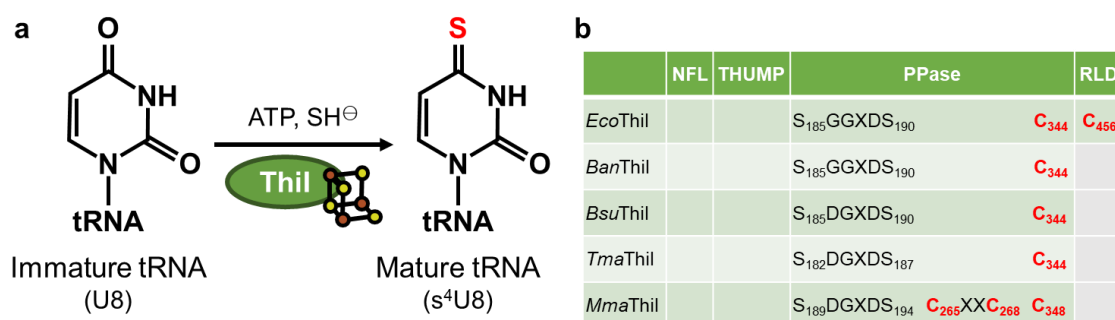


Fig. 1-25 Scheme of 4-thiouridine biosynthesis and characterization of ThiI.

(a) ThiI catalyzes s^4 U8 synthesis using ATP and inorganic sulfur source. The type of Fe-S cluster is still under discussion, but is shown as [3Fe-4S] in this figure. (b) Domain construct and the location of critical residues in typical ThiI. ThiI contains an N-terminal ferredoxin-like (NFL) domain, RNA binding domain (THUMP), catalytic PPase domain, and only *EcoThiI* contains the rhodanese-like domain (RLD) among the five sources shown in this figure. The catalytic cysteines are indicated in red. The abbreviations of the protein source are the same as in Fig. 1-13.

In 2016, spectroscopic and biochemical research showed that the catalytic form of ThiI from *Methanococcus maripaludis* (*Mma*ThiI) contains a [3Fe-4S] cluster^[69]. On the other hand, the preliminary report on the structural determination of [4Fe-4S]-*Mma*ThiI has been published under strictly anaerobic conditions^[67]. Therefore, the reaction mechanism of s⁴U8 synthesis catalyzed by ThiI involving an Fe-S cluster is still unclear.

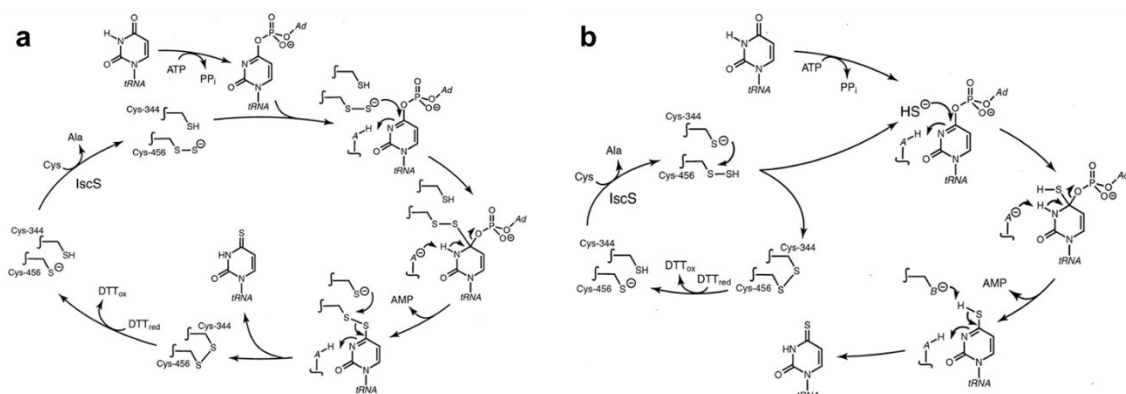


Fig. 1-26 Proposed reaction mechanism of s⁴U8 synthesis^[98].

(a) Direct pathway. Persulfide on Cys456 nucleophilically attacks the adenylated tRNA.
(b) Indirect pathway. Cys344 and Cys456 generate hydrogen sulfide, which serves as a nucleophile to attack the adenylated tRNA.

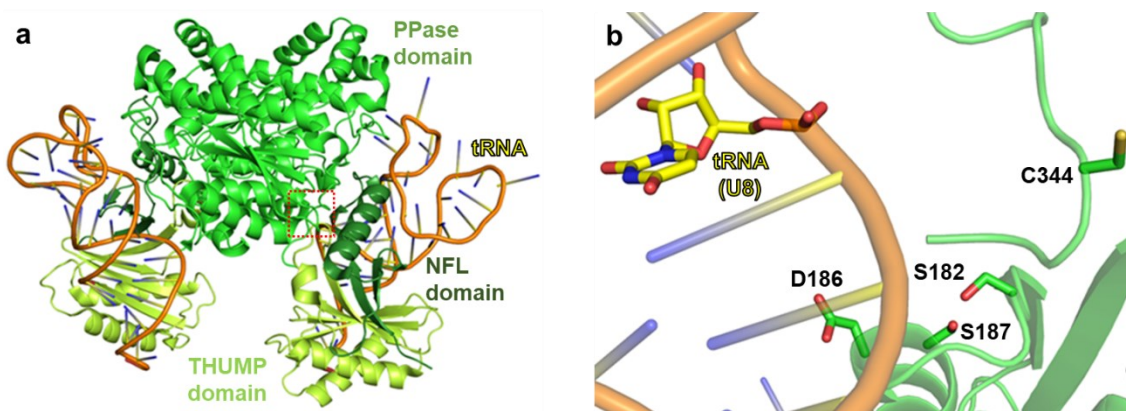


Fig. 1-27 Crystal structure of the *Tma*ThiI-tRNA^{Phe} (39-mer) complex^[105].

(a) Overall structure of the *Tma*ThiI-truncated tRNA^{Phe} complex (PDB ID: 4KR6). The catalytic site is marked by the red dotted frame. (b) Close-up view of the catalytic center of *Tma*ThiI. The side chains of the critical residues in the PPase domain are shown as a stick model.

1-6. The similarity of 2-thiouridine synthesis catalyzed by TtuA and Ncs6

TtuA and Ncs6 belong to the TtuA/Ncs6 family which contains zinc finger domains (CXXC...CXX[C/H] motif) and the PPase domain (SGGXD[S/T] motif) with [4Fe-4S]-binding site (CXXC...C motif) (Fig. 1-28a)^[92]. Two zinc fingers are predicted as tRNA binding sites and the PPase domain is identified as an ATP-binding site^[72,76]. Not only the amino acid sequence of *Tth*TtuA and *Mma*Ncs6, but their 3D structure is also similar (Cα RMSD = 1.649 Å, Fig. 1-28b). The inside of their catalytic pocket is positively charged, which is suitable for tRNA binding (Fig. 1-28c, d)^[92]. TtuA and Ncs6 also have two tunnels for sulfur donor protein and tRNA (Fig. 1-28e)^[72]. These structural similarities strongly support the similar catalytic mechanism of tRNA-thiolation.

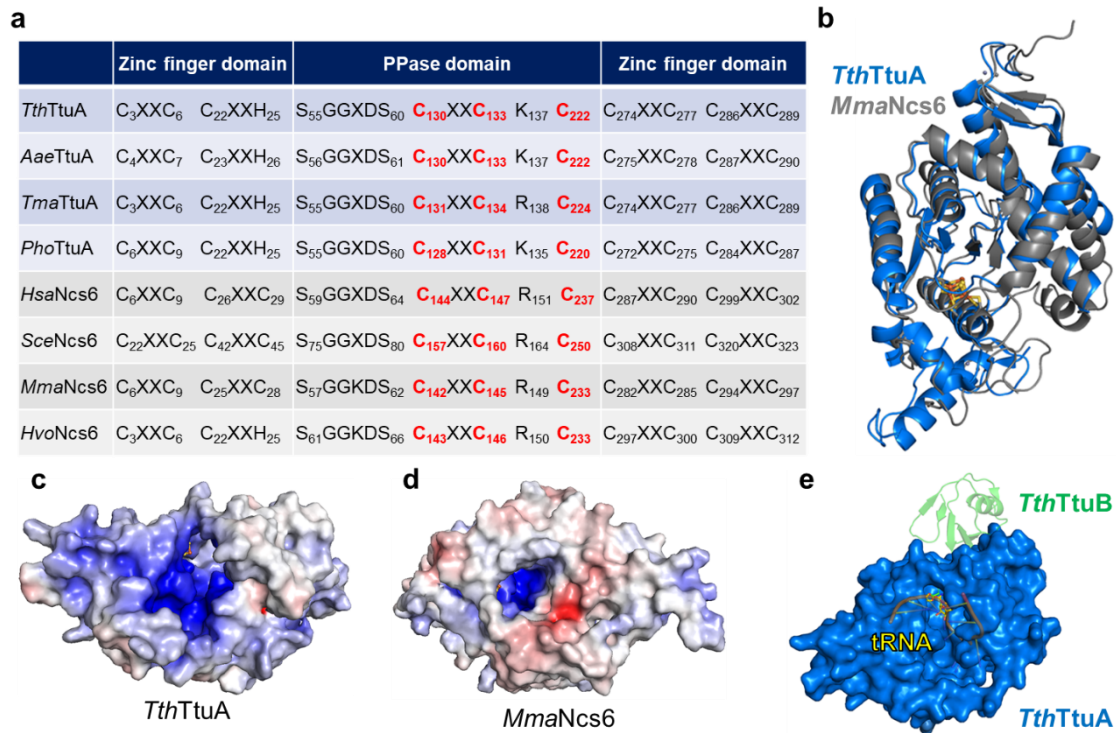


Fig. 1-28 Sequential and structural comparisons of TtuA and Ncs6.

(a) Domain construct and the location of critical residues in typical TtuA (blue) and Ncs6 (gray). TtuA and Ncs6 contain Zinc finger domains in the N and C-terminus regions. The [4Fe-4S]-binding cysteines are indicated in red. The abbreviations of the protein source are the same as in Fig. 1-13. (b) Superposition of the overall structures of [4Fe-4S]-*Tth*TtuA (PDB ID: 5B4F) and [4Fe-4S]-*Mma*Ncs6 (PDB ID: 6SCY). The zinc atoms are shown as gray balls. Electrostatic potential of the (c) *Tth*TtuA (PDB ID: 5B4F) and (d)

MmaNcs6 (PDB ID: 6SCY). For clearness, [4Fe-4S] is also shown after the calculation of the potential with the APBS plugin installed in PyMOL^[106]. The positively and negatively charged surface is colored blue (+10 kT/e) and red (-10 kT/e), respectively. (e) The [4Fe-4S]-TtuA-TtuB-tRNA complex model^[72]. The two tunnels through the catalytic site of TtuA allow TtuB and tRNA to enter the active site simultaneously.

Furthermore, some organisms employ TtuA or Ncs6 with sulfur donor protein TtuB or Urm1, respectively. Interestingly, the sulfur relay system from free L-cysteine to tRNA is very similar (Fig. 1-29)^[107,108]. In the first step, cysteine desulfurases IscS/SufS or Nfs1 transfer sulfur from L-cysteine to the rhodanese-like domain of TtuD or Tum1 (Yor251c), respectively^[87,107]. Next, TtuD or Tum1 transfers sulfur to the C-terminus of adenylated sulfur donor proteins (adenylated TtuB or Urm1, respectively) activated by E1-like proteins TtuC or UBA4, respectively^[79,85]. Finally, ubiquitin-like sulfur donors (TtuB or Urm1) transfer sulfur to substrate tRNA, which is catalyzed by TtuA or Ncs6, respectively.

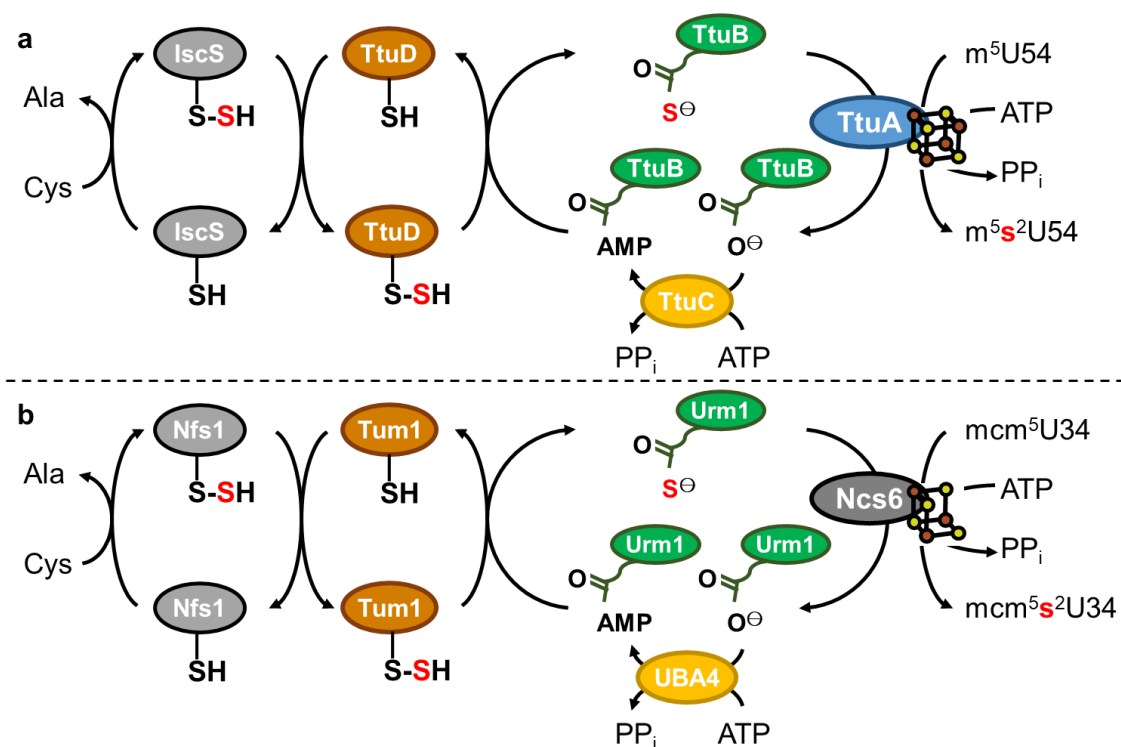


Fig. 1-29 Sulfur relay system in ubiquitin-like systems.

(a) m⁵s²U54 synthesis pathway. (b) mcm⁵s²U34 synthesis pathway. The type of Fe-S cluster in Ncs6 is still under discussion, but is shown as [3Fe-4S] in this figure.

1-7. Research question and purpose of this study

Considering the sequence identity (Fig. 1-13, Table 1-1), the similarity of 3D-structure (Fig. 1-28), and sulfur relay system (Fig. 1-29) among tRNA-thiolation enzymes, the opposite reports of [3Fe-4S]-Ncs6 and [4Fe-4S]-Ncs6 raised the question of whether TtuA only requires [4Fe-4S] or both [4Fe-4S] and [3Fe-4S] for its function. In previous studies on TtuA, [4Fe-4S] was reconstituted by excess iron and sulfur sources *in vitro* under anaerobic conditions^[72,76,83]. In other words, the possibility that TtuA contains [3Fe-4S] has not been examined. Furthermore, whereas the detailed mechanisms of sulfur transition catalyzed by Fe-S cluster-independent enzymes have been understood (Fig. 1-16), that of desulfurization from TtuB involved in Fe-S clusters was unknown.

In this study, I aimed to elucidate the detailed catalytic mechanism of tRNA thiolation based on the exact type of Fe-S clusters (Fig. 1-30). To achieve this purpose, I reconstituted [4Fe-4S]-TtuA and oxidized it to [3Fe-4S]-TtuA under strictly anaerobic conditions. Then, I analyzed the structure of the Fe-S cluster in the two types of TtuA using EPR spectroscopy in time-course and evaluated their activity. Furthermore, I analyzed the enzymatic activity of TtuA mutants and the condition of sulfur transition from TtuB to tRNA by biochemical assays. Then, I examined the structural similarity of tRNA-thiolation enzymes based on the bioinformatics approaches. Finally, I integrated all results and discuss the general catalytic mechanism of tRNA thiolation. Our findings would address the biochemical question of why all organisms employ oxygen-sensitive [4Fe-4S] in essential tRNA-thiolation.

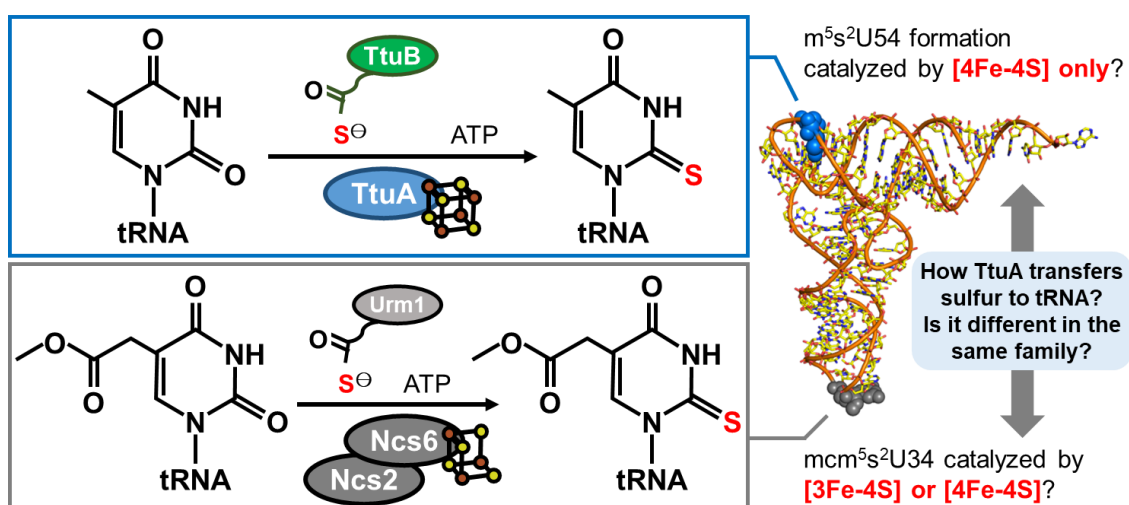


Fig. 1-30 Graphical abstract of this study.

Chapter 2. Materials and Methods

2-1. Expression of TtuA and TtuB

*Tth*TtuA (WT, UniProt ID: Q72LF3) was expressed as a C-terminal His6-tagged protein in *E. coli* (B834 DE3 strain) using the pET26 vector (Novagen)(Fig. 2-1a)^[72]. *Tth*TtuA (MT) was expressed as a N-terminal His6-tagged protein in *E. coli* (Rosetta DE3) using the pET15b vector (Fig. 2-1b)^[18]. Point mutations were introduced using the QuikChange II site-Directed Mutagenesis Kits (Stratagene) and designed primers (Sigma) (Table 2-1). I cultured recombinant *E. coli* in 3 L of lysogeny broth (LB) medium (Miller) containing 25 µg/mL kanamycin for C-His6-tagged TtuA or 100 µg/mL ampicillin for N-His6-tagged TtuA at 37°C and 150 rpm until the absorption at 600 nm (OD₆₀₀) reached 0.6. Then, I induced overexpression of TtuA with 1 mM isopropyl β-D-1-thiogalactopyranoside (IPTG) after 7°C cold shocks for 45 min, and cultured the cells at 25°C and 150 rpm for 16 hours. The cells were collected by centrifugation at 5,000 × *g* for 30 min and stored at -30°C.

*Tth*TtuB (UniProt ID: Q72LF4) was expressed as a C-terminal intein-tagged protein in *E. coli* (B834 DE3 strain) with the pTYB1 vector (New England BioLabs)(Fig. 2-1c)^[72]. I cultured recombinant *E. coli* in 3 L of LB containing 100 µg/mL ampicillin at 37°C and 150 rpm until OD₆₀₀ = 0.6. Then, I induced overexpression of TtuB and cultured the cells in the same manner as TtuA.

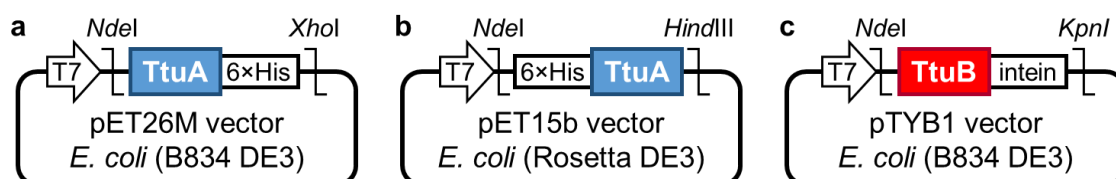


Fig. 2-1 Expression constructs of TtuA and TtuB.

(a) C-terminal His6-tagged TtuA for spectroscopic and biochemical analyses. (b) N-terminal His6-tagged TtuA for mutation assays. (c) C-terminal intein-tagged TtuB. All of these protein genes were inserted between the indicated multi-cloning site.

Sample Name	DNA sequence	Tm [°C]	GC ratio [%]	µg/OD	Mw
S55A_forward	CGGGGGGGAAGGACTCC	62	76	31.9	5301.5
S55A_reverse	CGACCGCCACCAGAACCC	59	72	32.7	5383.6
K58A_forward	CGGACTCCCTGGCCC	54	81	35.4	4819.2
K58A_reverse	CCCCCCCAGACCGC	63	87	34.7	4444.0
D59A_forward	GCTCCCTGGCCCTTTGG	56	72	36.7	5418.6
D59A_reverse	CTTCCCCCCCAGACCG	61	76	34.8	5052.4
S60A_forward	GCCCTGGCCCTTTGGGAC	61	72	35.3	5467.6
S60A_reverse	GTCCTTCCCCCCCAGAC	60	72	34.6	5356.5
K137A_forward	CGCGCTACATCATCAACCAGG	58	59	32.0	6689.4
K137A_reverse	GGAGAGGCCGCGAGGCG	64	81	31.5	4997.3
K137R_forward	CGCGCTACATCATCAACCAGG	58	59	32.0	6689.4
K137R_reverse	GGAGAGGCCGCGAGGCG	64	81	31.5	4997.3
D161A_forward	GCGAGGCCGCGCGTC	57	87	34.4	4555.0
D161A_reverse	GTCCAGGTTGTGCCCCGTG	62	67	34.0	5811.8
E203A_forward	GAGGGAGGTCCTCTCCTACAC	58	62	32.5	6407.2
E203A_reverse	CGCTAAAGCGGTAGAAGGGC	61	62	30.9	6545.3
E221A_forward	GGTGCCCGAACGCCAAG	58	72	32.5	5494.6
E221A_reverse	CTCGTGGAGGTAGCGGATCC	60	65	32.3	6174.1
K234A_forward	CGGAGGCCCTGAACCTG	55	72	33.1	5525.6
K234A_reverse	GTAGAGGAGGCTTTTCGCCCC	62	62	33.0	6438.2

Table 2-1 Designed primers for mutagenesis in the N-terminal His6-tagged TtuA.

Each primer is shown with its DNA sequence, melting temperature (T_m), GC ratio, conversion factor (µg/OD), and molecular weight (Mw). T_m values were calculated by a nearest neighbor thermodynamic algorithm with SnapGene Viewer version ver6.2 (<https://www.snapgene.com/>)^[109]. Nucleotides colored in red indicate mutation sites for the QuikChange method.

2-2. Anaerobic purification of TtuA

Lysis of recombinant *E. coli* and purification of TtuA were performed under strictly anaerobic conditions with 5% H₂ and 95% N₂ (Fig. 2-2a)^[72]. The collected cells cultured in 1 L of LB were lysed by sonication on ice for 45 min in the deoxidized purification buffer (50 mM HEPES-KOH (pH 7.6), 200 mM ammonium sulfate, 50 mM ammonium acetate, 5 mM magnesium chloride, 10% (v/v) glycerol, and 7 mM 2-mercaptoethanol) containing 0.1% Triton X-100. Then the cells were heat treated at 70°C for 20 min, and

the precipitates were removed by centrifugation at $7,000 \times g$ for 60 min and a 0.22- μm filter (Millipore). The supernatant was loaded onto a Ni-affinity chromatography (NiAC) column (1 mL His-Trap HP; GE Healthcare) equilibrated with the purification buffer. Non-specifically bound proteins were removed using the wash buffer (purification buffer containing 50 mM imidazole). TtuA was eluted with a gradient of 50-500 mM imidazole in the purification buffer (Fig. 2-2b). Further, I purified the eluted samples with a size exclusion chromatography (SEC) column (HiLoad 16/60 Superdex 200, GE Healthcare) equilibrated with the purification buffer (Fig. 2-2c). I checked the sample purity using 12.5% (v/v) SDS-PAGE at 40 mA for 60 min, then protein bands were visualized by CBB-G250 stain and detected using the Amersham Imager 680 (GE Healthcare) (Fig. 2-2d).

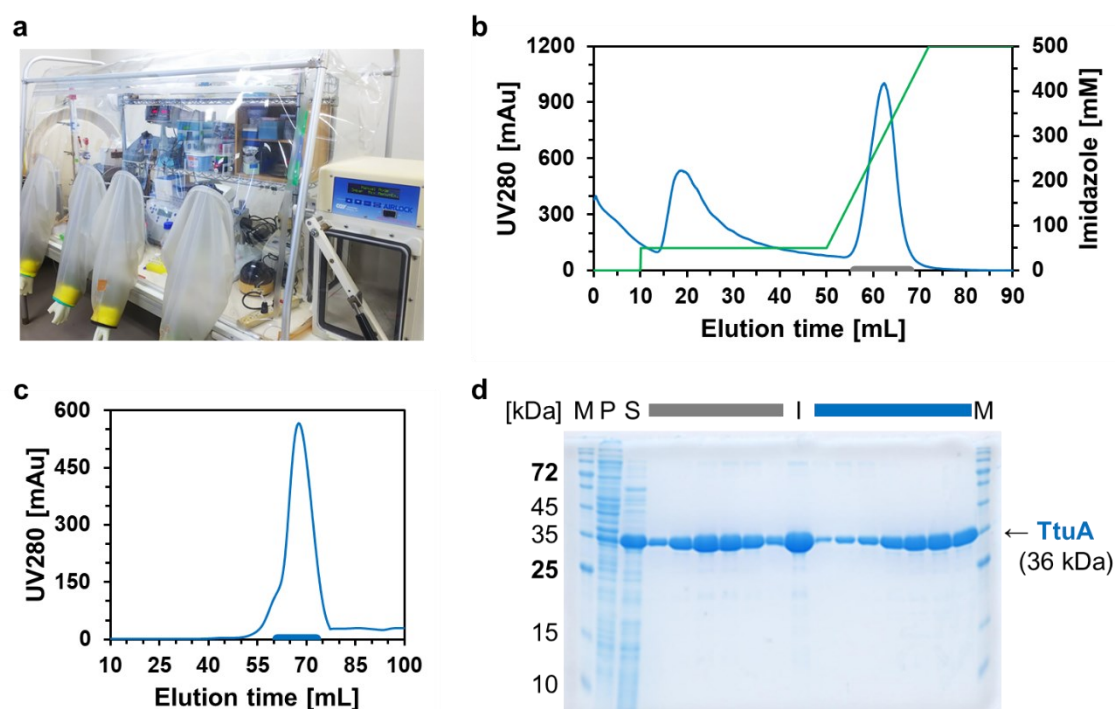


Fig. 2-2 Purification of TtuA (WT) in the anaerobic conditions.

(a) Vinyl Anaerobic Chamber (COY). (b) NiAC spectrum measured at wavelength of 280 nm (UV280). The collected sample was indicated in the gray band. (c) SEC spectrum measured at UV280. The collected sample was indicated in the blue band. (d) 12.5% (v/v) SDS-PAGE stained with CBB-G250. The two bold numbers indicate the molecular mass of reference bands of the PM1500 marker (SMOBIO). Sample M, P, S, and I are marker, precipitant of *E. coli* cells, supernatant, and injection to the SEC column, respectively.

2-3. Purification of thiocarboxylated TtuB (TtuB-COSH)

I sonicated the collected cells for 30 min in the purification buffer (20 mM Tris-HCl (pH 8.5 at 25°C) and 500 mM NaCl) with 0.1% Triton X-100, 0.5 mg/mL lysozyme (Sigma), and 0.1 mg/mL DNase I (Sigma). The precipitates were removed by centrifugation at $40,000 \times g$ for 30 min and a 0.22- μm filter (Fig. 2-3a, lane P). The supernatant was loaded onto a chitin resin (New England BioLabs) equilibrated with the purification buffer (Fig. 2-3a, lanes S and F). Non-specifically bound proteins were removed using 20 column volumes (CV) of the purification buffer (Fig. 2-3a, lane W). To cleave the intein tag from the TtuB-intein fusion protein, 1 CV of the purification buffer containing 50 mM ammonium sulfide ($(\text{NH}_4)_2\text{S}$) was added to the chitin resin and incubated at room temperature (RT) for 20 hours.

TtuB was eluted in a 7°C cold room with the purification buffer and then concentrated using an Amicon Ultra Centrifugal Filter (3-kDa cutoff). Excess $(\text{NH}_4)_2\text{S}$ was removed with the Sephadex PD-10 desalting column (GE Healthcare) equilibrated with the storage buffer (14 mM Tris-HCl (pH 8.5) at 25°C, 350 mM NaCl, and 30% (v/v) glycerol). The fractions containing TtuB were collected, concentrated using an Amicon Ultra Centrifugal Filter (3-kDa cutoff), and stored at -80°C. I confirmed the purity of TtuB by 15% (v/v) SDS-PAGE similar to TtuA (Fig. 2-3a, lane E). Note that the intein tag was removed from the C-terminus of TtuB without protease treatment^[110].

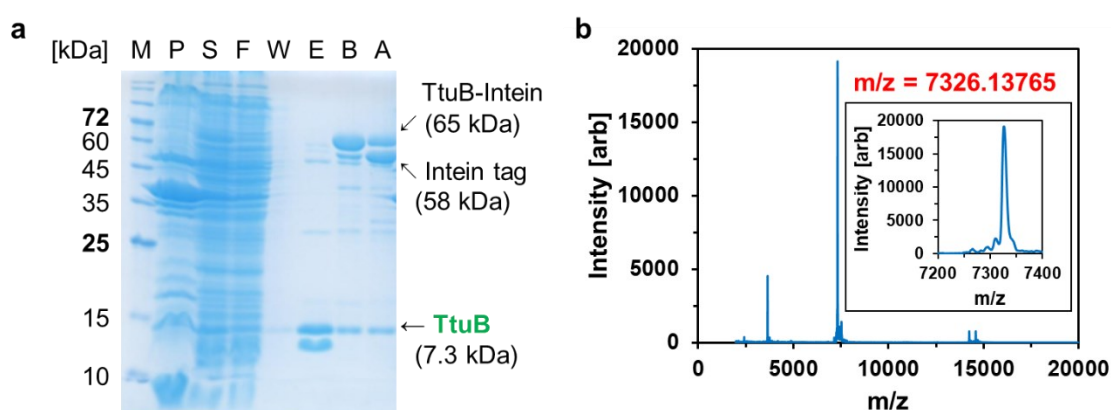


Fig. 2-3 Purification of TtuB-COSH.

(a) 15% (v/v) SDS-PAGE stained with CBB-G250. The two bold numbers indicate the molecular mass of reference bands of the PM1500 marker. Sample M, P, S, W, E, B, and A are a marker, a precipitant of *E. coli* cells, supernatant, flow-through, wash, elution,

before the cleavage reaction, and after the cleavage reaction, respectively. (b) Overall and enlarged MALDI-TOF MS spectra showed that the C-terminus of TtuB is not carboxylate group ($m/z = 7310.32$), but thiocarboxylate group ($m/z = 7326.43$).

To confirm the presence of sulfur at the C-terminus of TtuB, I removed salts from TtuB solution on ice with the C4 ZipTip (Millipore) and performed matrix-assisted laser desorption/ionization-time-of-flight mass spectrometry (MALDI-TOF MS) using UltraflexIII (Bruker) with sinapinic acid as the matrix^[72]. The concentration of TtuB was determined by the Bradford method^[111] using Protein Assay Dye Reagent Concentrate (Bio-Rad) because *Tth*TtuB does not contain tyrosine and tryptophan.

2-4. Reconstitution of [4Fe-4S]-TtuA and [3Fe-4S]-TtuA

All steps to reconstitute [4Fe-4S]-TtuA and [3Fe-4S]-TtuA were performed under strictly anaerobic conditions (Fig. 2-2a). Firstly, I measured the concentration of eluted TtuA after SEC with the Nanodrop DU 1000 U (Thermo Fisher). To incorporate [4Fe-4S] into TtuA, I incubated TtuA with 5 mM dithiothreitol (DTT) for 10 min at RT. Then, I added 9-fold molar excess of ferric chloride (FeCl_3) to the solution and incubated the mixture for a further 10 min at RT. Subsequently, 9-fold molar excess sodium sulfide (Na_2S) was added and the mixture was incubated for 3 hours at RT. The precipitate of iron sulfide was removed by centrifugation at $7,000 \times g$ for 10 min and a 0.22- μm filter. [4Fe-4S]-TtuA was concentrated using an Amicon Ultra Centrifugal Filter (30-kDa cutoff). Excess FeCl_3 and Na_2S were removed using the Sephadex PD-10 desalting column equilibrated with the purification buffer (Fig. 2-4a).

To remove the unique Fe and produce [3Fe-4S]-TtuA, I added 6-fold molar excess $\text{K}_3[\text{Fe}(\text{CN})_6]$ to [4Fe-4S]-TtuA and incubated the mixture at RT for 10 min. The precipitates were removed using centrifugation at $7,000 \times g$ for 30 sec and a 0.22- μm filter. $\text{K}_3[\text{Fe}(\text{CN})_6]$ and free Fe were removed with a Sephadex PD-10 desalting column equilibrated with the purification buffer (Fig. 2-4b). Notably, I used oxidized TtuA immediately for EPR spectroscopy or activity assay due to the instability of [3Fe-4S]-TtuA.

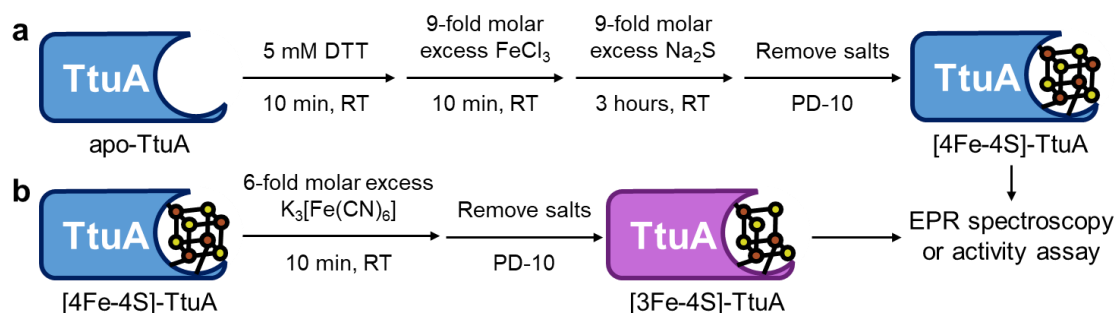


Fig. 2-4 Reconstitution of TtuA with an Fe-S cluster in the anaerobic chamber.

Reconstitution of (a) [4Fe-4S]-TtuA using an excess amount of iron and sulfur sources under reduced conditions and (b) [3Fe-4S]-TtuA using an excess amount of oxidant, K₃[Fe(CN)₆].

2-5. Structural determination of Fe-S clusters by time-course EPR spectroscopy

To analyze the structure of the Fe-S cluster bound to TtuA in time-course, I prepared EPR samples under strictly anaerobic conditions. Because [4Fe-4S]²⁺ and Fe²⁺ are EPR-silent, I evaluated [4Fe-4S]¹⁺, [3Fe-4S]¹⁺, and free Fe³⁺ using EPR spectroscopy to determine the structure of the Fe-S cluster in TtuA. I transformed reconstituted [4Fe-4S]²⁺-TtuA into [4Fe-4S]¹⁺-TtuA with a reductant, dithionite (DT) to obtain the spectra of [4Fe-4S]-TtuA^[72] and evaluated [3Fe-4S]¹⁺ and free Fe³⁺ for oxidized [4Fe-4S]-TtuA (Fig. 2-5).

Immediately after [3Fe-4S]-TtuA was prepared, I divided fresh 0.5 mM [3Fe-4S]-TtuA into 14 samples (200 µL/sample). For analyzing [3Fe-4S]-TtuA, DT was not added to the sample 5 min after preparation of [3Fe-4S]-TtuA (sample 1). To analyze [4Fe-4S]-TtuA, I added 5-fold molar excess DT to the sample 5 min after preparation of [3Fe-4S]-TtuA (sample 2) and incubated the sample at 25°C for 10 min. To prevent degradation of the Fe-S clusters, samples 1 and 2 were aliquoted into quartz EPR tubes (Agri) and frozen simultaneously with liquid N₂ in the anaerobic chamber. Similarly, I froze samples 10 min (samples 3 and 4), 20 min (samples 5 and 6), 30 min (samples 7 and 8), 1 hour (samples 9 and 10), 2 hours (samples 11 and 12), and 24 hours (samples 13 and 14) after preparation of [3Fe-4S]-TtuA (Fig. 2-6). Moreover, 200 µL of 0.5 mM [4Fe-4S]-TtuA was frozen as control to analyze the amount of [4Fe-4S]-TtuA before oxidation (sample 0).

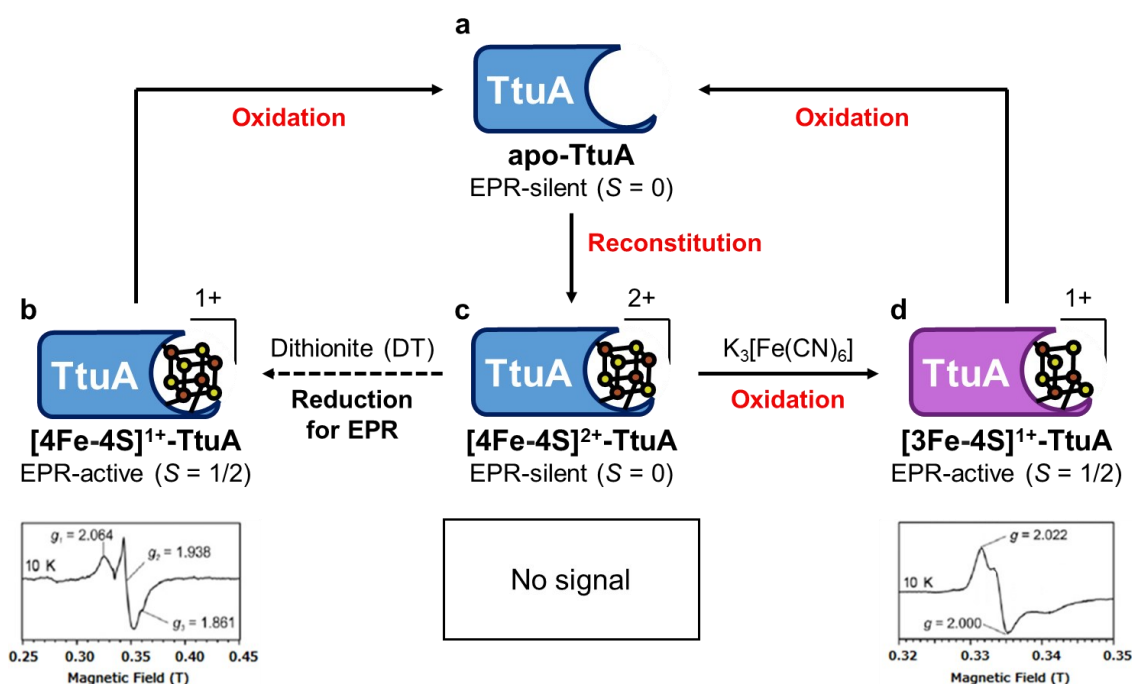


Fig. 2-5 Redox states of TtuA and their detectability by EPR spectroscopy.

(a) Reconstructed TtuA is +2 charged with spin quantum number $S = 0$ and EPR-silent. (b) Reduction using DT enables EPR detection of [4Fe-4S]-TtuA as +1 charged state. (c) The normal state of TtuA with [4Fe-4S]²⁺. (d) Oxidation with $K_3[Fe(CN)_6]$ enables EPR detection of [3Fe-4S]-TtuA. These EPR spectra were cited from our previous research^[72].

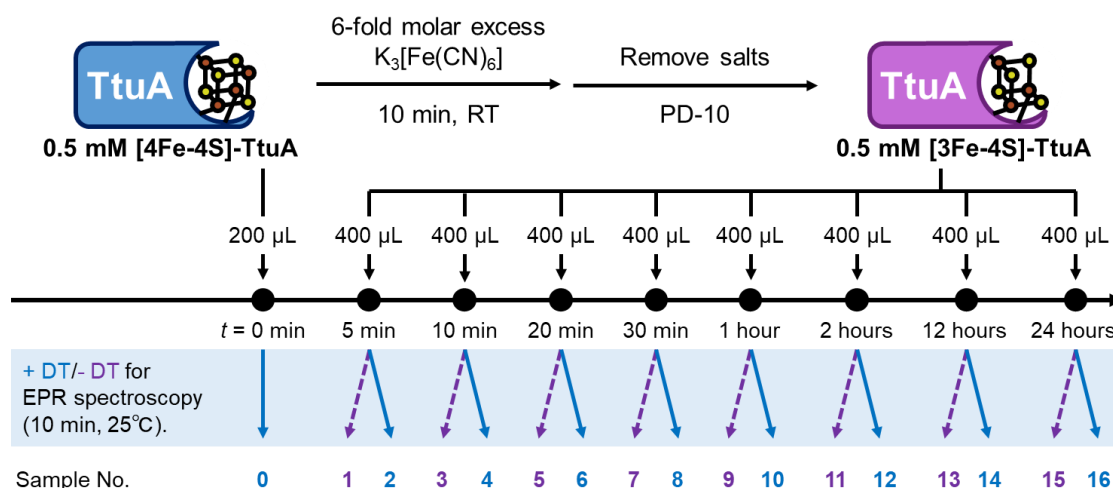


Fig. 2-6 Anaerobic preparation of EPR samples in time-course^[112].

Even-numbered EPR samples contain DT to evaluate [4Fe-4S] (blue solid arrows). Odd-numbered EPR samples do not contain DT to evaluate [3Fe-4S] (purple dashed arrows).

2-6. Interaction analysis between TtuA and TtuB by EPR spectroscopy

To analyze the interaction between [3Fe-4S]-TtuA and TtuB-COSH, I prepared EPR samples using a slightly modified version of our protocol^[83]. Fresh [3Fe-4S]-TtuA was mixed with TtuB to a final concentration of 0.5 mM and incubated at RT for 5 min under strictly anaerobic conditions. The sample mixture was incubated with a 5-fold molar excess DT at RT for 10 min, aliquoted into quartz EPR tubes, and frozen with liquid N₂ in the anaerobic chamber. The EPR samples were stored in liquid N₂ and transported to the Center for Experimental Science and Analysis at Saga University or Institute for Molecular Science in Okazaki. Continuous wave (CW) X-band EPR spectra were measured with parameters described in the original paper^[112].

To quantify [4Fe-4S] and [3Fe-4S] in each sample, I double-integrated the EPR spectra with the peak areas of [4Fe-4S] and [3Fe-4S]. For estimating the maximum amount of [3Fe-4S] regardless of the quick structural change of [3Fe-4S] to [4Fe-4S], I plotted the signal intensity of [3Fe-4S] versus time after oxidation, and then performed linear approximation by using samples up to 120 min after oxidation. I calculated the R² value with Microsoft Excel 2019 and found that the result (0.9276) was reliable.

2-7. Activity assay for [3Fe-4S]-TtuA and [4Fe-4S]-TtuA by HPLC

The formation of m⁵s²U54 on substrate tRNA was conducted under strictly anaerobic conditions^[83]. The standard assay was performed in 30 µL of reaction buffer (50 mM HEPES-KOH (pH 7.6), 100 mM KCl, 10 mM MgCl₂, and 0.1 mM DTT) containing 2.5 µM TtuA, 15 µM brewer's yeast total tRNA (Sigma), 15 µM TtuB, and 5 mM ATP at 60°C for 10 min or 30 min. Because [3Fe-4S] in TtuA quickly transforms into [4Fe-4S], it was necessary to initiate tRNA-thiolation immediately after removing K₃[Fe(CN)₆] from the TtuA solution and limit reaction time. In our previous study, I incubated 4-fold molar excess of TtuB (20 µM) with 5 µM of TtuA at 60°C for 30 min^[83]. In this study, I obtained a final concentration of 2.5 µM for TtuA and analyzed the time-course of enzymatic activity (Fig. 2-7). To detect enzymatic activity even at low concentrations of TtuA and short reaction times, I added 6-fold molar excess of TtuB (15 µM).

tRNA-thiolation was stopped with 120 µL stop buffer (75 µL Isogen (Nippon Gene) and 45 µL deionized water) and then frozen at -30°C until tRNA extraction. tRNA was

extracted with phenol:chloroform (5:1 (pH 4.5); Thermo Fisher), precipitated with ethanol, and digested at 37°C for 4 hours with digestion buffer (100 mM HEPES-NaOH (pH 7.5)) containing 6.2 mU/μL nuclease P1 (Yamasa) and 5.0 mU/μL bacterial alkaline phosphatase (Takara Bio). The digested samples were loaded onto the Inertsil ODS-3 column (2.1 mm × 150 mm × 3 μm; GL Science) equilibrated with high-performance liquid chromatography (HPLC) buffer (0.1% formic acid and 2% acetonitrile). m^5s^2U was eluted with a gradient of 2%-36% acetonitrile. The amount of m^5s^2U was detected with UV280 using the Extrema HPLC system (Jasco) (Fig. 2-8).

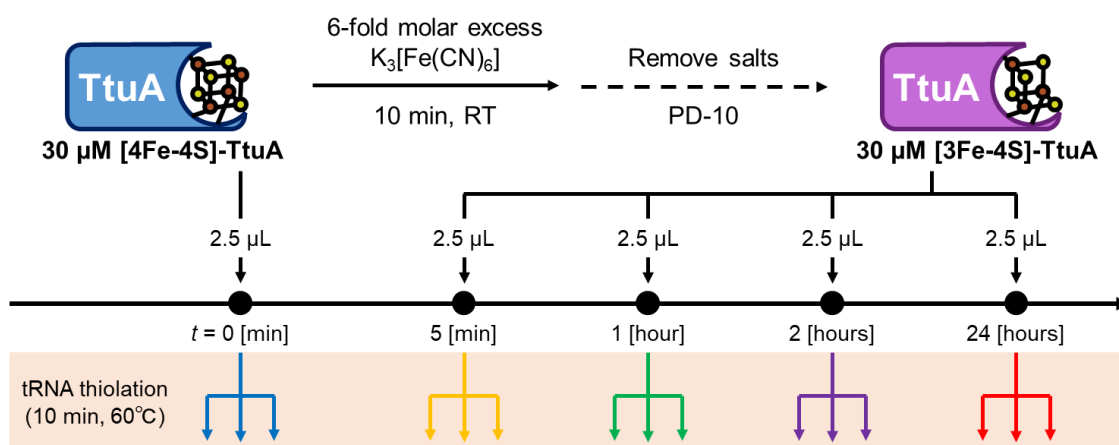


Fig. 2-7 Anaerobic preparation of samples for time-course activity assay.

The enzymatic reaction was started at 60°C for 10 min immediately after the preparation of [3Fe-S]-TtuA. Desalination is an optional step. All assays were performed three times.

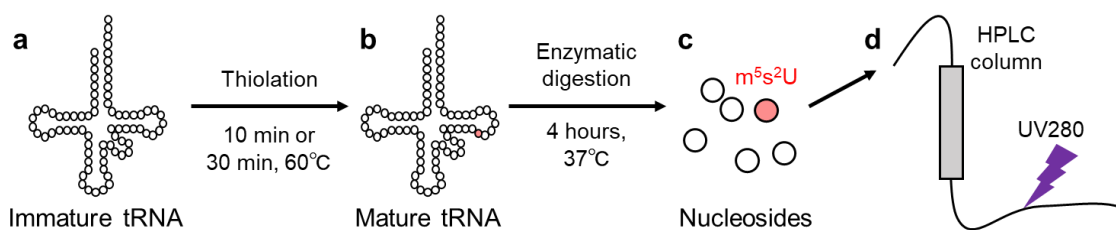


Fig. 2-8 Scheme of activity assay of TtuA by HPLC.

(a) tRNA without m^5s^2U 54 modification. (b) TtuA with an active form of Fe-S cluster catalyzes m^5s^2U 54 synthesis. The reaction time is 10 min for time-course assay using [3Fe-S]-TtuA and 30 min for mutation assays, respectively. (c) Enzymatic digestion of tRNA at 37°C for 4 hours. (d) Sample injection to the HPLC column and detect compounds by UV280.

2-8. Desulfurization assay by mercury-gel electrophoresis (APM-PAGE)

To reveal the condition when TtuB releases sulfur from its C-terminus, I performed desulfurization assays by 15% (v/v) SDS-PAGE with 12 $\mu\text{g/mL}$ of [(*N*-Acryloylamino) phenyl]mercuric chloride (APM; Toronto Research Chemicals) in the lower gel (Fig. 2-9a)^[104,113]. Since APM is toxic, I performed APM-PAGE using protective masks as well as gloves and lab coats, and experimental tools and waste liquid contaminated by APM were collected properly (Fig. 2-9b). Mercury in APM strongly interacts with sulfur atoms of TtuB, which enables to distinguish between carboxylated TtuB (TtuB-COOH) and thiocarboxylated TtuB (TtuB-COSH) (Fig. 2-9c).

APM-gel was prepared at 25°C (lower gel: 15% acrylamide/bis mixed solution, 360 mM Tris-HCl (pH 8.5), 0.1% SDS, 12 $\mu\text{g/mL}$ APM, upper gel: 4.5% acrylamide/bis mixed solution, 120 mM Tris-HCl (pH 6.0) at 25°C, 0.1% SDS) at RT. Then, I performed APM-PAGE at 20 mA for 80 min in 7°C cold room to stabilize TtuB-COSH. Protein bands were visualized by CBB-G250 stain and detected using the LAS-3000 imaging system (Fujifilm) (Fig. 2-9d).

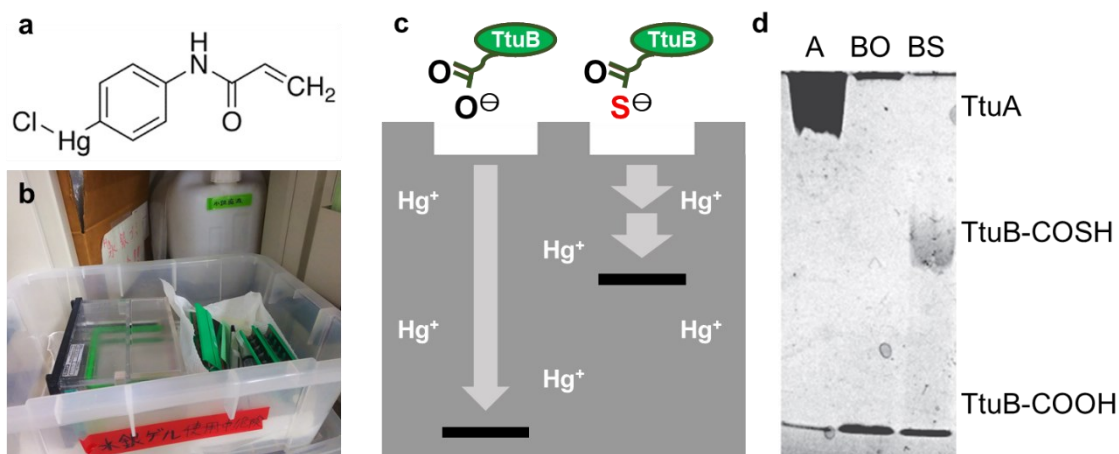


Fig. 2-9 Desulfurization assay by APM-PAGE.

(a) Structural formula of APM. (b) Experimental tools. (c) Concept of APM-PAGE. (d) Visualization of proteins in APM-gel. Sample A, BO, and BS are 600 pmol of [4Fe-4S]-TtuA, TtuB-COOH, and TtuB-COSH, respectively. The image was cited from our original paper^[83]. I prepared TtuB-COOH using DTT in the cleavage reaction instead of (NH₄)₂S. Note that TtuB-COSH contains a small amount of TtuB-COOH as a byproduct in the cleavage reaction.

2-9. Structural prediction and comparison of tRNA-thiolation enzymes

I downloaded 3D structures models of *HsaNcs6*, *EcoTtcA*, and *MmaThiI* predicted from the AlphaFold Protein Structure Database on 17th November 2022^[114]. Since these models do not contain Fe-S clusters, I superimposed the predicted models with [4Fe-4S]-*TthTtuA* (PDB ID: 5B4F) using the “align” command for Ncs6 and TtcA, whereas I used the “super” command for MnmA (PDB ID: 2DEU) and the PPase domain of *MmaThiI* (G182-H354) in PyMOL ver1.7. (Fig. 2-10).

Furthermore, I predicted the structure of *HsaNcs6-HsaUrm1* complex using local version of AlphaFold version 2.0 (AF2)^[115] with default parameters. I chose one model with the highest reliability based on the predicted local-distance difference test (pLDDT) score among the five predicted structures. To build enzyme-tRNA complex models, I superimposed tRNA-thiolation enzymes with a tRNA-modification enzyme TilS complexed with tRNA (PDB ID: 3A2K) in the same method as our previous study^[72]. TilS catalyzes lysidine synthesis at position C34 of tRNA^{Ile(CAU)}^[116,117].

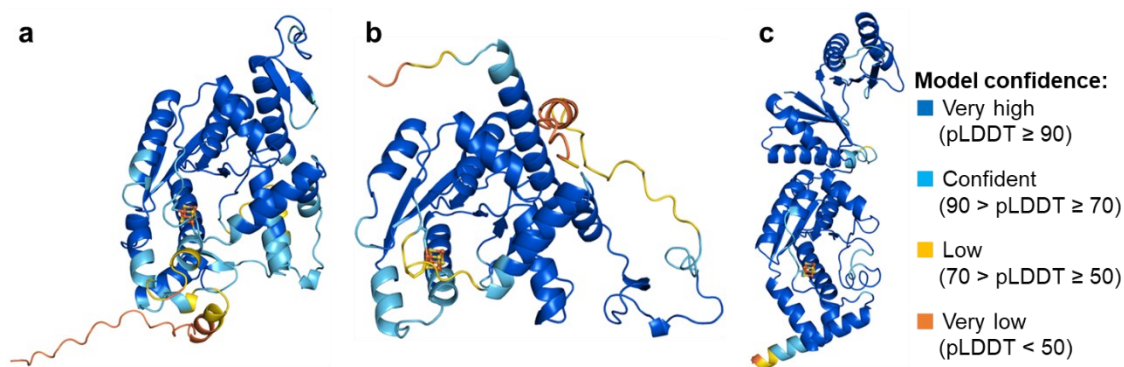


Fig. 2-10 Overall structure of tRNA-thiolation enzymes predicted by AlphaFold2.

(a) [4Fe-4S]-*HsaNcs6* model. (b) [4Fe-4S]-*EcoTtcA* model. (c) [4Fe-4S]-*MmaThiI* model. To visualize the reliability of predicted structures, regions other than [4Fe-4S] were colored based on the pLDDT score (pLDDT ≥ 90 in blue, $90 > \text{pLDDT} \geq 70$ in cyan, $70 > \text{pLDDT} \geq 50$ in yellow, pLDDT < 50 in orange, respectively). The C α RMSD between each AF2 model and *TthTtuA* is 1.314 Å (for *HsaNcs6*), 2.173 Å (for *EcoTtcA*), and 2.766 Å (for *MmaThiI*), respectively.

Chapter 3. Results

3-1. [3Fe-4S]-TtuA spontaneously and quickly transforms into [4Fe-4S]-TtuA

The EPR spectra of fresh oxidized TtuA showed a peak of $[3\text{Fe-4S}]^{1+}$ ($g = [2.02, 2.01, 1.97]$) (Fig. 3-1a)^[72] and that of free Fe^{3+} ($g = 4.3$) (Fig. 3-1b)^[118]. Considering that free Fe and excess $\text{K}_3[\text{Fe}(\text{CN})_6]$ were removed by desalination, free Fe should come from $[3\text{Fe-4S}]$ -TtuA. Notably, signal intensity of $[3\text{Fe-4S}]^{1+}$ and free Fe decreased with time after desalination under strictly anaerobic conditions (Fig. 3-1a, 3-1b). This observation indicated that $[3\text{Fe-4S}]$ cluster in TtuA is unstable and quickly transforms into $[4\text{Fe-4S}]$. Even I skipped the desalination in the preparation of $[3\text{Fe-4S}]$ -TtuA (which means that the solution contains free Fe and excess $\text{K}_3[\text{Fe}(\text{CN})_6]$), the signal intensity of $[3\text{Fe-4S}]^{1+}$ decreased similar to that after removing excess free Fe and $\text{K}_3[\text{Fe}(\text{CN})_6]$ (Fig. 3-2).

Furthermore, to evaluate the amount of $[4\text{Fe-4S}]$ in all the test samples, I reduced the samples with DT and analyzed their EPR spectra. The signal intensity of $[4\text{Fe-4S}]^{1+}$ increased with time (Fig. 3-1c). The DT addition to the solution did not affect the conformational change of Fe-S clusters in TtuA (Fig. 3-2c).

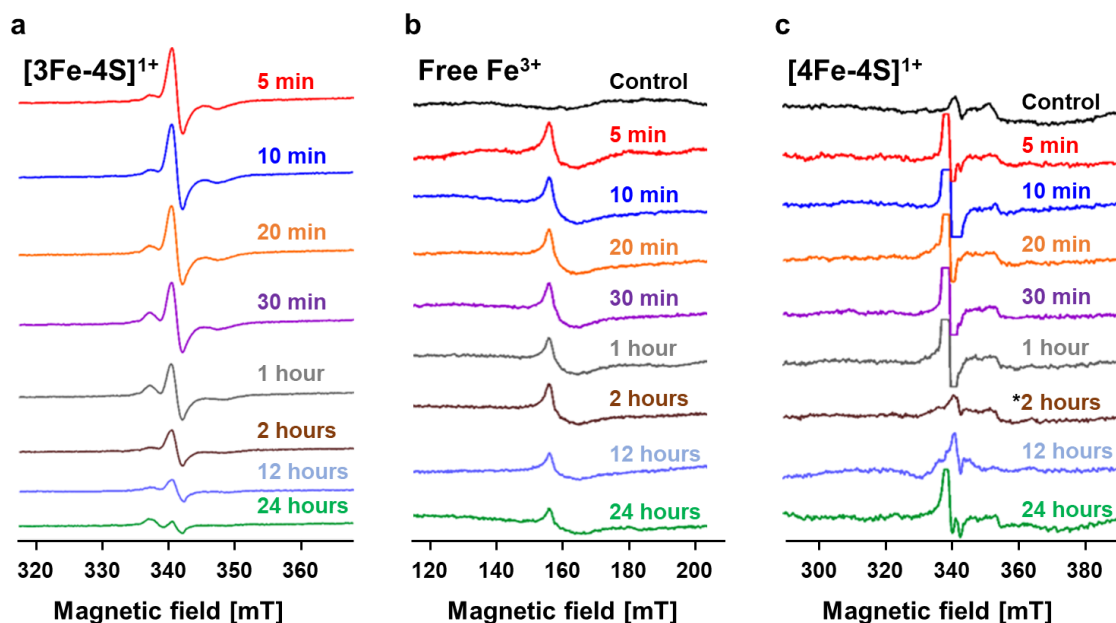


Fig. 3-1 EPR spectra of TtuA oxidized with $\text{K}_3[\text{Fe}(\text{CN})_6]$ ^[112].

Signal peak of (a) $[3\text{Fe-4S}]^{1+}$ measured at 40 K, (b) free Fe^{3+} measured at 40 K, and (c) $[4\text{Fe-4S}]^{1+}$ measured at 12 K. An asterisk (*) in Fig. 3-1c indicates a weak signal with a relatively larger experiment error.

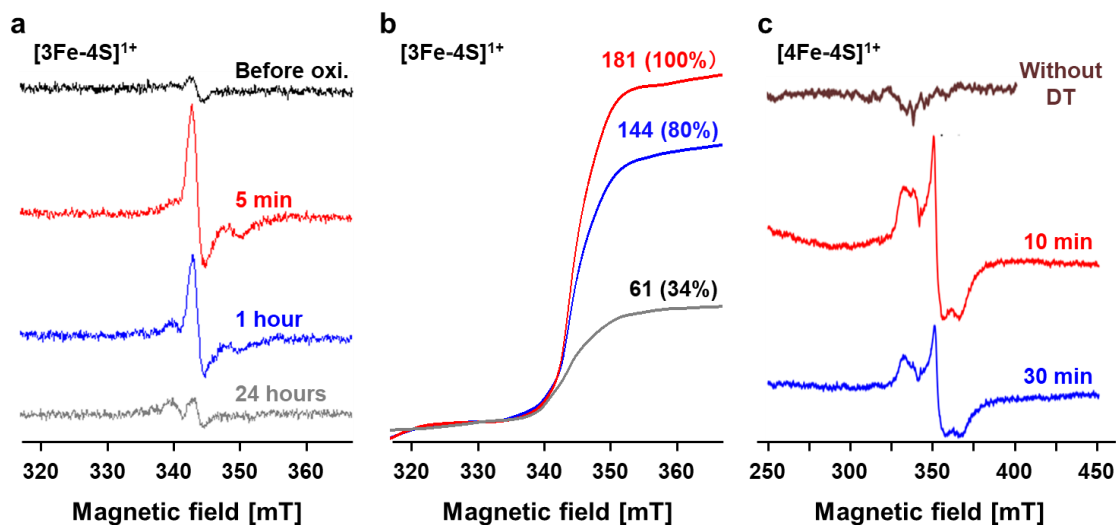


Fig. 3-2 Effect of excess $\text{K}_3[\text{Fe}(\text{CN})_6]$ and DT in EPR samples of TtuA^[112].

(a) $[4\text{Fe-4S}]\text{-TtuA}$ was oxidized to $[3\text{Fe-4S}]\text{-TtuA}$ with excess $\text{K}_3[\text{Fe}(\text{CN})_6]$ (without desalination) and incubated for 5 min (red), 1 hour (blue), and 24 hours (gray). The sample before oxidation ($[4\text{Fe-4S}]\text{-TtuA}$) was used as a reference (black). $[3\text{Fe-4S}]^{1+}$ was detected in the absence of DT at 40 K. (b) Double integration of Fig. 3-2a. The ratio of signal intensity is shown in parentheses, where the intensity at 5 min after oxidation is normalized to 100%. (c) Spectroscopic characterization of TtuA with excess DT. EPR spectra of $[4\text{Fe-4S}]\text{-TtuA}$ before DT reduction (brown) and reduced with DT and incubated for 10 min (red) and 30 min (blue). $[4\text{Fe-4S}]^{1+}$ were detected with DT at 12 K.

Next, I analyzed the amount of $[4\text{Fe-4S}]$ and $[3\text{Fe-4S}]$ in the time-course EPR spectroscopy (Fig. 3-3a). Since quantifying the amount of $[3\text{Fe-4S}]$ immediately (0 min) after oxidation is challenging due to experimental limitations, I estimated an approximate line correlation coefficient of 3.877 for the signal intensity of $[3\text{Fe-4S}]$ as a standard value to calculate the amount of $[3\text{Fe-4S}]$ (Fig. 3-3b). When the amount of $[4\text{Fe-4S}]$ is normalized to 100% in TtuA before oxidation, $[3\text{Fe-4S}]\text{-TtuA}$ was estimated to contain ~10% $[4\text{Fe-4S}]$ after 5 min of oxidation. Nearly one-third of $[3\text{Fe-4S}]\text{-TtuA}$ transformed into $[4\text{Fe-4S}]\text{-TtuA}$ after 1 hour of oxidation (Table 3-1). The collapse of $[3\text{Fe-4S}]$ and a corresponding increase in $[4\text{Fe-4S}]$ indicated that the unstable $[3\text{Fe-4S}]$ cluster of $[3\text{Fe-4S}]\text{-TtuA}$ spontaneously transformed into stable $[4\text{Fe-4S}]$ ($[4\text{Fe-4S}]\text{-TtuA}$) within one hour.

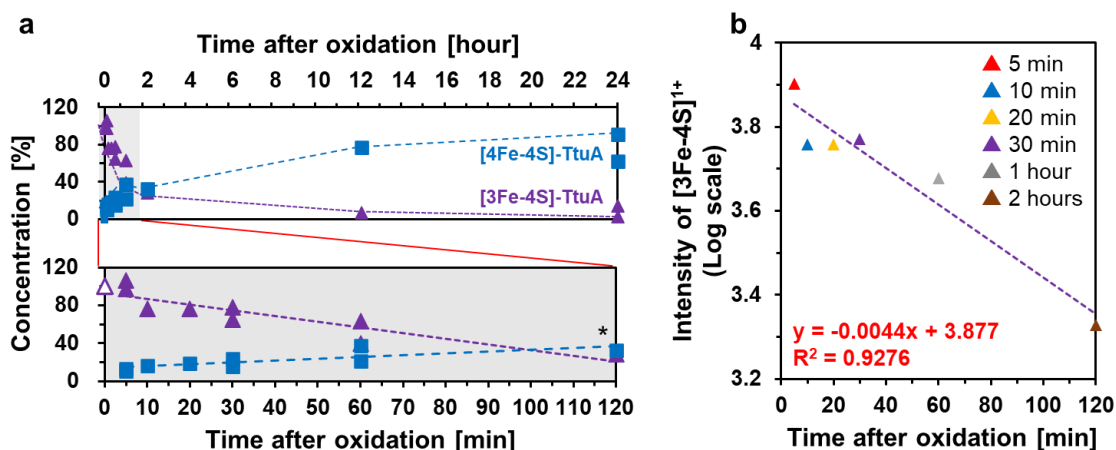


Fig. 3-3 Quantification of the structural changes in the Fe-S cluster in TtuA^[112].

(a) The plot of structural changes of Fe-S clusters in time-course, which was calculated from Fig. 3-1a and c. An asterisk (*) in Fig. 3-3a indicates a weak signal with a relatively larger experiment error. (b) Calibration curve of the signal intensity of $[3\text{Fe-4S}]^{1+}$ versus time after oxidation. Each point is colored in the same as Fig. 3-1.

Table 3-1. Spectroscopic quantification of the Fe-S cluster in TtuA^[112].

Time after oxidation	[3Fe-4S]	[4Fe-4S]
Before oxidation	—	100%
Immediately after oxidation (0 min)	(100%)	—
5 min [†]	99% ± 7%	12% ± 1%
10 min	76%	16%
20 min	76%	19%
30 min [†]	70% ± 8%	20% ± 4%
1 hour [†]	50% ± 13%	30% ± 8%
2 hours [*]	28%	33%
12 hours	7%	77%
24 hours [†]	9% ± 6%	77% ± 14%

EPR measurements were performed twice under condition marked with daggers (†) and one time without the dagger. The standard deviation was calculated by $N = 2$. The parentheses indicate the estimated amount from the calibration curve (Fig. 3-3b). The asterisk (*) indicates a weak signal with a relatively larger experiment error.

3-2. [3Fe-4S]-TtuA is an inactive form in tRNA thiolation

To analyze the enzymatic activity of [3Fe-4S]-TtuA, I performed activity assays of TtuA in time-course under strictly anaerobic conditions. Since [3Fe-4S] transforms into [4Fe-4S] within one hour (Fig. 3-3a), limiting reaction time is essential. I optimized the reaction conditions, including TtuA and TtuB concentrations and temperature, to monitor enzymatic activity in a relatively short reaction time.

I measured the yield of m^5s^2U using oxidized [4Fe-4S]-TtuA in the presence and absence of $K_3[Fe(CN)_6]$. Whereas m^5s^2U was detectable using [4Fe-4S]-TtuA, nearly no m^5s^2U was synthesized 5 min after oxidation, indicating that [3Fe-4S]-TtuA has no enzymatic activity. By contrast, the amount of m^5s^2U synthesized by TtuA increased with time (1, 2, and 24 hours after oxidation) regardless of whether excess $K_3[Fe(CN)_6]$ was removed from the solution or not (Fig. 3-4). When the enzymatic activity of TtuA before oxidation was normalized to 100%, it recovered 30%-50% in 1 hour, >50% in 2 hours, and 60%-80% in 24 hours (Table 3-2). Combining these results with EPR results, according to which [3Fe-4S] transformed into [4Fe-4S] with time (Fig. 3-3a, Table 1), it is clear that TtuA showed higher activity as the amount of [4Fe-4S] in TtuA increased.

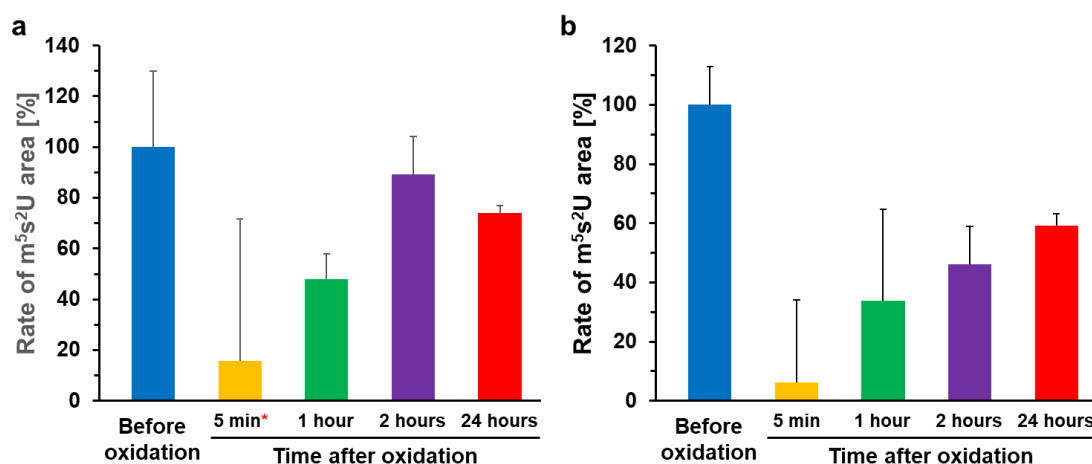


Fig. 3-4 Activity assay of [3Fe-4S]-TtuA and [4Fe-4S]-TtuA^[112].

Enzymatic activity of TtuA in the absence (a) and presence (b) of $K_3[Fe(CN)_6]$. The tRNA-thiolation activity of TtuA before oxidation was normalized to 100%. All data are presented with standard deviation (SD) values ($N = 3$, red asterisk (*): one time was measured from a different lot and scaling by the activity of [Before oxidation]). The quantification result of this assay is shown in Table 3-2.

Table 3-2. Quantification of the activity of [3Fe-4S]-TtuA and [4Fe-4S]-TtuA^[112].

Time after oxidation	Enzymatic activity without K ₃ [Fe(CN) ₆]	Enzymatic activity with K ₃ [Fe(CN) ₆]
Before oxidation	100% ± 30%	100% ± 13%
5 min	16 ± 56%*	6% ± 28%
1 hour	48% ± 10%	34% ± 31%
2 hours	89% ± 15%	46% ± 13%
24 hours	74% ± 3%	59% ± 4%

All data were measured three times and shown with standard deviation. Asterisk (*): one time was measured from a different lot and scaling by the activity of [Before oxidation].

3-3. The unique Fe in [4Fe-4S]-TtuA is required to bind the C-terminus of TtuB

To reveal the detailed function of the unique Fe in [4Fe-4S]-TtuA and evaluate why [3Fe-4S]-TtuA is inactive, I performed EPR spectroscopy of TtuA with TtuB-COSH and each type of Fe-S cluster. The EPR spectra of [3Fe-4S]-TtuA (freshly oxidized [4Fe-4S]-TtuA) showed a peak of [3Fe-4S]¹⁺ ($g \sim 2.01$), which supported that the unique Fe is absent (Fig. 3-5a). Then, I measured the EPR spectra of [3Fe-4S]-TtuA with TtuB-COSH and found that there is no significant difference in the spectra between [3Fe-4S]-TtuA alone and [3Fe-4S]-TtuA with TtuB-COSH (Fig. 3-5a). On the other hand, the addition of TtuB-COSH to [4Fe-4S]-TtuA changed the shape and intensity of the EPR spectra, showing the change in the redox potential when the thiocarboxylate group at the C-terminus of TtuB-COSH binds to [4Fe-4S]-TtuA (Fig. 3-5b)^[83].

Now, the structure of [4Fe-4S]-TtuA-TtuB complex shows that the C-terminus of TtuB directly coordinates with the unique Fe of [4Fe-4S]-TtuA (Fig. 1-17b)^[83]. Furthermore, the superposition of the structures of [4Fe-4S]-TtuA-TtuB and apo-TtuA-TtuB showed that the [4Fe-4S] cluster does not affect the interaction of TtuA and TtuB except for the C-terminus of TtuB-COSH (Fig. 3-6, Fig. 3-7). Therefore, the EPR spectra showed that [3Fe-4S]-TtuA cannot bind to the C-terminus of TtuB-COSH, indicating that the unique Fe of [4Fe-4S]-TtuA is necessary to coordinate with TtuB-COSH for transferring the sulfur atom to the substrate tRNA.

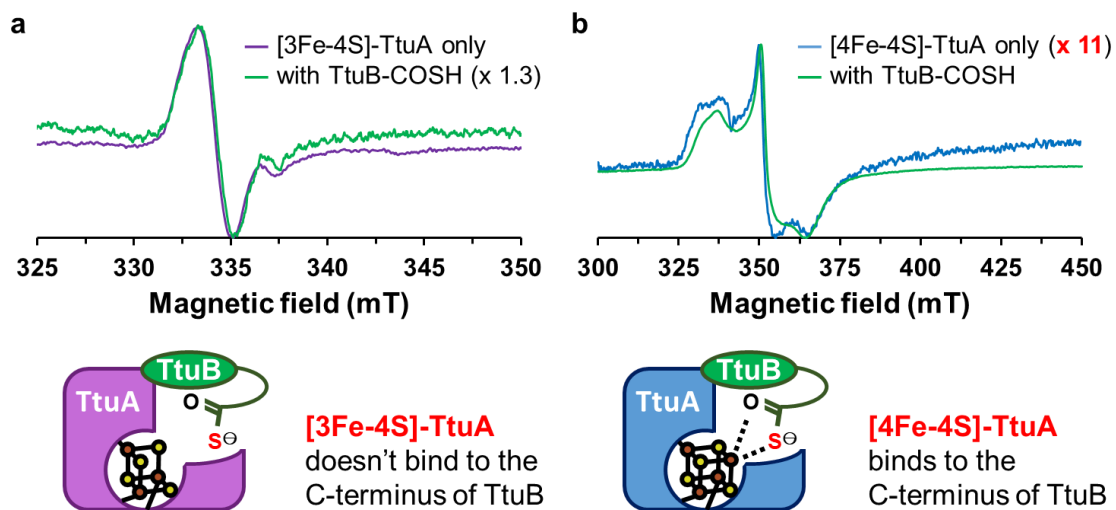


Fig. 3-5 Spectroscopic characterization of [3Fe-4S]-TtuA and [4Fe-4S]-TtuA with or without TtuB-COSH^[112].

(a) EPR spectra of [3Fe-4S]-TtuA in the absence (purple) and presence (green) of TtuB-COSH. (b) EPR spectra of [4Fe-4S]-TtuA in the absence (blue) and presence (green) of TtuB-COSH^[83]. The EPR spectra are normalized for comparison by indicated ratio.

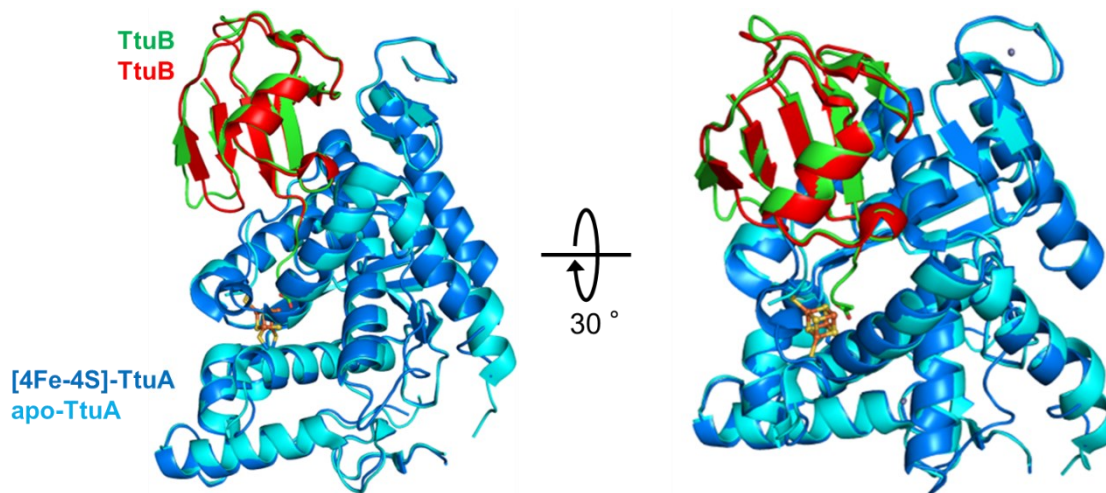


Fig. 3-6 Comparison of the overall structures of the [4Fe-4S]-TtuA-TtuB complex and the apo-TtuA-TtuB complex^[112].

TtuA and TtuB-COOH (WT) of the [4Fe-4S]-TtuA-TtuB complex (PDB ID: 5ZTB) are colored in blue and green, respectively^[83]. TtuA and TtuB-COOH (G65C) of the apo-TtuA-TtuB complex (PDB ID: 5GHA) are colored in cyan and red, respectively^[72]. α RMSD value is 0.483 Å, which was calculated by the “align” command of PyMOL ver1.7. The GG motif at the C-terminus of TtuB is disordered when [4Fe-4S] is absent.

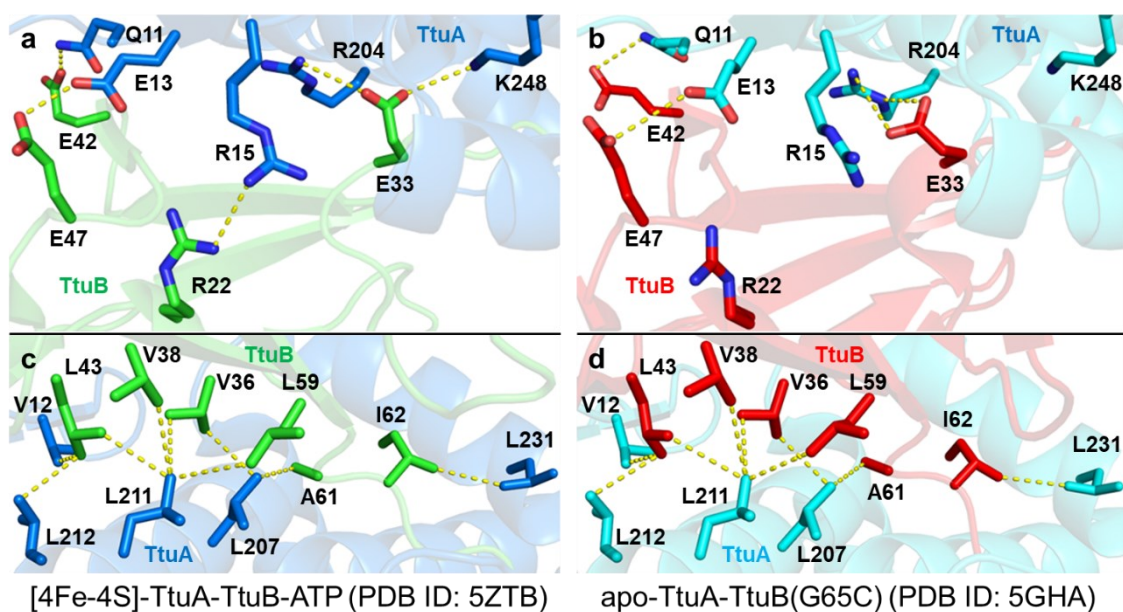


Fig. 3-7 Comparison of the binding site of the [4Fe-4S]-TtuA-TtuB complex and the apo-TtuA-TtuB complex^[112].

(a) The binding site I and (c) site II of the [4Fe-4S]-TtuA-TtuB complex (PDB ID: 5ZTB). TtuA and TtuB-COOH (WT) are colored in blue and green, respectively. (b) The binding site I and (d) site II of the apo-TtuA-TtuB complex (PDB ID: 5GHA). TtuA and TtuB-COOH (G65C) are colored in cyan and red, respectively. The yellow dashed lines indicate electrostatic or hydrophobic interactions.

3-4. TtuB does not release sulfur before tRNA activation

To understand the catalytic mechanism of [4Fe-4S]-TtuA, I analyzed the conditions under which TtuB releases sulfur from its C-terminus by APM-PAGE. I confirmed that the C-terminus of TtuB was thiocarboxylated (Fig. 3-8a, lane 1). Whereas TtuB did not release sulfur in the presence of [4Fe-4S]-TtuA and either tRNA or ATP (Fig. 3-8a lanes 1 and 2), TtuB was desulfurized when [4Fe-4S]-TtuA, tRNA, and ATP were added together and incubated for 10 min or longer (Fig. 3-8a, lanes 5 and 6). I also detected m^5s^2U in conditions that TtuB released sulfur by HPLC (Fig. 3-9b, c). These observations demonstrated that the binding of TtuB-COSH to TtuA is not sufficient for desulfurization, but TtuA, TtuB-COSH, substrate tRNA, and ATP are necessary.

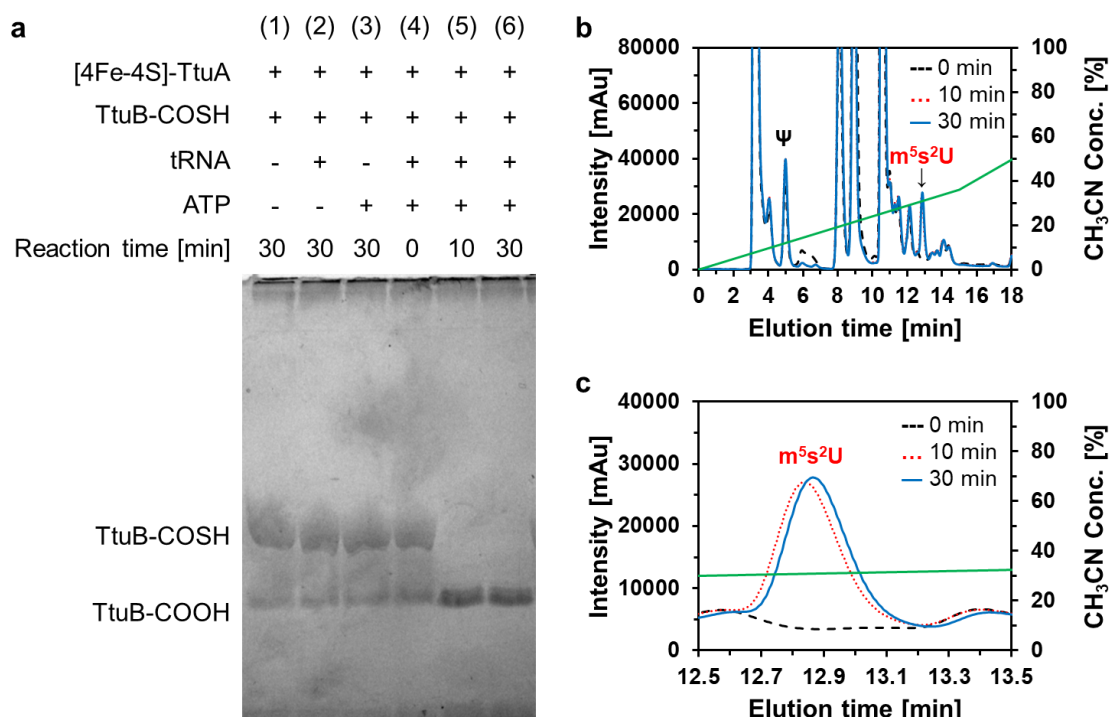


Fig. 3-8 Tracing sulfur atom from C-terminus of TtuB^[83].

(a) APM-gel for desulfurization assay. (b) Overall and (c) close-up view of HPLC spectra of desulfurization assay. The HPLC samples 0 min, 10 min, and 30 min were prepared in the same conditions in Fig. 3-8a lanes 4, 5, and 6, respectively. The reference peak of pseudouridine (Ψ) and the product peak of 5-methyl-2-thiouridine (m^5s^2U) are indicated. The green lines indicate the concentration of acetonitrile (CH_3CN).

3-5. Identification of critical residues of TtuA in tRNA thiolation

To identify the critical residues of TtuA for tRNA thiolation, I prepared 10 mutants of TtuA based on the structure of the [4Fe-4S]-TtuA-TtuB-ATP complex (Fig. 3-9a). These residues are conserved and located around the ATP-binding site ($S_{55}GGXDS_{60}$ motif) or the unique Fe of [4Fe-4S]-TtuA. The HPLC spectra showed that the enzymatic activities of S55A, D59A, K137A, and D161A were dramatically decreased (Fig. 3-9b). Furthermore, I evaluated the enzymatic activities of these four mutants in the presence of 10 mM Na_2S instead of TtuB-COSH to identify the catalytic residue of TtuA for desulfurization. Interestingly, only the K137A mutant recovered the activity, showing that Lys137 is essential for the sulfur transition from TtuB to tRNA (Fig. 3-9c). Notably, the

activity of Lys137 was not completely recovered even when Na₂S was supplied, suggesting that Lys137 might be involved in activation of tRNA. This conclusion is also supported by the result of APM-PAGE (Fig. 3-10a).

Interestingly, some organisms have TtuA or Ncs6 with Arg137 instead of Lys137 (the residue number is from *Tth*TtuA) (Fig. 1-28a) and our results of HPLC and AMP-PAGE also showed an enzymatic activity in K137R (Fig. 3-9b, Fig. 3-10). On the other hand, the enzymatic activity of S55A, D59A, and D161A did not recover in the presence of Na₂S. These residues are known to be responsible for ATP binding and hydrolysis^[119,120]. Therefore, I concluded that the positively charged residue at position 137 are critical for sulfur transition, and the other three residues activates tRNA (Fig. 3-11).

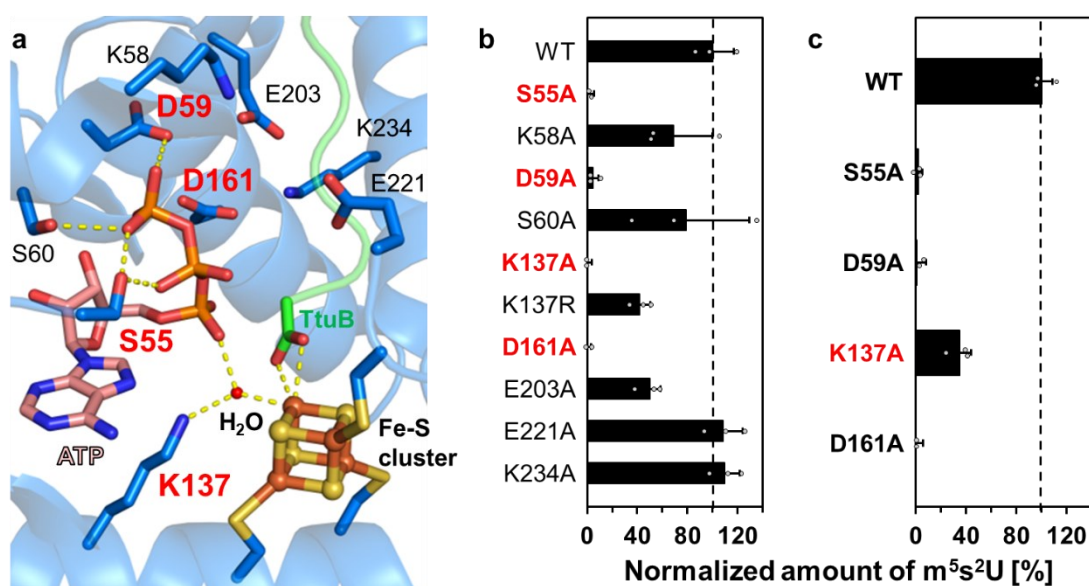


Fig. 3-9 Identification of critical residues of TtuA by HPLC^[83].

(a) Close-up view of the catalytic site of the [4Fe-4S]-TtuA-TtuB-ATP complex (PDB ID: 5ZTB). The dashed yellow lines indicate hydrogen bonds or coordination bonds. Mutational analysis of TtuA in the presence of (b) 20 μM TtuB-COSH and (c) 10 mM Na₂S. All activity assays were examined three times and these results were presented with data plots and SD values.

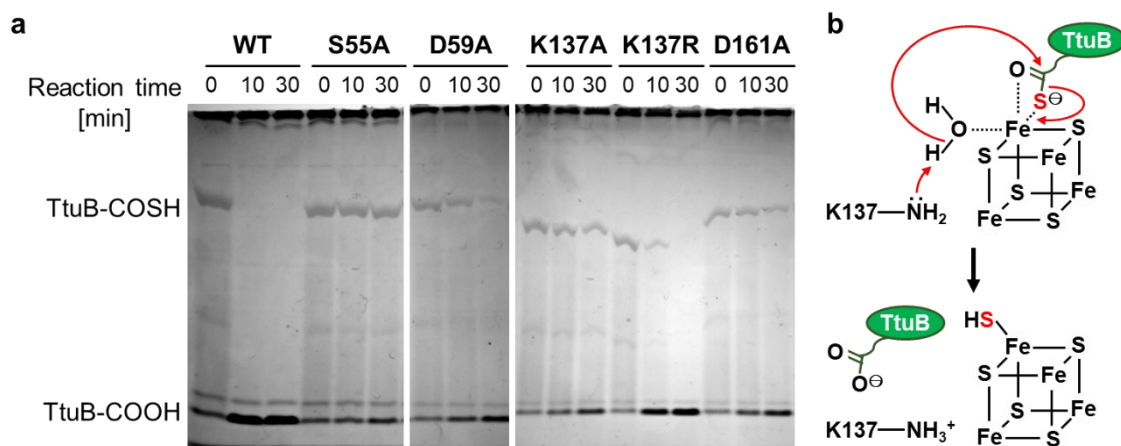


Fig. 3-10 Identification of critical residues of TtuA by APM-PAGE^[83].

(a) Desulfurization assay of TtuB catalyzed by TtuA mutants. (b) Proposed mechanism of desulfurization of TtuB catalyzed by Lys137 of TtuA.

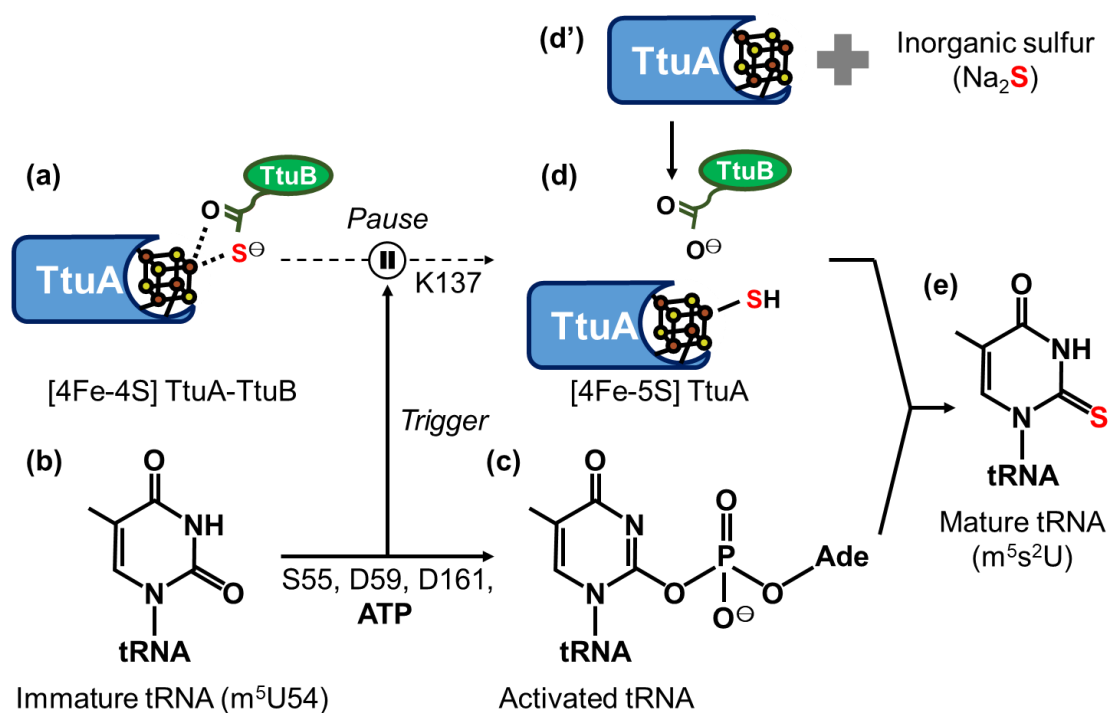


Fig. 3-11 Proposed scheme of 5-methyl-2-thiouridine (m^5s^2U54) biosynthesis^[83].

(a) Formation of the [4Fe-4S]-TtuA-TtuB complex. TtuB cannot release sulfur from its C-terminus until the activation of tRNA. (b) 5-methyl-uridine (m^5U54) is adenylated using ATP involving Ser55, Asp59, and Asp161. (c) Adenylated tRNA triggers sulfur transition from TtuB to TtuA. (d) [4Fe-5S]-TtuA intermediate is formed by Lys137 or (d') addition of inorganic sulfur source. (e) tRNA thiolation is completed.

Chapter 4. Discussion

4-1. [3Fe-4S]-TtuA stores Fe sources for the reconstitution of [4Fe-4S]-TtuA

Fe-S clusters are widely used cofactors and are responsible for various biological roles. Thus, it is necessary to identify the type of Fe-S cluster in the enzyme to understand the catalytic mechanism. To clarify whether TtuA uses [4Fe-4S] only or both [4Fe-4S] and [3Fe-4S], I performed time-resolved EPR spectroscopy under strictly anaerobic conditions. The results showed that [3Fe-4S]-TtuA was unstable and spontaneously transformed into [4Fe-4S]-TtuA within one hour (Fig. 3-3). Furthermore, interaction analysis by EPR spectroscopy demonstrated that [3Fe-4S]-TtuA cannot bind to the C-terminus of TtuB-COSH (Fig. 3-5). The enzymatic activity of [3Fe-4S]-TtuA (oxidized [4Fe-4S]-TtuA) recovered with an increase in [4Fe-4S]-TtuA, but did not completely recover (Fig. 3-4). These observations indicated that the reaction solution contained apo-TtuA, which was generated by the degradation of [3Fe-4S] (Fig. 4-1). Taking our results together, I concluded that only [4Fe-4S]-TtuA has enzymatic activity, and the unique Fe is essential as a binding site for TtuB-COSH for the sulfur transition to tRNA.

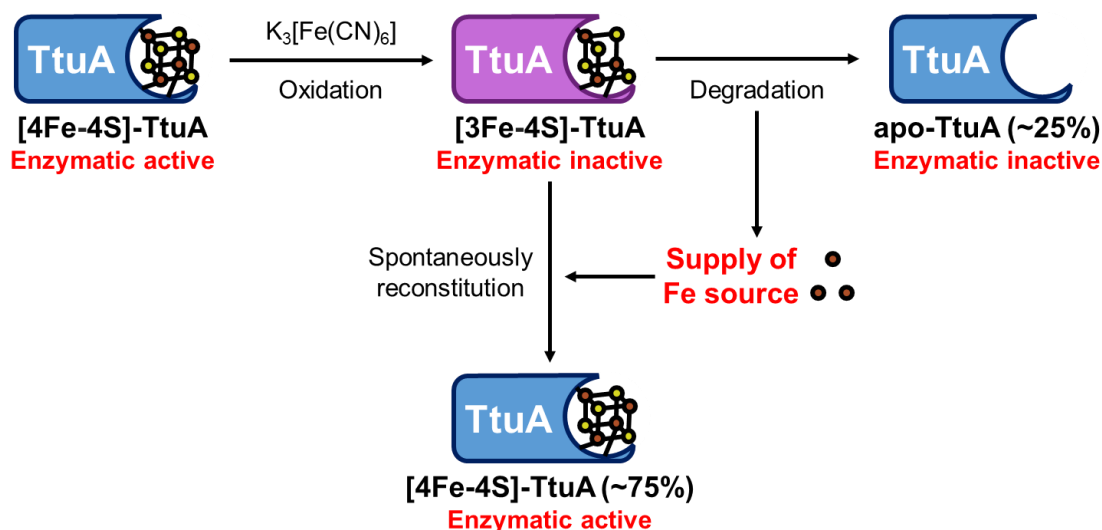


Fig. 4-1 Schematic diagram of the proposed changes of the Fe-S cluster in TtuA^[112]. One [3Fe-4S]-TtuA molecule provides three free Fe^{3+} ions with inactive apo-TtuA molecules to form active [4Fe-4S]-TtuA. The parentheses indicate the maximum ratio of apo-TtuA and [4Fe-4S]-TtuA.

Considering that TtuA catalyzes m^5s^2U biosynthesis which is essential for thermophiles to survive under high-temperature environments^[18], thermophiles eliminate inactive [3Fe-4S]-TtuA and reconstitute active [4Fe-4S]-TtuA. One of the biological roles of [3Fe-4S] is iron and sulfur source in iron regulatory protein 1 (IRP1)^[121]. Thus, the transformation of [3Fe-4S]-TtuA into [4Fe-4S]-TtuA maybe occur *in vivo*, but we need further study to demonstrate the possibility.

4-2. Time-course analyses are essential to identify the exact type of Fe-S clusters

Fe-S clusters are sensitive to oxidation and readily decayed. Their instability sometimes caused incorrect structural determination of Fe-S clusters even with the smallest amount of oxygen contaminant during the experiments (Fig. 1-20). Our findings demonstrated that the Fe-S cluster in TtuA is highly sensitive to redox levels and [3Fe-4S] clusters quickly and spontaneously transform into [4Fe-4S] clusters within one hour, even under strictly anaerobic conditions. Unexpectedly, [3Fe-4S]-TtuA is more unstable than [4Fe-4S]-TtuA even though all of the Fe atoms of [3Fe-4S] are coordinated with cysteines. Hence, misidentification of [3Fe-4S] and [4Fe-4S] easily lead to an improper proposal of the catalytic mechanism of the Fe-S enzyme without the correlation analysis between the structure of Fe-S clusters and the enzymatic activity in time-course under strictly anaerobic conditions.

Many proteins that employ the Fe-S cluster are overlooked as apo-type proteins because Fe-S clusters are degraded easily due to their oxygen-sensitivity. Today, 779 of [4Fe-4S]-binding proteins and 42 of [3Fe-4S]-binding proteins are deposited in the protein database (UniProt) from 21 organisms including *E. coli*, *S. cerevisiae*, and *H. sapiens*^[122]. However, some of them are annotated based on only sequence similarity, which means that we need experimental evidence to demonstrate the type of Fe-S clusters. Considering the recent advances in our understanding of expanding role of Fe-S clusters, we can expect the exponential discovery of new Fe-S proteins in the future. Therefore, we have to perform spectroscopic, structural, and biochemical experiments on Fe-S proteins under strictly anaerobic conditions to minimize the risk of proposing an incorrect catalytic mechanism.

4-3. Sequence similarity of critical residues of tRNA-thiolation enzymes

I found that only [4Fe-4S]-TtuA is the enzymatically active form. Interestingly, some groups have reported that the TtuA/Ncs6 family members (TtuA, Ncs6, and TtcA6) bind to [4Fe-4S]^[67,71,72,76,83], whereas others have indicated that the active form of Ncs6 and ThiI is [3Fe-4S]^[69]. Notably, the mechanism of sulfur transition catalyzed by TtuA and Ncs6 is believed to be similar^[107], because their sulfur relay systems involving ubiquitin-like sulfur donor proteins are similar (Fig. 1-29), despite the different positions of tRNA thiolation. To evaluate whether [4Fe-4S] is the unique ligand of TtuA or common among tRNA-thiolation enzymes, I compared amino acids around the active site of the TtuA/Ncs6 family members, MnmA, and ThiI. Although MnmA and ThiI do not belong to the TtuA/Ncs6 family (Fig. 1-13, Table 1-1), recent studies reported that *EcoMnmA* contains [4Fe-4S] and *MmaThiI* has [3Fe-4S], respectively^[69,77].

A previous study has shown that Asp59, Cys130, Cys133, Asp161, and Cys222 in apo-*TthTtuA* are critical in tRNA thiolation^[82], but the mutational analysis was performed in aerobic conditions because the presence of [4Fe-4S] in TtuA has not reported at that time. Our group previously reported that Cys130, Cys133, and Cys222 are [4Fe-4S]-binding residues^[72], and I identified that Ser55, Asp59, Lys137, and Asp161 are essential residues in strictly anaerobic conditions (Fig. 3-9)^[83]. Ser55 and Asp59 are located at the PP-loop motif (S₅₅GGXDS₆₀) which are responsible for ATP binding and hydrolysis^[119,120]. Lys137 catalyzes sulfur transition from TtuB to tRNA, stabilizing [4Fe-5S]-TtuA intermediate^[76], and potentially involved in tRNA adenylation^[83]. Asp161 is also predicted as a critical residue for ATP binding and hydrolysis from the structural insight of the PPase family although Asp161 is 5-Å away from ATP^[83,120].

Interestingly, these residues are highly conserved in not only the TtuA/Ncs6 family, but also in MnmA and ThiI with a few exceptions, such as D190 in the PP-loop of *MmaThiI* and D99/D101 coordinated to [4Fe-4S] in *EcoMnmA/BsuMnmA* (Fig. 4-2, Fig. 4-3). Notably, one of the [4Fe-4S]-binding residues is aspartic acid instead of cysteine in some MnmA similar to ferredoxin, Fnr, IscA, and BchB^[77,123,124].



Fig. 4-2 Sequence alignment of tRNA-thiolation enzymes.

The secondary structures of *TthTtuA* (PDB ID: 5B4F) are shown at the top. Completely conserved residues are highlighted in red and highly conserved residues are highlighted in yellow, respectively. Key residues in the ATP-binding motif (PP-loop, a green line); the three cysteine residues bound to the Fe-S cluster (red asterisks); the cysteine or histidine residues involved in the tRNA-binding motif (Zn finger, blue asterisks); and key residues of *TthTtuA* (magenta triangles)^[82,83]. The alignment was performed using Clustal Omega^[125] and ESPrint 3.0^[126] was used for illustration. Note that V345 of *TmaThiI* aligns with C222 of *TthTtuA* because their sequence identity is only 13.9%; however, C344 is the catalytic residue of *TmaThiI*^[105]. The abbreviations of protein names are the same as that in Fig. 1-13.

Enzymes	Organisms	Identified critical residues															
TtuA	<i>T. thermophilus</i>	S55	G56	G57	K58	D59	S60	...	C130*	...	C133*	...	K137	...	D161	...	C222*
TtuA	<i>A. aeolicus</i>	S56	G57	G58	K59	D60	S61	...	C130*	...	C133*	...	K137	...	D161	...	C222*
TtuA	<i>T. maritima</i>	S55	G56	G57	K58	D59	S60	...	C131*	...	C134*	...	R138	...	D162	...	C224*
TtuA	<i>P. horikoshii</i>	S55	G56	G57	K58	D59	S60	...	C128*	...	C131*	...	K135	...	D159	...	C220*
TtuA	<i>M. jannaschii</i>	S59	G60	G61	K62	D63	G64	...	C130*	...	C133*	...	K137	...	D161	...	C222*
TtcA	<i>E. coli</i>	S47	G48	G49	K50	D51	S62	...	C122*	...	C125*	...	R129	...	D153	...	C213*
TtcA	<i>P. putida</i>	S40	G41	G42	K43	D44	S45	...	C115*	...	C118*	...	R122	...	D146	...	C206*
TtcA	<i>S. oneidensis</i>	S45	S46	G47	K48	D49	S50	...	C120*	...	C123*	...	R127	...	D151	...	C211*
TtcA	<i>S. typhimurium</i>	S47	G48	G49	K50	D51	S52	...	C122*	...	C125*	...	R129	...	D153	...	C213*
Ncs6	<i>H. sapiens</i>	S59	G60	G61	K62	D63	S64	...	C144*	...	C147*	...	R151	...	D175	...	C237*
Ncs6	<i>S. cerevisiae</i>	S75	G76	G77	K78	D79	S80	...	C157*	...	C160*	...	R164	...	D188	...	C250*
Ncs6	<i>S. pombe</i>	S59	G60	G61	K62	D63	S64	...	C142*	...	C145*	...	R149	...	D173	...	C235*
Ncs6	<i>M. maripaludis</i>	S57	G58	G59	K60	D61	S62	...	C142*	...	C145*	...	R149	...	D173	...	C233*
Ncs6	<i>H. volcanii</i>	S61	G62	G63	K64	D65	S66	...	C143*	...	C146*	...	R150	...	D174	...	C242*
MnmA	<i>E. coli</i>	S13	G14	G15	V16	D17	S18	...	D99*	...	C102*	...	I106	...	R132	...	C199*
MnmA	<i>B. subtilis</i>	S15	G16	G17	V18	D19	S20	...	D101*	...	C104*	...	I108	...	V133	...	C200*
MnmA	<i>T. thermophilus</i>	S9	G10	G11	V12	D13	S14	...	C105*	...	C108*	...	V112	...	K137	...	C200*
ThiI	<i>E. coli</i>	S185	G186	G187	F188	D189	S190	...	G261	...	L264	...	M268	...	Q292	...	C344
ThiI	<i>M. maripaludis</i>	S189	D190	G191	I192	D193	S194	...	C265*	...	C268*	...	M272	...	Q296	...	C348*
ThiI	<i>T. maritima</i>	S182	G183	G184	I185	D186	S187	...	L260	...	Y263	...	M267	...	Q291	...	C344

Fig. 4-3 Sequence alignment of critical residues of TtuA for tRNA-thiolation^[112].

The abbreviations of source organisms are the same as that in Fig. 1-13. Residues with asterisks are coordinated to Fe-S clusters. The conserved residues are indicated in red.

4-4. [4Fe-4S] is an active form in tRNA-thiolation enzymes

Until now, only three 3D structures of *Tth*TtuA, *Pho*TtuA, and *Mma*Ncs6 has been deposited in the Protein Data Bank (PDB) among tRNA-thiolation enzymes complexed with Fe-S clusters (all of them are [4Fe-4S]). Furthermore, [4Fe-4S]-*Tth*TtuA is the only reported complex structure with sulfur donor protein, and no complex structures with tRNA has been reported among the TtuA/Ncs6 family. These limited structural information makes it difficult to discuss whether [4Fe-4S] is generally active form in tRNA thiolation. Recently, AlphaFold version 2 (AF2) enabled us to predict 3D structure of *Eco*TtcA and *Mma*ThiI, and compare the positions of critical residues Ser55, Asp59,

Cys130, Cys133, Lys137, Asp161, and Cys222 in tRNA-thiolation enzymes^[115]. The superimpositions showed that all key residues are located at similar positions with exceptions of Asp161 and Cys222 of TtuA (Fig. 4-4). Considering conservation of critical residues (Fig. 4-3) and structural similarity of the active site, [4Fe-4S] binding is likely shared by tRNA thiolation enzymes. Notably, ThiI from some organisms do not have [4Fe-4S]-binding cysteines at position 130 and 133 (Fig. 4-5), indicating that these conventual ThiI may catalyzes tRNA thiolation using persulfide as previously proposed (Fig. 1-26)^[98].

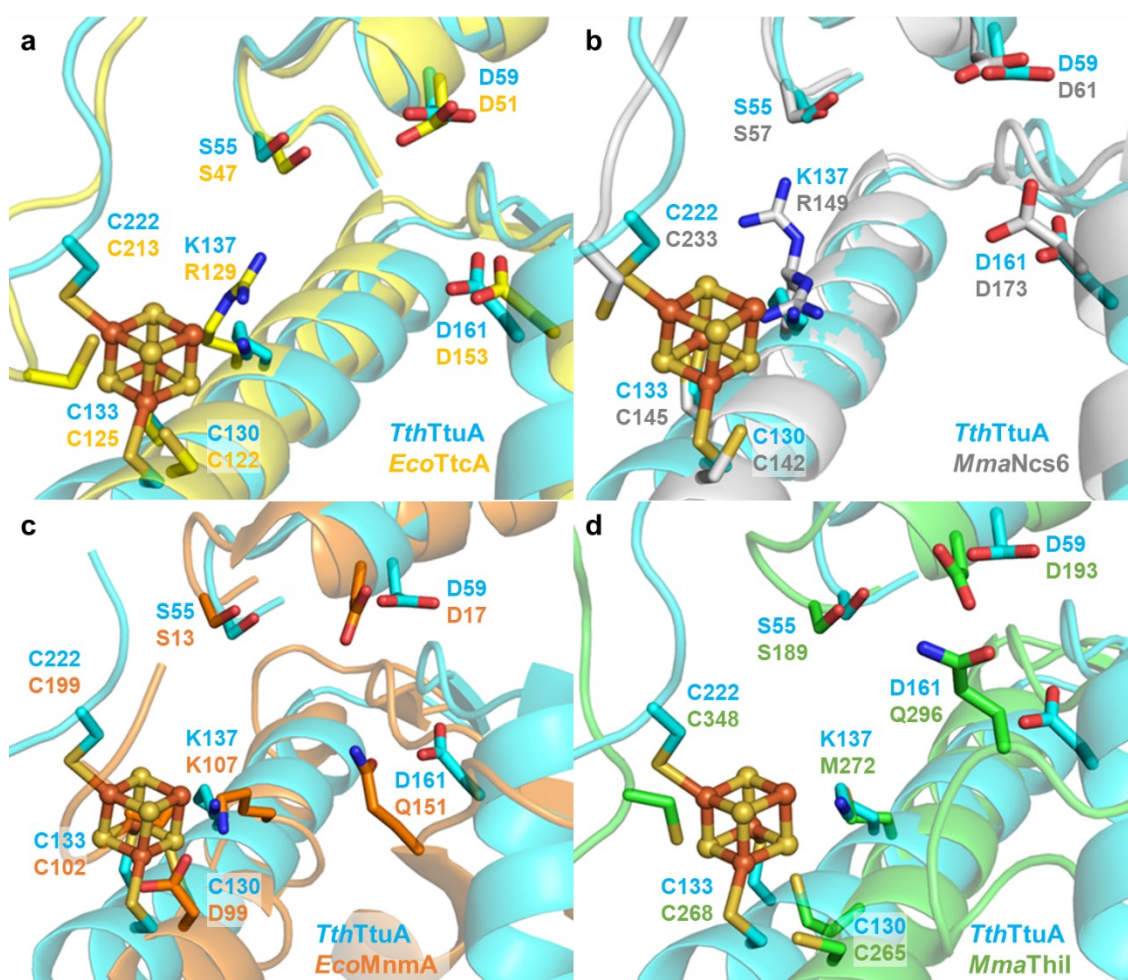


Fig. 4-4 Structural alignment of tRNA-thiolation enzymes^[112].

Superposition of *TthTtuA* (PDB ID: 5B4F) with the (a) *EcoTtcA* AF2 model, (b) *MmaNcs6* (PDB ID: 6SCY), (c) *EcoMnmA* (PDB ID: 2DEU), and (d) *MmaThiI* AF2 model. The AF2 models were superposed in the manner described in the Materials and Methods section.

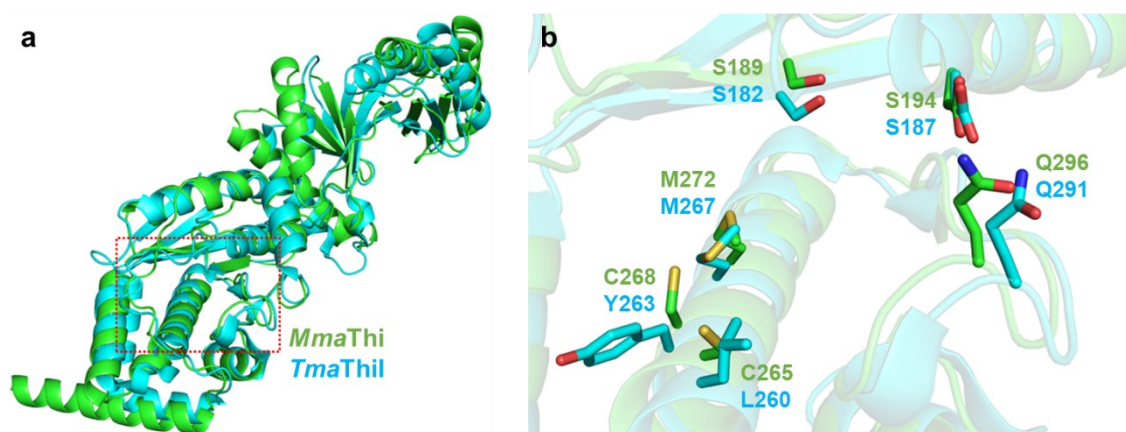


Fig. 4-5 Structural comparison of *MmaThiI* and *TmaThiI*.

(a) Superposition of overall structures of the *MmaThiI* AF2 model with *TmaThiI* (PDB ID: 4KR6). Their C α RMSD is 2.054 Å. The catalytic site is marked by the red dotted frame. (b) Close-up view of their catalytic center. The coordination of the critical residues matches well, but *TmaThiI* cannot bind to Fe-S clusters because Cys265 and Cys268 of *MmaThiI* are replaced by Y263 and M267, respectively.

4-5. [4Fe-4S] cluster is a potential scaffold of tRNA adenylation

In this study, our result of APM-PAGE showed that the binding of TtuB to TtuA is not sufficient for desulfurization, but TtuA, TtuB, tRNA, and ATP are necessary (Fig. 3-8). This observation indicated that sulfur transition from TtuA to tRNA occurs after activation of tRNA using ATP. Furthermore, I elucidated the sulfur transition mechanism from TtuB to tRNA catalyzed by Lys137 of [4Fe-4S]-TtuA. However, the catalytic mechanism of tRNA adenylation is unknown.

Previously, the structure of apo-*EcoMnmA* complexed with adenylylated tRNA (AdetRNA) has been determined (Fig. 1-23)^[95], but the [4Fe-4S] cluster was absent in *EcoMnmA*. Furthermore, our group reported the [4Fe-4S]-TtuA-TtuB-AMPPNP-tRNA complex model based on the structure of the TilS-tRNA complex (Fig. 4-6)^[72]. Although the C α RMSD between *TthTtuA* and TilS is 1.747 Å, we could not show the accurate model because the target base is position 54 of tRNA in TtuA whereas position 34 of tRNA in TilS. Therefore, the detailed catalytic mechanism of tRNA adenylation involving [4Fe-4S] clusters is still unknown.

Recently, the structure of [4Fe-4S]-*MmaNcs6* has been determined and the C α RMSD between *MmaNcs6* and TilS is 1.774 Å. The structural similarity allowed to build [4Fe-4S]-*MmaNcs6*-tRNA complex model (Fig. 4-7a). In many archaea including *M. maripaludis*, *urm1* gene is missing and assumed that SAMP2 (archaeal Urm1 homolog) is sulfur donor^[127]. On the other hand, typical eukaryotes have Urm1 as sulfur donor such as human and yeast, which enabled to predict the structure of the *HsaNcs6*-Urm1 complex using AF2 and build the [4Fe-4S]-*HsaNcs6*-Urm1-tRNA complex model (Fig. 4-7b).

These enzyme-tRNA complex models show that the unique Fe interacts with position 2 of the substrate uridine, which indicate that [4Fe-4S] cluster is not only a scaffold of sulfur transition, but also that of tRNA adenylation in tRNA-thiolation enzymes. Furthermore, it is suggested that Asp161 is a critical residue for tRNA adenylation similar to that of Asp131 in TilS^[116,117].

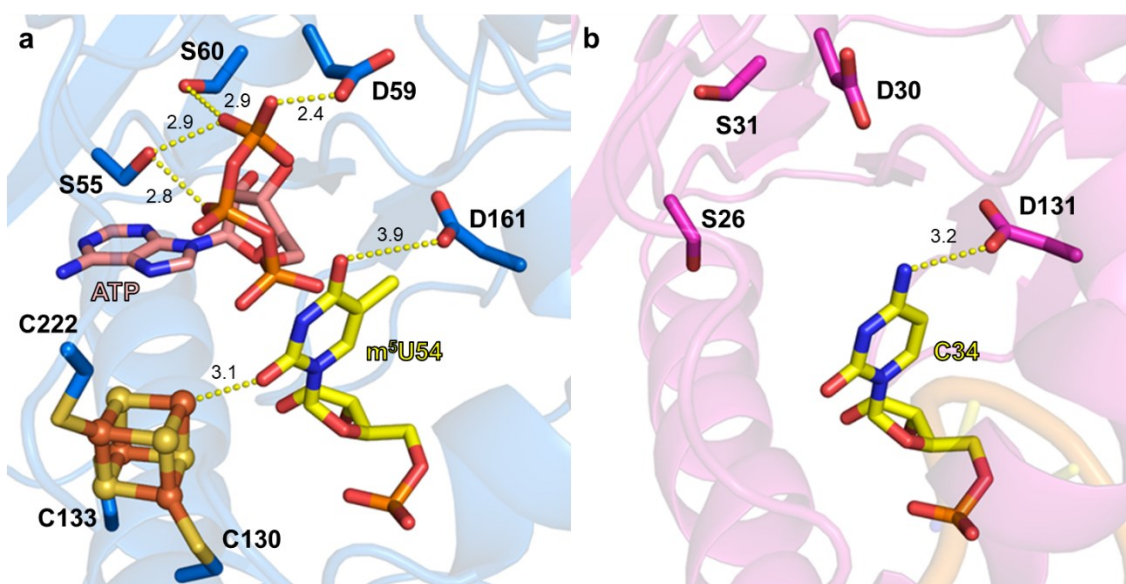


Fig. 4-6 Close-up view of the catalytic center of the TtuA model and TilS complexed with substrate tRNA.

(a) The [4Fe-4S]-TtuA-m⁵U54 complex model. This model was built by superposition of the [4Fe-4S]-TtuA-TtuB-ATP complex (PDB ID: 5ZTB), the TilS-tRNA complex (PDB ID: 3A2K), and *ScetRNA*^{Phe} (PDB ID: 1EHZ). TtuB and overall tRNA are not show. The substrate uridine is methylated similar to T₅₄ΨC₅₆ loop of *ScetRNA*^{Phe}. (b) The TilS-tRNA complex. Note that TilS is ATP-dependent, but [4Fe-4S]-independent. The yellow dotted lines indicate hydrogen bonds or coordination bonds with distance in angstrom units.

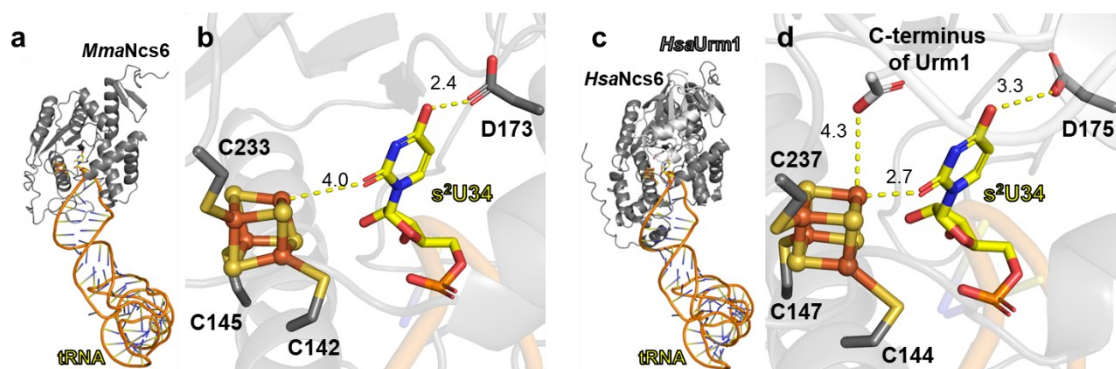


Fig. 4-7 The structure of [4Fe-4S]-Ncs6-tRNA complex model.

(a) Overall and (b) close-up view of the [4Fe-4S]-*MmaNcs6*-tRNA complex model, which was built by the TilS-tRNA complex (PDB ID: 3A2K) using the “Super” command in PyMOL ver.1.7. (c) Overall and (d) close-up view of the [4Fe-4S]-*HsaNcs6*-*HsaUrm1*-tRNA complex model, which was built by superposition of the [4Fe-4S]-TtuA-tRNA complex model using the “Super” command in PyMOL ver.1.7. The yellow dotted lines indicate hydrogen bonds or coordination bonds with distance in angstrom units. Note that the substrate nucleotide of TilS is cytidine and the substrate of Ncs6 is mcm⁵U, but I represented that of Ncs6 as uridine for clearness.

4-6. Proposed catalytic mechanism of tRNA-thiolation in the TtuA/Ncs6 family

Taking all results together, I proposed the detailed mechanism of tRNA thiolation catalyzed by [4Fe-4S]-binding enzymes. Firstly, tRNA is activated by using Ser55, Asp59, and Asp161 in the presence of tRNA and ATP (Fig. 4-8a). Secondly, TtuA transfers sulfur from the C-terminus of TtuB to the unique Fe using catalytic residue Lys137, which forms the [4Fe-5S]-TtuA intermediate (Fig. 4-8b). Thirdly, [4Fe-5S]-TtuA transfers sulfur to the substrate tRNA (Fig. 4-8c). Finally, tRNA thiolation is completed and AMP is released from AdetRNA (Fig. 4-8d). Owing to the [4Fe-4S]-TtuA-tRNA complex model, I can describe this schematic diagram in the level of organic chemistry (Fig. 4-9). I would like to emphasize that this study implied that not only TtuA but also tRNA-thiolation enzymes is likely to catalyze tRNA thiolation with [4Fe-4S].

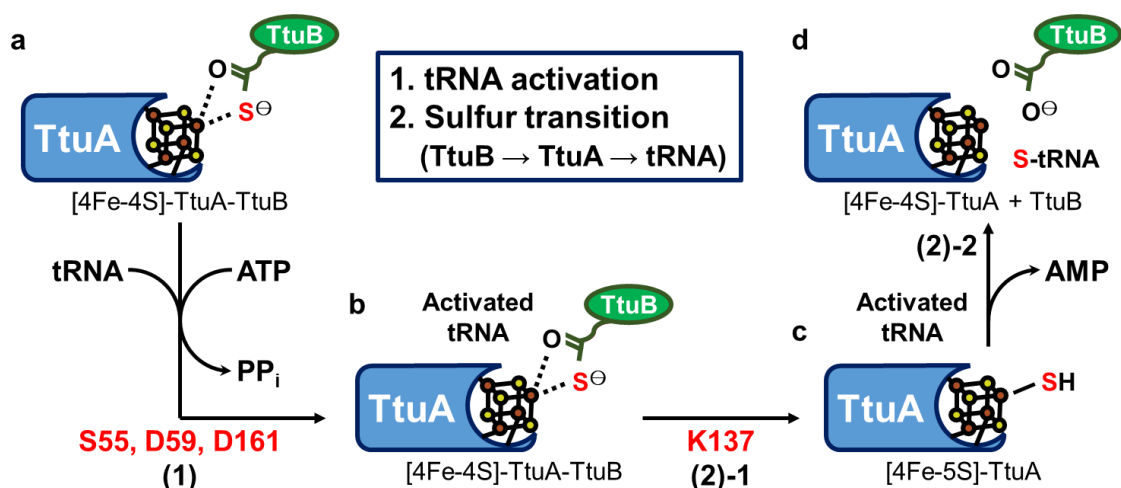


Fig. 4-8 Schematic diagram of the molecular mechanism of tRNA thiolation.

(a) The [4Fe-4S]-TtuA-TtuB complex catalyzes tRNA adenylation using ATP with critical residues Ser55, Asp59, and Asp161. (b) Lys137 of [4Fe-4S]-TtuA catalyzes sulfur transition from the C-terminus of TtuB to the unique Fe. (c) The [4Fe-5S]-TtuA intermediate transfers sulfur to AdetRNA. (d) tRNA thiolation is completed.

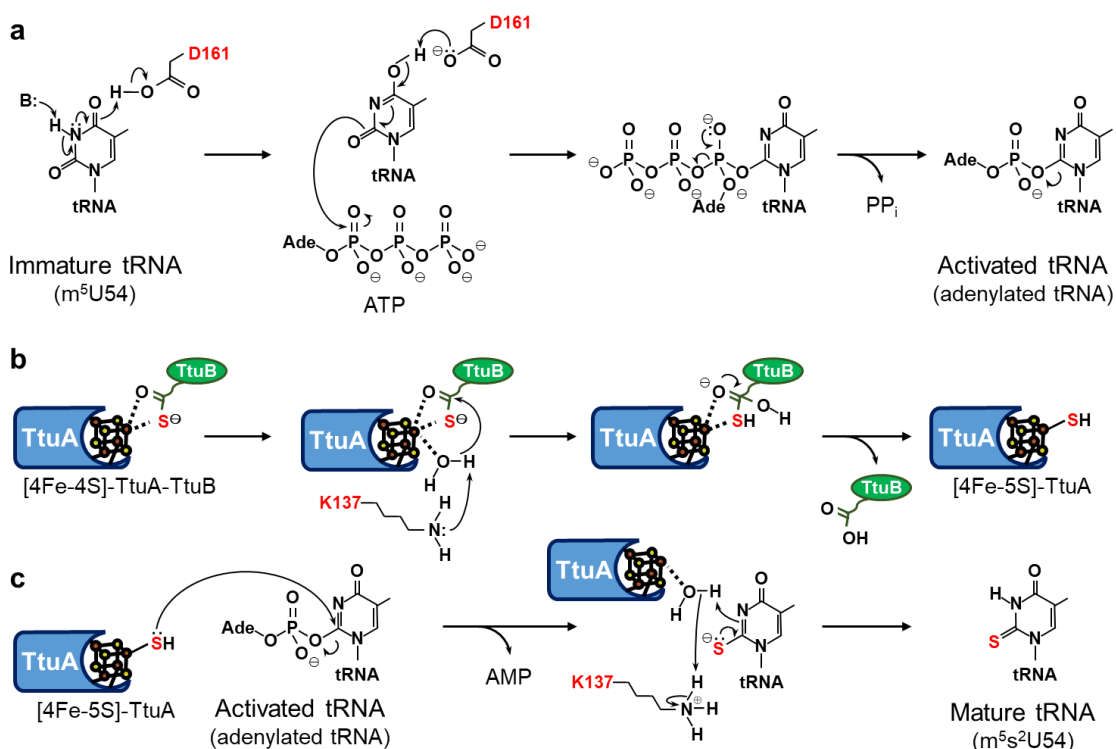


Fig. 4-9 Proposed mechanism of tRNA thiolation catalyzed by [4Fe-4S]-TtuA.

(a) tRNA adenylation catalyzed by Asp161. (b) Sulfur transition from TtuB to adenylylated tRNA catalyzed by Lys137. (c) Sulfur transition from [4Fe-5S]-TtuA to adenylylated tRNA.

Chapter 5. Perspective

In this study, I built the [4Fe-4S]-TtuA-m⁵U54 complex and the [4Fe-4S]-Ncs6-tRNA complex models, showing that the [4Fe-4S] cluster could potentially function as a scaffold of tRNA adenylation. However, the detailed catalytic mechanism of tRNA activation is still unclear. To address this problem, we need to determine the structure of the tRNA-thiolation enzymes with [4Fe-4S] and tRNA using X-ray crystallography, and perform mutation assays involving tRNA activation. This structural determination would be challenging because our preliminary experiments suggested that the binding strength of TtuA to tRNA is too weak to form the TtuA-tRNA complex even under anaerobic conditions (Data not shown).

In tRNA thiolation, it has been believed that the PP-loop motif is responsible for ATP binding/hydrolysis and releases AMP and Pyrophosphate (PP_i) via the AdetRNA intermediate^[72,76]. This tRNA activation mechanism is based on the crystal structure of the MnmA-AdetRNA complex (Fig. 1-23) and bioinformatics analysis that ThiI and TilS have conserved PP-loop motifs^[128,129]. In ThiI, furthermore, direct evidence for the formation of an AdetRNA intermediate and the release of AMP was reported using α-³²P-labeled ATP to quantify adenylated tRNAs^[130], and a preliminary observation showed that AMP is produced during s⁴U formation^[98]. However, I could not find any previous studies that analyze the release of ADP, PP_i, and inorganic phosphate (P_i). Therefore, we cannot exclude the possibility that ATP releases AMP and two P_i molecules via the ADP and AdetRNA intermediates.

Recently, we have determined the crystal structures of the [4Fe-4S]-TtuA-ATP complex and the [4Fe-4S]-TtuA-ADP complex (unpublished). We also found about 30% of the m⁵s²U54 synthesis activity of TtuA even when ADP was used instead of ATP (unpublished). Considering these observations and the finding that TtuB does not release sulfur until tRNA activation (Fig. 3-11), two major possibilities still remain in the molecular mechanism of tRNA activation: TtuA catalyzes ATP hydrolysis and releases AMP and PP_i (Fig. 5-1, hypothesis 1); ATP is trimmed to ADP and then releases AMP and two P_i molecules (Fig. 5-1, hypotheses 2 and 3).

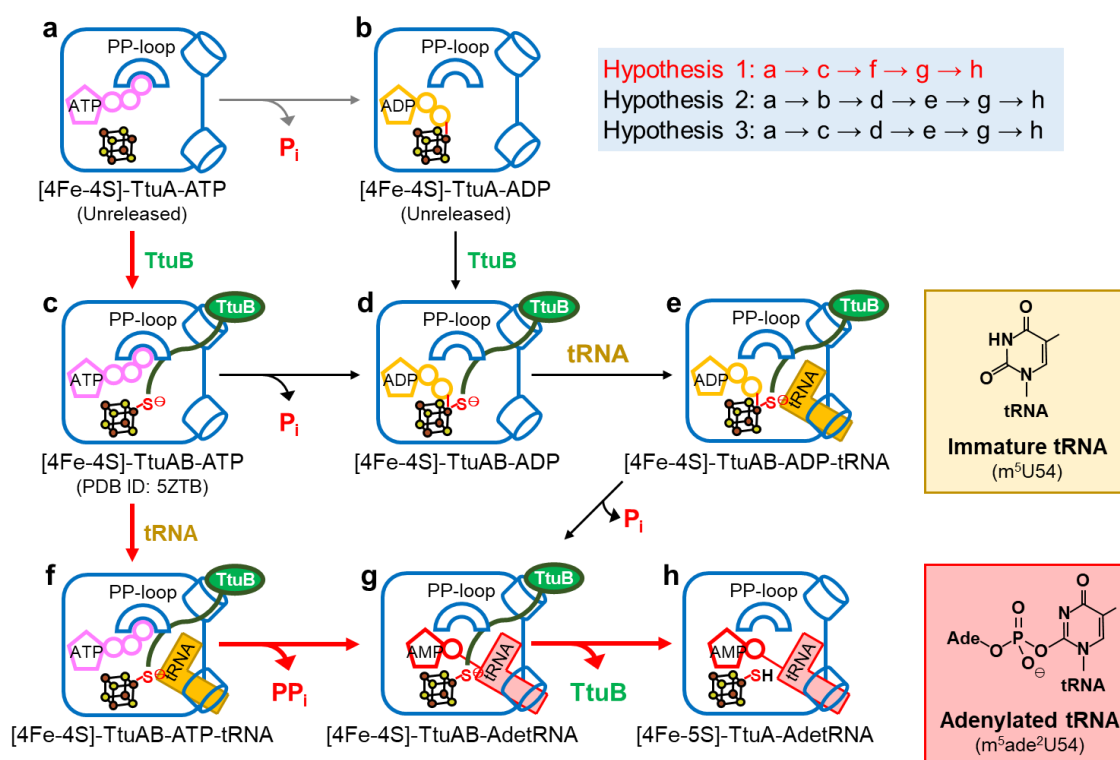


Fig. 5-1 Predicted mechanism of tRNA adenylation catalyzed by [4Fe-4S]-TtuA.

Hypothesis 1: (a) The PP-loop of [4Fe-4S]-TtuA interacts with ATP. (c) TtuB enters the catalytic site of TtuA and forms the [4Fe-4S]-TtuA-TtuB-ATP complex (PDB ID: 5ZTB). (f) Substrate tRNA enters the catalytic site of TtuA and forms the [4Fe-4S]-TtuA-TtuB-ATP-tRNA complex. (g) TtuA catalyzes tRNA adenylation and forms the [4Fe-4S]-TtuA-TtuB-AdetRNA complex with the release of PP_i . (h) TtuA catalyzes sulfur transition from the C-terminus of TtuB to the unique Fe, which forms the [4Fe-5S]-TtuA intermediate.

Hypothesis 2: (a) TtuA catalyzes ATP trimming and forms (b) the [4Fe-4S]-TtuA-ADP complex with release P_i . (d) TtuB enters the catalytic site of TtuA and forms the [4Fe-4S]-TtuA-TtuB-ADP complex. (e) Substrate tRNA enters the catalytic site of TtuA and forms the [4Fe-4S]-TtuA-TtuB-ADP-tRNA complex. (g) TtuA catalyzes tRNA adenylation and forms the [4Fe-4S]-TtuA-TtuB-AdetRNA complex with additional release of P_i .

Hypothesis 3: After the formation of the [4Fe-4S]-TtuA-TtuB-ATP complex, (d) TtuA catalyzes ATP trimming and form the [4Fe-4S]-TtuA-TtuB-ADP complex with release P_i . Then, (e) Substrate tRNA enters the catalytic site of TtuA, and (g) TtuA catalyzes tRNA adenylation with further release of P_i .

To elucidate the catalytic mechanism of tRNA thiolation, we also have to understand the regulatory mechanism of two different reactions: tRNA activation and sulfur transition. However, the experimental approaches are limited due to the oxygen sensitivity of [4Fe-4S]. Thus, I also expect that computational approaches such as RoseTTAFoldNA^[131] and molecular dynamics (MD) simulations. RoseTTAFoldNA is the newly developed tool that predicts the structure of the protein-nucleic acid complexes using deep learning approaches, which has the potential to provide the structure of the TtuA-tRNA complex.

Furthermore, MD simulations enable to reveal the structural dynamics of TtuA and the free-energy perturbation (FEP) method can calculate the binding energy of ligands^[132,133]. Although these approaches are powerful, we have to overcome the problem of force field parameters because [4Fe-4S]-TtuA contains two types of Zn fingers (Fig. 1-28a) and four Fe atoms including the unique Fe (Fig. 5-2). The missing residues of TtuA also make it difficult to perform MD simulations, but these residues maybe regulate the tRNA adenylation and sulfur transition.

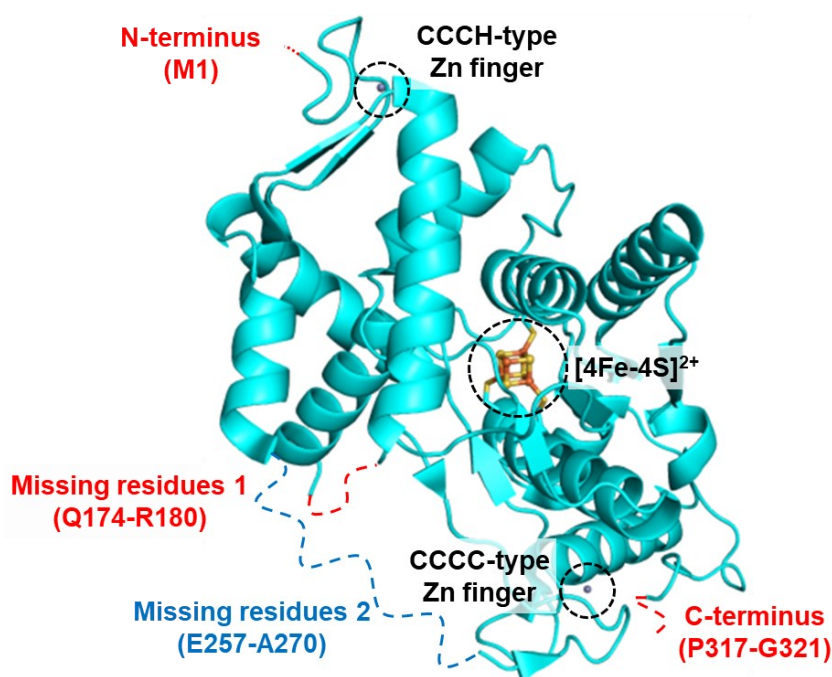


Fig. 5-2 Difficulties of MD simulations in [4Fe-4S]-TtuA.

TtuA (PDB ID: 5B4F) contains the CCCH-type zinc finger at the N-terminus, the CCCC-type zinc finger at the C-terminus, and four missing regions (Met1, Gln174-Arg180, Glu257-Ala270, and Pro317-Gly321).

Chapter 6. Summary

Thiolation at position 54 of tRNA (m^5s^2U54) is catalyzed by 2-thiouridine synthetase TtuA and sulfur donor TtuB, which allows thermophiles to survive in high-temperature environments. Our recent structural analysis of the [4Fe-4S]-TtuA-TtuB-ATP complex showed that an oxygen-sensitive [4Fe-4S] cluster with a non-cysteine coordinated Fe (the unique Fe) is required for the enzymatic activity of TtuA. On the other hand, it was reported that Ncs6, a homolog protein of TtuA, contains [3Fe-4S] or [4Fe-4S]. Therefore, the detailed mechanism of sulfur transition from TtuB to tRNA is unknown (Fig. 1-30).

In this study, I analyzed the structure of Fe-S clusters in TtuA in time-course and their enzymatic activities under strictly anaerobic conditions. As a result, I found that [3Fe-4S]-TtuA spontaneously transformed into [4Fe-4S] within one hour. I also revealed that [3Fe-4S]-TtuA could not bind to the C-terminus of TtuB. These observations supported that only [4Fe-4S] is the active form of TtuA (Fig. 4-1).

Furthermore, the result of APM-PAGE showed that [4Fe-4S]-TtuA does not transfer sulfur from TtuB to substrate tRNA until the tRNA is adenylated. In addition, the mutational analysis in this study revealed that Ser55, Asp59, and Asp161 of TtuA are responsible for tRNA adenylation and Lys137 is the catalytic residue of sulfur transition. These critical residues and the structure of catalytic pocket are conserved in not only the TtuA/Ncs6 family, but also MnmA and ThiI. Taking all result together, I proposed the reaction mechanism of tRNA-thiolation containing tRNA adenylation and sulfur transition via the unique Fe of [4Fe-4S] cluster (Fig. 4-8, Fig. 6-1). Notably, during the preparation of this dissertation, it was reported that some archaeal ThiI such as *Mma*ThiI contains [4Fe-4S] for sulfur transition^[134]. Our conclusion is consistent with this recent report that the unique Fe in [4Fe-4S]-*Mma*ThiI is essential to receive sulfur via the [4Fe-5S]-intermediate.

Finally, I showed that the time-course correlation analysis under strictly anaerobic conditions is required to reveal the correct reaction mechanism: the structures of an Fe-S cluster in the enzyme; the catalytic activity of the enzyme. I believe that this study contributes to the accurate understanding of Fe-S proteins which would exponentially be discovered in the future.

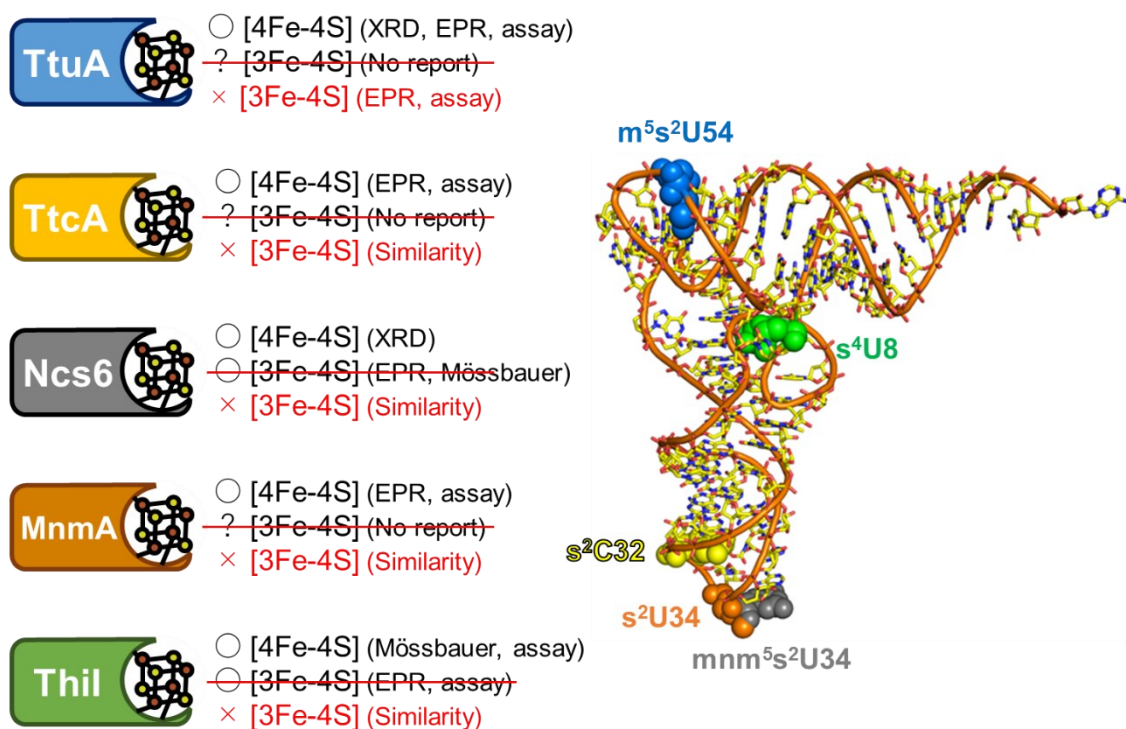


Fig. 6-1 Graphical summary of this study.

Previously, some research groups proposed that the [4Fe-4S] cluster is an active form in tRNA-thiolation enzymes whereas others reported that [3Fe-4S] cluster is a catalytic form in Ncs6 and ThiI. The possibility of the [3Fe-4S] was not considered in TtuA, TtcA, and MnmA (colored in black). This study demonstrated that the [4Fe-4S] cluster is the only active form of TtuA and proposed that all of the tRNA thiolation enzymes contain [4Fe-4S] cluster as the scaffold of tRNA adenylation and sulfur transition (colored in red).

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