Mechanism of aminoacyl-tRNA acetylation by an aminoacyl-tRNA acetyltransferase AtaT from enterohemorrhagic *E. coli*

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Toxin-antitoxin systems in bacteria contribute to stress adaptation, dormancy, and persistence. AtaT, a type-II toxin in enterohemorrhagic *E. coli*, reportedly acetylates the α-amino group of the aminoacyl-moiety of initiator Met-tRNAfMet, thus inhibiting translation initiation. Here, we show that AtaT has a broader specificity for aminoacyl-tRNAs than initially claimed. AtaT efficiently acetylates Gly-tRNA Gly, Trp-tRNA Trp, Tyr-tRNA Tyr and Phe-tRNA Phe isoacceptors, in addition to Met-tRNAfMet, and inhibits global translation. AtaT interacts with the acceptor stem of tRNAfMet, and the consecutive G-C pairs in the bottom-half of the acceptor stem are required for acetylation. Consistently, tRNA Gly, tRNA Trp, tRNA Tyr and tRNA Phe also possess consecutive G-C base-pairs in the bottom halves of their acceptor stems. Furthermore, misaminoacylated valyl-tRNAfMet and isoleucyl-tRNAfMet are not acetylated by AtaT. Therefore, the substrate selection by AtaT is governed by the specific acceptor stem sequence and the properties of the aminoacyl-moiety of aminoacyl-tRNAs.
A bacterial toxin–antitoxin (TA) module is a gene pair of a protein toxin that induces cell growth arrest and an antitoxin that counteracts the toxin\textsuperscript{1–4}. Toxins in TA-modules target pivotal cellular processes, including DNA replication, transcription, translation, and cell wall synthesis\textsuperscript{5}. TA modules have been implicated in bacterial stress adaptation, persistence, and dormancy to survive under environmental stresses, and are involved in bacterial pathogenesis\textsuperscript{6–8}. Based on the nature of antitoxins and the mechanism by which they neutralize the toxin activity, TA modules are classified into six different types\textsuperscript{9}. In the type II TA system, under normal conditions, aminoacylation of tRNAs is governed by both the specificity of aminoacyl-tRNAs, and the selection of substrate aminoacyl-tRNAs \textit{in vivo} and \textit{in vitro}. AtaT also recognizes the properties of the cognate antitoxin AtaR have been reported\textsuperscript{20,21}. The structural and biochemical studies revealed that the homodimerization of AtaT is required for both the enzymatic activity and the toxic effects of the toxin by direct protein–protein interactions. Since the type II antitoxins are susceptible to proteolytic degradation due to their inherently disordered regions, under stress conditions, antitoxin degradation is induced by proteases, such as Lon and ClpP, thus leading the activation of the toxins\textsuperscript{10,11}.

During the last few years, a new class of type II toxins belonging to the Gcn5-related N-acetyltransferase (GNAT) family has been identified\textsuperscript{12–17}. GNAT toxins are enzymes that acetylate the aminoacyl group of the aminoacyl-moiety of aminoacyl-tRNAs, using acetyl CoA (Ac-CoA) as the acetyl group donor, and inhibit cellular translation. The GNAT toxins reportedly have different substrate specificities for their target aminoacyl-tRNAs. TacT, TacT2, and TacT3 in \textit{Salmonella} can acetylate several isoaccepting aminoacyl tRNAs\textsuperscript{12,18}. TacT acetylates various aminoacyl-tRNA isoacceptors for glycine, isoleucine/leucine, tryptophan, and serine \textit{in vitro} and may inhibit global cellular translation\textsuperscript{18}. The TacT activity contributes to intracellular persistence in macrophages\textsuperscript{12,18}. ItaT, identified in the \textit{E. coli} HS strain, reportedly acetylates the isoaccepting aminoacyl tRNAs for isoleucine, valine, and methionine \textit{in vivo} and \textit{in vitro}, and may inhibit global cellular translation\textsuperscript{15,19}. On the other hand, AtaT of the enterohemorrhagic \textit{E. coli} O157:H7 strain specifically acetylates the initiator methionyl-tRNA\textsubscript{Met} (Met-tRNA\textsubscript{Met}) \textit{in vitro}\textsuperscript{13}. AtaT was assumed to block the translation initiation step, because acetylated Met-tRNA\textsubscript{Met} cannot interact with translation initiation factor 2 complexed with GTP (IF2-GTP). GNAT toxins have also been identified in \textit{Klebsiella pneumoniae} (KctA)\textsuperscript{14,17} and in the \textit{Shigella flexneri} pINV plasmid (GmvT)\textsuperscript{16}, although their aminoacyl-tRNA targets remain unknown.

Recently, the crystal structures of AtaT and its complex with the cognate antitoxin AtaR have been reported\textsuperscript{20,21}. The structural and biochemical studies revealed that the homodimerization of AtaT is required for both the enzymatic activity and the toxicity. The electrostatic surface potential of the dimeric AtaT showed that a positively charged patch is created upon dimerization\textsuperscript{14,16,20}. The electrostatic surface potential of the dimeric AtaT was experimentally determined\textsuperscript{18,22}, and its positive charge density is significantly higher than that of the monomeric AtaT\textsuperscript{22}. The positive charge density of AtaT is required for both the enzymatic activity and the toxicity\textsuperscript{14,16,20}.

To evaluate the recognition of the aminoacyl-moiety of Met-tRNA\textsubscript{Met} by AtaT, mutant tRNA\textsubscript{Met} molecules with a UAC anticodon (tRNA\textsubscript{Met}UAC) and a GAU anticodon (tRNA\textsubscript{Met}GAU) were prepared and charged with valine and isoleucine by valyl-tRNA\textsubscript{Met} synthase and isoleucyl-tRNA\textsubscript{Met} synthetase, respectively\textsuperscript{22,23}. The acetylations of valyl-tRNA\textsubscript{Met}UAC (Val-tRNA\textsubscript{Met}UAC) and isoleucyl-tRNA\textsubscript{Met}GAU (Ile-tRNA\textsubscript{Met}GAU) by AtaT were tested in \textit{in vitro}. Although Val-tRNA\textsubscript{Met}UAC and Ile-tRNA\textsubscript{Met}GAU have the same nucleotide sequence as Met-tRNA\textsubscript{Met}, except for the anticodon sequence, they were not acetylated by AtaT \textit{in vitro} (Fig. 1c). Since AtaT recognizes the acceptor region of tRNA\textsubscript{Met}, as described below, these results suggest that both the nucleotide sequence of tRNA and the side chain of the aminoacyl moiety of aminoacyl-tRNAs are important for the acetylation of aminoacyl-tRNAs by AtaT.

**Results**

AtaT recognizes both tRNA sequence and aminoacyl moiety. AtaT was previously shown to acetylate initiator methionyl-tRNA\textsubscript{Met} (Met-tRNA\textsubscript{Met}) specifically, but not the elongator methionyl-tRNA\textsubscript{Met} (Met-tRNA\textsubscript{Met}) \textit{in vitro}\textsuperscript{13}. Consistent with the previous report, the steady-state kinetics of Met-tRNA\textsubscript{Met} and Met-tRNA\textsubscript{Met} acetylation by AtaT showed different acetylation efficiencies between Met-tRNA\textsubscript{Met} and Met-tRNA\textsubscript{Met} (Fig. 1a, b). While the $K_a$ value of Met-tRNA\textsubscript{Met} is $\sim 29.6 \pm 3.8 \mu M$, the $K_a$ value of Met-tRNA\textsubscript{Met} is $172 \pm 43 \mu M$. The $V_{\text{max}}$ values of Met-tRNA\textsubscript{Met} and Met-tRNA\textsubscript{Met} acetylation were estimated to be $2.1 \pm 0.12$ and $0.38 \pm 0.07 \text{s}^{-1}$, respectively.

The overall reaction efficiency of Met-tRNA\textsubscript{Met} was $\sim 30$-fold higher than that of Met-tRNA\textsubscript{Met}, and thus Met-tRNA\textsubscript{Met} is a much better substrate than Met-tRNA\textsubscript{Met} for AtaT \textit{in vitro}. Since both Met-tRNA\textsubscript{Met} and Met-tRNA\textsubscript{Met} are charged with methionine, this result suggests that AtaT discriminates the tRNA sequence of Met-tRNA\textsubscript{Met} from that of Met-tRNA\textsubscript{Met}.

AtaT recognizes the acceptor region of tRNA. To identify the recognition elements in tRNA\textsubscript{Met} that allow AtaT to discriminate tRNA\textsubscript{Met} from tRNA\textsubscript{Met}, we first generated a chimeric tRNA\textsubscript{Met} transcript with the acceptor stem sequence of tRNA\textsubscript{Met} (fMet acceptor stem mMet) (Fig. 1a). Replacing the Met-tRNA\textsubscript{Met} acceptor stem with the tRNA\textsubscript{Met} acceptor stem conferred efficient acetylation of the mutant chimeric tRNA\textsubscript{Met} transcript, to almost 60% of that of the wild-type tRNA\textsubscript{Met} transcript (Fig. 1d, e), while the Met-tRNA\textsubscript{Met} transcript was not acetylated under these conditions. This result suggests that the elements recognized by AtaT reside in the acceptor stem region of tRNA\textsubscript{Met}.

The sequence differences in the acceptor stems between tRNA\textsubscript{Met} and tRNA\textsubscript{Met} are located at the top of the acceptor helix and the bottom half of the stem (Fig. 1a). The C1–G72 mispair at the top of the acceptor helix is one of the characteristic features of tRNA\textsubscript{Met}. Mutations of C1–A72 to the A1–C72 or G1–C72 base pair of the tRNA\textsubscript{Met} transcript decreased the acetylation efficiency to about 20% or 10% of that of the wild-type tRNA\textsubscript{Met}, respectively, while the mutation of C1–A72 to the C1–G72 base-pair decreased the acetylation efficiency to about 65% of the wild-type tRNA\textsubscript{Met} (Fig. 1d, e). Thus, the mispair at the top of the acceptor helix is not necessarily required for the acetylation of Met-tRNA\textsubscript{Met} by AtaT, but the nucleotide compositions at positions 1 and 72, in the context of the nucleotide sequences in the acceptor helix, are important for acetylation by AtaT, as described below. On the other hand, the mutation of the consecutive three G–C pairs, G4G5G6: G67C68C69, in the tRNA\textsubscript{Met} transcript to the corresponding sequence of tRNA\textsubscript{Met} (U4A5G6:G67U68A69) decreased the acetylation efficiency to about 10% of that of the wild-type...
Met-tRNA^{fMet} (Fig. 1d, e). Reversing the three consecutive G–C base pairs in the tRNA^{fMet} transcript to the three consecutive C–G pairs, C4C5C6:G67G68G69, reduced the acetylation efficiency to about 10% of that of wild-type Met-tRNA^{fMet} (Fig. 1d, e). The mutations of one of the three consecutive G–C pairs in the tRNA^{fMet} transcript to a C–G pair (C4–G69, C5–G68, or C6–G67) decreased the acetylation efficiency to 25–50% of that of the wild-type Met-tRNA^{fMet} (Fig. 1f, g). These results suggest that AtaT mainly recognizes the acceptor region of tRNA^{fMet} for discrimination from tRNA^{mMet}. In particular, the bottom-half
consecutive three G–C base-pairs in the tRNA\textsuperscript{Met} acceptor stem, rather than the C1–A72 mispair, are more important for the acetylation of Met-tRNA\textsuperscript{Met} by AtaT.

Structural determination of AtaT–Ac–Met–tRNA\textsuperscript{Met} complex. To uncover the molecular basis of Met-tRNA\textsuperscript{Met} recognition by AtaT, AtaT, and acetyl–Met–tRNA\textsuperscript{Met} (Ac–Met–tRNA\textsuperscript{Met}) were co-crystallized and the structure was determined. Since Ac–Met–tRNA\textsuperscript{Met} is chemically more stable than Met–tRNA\textsuperscript{Met}, Ac–Met–tRNA\textsuperscript{Met} was used instead of Met–tRNA\textsuperscript{Met} for crystallization. The structure should represent a post-reaction stage in which the acetyl group of acetyl-CoA (Ac-CoA) is transferred to the α-amino group of the aminoacyl moiety of Met–tRNA\textsuperscript{Met}, and the CoA byproduct is released from the AtaT catalytic site.

The crystal belongs to the space group P2\textsubscript{1}2\textsubscript{1}2\textsubscript{1} and contains eight AtaT molecules and two Ac–Met–tRNA\textsuperscript{Met} molecules in the asymmetric unit. The initial phase was determined by the molecular replacement method, using the AtaT homodimer (PDB: 6GTP)\textsuperscript{20} and tRNA\textsuperscript{Met} structures in the structure of the complex of Met–tRNA\textsuperscript{Met} formyltransferase and formyl–Met–tRNA\textsuperscript{Met} (PDB: 2PMT)\textsuperscript{24} as search models. The structure was model-built and refined to an R factor of 28.6% (R\textsubscript{free} = 34.1%) at 3.8 Å resolution. Detailed crystallographic data collection and refinement statistics are provided in Table 1.

Of the eight AtaT molecules in the asymmetric unit (AtaT1–8), six (AtaT3–8) formed three dimers (AtaT3–AtaT4, AtaT5–AtaT6, and AtaT7–AtaT8), and AtaT1 and AtaT2 each formed AtaT1–AtaT1′ and AtaT2–AtaT2′ dimers, respectively, through the crystallographic two-fold rotation axis (Supplementary Fig. 1). The 3′ single-stranded regions of the two tRNA molecules, tRNA\textsuperscript{Met}1 and tRNA\textsuperscript{Met}1′, tRNA\textsuperscript{Met}2 and tRNA\textsuperscript{Met}2′ interact with the regions proximal to the two catalytic sites of AtaT1–AtaT1′ and AtaT2–AtaT2′, respectively, through the crystallographic two-fold rotation axis (Supplementary Fig. 1). The 3′ single-stranded regions of the two tRNA molecules, tRNA\textsuperscript{Met}1 and tRNA\textsuperscript{Met}1′, tRNA\textsuperscript{Met}2 and tRNA\textsuperscript{Met}2′ interact with the regions proximal to the two catalytic sites of AtaT1–AtaT1′ and AtaT2–AtaT2′, respectively.

Interaction of tRNA with AtaT homodimer. In the structure of AtaT complexed with Ac–Met–tRNA\textsuperscript{Met}, two tRNA\textsuperscript{Met} molecules (tRNA\textsuperscript{Met}a and tRNA\textsuperscript{Met}b) interact with an AtaT homodimer (AtaTα and AtaTβ) (Fig. 2a). The 3′-terminal single-stranded region and acceptor stem of each tRNA\textsuperscript{Met} interact with the positively charged interface between the two subunits of the AtaT dimer. The 3′-single-stranded termini of tRNA\textsuperscript{Met}a and tRNA\textsuperscript{Met}b are directed toward the active sites of AtaTα and AtaTβ in the dimer, respectively (Fig. 2b). The binding of the acceptor stem region to the interface between the AtaT dimer subunits explains the previous results showing that the AtaT homodimerization is required for both the enzymatic and toxic activities of AtaT\textsuperscript{11}. The structure also revealed that the recognition elements in tRNA\textsuperscript{Met} for acetylation by AtaT reside in its acceptor stem region (Fig. 1d, e).

In the complex structure, the a3 in AtaT\textsubscript{β} penetrates the major groove of the acceptor helix of tRNA\textsuperscript{Met}a, and the a1 in AtaT\textsubscript{α} also penetrates the groove from another angle (Fig. 2c). As a result, C1–A72 at the top of the acceptor helix of tRNA\textsuperscript{Met} are completely separated and the 5′-terminal C1 nucleoside is flipped. The 5′-phosphate of C1 interacts with K82 and K165 in AtaT\textsubscript{β} and with N41 and R37 in AtaT\textsubscript{α} (Fig. 2c, f). On the other hand, although 3′-C75A76–Ac–Met of Ac–Met–tRNA\textsuperscript{Met} were not visible in the present structure (Supplementary Fig. 2), the nucleobases of A72, A73, and C73 are continuously stacked and the 3′-end is directed toward the catalytic site, where G108 and Y114 participate in Ac-CoA binding and catalysis, respectively.

![Fig. 1 Elements in Met-tRNA\textsuperscript{Met} required for acetylation by AtaT.](https://doi.org/10.1038/s41467-020-19281-z)

- **a**: Secondary structures of E. coli initiator tRNA\textsuperscript{Met} and elongator tRNA\textsuperscript{Met} in cloverleaf forms. The nucleotide substitutions in the tRNA\textsuperscript{Met} and tRNA\textsuperscript{Met} transcripts used for assays in d and f are shown by boxes and arrows.
- **b**: Steady-state kinetics of methionyl-tRNA\textsuperscript{Met} (Met-tRNA\textsuperscript{Met}) and methionyl-tRNA\textsuperscript{Met} (Met-tRNA\textsuperscript{Met}) acetylation by AtaT. The initial velocities of acetylation of Met-tRNA\textsuperscript{Met} and Met-tRNA\textsuperscript{Met} were measured with various concentrations of Met-tRNA\textsuperscript{Met} and Met-tRNA\textsuperscript{Met} (5–90 μM). Magnified view of the graph for the acetylation of Met-tRNA\textsuperscript{Met} and Met-tRNA\textsuperscript{Met} and tRNA\textsuperscript{Met} were overexpressed in E. coli and purified.
- **c**: In vitro acetylation of misaminoacylated Val-tRNA\textsuperscript{Met}UAC and Ile-tRNA\textsuperscript{Met}GAU, tRNA\textsuperscript{Met}UAC and tRNA\textsuperscript{Met}GAU were overexpressed in E. coli, purified and charged with valine and isoleucine by valyl-tRNA and isoleucyl-tRNA synthetases, respectively. Aminoacyl-tRNA\textsuperscript{s} (10 μM) were used for the acetylation by AtaT and incubated for 12 min at 37 °C.
- **d**: Time courses of the acetylation of Met-tRNA\textsuperscript{Met}, Met-tRNA\textsuperscript{Met}, and Met-tRNA\textsuperscript{Mut}a, and Met-tRNA\textsuperscript{Mut}b, respectively.
- **e**: Quantification of the acetylation efficiencies in d. The initial velocities of the acetylation of Met-tRNA\textsuperscript{Met}, Met-tRNA\textsuperscript{Met}, and Met-tRNA\textsuperscript{Mut} were measured. The bars in the graph are SD of three independent (n = 3) experiments, and the data are presented as mean values ± SD.

### Table 1 Data collection and refinement statistics.

| Data collection | AtaT-Ac-Met-tRNA\textsuperscript{Met} |
|-----------------|--------------------------------------|
| Space group     | P2\textsubscript{1}2\textsubscript{1}2 |
| Cell dimensions | a, b, c (Å)                          |
| Wavelength (Å)  | 0.98000                               |
| Resolution (Å)  | 48.21–3.804                           |
| R\textsubscript{sym} (Å) | 0.449 (3.45)               |
| R/α             | 8.4 (1.3)                             |
| Completeness (%)| 99.64 (99.27)                         |
| Redundancy (Å)  | 26.2 (26.6)                           |
| Resolution (Å)  | 48.21–3.804                           |
| No. of reflections | 25,488                               |
| No. of atoms    | 10,968                                |
| Protein         | 3,206                                 |
| RNA             | 142.23                                |
| r.m.s. deviations | 131.99                             |
| Bond length (Å) | 0.005                                 |
| Bond angles (°) | 1.09                                  |

*Values in parentheses are for the highest-resolution shell.*
The ribbon models of the AtaTa, AtaTb, and AcMet molecules (tRNAs and tRNAb) are colored as in Fig. 2d. The loop between α3 and β4 in AtaTb and α1 in AtaTa form the path for the 3′-end of tRNA to enter the catalytic site and interact with the 3′-single-stranded region of tRNAfMet (Fig. 2d). The ribose 2′-OH groups of C74 and A73 in tRNAfMet interact with the main-chain carbonyl oxygens of P84 in AtaTb and H34 in AtaTa, respectively, and the O2 atom of the C74 nucleobase of tRNAfMet interacts with R71 in AtaTb. Residues S76, K77, S78, and K79 in α3 of AtaTb interact with the phosphate backbones of C67C68C69, at the bottom half of the acceptor helix. K81 in α3 is positioned so its side chain can interact with the phosphate backbones and nucleobases of consecutive G–C pairs, at the top-half of the acceptor helix (Fig. 2d, f). Although an in vitro analysis demonstrated the importance of the (G4G5G6:C67C68C69) sequence at the bottom half of the acceptor helix (Fig. 1d, e), no base-specific interactions were observed in the present structure, as described below.

The 3′-C75A76–Ac–Met region of AcMet–tRNAfMet was disordered and not visible in the present structure. Y85 in the...
loop between α3 and β4 in AtaTb, and R95, which stacks with F30 in α1, in β4 of AtaTa form a hydrogen-bond, and these interactions would block the access of the 3′-end of Ac–Met–tRNA\textsuperscript{Met} to the catalytic site (Fig. 2e). No significant structural differences were detected in a comparison of the catalytic pocket structure of AtaTb complexed with Ac–Met–tRNA\textsuperscript{Met} with that of apo AtaT (Supplementary Fig. 3). Thus, the present structure represents a post-reaction stage of Met–tRNA\textsuperscript{Met} acetylation, where the Ac–Met–tRNA\textsuperscript{Met} product is ready for release from the enzyme after the completion of the reaction. This would explain why no specific interaction between α3 and the bottom half of the acceptor helix was observed in the present structure. During the reaction stage of the acetylation of Met–tRNA\textsuperscript{Met}, the basic residues in α3 would specifically recognize the sequence and structure at the bottom half of the tRNA\textsuperscript{Met} acceptor helix.

To verify the effects of the above-mentioned residues on the activity of AtaT, AtaT mutants were assessed for the ability to inhibit E. coli cell growth (Fig. 3). G108D and Y144F mutants were used as positive controls, since the mutation of G108 to D108 causes AtaT to lose its toxicity by blocking Ac-CoA binding\textsuperscript{20,21}, and the mutation of the catalytic residue Y144 to F144 also decreases AtaT toxicity in vivo\textsuperscript{20,21,25}. As expected, the substitution of positively charged residues in helix α1 (H34A/R37A/R40A) attenuated the cell growth inhibition by AtaT. R71A slightly decreased the AtaT cytotoxicity (Fig. 3a). S76A and S78A also decreased the AtaT toxicity, and the mutation of lysines to alanines in α3 (K77A/K79A/K81A/K82A) reduced the AtaT toxicity to the same extent as Y144F. R95A decreased the AtaT toxicity to a similar level as Y144F, and F30A also affected the AtaT toxicity. Y85A abolished the AtaT toxicity (Fig. 3b). These results suggest that the above-mentioned residues are important for tRNA\textsuperscript{Met} recognition during the acetylation of Met–tRNA\textsuperscript{Met}. The effects of the mutations on the suppression of growth arrest in liquid medium (Fig. 3a, b) were also confirmed by the suppression of growth inhibition on agar plates (Fig. 3c).

**Acetylation of aminoacyl-tRNAs by AtaT in vivo.** Although AtaT was initially shown to acetylate the α-amino group of Met–tRNA\textsuperscript{Met} in vitro (Fig. 1b)\textsuperscript{13}, there is no direct evidence for Met–tRNA\textsuperscript{Met} acetylation by AtaT in vivo. To evaluate the in vivo target of aminoacyl–tRNAs for acetylation by AtaT, AtaT was expressed in E. coli and the intracellular aminoacyl–tRNAs were analyzed by LC/MS\textsuperscript{26} (Fig. 4a). E. coli cells were harvested after the induction of AtaT expression, using the arabinose-inducible pBAD system\textsuperscript{27}. Upon arabinose addition, the growth of AtaT-expressing cells was suppressed, as compared to that of E. coli cells bearing the empty pBAD plasmid.

To analyze intracellular aminoacyl-tRNAs, the RNA fraction was extracted under acidic and cold conditions, and subsequently acetylated by stable isotopic acetic anhydride-D\textsubscript{6} ([CD\textsubscript{3}CO])\textsubscript{2}O; D is deuterium) to chemically acetylate the remaining aminoacyl-tRNAs.

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**Fig. 3 Toxicity of AtaT variants.** a, b Growth curves of E. coli MG1655 transformed with pBAD-ataT and its variants. E. coli MG1655 transformed by pBAD-ataT or its variants was inoculated in LB containing 50 μg mL\textsuperscript{−1} ampicillin and supplemented with 0.2% (w/v) arabinose at 37 °C. Bars in the graphs indicate the SD of three independent experiments. c Overnight cultures of E. coli MG1655 transformed with pBAD-ataT and its variants were serially diluted, and the dilutions were spotted on LB agar plates containing 50 μg mL\textsuperscript{−1} ampicillin and supplemented with 0.2% (w/v) arabinose or without arabinose. The plates were incubated at 37 °C overnight. The bars in the graph are SD of three independent (n = 3) experiments, and the data are presented as mean values ± SD.
As a result, all aminoacyl-tRNAs were converted into either intracellularly acetylated aminoacyl-tRNAs (Ac-aa-tRNAs) by AtaT or chemically acetylated aminoacyl-tRNAs (D3Ac-aa-tRNAs). The tRNAs were then digested with RNase I, and Ac-aa-A76 (A76 is the 3′-terminal adenosine of the tRNA molecule) and D3Ac-aa-A76 of each amino acid species were detected by LC/MS. The scheme of the analysis is illustrated in Fig. 4a.

The LC/MS analyses of RNase I-digested RNAs prepared from cells with AtaT induction revealed the molecular masses corresponding to Ac-Gly-A76, Ac-Trp-A76, Ac-Tyr-A76, Ac-Phe-A76, Ac-Gln-A76, Ac-Asn-A76, and Ac-Met-A76, derived from Ac-Gly-tRNAGly, Ac-Trp-tRNATrp, Ac-Tyr-tRNATyr, Ac-Phe-tRNAPhe, Ac-Gln-tRNAGln, Ac-Asn-tRNAAasn, and Ac-Met-tRNAMet isocellotransferringos, respectively. Thus, AtaT acetylates not only Met-tRNAMet, but also Gly-tRNAGly, Trp-tRNATrp, Tyr-tRNATyr, Phe-tRNAPhe, Gln-tRNAGln, and Asn-tRNAAasn (Fig. 4b, Supplementary Fig. 4).

Quantification of the ratio of Ac-aa-A76 and D3Ac-aa-A76 enables an estimation of the fraction of aminoacyl-tRNAs acetylated by the action of AtaT in vivo. The amounts of Ac-aa-A76 and D3Ac-aa-A76, after 15 min of AtaT expression induction, were quantified, and the fractions of individual aminoacyl-tRNAs acetylated by AtaT in vivo were estimated as the ratio of Ac-aa-A76 to the sum of the amounts of Ac-aa-A76 and D3Ac-aa-A76. The bars in the graph are SD of three independent (n = 3) experiments, and the data are presented as mean values ± SD. c Time courses of the fractions of aminoacyl-tRNAs acetylated by AtaT in vivo. The bars in the graphs are SD of three independent experiments.

Fig. 4 Acetylation of aminoacyl-tRNAs by AtaT in vivo. a Schematic diagram of the LC/MS detection and quantification of acetyl-aminoacyl-tRNAs, produced by the action of AtaT in vivo. b LC/MS analysis of RNase I-digested fragments of acetyl-aminoacyl-tRNAs after 15 min of induced AtaT expression (pBAD-AtaT) and the negative control (pBAD). Extracted ion chromatograms of the proton adducts corresponding to Ac-Gly-A76 (m/z = 367.13), Ac-Trp-A76 (m/z = 496.19), Ac-Phe-A76 (m/z = 457.18), Ac-Tyr-A76 (m/z = 473.17), Ac-Met-A76 (m/z = 441.15), and Ac-Ala-A76 (m/z = 381.15), derived from Ac-Gly-tRNAGly, Ac-Trp-tRNATrp, Ac-Phe-tRNAPhe, Ac-Tyr-tRNATyr, Ac-Met-tRNAMet, and Ac-Ala-tRNAAala isocellotransferringos, respectively. The two peaks observed for each Ac-aa-A76 are likely attributable to structural isomers of 3′-acetyl-aminoacyl-A76 and 2′-acetyl-aminoacyl-A76, as 3′-O-methyl and 2′-O-methyl nucleotides are known to be separated in a similar manner. c Fractions of aminoacyl-tRNAs acetylated by the action of AtaT in vivo. The amounts of Ac-aa-A76 and D3Ac-aa-A76, after 15 min of AtaT expression induction, were quantified, and the fractions of individual aminoacyl-tRNAs acetylated by AtaT in vivo were estimated as the ratio of Ac-aa-A76 to the sum of the amounts of Ac-aa-A76 and D3Ac-aa-A76. The bars in the graph are SD of three independent (n = 3) experiments, and the data are presented as mean values ± SD. d Time courses of the fractions of aminoacyl-tRNAs acetylated by AtaT in vivo. The bars in the graphs are SD of three independent experiments.
Acetylation of aminoacyl-tRNAs by AtaT in vitro. The LC/MS analysis of the intracellular aminoacyl-tRNAs upon AtaT expression revealed that, in addition to Met-tRNA\textsubscript{Met}, Gly-tRNA\textsubscript{Gly}, Thr-tRNA\textsubscript{Thr}, Tyr-tRNA\textsubscript{Tyr}, and Phe-tRNA\textsubscript{Phe} are efficiently acetylated in vivo by AtaT (Fig. 4c). To verify the results, the acetylation of those aminoacyl-tRNA species (10 μM) by AtaT was assessed in vitro before AtaT was estimated to be ~30 μM (Fig. 1b). Thus, the 10 μM concentration of various aminoacyl-tRNAs for the in vitro assays would be sufficiently sensitive to evaluate the specificity of AtaT for aminoacyl-tRNAs. The results revealed the efficient acetylation of the isoacceptors of Gly-tRNA\textsubscript{Gly}, Thr-tRNA\textsubscript{Thr}, Tyr-tRNA\textsubscript{Tyr}, and Phe-tRNA\textsubscript{Phe} by AtaT (Fig. 5a, b). The acetylation efficiencies of Gly-tRNA\textsubscript{Gly} and Thr-tRNA\textsubscript{Thr} by AtaT were about two-times and three-times higher than that of Met-tRNA\textsubscript{Met}, respectively. The acetylation efficiencies of the Tyr-tRNA\textsubscript{Tyr} and Phe-tRNA\textsubscript{Phe} isoacceptors were almost the same as that of Met-tRNA\textsubscript{Met}, Gln-tRNA\textsubscript{Gln} isoacceptors were also acetylated by AtaT, but with a relatively lower efficiency of <25% of Met-tRNA\textsubscript{Met}. Since tryptophan, tyrosine, and phenylalanine, but not glycine, are aromatic hydrophobic amino acids (Fig. 5c), we tested the acetylation of other aminoacyl-tRNAs charged with hydrophobic amino acid residues, such as Val-tRNA\textsubscript{Val}, Ile-tRNA\textsubscript{Ile}, Ala-tRNA\textsubscript{Ala}, and Leu-tRNA\textsubscript{Leu} isoacceptors. Neither Val-tRNA\textsubscript{Val}, Ile-tRNA\textsubscript{Ile}, Ala-tRNA\textsubscript{Ala} nor Leu-tRNA\textsubscript{Leu} was acetylated by AtaT in vitro, and these aminoacyl-tRNAs were also not acetylated by AtaT in vivo (Figs. 4c and 5b). The in vitro acetylation analyses showed similar substrate preferences of AtaT as those in vivo, and thus AtaT has a broader specificity toward aminoacyl-tRNAs than initially described\textsuperscript{13}.

The superimposition of the AtaT structure onto the structure of the protein N-terminal acetyltransferase, NatF, complexed with a bisubstrate analog, Coa-\textsubscript{Ac-MKAV}\textsuperscript{28}, identified the residues that may constitute the pocket for the aminoacyl moiety side chain in AtaT. In the model of AtaT complexed with Coa-\textsubscript{Ac-MKAV} (Fig. 5d), the methionine (M) side chain of Coa-\textsubscript{Ac-MKAV} is proximal to A26, L27, and L136. These hydrophobic residues would constitute the pocket for the aminoacyl-moiety of aminoacyl-tRNAs. The L27E mutation suppressed the toxicity of AtaT on the solid and liquid LB liquid media to the same extent, as the Y144F catalytic mutation, and the L136E mutation weakly suppressed the toxicity of AtaT in the liquid medium (Fig. 5e, f). Thus, the hydrophobic and aromatic side chains of the aminoacyl moieties of aminoacyl-tRNAs, Trp-tRNA\textsubscript{Trp}, Tyr-tRNA\textsubscript{Tyr}, Phe-tRNA\textsubscript{Phe}, and Met-tRNA\textsubscript{Met}, would be recognized by these hydrophobic residues in the active site of AtaT. The recognition of Gly-tRNA\textsubscript{Gly} is discussed below.

Discussion

In this study, we showed that, in addition to Met-tRNA\textsubscript{Met}, AtaT can acetylate isoacceptors of Gly-tRNA\textsubscript{Gly}, Thr-tRNA\textsubscript{Thr}, Tyr-tRNA\textsubscript{Tyr}, and Phe-tRNA\textsubscript{Phe} efficiently, and Gln-tRNA\textsubscript{Gln} to some extent, in vivo and in vitro (Figs. 4 and 5). In particular, Gly-tRNA\textsubscript{Gly} and Thr-tRNA\textsubscript{Thr} are more efficiently acetylated than Met-tRNA\textsubscript{Met}, and Tyr-tRNA\textsubscript{Tyr} and Phe-tRNA\textsubscript{Phe} are acetylated as efficiently as Met-tRNA\textsubscript{Met} in vitro. Therefore, the substrate specificity of AtaT toward aminoacyl-tRNAs is broader than initially described. Moreover, after AtaT expression in \textit{E. coli}, the level of formyl-Met-tRNA\textsubscript{Met} in the cells did not change (Supplementary Fig. 5). The previous study suggested that the acetylation of the α-amino group of the aminoacyl-moiety of Met-tRNA\textsubscript{Met} by AtaT inhibits the formation of the α-amino group of the aminoacyl-moiety of Met-tRNA\textsubscript{Met} by formylase\textsuperscript{13}. Thus, AtaT was assumed to block the translation initiation step, because acetylated Met-tRNA\textsubscript{Met} cannot interact with translation initiation factor 2. After methionylation of tRNA\textsubscript{Met} by methionyl-tRNA synthetase, Met-tRNA\textsubscript{Met} is subsequently formylated and used for translation initiation. Under the conditions in which AtaT is expressed in \textit{E. coli}, AtaT and formylase would both compete for the Met-tRNA\textsubscript{Met} substrate. The \textit{K}_m values of Met-tRNA\textsubscript{Met} toward the formylase and AtaT are 0.2 ± 0.122 and 29.6 ± 3.8 μM (Fig. 1b), respectively. Thus, the affinity of Met-tRNA\textsubscript{Met} toward the formylase is around two orders of magnitude larger than that of Met-tRNA\textsubscript{Met} toward AtaT for acetylation. Therefore, even after the induction of AtaT in \textit{E. coli}, the level of fMet-tRNA\textsubscript{Met} would not be drastically decreased and the translation initiation would not be inhibited. Consistent with this notion, as revealed by the LC/MS analyses of aminoacyl-tRNAs isolated from \textit{E. coli} with AtaT induction, AtaT does not change the level of fMet-tRNA\textsubscript{Met} in vivo (Supplementary Fig. 5). Thus, AtaT inhibits global translation, rather than the translation initiation step. In the previous report\textsuperscript{13}, to identify the target aminoacyl-tRNA of AtaT, various synthetic tRNA transcripts were used as substrates for aminoacylation by the \textit{E. coli} S100 fraction, followed by acetylation by AtaT. Some of the synthetic tRNA transcripts would not have been aminoacylated efficiently, as they lacked modified nucleosides in them, and thus only the acetylation of initiator Met-tRNA\textsubscript{Met} was significantly observed.

The structural analysis of AtaT complexed with Ac-Met-tRNA\textsubscript{Met} and the biochemical analysis of AtaT using tRNA\textsubscript{Met} variant transcripts suggested that the acceptor stem region of tRNA interacts with the dimeric AtaT, and the consecutive three G–C pairs in the bottom half of the acceptor stem are important for the acetylation by AtaT (Figs. 1d, g, 2a). Consistent with this
Fig. 5 Acetylation of aminoacyl-tRNAs by AtaT in vitro. a Time courses of the acetylations of Gly-tRNAGly (Gly), Trp-tRNATrp (Trp), Tyr-tRNATyr (Tyr), Phe-tRNAPhe (Phe), Met-tRNAfMet (f-Met), Met-tRNAmMet (m-Met), Leu-tRNALeu (Leu), Ala-tRNAAla (Ala), Ile-tRNAIle (Ile), and Val-tRNAVal (Val) by AtaT in vitro. The respective aminoacyl-tRNAs (10 μM) were used for the acetylation by AtaT. b Quantification of the acetylation efficiencies in a. The relative initial velocities of the acetylation of various aminoacyl-tRNAs were measured. Acetylation of the Met-tRNAfMet transcript was taken as 1.0. The bars in the graph are SD of three independent (n = 3) experiments, and the data are presented as mean values ± SD. c Nucleotide sequences of acceptor helices and chemical structures of aminoacyl-moieties of aminoacyl-tRNAs acetylated by AtaT. d Model of CoA-Ac-MKAV in the catalytic pocket of the AtaT structure. CoA-Ac-MKAV is shown in a stick model. e Growth curves of E. coli MG1655 transformed with pBAD-ataT and its variants with mutations in the putative aminoacyl moiety binding site. f Overnight cultures of E. coli MG1655 transformed with pBAD-ataT and its variants in e were serially diluted, and the dilutions were spotted on LB agar plates containing 50 μg mL⁻¹ ampicillin supplemented with 0.2% (w/v) arabinose or without arabinose. The plates were incubated at 37 °C overnight.
finding, the aminoacyl-tRNAs acetylated by AtaT in vivo, other than Met-tRNAMet, also carry two or three G–C pairs in the bottom halves of their acceptor stems (Fig. 5c). At the reaction stage, α3 in AtaT should specifically interact with the G–C pairs in the bottom half of the acceptor stem. The mutation of C1–A72 to C1–G72 in the tRNAMet transcript did not abolish acetylation by AtaT (Fig. 1d, e). Therefore, the mispair at the top of the acceptor helix is not necessarily required for the acetylation of aminoacyl tRNAs. On the other hand, mutations of C1–A72 to A1–C72 or G1–C72 in the tRNAMet transcript decreased the acetylation efficiency by AtaT (Fig. 1d, e). A purine base at position 72 might strengthen the base stacking with the base at 73 to facilitate the relocation of the aminoacyl-moiety in the active site of AtaT. Gly-tRNAGly, Tyr-tRNATyr, and Phe-tRNAPhe have G1–C72 base pairs at the top of their acceptor stems and are efficiently acetylated by AtaT in vivo and in vitro (Figs. 4c, d and 5a, b). The G–C pairs in the bottom half of the acceptor stems of tRNAGly, tRNATyr, or tRNAPhe might be sufficient to overcome the negative effect of the G1–C72 base pair at the top of the acceptor helix for acetylation by AtaT.

While Met-tRNAMet is acetylated by AtaT, misaminoacylated tRNAMet, such as Val-tRNAMetUAC and Ile-tRNAMetGAU, are not acetylated (Fig. 1c), suggesting that the aminoacyl-moieties of aminoacyl-tRNAs are also recognized by AtaT. The structure of the binding pocket for the aminoacyl-moiety of aminoacyl-tRNA in AtaT would be suitable for the accommodation of hydrophobic aromatic residues (Fig. 5d–f). The side chains of tryptophan, tyrosine, and phenylalanine would snugly fit into the hydrophobic pocket of the AtaT catalytic site. The hydrophobic side chain of methionine would also be accommodated in the hydrophobic pocket. While the side chain of the aminoacyl-moiety of Gly-tRNAGly is a hydrogen atom and thus not hydrophobic, the small hydrogen atom could enter the hydrophobic pocket of AtaT without any restraints. On the other hand, although the valine and isoleucine side chains of mischarged Val-tRNAMetUAC and Ile-tRNAMetGAU, respectively, are hydrophobic, they are small and cannot be snugly accommodated within the hydrophobic pocket, and thus neither Val-tRNAMet nor Ile-tRNAMet is acetylated by AtaT (Fig. 1c). Since glutamine is not a hydrophobic residue, the side chain of the aminoacyl-moiety of Gln-tRNAGln cannot snugly fit in the hydrophobic pocket of AtaT, and tRNAGln has two G–C pairs and one U–A pair in the bottom half of the acceptor stem (Fig. 5c). Thus Gln-tRNAGln is not acetylated as efficiently as other substrate aminoacyl-tRNAs by AtaT. The selection mechanism of the aminoacyl-moiety side-chain of an aminoacyl-tRNA is homogeneous to that of ItaT26 and other related aminoacyl-tRNA protein transferases, such as leucyl/phenylalanyl-tRNA protein transferase29,30. The size and shape of the binding pocket of AtaT for the aminoacyl moiety of an aminoacyl-tRNA are suitable for the accommodation of hydrophobic aromatic residues, except Gly-tRNAGly.

Taken together, the selection of an aminoacyl-tRNA for acetylation by AtaT is governed by both the specific sequence in the acceptor stem and the properties of the aminoacyl-moiety side chains of aminoacyl-tRNAs. The mechanism governing the affinity of AtaT for an aminoacyl-tRNA is analogous to that observed in the interaction between an aminoacyl-tRNA and translation elongation factor Tu (EF-Tu). The affinity of EF-Tu for an aminoacyl-tRNA is modulated by the aminoacyl moiety and tRNA sequence, and the sum of the affinities for the aminoacyl-moiety and the RNA body determines the total affinity of EF-Tu for an aminoacyl-tRNA31–33. While EF-Tu binds with approximately the same affinity to all elongator aminoacyl-tRNAs, AtaT binds with different affinity to aminoacyl-tRNAs. The sum of the affinities of AtaT for the hydrophobic aminoacyl-moiety and the tRNA acceptor stem sequence would determine the suitability of a substrate to be acetylated by AtaT.

Methods

Plasmas. The plasmas pET-ataT(G108D) for overexpression of AtaT (G108D), and the plasmid pET-ataRT for overexpression of the AtaRT complex were previously described21. The pBAD-ataT plasmid, for evaluation of the toxic activity of AtaT in vivo, was described previously21. Mutations in pBAD-ataT were introduced by the overlap PCR method. The DNA fragment encoding glutaminyl-tRNA synthetase, glnS, was PCR amplified from the E. coli genome and cloned into the Ndel and XhoI sites of pET-22b. The plasmids for expression of methionyl-tRNA synthetase and isoleucyl-tRNA synthetase were described previously26. The plasmas for the overexpression of aminoacyl-tRNA synthetases (ARSs: AlaRS, ValRS, LeuRS, GlyRS, ThrRS, TyrRS, and PheRS) were kindly gifted from Dr. Shirimizu (RIKEN, Japan). The oligonucleotide sequences used for the plasmid constructions or mutageneses are listed in Supplementary Table S1.

Protein expression and purification. The AtaT (G108D) protein was overexpressed in E. coli BL21(DE3) cells (Novagen-Merk Millipore) and purified as described21. The wild-type AtaT was expressed with AtaR in E. coli BL21(DE3) cells and purified from the AtaT–AtaR complex as described21.

In vitro transcription of mutant tRNAMet. A precursor tRNA bearing a 5′ leader was synthesized by T7 RNA polymerase and processed by RNaseP, comprising the M1 RNA and C5 protein. The synthetic rnpG gene encoding M1 RNA under the T7 promoter was synthesized and purchased from Eurofins, Japan, and M1 RNA was synthesized by T7 RNA polymerase. The synthetic rnpG gene encoding the C5 protein was also synthesized and purchased from Eurofins, Japan, and cloned into the Ndel and XhoI sites of pET-22b. The C5 protein was overexpressed in E. coli BL21(DE3) and purified essentially as described. The pre-tRNA transcript was processed in a mixture containing 100 nM M1 RNA, 100 nM C5 protein, 30 mM Tris–HCl, pH 7.4, 10 mM MgCl2, and 100 mM NH4Cl at 37 °C for 3 h, followed by phenol-extraction and isopropanol precipitation. The processed mature tRNAs...
were dissolved in 20 mM Tris–HCl, pH 7.4, buffer containing 200 mM NaCl, applied to a Resource Q column (GE Healthcare, Japan), and separated by a linear NaCl gradient (0.2–1.0 M) in the buffer.

The DNA fragments encoding template sequences for in vitro transcription, and the sequences of the synthetic rnpA and rnpB genes, were listed in Supplementary Table 2.

**Purification of overexpressed tRNAs in E. coli cells.** tRNA<sup>Met</sup>, tRNA<sup>Glu</sup>, tRNA<sup>Thr</sup>, tRNA<sup>Ser</sup>, tRNA<sup>Arg</sup>, tRNA<sup>Ile</sup>, tRNA<sup>Leu<sup>UAA</sup></sup>, tRNA<sup>Leu<sup>UAG</sup></sup>, tRNA<sup>Pro</sup>, tRNA<sup>Gln</sup>, tRNA<sup>Ala</sup>, tRNA<sup>Val</sup>, and tRNA<sup>Gly</sup>, tRNA<sup>Tyr</sup>, tRNA<sup>Asp</sup>, tRNA<sup>Asn</sup>, tRNA<sup>Ala</sup>, tRNA<sup>Trp</sup>, and tRNA<sup>Met</sup> were overexpressed. E. coli transformed with the pBSTNAV3 plasmids encoding the respective tRNA genes, and purified as described previously. The synthetic DNA fragments encoding the E. coli tRNA<sup>Met</sup>, tRNA<sup>Glu</sup>, tRNA<sup>Thr</sup>, and tRNA<sup>Ser</sup> genes were inserted between the ScaI and PstI sites of the pBSTNAV3 plasmids. DNA fragment sequences are listed in Supplementary Table 2. The pBSTNAV3 plasmids encoding tRNA<sup>Met</sup>, tRNA<sup>Glu</sup>, tRNA<sup>Thr</sup>, tRNA<sup>Ser</sup>, tRNA<sup>Arg</sup>, tRNA<sup>Ile</sup>, tRNA<sup>Leu<sup>UAA</sup></sup>, tRNA<sup>Leu<sup>UAG</sup></sup>, tRNA<sup>Pro</sup>, tRNA<sup>Gln</sup>, tRNA<sup>Ala</sup>, tRNA<sup>Val</sup>, tRNA<sup>Gly</sup>, tRNA<sup>Tyr</sup>, tRNA<sup>Asp</sup>, tRNA<sup>Asn</sup>, tRNA<sup>Ala</sup>, tRNA<sup>Trp</sup>, and tRNA<sup>Met</sup> were described previously. The E. coli JM110Tr strain was transformed with the pBSTNAV3 plasmid and cultured in 2× YT medium containing 50 μg·ml<sup>−1</sup> ampicillin, at 37 °C for 24 h. The total tRNA fraction was prepared as described, with modifications. After deacylation of the tRNAs, tRNAs were dissolved in buffer containing 20 mM Tris–HCl, pH 7.4, 150 mM KCl, 7 mM MgCl<sub>2</sub>, 10 mM β-mercaptoethanol, 15 μM tRNA, 200 μM methionine, and 2 μM MetRS, at 37 °C for 60 min. The acetylation reaction was started by adding a mixture containing 160 μM [acetyl-<sup>3</sup>H] Coenzyme A, 1 μM 3-mercaptopropionic acid, 2 μM MetRS, at 37 °C for 15 min, the cells were harvested and suspended in buffer containing 50 mM NaOAc, pH 5.0, 0.5 mM EDTA, and 1 μM MetRS. The mixture was subjected to an LC/MS analysis using a Q Exactive Hybrid Quadrupole-Orbitrap Mass Spectrometer (Thermo Fisher Scientific) equipped with a Dionex UltiMate 3000 LC System (Thermo Fisher Scientific) and the initial reaction velocities were measured. For analysis of aminoacyl-tRNA specificity of AtaT target in vitro, 10 μM of aminoacyl-tRNA were used for the assay.

**Crystallization and structural determination of AtaT-dimer complexed with acetyl-methionyl-tRNA<sup>Met</sup>**. The purified tRNA<sup>Met</sup> was charged with methionine by MetRS, in a reaction mixture containing 20 mM Tris–HCl, pH 7.4, 150 mM KCl, 7 mM MgCl<sub>2</sub>, 10 mM β-mercaptoethanol, 20 mM L-Asp, 1 μM tRNA<sup>Met</sup>, 1 mM methionine, 2 mM ATP, and 1 μM MetRS, at 37 °C for 1 h. The reaction product was extracted by cold phenol saturated with 300 mM NaOAc, pH 5.2, and applied to a NAP-5 column (GE Healthcare, Japan) for removal of the remaining ATP, and then the eluate was precipitated by ethanol. Met-tRNAs were dissolved in 5 mM NaOAc, pH 5.2, quantified by measuring the A<sub>260</sub> values, and diluted to a series of concentrations. Met-tRNA<sup>Met</sup> was incubated with 20 and 250 μM AtaT, respectively, in a mixture containing 10 mM Tris–HCl, pH 7.0, 100 mM KCl, 5 mM MgCl<sub>2</sub>, 10 mM β-mercaptoethanol, and 160 μM [acetyl-<sup>14</sup>C] Coenzyme A (10 Ci·mol<sup>−1</sup>, Perkin Elmer, Japan), and the initial reaction velocities were measured. For analysis of aminoacyl-tRNA specificity of AtaT target in vitro, 10 μM of aminoacyl-tRNA were used for the assay.

**In vitro acetylation assay.** tRNA was first aminoacylated in a mixture containing 20 mM Tris–HCl, pH 7.4, 150 mM KCl, 7 mM MgCl<sub>2</sub>, 10 mM β-mercaptoethanol, 15 μM tRNA, 200 μM methionine, and 2 μM MetRS, at 37 °C for 60 min. The acetylation reaction was started by adding a mixture containing 160 μM [acetyl-<sup>14</sup>H] Coenzyme A, 1 μM 3-mercaptopropionic acid, 2 μM MetRS, at 37 °C for 15 min, and the eluate was precipitated by ethanol. Methionyl tRNA<sup>Met</sup> was dissolved in buffer containing 400 mM NaCl, 10 mM MgOAc, and 0.5 mM EDTA, and dissolved in buffer containing 400 mM NaCl, 10 mM MgOAc, and 0.5 mM EDTA, and dissolved in buffer containing 400 mM NaCl, 10 mM MgOAc, and 0.5 mM EDTA.

**Preparation of acetylated aminoacyl-tRNAs from E. coli upon AtaT induction.** E. coli strain MG1655 was transformed with pBAD-AtaT or empty pBAD-Myc-His A (Invitrogen) and cultured overnight. The overnight cultures were diluted to an OD<sub>660</sub> of 0.02 in fresh liquid LB (4 mL) containing 50 μg·ml<sup>−1</sup> ampicillin, and cultured at 37 °C until the A<sub>660</sub> reached 0.2. Then, 0.02% (v/v) arabinose was added. After a 15, 30, or 60 min incubation, the cells were harvested and suspended in buffer containing 50 mM NaOAc, pH 5.0, 0.5 mM EDTA, and 0.2 M NaCl. RNA was extracted by phenol saturated with 300 mM NaOAc, pH 5.2, followed by isopropanol precipitation. The RNA was dissolved in 250 μL of 200 mM NaOAc, pH 5.0, acetylated by adding acetic anhydride-D<sub>3</sub> (Sigma Aldrich, Japan), and then ethanol precipitated and rinsed with 70% cold ethanol. The RNA was dissolved in cold buffer containing 50 mM NaOAc, pH 5.0, 0.5 mM EDTA, and 0.2 M NaCl, and incubated for 100 μL of 0.3 M NaOAc, pH 5.0, 0.5 mM EDTA, and 0.2 M NaCl. tRNA was eluted with buffer containing 50 mM NaOAc, pH 5.0, 0.5 mM EDTA, and 0.2 M NaCl, ethanol precipitated, and rinsed with 70% (v/v) ethanol.

**LC/MS spectrometry.** The purified acetylated aminoacyl-tRNA<sup>Met</sup> were digested with Rnase One Ribonuclease (Promega, Japan), in a reaction mixture (25 μL) containing 25 mM NH₄OAc and 2.5 units enzyme, at 37 °C for 60 min. The digests were subjected to an LC/MS analysis using a Q Exactive Hybrid Quadrupole-Orbitrap Mass Spectrometer (Thermo Fisher Scientific) equipped with a Dionex UltiMate 3000 LC System (Thermo Fisher Scientific) and an InertSustain C18 column (5 μm, 2.1 × 250 mm, GL Sciences). Elution was carried out using a multi-linear gradient with 5 mM ammonium acetate (pH 5.3) (solvent A) and 60% acetonitrile (solvent B). The following gradient was used: 1–35% B from 0 to 35 min; 35–99% B from 35 to 40 min; 99% B from 40 to 50 min; 99–1% B from 50 to 50.1 min; and 1% B from 50.1 to 60 min. Ions were scanned by the use of the positive polarity mode over an m/z range of 110–950.<ref>

**Reporting summary.** Further information on research design is available in the Nature Research Reporting Summary linked to this article.

**Data availability** Coordinates and structure factors of AtaT-dimer (PDB ID: 6GTP) complex have been deposited in the Protein Data Bank, under the accession code 7CHD. As the search model, the structures of the AtaT dimer (PDB ID: 6GTP) and Met-tRNA<sup>Met</sup> (PDB ID: 2FMT) were used. All data are available from the corresponding author upon reasonable request. Source data are provided with this paper.

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Author contributions

K.T. planned and designed the research; Y.Y. performed the protein purification, crystallization, structure determination, and biochemical and functional analyses. Y.S. and T.S. assisted with the LC/MS spectrometric analysis. Y.Y. and K.T. analyzed the data and wrote the paper.

Competing interests

The authors declare no competing interests.

Additional information

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