

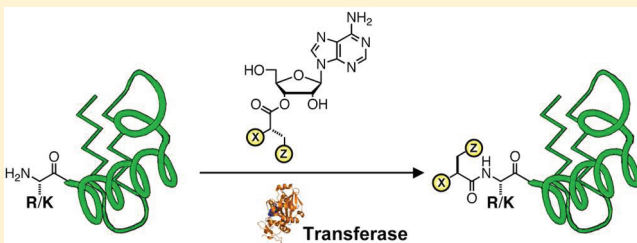
N-Terminal Protein Modification Using Simple Aminoacyl Transferase Substrates

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 Supporting Information

ABSTRACT: Methods for synthetically manipulating protein structure enable greater flexibility in the study of protein function. Previous characterization of the *Escherichia coli* aminoacyl tRNA transferase (AaT) has shown that it can modify the N-terminus of a protein with an amino acid from a tRNA or a synthetic oligonucleotide donor. Here, we demonstrate that AaT can efficiently use a minimal adenosine substrate, which can be synthesized in one to two steps from readily available starting materials. We have characterized the enzymatic activity of AaT with aminoacyl adenosyl donors and found that reaction products do not inhibit AaT. The use of adenosyl donors removes the substrate limitations imposed by the use of synthetases for tRNA charging and avoids the complex synthesis of an oligonucleotide donor. Thus, our AaT donors increase the potential substrate scope and reaction scale for N-terminal protein modification under conditions that maintain folding.



INTRODUCTION

The conjugation of synthetic molecules to proteins contributes to biomedical research by enabling the immobilization of proteins on surfaces, modifications with chromophores to make *in vitro* sensors, or tagging with *in vivo* imaging agents.^{1–5} In most cases, the formation of well-defined conjugates is valuable, if not essential. The protein termini are attractive targets for conjugation, because in many cases appending synthetic molecules at the termini will have minimal undesired effects on protein folding and function.^{6–26} This is particularly applicable in cases where the terminus is relatively unstructured. As part of our research on minimalist labeling strategies for biophysical studies, we sought a method for N-terminal modification that could be carried out easily under conditions that maintain folding without substantial prior protein manipulation.²⁷

Selective N-terminal modification has been achieved by a variety of chemical and enzymatic methods, each with benefits and drawbacks. Small molecule strategies permit the attachment of a variety of molecules, but they are subject to side reactions and incomplete specificity for the N-terminus and may need to be carried out in organic solvent mixtures.^{16,28–37} Reverse proteolysis methods can be used to modify the N-terminus under conditions that do not require protein unfolding, but the reaction can be difficult to drive to completion without high protein concentrations.^{38–51} Several other chemoenzymatic methods are valuable in functioning under benign conditions but have moderate-sized target sequences that must be appended to the protein.^{52–59} Here, we describe a minimal system for N-terminal protein labeling that utilizes only adenosine esters of natural or

unnatural amino acids and a single, readily available enzyme. This process gives high yields of modified proteins under nondenaturing conditions and requires only a single basic amino acid for specific recognition.

Aminoacyl tRNA transferases (AaTs) are members of a growing class of enzymes that use aminoacyl tRNAs in secondary metabolism.⁶⁰ The *E. coli* AaT catalyzes the transfer of Leu, Phe, or Met from an aminoacyl tRNA to a protein bearing an N-terminal Arg or Lys.^{61–63} The addition of Leu or Phe targets that protein for degradation by ClpA as part of the N-end rule pathway.^{64–67} Kaji et al. first observed AaT aminoacylation activity in crude *E. coli* preparations.^{61,68,69} Soffer and Leibowitz subsequently reconstituted the purified enzyme and characterized its specificity for both the RNA and amino acid components of the donor molecule, demonstrating its first use in transferring an unnatural amino acid, *p*-fluorophenylalanine^{70,71} (Figure 1, top).

Recently, purified AaT has been used to modify proteins *in vitro* with a variety of unnatural amino acids charged onto tRNAs, either by chemical semisynthesis or the use of a mutant aminoacyl tRNA synthetase (aaRS).^{19,72–75} Abramochkin and Shrader discovered that AaT was tolerant of variation in the acceptor stem (aminoacylation site) of the tRNA, and Sisido and co-workers have shown that much shorter oligonucleotides (Figure 1, bottom, $R_n = 2–22$ nucleotides) can act as donors.^{74,76,77} As part of their structural characterization of AaT, Tomita and co-workers found that AaT could bind phenylalanyl adenosine and

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transfer Phe to peptides in trace amounts.⁶³ However, the use of adenosine mononucleoside as a donor substrate for AaT has not, to our knowledge, been further explored.

We have found that a variety of hydrophobic amino acids can be transferred from adenosine donors with high yields at relatively low protein concentrations. (Figure 1, bottom, $R_n = H$) These donors can be synthesized from commercially available active esters of the amino acids. Reduction of the two step synthetase/AaT reaction sequence to the one step AaT-only sequence allows us to explore the substrate specificity of AaT without limitation by the specificity of the synthetase in the first step. While this is also possible with Sisido's pdCpA dinucleotide donors, the shorter synthesis of the simple adenosine donors (1–2 steps vs 7 steps) makes this exploration easier. We have characterized the substrate scope of the wild type AaT enzyme using a reporter peptide, determined the kinetic parameters for

the adenosyl substrates, and demonstrated the efficient modification of a full-sized protein, 23 kDa α -casein.

RESULTS AND DISCUSSION

The synthesis of adenosyl donor compounds, shown in Scheme 1, can be carried out on any amino acid with appropriate acid-labile protecting groups (e.g., *N*-Boc). This synthesis is not limited to α -amino acids, and our donor synthesis has also enabled us to test the transfer of analogs such as α -azidophenylalanine (**N₃f**, **4g**). Conversion to the cyanomethyl ester (**1a–h**) is typically high-yielding, although yields for the subsequent acylation of 5'-*O*-dimethoxytrityl-adenosine ((DMT)-A) varied between 60 and 90%. Acylation yields generally decreased with side chain bulk; derivatives such as *N*-1,8-naphthyldiaminopropionic acid coupled poorly (data not shown). Uncatalyzed reactions took 1–5 days, but in all cases, addition of tetrabutylammonium acetate increased reaction rates.

The Phe donor **4a** was synthesized from both the cyanomethyl ester (**1a**) and *N*-hydroxysuccinimidyl (OSu) ester (**2a**). All other adenosyl donors were synthesized from the cyanomethyl ester (**1b–h**). After trifluoroacetic acid (TFA) deprotection and CH_2Cl_2 washing (or Et_2O precipitation) to remove DMT byproducts, the compounds could be taken on directly to AaT enzymatic reactions (contaminants were limited to incompletely deprotected compounds which are not transferred by AaT, see Supporting Information). However, in most cases the deprotected adenosyl donor was purified by HPLC, and higher transfer yields were generally observed with these purified donors.

Transfer extent was evaluated by HPLC analysis of reactions with adenosyl donor, AaT, and an aminocoumarin-labeled reporter peptide used previously by Tirrell and co-workers (LysAlaAcm, Scheme 1).⁷⁸ Monitoring LysAlaAcm acylation in crude reaction mixtures using absorbance at 325 nm conveniently eliminates background due to other proteins and nucleic acids, although some unnatural amino acids such as naphthylalanine (Nap) and methoxycoumarinylalanine (Mcm) also absorb in this range. Example HPLC chromatograms from Phe ligations are shown in Figure 2. LysAlaAcm starting material eluted at 11.8 min (Figure 2A); hydrophobic products eluted at 12 to

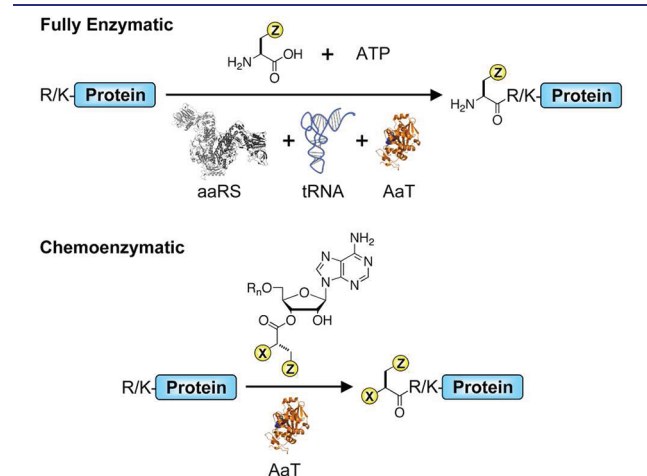
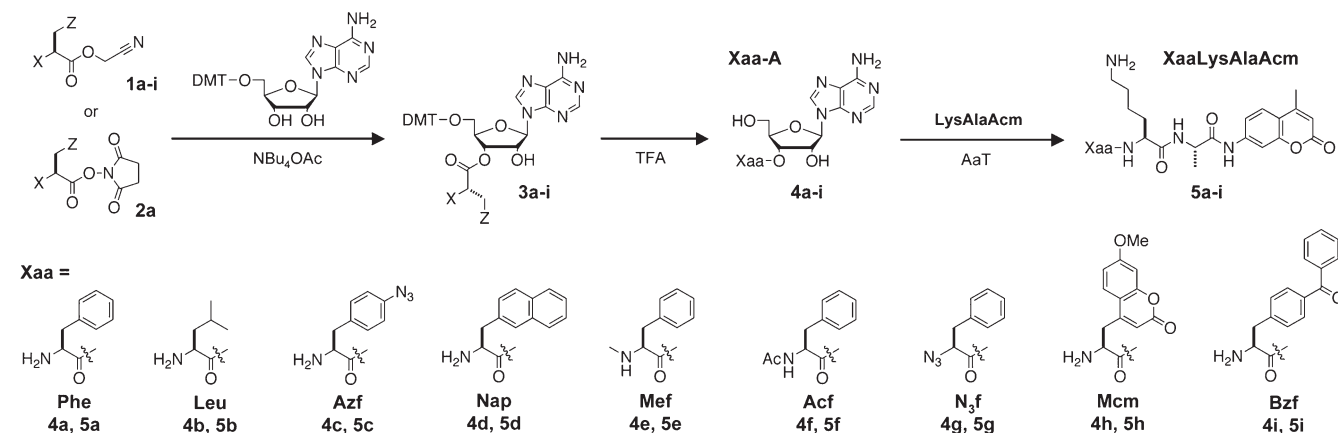


Figure 1. Transferase-mediated N-terminal protein modification. (Top) Fully enzymatic methods use aminoacyl tRNA synthetase (aaRS), tRNA, aminoacyl transferase (AaT), amino acid, and ATP. (Bottom) Chemoenzymatic methods use only a synthetic nucleic acid donor and transferase. In this work, an aminoacyl mononucleoside ($R_n = H$) was used.

Scheme 1. Synthesis of Adenosyl Donors and Use in AaT-Catalyzed Protein Modification^a



^a Donor synthetic intermediates: **1a**, **2a**, **3a** $X = \text{NH}(\text{Boc})$, $Z = \text{Ph}$; **1b**, **3b** $X = \text{NH}(\text{Boc})$, $Z = i\text{-Pr}$; **1c**, **3c** $X = \text{NH}(\text{Boc})$, $Z = p\text{-Azidophenyl}$; **1d**, **3d** $X = \text{NH}(\text{Boc})$, $Z = 2\text{-naphthyl}$; **1e**, **3e** $X = \text{NH}(\text{MeBoc})$, $Z = \text{Ph}$; **1f**, **3f** $X = \text{NH}(\text{Ac})$, $Z = \text{Ph}$; **1g**, **3g** $X = \text{N}_3$, $Z = \text{Ph}$; **1h**, **3h** $X = \text{NH}(\text{Boc})$, $Z = 7\text{-Methoxycoumarinyl}$; **1i**, **3i** $X = \text{NH}(\text{MeBoc})$, $Z = p\text{-Benzoylphenyl}$

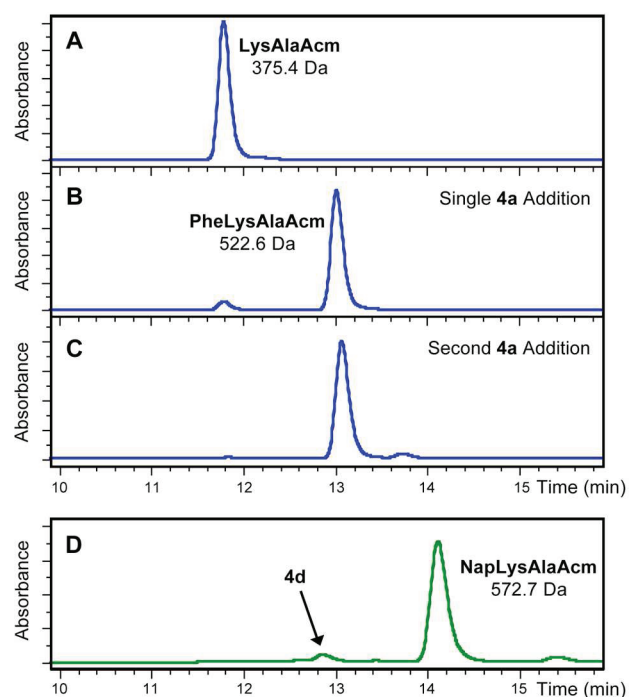


Figure 2. Reversed-phase HPLC Analysis of Transferase Reactions. Top: AaT-mediated transfer of Phe from adenosyl donor **4a** to LysAlaAcm reporter peptide. HPLC chromatograms obtained after (A) 0 h or (B) 4 h show conversion of LysAlaAcm (11.8 min retention time) to PheLysAlaAcm (13.0 min retention time). Conversion after one addition of donor **4a** is 92% (B). A second addition of 1 mM **4a** drives the reaction to completion with a small amount of double Phe addition (C). Bottom: HPLC analysis of transfer of Nap from adenosyl donor **4d** to LysAlaAcm after (D) 4 h shows conversion to NapLysAlaAcm (14.1 min retention time). Product identities confirmed by observation of indicated masses by MALDI MS.

15 min (PheLysAlaAcm **5a** elutes at 13.0 min in Figure 2B, NapLysAlaAcm **5d** elutes at 14.1 min in Figure 2D).

To evaluate substrate scope, reactions were analyzed after 4 h, when transfer should be complete for the fully enzymatic reaction with Phe tRNA synthetase.⁷⁸ Phe (**4a**) and Leu (**4b**) were transferred efficiently. (Table 1) Our PheLysAlaAcm yields from donor **4a** were comparable to yields from Phe-pdCpA, the dinucleotide used by Sisido. Transfer yields for other substrates varied with side chain size, where Mcm (**4h**) is too large and benzoylphenylalanine (Bzf, **4i**) is a poor substrate (see Table 1). Sisido has shown that AaT mutants can use substrates with larger side chains; we are currently exploring some of these mutations.⁷⁴ While transfer yields after 4 h varied, most reactions could be driven to completion by a bolus of donor molecule. (Figure 2C)

One of the advantages of adenosyl donors over *in situ* tRNA aminoacylation with a tRNA synthetase is that we are not limited by the substrate specificity of the synthetase. In particular, nonamino acid substrates can be attached to adenosyl donors to assess their transferability by AaT. *N*-Methyl phenylalanine (Mef, **4e**) can be transferred, albeit with low yield. We have also tested *N*-acetyl phenylalanine (Acf, **4f**) and α -azidophenylalanine (**4g**) and found them to be poor substrates. In some experiments, substantial conversion of LysAlaAcm was found when incubated with **4g**, but MALDI MS analysis has shown the product to be **5a**, presumably obtained by transfer of Phe after prior reduction (i.e., Phe donor **4a** is formed from **4g** under the

Table 1. N-Terminal Modification of Reporter Peptide After Single Addition of Adenosyl Donor

donor	additives ^a	yield (%) ^b
4a (Phe)		92.7 \pm 5.2
Phe-pdCpA		95.2 \pm 4.3
4a (Phe)	1 mM ATP	86.1 \pm 0.8
4a (Phe)	5 mM ATP	79.1 \pm 0.4
4a (Phe)	5 mM AMP	88.2 \pm 0.8
4a (Phe)	5 mM A	93.1 \pm 1.1
4a (Phe)	1 mM pdCpA	90.4 \pm 0.9
4b (Leu)		80.4 \pm 0.9
4c (Azf)		78.5 \pm 5.2
4d (Nap)		95.4 \pm 0.4
4d (Nap)	100 μ M PheLysAlaAcm	93.8 \pm 0.3
4e (Mef)		0.6 \pm 0.1
4f (Acf)		5f not observed
4g (N ₃ f)		5g not observed ^c
4h (Mcm)		5h not observed
4i (Bzf)		9.2 \pm 0.3

^a Standard AaT-catalyzed transfer conditions with or without potential inhibitor added. See description under Experimental Procedures. ^b Yield determined by integration of HPLC chromatogram peak areas at 325 nm, described in Experimental Procedures. Standard deviations are reported for an average of at least 6 experiments with at least 2 different AaT preparations. ^c HPLC and MALDI MS analysis indicate that the observed product is **5a** not **5g**.

reaction conditions). Steps taken to alter the concentration of β -mercaptoethanol reductant did not improve the yield of **5g** (data not shown). We believe that disproportionation of the azide is incomplete, but that AaT is selective for the α -amine substrate (Phe-A, **4a**), so far more **5a** than **5g** is formed. This may be due to an important interaction of the α -amine with Q188, which also acts as a catalytic base to deprotonate the peptide substrate for amide bond formation. The Q188/ α -amine interaction was defined by the structural and enzymological work of Watanabe and co-workers.⁶³ We note that the preference of AaT for unmodified N-termini was previously documented with full-length tRNA by Soffer and Leibowitz.⁷¹

To better understand limitations on the yield from AaT reactions, we assessed the possibilities of inhibition by hydrolyzed substrate or aminoacylated product. Since the acyl adenosine donor substrate is not regenerated during the reaction (unlike the aaRS/tRNA/AaT reaction), we hypothesized that the adenosine byproduct might inhibit further acylation by binding to AaT. To investigate this possibility, we added adenosine to the reaction at varying concentrations and monitored the reaction timecourse by fluorescence spectroscopy and by HPLC. Addition from the Phe donor **4a** was not inhibited by the addition of 5 mM concentrations of adenosine (Table 1). A comparison of adenosine compounds, including pdCpA, the dinucleotide donor used by Sisido, showed that only adenosine triphosphate (ATP) inhibited the reaction.

We were also able to monitor product formation in real time based on partial quenching of coumarin fluorescence upon addition of Nap to the LysAlaAcm peptide to form **5d**. The fluorescence intensity at 390 nm of NapLysAlaAcm is 28% of the fluorescence of LysAlaAcm, and the overall fluorescence of mixtures of the two peptides can be used to determine the proportions of each peptide if the total concentration is known

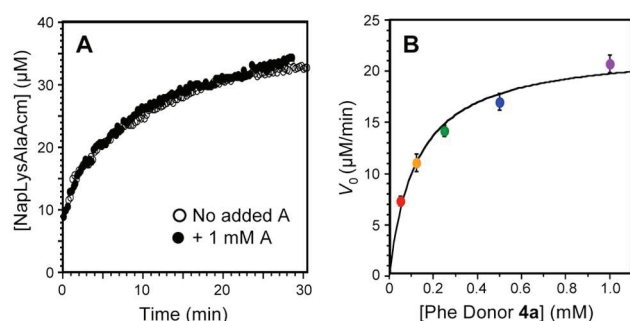


Figure 3. Kinetic analysis of AaT reactions. (A) Real time monitoring of Nap transfer to LysAlaAcm from **4d** by quenching of Acm fluorescence. No significant inhibition is observed in the presence of 1 mM adenosine. (B) Saturation curve used to determine Michaelis–Menten kinetic parameters for LysAlaAcm modification by Phe donor **4a**.

(see Supporting Information). Thus, the change in fluorescence intensity was used to monitor NapLysAlaAcm formation in real time, and HPLC injection of the reaction end points were used to confirm the final product distribution. We found that exogenous adenosine did not significantly inhibit the reaction, even at concentrations up to 1 mM (Figure 3A).

We also assessed the possibility of product inhibition by the aminoacylated LysAlaAcm product. In our standard 4 h transfer assay, addition of 100 μ M (1 equiv) PheLysAlaAcm did not significantly inhibit production of NapLysAlaAcm (**5d**) when Nap-A (**4d**) was used as a donor. (Table 1) MALDI MS analysis and differences in HPLC retention time allowed us to easily distinguish LysAlaAcm and PheLysAlaAcm from NapLysAlaAcm (see Figure 2 and Supporting Information). Finally, we investigated the question of whether donor hydrolysis slowed reaction rates after 30–60 min. Analysis of mock reactions lacking transferase or LysAlaAcm indicates that donor hydrolysis may contribute significantly to slowing reactions, which is why a donor bolus can be used to drive the reaction to completion (see Supporting Information).

To characterize the enzymatic activity of AaT toward our acyl adenosyl substrates, we used our HPLC assay to determine initial reaction rates to be fit to a standard Michaelis–Menten kinetic model. Equations describing this kinetic analysis are included in the Supporting Information. Since adenosine does not inhibit the enzyme, by keeping LysAlaAcm concentrations high (100 μ M), we measured the rate of transfer as a function of the concentration of Phe donor **4a** and fit this to a single-substrate kinetic model to determine a k_{cat} of $1.68 \pm 0.09 \times 10^{-1} \text{ s}^{-1}$ and a K_M of $1.24 \pm 0.23 \times 10^{-4} \text{ M}$. The enzyme efficiency ($k_{\text{cat}}/K_M = 1.35 \times 10^3 \text{ M}^{-1} \text{ s}^{-1}$) is relatively low.⁷⁹ While K_M is higher for Phe-A than what Shrader reported for aminoacyl tRNA^{Leu-4} (0.3 μ M), the k_{cat} is comparable ($0.13 \times 10^{-1} \text{ s}^{-1}$).⁷⁷ Since it is trivial to run our reactions at donor concentrations well above K_M , the lower apparent affinity for our substrates should not affect the utility of the reaction. We note that the binding affinity, which gives a K_M of 124 μ M for **4a** must arise primarily from Phe binding interactions since the K_I for adenosine inhibition is greater than 5 mM.

It is possible that larger protein substrates are modified with different efficiencies than peptide substrates.^{71,74,77} To address this question and to demonstrate the utility of our method in tagging full-sized proteins with useful fluorophores or affinity purification tags, we modified the milk protein α -casein (bearing

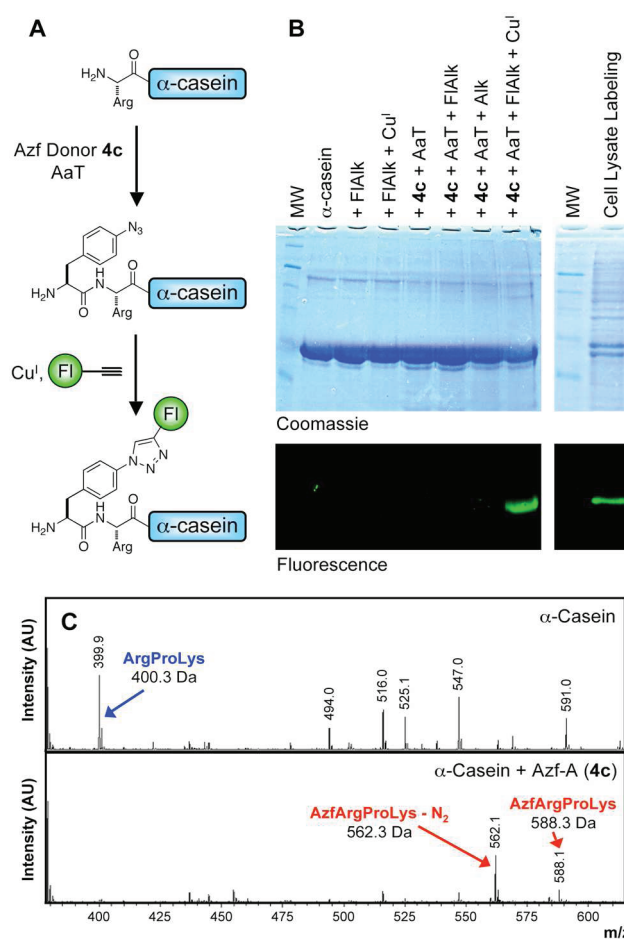


Figure 4. AaT-Catalyzed Modification and “Click” Reaction of α -Casein N-terminus. (A) α -Casein modification scheme. (B) PAGE gel analysis of α -casein modification. Lanes (left to right): (1) molecular weight (MW) markers (Masses in kDa: 17, 25, 30, 46, 58, 80, 175); (2) α -casein; (3) α -casein mixed with fluorescein alkyne (FlAlk); (4) α -casein mixed with FlAlk, CuSO_4 , THPTA, and sodium ascorbate; (5) α -casein mixed with Azf-A (**4c**) and AaT; (6) α -casein mixed with **4c** and AaT, then FlAlk; (7) α -casein mixed with **4c** and AaT, then propargylamine (Alk), CuSO_4 , Tris-(3-hydroxypropyl)triazolylmethyl amine (THPTA), and sodium ascorbate; (8) α -casein mixed with **4c** and AaT, then FlAlk, CuSO_4 , THPTA, and sodium ascorbate; (9) MW markers; and (10) cell lysate labeling, α -casein reaction carried out using conditions of lane 8 with unpurified AaT in cleared *E. coli* lysate. (C) MALDI MS analysis of trypsinized N-terminal fragment of α -casein with or without modification by Azf using AaT and **4c** (double addition).

a native N-terminal Arg after proteolysis of a leader sequence).⁸⁰ First, modification using Phe donor **4a** was assessed by Edman degradation, where successful, quantitative addition of Phe was observed (see Supporting Information).

In a more complex set of experiments, we first modified α -casein with Azf using donor **4c** and then used a Cu-catalyzed Huisgen cycloaddition (“click” reaction) to append a fluorescein label^{81–84} (Figure 4A). Prior to fluorescent-labeling, the initial transfer of Azf was analyzed by trypsin digest and MALDI MS of α -casein (Figure 4C). The N-terminal peptide ArgProLys disappeared from the mass spectrum following modification and a peptide corresponding to AzfArgProLys was observed (also a mass corresponding to AzfArgProLys after loss of N_2 during ionization; N_2 loss was also observed for AzfLysAlaAcm).

Complete conversion of α -casein was observed in MALDI MS due to double addition of **4c** to drive the reaction.

After AaT-catalyzed modification of α -casein with Azf, residual **4c** was removed by dialysis, AaT (His₁₀-tagged) was removed by treatment with Ni beads, and the azide-bearing protein was reacted with fluorescein alkyne (FlAlk) in the presence of Cu(I). The results of this labeling experiment, as well as appropriate control reactions, are shown in Figure 4B. One can see that fluorescent α -casein is only observed under the proper conditions (Lane 8), where both the chemoenzymatic labeling by AaT with **4c** and the subsequent click reaction are successful. Finally, we carried out a similar labeling experiment in crude *E. coli* lysate with unpurified AaT. Azf transfer was successful, as shown by the observation of fluorescence after labeling with FlAlk. Control reactions (shown in Supporting Information) show no observable labeling of any protein, implying that endogenous levels of proteins terminating in Arg or Lys are too low to be observed here.

CONCLUSION

Substantial recent effort has been devoted to developing methods for labeling proteins under conditions that maintain protein folding and activity.^{85–87} *E. coli* aminoacyl tRNA transferase can work under these conditions and specifically modifies the N-terminus, a useful point for conjugation. However, certain aspects of its native mechanism are nonideal for its use as a preparative tool (i.e., a multistep, multienzyme reaction sequence). The work of Leibowitz, Shrader, and Sisido showed that *E. coli* aminoacyl tRNA transferase can modify the N-terminus of a protein ending in Arg or Lys with amino acids from an aminoacyl oligonucleotide donor.^{71,74,77} Watanabe et al. crystallized AaT with Phe-A bound and showed that Phe could be transferred from Phe-A to the N-terminus of the peptide RYLGYL in trace amounts. These two results provide crucial precedent for our own work, in which we show that AaT can efficiently use a variety of easily synthesized acyl adenosine substrates. Enzymological analysis demonstrates that AaT is able to use the Xaa-A substrates with turnover numbers comparable to full-length aminoacyl tRNAs. Furthermore, we have shown that neither the adenosine reaction byproduct nor aminoacylated protein product (at least in the case of our reporter peptide) appreciably inhibit the reaction. In fact, the major obstacle to efficient transfer is donor hydrolysis, but this can be overcome by a second addition of donor molecule since the K_i for inhibition by adenosine is greater than 5 mM.

Our AaT-only reaction sequence allows us to explore the compatibility of Phe analogs lacking a primary α -amine with transfer by AaT. The wild type transferase seems to have a strong preference for the α -amine, as only *N*-methylphenylalanine was transferred. In contrast, the side chain pocket is quite permissive: we have observed the transfer of bicyclic amino acid side chains and Sisido has reported mutants capable of transferring tricyclic amino acids.⁷⁴ We are currently investigating the transfer of larger side chains such as benzophenone, biotin, and fluorescein derivatives.

Our reaction conditions and yields compare favorably to two well-established methods in the field. Subtiligase (reverse proteolysis) reactions can be carried out in a few hours, but rarely go to completion.^{44,45,51} The pyridoxyl-5-phosphate (PLP) transamination reactions of Francis and co-workers can require elevated temperatures to achieve high yields and are subject to a number of side reactions.^{36,37} AaT transfer reactions are highly

specific (no Lys side chain modifications were observed in our work or the work of others) and can be driven to completion by subsequent repeated additions of donor. Our discovery of efficient transfer from simple adenosyl donors removes the limitations imposed by the selectivity of synthetases on the range of molecules that can be AaT substrates without restricting reaction scale by necessitating the synthesis of oligonucleotide donors. Our yields are comparable to those obtained with pdCpA donors, but pdCpA synthesis requires 6 steps, and overall yields are typically 25%. We see our methods as complementary to the fully enzymatic synthetase-based methods (Figure 1, Top). Fully enzymatic methods do not require prior synthesis and aminoacyl tRNAs are efficient substrates for AaT. On the other hand, our methods can be easily scaled up to modify larger quantities of protein since they do not require the use of purified tRNA.

We have shown that our method can be used to transfer reactive handles to the N-terminus under “protein friendly” (pH \sim 7, high salt, 37 °C) conditions and that the labeled protein product can be easily purified afterward. Given the previous success of Sisido and co-workers, we expect that further mutation of AaT will allow us to transfer larger side chains that may permit direct transfer of fluorescent probes or affinity tags. We also hope to use the available crystal structures to redesign AaT to act on other N-terminal sequences through a combination of rational design and selection. Finally, our demonstration that AaT can modify α -casein in crude lysates, coupled with the cell-permeability of our moderately polar donors raises the exciting possibility of using AaT as part of an *in vivo* labeling strategy.

EXPERIMENTAL PROCEDURES

General Information. Chloroacetonitrile, *N,N*-diisopropylethylamine (DIPEA), 5'-O-(4,4'-Dimethoxytrityl) adenosine ((DMT)-A), tetrabutylammonium acetate (TBAAC), trifluoroacetic acid (TFA), and triisopropyl silane (TIPSH) were purchased from Sigma-Aldrich (St. Louis, MO). Note: (DMT)-A production was discontinued by Sigma-Aldrich, after which it was purchased as a custom order from ChemGenes Corporation (Wilmington, MA) and was additionally synthesized in-house following the protocol outlined by Ogilvie et al.⁸⁸ *N*-Boc-L-phenylalanine-O-succinimide (Boc-Phe-OSu) and Lysylalanylaminomethylcoumarin (LysAlaAcm) were purchased from Bachem (Torrance, CA). *N*-Boc-L-phenylalanine (Boc-Phe-OH) and all solvents were purchased from Fisher Scientific (Pittsburgh, PA). All deuterated solvents were purchased from Cambridge Isotopes Laboratories, Inc. (Andover, MA). *E. coli* BL21(DE3) cells were purchased from Stratagene (La Jolla, CA). The pEG6 plasmid, containing His₁₀-tagged *E. coli* AaT, was a gift from Alexander Varshavsky (California Institute of Technology). Sequencing-grade trypsin was purchased from Promega (Madison, WI). All other reagents were purchased from Fisher Scientific (Pittsburgh, PA).

Milli-Q filtered (18 M Ω) water was used for all aqueous solutions (Millipore; Billerica, MA). Matrix-assisted laser desorption/ionization (MALDI) mass spectra were collected using a Bruker Ultraflex III MALDI-TOF-TOF mass spectrometer (Billerica, MA). UV absorbance spectra were obtained with a Hewlett-Packard 8452A diode array spectrophotometer (currently Agilent Technologies; Santa Clara, CA). Donor molecule purification was conducted on a BioCad Sprint FPLC (GMI Inc.; Ramsey, MN; originally from Perseptive Biosystems) with a Waters Sunfire Prep C18-prep OBD column, 5 μ m, 17 \times 150 mm (Milford, MA). Analytical HPLC assays were performed on an Agilent 1100 HPLC using a Waters Symmetry Shield C18 column. NMR spectra, ¹H and ¹³C, were collected with a Bruker DRX 500 MHz instrument. “Low resolution” electrospray ionization (ESI) mass spectra

(LRMS) were obtained on a Waters Acquity Ultra Performance LC connected to a single quadrupole detector (SQD) mass spectrometer. "High resolution" ESI mass spectra (HRMS) were obtained on a Waters LCT Premier XE LC/MS. DNA sequencing was performed at the University of Pennsylvania DNA sequencing facility. Fluorescence spectra were collected with a Varian Cary Eclipse fluorescence spectrophotometer fitted with a Peltier multicell holder (currently Agilent Technologies).

Donor (4a–h) Synthesis. The synthesis of Phe donor **4a** is given as a general procedure, starting from commercially available Boc-Phe-OH or Boc-Phe-OSu. The carboxy-terminus of the amino acid was activated as a cyanomethyl ester using chloroacetonitrile to give **1a**. Next, the protected amino acid was attached to one of the available hydroxyls (2' OH or 3' OH) of (DMT)-A without preference. The resulting protected adenylate (**3a**) was deprotected using a 50/50 mixture of TFA and THF with TIPSH present as a scavenger. The final product, deprotected adenylate (**4a**), was purified on C18 reverse-phase HPLC column. The synthesis and characterization of donor **4a** is provided below, characterization of other donor compounds (**4b–h**) and precursors is provided in the Supporting Information.

(S)-Cyanomethyl 2-((tert-butoxycarbonyl)amino)-3-phenylpropanoate (Boc-Phe-OCH₂CN, 1a). Boc-Phe-OH (2.01 g, 7.57 mmol) was dissolved in 10 mL tetrahydrofuran under ambient conditions. Ten equivalents of chloroacetonitrile (4.78 mL, 75.4 mmol) and 1.1 equiv of DIPEA (8.30 mmol, 1.45 mL) were added to the reaction mixture and allowed to stir overnight. **1a** was purified on silica gel (20% EtOAc in hexanes) to give 2.474 g (100%) of a pale-yellow oil after evaporation. *R*_f 0.4 in 20% EtOAc in hexanes; ¹H NMR (500 MHz, CDCl₃) δ 7.26 (t, *J* = 7.0 Hz, 2H), 7.20 (t, *J* = 7.1 Hz, 1H), 7.11 (d, *J* = 7.3 Hz, 2H), 5.11 (d, *J* = 7.8 Hz, 1H), 4.69–4.57 (m, 3H), 3.09–2.98 (m, 2H), 1.36 (s, 9H); ¹³C NMR (125 MHz, CDCl₃) δ 170.8, 155.1, 135.4, 129.2, 128.7, 127.3, 114.0, 80.9, 80.2, 54.4, 48.8, 37.8, 28.2; HRMS (ESI) calcd *m/z* for C₁₆H₂₀N₂O₄Na [M + Na]⁺ 327.132, found 327.133.

(S)-(2R,3R,4R,5R)-2-(6-Amino-9H-purin-9-yl)-5-((bis(4-methoxyphenyl)(phenyl)methoxy) methyl)-4-hydroxytetrahydrofuran-3-yl 2-((tert-butoxycarbonyl)amino)-3-phenylpropanoate (Boc-Phe-(DMT)-A, 3a). (DMT)-A (25 mg, 44 μmol) was dissolved in 5 mL of tetrahydrofuran (dried with 4 Å molecular sieves) in an oven-dried 4 dram reaction vessel. Twenty equivalents of **1a** (272 mg, 0.895 mmol) were added to the reaction mixture. TBAAC (4.9 mg, 16 μmol), the reaction catalyst, was added last to the reaction mixture. In reproductions of this reaction, TBAAC additions varied according to reaction progress. The reaction stirred under argon overnight at room temperature for 1–4 days. After evaporation of solvent, the product was purified on silica gel (gradient solvent system: 20–0% petroleum ether in EtOAc, 5% methanol in EtOAc) to give 27 mg (75%) of a pale-yellow foam after evaporation. *R*_f 0.5 in 5% methanol in EtOAc. ¹H NMR (500 MHz, THF-*d*₈) δ 8.08 (s, 1H), 8.00 (s, 1H), 7.44 (d, *J* = 3.68 Hz, 2H), 7.32 (d, *J* = 8.85 Hz, 4H), 7.20 (dd, *J* = 8.33 Hz, *J* = 6.66, 2H), 7.13 (t, *J* = 7.25 Hz, 1H), 7.21–7.10 (m, 4H), 6.77 (d, *J* = 4.30 Hz, 2H), 6.75 (d, *J* = 4.29 Hz, 2H), 5.92 (d, *J* = 4.87 Hz, 1H), 5.49 (t, *J* = 4.34 Hz, 1H), 5.13 (t, *J* = 5.54 Hz, 1H), 4.51–4.47 (m, 1H), 4.14 (t, *J* = 4.07 Hz, 1H), 3.71 (s, 6H) (S), 3.42–3.32 (m, 2H), 3.12–2.96 (m, 3H), 1.37 (s, 9H); ¹³C NMR (125 MHz, THF-*d*₈) δ 172.2, 159.9, 157.6, 156.7, 153.8, 146.3, 138.3, 146.3, 138.3, 136.7, 136.9, 131.2, 131.1, 130.3, 129.3, 129.2, 129.1, 128.6, 128.5, 127.5, 114.0, 113.9, 89.7, 87.5, 82.5, 79.7, 75.1, 73.5, 64.7, 56.4, 55.5, 38.9, 30.6, 28.8; LRMS (ESI) calcd *m/z* for C₄₅H₄₈N₆O₉H (M + H)⁺ 816.9, C₄₅H₄₈N₆O₉Na (M + Na)⁺ 839.3, found 817.3, 839.3.

(S)-(2R,3R,4R,5R)-2-(6-Amino-9H-purin-9-yl)-5-((bis(4-methoxyphenyl)(phenyl)methoxy) methyl)-4-hydroxytetrahydrofuran-3-yl 2-((tert-butoxycarbonyl)amino)-3-phenylpropanoate (3a) from 2a. (DMT)-A (26 mg, 45 μmol) was dissolved in 5 mL of tetrahydrofuran (dried with 4 Å molecular sieves) in an oven-dried 4-dram reaction vessel. Twenty equivalents of **2a** (321 mg, 0.885 mmol) were

added to the reaction mixture. TBAAC (19 mg, 64 μmol), the reaction catalyst, was added last to the reaction mixture in two separate additions, 6 mg, at the beginning of the reaction, and 13 mg after the first 24 h of reaction. The reaction stirred under argon overnight at room temperature for 2 days. After evaporation of solvent, the product was purified on silica gel (gradient solvent system: 20–0% petroleum ether in EtOAc, 5% methanol in EtOAc) to give 19.1 mg (52.1%) of a pale-yellow foam after evaporation. LRMS (ESI) calcd *m/z* for C₄₅H₄₈N₆O₉H (M + H)⁺ 816.9, C₄₅H₄₈N₆O₉Na (M + Na)⁺ 839.3, found 817.4, 839.4.

(S)-(2R,3R,4R,5R)-2-(6-Amino-9H-purin-9-yl)-4-hydroxy-5-(hydroxymethyl)tetrahydrofuran-3-yl 2-amino-3-phenylpropanoate (Phe-A, 4a). **3a** (50 mg) was dissolved in 1 mL of THF and 1 mL trifluoroacetic acid (TFA) and reacted overnight with four equiv of TIPSH. Upon addition of TFA, the reaction turned a bright-orange color and at the completion of the reaction, the color had become a dark brown. The reaction mixture was reduced to dryness by rotary evaporation and extracted using 1 mL dichloromethane and 1 mL water twice with a 1 mL water final back-extraction against the dichloromethane layer. The water-soluble layer containing **4a** was then HPLC-purified on a C18-prep column using an increasing acetonitrile gradient (Gradient 2: 1–100% acetonitrile with water and 0.1% TFA, 1% per min) to remove any protected material. Prep. HPLC purified to give 13.6 mg (53.5%) of a white solid after evaporation. MALDI MS calcd *m/z* for C₁₉H₂₂N₆O₅H (M + H)⁺ 415.4, C₁₉H₂₂N₆O₅Na (M + Na)⁺ 437.4, found 415.0, 437.0.

LysAlaAcm Ligation Assay. Each ligation reaction, 125 μL total volume, contained the following reagents: aminoacyl adenosine donor (1 mM), recombinant His₁₀-tagged *E. coli* aminoacyl transferase (2.26 μM), and LysAlaAcm (100 μM) in the AaT Ligation Buffer (50 mM HEPES pH 8.0, 150 mM KCl, 10 mM MgCl₂). The reaction mixtures were incubated at 37 °C for four hours and quenched with 1% acetic acid. The proteins were extracted from the reactions via acetone precipitation. The reactions were precipitated using 4X reaction volume of acetone and cooled at –20 °C for 1 h. Next, the reactions were centrifuged at 13, 200 rpm at 4 °C for 20 min to separate the reaction from precipitated protein. The supernatant was transferred to fresh 1.5 mL centrifuge tubes and allowed for acetone evaporation overnight at room temperature. After acetone evaporation, the supernatant was dried in a Speedvac (Savant, Thermo Scientific, Fisher Inc.) for 30 min to remove residual acetone. The resulting reaction volume was dissolved up to 1.2 mL using Milli-Q water and analyzed by HPLC (gradient below) to determine ligation yield by integration of separated peak intensities monitored at 325 nm. Collected HPLC fractions were characterized through MALDI MS analysis. Reactions were performed with 6 trials on at least two protein preparations for all successful reactions, 3 trials for failed reactions. Reactions using Phe-pdCpA as donor were carried out in an identical fashion.⁸⁹

A/AMP/ATP/pdCpA Inhibition Studies. Each ligation reaction was performed as described above. The addition of four types of adenosyl compounds (adenosine, AMP, ATP, or pdCpA) were monitored individually in ligation reactions with Phe donor **4a**. Adenosyl compound concentrations tested were 1 mM, 2.5 mM, and 5 mM. HPLC analysis was used to determine inhibition of ligation by integration of HPLC reagent and product peaks. The LysAlaAcm ligation assay was used to analyze the kinetics of AaT with substrate **4a**. The ligation assay was modified for ease of analysis as follows: The total reaction volume was scaled up to 590 μL and all reagents were maintained at the same concentrations as mentioned above. Five concentrations of **4a** were monitored for a total reaction time of 30 min, with concentrations ranging from 0.05 to 1 mM.

Each 30 min kinetic experiment was done in triplicate with a total of fourteen 40 μL aliquots for all substrate concentrations. The synthesis of pdCpA was carried out essentially as previously described.⁸⁹

AaT Ligation Reaction Analysis by HPLC. All HPLC analyses of LysAlaAcm ligations were monitored on an Agilent HPLC using a

Waters C18 column (Milford, MA). The solvents used for peptide purification were the following: 0.1% trifluoroacetic acid in water (Solvent A) and 0.1% trifluoroacetic acid in acetonitrile (Solvent B). The HPLC method had the following solvent gradient (Gradient 1): 0 min 1% B, 5 min 1% B, 10 min 30% B, 15 min 40% B, 20 min 100% B, 25 min 100% B, 27 min 1% B, 30 min 1% B. Peptides were monitored at two absorption wavelengths during HPLC analysis, 215 nm for peptide absorption, and 325 nm for Acm absorption. MALDI MS analyses confirmed identity of peptide, Xaa-A donor, and ligated products.

Fluorometric Analysis of Transferase Experiments. Real time fluorescent monitoring of Nap addition to LysAlaAcm was carried out using reaction solutions prepared on the 500 μ L scale in stirred quartz cuvettes with 1.00 cm path lengths. Reagent concentrations were as described above, except that donor **4d** was withheld until the cuvette was placed in the fluorometer and temperature equilibrated. Reactions were then initiated by addition of 20 μ L of 25 mM Nap-A (**4d**) stock or water (for negative controls). Fluorescence emission at 390 nm was monitored after excitation at 325 nm. Prior calibration had shown that NapLysAlaAcm (**5d**) concentrations could be estimated from fluorescence intensity readings (F_{5d}) using the equation [**5d**] = $(1 - (F_{5d}/F_{\text{Control}}))1.39[\text{LysAlaAcm}]$, where F_{Control} is the fluorescence reading of the reaction with water added instead of **4d**. Calibration details are given in Supporting Information. Measurements were taken on the Cary Eclipse fluorometer in the "Kinetics" mode at 37 °C, acquiring data every 15 s. The excitation and emission slit widths were 5 nm and the averaging time was 1 s.

α -Casein N-Terminal Modification. α -Casein (4.8 mg) was modified with Azf-A (**4c**) in a reaction volume of 1 mL in modified AaT buffer (50 mM HEPES pH 8.0, 150 mM KCl, 10 mM MgCl_2) and AaT (0.05 mg). The reaction mixture was incubated at 37 °C for 12 h, AaT was removed using Ni^{2+} resin (Ni-NTA Superflow, Qiagen), and then buffer exchanged four times into phosphate buffered saline (PBS, 12 mM NaH_2PO_4 , 50 mM NaCl, 4.7 mM KCl, pH 8.0) by Spectra/Por 1 dialysis tubing (Spectrum Laboratories; Rancho Dominguez, CA). Azf-modified α -casein was used directly after buffer exchange into PBS. Aliquots (1 nmol) of α -casein were diluted into PBS (22 μ L) containing either fluorescein-alkyne (FlAlk, 91 μ M) or propargylamine (Alk, 227 μ M) for copper-catalyzed azide-alkyne cycloaddition (CuAAC). The CuAAC reaction was initiated by addition of CuSO_4 (100 μ M), tris-(3-hydroxypropyl)triazolylmethyl amine (THPTA, 500 μ M), and sodium ascorbate (5 mM). For control reactions, equivalent volumes were replaced with either PBS or water. The reaction mixtures were incubated at 4 °C for 2 h. Reactions were boiled with gel loading dye LDS (Pierce; Rockford, IL) for 10 min at 95 °C and analyzed by SDS-PAGE. Fluorescence images were obtained with a Geldoc (BioRAD; Hercules, CA) using an excitation wavelength of 302 nm for detection of fluorescein. α -Casein was also directly visualized by Coomassie Brilliant Blue staining.

Tryptic Digest Analysis of α -Casein Modification. α -Casein-Azf (33 μ g), or α -casein (33 μ g), was digested with sequencing grade modified Trypsin (0.6 μ g) in 27 μ L of 25 mM ammonium bicarbonate pH 7.5 (freshly prepared). Digestions were carried out at 37 °C for 14 h. Trypsin digest aliquots (1 μ L) were combined with α -cyano-4-hydroxycinnamic acid (1 μ L of a saturated solution in 1:1 $\text{H}_2\text{O}/\text{CH}_3\text{CN}$ with 1% TFA) and analyzed by MALDI MS.

α -Casein Labeling in Cleared Cell Lysate. AaT was expressed in *E. coli* BL21-Gold (DE3) cells as previously described. A cleared cell lysate was obtained by centrifugation following cell lysis using sonication. Protein were modified in a final reaction volume of 110 μ L with Azf-A (**4c**, 3 mM) using AaT (25 μ L) from cleared lysate in AaT buffer (50 mM HEPES pH 8.0, 150 mM KCl, 10 mM MgCl_2). α -Casein (0.012 mg or 0.12 mg) was added to the reaction and incubated at 37 °C for 2 h, after 1 h, an additional 0.3 μ mole of **4c** was added. For control reactions, equivalent volumes were replaced with water. Excess **4c** was removed by

buffer exchange four times against phosphate buffered saline (PBS, 12 mM NaH_2PO_4 , 50 mM NaCl, 4.7 mM KCl, pH 8.0) by Spectra/Por 1 dialysis tubing. Aliquots (18 μ L) of **4c** modified lysates were diluted to a final volume of 22 μ L using PBS containing FlAlk (182 μ M) for copper-catalyzed "click" reactions. The "click" reaction was initiated by addition of CuSO_4 (100 μ M), THPTA (500 μ M), and sodium ascorbate (5 mM). The reaction mixtures analyzed by PAGE gel using Coomassie staining and fluorescence as above. The full gel is shown in the Supporting Information.

■ ASSOCIATED CONTENT

S Supporting Information. Synthesis and characterization of **1b–h**, **3b–h**, and **4b–h**; synthesis of reagents for "click" chemistry; AaT expression and purification; analysis of transferase reactions with some representative HPLC chromatograms; kinetics derivations for enzymology; analysis of α -casein N-terminal modification by Edman degradation; full PAGE gel analysis of Azf labeling in cell lysates; and descriptions of AaT mutation experiments. Complete author lists for references¹³ and¹⁵ This material is available free of charge via the Internet at <http://pubs.acs.org>.

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