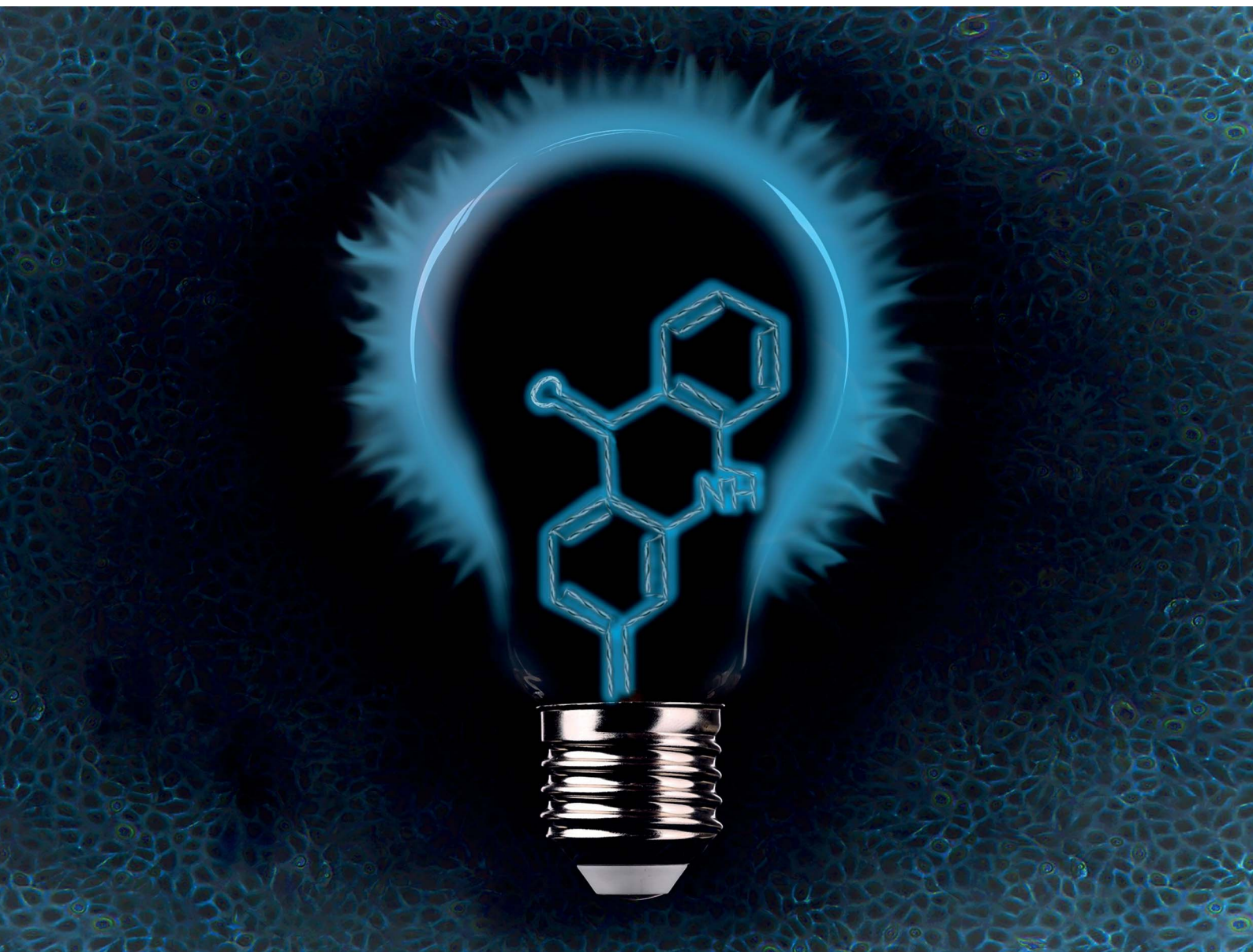


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EDGE ARTICLE

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Genetic encoding of a highly photostable, long lifetime fluorescent amino acid for imaging in mammalian cells†

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Acridonylalanine (Acd) is a fluorescent amino acid that is highly photostable, with a high quantum yield and long fluorescence lifetime in water. These properties make it superior to existing genetically encodable fluorescent amino acids for monitoring protein interactions and conformational changes through fluorescence polarization or lifetime experiments, including fluorescence lifetime imaging microscopy (FLIM). Here, we report the genetic incorporation of Acd using engineered pyrrolysine tRNA synthetase (RS) mutants that allow for efficient Acd incorporation in both *E. coli* and mammalian cells. We compare protein yields and amino acid specificity for these Acd RSs to identify an optimal construct. We also demonstrate the use of Acd in FLIM, where its long lifetime provides strong contrast compared to endogenous fluorophores and engineered fluorescent proteins, which have lifetimes less than 5 ns.

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Introduction

Fluorescence spectroscopy provides a powerful method to monitor protein folding and dynamics in real-time under physiological conditions. One challenge for monitoring these dynamics is the ability to fluorescently label the protein of interest without disrupting its folding and function.^{1,2} Although widely used, large fluorescent protein tags such as green fluorescent protein (GFP)^{3,4} can hinder native protein folding and function,^{1,5,6} and affinity labeling such as bioorthogonal (“click”) chemistry⁷ and SNAP⁸ or Halo⁹ tags can be limited by biocompatibility, labeling efficiency and solvent accessibility of the labeling location.^{1,10} To avoid these issues, a fluorescent label can be incorporated co-translationally, eliminating the need for additional post-translational manipulation and allowing one to label proteins at interior sites.¹¹ Incorporation of the labels in living cells is made possible by an engineered aminoacyl tRNA synthetase (RS) that selectively charges an orthogonal tRNA with a non-canonical amino acid (ncAA).¹² The

anticodon loop recognizes an amber stop codon that is introduced in the mRNA chain, leading to the insertion of the ncAA in the desired position of a protein of interest.

The Petersson and Mehl laboratories have previously incorporated acridonylalanine (Acd or δ) in *E. coli* using these amber codon suppression methods.^{13–18} Acd is a small fluorescent amino acid (222 Å³), with an emission maximum at 420–450 nm, an extinction coefficient of 5700 M^{−1} cm^{−1} and a quantum yield of 0.98 in aqueous buffer (ESI, Fig. S1 and Table S1†) and high photostability (ESI, Fig. S5 and Table S2†).^{19,20} We have previously demonstrated Acd's utility as a minimally perturbing biophysical probe (Fig. 1). Acd's fluorescence can be used to quantify protein amounts¹⁸ and as part of a short-range Förster resonance energy transfer (FRET) pair.^{13,14,18,21} One key advantage of Acd is its long fluorescence lifetime (τ), which varies with the polarity of the environment and has been used in fluorescence polarization (FP) assays to investigate the dynamics of protein–protein interactions.¹⁷ Compared to other visible wavelength fluorescent ncAAs used for *in cellulo* genetic incorporation,^{22–25} hydroxycoumarin ethylglycine (Hco), coumarinyl lysine (CouLys), dansylalanine (DanAla), and acetylnaphthalenylaminoalanine (Anap), Acd has the highest quantum yield, longest fluorescence lifetime, and greatest photostability in water (ESI, Fig. S1–S9 and Tables S1–S6†). The longer lifetime of Acd makes the fluorophore more responsive to molecular weight changes in FP experiments.^{17,26} It also permits better resolution of multiple populations in lifetime experiments, including single photon counting experiments²⁷ and in fluorescence lifetime imaging microscopy (FLIM, Fig. 1).²⁸

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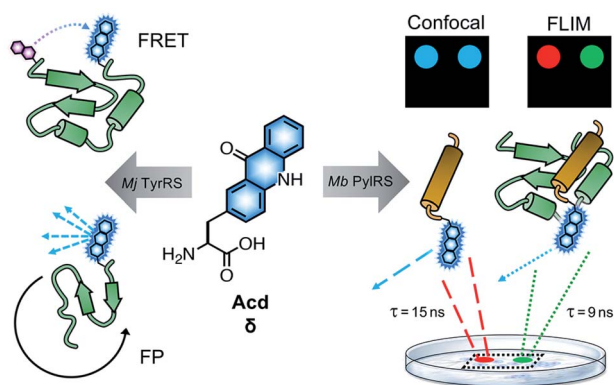


Fig. 1 Acrid Applications. Acrid (or δ) has been previously used in Förster resonance energy transfer (FRET) and fluorescence polarization (FP) to monitor conformational changes and protein binding interactions among purified proteins expressed in *E. coli* using RSs based on *Mj* TyrRS. Development of a *Mb* PylRS-based system for mammalian cell incorporation enables confocal microscopy and fluorescence lifetime imaging (FLIM) studies to observe processes like protein association in live cells. FLIM pixels are coloured by lifetime, (red = 15 ns, green = 9 ns) which is related to the Acrid local environment.

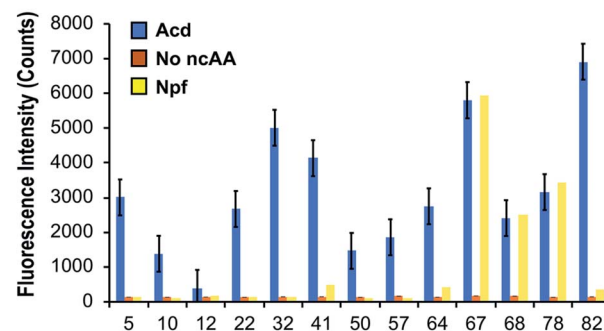
To date, all Acrid studies relied on *E. coli* protein expression, as our engineered RSs were based on an *Methanocaldococcus jannaschii* (*Mj*) tyrosyl RS and thus were only orthogonal to the translation machinery in prokaryotes. In order to use Acrid in mammalian cell imaging experiments, including FLIM, we needed an RS that is orthogonal in mammalian cells.

Here, we report the development of an Acrid incorporation method that relies on an engineered pyrrolysine (Pyl) RS for site specific protein labeling. PylRS variants from *Methanosarcina bakeri* (*Mb*),²⁹ *Methanosarcina mazei* (*Mm*),²⁹ and *Methanomethylophilus alvus* (*Ma*)^{30,31} have been shown to be orthogonal in both *E. coli* and mammalian cells. Using engineered *Mb* PylRS variants that specifically and efficiently incorporate Acrid in both *E. coli* and mammalian cells, we also demonstrate that this new labelling platform can be used for FLIM studies in mammalian cells.

Results

Acrid synthetase selection

To identify a *Mb* PylRS/tRNA_{CUA} pair capable of site-specifically incorporating Acrid into proteins in response to an amber stop codon (TAG) we screened a library of *Mb* PylRS variants in which five active-site residues were randomized to all 20 amino acids (N₃₁₁, C₃₁₃, V₃₆₆, W₃₈₂, G₃₈₇) using a standard life/death selection in *E. coli*.^{32,33} Following a round of positive selection in the presence of Acrid and a round of negative selection against canonical amino acids, 96 colonies were assessed for their ability to suppress a TAG codon interrupted sfGFP gene (sfGFP-TAG₁₅₀) in the presence of Acrid and suppressor tRNA_{CUA}. The top 25 performing clones were sequenced and 13 unique RS clones were identified with highly similar active site sequences (Fig. 2 top and ESI, Table S7†). These 13 clones were evaluated for (1) their efficiency (full-length protein yield in presence of Acrid), (2)



Mb AcridRS 82

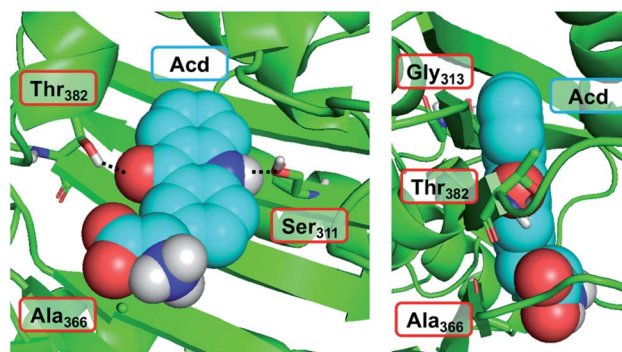


Fig. 2 PylRS mutants for Acrid incorporation. Top: fluorescence measurements of sfGFP reporter to evaluate RS efficiency. Orange, blue, and yellow represent fluorescence from colonies induced in media containing no ncAA, 1 mM Acrid, or 1 mM Npf, respectively. Cultures of 500 μ L were grown for 48 hours before 4-fold dilution of suspended cells with buffer for fluorescence measurement at 528 nm (485 nm excitation) using a plate reader. RSs enabling high sfGFP fluorescence in Acrid conditions with low fluorescence in no ncAA or Npf conditions were selected for further characterization. Additional data showing sfGFP fluorescence for all clones are shown in ESI, Fig. S10.† Bottom: a homology model of *Mb* AcridRS 82 based on the X-ray crystal structure of *Mm* PylRS (PDB ID: 2ZIN)³⁴ with Acrid docked in the active site using Rosetta. Key points of interaction with Acrid are shown from two angles. Sites of mutation relative to parent *Mb* PylRS are labeled in red boxes. A model of *Mb* AcridRS 41 is shown in ESI, Fig. S11.†

fidelity (level of canonical amino acid misincorporation in absence of Acrid) and (3) their efficiency in producing full length protein using *N*-phenyl-amino phenylalanine (Npf), an undesired side product of Acrid synthesis that was recognized by first-generation AcridRSs.^{13,14} Three of these clones showed permissivity for Npf, and were not considered further.

The most active and selective variants (32, 41, 82) had similar mutations in the active site (N₃₁₁S and V₃₆₆A, as well as W₃₈₂T or W₃₈₂V). *Mb* AcridRS 32 and 82 also had an additional active site mutation of C₃₁₃A or C₃₁₃G, respectively, and *Mb* AcridRS 82 also contained a non-active site mutation, L155V. AcridRS 41 and AcridRS 82 were selected for further evaluation as 82 was most active and 41 featured the same active site mutations, except it retained Cys₃₁₃ of the parent RS. The two RSs were cloned into both pDULE2 (ref. 35) and pUltra³⁶ plasmids, under respective lpp and tacI promoters, each also containing tRNA_{CUA}. We did not retain the L155V mutation since other active sequences (e.g.

Mb PylRS clone 75) showed activity without the mutation. If necessary, Npf incorporation concerns can be eliminated by using an alternative Acd synthetic route.¹³

Structural modeling

In order to better understand the effects of the mutations in the selected mutants, we built a homology model of *Mb* AcdRS 82 based on a X-ray crystal structure of substrate-bound *Mm* PylRS (PDB ID: 2ZIN).³⁴ With the exception of one loop region, there is very high active site sequence homology between the *Mm* and *Mb* PylRS (ESI, Fig. S12†), and it has been demonstrated that the conservation of the active site allows mutations for nAA-selective variants to be swapped between the two types of PylRS.^{37–41} Thus, we expect the homology model to be quite accurate. Acd was docked into the *Mb* AcdRS 82 active site in Rosetta, followed by relaxation of the structure (Fig. 2 bottom). A model of *Mb* AcdRS 41 was generated by a G₃₁₃C mutation of *Mb* AcdRS 82, followed by relaxation (ESI, Fig. S11†).

One can rationalize the effects of the four mutations in AcdRS 82 as follows: Asn₃₁₁ to Ser creates space in the active site and provides a hydrogen-bond acceptor for the Acd sidechain N–H, Trp₃₈₂ to Thr removes a steric clash and provides a hydrogen-bond donor for the Acd ketone, Cys₃₁₃ to Gly removes a steric clash with the distal ring of Acd and allows it to hydrogen-bond with both Ser₃₁₁ and Thr₃₈₂ together, and Val₃₆₆ to Ala removes a steric clash. The existence of the hydrogen bond bridge in the AcdRS 82 model nicely explains its superior activity relative to AcdRS 41.

Protein expression in *E. coli* and characterization of Acd incorporation

From the initial screen, *Mb* AcdRS 41 and *Mb* AcdRS 82 were subjected to further investigation of their incorporation efficiency and ability to mis-incorporate canonical amino acids at the TAG site. To this end we expressed TAG-bearing constructs of α -synuclein (α S) and calmodulin (CaM) to investigate the RSs in the context of an intrinsically disordered protein (α S) where the tertiary structure of the protein should not affect incorporation, and a structured protein (CaM) where destabilization by Acd incorporation could decrease protein expression.¹⁸ We have previously studied conformational changes and protein/protein interactions in both proteins using Acd.^{13,16} Each protein construct was fused to a C-terminal GyrA intein containing a six histidine (His₆) tag. This fusion construct provides a way to eliminate potential protein truncations, as the intein with the His₆ tag is only produced for the full length protein and can be cleaved tracelessly using β -mercaptoethanol.⁴²

To compare RS efficiency and specificity, we first expressed α S using a plasmid encoding α S with a TAG codon replacing Phe94 or Glu114 as well as a plasmid encoding either *Mb* AcdRS 41 *Mb* AcdRS 82 and the cognate tRNA_{CUA}. We compared incorporation efficiency, based on isolated yields, and amino acid specificity, based on matrix-assisted laser desorption/ionization (MALDI) mass spectrometry (MS) data. Phe94 and Glu114 were chosen as the incorporation sites as they have been shown to tolerate a variety of bulky fluorophore substitutions,

including both nAA incorporation and Cys modification.^{16,43–50} As a positive control, we expressed the same protein construct using an *Mj* RS previously optimized for Acd incorporation in *E. coli* (*Mj* AcdRS A9).¹⁴ As an additional comparison of anticipated masses and yields, we also expressed the wild-type (WT) α S sequence fused to the His₆ intein tag. This serves as an approximate reference mass for any mis-incorporation at the TAG site due to charging of canonical amino acids. Given the structural similarity of Acd to the aromatic amino acids Phe, Trp, and Tyr we anticipated that there was a potential for mis-incorporation of these amino acids. Indeed, Phe mis-incorporation has been previously observed in PylRS mutants selected for incorporation of aromatic unnatural amino acids,⁵² although the selectivity of engineered PylRSs has been reported to be generally good.⁵³

Protein expressions in *E. coli* were performed with either pDULE2 or pUltra plasmids, using BL21 DE3 or C321.deltaA cells (release factor 1 removed and all 321 UAG codons changed to improve nAA incorporation),⁵⁴ respectively. Protein levels were monitored *via* gel throughout cell lysis, nickel purification and intein cleavage. SDS-PAGE analysis confirmed that full-length Acd-containing α S was produced for both *Mb* AcdRS 41 and *Mb* AcdRS 82 using either the pDULE2/BL21 and pUltra/C321 expression methods. However, MALDI MS analysis of purified whole protein constructs demonstrated that significant mis-incorporation of a canonical amino acid occurred with pUltra/C321 with either *Mb* AcdRS 41 and *Mb* AcdRS 82 (ESI, Fig. S19†). In contrast, MALDI MS data confirmed that both *Mb* AcdRS 41 and *Mb* AcdRS 82 exclusively charge tRNA_{CUA} with Acd for pDULE2/BL21 expressions (Fig. 3, S17 and S18, ESI †). Whole protein masses matching the anticipated mass for Acd for incorporation at position 114 in α S are shown in Table 1. No evidence of Npf incorporation was observed.

Final protein yields were calculated based on molar absorptivity and confirmed by detergent compatible (DC) protein quantification assay, showing that *Mb* AcdRS 82 incorporates Acd into α S more efficiently than *Mb* AcdRS 41 (Tables 1, S8 and S9, ESI,† gel analysis in Fig. S13–S16†). The mis-

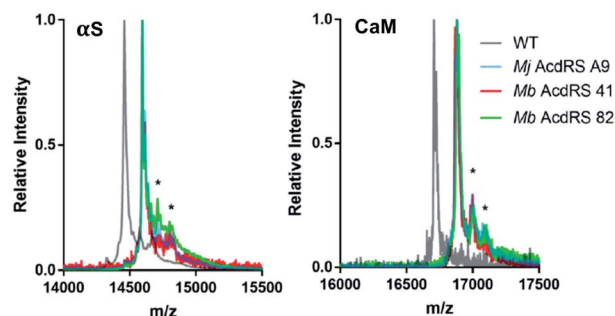


Fig. 3 MALDI MS analysis of Acd incorporation. Expressions of α S-E₁₁₄ δ or CaM-L₁₁₃ δ using *Mj* AcdRS A9, *Mb* AcdRS 41, or *Mb* AcdRS 82 were performed in parallel. WT α S and CaM were also expressed for comparison. MALDI MS data showed masses corresponding to Acd incorporation with no peaks corresponding to mis-incorporation of a canonical amino acid (approximately equal to WT mass). *Indicates a previously described MALDI matrix adduct.⁵¹



^a Yields based on Acd absorbance normalized to WT α S or CaM expression, determined by Tyr absorbance, and confirmed by DC assay, as described in ESI.

Mammalian cell expression screening

Figure 1: Fluorescence intensity of AcdRS variants.

Top Panel: Fluorescence intensity (Counts) vs. Variant

Variant	Acd (Counts)	No ncAA (Counts)
AcdRS 41	~3000	~2000
AcdRS 82	~8000	~2000
- RS	~1500	~1500
WT	~23500	~23500
EGFP	~23500	~23500

Bottom Panel: Fluorescence intensity (Counts) vs. Acid Concentration (μM)

Acid Concentration (μM)	Fluorescence Intensity (Counts)
600	~9000
300	~11000
200	~10500
100	~8000
50	~5000
25	~3500

Fig. 4 Lysate fluorescence analysis of Acd incorporation in mammalian cells. Top: fluorescence measured from EGFP-Y₄₀Δ HEK cell lysates after 36 h using *Mb* AcdRS 41 or 82 with 300 μM Acd or no ncAA added. As a negative control, the EGFP plasmid was transfected without the AcdRS and tRNA_{CUA} plasmid (–aaRS). WT EGFP was separately transfected as a positive control. In all conditions, DN-eRF1 was also transfected. *n* = 3. Protocols are described in detail the ESI† comparison of AcdRS incorporation efficiency section. Cell images are shown in ESI, Fig. S21.† Bottom: expression of EGFP-Y₄₀Δ using *Mb* AcdRS 82 was performed as above, with varying concentrations of Acd. *n* = 3. Protocols are described in the ESI† optimizing expression conditions section. Cell images are shown in ESI, Fig. S3.†

Mb AcdRS 82 was 3-fold higher than *Mb* AcdRS 41. Additionally, both RSs showed low EGFP fluorescence in the absence of Acd, indicating little to no mis-incorporation of natural amino acids (Fig. 4, top and images in and S21, ESI†). We also purified EGFP using a C-terminal His₆ tag and analysed incorporation fidelity using electrospray ionization (ESI) MS. We found no evidence of mis-incorporation of Npf with either RS, but trace incorporation of a natural amino acid using AcdRS41 (ESI, Fig. S26†). Based on these results and our studies in *E. coli*, we selected *Mb* AcdRS 82 for further mammalian cell experiments.

To determine optimal Acd concentrations for imaging experiments, we expressed EGFP-Y₄₀δ using AcdRS82 with varying concentrations of Acd. We found that maximal EGFP fluorescence was observed with 300 μM Acd (Fig. 4, bottom and images in ESI, S22†). We also performed a cell viability study with varying Acd concentrations and found that Acd was toxic to cells at concentrations above 600 μM, possibly explaining the decrease in EGFP expression at 600 μM Acd (ESI, Fig. S20†). Cell imaging revealed that the cell morphology was normal at 300 μM (ESI, Fig. S25†). Finally, we varied the concentrations of transfection reagents and plasmids and tested the effect of DN-eRF1 to find optimal expression conditions (ESI, Fig. S23 and S24†). Comparing to WT EFGP expression based on both imaging and lysate fluorescence measurements in these experiments demonstrated that suppression was 29% efficient under

optimal conditions, using *Mb* AcdRS 82 with DN-eRF1, 300 μ M Acd, and 1.6 μ g DNA (for a 6 well dish) at a 1 : 5 DNA to Lipofectamine 2000 ratio, with imaging after 24–48 h.

Mammalian cell fluorescence microscopy

To investigate whether Acd can be used as a fluorescence reporter for protein localization in mammalian cells, we incorporated Acd at TAG mutants in three fusion proteins of varying localization, insulin receptor fused to GFP (IR-K₆₇₆ δ -GFP), hyperpolarization-activated cyclic nucleotide-gated ion channel fused to yellow fluorescent protein (spHCN-W₃₅₅ δ -YFP), and maltose binding protein (MBP-E₃₂₂ δ). The GFP and YFP fusions are C-terminal, so production of the fluorescent protein ensures that suppression of the stop codon was successful. Similar to the GFP screening transfection, *Mb* AcdRS 82 and DN-eRF1 plasmids were co-transfected into HEK293T/17 cells along with a vector encoding the protein of interest with a TAG codon at a site previously shown to permit bulky amino acids.^{57–59} After washout of unincorporated Acd, we imaged Acd-containing insulin IR-K₆₇₆ δ -GFP, spHCN-W₃₅₅ δ -YFP, and MBP-E₃₂₂ δ in live cells with confocal microscopy.

The localization of these proteins is known to be membrane bound in the case of IR-GFP and spHCN-YFP, and cytosolic for MBP. Confocal imaging for IR-K₆₇₆ δ -GFP and spHCN-W₃₅₅ δ -YFP revealed that Acd fluorescence was localized to the membrane and colocalized with YFP or GFP fluorescence, respectively (Fig. 5). This demonstrates Acd's ability to be used as a fluorescent probe for protein localization. We additionally imaged cells containing the protein construct and Acd in the absence of the pAcBac1 tR4-MbAcdRS82 plasmid (Fig. 5, no RS conditions). Confocal imaging revealed little or no fluorescence in the GFP or YFP channels, which indicates that no substantial read-through of the TAG codon occurred. We observed only minor

fluorescence in the Acd channel. This indicates that our washout protocols are effective and the remaining fluorescence may be due to endogenous fluorophores such as NADH or tRNA loaded with Acd, which cannot be washed out of the cell (wider fields of view shown in ESI, Fig. S27†). Note that the rounded morphology of the cells is not a result of Acd toxicity, but of the extensive washing to remove Acd, as demonstrated by applying the same protocol to cells in the absence of Acd (ESI, Fig. S27†). From these studies, we conclude that Acd incorporation is specific to our protein of interest and requires *Mb* AcdRS 82, as expected.

Mammalian cell fluorescence lifetime imaging

Since Acd is particularly useful as a FLIM probe, as a proof of concept, we monitored the localization of Acd labeled insulin receptor (IR-K₆₇₆ δ -GFP) in HEK293T/17 cells using FLIM. FLIM spatially resolves the fluorescence decay rate of a sample to produce an image based on its fluorescence lifetime. Endogenous fluorophores typically found in a cell have a fluorescence lifetime in the range of 0.3–7.5 ns.²⁸ Since Acd has a fluorescence lifetime of 15–16 ns, our imaging method is free of background from endogenous cellular fluorophores. For this study, we transfected vectors containing *Mb* AcdRS 82, DN-eRF1, and IR-K₆₇₆ δ -GFP into HEK293T/17 cells. Imaging the cells using intensity alone while exciting at 375 nm (Fig. 6) shows Acd fluorescence as well as autofluorescence, likely from NAD and NADH. Highlighting the pixels in which the majority of the intensity was due to Acd (lifetime >15 ns) in red allows the membrane localization of IR-K₆₇₆ δ -GFP to be discerned and also eliminates the autofluorescence signal. Highlighting the pixels with a short lifetime (<4 ns) in blue reveals the autofluorescence in all of the cells. The long lifetime pixels (red) compare well with emission from the GFP tag (excitation at 488 nm). The

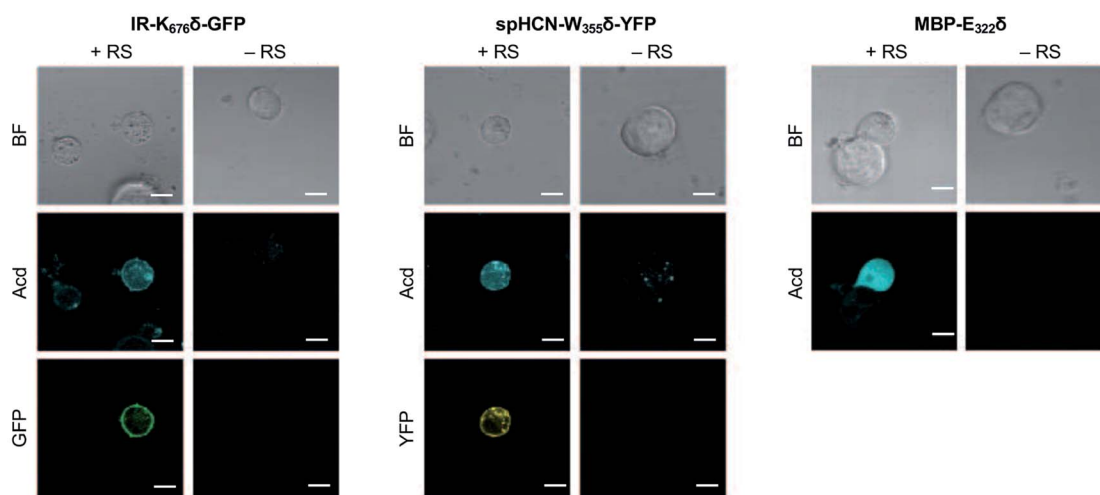


Fig. 5 Acd Incorporation into proteins in Live HEK Cells. Cells were transfected with the indicated plasmid for the protein of interest with a TAG codon and a plasmid encoding a dominant negative release factor to suppress protein truncation, with (+RS) or without (–RS) a plasmid encoding *Mb* AcdRS 82 and tRNA_{CUA}, confocal microscopy images show brightfield (BF), the Acd fluorescence channel (405 nm excitation; 409–480 nm emission), and the GFP (488 nm excitation; 500–621 nm emission) or YFP (514 nm excitation; 519–621 nm emission) channels. Scale bar = 20 μ m. Transfection and imaging conditions are described in the ESI† confocal microscopy and FLIM section.



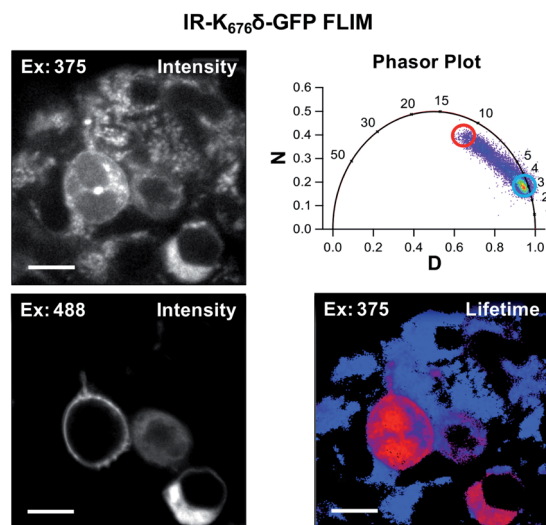


Fig. 6 Acrid fluorescence lifetime imaging (FLIM). FLIM allows Acrid fluorescence to be separated from autofluorescence in living cells. IR-K₆₇₆δ-GFP was expressed with *Mb* AcdRS 82 in HEK cells. Intensity plots (left) show emission after excitation at 375 or 488 nm. Phasor plot (top right) shows phase vectors D and N (see ESI† for details) plotted for excitation at 375 nm with 10 MHz modulation and points representing lifetimes of pixels. Circles indicate pixels with a large fraction of lifetime ≥ 15 ns (red) and pixels with a large fraction of lifetimes < 4 ns (blue) in the lifetime image (bottom right). Scale bar = 20 μ m. Transfection and imaging conditions are described in the ESI† confocal microscopy and FLIM section.

pronounced autofluorescence signal in Fig. 6 compared to Fig. 5 is due to differences in the excitation wavelength used in each experiment. We note that the images in Fig. 6 seem to capture IR-K₆₇₆δ-GFP in different stages of maturation where it is observed in what appears to be ER and/or Golgi in addition to the plasma membrane. Additional FLIM data with spHCN-W₃₅₅δ-YFP highlighting long lifetime Acrid emission are shown in Fig. S28 in ESI†.

Discussion

The development of a PylRS-based system for incorporating Acrid is valuable for a number of reasons, including the applications in confocal microscopy and FLIM described here, as well as the potential for double-labelling of proteins using mutually orthogonal RSs deriving from PylRS and *Mj* TyrRS. Acrid has a higher quantum yield and greater photostability in buffer than any of the other genetically encodable ncAAs (ESI, Fig. S1–S9 and Tables S1–S6,† quantum yield is comparable to Hco). Thus, Acrid is in many ways superior for imaging, although this depends on the application. For example, the high level of environmental sensitivity depending on the protonation state of Hco is valuable for probing changes in local environment, but may confound other types of studies like FRET. Alternatively, the large Stokes shift of DanAla is useful for eliminating background scatter in spite of DanAla's very low quantum yield in water (0.10, see ESI, Table S2†). Acrid is particularly well-suited to FLIM studies because of its 15–16 ns lifetime in water, as most

endogenous fluorophores have lifetimes ≤ 7 ns allowing one to exclude their fluorescence by gating lifetime data.⁶⁰ This makes Acrid a superior FLIM probe compared to GFP variants, none of which have comparably long lifetimes (< 5 ns).^{61,62} The other genetically-encodable ncAAs all have shorter lifetimes, except DanAla in hydrophobic environments. However, the variation of the DanAla lifetime from 3 ns in water to 10–12 ns in MeOH, DMSO, and THF (ESI, Table S5†) is a less useful range than that of Acrid, which varies from ~ 15 ns in water to ~ 8 ns in organic solvent, only dropping to ~ 4 ns in the very hydrophobic environment of THF (ESI, Table S3†). Although we did not demonstrate it here, Acrid's lifetime also makes it useful for anisotropy imaging, where the size range distinguishable by a fluorophore increases with fluorophore lifetime.⁶³

Beyond FLIM applications, *Mb* AcdRS 82 can potentially be used to incorporate a photo-activatable version of Acrid developed by Xiao and coworkers.⁶⁴ *Mb* AcdRS 82 could also be paired with an *E. coli* TyrRS engineered to encode *p*-azidophenylalanine in mammalian cells.⁶⁵ This would allow *in cellulo* fluorescent labelling of a second site with “clickable dyes” opening access to site-specific controlled addition of FRET partners for Acrid. Several strategies for encoding two amino acids have been published, including the use of ochre or opal codons, as well as four base codons.^{65–67} Chin and coworkers have also shown that one can generate mutually orthogonal RS/tRNA pairs from PylRS variants.⁶⁸ We will pursue these strategies where a coumarin derivative (Acrid FRET donor) or BODIPY derivative (Acrid FRET acceptor) could be attached.¹³ Finally, while Acrid's properties are valuable, its low extinction coefficient and blue wavelength emission are not ideal for some microscopy applications, and we will attempt to obtain *Mb* AcdRS 82 variants that can encode brighter and/or red-shifted Acrid analogs such as aminoacridonylalanine.²¹ We expect that the PylRS may be better for this than the *Mj* TyrRS as it has been shown to accommodate many large amino acids.⁵³

Conclusion

Acrid has been previously demonstrated to be useful as a fluorescent marker and as a biophysical probe to monitor protein conformations without perturbing protein structure. Prior to this study, Acrid labelling was limited to proteins expressed in *E. coli* and these proteins were used for *in vitro* biophysical experiments. Here, we have demonstrated the capability of two PylRS variants to efficiently and selectively incorporate Acrid into several proteins in *E. coli* and mammalian cells. Notably, transfection and imaging experiments were independently conducted in the investigators' laboratories and consistent results were observed. *Mb* AcdRS 82 afforded higher levels of protein expression in both cell types with excellent fidelity, as confirmed by MALDI and ESI MS, and was used for proof-of-concept fluorescence microscopy and FLIM studies. FLIM has advantages over confocal microscopy because it provides an additional dimension with which to discriminate changes in intensity due to the number of proteins from intensity changes due to solvatochromic effects resulting from conformational changes or interactions with other proteins. To our knowledge,



the work described here is the first example of the direct use of a fluorescent ncAA in FLIM (the lone previous example used FRET with GFP).⁶⁹ The long lifetime of Acd provides a large dynamic range for FLIM measurements, making it superior to GFP variants or the other genetically encodable ncAAs. This, in addition to Acd's high quantum yield and photostability, make *Mb* AcdRS 82 a valuable new tool for the protein imaging community, enabling facile labelling of proteins in mammalian cells with a small, non-perturbing fluorophore that has unique properties.

Experimental

Aminoacyl tRNA synthetase (RS) screening

Acd RS selection was performed on a library of *Mb* PylRS mutants a life/death selection as previously described.⁷⁰ Briefly, RS library members in a pBK screening plasmid were transformed into *E. coli* cells with a positive selection plasmid, pREP-pylT, containing an amber codon-disrupted chloramphenicol acetyltransferase gene and pylRS tRNA (pylT). Cells containing the pBK-RS library and pREP-pylT were cultured in the presence of chloramphenicol and Acd. Surviving pBK RS library members were isolated and the pBK plasmid was transformed into cells containing a negative selection plasmid, pYOB2-pylT, encoding an amber codon-disrupted barnase gene under control of an arabinose promoter as well as pylT. Cells containing pBK-RS library and pYOB2-pylT were cultured in the presence of arabinose, but in the absence of Acd. The remaining pBK-RS library from the surviving colonies was transformed into cells containing a GFP reporter plasmid, pALS-GFP-TAG₁₅₀. Cells containing pBK-RS library and pALS-GFP-TAG₁₅₀ plasmids were grown in 96 well plates in autoinducing media (see ESI†) containing 1 mM Acd or Npf, or autoinducing media with no ncAA. GFP fluorescence was measured after 48 h. Additional selection analysis and sequence information are provided in ESI.†

Computational modeling of AcdRS 82 active site

M. mazei homologues of AcdRS 82 and 41 were identified by ClustalOmega alignment of *Mb*PylRS and *Mm*PylRS. A crystal structure of *Mm*PylRS bound to *N*_ε-Boc-L-Lysine and an adenosine triphosphate (ATP) analogue (PDB 2ZIN) was used as the basis for Rosetta modeling.³⁴ Firstly, the ATP analogue was removed, and the amino acid ligand was replaced with Ala. Subsequently, missing loops were modeled using Rosetta Remodel and the structure was relaxed using the beta_{nov16} scorefunction.⁷¹ Upon active site mutations, Acd was initially positioned using DARC and several relaxations of the structure were performed.⁷² The final structure was analyzed in PyMol.⁷³

Protein expression and characterization of Acd incorporation

Plasmids for each RS (*Mj* AcdRS A9, *Mb* AcdRS 41, or *Mb* AcdRS 82) in the previously described pDULE2 vector,³⁵ containing tRNA_{CUA}, were separately transformed into BL21 cells that also contained the pTXB1-αS-TAG₁₁₄-Mxe-H₆ plasmid. After growth in auto-inducing media (see ESI† for details), cells were harvested by centrifugation. The supernatant was discarded and

the pellet was resuspended, sonicated and pelleted, then the supernatant was collected and the protein purified with a Ni²⁺-NTA column. After elution, the protein was subjected to intein cleavage. Protein production and cleavage yields were analyzed by SDS PAGE. Each construct was further purified by fast protein liquid chromatography and analyzed by MALDI MS. The protein concentrations were analyzed by DC assay and UV-Vis absorbance. See ESI† for CaM expression and purification details.

Mammalian cell expression (EGFP screening)

Mb AcdRS 41 and *Mb* AcdRS 82 were separately cloned into the pAcBac1 tR4-MbPyl vector to form pAcBac1 tR4-MbAcdRS41 and pAcBac1 tR4-MbAcdRS82.⁵⁵ These plasmids were transfected into HEK293T/17 cells using Lipofectamine 2000, along with DN-eRF1 (peRF1-E55D.pcDNA5-FRT) and pEGFP-Y40TAG-C1 to evaluate suppression efficiency. Transfections were also performed in the absence of pAcBac1 tR4-MbAcdRS41 or pAcBac1 tR4-MbAcdRS82 as negative (–RS) controls. Transfections with DN-eRF1 and pEGFP-C1 (WT EGFP) were used as positive controls. Transfections were performed with and without DN-eRF1 as well as with varying Acd concentrations, incubation times, and Lipofectamine 2000/DNA ratios to identify optimal expression conditions. Transfections were evaluated by imaging using an Olympus CKX53 microscope and by measuring the EGFP fluorescence of cell lysates using a Tecan M1000 plate reader.

Mammalian cell expression (confocal microscopy)

pAcBac1-tR4-MbAcdRS82 and DN-eRF1 plasmids were co-transfected into HEK293T/17 cells along with either pIR-TAG676-GFP, pcDNA3.1HE-spHCN-TAG355-YFP, or pcDNA3-FLAG-MBP1-E322TAG (FLAG-MBP1-E322TAG-W340W.pcDNA3-k) plasmids, using a total of 1.6 μg DNA. Confocal imaging was performed on a Zeiss LSM 710 microscope in and analysed using Fiji software.⁷⁴ Transfection and imaging conditions are described in detail in the ESI† confocal microscopy and FLIM section.

Fluorescence lifetime imaging microscopy (FLIM)

Cells were prepared and transfected as described for confocal microscopy experiments. FLIM experiments were performed at room temperature in DMEM supplemented with 10% FBS, penicillin, and streptomycin using an ISS Q2 laser scanner and ISS A320 FastFLIM system mounted on a Nikon TE2000U microscope. Acd was excited using a 375 nm laser and GFP and YFP were excited using bandpass filters to select lines of a YSL SC-PRO 7 supercontinuum laser. FLIM data were acquired and analysed using ISS VistaVision software.

Plasmid and Acd availability

Sequences of the pDULE2-MbAcdRS82, for expression in *E. coli*, and pAcBac1-tR4-MbAcdRS82, for expression in mammalian cells, are given in the ESI.† The plasmids will be made available



via Addgene. Acd is available upon request to Prof. Petersson and is commercially available from Watanabe Chemicals.

Data availability

Primary data for photophysical measurements and synthetase screening, as well as computational modeling code and atomic coordinates, are provided as ESI.†

Author contributions

Experiments were designed collaboratively by C. M. J., E. J. P., W. N. Z., S. E. G., and R. A. M. C. M. J. performed the photophysical characterization of all Uaas, *E. coli* incorporation studies, and optimization of mammalian cell incorporation. R. J. B. screened *Mb* PylRS mutants. A. J. E. performed mammalian cell mass spectrometry analysis of Acd incorporation. M. M., W. N. Z. and S. E. G performed confocal imaging and FLIM experiments. D. M. R. assisted in the synthetase screen and performed the initial *E. coli* experiments in C321 cells. T. S. M. provided computational docking models. Y. V. synthesized Acd. C. M. J and E. J. P. wrote the manuscript with input from all authors.

Conflicts of interest

There are no conflicts to declare.

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