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Structural insights into latency of the metallopeptidase ulilysin (lysargiNase) and its unexpected inhibition by a sulfonyl-fluoride inhibitor of serine peptidases

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Peptidases are regulated by latency and inhibitors, as well as compatibilization and cofactors. Ulilysin from *Methanoscincus acetivorans*, also called lysargiNase, is an archaeal metallopeptidase (MP) that is bio-synthesized as a zymogen with a 60-residue N-terminal prosegment (PS). In the presence of calcium, it self-activates to yield the mature enzyme, which specifically cleaves before basic residues and thus complements trypsin in proteomics workflows. Here, we obtained a low-resolution crystal structure of proulilysin, in which 28 protomers arranged as 14 dimers form a continuous double helix of 544 Å pitch that parallels cell axis *b* of the crystal. The PS includes two α -helices and obstructs the active-site cleft of the catalytic domain (CD) by traversing it in the opposite orientation of a substrate, and a cysteine blocks the catalytic zinc according to a "cysteine-switch mechanism". Moreover, the PS interacts through its first helix with an "S-loop" of the CD, which acts as an "activation segment" that lacks one of two essential calcium cations. Upon PS removal during maturation, the S-loop adopts its competent conformation and binds the second calcium ion. Next, we found that in addition to general MP inhibitors, ulilysin was competitively and reversibly inhibited by 4-(2-aminoethyl)benzenesulfonyl fluoride (AEBSF; $K_i = 4 \mu\text{M}$). This is a compound that normally forms an irreversible covalent complex with serine peptidases but does not inhibit MPs. A high-resolution crystal structure of the complex revealed that the inhibitor penetrates the specificity pocket of ulilysin. A primary amine of the inhibitor salt-bridges an aspartate at the pocket bottom, thus mimicking the basic side chain of substrates. In contrast, the sulfonyl fluoride warhead is not involved and the catalytic zinc ion is freely accessible. Thus, the usage of inhibitor cocktails of peptidases, which typically contain AEBSF at ~ 25 -fold higher concentrations than the determined K_i , should be avoided when working with ulilysin. Finally, the structure of the complex, which occurred as a crystallographic dimer recurring in previous mature ulilysin structures, unveiled an N-terminal product fragment that delineated the non-primed side of the cleft. These results complement prior structures of ulilysin with primed-side product fragments and inhibitors.

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1. Introduction

The pappalysins are a family of zinc-dependent metallopeptidases (MPs) whose founding member is a ~ 180 kDa multidomain protein first found in the bloodstream of pregnant women termed pregnancy-associated plasma protein-A or pappalysin-1.^{1,2} They form family M43 in the MEROPS peptidase database (<https://www.ebi.ac.uk/merops>;³) and are included within the metzincin clan of mononuclear MPs, which eng-

lobes at least 12 structurally-distinct families from all kingdoms of life.⁴⁻⁸ Metzincin catalytic domains (CDs) span ~ 130 –350 residues and share structural features including a five-stranded β -sheet, a "backing helix" and an "active-site helix" within an N-terminal subdomain (NSD); and a "Met-turn" centred on a conserved methionine residue plus a "C-terminal helix" within a C-terminal subdomain (CSD).⁷⁻¹⁰ The two subdomains frame the top and bottom of the active-site cleft, respectively, when viewed in the typical standard orientation of MPs,¹¹ with the lowermost β -strand of the five-stranded β -sheet featuring the upper rim of the active-site cleft. Moreover, a conserved zinc-binding consensus sequence (H-E-x-x-H-x-x-G/N-x-x-H/D; amino-acid one-letter-code, x for any residue), which is contained in the second half of the

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active-site helix and beyond, forms the interface between the NSD and the CSD. It includes three histidine/aspartate zinc ligands and the general base/acid glutamate for catalysis.^{8,9}

The structure of pappalysin-1 has been recently elucidated.^{10,12} It reveals that its CD resembles that of the subfamily of the “lower pappalysins”, formerly the “unicellular pappalysins”,^{13,14} which encompasses related sequences from archaea, bacteria, and, as more recently ascertained, also unicellular and multicellular eukaryotes like fungi and algae. They span a short N-terminal prosegment (PS) engaged in latency, as described for many MPs,⁹ followed by the CD and, optionally, downstream segments and domains. Several subfamily members have been biochemically, functionally or transcriptionally characterized (Table 1). In contrast, experimental structural information of these pappalysins is restricted to mature ulilysin from the archaeon *Methanosarcina acetivorans*, as well as mature and latent mirolysin from the periodontopathogenic bacterium *Tannerella forsythia*.¹³⁻¹⁸

Ulilysin is biosynthesized as a zymogen with a 60-residue PS for latency that is self-activated *in trans* at bond S⁶⁰–R⁶¹ (ulilysin residue numbers in superscript according to UniProt database entry [UP] Q8TL28) in the presence of calcium as the indispensable cofactor.¹³ In addition, a 20-residue negatively-charged flexible C-terminal peptide is further removed upon cleavage at bond A³²²–R³²³, eventually releasing the mature 262-residue CD (R⁶¹–A³²²). Functionally, ulilysin efficiently cleaved insulin, insulin-like growth-factor-binding proteins 2 to 6, several types of denatured collagen and gelatin, as well as other model and extracellular-matrix proteins.^{13,15} Regarding inhibitors, ulilysin was only inhibited by the general metal chelators 1,10-phenanthroline and EDTA, as well as excess zinc.¹³ Some inhibition (~75%) was observed with the broad-spectrum inhibitor of matrix metallopeptidases, batimastat, at ~50-fold molar excess.^{13,16} A comprehensive analysis of ulilysin’s substrate specificity revealed strict N-terminal selectivity for lysine and arginine residues, including methylated and di-

methylated species, and further proved the suitability of the MP, which is also referred to as “lysargiNase”, as a tool for proteomics.¹⁹ It is currently marketed by Merck/Millipore and, by generating peptides with N-terminal positively-charged side chains, it is complementary to trypsin, which generates peptides with C-terminal positively-charged residues. Thus, by using both enzymes, more peptides can be covered during proteomics studies. Moreover, ulilysin operates at temperatures of up to 55 °C while proteomics-grade trypsin usually works at 37 °C, which further enlarges the palette of biotechnological applications of the archaeal MP.

Structurally, mature ulilysin includes all the aforementioned hallmarks of the metzincins, with H²²⁸–H²³⁸ featuring the zinc-binding consensus sequence.^{13,16,17} Moreover, like astacins and serralysins, ulilysin possesses a fourth zinc ligand in the form of a tyrosine (Y²⁹²) two positions downstream of the Met-turn methionine (M²⁹⁰), which participates in substrate and reaction–intermediate binding following a “tyrosine switch”.²⁰ Among the specific characteristics of lower pappalysins, ulilysin further contains a Lin12-Notch-repeat-like loop within the upper β-strand of the NSD β-sheet, a β-ribbon inserted after the central β-strand of this sheet, and a segment encompassing two short α-helices, which is found between the backing helix and the top β-strand of the sheet and lines the back of the CSD. A C-terminal product fragment bound to the primed-side of the cleft revealed that the basic side chains in the P₁ position of substrates (active-site cleft sub-site nomenclature according to ref. 11 and 21) are bound by D²⁹⁵ at the bottom of the deep S₁ pocket. This pocket is the main driver of specificity in most MPs¹¹ and in ulilysin it is further lined by the L¹⁸⁸, F²²⁰, T²²⁵ and M²⁹⁸ side chains and the main chain of segment Y²⁹²–D²⁹⁵.¹⁵

Sulfonyl fluorides (SFs) were identified in the 1930s as toxic to animals through their capacity to inhibit serine esterases including acetylcholinesterase.^{22,23} They were found to also inhibit the serine peptidases α-chymotrypsin, trypsin, neutro-

Table 1 Biochemically, functionally or transcriptionally analysed lower pappalysins

Name	Organism	Ref.
Cytophagalsin	Collagenolytic bacterium <i>Cytophaga</i> sp. L.43-1	69–71
Fpp2, Fpcol and related MPs	Bacterial fish pathogen <i>Flavobacterium psychrophilum</i> and the related psychrophilic Antarctica bacterium <i>Flavobacterium frigidimar</i>	72–76
PoMTP	Edible hiratake mushroom <i>Pleurotus ostreatus</i>	77
MEP1	Fungal pathogen <i>Coccidioides posadasii</i> , which causes the respiratory San-Joaquín-Valley fever	78
Ulilysin (lysargiNase)	Archaeon <i>Methanosarcina acetivorans</i>	13, 15–17 and 19
Mirolysin	periodontopathogenic bacterium <i>Tannerella forsythia</i>	14, 18, 43 and 79
Mrmep1, Mrep2 and orthologues	Filamentous entomopathogenic fungi <i>Metarhizium robertsii</i> (formerly <i>M. anisopliae</i>), which is utilized in pest control, and <i>Metarhizium rileyi</i>	80 and 81
RcMEP1	Fungal plant pathogen <i>Rhizonia cerealis</i>	82
Mep2	Fungal plant pathogen <i>Verticillium longisporum/dahliae</i>	83
Tryp-N	Thermophilic filamentous fungus <i>Chaetomium thermophilum</i>	84
Leptolysin	Spirochaete bacterium <i>Leptospira interrogans</i> , which causes leptospirosis and Well’s disease	46
M43-type MP	Human dermatophytic fungal pathogen <i>Trichophyton violaceum</i>	85
M43-type MP	Mutualistic root-symbiotic fungus <i>Piriformospora indica</i>	86
M43-type MP	Hemibiotrophic fungal plant pathogens <i>Colletotrichum</i> spp.	87 and 88
M43-type MP	Nematodeegg-parasitic fungus <i>Pochonia chlamydosporia</i>	89



phil elastase, cathepsin G and *Staphylococcus aureus* V8 protease, as well as complement, fibrinolytic and coagulation serine peptidases.^{24–27} These enzymes operate through covalent acyl-enzyme reaction intermediates resulting from the nucleophilic attack of the Oy atoms of the catalytic residue onto the scissile carbonyl carbon of a substrate and the concomitant release of the C-terminal product half.^{28,29} In the second part of the reaction, this intermediate is resolved by the attack of a solvent molecule, which frees the N-terminal half of the product and restores the catalytic residue. SFs mimic the first part of the reaction inasmuch as the catalytic serine performs a nucleophilic attack onto the fluorosulfonyl group of the inhibitor, which produces a sulfonic ester and causes a fluoride anion to leave.²⁷ In contrast to substrates, however, this reaction product is highly stable and cannot be hydrolysed, which causes irreversible inhibition of the enzyme.^{23,27} This property has been taken advantage of to develop probes for target identification and validation, as well as the mapping of protein–protein interactions, substrates and binding sites.³⁰ Finally, in addition to catalytic serine residues, reactive threonine, lysine, tyrosine, cysteine and histidine residues, as well as N-terminal α -amino groups of proteins, may likewise be sulfonylated by SFs.³⁰

Among the most common SFs in protein chemistry are phenylmethylsulfonyl fluoride (PMSF) and 4-(2-aminoethyl)benzenesulfonyl fluoride (AEBSF), which is commercialized under the brand Pefabloc® SC (Fig. 1A).^{27,30–32} AEBSF is more soluble and stable in water than PMSF, which results in higher inhibitory efficiency and duration, as well as storability in solution.^{26,30,32,33} Moreover, contrary to PMSF, AEBSF is not toxic and does not inhibit cysteine peptidases. Currently, AEBSF and/or PMSF are included in commercial peptidase inhibitor cocktails that are routinely employed during protein purification protocols to prevent degradation.^{30,34}

Here, we aimed to understand the regulation of ulilysin activity through zymogenicity, and, in the process, we serendipitously discovered that AEBSF – but not PMSF – leads to reversible inhibition, which we further explored through biochemical and structural approaches. Finally, we dissected the *in crystallo* oligomerization behaviour of the mature enzyme and the zymogen.

2. Results and discussion

2.1. Molecular structure of proulilysin

We prepared proulilysin with the E²²⁹A mutation in which the general base/acid for catalysis was replaced with alanine. This is a common strategy to prevent autolysis and obtain intact MP zymogens for structural studies because this mutation generally does not affect the fold.⁹ The construct spanned segment M¹–R³⁴², which thus included the aforementioned 20-residue flexible C-terminal peptide, and was flanked by a disordered 20-residue tail on the N-terminus owing to the cloning strategy. This notwithstanding, we managed to obtain well-shaped crystals (Fig. 1B, left), which diffracted to 4.5 Å resolution (see

Table 2). Exhaustive efforts to obtain higher-diffracting crystals with a shorter construct lacking these tails (S¹⁴–L³²¹) were unsuccessful. Next, the structure solution proved challenging owing to highly anisotropic diffraction, low resolution of the diffraction data, presence of translational non-crystallographic symmetry, very high solvent content (71%) and large cell constants (max. 544 Å). Eventually, 14 protomers totalling 4284 residues (33 586 atoms) were found in the asymmetric unit (a.u.) of the crystals (see section 2.5) by recursive molecular replacement employing a predicted model for the searches (see section 3.5).

The final refined structure of proulilysin (see Table 2 for statistics) includes residues P¹⁸–R³²³ plus a zinc and a calcium cation for each protomer (A–N). No further features were modelled due to the low resolution of the data. Segment P¹⁸–S⁶⁰ constitutes the PS, which runs across – and thus blocks – the active-site cleft in the opposite orientation of a substrate (Fig. 2A). This is a general strategy found in MP zymogens to prevent untimely autolytic cleavage *in cis* as the PS:CD interaction does not conform to a Michalis complex.⁹ The PS includes two α -helices (α 1p and α 2p) and a cysteine (C²³), which blocks the catalytic zinc following a “cysteine-switch mechanism”. Within metzincins, such mechanisms have been previously reported for matrix metallopeptidases,^{35,36} a-disintegrin-and-metallopeptidase (ADAM) enzymes³⁷ and bacterial – but no holozoan – astacins.^{38,39} The mature part of proulilysin (R⁶¹–R³²³) is largely indistinguishable from the mature CD (Fig. 2B), which has been previously reported.^{13,15,16} The only exception is an “S-loop” (I²³⁹–D²⁶⁴) within the CSD, which adopts a different trace in the zymogen for the first “S” turn (G²⁴¹–P²⁴⁹). It is hereafter termed “activation segment” in analogy to similar elements for latency in peptidases from the astacin family within metzincins^{39–41} and trypsin-type serine peptidases in general.⁴² The S-loop is a key regulatory element for activity of mature ulilysin as it binds two nearby calcium cations, which are ~9 Å apart (Ca⁹⁹⁷ and Ca⁹⁹⁸ in Fig. 2C) and are essential for protein stability as a “calcium switch”. Indeed, treatment with EDTA, a broad-spectrum calcium chelator, abolishes activity, but it can be reversibly restored by removing the chelate and adding calcium.¹³ In the zymogen, the activation segment lacks Ca⁹⁹⁷ (Fig. 2C) and is partially flexible. Through segment D²⁴²–R²⁴⁵, it participates in interactions with M²⁶–H³⁰ from α 1p (Fig. 2A). Moreover, it is engaged in packing contacts in the crystal, which would at least partially explain the low resolution of the diffraction data.

2.2. Mechanism of maturation and comparison with mirolysin

Maturation cleavage before R⁶¹ releases the PS so that the interaction between α 1p and the activation segment is disrupted. A subsequent rotation of the latter under maximal displacement of ~9 Å (at R²⁴⁵) reorients E²⁴³ from its surface location to a position compatible with Ca⁹⁹⁷ binding and adoption of an overall competent conformation of the calcium switch (Fig. 2C). In the mature enzyme, the cation is further tightly bound by the main-chain oxygens of W²⁴⁰, P²⁴⁹, Q²⁶²



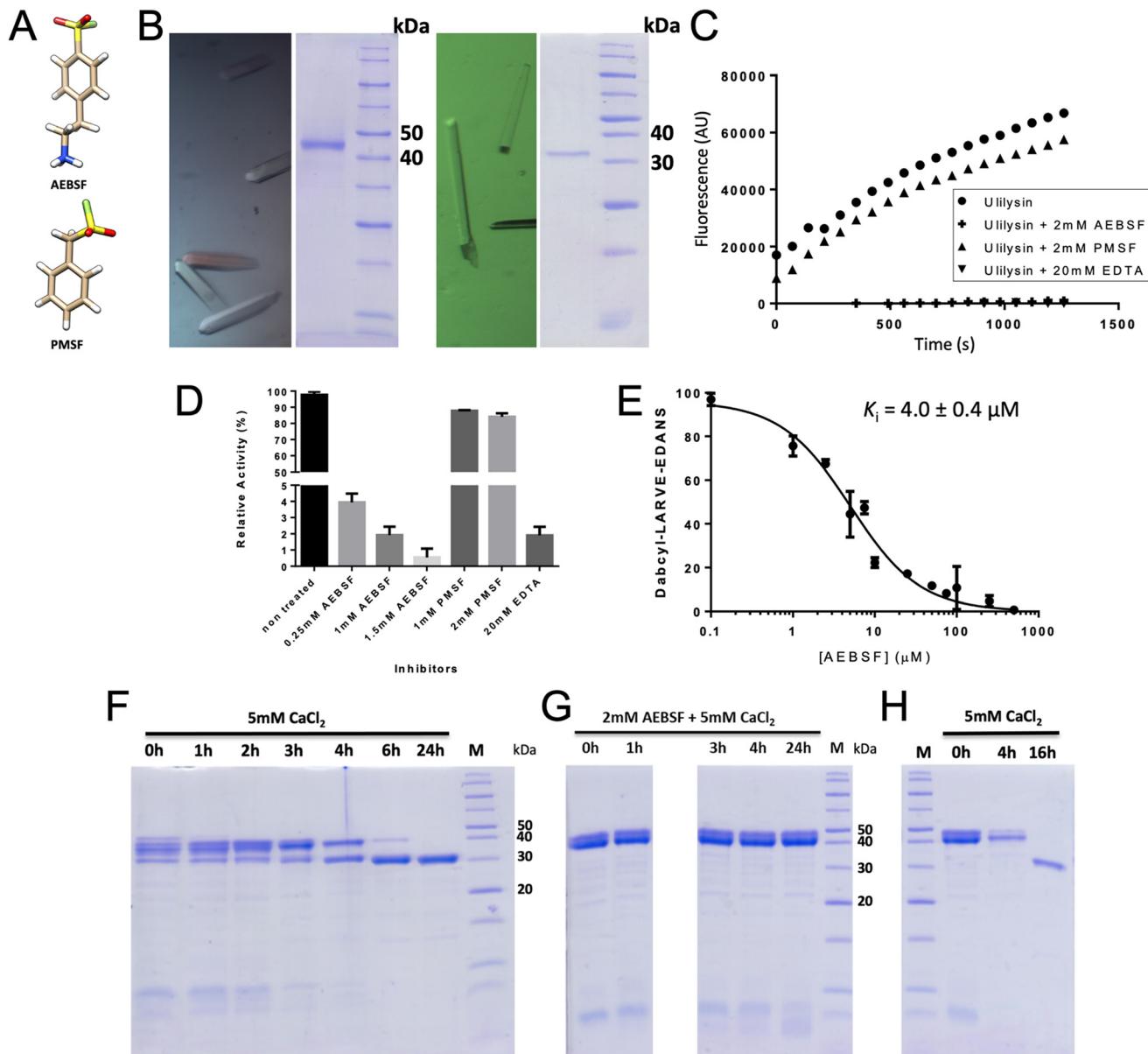


Fig. 1 Biochemical studies. (A) Chemical structures of 4-(2-aminoethyl)benzenesulfonyl fluoride (AEBSF) and phenylmethylsulfonyl fluoride (PMSF). (B) Centred orthorhombic crystals of proulilysin E²²⁹A (left panel) contained the intact zymogen (middle-left panel) while primitive orthorhombic crystals of ulilysin C²⁶⁹A (middle-right panel) contained the mature enzyme only (right panel). (C) Inhibition of ulilysin cleavage of the fluorogenic peptide Dabcyl-L-A-R-V-E-Edans by AEBSF or EDTA but not by PMSF as shown by the fluorescence response. (D) Relative residual activity plot of ulilysin showing that AEBSF inhibition is concentration dependent. (E) Graphical determination of the apparent inhibition constant K_i . (F) Time-dependent autolytic activation of proulilysin to mature ulilysin in the presence of 5 mM calcium chloride. Conversion is complete after 6–24 h under the conditions assayed. (G) Activation of proulilysin (at 1 mg mL⁻¹) is abrogated by AEBSF at 1 mM. (H) Removal of AEBSF from (G) in a PD-10 desalting column results in complete activation before 16 h in the presence of calcium chloride. Thus, AEBSF inhibition is reversible.

and A²⁶³, which are in a comparable position in the incompetent zymogen (Fig. 2C). Accordingly, calcium binding and the associated enzyme activation is mainly triggered by the repositioning of E²⁴³, which thus functions as a “glutamate switch”.

To date, structural information on lower pappalysin zymogens is restricted to promirolysin^{14,18} whose CD shares ~50% sequence identity with ulilysin.⁴³ Both proulilysin E²²⁹A and promirolysin E₂₂₅A (residue numbers in subscript according to UP A0A0F7IPS1/G8ULV1) were crystallized in the absence of

calcium (see section 3.4 and ref. 14) but both structures contain the calcium site around Ca⁹⁹⁸ (Fig. 2C). In contrast to the archaeal zymogen, however, the *Tannerella* zymogen further bears the calcium site around Ca⁹⁹⁷, which in turn is found in mature ulilysin and mirolysin.¹⁴ This results from the S-loop adopting an equivalent conformation in mature and latent mirolysin. When compared with mature ulilysin (Fig. 2C), the chain trace is also comparable bar a three-residue insertion within the first turn of the S-loop, which

Table 2 Crystallographic data

Dataset	Proulilysin E ²²⁹ A	Ulilysin C ²⁶⁹ A/AEBSF
Beam line (synchrotron)	XALOC (ALBA)	XALOC (ALBA)
Space group/protomers per a.u. ^a	C222 ₁ /14	P2 ₁ 2 ₁ /2
Cell constants (<i>a</i> , <i>b</i> , and <i>c</i> in Å)	192, 544, 186	50.13, 124.66, 86.90
Wavelength (Å)	0.97926	0.97866
Measurements/unique reflections	254 017/57 824	874 866/66 338
Resolution range (Å) (outermost shell) ^b	94.7–4.50 (4.77–4.50)	86.9–1.65 (1.75–1.65)
Completeness (%)/ <i>R</i> _{merge} ^c	99.5 (98.9)/0.277 (0.934)	99.9 (89.2)/0.120 (1.655)
<i>R</i> _{meas} ^c /CC(1/2) ^c	0.314 (1.054)/0.994 (0.874)	0.125 (1.723)/0.999 (0.892)
Average intensity ^d	5.3 (2.0)	11.8 (1.8)
<i>B</i> -Factor (Wilson) (Å ²)/Aver. multiplicity	121.7/4.4 (4.5)	53.7/13.2 (13.1)
Resolution range used for refinement (Å)	94.7–4.50	62.3–1.65
Reflections used (test set)	56 408 (842)	65 256 (752)
<i>R</i> _{cryst} (<i>R</i> _{free}) ^c	0.289 (0.302)	0.207 (0.243)
Non-H protein atoms/ionic ligands /waters/non-ionic ligands per a.u.	33 586/14 Zn ²⁺ , 14 Ca ²⁺ /-/	4120/2 Zn ²⁺ , 5 Ca ²⁺ /688/2 GOL, 2 AES, 2 GSS
Rmsd from target values		
bonds (Å)/angles (°)	0.008/0.875	0.006/0.901
Average <i>B</i> -factor (Å ²)	171.2	29.1
Analysis of protein contacts and geometry ^e		
Ramachandran favoured/outliers/all analysed	4026 (89%)/0/4256	509 (97%)/0/524
Bond-length/bond-angle/chirality/plan. outliers	1/0/0/0	0/0/0/0
Side-chain outliers	412 (11.2%)	7 (1.5%)
All-atom clashes/clashscore ^e	1446/22.1	15/1.7
RSRZ outliers ^e <i>F</i> _o : <i>F</i> _c correlation	176 (4.1%)/0.86	6 (1.1%)/0.98
PDB access code	8CDB	8CD8

^a Abbreviations: AES, 4-(2-aminoethyl)benzenesulfonyl fluoride; a.u., crystallographic asymmetric unit; GOL, glycerol; GSS, tripeptide of sequence glycine-serine-serine; RSRZ, real-space *R*-value Z-score. ^b Values in parentheses refer to the outermost resolution shell. ^c For definitions, see ref. 90. ^d Average intensity is $\langle I/\sigma(I) \rangle$ of unique reflections after merging according to Xscale.⁴⁹ ^e According to the wwPDB validation Service (<https://www.pdb-validation.wwpdb.org/validservice>).

causes the tip of this turn to flip outward in the archaeal enzyme without affecting the calcium-binding sites. Most noteworthy, this state of the arts obeys to deviating PS structures in promirolysin and proulilysin (Fig. 2D). Indeed, while both zymogens share the general topology of two tandem helices (α 1p and α 2p) blocking the left, non-primed side of the cleft and a cysteine binding the catalytic zinc ion, the particular arrangement of the helices largely deviates (Fig. 2D). This causes helix α 1p of the bacterial enzyme to be incompatible with the activation segment of the archaeal zymogen. In mirolysin, the topologically equivalent segment, which is three residues shorter (see above), adopts the orientation of the mature CD, *i.e.*, it is preformed in the zymogen.

This remarkable difference in so closely related enzymes correlates with slight but significant differences in function: in contrast to ulilysin (see section 2.1), in mirolysin calcium acts first as an activator of the zymogen to yield the mature form by autolysis and then as a protector from further degradation, thus providing structural – though not thermal – stabilization to the CSD.⁴³ Moreover, calcium is an activity enhancer but it is not essential.

2.3. AEBSF is a ulilysin inhibitor

During purification of the distinct proulilysin variants, an EDTA-free inhibitor cocktail was added after cell lysis to prevent degradation by most peptidases except MPs.^{30,34} Thus, this crude extract should have permitted proulilysin autoactivation upon addition of calcium, as observed for the zymogen

once purified,¹³ but we found this was not the case. We hypothesized that any of the cocktail components might have an unexpected inhibitory effect on ulilysin, which led us to analyse SFs. These covalently target the catalytic O γ atoms of serine peptidases and are normally not inhibitors of MPs, which, like aspartic and glutamic peptidases, operate through a polarized solvent as the attacking nucleophile.^{9,44,45} This notwithstanding, we found that mature ulilysin was inhibited by AEBSF in a competitive dose-dependent manner, with an apparent inhibition constant (K_i) of 4.0 μ M (Fig. 1C–E). Similarly, the enzyme was inhibited by the general metal chelator EDTA (Fig. 1C and D), as previously reported.¹³ In contrast, it was not inhibited by other reversible and irreversible serine- and cysteine-peptidase inhibitors such as aprotinin, E-64, iodacetamide or benzamidine (Fig. 1C, D and ref. 13). Moreover, PMSF, which is also an SF and shares with AEBSF the warhead and an aromatic ring (Fig. 1A), did not inhibit ulilysin (Fig. 1C and D), thus confirming earlier findings.¹³ Overall, this contrasts with mirolysin, which was not inhibited by either PMSF or AEBSF (M. Ksiazek and J. Potempa, personal communication, and ref. 43). Finally, another lower pappalysin, leptolyisin, was also insensitive to PMSF but AEBSF was not tested.⁴⁶

We next studied the effect of AEBSF on the autolytic activation of proulilysin *in vitro*. In the presence of 5 mM calcium chloride, the MP zymogen underwent activation, which was complete after <6 hours (Fig. 1F). In contrast, activation was ablated if a further 2 mM AEBSF was present in the reaction mixture (Fig. 1G). This inhibition was reverted upon removal



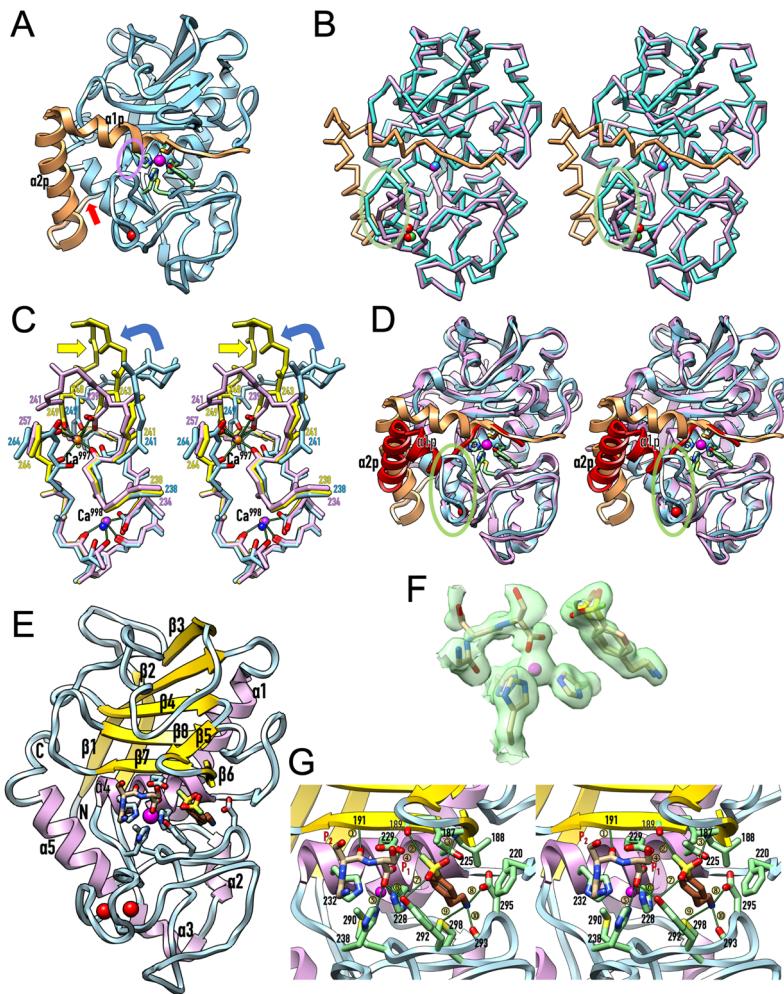


Fig. 2 Protomer structures. (A) Ribbon-type plot of proulysin, with the prosegment featuring helices $\alpha 1p$ and $\alpha 2p$ in orange and the mature moiety in light blue. The “cysteine-switch” cysteine C²³ of the prosegment and the zinc-binding residues, as well as M²⁹⁰ and Y²⁹², are shown for their side chains. The calcium cation (red sphere) and the catalytic zinc ion (magenta sphere) are further displayed. A red arrow pinpoints the final maturation site (S⁶⁰–R⁶¹) and a semi-transparent ellipse in purple highlights the interaction between $\alpha 1p$ and the “activation segment” at the tip of the “S-loop”. (B) Superposition in cross-eye stereo of proulysin (colours as in (A); calcium ion in green, zinc in blue) and the mature ulysin CD (in plum; calcium ions in red, zinc in magenta) as C α -atom traces. The only noteworthy differences occur in the activation segment (encircled by a semi-transparent green ellipse) and the associated absence of the second calcium cation in the zymogen. (C) Superposition in stereo of the main chain of the S-loops of proulysin (carbons and residue numbers in light blue, the calcium ion as a dark blue sphere) and mature ulysin (carbons and residue numbers in yellow, calcium ions as orange spheres), both spanning segment H²³⁸–D²⁶⁴, as well as the S-loop of promirolysin (carbons and residue numbers in plum, calcium ions as purple spheres), which spans segment H₂₃₄–G₂₅₇. Side chains participating in calcium binding are further displayed, all calcium-binding oxygens from the protein moieties are coloured red and green lines denote the calcium–oxygen bonds of ulysin for reference. The metal-coordinating solvent molecules have been omitted for clarity. Both ulysin and promirolysin share both metal-binding sites (Ca⁹⁹⁷ and Ca⁹⁹⁸) and possess equivalent chain traces for their S-loops, except for E²⁴³–D²⁴⁸ (ulysin) and D₂₃₉–G₂₄₁ (promirolysin) owing to a three-residue insertion in the archaeal enzyme (yellow arrow). In contrast, proulysin lacks Ca⁹⁹⁷ and adopts a chain trace differing from the mature enzyme for the activation segment (G²⁴¹–P²⁴⁹). Upon binding of the second calcium and maturation, the enzyme acquires its competent conformation (blue curved arrow). In turn, mirolysin already adopts the competent conformation of the mature moiety in the zymogen, and activation entails mere removal of the blocking prosegment (see also Fig. 4 in ref. 14). (D) Superposition in stereo of proulysin (colouring as in panel (A)) and promirolysin (prosegment in red, mature moiety in plum) in the same view as in (A). The main differences are found in the arrangement of the two prosegment helices (labelled $\alpha 1p$ and $\alpha 2p$ for the *Tannerella* zymogen only) and, concomitantly, the S-loop (semi-transparent green ellipse). (E) Ribbon-type plot of mature ulysin (protomer A) in complex with an upstream product fragment in S₁, S₂ and one position upstream (stick model with carbons in tan), as well as AEBPS (present in double occupancy; carbons in sienna) fitting into the S₁ pocket. The view follows the standard orientation of MPs.¹¹ Selected protein side chains are depicted with carbons in light blue, the zinc and calcium cations are shown as magenta and red spheres, respectively. The regular secondary structure elements of the peptidase are presented as arrows for β -strands (labelled $\beta 1$ – $\beta 8$) and ribbons for α -helices ($\alpha 1$ – $\alpha 5$). The N- and the C-terminus are labelled. (F) Experimental (2mF_{obs} – DF_{calc})-type Fourier omit map contoured at 0.7 σ above threshold superposed on the catalytic zinc, the three zinc-binding histidine residues, the AEBSF molecule (present in double conformation) and the upstream product fragment. (G) Close-up of (E) in stereo centred on the active site. Selected protein residues are shown for their side chains or relevant main-chain atoms with carbons in light green, and labelled with their residue number. Product residues in P₁ and P₂ are likewise labelled. Only the conformation of AEBSF that performs most interactions with the protein is shown for clarity. Thin dark green lines labelled ① to ⑩ denote electrostatic interactions (see Table 3 for the numerical values).



of the inhibitor (Fig. 1H). Thus, both mature and latent ulilysin are efficiently inhibited by AEBSF in a dose-dependent, competitive and reversible fashion, *i.e.* not according to the conventional covalent mechanism of SFs.

2.4. Structure of the ulilysin/AEBSF complex

To study the modus of inhibition of AEBSF, we obtained its high-resolution structure in complex with mature ulilysin by soaking of unbound protein crystals (Fig. 1B, right), which revealed that the inhibitor molecule is present in two very similar conformations (Fig. 2E and F). It is tightly bound in the deep S_1 specificity pocket of the enzyme, so that the terminal primary amine establishes a salt bridge with D^{295} at the bottom of the pocket and further interacts with $M^{298}S\delta$ and $V^{293}O$ (see Fig. 2G and Table 3). The aromatic/aliphatic part of AEBSF is sandwiched by the hydrophobic side chains of L^{188} , F^{220} , T^{225} , H^{228} and Y^{298} , which frame the S_1 pocket. The SF warhead is not involved in covalent binding but establishes electrostatic interactions with $L^{188}N$, $G^{189}O$ and $Y^{292}O\eta$ (Fig. 2G and Table 3). Most noteworthy, the catalytic zinc is not targeted, so AEBSF operates through competitive inhibition of substrate binding but not catalytic hindrance. Overall, this inhibitory modus also explains the lack of inhibition by PMSF, as this SF does not possess an amine on the end of the aromatic ring opposite to the warhead to block the S_1 pocket (Fig. 1A).

The structure of the complex further revealed a tripeptide of tentative sequence G-S-S in the non-primed side of the cleft, thus mimicking a left-behind upstream product fragment after catalysis (Fig. 2E, G and Table 3). The two C-terminal serine residues occupy positions S_2 and S_1 of the cleft, respectively, in extended conformation owing to an inter-main-chain interaction of the P_2 -serine with $A^{191}N$ from the upper-rim β -strand, as expected for a product. The C-terminal carboxylate binds the zinc in a (distorted) bidentate fashion, again as expected for a product. However, the upstream glycine of the tripeptide points out of the cleft owing to a 90° -rotation of the polypeptide chain at the P_2 serine. This is reminiscent of the chain trace of the prosegments of proulilysin and promirolysin (Fig. 2D), although in reverse orientation (see sections 2.1 and

Table 3 Electrostatic interactions between ulilysin and AEBSF or the product peptide

	$A + a$	$A + b$	$B + a$	$B + b$
①	2.83 Å	—	2.84 Å	—
②	2.76 Å	—	2.83 Å	—
③	2.66 Å	2.79 Å	2.91 Å	2.93 Å
④	2.48 Å	—	2.47 Å	—
⑤	2.58 Å	—	2.59 Å	—
⑥	2.87 Å	—	2.85 Å	—
⑦	3.15 Å	3.34 Å	3.06 Å	3.29 Å
⑧	2.70 Å	2.59 Å	2.58 Å	2.65 Å
⑨	3.42 Å	3.22 Å	3.32 Å	3.22 Å
⑩	2.62 Å	3.20 Å	2.82 Å	3.08 Å

Indicated are the values in Å of the ten distances (①–⑩) displayed in Fig. 2G for the two ulilysin protomers (A and B) and the two alternate conformations of AEBSF (a and b), respectively.

Table 4 Reported (pro)ulilysin crystals

PDB code [ref.]	Symmetry	Cell constants (a, b, c in Å)	Content of a. u.
Mature ulilysin (all $C^{269}A$)			
2CKI ¹³	$P2_12_12$	50, 126, 87	One dimer ^a
8CD8	$P2_12_12$	50, 126, 87	One dimer
2J83 ¹⁶	$C222_1$	119, 61, 169	One dimer
3LUM ¹⁷	$C222_1$	97, 174, 125	Two dimers
3LUN ¹⁷	$C222_1$	59, 116, 165	One dimer
Prolulilysin ($E^{229}A$)			
8CDB	$C222_1$	192, 544, 186	Seven dimers ^a

^aThe mature dimers are all equivalent and differ from zymogen dimers, which are also all equal.

2.2). Overall, this upstream product complex complements prior structures of ulilysin with downstream product fragments and primed-side inhibitors.^{13,15}

2.5. Considerations about ulilysin oligomerization

Despite migrating as a monomer in size-exclusion chromatography,^{13,15,16} all previously reported and present crystal structures of mature ulilysin (Table 4) revealed identical types of dimers (Fig. 3A). Indeed, the protein has been previously crystallized indistinctly in four crystal forms from space groups $P2_12_12$ and $C222_1$, with disparate cell constants, and crystals contained either two or four protomers per a.u. The crystals with four protomers (Protein Data Bank entry [PDB] 3LUM;¹⁷) included one dimer formed by chains A and C, while chains B and D formed respective dimers with crystallographic symmetry mates, thus totalling three identical dimers formed by variable combinations of crystallographic and non-crystallographic chains. Dimers are generated by a vertical twofold relating the two protomers joined by the respective front surfaces of Fig. 2E after a relative rotation in the plane of $\sim 35^\circ$. Thus, mature ulilysin has a strong propensity to associate symmetrically in an equivalent fashion despite differences in crystal packing.

With respect to the proulilysin crystals, fourteen protomers were found in the a.u. forming seven dimers (Fig. 3B). These dimers, which follow a twofold at the interface between respective PS helices $\alpha 2p$, are distinct from the mature dimers owing to the presence of the PS, which would interfere in the latter dimerization (Fig. 3C). Here, protomers do not associate over the front surface of Fig. 2A but rather through the lower left surface, and the twofold has its axis rotated 45° when compared with the axis relating mature dimers. Two dimeric heptamers from vertically vicinal a.u.s associate to a continuous left-handed double helix of ~ 544 Å pitch and ~ 95 Å width, whose straight axis is parallel to and spans the entire length of the crystal cell axis b (Fig. 3D). The two congruent intertwined helices are translated relative to each other along the helix axis by half the helix pitch (~ 272 Å). Moreover, the two protomers from each of the two intertwined helices that interact and give rise to each step of the double helix are not perpendicular to the helix axis but inclined by $\sim 15^\circ$. In addition, the double helix extends ~ 39 Å and rotates $\sim 26^\circ$ per step. Overall, the

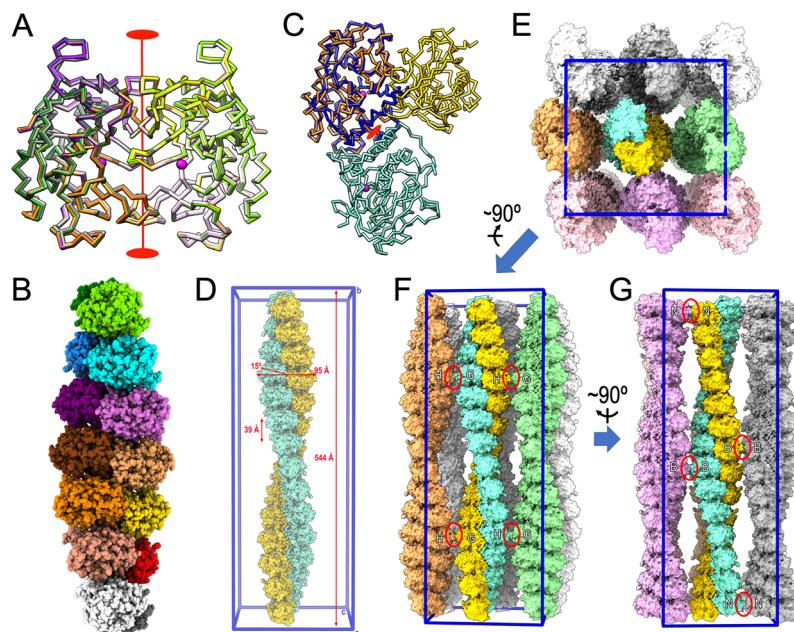


Fig. 3 Oligomerization within the crystals. (A) Superposition of the Ca -traces of the mature ulilysin dimers found in PDB 2J83 (dark green/chartruese), PDB 2LUM (plum/purple), PDB 2LUN (white/grey) and the complex with upstream product and AEBSF (protomers in gold/orange; PDB 8CD8). See also Table 4. The view is perpendicular to the twofold axis (in red) relating both protomers. The overlapping zinc cations are shown as magenta spheres to mark the active site. (B) Full-atom model of the seven identical dimers (protomers in, bottom to top, white/gray, salmon/red, orange/gold, brick/coral, orchid/magenta, dodger blue/cyan and dark green/chartruese) present in the asymmetric unit of the proulilysin crystals. (C) Same as (A) showing only the ulilysin dimer of the product + AEBSF complex (gold/orange) after a horizontal 45°-rotation downward. The orange protomer was superposed onto the mature moiety of one of the protomers of the proulilysin dimer (dodger blue), which interacts with a second protomer (in aquamarine) over a different interface that is created by its own twofold (red ellipse). This twofold has its axis rotated 45° towards the reader when compared with the twofold of mature dimers. (D) Two dimeric heptamers of (B) from vertically vicinal asymmetric units associate to a continuous left-handed double helix of $\sim 544 \text{ \AA}$ pitch and $\sim 95 \text{ \AA}$ width, whose unbent axis is parallel to and spans the entire length of cell axis b of the crystal. The unit cell is further shown in blue for reference (0 stands for the origin; a , b and c are the unit-cell axes). The two congruent intertwined helices (shown as gold and aquamarine full-atom models) are translated relative to each other along the helix axis by half the helix pitch ($\sim 272 \text{ \AA}$). The two protomers give rise to each step of the double helix are not perpendicular to the helix axis but inclined by $\sim 15^\circ$. The double helix extends $\sim 39 \text{ \AA}$ and rotates $\sim 26^\circ$ per step. (E) View resulting from a horizontal 90°-rotation towards the reader of (D), i.e., along cell axis b , of the full content of the unit cell (framed in blue), which includes eight helices further to the one shown in (D). (F) View roughly perpendicular to (E) from which the three bottom double helices (in pink tones) were removed to provide insight into crystal packing contacts. The only lateral interactions are observed between G and H protomers from vicinal double helices (red ellipses). (G) Roughly perpendicular view to (F) depicting the two left columns of double helices of (E). The only interactions are observed between B protomers and N protomers, respectively, from vicinal double helices (red ellipses).

crystal is formed through the parallel association of such continuous double helices (Fig. 3E–G). Superhelices establish only very few contacts with neighbours, namely among G and H protomers from laterally contacting double helices (Fig. 3F) and among B and N protomers from frontally/posteriorly interacting double helices (Fig. 3G). These findings explain the low overall diffraction resolution (few crystal contacts) and the strongly anisotropic diffraction of the crystals (protein fibers within the crystals).

3. Materials and methods

3.1. Protein production and purification

Wild-type proulilysin and the E^{229}A and C^{269}A mutants, all spanning residues $\text{M}^1\text{--R}^{342}$ (see UniProt database [UP] entry Q8TL28 for residue numbering), were expressed in BL21 (DE3)

Star *Escherichia coli* cells transformed with a pET28a-derived vector encoding the respective protein fused with an N-terminal hexahistidine (His_6)-tag, similarly to reported previously for the C^{269}A variant.^{13,15,16,47} The E^{229}A mutation was introduced by site-directed mutagenesis using the Phusion High-Fidelity DNA Polymerase (Thermo Fisher Scientific) employing 5'-CCATGCAATCGGACACTGGTTA-3' as the forward primer and 5'-CCGATTGCATGGTTGCCGT-3' as the reverse primer with wild-type proulilysin as the template. A shorter construct of proulilysin E^{229}A spanning residues $\text{S}^{14}\text{--L}^{321}$ was obtained by PCR using the aforementioned polymerase with 5'-ATATCATATGTCATCTGAAGTGCAA-3' as the forward primer and 5'-ATATCTCGAGTTACAGGAACGATGATCTTGGTCC-3' as the reverse primer, and the vector encoding proulilysin E^{229}A as the template.

Single bacterial colonies transformed by heat shock with the respective vectors were grown overnight in 5 mL starter

pre-cultures of Luria–Bertani medium (LB) supplemented with 50 $\mu\text{g mL}^{-1}$ kanamycin at 20 °C under gentle agitation at 200 rpm in an INFORS orbital incubator. Subsequently, 500 mL of medium were inoculated with a single pre-culture, and cells were grown at 37 °C under agitation until the culture reached an OD_{600} of ~0.8. Protein expression was then induced by adding 0.1 mM of isopropyl- β -D-1-thiogalactopyranoside, and the culture was further incubated overnight at 20 °C under agitation. Cells were harvested by centrifugation (3500g; 20 min; 4 °C) and washed twice with washing buffer (50 mM Tris-HCl, 250 mM sodium chloride, pH 7.5), resuspended in the same buffer plus 10–20 mM imidazole and “cComplete” EDTA-free protease inhibitor cocktail tablets (Roche Life Sciences), and lysed with a cell disruptor (Constant Systems) operated at 1.35 kbar. The cell debris was removed by centrifugation (50 000g; 30 min; 4 °C), and the supernatant containing proulilysin was subsequently filtered, loaded onto a 5 mL nickel-nitrilotriacetic acid HisTrap HP column (GE Healthcare) and washed with washing buffer plus 20–40 mM imidazole. The protein was subsequently eluted by passing 2–3 times 6 mL elution buffer (washing buffer plus 250–300 mM imidazole). After this affinity chromatography step, protein containing fractions were concentrated using centricons of 10 kDa molecular-mass cutoff and applied onto a Superdex 75 Increase 10/300 GL column attached to an ÄKTA Purifier chromatography apparatus (both from Cytiva) for size-exclusion chromatography at 20 °C with purification buffer (20 mM Tris-HCl, 150 mM sodium chloride, pH 7.5). The purified protein was collected in 0.5 mL fractions, and its purity was assessed by SDS-PAGE on custom-made 14% glycine gels followed by staining with Coomassie Brilliant Blue (Sigma-Aldrich).

3.2. Autolytic activation studies of proulilysin

Purified proulilysin E²²⁹A was incubated with activation buffer (purification buffer plus 5 mM calcium chloride), with or without 2 mM 4-(2-aminoethyl)benzenesulfonyl fluoride (AEBSF), for 0 h, 1 h, 2 h, 3 h, 4 h, 6 h and overnight, and the reactions were stopped by boiling the samples in SDS reducing buffer for SDS-PAGE analysis. In a second round of experiments, protein was first incubated with activation buffer containing AEBSF for 24 h, then buffer-exchanged to purification buffer in a PD-10 desalting column, and incubated again with activation buffer without AEBSF. Aliquots were taken after 0 h, 4 h and overnight, and reactions were stopped by boiling the samples with SDS reducing buffer for SDS-PAGE monitoring.

3.3. Activity, inhibition and kinetic analyses

Enzymatic activity of ulilysin C²⁶⁹A was measured with the fluorescence-based EnzCheck Assay Kit (Thermo Fisher Scientific) at 4 $\mu\text{g mL}^{-1}$ (140 nM) mature enzyme concentration, as well as with the fluorescein-conjugate DQ Gelatin from Pig Skin at 5 μM ($\lambda_{\text{exc}} = 480 \text{ nm}$; $\lambda_{\text{em}} = 520 \text{ nm}$; Thermo Fisher Scientific). Reactions were performed in the presence of 10 μM zinc chloride and 5 mM calcium chloride and monitored for the fluorescence response at 37 °C in an Infinite M200 microplate fluorimeter (Tecan). Inhibition of proteolytic

activity against DQ Gelatin was assessed by incubation of the enzyme with EDTA (at 20 mM), AEBSF (at 0.25 mM, 1 mM, 1.5 mM and 2 mM) or phenylmethylsulfonyl fluoride (PMSF; at 1 mM and 2 mM) 5 min at 37 °C before adding the substrate.

The apparent constant (K_i) of ulilysin (at 140 nM) inhibition by AEBSF (at 0.1 mM, 1 mM, 2.5 mM, 10 mM, 50 mM, 250 mM and 500 mM) was determined with the fluorogenic peptide Dabcyl-L-A-R-V-E-Edans ($\lambda_{\text{exc}} = 340 \text{ nm}$; $\lambda_{\text{em}} = 460 \text{ nm}$) as the substrate (at 18.4 μM) after preincubating the enzyme with the inhibitor for 1 h at room temperature.

All activity and inhibition experiments were carried out in triplicate, and data were analysed with the GraphPad software.⁴⁸

3.4. Crystallization and diffraction data collection

Crystals of mature ulilysin C²⁶⁹A formed at 20 °C in crystallization sitting drops containing 1 μL zymogen solution at 5.5 mg mL^{-1} and 1 μL reservoir solution (100 mM 2-(*N*-morpholino) ethanesulfonic acid pH 6.5, 200 mM calcium chloride, 18% polyethylene glycol 8000), as reported previously.^{13,16} These crystals, which resulted from autolytic maturation at bonds S⁶⁰–R⁶¹ and A³²²–R³²³ in the crystallization drops, were incubated for 16 h in soaking solution (reservoir solution plus 10 mM AEBSF) to form a complex with AEBSF. Crystals were harvested with 0.1–0.2 mm nylon loops (Hampton Research), cryoprotected by rapid passage through a drop containing soaking solution plus 20% (v/v) glycerol, and flash-vitrified in liquid nitrogen prior to storage and transport for diffraction data collection.

For proulilysin E²²⁹A, we searched for initial crystallization conditions through the sitting-drop vapour diffusion method at the joint IBMB/IRB Automated Crystallography Platform (<https://www.ibmb.csic.es/en/platforms/automated-crystallographic-platform>). Reservoir solutions were prepared with a Tecan Freedom EVO robot and dispensed into the reservoir wells of 96 \times 2-well MRC plates (Innovadyne Technologies). Crystallization nanodrops containing 100 nL each of protein and reservoir solution were dispensed by a Phoenix/RE robot (Art Robbins) into the respective protein wells. Subsequently, crystallization plates were incubated at 4 or 20 °C in thermostatic crystal farms (Bruker) and remotely inspected for crystal growth. Whenever possible, successful conditions were scaled up to the microliter range in 24-well Cryschem crystallization dishes (Hampton Research). The best crystals of proulilysin E²²⁹A were obtained at 20 °C in drops consisting of 1 μL each of protein solution at 5.5 mg mL^{-1} and reservoir solution (100 mM sodium acetate pH 4.5, 1 M diammonium hydrogen phosphate, 10 mM AEBSF). Crystals were harvested as above and cryoprotected by rapid passage through a drop containing reservoir solution plus 20% glycerol [v/v] prior to flash-vitrification.

X-ray diffraction data were recorded on a Pilatus 6 M pixel detector (Dectris) at 100 K at the XALOC beamline of the ALBA synchrotron (Cerdanyola, Catalonia) and processed using Xds⁴⁹ and Xscale. Structure-factor amplitudes were trans-



formed with Xdsconv to MTZ-format for the Phenix⁵⁰ and CCP4⁵¹ program suites. The asymmetric unit (a.u.) of the ulilysin/AEBSF crystals contained one non-crystallographic dimer and that of the proulilysin E²²⁹A crystals encompassed 14 molecules arranged as seven non-crystallographic dimers. Analysis of the diffraction data with Pointless⁵² within CCP4 and Xtriage⁵³ within Phenix discarded twinning but revealed significant off-origin peaks in both cases, at fractional cell coordinates 0.50, 0.50, 0.50 (ulilysin C²⁶⁹A/AEBSF; 95% of the origin peak, $p = 3 \times 10^{-9}$) and 0.62, 0.00, 0.50 (proulilysin E²²⁹A; 33% of the origin peak, $p = 9 \times 10^{-4}$). These accounted for the presence of (pseudo)translational non-crystallographic symmetry (NCS), which results in distortions of the mean intensity distribution.^{54,55} Table 2 provides essential statistics on data collection and processing.

3.5. Structural solution and refinement

The structure of the ulilysin C²⁶⁹A/AEBSF complex, which was obtained in the same space group ($P2_12_12$) of and similar cell constants to Protein Data Bank (PDB) entry 2CKI,¹³ was solved by Fourier synthesis employing the dimer of protomers A and B found in the a.u. of this structure. The coordinates were subjected to crystallographic refinement using the Phenix⁵⁶ and Buster/Tnt⁵⁷ programs, which included translation/libration/screw-motion refinement. Subsequently, several rounds of refinement alternated with manual model building with Coot⁵⁸ until the final model was obtained. The latter comprised ulilysin residues R⁶¹–A³²², one zinc and two calcium cations, one AEBSF moiety (in alternate occupancy) and a tripeptide of tentative sequence G–S–S for each ulilysin protomer. An extra, functionally irrelevant calcium cation was found at the dimerization interface, and 688 solvent molecules plus two glycerol molecules were placed on the accessible surfaces of the protein molecules. The final structure displays slightly above-average R_{cryst} and R_{free} values considering the high resolution of the data and the quality of the final Fourier map, as previously reported for other cases with translational NCS.^{59–62} Table 2 provides essential statistics of the final model.

The structure of proulilysin E²²⁹A was solved by molecular replacement using the Phaser crystallographic software⁶³ and a computational model for residues V¹⁷–L³²¹ obtained with AlphaFold.⁶⁴ The prediction suggested that, out of the full-length sequence, this segment would correspond to a well-defined structure based on the per-residue values of the predicted local-distance difference test.⁶⁵ Superposition of this model onto the experimental structure of PDB 2J83,¹⁶ which spans the mature moiety (E⁶³–L³²¹), enabled us to implement the catalytic zinc ion and one of the structural calcium cations into the searching model. Molecular replacement calculations with this model found 12 solutions, which formed a helical staircase with seven steps whose axis paralleled the cell axis c . Each step featured an identical dimer bar two, which only evinced one protomer. Superposition of one such dimer onto the two isolated protomers allowed us to complete the 14 protomers actually present in the a.u. Given the low resolution of

the data (4.50 Å), only protomer A was manually adjusted to the electron density with Coot and then copied onto the other 13 protomers. Crystallographic refinement was painstakingly carried out with Phenix applying Cartesian-based NCS restraints, secondary-structure restraints, restraints to the AlphaFold model to steer refinement, translation/libration/screw-motion refinement considering each protomer a separate group, and one thermal displacement parameter per residue. The final model consisted of residues P¹⁸–R³²³ plus a zinc and a calcium cation for each of the 14 protomers (chains A–N). As for the ulilysin C²⁶⁹A/AEBSF structure, the final R_{cryst} and R_{free} values were also artificially higher than expected due to the aforementioned strong translational NCS. Moreover, albeit reflections for R_{free} monitoring had been selected in thin shells, the final values of R_{cryst} and R_{free} were nonetheless partially coupled owing to the 14-fold NCS. Table 2 provides essential statistics of the final model.

3.6. Miscellaneous

Structural superpositions and structure-based sequence alignments were calculated using the SSM routine⁶⁶ in Coot. Figures were prepared using Chimera⁶⁷ and/or ChimeraX.⁶⁸ The final refined models of ulilysin C²⁶⁹A/AEBSF and proulilysin E²²⁹A were validated using the wwPDB Validation Service at <https://validate-rscb-1.wwpdb.org/validservice> (see Table 2) and deposited with the PDB at <https://www.pdb.org> (access codes 8CD8 and 8CDB).

Author contributions

F.X.G.R. and U.E. conceived and supervised the project; A.R.B. and M.M.C. produced and purified proteins, A.R.B. generated mutants, A.R.B. and M.M.C. performed *in vitro* studies, A.R.B. and S.R.M. crystallized proteins, A.R.B. and U.E. collected diffraction data, analysed data and supervised workers; F.X.G. R. solved and refined crystal structures and wrote the manuscript with contributions from all authors.

Conflicts of interest

The authors declare no financial or non-financial conflicts of interest with the contents of this article.

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137827OB-I00 from the State Agency of Research [MCIN/AEI/10.13039/501100011033; may include "Next Generation EU/PRTR" and/or "cofunded by the EU" and/or "ERDF – A way of making Europe"]; and 2021SGR00423 from the Catalan Government).

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