

Crystallization and preliminary X-ray diffraction studies of pyrrolidone carboxyl peptidase from the hyperthermophilic archaeon *Thermococcus litoralis*

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Pyrrolidone carboxyl peptidase from the hyperthermophilic archaeon *Thermococcus litoralis* has been crystallized in a form suitable for X-ray diffraction from ammonium sulfate or ammonium dihydrogen orthophosphate using the vapour-phase diffusion method. Crystals from both precipitants are of the orthorhombic space group $P2_12_12$ with unit-cell dimensions $a = 94.06$, $b = 149.06$, $c = 73.54$ Å. A complete data set to 2.8 Å resolution has been collected from crystals grown from ammonium sulfate

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1. Introduction

Pyrrolidone carboxyl peptidases (pcps; E.C. 3.4.19.3) catalyse the hydrolysis of amino-terminal pyroglutamate residues from peptides and proteins (Doolittle & Armentrout, 1968). They form a subgroup of the large aminopeptidase family of enzymes which demonstrate a wide range of substrate specificity and catalytic mechanisms (Gonzales & Robert-Baudouy, 1994). Pcps have been identified in mammalian and plant species (Szewczuk & Kwiatkowska, 1970), as well as from a variety of bacterial sources (Awadé *et al.*, 1992).

Mechanistically, pcps have been divided into two groups: cysteine peptidases and metallo-peptidases. The former group includes all the bacterial enzymes studied so far and one class of mammalian enzyme. The other class of mammalian enzyme is a zinc-containing metallopeptidase and has been shown to regulate the activity of the neuropeptide, thyrotropin-releasing hormone.

The bacterial pcps are the most extensively studied, with *pcp* genes being cloned from the bacteria *Pseudomonas fluorescens* (Gonzales & Robert-Baudouy, 1994), *Bacillus subtilis* (Awadé *et al.*, 1992), *Bacillus amyloliquefaciens* (Yoshimoto *et al.*, 1993), *Streptococcus pyogenes* (Cleuziat *et al.*, 1992) and *Staphylococcus aureus* (Patti *et al.*, 1995). The enzymes have been biochemically characterized and are either homodimers or tetramers with subunit molecular weights between 22 and 30 kDa. They possess high sequence homology; however, they show no similarity to any other type of peptidase and the pcps are therefore thought to constitute a distinct class of enzyme (Cleuziat *et al.*, 1992). A common feature of all these sequences is the totally conserved cysteine- and histidine-containing motifs, which are proposed to be catalytic residues, as in the cysteine proteases exemplified by papain

(Polgar, 1973). Inhibition studies (Sullivan *et al.*, 1977) and site-directed mutagenesis experiments (Yoshimoto *et al.*, 1993) support this hypothesis. A third, acidic residue has recently been proposed to stabilize the imidazolium ring of the histidine during catalysis in an analogous manner to the serine protease triad, although the exact sequence position of this residue has not been established (Lesaux *et al.*, 1996).

The pyrrolidone carboxyl peptidase from the anaerobic hyperthermophile *Thermococcus litoralis* has recently been cloned and overexpressed in *Escherichia coli* (Singleton *et al.*, unpublished data). The enzyme shows high sequence homology to the bacterial pcps and has a similar quaternary structure of 4×24 kDa subunits. In addition, the protein shows the extreme thermotolerance exhibited by enzymes derived from such sources, with a half-life of more than 0.5 h at 363 K. Structural investigations should help to further understand the mechanism of these enzymes and provide some insights into the property of protein thermostability. Here, we describe the crystallization of the enzyme in a form suitable for high-resolution X-ray studies.

2. Experimental

The *pcp* gene from *Thermococcus litoralis* strain NS-C was isolated and overexpressed in *E. coli* as described previously (Singleton *et al.*, unpublished data). For crystallization, the protein was concentrated to 10 ml^{-1} in 50 mM potassium phosphate pH 7.5, 10 mM DTT by ultrafiltration using Centricon 10 concentrators (Amicon) in the same buffer, and crystallization trials were set up using the hanging-drop vapour-diffusion method. Initial screening was carried out with the sparse-matrix conditions described by Jancarik & Kim

Table 1
Data-collection statistics.

Resolution shell (Å)	Number of reflections	Completeness (%)	R_{sym}^\dagger	$I/\sigma(I)$	$I > 2\sigma$ (%)
5.94	2882	96.42	0.075	18.13	86.0
4.72	5646	97.08	0.096	10.27	84.4
4.12	8323	96.87	0.094	9.32	86.9
3.75	10942	96.18	0.138	6.16	79.3
3.48	13526	95.87	0.203	4.25	70.5
3.27	16093	95.51	0.241	3.43	64.1
3.11	18681	95.27	0.312	2.53	53.8
2.98	21139	94.70	0.361	2.17	48.7
2.86	22766	90.90	0.340	2.05	46.7

$^\dagger R_{\text{sym}} = (\sum |I - \langle I \rangle|) / \sum I$, where I is the observed intensity and $\langle I \rangle$ is the average intensity of symmetry-related reflections.

(1991) using a Crystal Screen kit (Hampton Research, Laguna Hills, CA) in addition to a range of ammonium sulfate concentrations.

Diffraction studies were carried out using a Siemens Hi-Star area detector with graphite-monochromated $\text{Cu K}\alpha$ radiation from a Siemens rotating-anode generator operating at 50 kV and 100 mA. Frames were indexed and processed using the programs *FRAMBO* and *SAINT* (Siemens Industrial Automation, Inc.).

3. Results and discussion

Needle-shaped crystals of dimensions $0.8 \times 0.1 \times 0.05$ mm grew from 1 M ammonium dihydrogen orthophosphate, 0.1 M citrate pH 5.6 in approximately two weeks. Crystals of similar morphology but with larger dimensions ($0.5 \times 0.1 \times 0.1$ mm) also grew from 35% ammonium sulfate, 50 mM

potassium phosphate pH 7.5. By reducing the reservoir buffer to pH 6.5, still larger crystals of $1.0 \times 0.3 \times 0.15$ mm could be grown. X-ray analysis showed crystals from both precipitants to have identical cell dimensions and space group, so the larger crystals grown from ammonium sulfate were used for all further work.

These crystals showed diffraction to 2.7 \AA , with weak reflections visible at 2.1 \AA on the

rotating-anode generator. The unit cell was determined to be orthorhombic with refined lattice dimensions of $a = 95.06$, $b = 150.06$, $c = 73.54 \text{ \AA}$. An examination of systematic absences indicated space group $P2_12_12$. This allows two or four subunits per asymmetric unit. Assuming four subunits gives a Matthews coefficient of $2.73 \text{ \AA}^3 \text{ Da}^{-1}$ and a solvent content of 55%, while two subunits give values of $5.46 \text{ \AA}^3 \text{ Da}^{-1}$ and 77% for the Matthews coefficient and solvent content, respectively, which are extreme values for a water-soluble protein (Matthews, 1968). Thus, it is most likely that the asymmetric unit contains one tetramer of the enzyme.

A complete data set was collected at room temperature using 0.2° frames comprising 49 152 observations merging to 23 309 unique reflections with an overall $R_{\text{sym}} = 11.2\%$. Some statistics of the data collection are shown in Table 1.

The crystals have been shown to diffract to beyond 1.8 \AA resolution on station 9.5 of the Daresbury Synchrotron Radiation Source, and a search for suitable heavy-atom derivatives for phasing purposes is currently in progress.

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