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## Studies on the Active Site of Deacetoxycephalosporin C Synthase

Matthew D. Lloyd<sup>1\*</sup>, Hwei-Jen Lee<sup>1</sup>, Karl Harlos<sup>2</sup>, Zhi-Hong Zhang<sup>1</sup>  
 Jack E. Baldwin<sup>1</sup>, Christopher J. Schofield<sup>1</sup>, John M. Charnock<sup>3</sup>  
 C. David Garner<sup>3</sup>, Takane Hara<sup>4</sup>, Anke C. Terwisscha van Scheltinga<sup>5</sup>  
 Karin Valegård<sup>4</sup>, Jenny A. C. Viklund<sup>4</sup>, Janos Hajdu<sup>4</sup>, Inger Andersson<sup>5</sup>  
 Åke Danielsson<sup>6</sup> and Rama Bhikhabhai<sup>6</sup>

<sup>1</sup>The Dyson Perrins Laboratory and the Oxford Centre for Molecular Sciences, South Parks Road, Oxford OX1 3QY UK

<sup>2</sup>The Laboratory of Molecular Biophysics and Oxford Centre for Molecular Sciences, South Parks Road, Oxford OX1 3QU UK

<sup>3</sup>Department of Chemistry University of Manchester Manchester M13 9PL, UK

<sup>4</sup>Department of Biochemistry Uppsala University, Box 576 S-751 23 Uppsala, Sweden

<sup>5</sup>Department of Molecular Biology, Swedish University of Agricultural Sciences, P.O. Box 590, S-751 24 Uppsala, Sweden

<sup>6</sup>Amersham Pharmacia Biotech S-751 84 Uppsala, Sweden

The Fe(II) and 2-oxoglutarate-dependent dioxygenase deacetoxycephalosporin C synthase (DAOCS) from *Streptomyces clavuligerus* was expressed at ca 25% of total soluble protein in *Escherichia coli* and purified by an efficient large-scale procedure. Purified protein catalysed the conversions of penicillins N and G to deacetoxycephems. Gel filtration and light scattering studies showed that in solution monomeric apo-DAOCS is in equilibrium with a trimeric form from which it crystallizes. DAOCS was crystallized  $\pm$ Fe(II) and/or 2-oxoglutarate using the hanging drop method. Crystals diffracted to beyond 1.3 Å resolution and belonged to the R3 space group (unit cell dimensions:  $a = b = 106.4$  Å,  $c = 71.2$  Å;  $\alpha = \beta = 90^\circ$ ,  $\gamma = 120^\circ$  (in the hexagonal setting)). Despite the structure revealing that Met180 is located close to the reactive oxidizing centre of DAOCS, there was no functional difference between the wild-type and selenomethionine derivatives. X-ray absorption spectroscopic studies in solution generally supported the iron co-ordination chemistry defined by the crystal structures. The Fe K-edge positions of 7121.2 and 7121.4 eV for DAOCS alone and with 2-oxoglutarate were both consistent with the presence of Fe(II). For Fe(II) in DAOCS the best fit to the Extended X-ray Absorption Fine Structure (EXAFS) associated with the Fe K-edge was found with two His imidazolate groups at 1.96 Å, three nitrogen or oxygen atoms at 2.11 Å and one other light atom at 2.04 Å. For the Fe(II) in the DAOCS-2-oxoglutarate complex the EXAFS spectrum was successfully interpreted by backscattering from two His residues (Fe-N at 1.99 Å), a bidentate O,O-co-ordinated 2-oxoglutarate with Fe-O distances of 2.08 Å, another O atom at 2.08 Å and one at 2.03 Å. Analysis of the X-ray crystal structural data suggests a binding mode for the penicillin N substrate and possible roles for the C terminus in stabilising the enzyme and ordering the reaction mechanism.

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**Keywords:** antibiotic biosynthesis; dioxygenase; EXAFS; iron; 2-oxoglutarate

\*Corresponding author

Abbreviations used: ACV, L- $\delta$ -( $\alpha$ -aminoadipoyl)-L-cysteinyl-D-valine; ADH, alcohol dehydrogenase; AUFS, absorbance units full scale; 7-ADCA, 7-aminodeacetoxycephalosporanic acid; 6-APA, 6-aminopenicillanic acid; D-AA, D- $\delta$ -( $\alpha$ -aminoadipoyl)-; BSA, bovine serum albumin; DAC, deacetylcephalosporin C; DACS, deacetylcephalosporin C synthase; DAOC, deacetoxycephalosporin C; DAOCS, deacetoxycephalosporin C synthase; DTNB, 5,5'-dithio-bis(2-nitrobenzoic acid); ESI MS, electrospray ionisation mass spectrometry; EXAFS, extended X-ray absorption fine structure; G-7-ADCA, phenylacetyl-7-aminodeacetoxycephalosporanic acid; IU, nmol/minute; L-AA, L- $\delta$ -( $\alpha$ -aminoadipoyl)-; N/D, not determined; PMSF, phenylmethylsulfonyl fluoride; PEI, polyethylenimine; TCA, trichloroacetic acid; TSP, 2,2',3,3'-[<sup>2</sup>H<sub>4</sub>]trimethylsilylpropanoate, sodium salt; XAS, X-ray absorption spectroscopy; ' denotes residues derived from a neighbouring molecule in the crystal structure.

E-mail address of the corresponding author: [matthew.lloyd@chem.ox.ac.uk](mailto:matthew.lloyd@chem.ox.ac.uk)

## Introduction

The first committed step in the biosynthesis of cephalosporins is expansion of the five-membered thiazolidine ring of penicillin N (1) to deacetoxycephalosporin C (DAOC, 2), which contains the characteristic six-membered cephem ring. Subsequent hydroxylation of DAOC (2) leads to deacetylcephalosporin C (DAC, 3; Figure 1). In prokaryotes, e.g. *Streptomyces clavuligerus*, the conversions of (1) to (2) and of (2) to (3) are catalysed (at least predominately) by separate enzymes, deacetoxycephalosporin C synthase (DAOCS) and deacetylcephalosporin C synthase (DACS). In eukaryotes, e.g. *Cephalosporium acremonium*, both steps are catalysed by a single, bifunctional enzyme (DAOC/DAC synthase). In *C. acremonium* DAC (3) is converted by acetylation into cephalosporin C, whilst in prokaryotes alternative pathways lead to other cephems, e.g. in *S. clavuligerus* DAC (3) is converted *via* several steps into cephamycin C (4) (for reviews see: Baldwin & Abraham, 1988; Baldwin & Schofield, 1992). Medicinally useful cephalosporins are produced by *in vitro* removal of the D- $\delta$ -( $\alpha$ -aminoadipoyl)-side-chain from a fermented cephalosporin followed by reacylation with the requisite side-chain (see e.g. Crawford *et al.*, 1995). Alternatively, useful cephems may be obtained by the chemical ring expansion of the corresponding penicillin (Morin *et al.*, 1963; Chauvette *et al.*, 1971; Colvin, 1992).

The three oxygenases involved in the conversion of penicillin N (1) to DAOC (2) and DAC (3) are

members of the non-heme Fe(II)-dependent family of dioxygenases (Prescott, 1993; Que & Ho, 1996; Hegg & Que, 1997). Each reaction has a stoichiometric requirement for 2-oxoglutarate and (presumably) dioxygen and *in vitro* optimum substrate conversion requires the presence of DTT and ascorbate. The hydroxylation of (2) to give (3) may be regarded as a typical type of reaction catalysed by 2-oxoglutarate utilising dioxygenases, but the oxidative ring expansion of (1) to give (2) is unique. Labelling studies on the mechanism of ring expansion indicate that cleavage of the C-H bond of the  $\beta$ -methyl group precedes that of the C-2 C-H bond (Baldwin *et al.*, 1991) and have provided evidence for a methylene radical intermediate (Pang *et al.*, 1984; Townsend *et al.*, 1985).

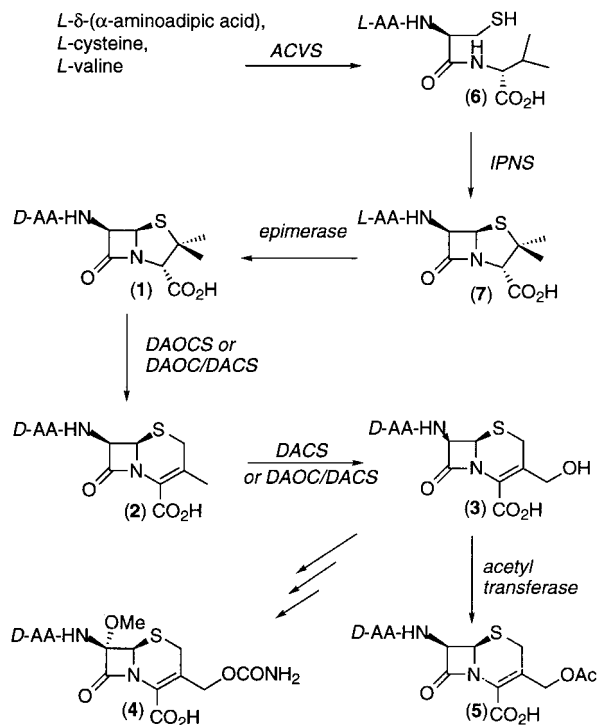
Isopenicillin N synthase (IPNS), for which crystal structures have been described (Roach *et al.*, 1995, 1997), catalyses the oxidative bicyclisation of the tripeptide, L- $\delta$ -( $\alpha$ -aminoadipoyl)-L-cysteinyl-D-valine (ACV, 6) to isopenicillin N (7), the immediate precursor of penicillin N (1) (Figure 1). IPNS does not require a 2-oxoglutarate cosubstrate since it catalyses a four-electron oxidation of (6). The cephalosporin oxygenases clearly have a close evolutionary relationship with IPNS (Cooper, 1993) and comparison of their active sites is of interest, particularly with regard to defining the interactions responsible for 2-oxoglutarate binding and turnover.

DAOCS from *S. clavuligerus* was first cloned and over-expressed in *Escherichia coli* as insoluble protein, which was refolded to give soluble, active enzyme for characterization but was unsuitable for crystallization trials (Dotzlar & Yeh, 1989; Kovacevic *et al.*, 1989). Subsequently, soluble DAOCS was expressed using a plasmid containing the *trc* promoter, but the level of expression and purification methods were not optimal (Morgan *et al.*, 1994). Herein, we report the over-production, efficient purification, crystallization and characterization of DAOCS. The results have implications for the interpretation of the recently reported DAOCS crystal structure (Valegård *et al.*, 1998) and suggest roles for the C termini of DAOCS and related 2-oxoglutarate-dependent dioxygenases.

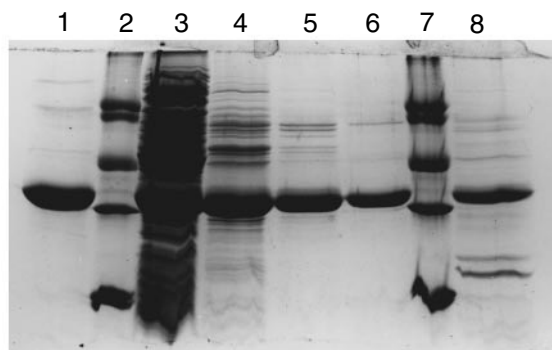
## Results and Discussion

### Expression and purification of recombinant DAOCS

High-level expression of the DAOCS gene to produce soluble protein was achieved *via* sub-cloning of the DNA fragment containing the DAOCS gene into the pET11a and pET24a vectors. The resultant plasmids (pML1 and pHL1, respectively) were transformed into *E. coli* BL21 (DE3). In both cases expression at >25% of the soluble cell protein was observed two to three hours after induction by IPTG when grown at 27 for 30°C. This level of expression was ca ten times better than that previously obtained using *E. coli* NM554/pNM88



**Figure 1.** The biosynthesis of penicillins, cephalosporins and cephamycins (names of compounds (1) to (7) are given in the text).



**Figure 2.** SDS-PAGE analysis of wild-type DAOCS purified by ion-exchange, hydrophobic interaction and gel filtration chromatographies: 1, DAOCS standard (15  $\mu\text{g}$ ); 2, markers (9  $\mu\text{g}$ ); 3, crude extract (125  $\mu\text{g}$ ); 4, Q-Sepharose<sup>®</sup> HP fractions (34  $\mu\text{g}$ ); 5, hydrophobic interaction fractions (26  $\mu\text{g}$ ); 6, S-75 fractions (10  $\mu\text{g}$ ); 7, markers (9  $\mu\text{g}$ ); 8, DAOCS purified from *E. coli* NM554/pNM88(10  $\mu\text{g}$ ) (Morgan *et al.*, 1994). Markers (BDH, 10  $\mu\text{g}$  of each protein) were: ovotransferrin (76 and 78 kDa); BSA (66 kDa); ovalbumin (42.7 kDa); carbonic anhydrase (30 kDa); myoglobin (17.2 kDa); cytochrome c (12.3 kDa).

(Morgan *et al.*, 1994) as judged by SDS-PAGE. Soluble expression was not observed when cells were grown at 37 °C. The pET24a vector was preferred since it allows the use of single-stranded mutagenesis techniques. Unlike the pET11a vector, the pET24a vector uses kanamycin rather than ampicillin (i.e.  $\beta$ -lactamase mediated) resistance as the selection method and although we could not find any evidence for contamination of purified DAOCS by  $\beta$ -lactamase, the use of this vector reduces this possibility.

In the optimised method, crude extracts were purified using anion-exchange, hydrophobic interaction and gel filtration chromatographies (Figure 2). The use of SOURCE<sup>®</sup> 15ISO resin enabled very rapid (less than 30 minutes) concentration and purification of the pool from a Q-Sepharose<sup>®</sup> HP column. It was possible to purify 150–200 mg of active, crystallography-grade DAOCS within 24 hours (Table 1).

Electrospray ionisation mass spectrometry (ESI MS) of DAOCS gave a mass of 34,554 ( $\pm 3$ ) Da in agreement with the calculated value (34,551 Da; Figure 3(a)). The predicted N-terminal amino acid sequence (including the N-terminal methionine) was obtained for the first 15 residues (data not shown). The UV-visible spectrum of wild-type

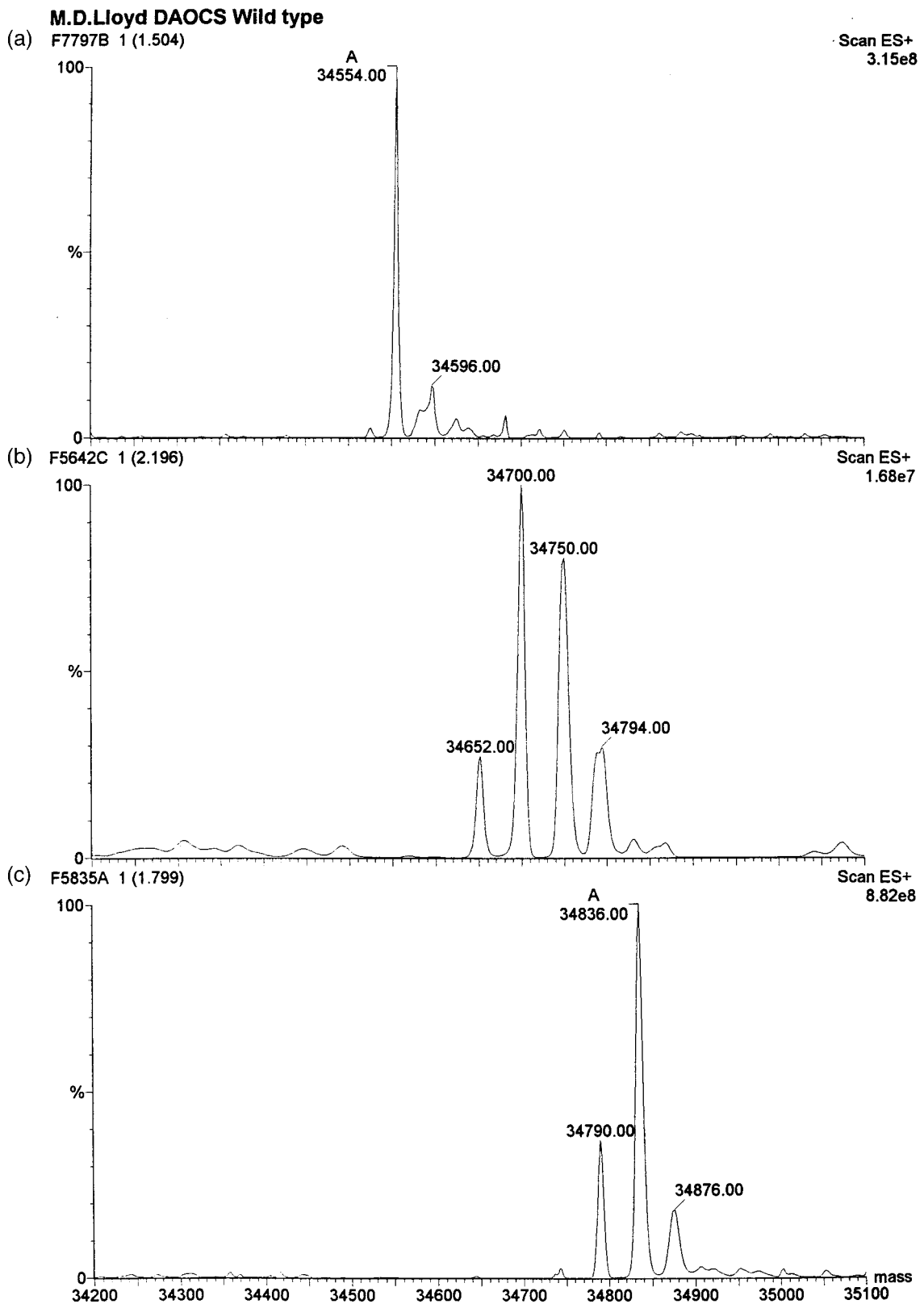
apo-DAOCS showed a absorbance maximum at 278 nm ( $\epsilon = 33,578 \text{ M}^{-1} \text{ cm}^{-1}$ ), with no significant absorbance above 300 nm. DTNB titrations (Habeeb, 1974) in the presence/absence of SDS were consistent with the presence of seven cysteine residues as predicted. In the absence of SDS, four of the cysteine residues were derivatized significantly faster than the other three. Examination of the DAOCS structure (Valegård *et al.*, 1998; K. Harlos & M. D. Lloyd, unpublished results) suggests that all of the cysteine residues are relatively exposed to solvent but it was not possible to correlate specific residues with the observed DTNB titration data.

As noted previously (Sami *et al.*, 1997; Zhang *et al.*, 1997) the errors in apparent  $K_m$  and  $k_{cat}$  values determined from preliminary steady-state kinetic analysis for Fe(II)-dependent oxidizing enzymes may well be significant and caution should be exercised in drawing mechanistic conclusions from relatively small (i.e. less than an order of magnitude) differences. Purified DAOCS had the following apparent kinetic parameters for penicillin N (1):  $K_m = 6.6 (\pm 0.8) \mu\text{M}$  and  $k_{cat} = 0.42 (\pm 0.05) \text{ s}^{-1}$  (specific activity = 727 ( $\pm 86$ ) nmol/min per mg). Adipoyl-6-APA (8) has been previously shown to be a (poor) substrate for the enzyme (Shibata *et al.*, 1996), being converted into adipoyl-7-ADCA (9) with an apparent  $K_m = 1.30 (\pm 0.10) \text{ mM}$  and  $k_{cat} = 0.073 (\pm 0.0003) \text{ s}^{-1}$  (127 ( $\pm 5.3$ ) nmol/min per mg; Figure 4). Penicillin G (10) was also converted to the corresponding cephem (11) with the following apparent kinetic parameters:  $K_m = 2.03 (\pm 0.12) \text{ mM}$ ;  $k_{cat} = 0.062 (\pm 0.003) \text{ s}^{-1}$  (specific activity = 108 ( $\pm 5.2$ ) nmol/min per mg). These are similar to the previously reported values ( $K_m = 1 \text{ mM}$ ;  $V_{max} = 87 \text{ nmol/min per mg}$ ; Morgan, 1994). In preparative conversions using a higher enzyme: substrate ratio, ca 20–30% conversion of penicillin G (10) to cephem (11) was observed. Analogous preparative experiments with penicillin N (1) demonstrated >95% conversion to DAOC (2).

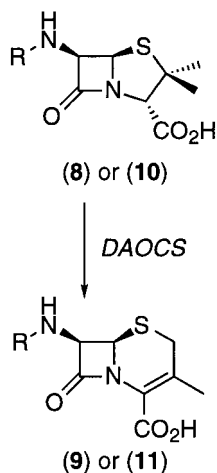
Based on the present analyses the differences in  $k_{cat}$  for the three substrates analysed are probably insignificant. The differences in the apparent  $K_m$  values for penicillin N (1) compared to penicillin G (10) and adipoyl-6-APA (Shibata *et al.*, 1996) are greater being ca 100-fold. Interpretation of these apparent  $K_m$  values as indicators of binding ability for different substrates to DAOCS should be regarded as tentative, but the available data, implying that penicillin G (10) and adipoyl-6-APA (8) bind to DAOCS with similar affinities, suggest

**Table 1.** Purification of DAOCS from *E. coli* BL21 (DE3)/pML1

Purification step	Protein (mg/ml)	Total protein (mg)	IU/mg	Total IU
Crude extract	5.70	1727	N/D	N/D
Q-Sepharose <sup>®</sup> HP (13–22)	1.60	400	42	16,800
SOURCE <sup>®</sup> 15ISO (3–7)	2.08	208	39	8112
Superdex <sup>®</sup> 75 (18–32, 49–62)	1.11	169	66	11,154



**Figure 3.** (a) ESI MS of recombinant wild-type DAOCS (top); (b) ESI MS of the recombinant selenomethionyl-DAOCS (grown with 25  $\mu\text{g}/\text{ml}$  SeMet) (centre); (c). ESI MS of the recombinant selenomethionyl-DAOCS (grown with 50  $\mu\text{g}/\text{ml}$  SeMet) (below). Note the presence of a heavier species at 34,875 ( $\pm 2.3$ ) Da. The mass difference (ca 40 Da) is inconsistent with the incorporation of an additional selenium atom (ca 47 Da) or oxidation (32 or 48 Da for two or three additional oxygen atoms). It is possible that this species represents an alternative modification(s) or a contaminant.



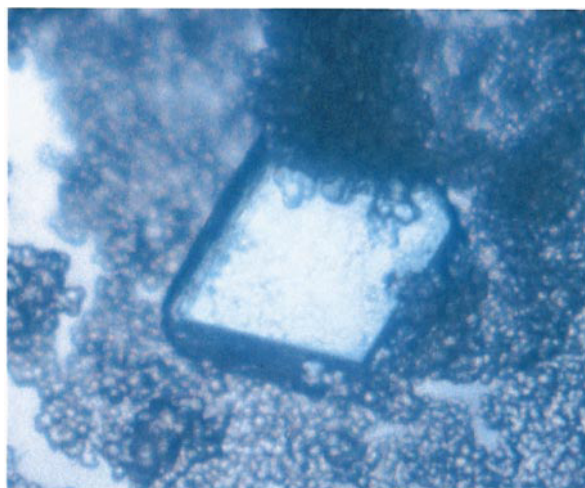
- (8), (9)      R = adipoyl  
 (10), (11)    R = phenylacetyl

**Figure 4.** Conversion of adipoyl-6-APA (8) to adipoyl-7-ADCA (9), and penicillin G (10) to phenylacetyl-7-ADCA (11) by DAOCS.

that the amino group plays an important role in the binding of the  $\alpha$ -aminoadipoyl-side-chain of penicillin N (1).

IPNS has a very lax substrate selectivity and is able to catalyse the cyclisation of many analogues of ACV (6) (Baldwin & Bradley, 1990), including the relatively efficient conversion of adipoyl-L-cysteinyl-D-valine to adipoyl-6-APA (8) and D-( $\alpha$ -aminoadipoyl)-L-cysteinyl-D-valine to penicillin N (1) (Baldwin *et al.*, 1984, 1990). In contrast DAOCS apparently has tighter selectivity, at least with respect to the presence of the amino group in the side-chain of its substrates. This may be related to the evolution of the pathway and the fact that penicillin N (1) but not isopenicillin N (7) is a substrate for DAOCS and DAOC/DACS.

Various metals, including cadmium, cobalt, chromium, copper and manganese salts were screened for their ability to catalyse the conversion of penicillin N (1) to DAOC (2) and as inhibitors of the reaction (data not shown). Catalysis was only observed in the presence of added  $\text{Fe}^{2+}$  or  $\text{Fe}^{3+}$  (the latter has ca 34% relative specific activity to the former under standard incubation conditions), with the activity observed in the latter case presumably resulting from reduction in solution. When 2 mM  $\text{Fe(III)}$  was added to the standard incubation mixture containing 2 mM  $\text{Fe(II)}$  it was inhibitory (>80% inhibition).  $\text{Cd(OAc)}_2$  and  $\text{CuCl}_2$  effected complete inhibition at 2 mM, whilst 2 mM  $\text{CoCl}_2$  caused ca 50% inhibition and  $\text{NiSO}_4$  caused ca 25% inhibition. The other metal salts tested ( $\text{CrK(SO}_4)_2$ ,  $\text{MnCl}_2$  and  $\text{SnCl}_2$ ) showed little or no effect on activity.



**Figure 5.** An R3 crystal of wild-type DAOCS.

### Crystallization of DAOCS

The first conditions leading to DAOCS crystals used 100 mM Tris-HCl pH 8.5 buffer containing 2 mM 2-oxoglutarate and ammonium sulphate. These were refined to pH 7.0 and 100 mM Hepes-NaOH buffer containing 5 mM monopotassium 2-oxoglutarate. Crystals with dimensions of 0.2 mm  $\times$  0.2 mm  $\times$  0.2 mm appeared after three to eight weeks (Figure 5). The space group of the crystal was R3 and with a single DAOCS molecule in the asymmetric unit, with the following unit cell dimensions:  $a = b = 106.4 \text{ \AA}$ ,  $c = 71.2 \text{ \AA}$ ;  $\alpha = \beta = 90^\circ$ ,  $\gamma = 120^\circ$  (in the hexagonal setting). The predicted density  $V_M$  was  $2.65 \text{ \AA}^3/\text{Da}$  (Matthews, 1968). All crystals obtained so far were merohedrally twinned with twinning ratios between 0.06 and 0.45 (Terwisscha van Scheltinga *et al.*, unpublished results) and diffracted X-rays beyond 1.3  $\text{\AA}$  resolution (Valegård *et al.*, 1998).

### EXAFS

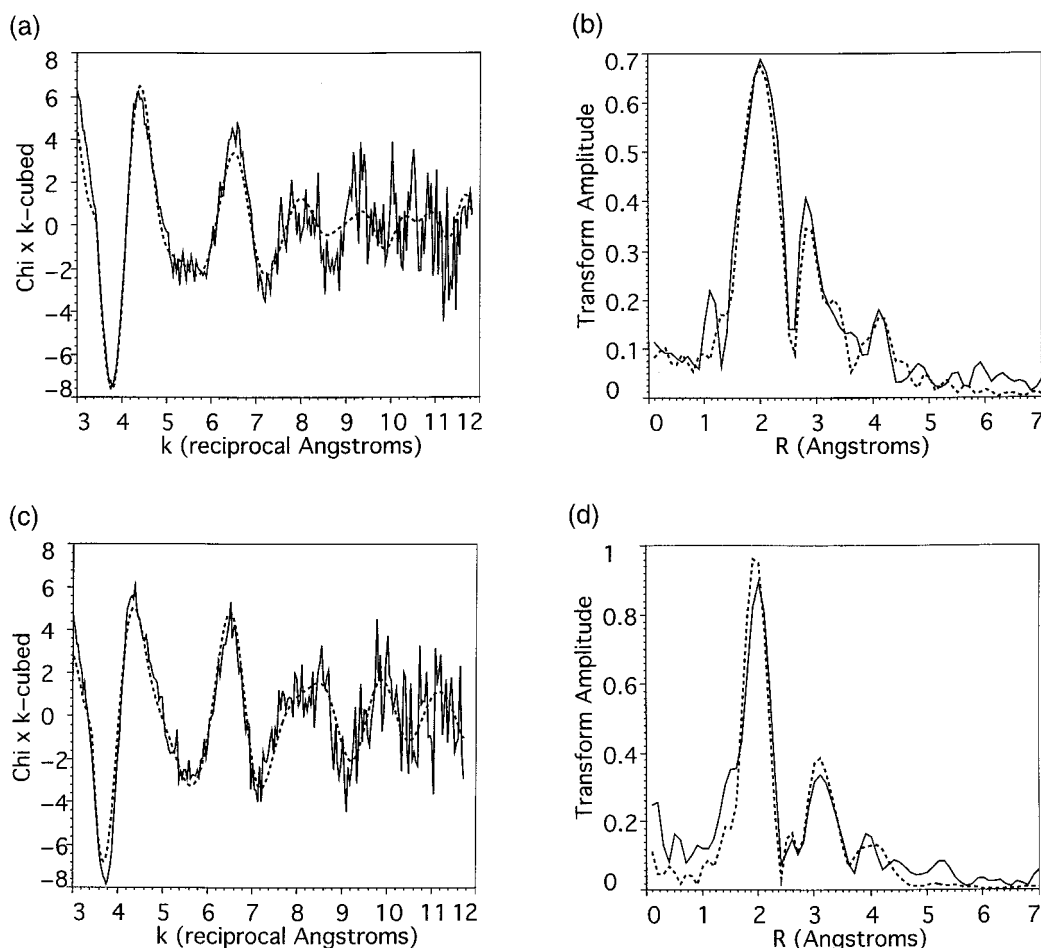
EXAFS studies were carried out in order to compare the co-ordination chemistry of DAOCS in solution and crystalline states. This is important because studies on IPNS (Roach *et al.*, 1995, 1997) indicate that significant conformational changes affecting the active site and possibly the co-ordination chemistry of iron(II)-dependent oxidizing enzymes occur upon substrate binding. Significant aspects of the mechanism of 2-oxoglutarate-dependent dioxygenases also remain unclear; for example, it is not known if the 2-oxoglutarate is ligated to the iron in a monodentate or bidentate fashion upon binding and reaction of dioxygen, and at what stage carbon dioxide is released from the iron during catalysis.

The Fe K-edge positions for  $\text{Fe(II)}$  DAOCS alone and in the presence of 2-oxoglutarate of 7121.2 and 7121.4 eV, respectively, are both consistent with the  $\text{Fe(II)}$  oxidation state (Randall *et al.*, 1993). The

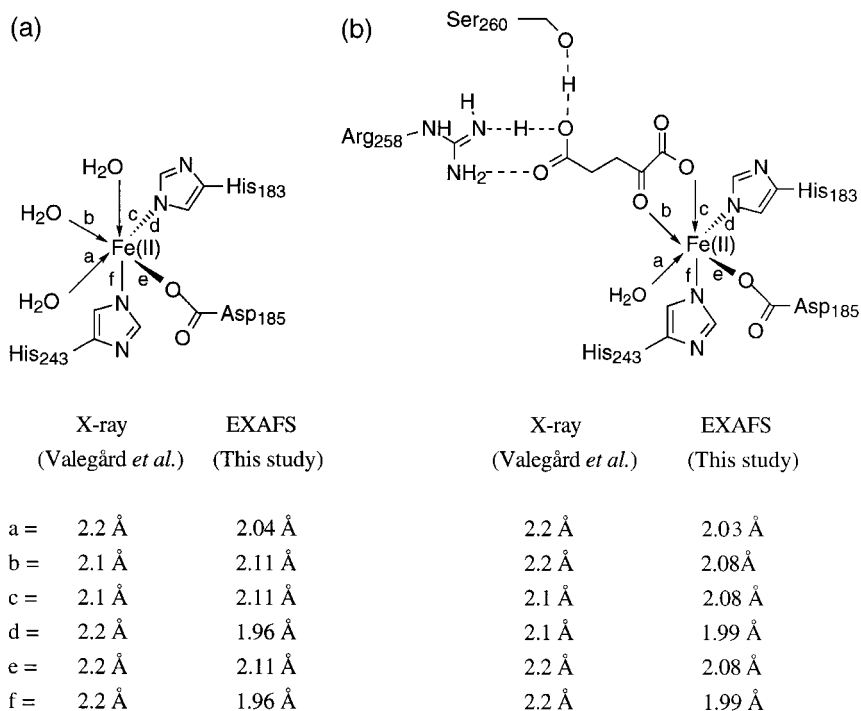
Fe K-edge EXAFS and Fourier transform data (Figure 6(a)) from the Fe(II)-DAOCS complex showed contributions from several shells of backscatterers. The backscattering contributions from the inner shell were successfully simulated using light atoms (C, N, O) only, with no evidence of coordination from heavier elements such as sulphur. Features in the Fourier transform at ca 3 Å and at ca 4.2 Å are characteristic of imidazole groups (Figure 6(a) and (b)). Fits were attempted using 1-3 imidazole groups, including multiple scattering, with other light atoms to complete the inner coordination sphere. The best fit was found with two imidazole groups co-ordinating to the iron with Fe-N of 1.96 Å, three oxygen (or nitrogen) atoms at 2.11 Å, and one other light atom at 2.04 Å. A Fe-N bond distance of 1.96 Å is more consistent with an imidazolate ligand rather than imidazole. These conclusions are consistent with the results obtained

from the crystal structure of the DAOCS-Fe(II) complex (Valegård *et al.*, 1998), which show octahedral ligation of the iron by His183, His243, Asp185 and three water molecules (Figure 7(a)).

However, the experimental and simulated EXAFS of the DAOCS-Fe(II) complex were not in complete agreement, in particular a feature in the Fourier transform at ca 3.3 Å was not readily fitted by the data analysis given above. We attempted to reproduce this feature by including backscattering from a single heavy atom such as iron or by several light atoms. However, reproduction of the magnitude of the peak using light atoms required unrealistically low or negative Debye-Waller factors. The use of a heavier backscatterer gave the correct amplitude in the Fourier transform but the improvement in fit of the EXAFS was insufficient to justify the inclusion of the additional parameters. The best fit was found using a light atom



**Figure 6.** Fe K-edge EXAFS data in 10 M Tris-HCl at ca 80 K and its Fourier transformations (continuous lines) together with the optimum simulation achieved (broken lines). (a) DAOCS-Fe(II) complex (above), optimum simulation: three O atoms at 2.11 Å (Debye-Waller parameter,  $2\sigma^2$ , 0.008 Å<sup>2</sup>); two imidazole groups with Fe-N of 1.96 Å (0.002 Å<sup>2</sup>), Fe-C of 2.92 Å (0.004 Å<sup>2</sup>), Fe-C/N of 4.16 Å (0.008 Å<sup>2</sup>); plus a linear Fe-X-Y group with X atom at 2.04 Å (0.004 Å<sup>2</sup>) and Y atom at 3.27 Å (0.005 Å<sup>2</sup>). (b) DAOCS-Fe(II)-2-oxoglutarate complex (below), optimum simulation: one atom at 2.08 Å (Debye-Waller parameter,  $2\sigma^2$ , 0.004 Å<sup>2</sup>); two imidazole groups with Fe-N of 1.99 Å (0.001 Å<sup>2</sup>), Fe-C of 2.92 Å (0.004 Å<sup>2</sup>), Fe-C/N of 4.10 Å (0.012 Å<sup>2</sup>); a 2-oxocarboxylate group with two Fe-O of 2.08 Å (0.002 Å<sup>2</sup>), Fe-C of 2.80 Å (0.007 Å<sup>2</sup>), Fe-C of 4.41 Å (0.004 Å<sup>2</sup>); plus a linear Fe-X-Y group with X at 2.03 Å (0.003 Å<sup>2</sup>) and Y atom at 3.33 Å (0.004 Å<sup>2</sup>). The estimated uncertainty in the inner shell distances is  $\pm 0.03$  Å.



**Figure 7.** Co-ordination of the Fe(II) atom in DAOCS (a) without 2-oxoglutarate (left); (b) with 2-oxoglutarate (right). The ligand opposite His183 is shown as a water molecule as in the crystal structures. The EXAFS data, however, suggest the possible presence of an alternative ligand in this position; see the text for details. This species is assigned as being *trans* to His183 since the EXAFS data indicate its presence in both the DAOCS-Fe(II) and DAOCS-Fe(II)-2-oxoglutarate complexes.

(X = O/N/C) at 2.04 Å and another light atom (Y) at 3.27 Å with the inclusion of multiple scattering for a linear Fe-X-Y group (see below).

The EXAFS associated with the Fe K-edge of the DAOCS-Fe(II)-2-oxoglutarate complex and its Fourier transform (Figure 6(b)), whilst similar to the corresponding information for the DAOCS-Fe(II) complex, showed significant differences (notably in the EXAFS between 8 and 10 Å<sup>-1</sup>), indicating that the Fe(II) site has been perturbed by the presence of 2-oxoglutarate. The EXAFS was successfully interpreted by backscattering from two histidine groups (Fe-N at 1.99 Å) with a bidentate O,O-co-ordinated 2-oxoglutarate with Fe-O distances of ca 2.08 Å, with another O atom at this distance and a light (O or N) atom at 2.03 Å (Figure 6(b)). As for the DAOCS-Fe(II) complex, a successful interpretation of the Fe K-edge EXAFS and its Fourier transform required inclusion of backscattering from a linear Fe-X-Y arrangement with the light atoms X and Y being 2.03 and 3.33 Å from the iron. Thus, the Fe K-edge EXAFS imply the 2-oxoglutarate replaces two water molecules bound to the iron in the Fe(II) DAOCS complex, which is entirely consistent with the crystal structure of the same complex (Valegård *et al.*, 1998) which shows bidentate binding of 2-oxoglutarate to the iron, *via* the keto-oxygen and one oxygen of the carboxylate group (Figure 7(b)). Since the unusual feature in the EXAFS at ca 3.3 Å is present both before and after 2-oxoglutarate binding the simplest explanation is that it results from a ligand binding to the iron *trans* to His183, the proposed binding site for dioxygen. However, in the crystal structure this ligand is clearly a water molecule. The identity of the Fe-X-Y group proposed on the basis of the EXAFS studies is unknown; an average

X-Y distance of ca 1.27 Å is indicative of a bonded contact but the linearity of the arrangement is surprising and appears to preclude X and Y coming from the carboxylate group of Asp185. Contaminants derived from dioxygen (such as superoxide or peroxide) can probably be excluded, since the Fe K-edge position indicates the presence of iron(II) at the active site. The possibility that this feature arises due to ligation of sodium azide (derived from the buffer) by the iron is also unlikely because a non-linear Fe-N bond would be expected. This discrepancy between the crystallographic and frozen solution EXAFS studies will have to be resolved by further experimentation and analysis.

Thus, the EXAFS study provides evidence that the crystal structure reflects solution binding of 2-oxoglutarate (Figure 7(b)). Although a reasonable proposal, care should be taken in assuming that the 2-oxoglutarate remains bound to the iron in a bidentate manner at all times during the catalytic cycle prior to fragmentation of 2-oxoglutarate into carbon dioxide and succinate. It is possible that binding of dioxygen and/or substrate triggers a rearrangement to monodentate binding. Spectroscopic studies on 2,4-(dichlorophenoxy)acetate/2-oxoglutarate dioxygenase complexed to Cu(II) (substituting for Fe(II)) and 2-oxoglutarate indicate monodentate binding of 2-oxoglutarate and the carboxylate of the substrate (Whiting *et al.*, 1997). However, it is likely that the substitution of Cu(II) for Fe(II) significantly perturbs the co-ordination chemistry and more recent spectroscopic studies on complexes formed between clavaminatase synthase, iron and 2-oxoglutarate indicate bidentate binding of the latter (Pavel *et al.*, 1998).



The largest discrepancies between the bond lengths obtained from X-ray crystallographic and EXAFS measurements are for the iron(II)-histidinyll residue bonds. Bond lengths obtained by X-ray crystallography suggest imidazole ligands, whilst the EXAFS measurements are more consistent with imidazolate ligands. These differences may be partially a consequence of the different buffer conditions used in these studies, Hepes-NaOH (pH 7.0) with ammonium sulphate in the former case and 50 mM Tris-HCl (pH 7.5), 0.3 mM NaN<sub>3</sub> in the latter. However, in general the atomic distances between a metal and lighter ligands are usually over-estimated in refined X-ray structures of proteins. An analysis of available data demonstrates these distances are systematically longer than those obtained from EXAFS measurements. This suggests that the differences arise from technical procedures rather than inherent structural differences between crystalline and solution states. There may be a number of reasons for the apparent discrepancies. During refinement, in some cases, the iron-ligand distances are treated as non-bonded atoms without additional restraints. Under these conditions, non-bonded contact parameters dominate the refinement, which results in increased bond lengths. This mistake can be corrected for, e.g. by removing such restraints around the metal and its immediate environment, and fixing the distances in subsequent restrained refinement to the values obtained. This procedure often produces bond lengths which agree more closely with values obtained from EXAFS measurements. However, there is another factor to be considered. At the usual resolutions of protein structures (1.5-3.0 Å), series termination errors in Fourier synthesis can produce sizeable ripples around heavy metal scatterers, pushing apparent electron densities from lighter ligands to somewhat longer distances than they may be in reality. The number, depth and location of such ripples depends on the resolution and completeness of the data set. At very high resolutions (such as those in small molecule structures) these ripples become negligible. However, at the resolutions of most protein structures they will be present and systematically produce slightly longer distances than EXAFS measurements, even without the restraints in the X-ray refinement. A source of error in EXAFS measurements should also be considered in situations such as those described herein, where the inner co-ordination sphere of the metal involves atoms with very similar backscattering characteristics at similar but not identical distances. In the interpretation of the EXAFS it is difficult to unambiguously determine the individual contributions to the overall backscattering. Useful clarification can be obtained when outer shells provide multiple scattering contributions, e.g. those observed with imidazole groups.

### Role of methionine-180

The DAOCS crystal structures (Valegård *et al.*, 1998; K. Harlos & M.D. Lloyd, unpublished results) reveal that all (but one) methionine residues are located relatively close to the surface of the protein. The exception, methionine-180, is located within the active site and its side-chain becomes ordered upon binding of 2-oxoglutarate and/or iron(II). In the DAOCS-Fe(II)-2-oxoglutarate complex Met180 is in van der Waals contact with the C-2 atom of the 2-oxoglutarate (Valegård *et al.*, 1998; K. Harlos & M. D. Lloyd, unpublished results). The presence of an easily oxidizable side-chain so close to the reactive centre of DAOCS is somewhat surprising and it is probable that its oxidation to a sulphoxide or sulphone would affect the binding of 2-oxoglutarate and hence catalysis. Despite the observations that selenomethionine residues are more easily oxidized than methionine residues and crystallization of the selenomethionine derivative required anaerobic conditions, there were no apparent functional differences between the wild-type and selenomethionine derivative (Materials and Methods, Figure 3), nor was there any clear ESI MS evidence for oxidative modifications of the derivative. Similar specific activities were obtained for both, suggesting that the oxidative chemistry occurring within the DAOCS active site is tightly controlled and directed away from Met180, making the possibility of a methionine sulphoxide or radical intermediate during catalysis unlikely.

### Significance of the trimeric nature of DAOCS

The solution of the DAOCS structure (Valegård *et al.*, 1998; K. Harlos & M. D. Lloyd, unpublished results) demonstrates that it crystallizes as a trimer with one molecule in the asymmetric unit (Figure 8). The C-terminal residues from one DAOCS molecule project towards the active site of an adjacent sub-unit in a cyclic manner, forming a trimer. The key intermolecular interactions resulting from trimerization are shown in Table 2. Gel filtration analysis indicated that in solution apo-DAOCS exists as an equilibrium mixture of monomeric (est. 28.9 kDa) and trimeric (est. 92.9 kDa) forms. The presence of 5 mM 2-oxoglutarate had no apparent effect on the equilibrium position. Dynamic laser light scattering studies showed that apo-DAOCS is predominately trimeric at 15-20 °C, pH 7.0. At 20 °C the mean hydrodynamic radius was 4.2 nm, corresponding to an apparent mass of 96 kDa. The protein showed a bimodal size distribution in which the trimeric and monomeric forms are in equilibrium. This equilibrium was temperature-sensitive, and shifted towards the monomeric form at 15 °C (mean radius = 3.4 nm, 57 kDa). Addition of 10-100 mM 2-oxoglutarate alone did not effect the equilibrium position. Addition of 5-10 mM FeSO<sub>4</sub> alone to DAOCS induced a shift towards the monomeric form of the enzyme (mean

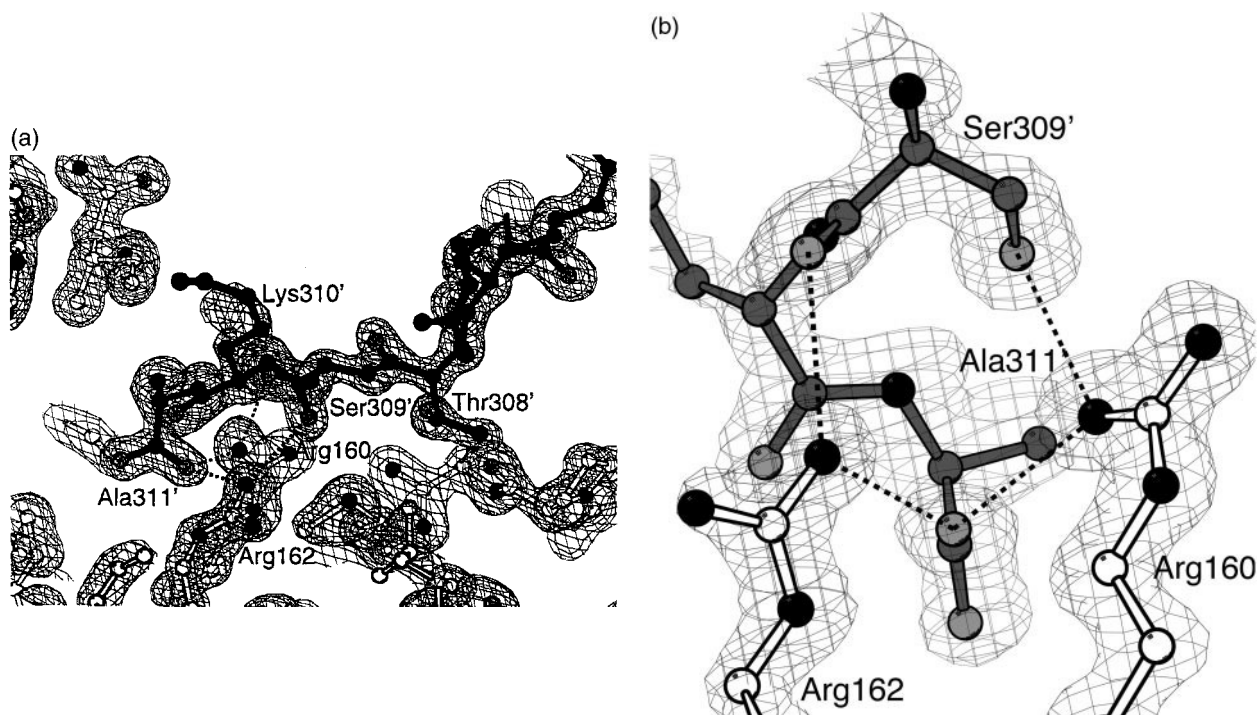
hydrodynamic radius 3.2 nm, 48 kDa at 20°C). Addition of 5-10 mM FeSO<sub>4</sub> with 25-50 mM 2-oxoglutarate resulted in almost complete dissociation to the monomeric form at 20-37°C (mean radius 2.9 nm, 39 kDa). These studies imply that it is the monomeric form of DAOCS which is catalytically active. Indirect support for this proposal came from X-ray crystallographic results (Valegård *et al.*, 1998), which show a clear movement of the C terminus upon binding of iron(II), which was predicted to cause dissociation into the monomer. Further oligomerisation *via* active site and C-terminal tail interactions may be partially responsible for the observation that DAOCS forms inclusion bodies when over-produced, particularly at 37°C. During refolding studies (data not shown) on DAOC/DACS an apparent temperature-dependent tendency to form oligomers was also noticed and this may also reflect the operation of an oligomerization process.

### Role of the C terminus and substrate binding

Comparison of the DAOCS and IPNS-Mn structures reveals strikingly similar general relationships between their active sites and C termini. In the structure of IPNS complexed to manganese (substituting for Fe(II)) Gln330 ligates to the metal, and thus the C terminus forms a "lid" over the active site, an arrangement also observed in the case of DAOCS. However, in the case of the DAOCS crystal structure the C-terminal lid is derived not from the same molecule, but from a neighbour in the trimeric unit. The active sites of

DAOCS and IPNS are unusual in that they are located within, rather than external to, the jelly roll barrel which forms the core of their structures. It is possible the presence of a C-terminal lid (which combined with other conformational changes would constitute a closed form of the enzyme) on the active site (in either monomeric or trimeric forms) stabilizes the jelly roll core of the resting enzyme, thereby reducing the possibility of unfolding and subsequent oxidative or proteolytic damage. Formation of the oligomers *in vivo* may also be significant if high concentrations of enzyme are present such as in an antibiotic production strain. The potential of the C terminus to adopt different conformations and its involvement in oligomerization should also be noted in respect of crystallization experiments on enzymes of the DAOCS family, in particular with regard to choice of mutant or isoenzyme for study.

Both IPNS and DAOCS are unstable *in vitro*, particularly under catalytic conditions. In part inactivation results from oxidative damage resulting from reactive oxidizing species (e.g. hydrogen peroxide) generated from the requisite cofactors (iron, ascorbate, dioxygen). ACCO is particularly sensitive to this type of oxidative damage, undergoing extensive fragmentation (Barlow *et al.*, 1997). However, it is clear that oxidative damage to a single form of the enzyme is not the only mode of inactivation and it is also unlikely that oxidation (which is likely to be significantly irreversible) explains the differences between the *in vitro* and *in vivo* inactivation rates. Upon standing in buffer recombinant ACCO from tomato fruit undergoes a confor-



**Figure 8.** (a) Close up view of the C-terminal arm interactions with the neighbouring sub-unit. (b) Close up view of #Ala311 and its interactions with Arg160 and Arg162. View is perpendicular to that in (a).

**Table 2.** Intermolecular hydrogen bonding interactions in R3 DAOCS crystals

Atom 1	Atom 2	Distance (Å)
O Pro69	N Ala240	2.9
O <sup>γ</sup> Thr72	O Gly282	2.6
O Gly79	N <sup>n1</sup> Arg242	3.0
O Gly79	N <sup>n2</sup> Arg242	3.0
N <sup>c</sup> Arg125	O <sup>δ2</sup> Asp214	3.0
N <sup>n1</sup> Arg160	O <sup>γ</sup> Ser309	2.9
N <sup>n1</sup> Arg160	OT2Ala311	2.8
N <sup>n2</sup> Arg162	O Ser309	2.9
N <sup>n2</sup> Arg162	OT2Ala311	2.7
N <sup>n1</sup> Arg266	O <sup>δ2</sup> Asp284	2.9
N Ser286	O Gly300	2.9
O <sup>γ</sup> Ser286	O-Gly299	2.8
N <sup>n1</sup> Trp297	O <sup>δ1</sup> Asn301	2.9

mational change to a less active form in which the main secondary structure elements are conserved. Supporting evidence for the proposal that a conformational change contributes to inactivation includes mutagenesis experiments which demonstrate that oxidative fragmentation of ACCO, which occurs as a burst upon addition of substrates/cofactors, only occurs in a form of the enzyme in which the iron is “incorrectly” ligated for catalysis (Zhang *et al.*, 1997).

The predicted inherent mobility in the C-terminal region of DAOCS and related enzymes, specifically the interconversion between open and closed forms, and the possibility of misfolding (and enhanced susceptibility to oxidative damage) in the former or intermediate forms, may well be a major cause of the notorious instability of this family of enzymes. It also explains the discrepancy between the *in vitro* and *in vivo* inactivation rates, since refolding mechanisms will exist in the latter case.

In the IPNS-Mn structure Gln330, the penultimate residue on the C terminus, acts as a fourth protein-based iron ligand (Roach *et al.*, 1995), which is displaced upon ACV (6) binding (Roach *et al.*, 1997). In contrast, the DAOCS structure reveals only three protein-derived iron ligands and the EXAFS data do not provide evidence for a fourth protein ligand. Thus, the DAOCS studies support mutagenesis studies with IPNS, concluding that iron ligation by a C-terminal residue is not required for catalysis (Sami *et al.*, 1997; Borovok *et al.*, 1996). The structures suggest that the DAOCS C terminus may assist in ordering binding of substrates to the active site. Thus, whilst 2-oxoglutarate can co-ordinate to the iron in the closed form, in order for penicillin N (1) to bind to the active site of DAOCS, the C terminus must be displaced to give a more open form of the enzyme. Thus, although DAOCS crystallizes as a trimer and can exist in this form in solution, it is the monomer which is the catalytically active form. A possible hinge allowing movement of the C terminus with respect to the active site is between residues 298 and 302 (IGGNY), a well conserved sequence in DAOCS (*S. clavuligerus*, *Norcardia lactamdurans*),

DACS (*S. clavuligerus*, *Norcardia lactamdurans*) and DAOCS/DACS (*C. acremonium*) (Cooper, 1993), suggesting that the C terminus is also important in the DACS and DAOCS/DACS catalytic cycles (see below).

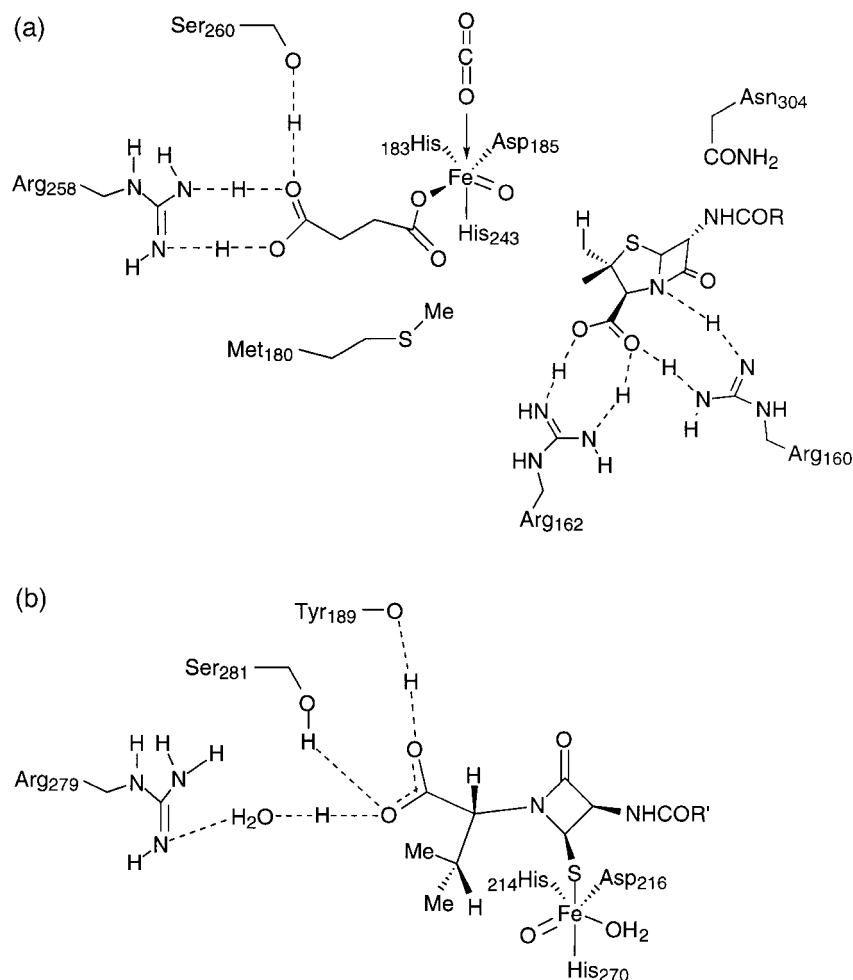
Examination of the interactions between the C-terminal residues (Asn304' to Ala311') of one DAOCS molecule and the active site of another (Figure 8) suggests a binding mode for the penicillin nucleus (Figure 9(a)). The penam C-3 carboxylate group probably occupies an analogous position to that of Ala311' from a symmetry-related molecule in the active site (Figure 8), forming electrostatic interactions with Arg162 and Arg160 (Figure 9 and Table 2). The side-chain of Arg160 may also be in position to form a hydrogen bonding interaction with the β-lactam nitrogen/carbonyl. Asn304, close to the C terminus of the enzyme, may form a hydrogen bond with the amide bond linking the side-chain to the penicillin nucleus.

Although anticipated from sequence analyses, the observation that the side-chains of the highly conserved Arg258 and Ser260 residues bind to the 5-carboxylate of 2-oxoglutarate is of interest, since the analogous residues in IPNS (together with Tyr189 and a water molecule) bind the ACV (6) valine carboxylate. Within the family of known iron-dependent oxidizing enzymes clearly related by sequence to DAOCS only two, IPNS and ACCO, do not use a 2-oxoglutarate cosubstrate with ACCO apparently using ascorbate in the place of 2-oxoglutarate. Thus, it seems most probable that IPNS evolved from a 2-oxoglutarate-dependent oxygenase rather than *vice versa*.

The presence of Arg160 and Arg162 in the DAOCS active site is a significant difference compared to that of IPNS (Figure 9(b)). In the structure of IPNS complexed to iron(II) and ACV (6), apart from the residues complexing to the iron and those forming electrostatic interactions with the valine carboxylate, the active site is largely devoid of polar residues. In the proposed mechanism for IPNS (Roach *et al.*, 1997) the iron-bound oxidizing species are isolated from the environment in part by the presence of the ACV (6) substrate, which is wrapped around them. The presence of the extra electrostatic interactions in the DAOCS active site relative to that of IPNS reflect its requirement to bind and position an extra substrate with a carboxylate group, penicillin N (1).

### Mechanistic implications

Kinetic studies on other 2-oxoglutarate-dependent oxygenases (Holme, 1975) have demonstrated ordered sequential binding of 2-oxoglutarate and the prime substrate followed by that of dioxygen, as shown for DAOCS in Figure 10. Binding of penicillin N (1) (and 2-oxoglutarate) probably occurs to an open form of the enzyme. Binding of penicillin N (1) may also assist dioxygen binding *via* displacement of the water molecule ligated to



**Figure 9.** (a) Proposed binding site for the penicillin N (**1**) in the active site of DAOCS, showing the proximity of the reactive ferryl species and the  $\beta$ -methyl group of the substrate. The side-chain of Arg160 may also bind to the  $\beta$ -lactam oxygen. R=D- $\alpha$ -aminoadipoyl. Note the presence of the Met180 side-chain, which is prevented from approaching the iron by the presence of the penicillin. (b) Possible structure for the ferryl/monocyclic  $\beta$ -lactam intermediate in IPNS catalysis as proposed by Roach *et al.* (1997, 1995) on the basis of the structures of the IPNS-Fe(II)-ACV and IPNS-Fe(II)-ACV-NO complexes. ACV=L- $\delta$ - $\alpha$ -aminoadipoyl)-L-cysteinyl-D-valine (**6**). R=L- $\alpha$ -aminoadipoyl. Note in the intermediate the C $^{\alpha}$ -C $^{\beta}$  bond of the isopropyl group of the valine is rotated relative to the tripeptide structures where the C $^{\beta}$ -H bond, which must be cleaved, projects away from the iron centre.

the iron *trans* to His183. Formation of a ferryl complex may then occur as proposed previously (Valegård *et al.*, 1998) *via* superoxide or peroxide intermediates in an isolated environment, i.e. a closed form of the enzyme. Carbon dioxide may remain co-ordinated to the ferryl iron, possibly as a bicarbonate ion after reaction with water. Together with the carboxylate derived from the succinate continued ligation of the carbon dioxide/bicarbonate may serve to modulate or direct the reactivity of the ferryl species.

Isotopic labelling experiments have demonstrated that cleavage of a C-H bond of the penicillin *pro-S* methyl group occurs before cleavage of the C-3 C-H bond (Baldwin *et al.*, 1991): Binding of penicillin N (**1**) in the proposed manner projects this methyl group directly towards the iron centre whilst the C-3 C-H bond is further away. The isolated nature of the active site in the closed form, the distance of the thiazolidine sulphur from the iron and the requirement for (at least) one of the iron ligands to be decomplexed renders mechanisms involving complexing of the penam sulphur to the iron unlikely. We presently favour a mechanism in which a ferryl intermediate, generated as previously proposed (Valegård *et al.*, 1998), reacts with the *pro-S* methyl group to generate a primary

radical (Pang *et al.*, 1984; Townsend *et al.*, 1985). Biomimetic studies (Baldwin *et al.*, 1988) suggest that this species enters an equilibrating manifold of radicals in which a tertiary cepham radical predominates. This may react with the Fe-OH species, either to give DAOC (**2**) and water (Figure 10, path A) or to form the 3 $\beta$ -hydroxymethylcepham (**12**) (Figure 10, path B; Baldwin *et al.*, 1991).

Alternatively, single electron transfer from the intermediate radical(s) may occur (Figure 10, path C) to give a cation, which must lose a proton to give DAOC (**2**) (Figure 10, path D) or which reacts with water/hydroxide to give (**12**) (Figure 10, path E). Since no protein derived functionality, including the side-chains of both Arg162 and Arg160 which are on the "wrong" face of the intermediates, is in position to act as a general base to deprotonate the C-2 C-H, it seems likely that (if a cationic mechanism is in operation) one of the three substrate-derived iron ligands acts in this manner. The binding model implies the position *trans* to His183 as the likely location for an iron ligated base. Further, use of a succinate oxygen as a general base would seem unlikely, since it is locked into a highly electron-deficient iron centre. It is possible that the CO<sub>2</sub> derived from 2-oxoglutarate reacts with water to form bicarbonate, which



acts as a base. In our mechanism this would place the bicarbonate *trans* to His243, possibly rendering it too far away to cleave the C-3 C-H bond directly. Thus, the bicarbonate may serve to increase the basicity of the hydroxyl *trans* to His183, which may act as a base, deprotonating the cationic intermediate to give (2) or quench it to give (12). Migration of the CO<sub>2</sub> to the hydroxyl group *trans* to His183 is another, less likely, possibility.

Kinetic studies imply that carbon dioxide is the first released product in 2-oxoglutarate-dependent dioxygenase catalysis (Holme, 1975) and it may be that return of the metal to the iron (II) oxidation state facilitates release of CO<sub>2</sub> (or bicarbonate) from the metal, which in turn triggers conformational changes, including movement of the C-terminal region, to give a more open form of the enzyme, allowing release of DAOCS and succinate.

DAOCS like other 2-oxoglutarate-dependent oxygenases can catalyse the "uncoupled" turnover of 2-oxoglutarate to succinate and carbon dioxide, albeit at a much reduced rate, in the absence of the prime substrate (Myllyharju & Kivirikko, 1997). The uncoupled turnover process has been studied in detail in the case of prolyl-4-hydroxylase, where in the absence of collagen substrate the turnover of 2-oxoglutarate is apparently stoichiometric with that of ascorbate in the presence of excess of the latter (Myllyharju & Kivirikko, 1997). Some substrate analogues can stimulate the uncoupled turnover process without themselves being oxidized and certain mutations can significantly alter the ratio of coupled:uncoupled turnover (Myllyharju & Kivirikko, 1997). The uncoupled turnover process may represent an editing process contributing to the attainment of both substrate and product (with respect to the type of oxidation reaction catalysed) selectivity by the 2-oxoglutarate-dependent enzymes. Conformational flexibility between (partially) open and closed enzyme forms may hinder binding and reaction of dioxygen to an enzyme-Fe(II)-2-oxoglutarate complex to which either the wrong "prime" substrate is bound or to one in which the correct prime substrate is incorrectly bound. Furthermore, such flexibility may also allow release of an undesirable or incorrectly bound prime substrate after formation of the ferryl species, which may be reduced back to the iron (II) oxidation state by a relatively non-specific reductant, e.g. ascorbate in the case of prolyl-4-hydroxylase<sup>†</sup>. The need for such an editing process may be particularly important in the case of prolyl-4-

hydroxylase, where the natural substrate is a polymer, collagen, with a considerable degree of secondary structure, which may make precise arrangement of the proline residues within the active sight of prolyl-4-hydroxylase awkward.

The mechanistic complexity of the 2-oxoglutarate-dependent dioxygenases ensures that many questions remain unanswered, but the present work, together with that of the recently reported crystal structures (Valegård *et al.*, 1998), places previous mechanistic studies within a structural context and provides new detailed proposals. It is important that future studies are now directed towards the structural characterization of intermediates in the catalytic cycle.

## Materials and Methods

### Materials

All chemicals were supplied by the Sigma-Aldrich Chemical Co., unless otherwise stated, and were of analytical grade or higher. Penicillin N (1) was synthesized by an extension of the reported method (Baldwin *et al.*, 1987b). Chromatography systems and columns were obtained from Amersham Pharmacia Biotech. Protein purification was performed on ÄKTA explorer<sup>®</sup>, FPLC<sup>®</sup> or BioPilot System. Q-Sepharose<sup>®</sup> HP (254 ml) was packed into a 60/100 column (I.D. = 6 cm × 9 cm). DEAE Sepharose<sup>®</sup> FF (30 ml) was packed into a XK16 column (I.D. = 1.6 cm × 15 cm) or 250 ml into a XK50 column (I.D. = 5.0 cm × 12.7 cm). SOURCE<sup>®</sup> 151SO (33 ml) was packed into a FineLINE<sup>®</sup> Pilot 35 column. Superdex<sup>®</sup> 75 was packed into an G32 × 1000 column (Amicon, I.D. = 3.2 cm × 89 cm, 716 ml and a XK26 column (I.D. = 2.6 cm × 60 cm, 320 ml). The following pre-packed columns were used: RESOURCE<sup>®</sup> ISO (1 ml); RESOURCE<sup>®</sup> Q (6 ml); Mono Q<sup>®</sup> HR 16/10; Superdex<sup>®</sup> 200 10/30; NAP-5. SDS-PAGE was performed using the Mini-Protean system (Bio-Rad). Gels were stained using Coomassie brilliant blue and Bismark brown (Lloyd, 1996), and were calibrated with the molecular mass marker kit 12,300-78,000 (BDH). <sup>1</sup>H NMR spectra were obtained in <sup>2</sup>H<sub>2</sub>O and referenced to TSP. The anaerobic glove box was supplied by Belle Technology and was supplied with an atmosphere of oxygen-free nitrogen.

### Construction of expression vectors

Expression vector pML1 was constructed according to standard protocols (Sambrook *et al.*, 1989). Sub-cloning of the fragment of DNA containing the DAOCS gene was performed by digestion of pNM88 (Morgan *et al.*, 1994) and pET 11a vectors using *Bam*HI and *Nde*I. The appropriate fragments were isolated by agarose gel electrophoresis, ligated, transformed into *Epicurian coli* XL1 Blue (Stratagene) and plated onto LB-agar containing 100 mg/ml ampicillin. Individual colonies were screened by agarose gel electrophoresis following initial digestion with *Bam*HI/*Nde*I. The required clones were repurified using the Wizard Mini-prep method (Promega), transformed by heat-shock into competent *E. coli* BL21 (DE3) and plated as before. Single colonies were screened for expression in 2TY containing 50 µg/ml ampicillin at 27, 30, and 37°C. The DNA encoding the DAOCS gene was sub-cloned from pML1 into the pET24a vector using an analogous procedure with kanamycin sulphate

<sup>†</sup> The release of substrate after formation of the ferryl species offers a plausible explanation for kinetic studies measuring  $V_{\max}/K_m$  isotope effects using deuterium labelled penicillin Ns (Baldwin *et al.*, 1987a). These were interpreted previously as indicating a mechanism involving reversible binding of penicillin N (1) to a ferryl intermediate formed by reaction of Fe(II) and 2-oxoglutarate with dioxygen. Note also that the possibility of more than one kinetic mechanism cannot be excluded.

(50 µg/ml) as the selection antibiotic. This vector was designated pHL1.

### Fermentation of *E. coli* BL21 (DE3)/pML1

*E. coli* BL21 (DE3)/pML1 was inoculated from a glycerol freeze into 4 × 100 ml of 2TY supplemented with 50 µg/ml ampicillin and grown overnight at 33 °C. The starter culture was used to inoculate 30 l of the same medium supplemented with 5 ml of polypropylene glycol 2025 in a New Brunswick Scientific MPP40 fermenter. The culture was grown at 30 °C, 300 r.p.m. and 30 l/min air for four hours ( $A_{600\text{ nm}} = 2.96$ ). IPTG (0.4 mM, Melford) was added and the culture was grown for a further three hours under the same conditions. Cells were concentrated to ca three litres using a tangential flow concentrator (Satorius) and centrifuged (JA-10 rotor, 10 000 r.p.m., 17,500 g, 15 minutes) and stored at -80 °C. Cells containing pHL1 were grown under identical conditions using kanamycin sulphate (50 µg/ml) in place of ampicillin.

Production of the selenomethionine derivative required re-optimization of the fermentation and purification conditions. The methionine auxotrophs B834 (DE3) and B834 (LyS) (DE3) (Pappa *et al.*, 1996) containing the pML1 plasmid gave soluble expression of DAOCS in 2TY medium, but soluble expression was not observed in LeMaster medium (by SDS-PAGE) and consequently *E. coli* BL21 (DE3) was used as the host cell. Small-scale incorporation trials showed that lower incorporation occurred for cultures induced at high  $A_{600}$ , increasing post-induction growth times and decreasing selenomethionine concentrations (Figure 3). In the optimized procedure a single colony was inoculated into culture medium consisting of LeMaster media (Hendrickson *et al.*, 1990) (5 ml), non-autoclavable LeMaster medium (0.5 ml), ampicillin (50 µg/ml), 1 × Kao and Michayluk vitamin supplement (growth medium) supplemented with 2TY (0.5 ml) and DL-methionine (40 µg/ml) and grown at 37 °C and 250 r.p.m. This culture was used to inoculate growth medium (2 × 100 ml) supplemented with DL-methionine and grown as before. This culture was used to inoculate a ten litre fermenter containing growth medium and 50 µg/ml DL-selenomethionine and was grown at 37 °C to  $A_{600} = 0.8$ , induced and allowed to grow for a further two hours before harvesting.

### Purification of DAOCS

All manipulations were performed on ice or at 4 °C. All fractions were analysed using 12.5% (w/v) SDS-PAGE, following initial TCA precipitation for those samples containing ammonium sulphate. Cells expressing wild-type DAOCS (*E. coli* BL21 (DE3)/pML1, 70.15 g) were thawed and resuspended in 280 ml of 50 mM Tris-HCl, 1 mM EDTA (pH 7.5), 2 mM DTT, 1 mM benzamidine-HCl, 1 mM PMSF, 1 µM leupeptin (lysis buffer). Lysozyme (155 mg) was added and the solution was stirred for 15 minutes. Complete cell disruption was achieved by sonication using 4 × 20 second burst on power 5 with 60 seconds cooling between each burst (W-380 sonicator, Ultrasonics Inc.). DNA was removed by precipitation with 1% (w/v) streptomycin sulphate and 0.1% (w/v) PEI. After 15 minutes stirring the extract was centrifuged (Beckman, JA-14 rotor, 12,000 r.p.m., 22,000 g, ten minutes), loaded onto a Q-Sepharose<sup>®</sup> HP column which had been equilibrated in

50 mM Tris-HCl, 1 mM EDTA, (pH 7.5), 2 mM DTT, 0.2 mM benzamidine-HCl, 0.2 mM PMSF (column buffer) and washed with 1000 ml of the same buffer at 25% ml/min. Protein was eluted with a 25-75% 0.3 M NaCl gradient in the same buffer over 600 ml. All fractions (25 ml) were analysed by 12.5% SDS-PAGE. Fractions 13-22 (150-220 mM NaCl) were pooled and 260 ml of 3.2 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> was added slowly with stirring. This pool was loaded onto a SOURCE<sup>®</sup> 151SO column which had been equilibrated with 1.6 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> in column buffer at 40 ml/min. The column was washed with 100 ml of the same buffer and eluted with a 90-50% gradient using column buffer supplemented with 20% (v/v) glycerol over 300 ml with 20 ml fractions collected. The inclusion of glycerol in this buffer is essential for efficient elution of purified protein. Fractions 3-7 (1.3-1.1 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>) were pooled, concentrated in a 50 ml Amicon stirred cell to ca 13 ml and loaded as 2 × 6.5 ml onto a Superdex<sup>®</sup> 75 column (716 ml) which had been equilibrated in 100 mM Tris-HCl, 1 mM EDTA (pH 7.5), 2 mM DTT, 0.2 mM benzamidine-HCl, 0.2 mM PMSF (gel filtration buffer). The column was eluted with the same buffer at 2 ml/min and fractions (5 ml) were collected between 280 and 440 ml. Fractions 18-32 (370-440 ml and 49-62 (360-430 ml) were pooled, concentrated to ca 9 ml and exchanged into 10 mM Hepes-NaOH (pH 7.0) using Econo-pak columns (Bio-Rad). Protein was concentrated to 50 mg/ml and frozen on dry ice before storage at -80 °C. It was not possible to measure accurately DAOCS activity in crude *E. coli* extracts, due to interference with the holed-plate assay (Baldwin *et al.*, 1987a,b) by the antibiotic activity of streptomycin sulphate. Assay of DAOCS activity in crude extracts, by HPLC was also difficult because of the presence of other materials which absorb at  $\lambda = 254$  nm.

The molecular mass of native apo-DAOCS was determined by exchanging into gel filtration buffer and loading 0.38 mg (in 100 µl) onto a Superdex<sup>®</sup> 200 10/30 column which had been equilibrated in the same buffer at 0.5 ml/min. The column was calibrated using the following proteins (0.125 mg unless otherwise stated):  $\beta$ -amylase (200 kDa); ADH (150 kDa); BSA (66 kDa); ovalbumin (42.7 kDa, 0.31 mg); carbonic anhydrase (29 kDa). An identical analysis was also performed using 5 mM monopotassium 2-oxoglutarate in the buffer and sample.

The expression level for the selenomethionyl-DAOCS was ca threefold less than for the wild-type, possibly due to the toxic effects of the selenomethionine and/or the relatively short induction time, and its purification was performed in the presence of 5 mM DTT to avoid oxidative damage (Hendrickson *et al.*, 1990). Two different ion-exchange column chromatographies (DEAE Sepharose<sup>®</sup> FF and RESOURCE<sup>®</sup> Q) at different pH values were required in order to improve the final level of purification. Thus, selenomethionyl-DAOCS was purified from 22 g cells using an identical extraction procedure with 5 mM DTT. The extract was loaded onto a DEAE Sepharose<sup>®</sup> FF column (30 ml) in column buffer at pH 8.5 and eluted with a 30-70% 0.4 M NaCl gradient over 135 ml with 10 ml fractions. Further purification of fractions 3-5 was achieved using five runs with a RESOURCE<sup>®</sup> ISO column (1 ml), eluting with a 90-35% 1.6 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> gradient over 12 ml at pH 7.5 with 1 ml fractions. Fractions 2-6 were pooled and concentrated to 6 ml, loaded as 2 × 3 ml onto a Superdex<sup>®</sup> 75 column (320 ml) and purified as before with 3 ml fractions. Fractions 20-28 and 70-78 were pooled, diluted with one volume of Milli-Q water and loaded onto a RESOURCE<sup>®</sup> Q column (6 ml) in column buffer at

pH 7.5. The column was eluted with a 25-75% 0.3 M NaCl gradient over 27 ml with 3 ml fractions. Fractions 3-5 were pooled, buffer exchanged and concentrated to 26.4 mg/ml before storage at  $-80^{\circ}\text{C}$ . The derivative was ca 95% (i.e. less than the wild-type) pure by SDS-PAGE analysis and had a similar specific activity to wild-type DAOCS using penicillin N (**1**) as substrate in the holed plate assay (232 nmol/min per mg compared to 208 nmol/min per mg for wild-type DAOCS control).

### Synthesis of G-7-ADCA (**11**)

(**11**) was prepared from 7-ADCA (5.1 mmol, 1.02 g) and phenylacetyl chloride (7.56 mmol, 1 ml) (Chauvette *et al.*, 1971) using THF as the organic solvent and NaOH as the base. The product was solvent extracted into ethyl acetate to give a pale yellow solid (475 mg, 26%). HPLC purification ( $\text{C}_4$  Hypersil (250  $\times$  4.6 mm), 25 mM  $\text{NH}_4\text{HCO}_3$  in 5% (v/v) methanol at 2 ml/min,  $\lambda = 254$  nm) gave (**11**) (>95% pure by  $^1\text{H}$  NMR analysis) with a retention volume of 34-42.8 ml:  $^1\text{H}$  NMR (500 MHz)  $\delta_{\text{H}}$  1.9 (3 H, s,  $\text{CH}_3$ ); 3.25 (1 H, d,  $J = 7.5$  Hz, H-4); 3.35 (3 H, s, residual MeOH); 3.55 (1 H, d,  $J = 7.5$  Hz, H-4); 3.70 (2 H, ABq,  $J = 16$  Hz,  $\text{PhCH}_2$ -); 5.05 (1 H, d,  $J = 5.5$  Hz, H-6); 5.60 (1 H, d,  $J = 5.5$  Hz, H-7);  $m/z$  (negative ESI MS) = 331 (M-H).

### Incubation of penicillin G (**10**)

DAOCS (6.14 mg, 119 units/mg, 730 units) and penicillin G (**10**, 7.56 mg, 20  $\mu\text{mol}$ ) were incubated in a final volume of 2 ml (Shibata *et al.*, 1996) for two hours. The reaction was quenched with 5 ml of acetone, which was removed *in vacuo* before freeze-drying.  $^1\text{H}$  NMR (500 MHz) analyses implied a ca 20-30% conversion of penicillin G (**10**) to G-7-ADCA (**11**), as judged by integration of the resonance at 5.60 p.p.m. against TSP. Purification of the crude incubation mixture by HPLC (as above with 10% (v/v) methanol in the eluent) gave G-7-ADCA (**11**) (45  $\mu\text{g}$ ) with a retention volume of 23.2-26.4 ml.  $^1\text{H}$  NMR and ESI MS analyses were the same as those reported above for synthetic (**11**).

### Kinetic assays

Protein (10  $\mu\text{l}$ ) was mixed with 10  $\mu\text{l}$  of  $10 \times$  stock solution of cofactors (500 mM  $(\text{NH}_4)_2\text{SO}_4$ , 10 mM 2-oxoglutarate, 20 mM DTT, 10 nM ascorbate, 20 mM  $\text{FeSO}_4$ , NaOH to pH 7.5 in 50 mM Tris-HCl, p 7.5) and incubated at ambient temperature for two minutes. The mixture was diluted with 70  $\mu\text{l}$  of 50 mM Tris-HCl (pH 7.5) and the reaction initiated by addition of 1.0 mM penicillin N (**1**, 10  $\mu\text{l}$ ). After 5 minutes methanol (100  $\mu\text{l}$ ) was added and 50 or 100  $\mu\text{l}$  was used to measure the DAOC present using the holed-plate method with *E. coli* ESS as the indicator organism (Baldwin *et al.*, 1987a,b). Plates were calibrated using DAOC (0.5-1.0 nmol). Alternatively, DAOC (**2**) was quantified using anion-exchange HPLC (Morgan *et al.*, 1994; Shibata *et al.*, 1996). Protein concentrations were determined by the method of Bradford (1976) using BSA as a calibration standard.

Kinetic parameters for penicillin G (**10**) were determined with 1, 2, 3, 4, 5, 6, 8 and 10 mM substrate using 18.8  $\mu\text{g}$  DAOCS/assay in a total volume of 200  $\mu\text{l}$ . After quenching the reaction with 200  $\mu\text{l}$  methanol, 200  $\mu\text{l}$  of water was added. Samples were centrifuged and analysed by HPLC (150  $\mu\text{l}$  injections, as above using 15% (v/v) methanol in the eluent). Product was quantified by

peak height at 0.05 AUFS using synthetic G-7-ADCA (**11**, 0.5-3.5 nmol) as standard. Kinetic parameters ( $\pm\text{SE}$ ) were determined using Leonora by directly fitting of the data to the appropriate kinetic expression following initial graphical analysis.

Preliminary experiments with metal ions used the HPLC assay with penicillin N (**1**) as substrate. Enzyme was preincubated for two minutes with cofactors with the appropriate metal ion at 2 mM final concentration replacing Fe(II). Analysis of metal ions as potential inhibitors was performed by the same method using the standard cofactor mixture with Fe(II).

UV-visible spectra were recorded using a Phillips Pye-Unicam 8800 UV-visible spectrophotometer in 10 mM  $\text{Na}_2\text{HPO}_4\text{-HCl}$  (pH 7.0).

### DTNB titration of DAOCS

DAOCS (ca 1 mg) in 200  $\mu\text{l}$  of 50 mM Tris-HCl (pH 7.5) was incubated on ice with 2.5 mM DTT for two hours 15 minutes. The sample was diluted to 500  $\mu\text{l}$  and exchanged into 50 mM Tris-HCl (pH 8.0) with a NAP-5 column. The sample was diluted to 4 ml and 0.7 ml was assayed at  $A_{412}$  (Habeb, 1974) in the presence or absence of 2% SDS (w/v) with 1 mM DTNB.

### Dynamic laser light scattering

Dynamic laser light scattering measurements were performed in a DynaPro-801 TC instrument at  $8-37^{\circ}\text{C}$  using DAOCS (1 mg/ml) in 10 mM Hepes-NaOH (pH 7.0) Samples containing iron(II) were prepared and measured under anaerobic conditions.

### Preparation and analysis of EXAFS samples

A crude extract of *E. coli* BL21 (DE3)/pML1 was prepared and purified by anion-exchange chromatography on DEAE Sepharose<sup>®</sup> FF and gel filtration chromatography as described above. Final purification was achieved with Mono Q<sup>®</sup> 16/10 column chromatography using a 0-0.4 M NaCl gradient over 13 ml. Purified DAOCS were exchanged into 10 mM Tris-HCl, 0.3 mM  $(\text{NaN}_3)$  (pH 7.5) and concentrated to 74 mg/ml (ca 2.15 mM). Iron(II) sulphate (100 mM, 5  $\mu\text{l}$ ) or  $\text{FeSO}_4$  and potassium 2-oxoglutarate (200 mM, 2.5  $\mu\text{l}$ ) were added to 250  $\mu\text{l}$  of enzyme under anaerobic conditions and frozen in liquid nitrogen in the EXAFS cell.

The EXAFS data (Randall *et al.*, 1993) were recorded at ca 80 K on station 8.1 of the CCLRC Daresbury SRS, operated at 2 GeV with an average current of 150 mA, using a liquid nitrogen-cooled cold finger. Twelve scans were recorded and averaged for each sample. The Fe K-edge spectra were calibrated relative to the first peak in the derivative for an iron foil Spectrum at 7112 eV. Spectra were analysed, including multiple scattering, in EXCURV92 (Binsted *et al.*, 1991) using exact wave theory (Lee & Pendry, 1975; Gurman *et al.*, 1984, 1986). Phase shifts were calculated *ab initio* using Hedin-Lundqvist potentials (Hedin & Lundqvist, 1969). Theoretical fits were generated by adding shells of scatterers around the central iron atom and iterating the bond lengths and Debye-Waller type parameters to achieve the optimum fit to the experimental data. To reduce the number of parameters used to interpret the EXAFS, the occupation number of each shell of backscattering atoms was fixed at an integer value and a restrained refinement approach (Binsted *et al.*, 1992) used to treat the backscattering of



the imidazole rings of the histidine ligands. Thus, these groups were each treated as single units constrained at the known geometry, including multiple scattering pathways and keeping Debye-Waller parameters the same for atoms at similar distances from the metal (Binsted *et al.*, 1992). Backscattering from the 2-oxoglutarate group was similarly treated by restrained refinement using the geometry previously determined for sodium 2-oxoglutarate (Lis & Matuszewski, 1984).

### Crystallization of DAOCS

Crystals of DAOCS were grown by vapour-diffusion using the hanging drop method at ca 20 °C. The precipitating solution consisted of 100 mM Hepes-NaOH (pH 7.0), 5 mM potassium 2-oxoglutarate and 1.6-1.7 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>. Drops (3 µl) consisted of 55 mM Hepes-NaOH (pH 7.0), 2.5 mM potassium 2-oxoglutarate and 0.8-0.85 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>. Selenomethionyl-DAOCS could only be crystallized under these conditions in the anaerobic glove box with dioxygen levels maintained at 0.1-0.3 p.p.m. throughout. These crystals were used to determine the structure of DAOCS (Valegård *et al.*, 1998) using isomorphous replacement and a detwinning strategy (A.C. Terwisscha van Scheltinga *et al.*, unpublished results). An independent structure of the DAOCS, 2-oxoglutarate complex was also determined to 2.5 Å by isomorphous replacement (K. Harlos & M. D. Lloyd, unpublished results). Both structures were used during the analysis of the DAOCS reaction mechanism.

### Accession numbers

Co-ordinates and structure factors have been deposited with the Protein Data Bank (entries 1dcs and r1dcssf for apoenzyme, 1rxf and r1rxsf for the iron complex, and 1rxg and r1rxgsf for the iron, 2-oxoglutarate complex).

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