Dihydroxy-Acid Dehydratases From Pathogenic Bacteria: Emerging Drug Targets to Combat Antibiotic Resistance

Tenuun Bayaraa,[a] Jose Gaete,[a] Samuel Sutiono,[b] Julia Kurz,[a] Thierry Lonhienne,[a] Jeffrey R. Harmer,[c] Paul V. Bernhardt,[a] Volker Sieber,[a, b] Luke Guddat,[a] and Gerhard Schenk*[a, d, e]

Abstract: There is an urgent global need for the development of novel therapeutics to combat the rise of various antibiotic-resistant superbugs. Enzymes of the branched-chain amino acid (BCAA) biosynthesis pathway are an attractive target for novel anti-microbial drug development. Dihydroxy-acid dehydratase (DHAD) is the third enzyme in the BCAA biosynthesis pathway. It relies on an Fe–S cluster for catalytic activity and has recently also gained attention as a catalyst in cell-free enzyme cascades. Two types of Fe–S clusters have been identified in DHADs, i.e. [2Fe–2S] and [4Fe–4S], with the latter being more prone to degradation in the presence of oxygen. Here, we characterise two DHADs from bacterial human pathogens, Staphylococcus aureus and Campylobacter jejuni (SaDHAD and CjDHAD). Purified SaDHAD and CjDHAD are virtually inactive, but activity could be reversibly reconstituted in vitro (up to ~19,000-fold increase with kcat as high as ~6.7 s⁻¹). Inductively-coupled plasma-optical emission spectroscopy (ICP-OES) measurements are consistent with the presence of [4Fe–4S] clusters in both enzymes. N-isopropylxoy-aryl hydroxamate (IpOHA) and aspartic acid are both potent inhibitors for both SaDHAD (Kᵢ = 7.8 and 51.6 μM, respectively) and CjDHAD (Kᵢ = 32.9 and 35.1 μM, respectively). These compounds thus present suitable starting points for the development of novel anti-microbial chemotherapeutics.

Introduction

The rapid increase in antibiotic resistance among human pathogens has become a major global threat. It is well established that new lines of drugs are urgently needed to combat resistance. The branched chain amino acids (BCAAs), valine, leucine and isoleucine are synthesized de novo through the BCAA pathway (Figure 1) in bacteria, plant and fungi.[1,2] The BCAAs are crucial building blocks of nearly all proteins and are thus necessary for the survival of all organisms. However, this pathway is not present in animals, including humans, which thus necessary for the survival of all organisms. However, this pathway is not present in animals, including humans, which obtain BCAAs directly from their diet.[1,3] Therefore, the enzymes of the BCAA biosynthesis pathway are excellent targets for antimicrobial drug discovery and herbicide development since their absence in the animal kingdom is anticipated to minimize side effects caused by their inhibition by biocides.[1,2] The first three enzymes of this pathway are acetohydroxyacid synthase (AHAS)[5] ketol-acid reductoisomerase (KARI)[6–9] and dihydroxy- acid dehydratase (DHAD).[10–13] AHAS has been used effectively as a target of more than 50 commercial herbicides with some of the inhibitors also showing antimicrobial activity.[1,4] Importantly, sulfometuron methyl, a commercial AHAS herbicide, has been shown to have anti-tuberculosis (TB) activity.[14] Other novel AHAS inhibitors have also been shown to have anti-TB activity within the range 0.38 to 200 μM in vitro assays.[15] More recently, KARI has also emerged as a promising drug target,[15,16,17] in both Mycobacterium tuberculosis (Mt) and uropathogenic Escherichia coli (Ec), KARI has been shown to be essential for growth and survival.[16,17] Consequently, diverse compounds that significantly inhibit KARI have recently been developed as potential anti-microbial drug candidates.[18–22]
DHAD, the third enzyme in the BCAA biosynthesis pathway, has received less attention than AHAS and KARI. It belongs to the ilvD/EDD superfamily that also includes sugar acid dehydratases (DHTs).\textsuperscript{12,13,23,24} One of the main characteristics of this family is the presence of an Fe$\textsubscript{S}$ cluster in the active site, that is essential for the catalytic function of these enzymes.\textsuperscript{23} Two types of Fe$\textsubscript{S}$ clusters have been observed, i.e. $[2\text{Fe}_{2}\text{S}]$ and $[4\text{Fe}_{4}\text{S}]$.\textsuperscript{10,11,25} The $[4\text{Fe}_{4}\text{S}]$ cluster was first observed in DHAD from \textit{E. coli},\textsuperscript{11} while the $[2\text{Fe}_{2}\text{S}]$ cluster was first reported for spinach DHAD.\textsuperscript{10} The $[4\text{Fe}_{4}\text{S}]$ cluster is sensitive to oxygen, with oxidation leading to enzyme inactivation.\textsuperscript{13} The $[2\text{Fe}_{2}\text{S}]$ cluster, on the other hand, is more stable in the presence of oxygen.\textsuperscript{10,23,24,26}

DHAD catalyses the dehydration of dihydroxy-isovalerate (DHIV) or dihydroxy-methylvalerate (DHMV) to keto-isovalerate (KIV) and keto-methylvalerate (KMV), respectively (Figure 2).\textsuperscript{25,27} It is proposed that the dehydration reaction is initiated by the abstraction of the proton at the C2 position of the substrate by the alkoxide side chain of a conserved serine residue in the active site. The resulting carbanion may be stabilized by the

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Mg$^{2+}$ ion in the vicinity of the Fe–S cluster. This Mg$^{2+}$ ion is also essential for the catalytic activity of DHADs.$^{[11,12,13,28]}$ The combined effect of this proton abstraction and the coordination of the alcohol moiety at position C3 of the substrate to one of the iron atoms of the Fe–S cluster leads to a weakening of the C3–OH bond and its subsequent cleavage (Figure 2). A tautomerization completes the catalytic reaction, leading to the keto product.

Recent studies have also highlighted DHAD to be an essential enzyme for the growth and survival of microbes and plants; its combined effect of this proton abstraction and the coordination of the alcohol moiety at position C3 of the substrate to one of the iron atoms of the Fe–S cluster leads to a weakening of the C3–OH bond and its subsequent cleavage (Figure 2). A tautomerization completes the catalytic reaction, leading to the keto product.

DHAD and DHT have been shown to play a central role in cell-free enzyme cascades that facilitate the conversion of glucose to high-value products such as isobutanol.$^{[12,13,30–32]}$ Recent studies have also highlighted DHAD to be an essential enzyme for the growth and survival of microbes and plants; its inhibition by the natural product aspterric acid ($K_i = 9 \text{nM}$) greatly impairs the growth of harmful cyanobacteria.$^{[26]}$ Aspterric acid is also a potent inhibitor of both AtDHAD ($K_i = 0.3 \mu \text{M}$) and MtDHAD ($K_i = 10.1 \mu \text{M}$) DHAD.$^{[28,29]}$ Furthermore, the deletion of the gene encoding DHAD in *Aspergillus fumigatus* was shown to strongly reduce the virulence of this fungal pathogen.$^{[33]}$

Here, the DHADs from the two bacterial pathogens, *Staphylococcus aureus* and *Campylobacter jejuni* (SaDHAD and CjDHAD), were recombinantly expressed, purified and their catalytic properties investigated and compared to those of other DHADs. *S. aureus* is responsible for one of the most common causes of bacterial infections around the world. It can cause numerous medical conditions that range from minor skin conditions to necrotizing fasciitis, sepsis and even death.$^{[36–38]}$ In particular, the recent emergence of methicillin-resistant *S. aureus* (MRSA) has caused significant problems in the health care sector.$^{[39]}$ *C. jejuni* is a food-borne pathogen that causes gastrointestinal infections and can lead to Guillan-Barre syndrome.$^{[37–39]}$ The haphazard use of antibiotics fed to farm animals has led to a rise of quinolone- and macrolide-resistant *C. jejuni*, diminishing the potency and number of available antibiotics.$^{[37,39]}$ Therefore, there is an urgent need to develop novel antimicrobials to combat the rise of antibiotic-resistant *S. aureus* and *C. jejuni* as well as other emerging superbugs.

**Results and Discussion**

**Expression and purification**

*E. coli* BL21 strains were transformed with pET28a vectors (Figure S1) containing the genes that encode DHADs from *S. aureus* or *C. jejuni* (SaDHAD or CjDHAD, respectively). The DNA and protein sequences of SaDHAD and CjDHAD are also included in the Supporting Information. Successful transformants were selected based on their resistance to kanamycin. The recombinant expression (Figure S2) of the target enzymes was performed in autoinduction media, and the proteins were purified as described in the experimental section (Figures S3 and S4), resulting in a yield of approximately 50 mg/L for both SaDHAD and CjDHAD. The calculated molecular weights of SaDHAD and CjDHAD are 60.021 kDa and 60.104 kDa, respectively, in good agreement with SDS PAGE analyses (Figures S3 and S4).

![Figure 3. Crystal structure of a DHAD monomer from *A. thaliana*.](image)

The active site is zoomed in on the right (red circle). The Fe–S cluster is shown in orange and yellow spheres. The Mg$^{2+}$ ion is shown as a green sphere. The Fe–S cluster-coordinating cysteine residues are shown as green sticks. The Mg$^{2+}$-coordinating residues are shown as magenta sticks. The catalytically essential serine residue is shown in cyan sticks.
and S4). Protein samples were concentrated to ~20 mg/mL and frozen at −80 °C in the presence of 10% glycerol.

**Characterisation of SaDHAD and CjDHAD as initially purified**

The reported catalytic rates ($k_{\text{cat}}$ values) for the reaction of DHADs with their natural substrate, DHIV (Figure 2), vary from values as low as 0.3 s$^{-1}$ for DHAD from *Sulfolobus solfataricus* (SsDHAD) to 70 s$^{-1}$ for the DHAD from *E. coli* (EcDHAD).\[11,30\] By comparison, the measured catalytic rates of SaDHAD and CjDHAD indicate that both enzymes are very slow, with $k_{\text{cat}}$ values of $7.6 \times 10^{-4}$ s$^{-1}$ and $3.5 \times 10^{-4}$ s$^{-1}$, respectively (Figure 4). It should, however, be pointed out that previous studies with SsDHAD demonstrated that the enzyme activity may be reversibly lost during the purification procedure.\[30\] Subsequent incubation of that enzyme with the reducing agent, 2-mercaptoethanol (2-ME), led to a ~threefold increase in the activity of that enzyme.\[30\] However, further activation attempts in the presence of added Fe$^{2+}$ led to a 78% loss of enzymatic activity, and it was proposed that storage and purification leads to a gradual disintegration/dissociation of the Fe–S cluster from the active site.\[30\] The difficulty in obtaining and maintaining a fully active DHAD was further illustrated by a recent study with a [2Fe–2S] cluster-containing DHAD from the cyanobacterium *Synechocystis sp. PCC 6803* (SnDHAD).\[26\] As purified, its catalytic rate for the reaction with DHIV is $k_{\text{cat}} \sim 0.47$ s$^{-1}$.\[26\] Metal analysis indicated that the Fe content per protein monomer is only 0.62 atoms. The enzyme was thus anaerobically incubated with sodium sulfide and ammonium ferrous sulfate, which led to an increase in the Fe content to 2.29 atoms per monomer, suggesting the presence of an active site with a fully loaded [2Fe–2S] cluster. However, despite this increase in Fe loading the activity of SnDHAD dropped to 30% of the value recorded for enzyme before its incubation.\[26\]

We suspected that the low activity of SaDHAD and CjDHAD may be due to the absence of functional Fe–S clusters after purification. We therefore determined their iron content using inductively-coupled plasma optical emission spectrometry (ICP-OES). SaDHAD and CjDHAD contain sub-stoichiometric amounts of iron, 0.30 and 0.22 atoms per protein monomer, respectively (Table 1). This suggests that functional Fe–S clusters are indeed absent in SaDHAD and CjDHAD when initially purified, in correlation with the minimal activity of these enzymes. However, there is a large difference between the enzymes in this study and SnDHAD (see preceding paragraph). While the iron content of these three enzymes as purified is comparable, their catalytic proficiency differs by several orders of magnitude. This observation indicates that other factors, for instance the nature of the Fe–S clusters (i.e. [2Fe–2S] vs. [4Fe–4S]) and their oxidation state(s), may contribute to both the activity and stability of these enzymes.

Figure 4. Michaelis-Menten plots of the reaction rates of SaDHAD (A) and CjDHAD (B). Catalytic parameters ($K_m$ and $k_{\text{cat}}$) were measured using the natural substrate, DHIV, for both the enzymes as initially purified and their activated counterparts. Note that the catalytic activities (shown on the y-axes) of the initially purified and activated enzymes differ by four orders of magnitude.
Enzyme activation

To activate SaDHAD and CjDHAD, we modified a protocol previously developed for the activation of SsDHAD[30] and incubated both enzymes for one hour (aerobically or anaerobically) with the reducing agents, sodium dithionite and 2-ME, in the presence of ammonium ferrous sulfate in 50 mM HEPES, pH 8, at 37 °C. The optimal concentrations for the reducing agents and Fe²⁺ were established by comparing the effect of three different mixtures on the catalytic activities of SaDHAD and CjDHAD (Figure S5). The mixture with 50 mM sodium dithionite, 200 mM 2-mercaptoethanol and 10 mM (NH₄)₂Fe-(SO₄)₂ resulted in the highest catalytic activity and was thus used for all subsequent activations of the enzymes. Both the aerobic and anaerobic activation procedures led to a significant increase in both the iron content and catalytic activity (Figure 4 and Table 1). These results suggest that SaDHAD and CjDHAD are fully activated with reconstituted Fe–S clusters, likely of the [4Fe–4S] form (discussed below).

Stability of activated SaDHAD and CjDHAD under aerobic conditions

In an aerobic environment at room temperature, activated SaDHAD and CjDHAD rapidly lost activity (Figure 5). Irrespective of how the enzymes were activated (aerobic or anaerobic), their activities dropped by at least 30% after one hour with only residual activity left after 24 h. The Fe content of each enzyme sample was re-measured at the end of the 24-hour incubation period. For the anaerobically activated enzymes, the metal content decreased modestly (SaDHAD: from 5.45 ± 0.31 to 5.03 ± 0.06 atoms per monomer; CjDHAD: from 5.94 ± 0.19 to 4.33 ± 0.12 atoms per monomer), however it decreased more significantly for the aerobically activated enzymes (SaDHAD: from 4.66 ± 0.57 to 1.14 ± 0.02 atoms per monomer; CjDHAD: from 4.71 ± 0.01 to 1.95 ± 0.06 atoms per monomer). It is thus likely that anaerobic and aerobic activations lead to Fe–S clusters with different oxidation states, with the more oxidized form (i.e. after aerobic activation) being more labile. However, it is interesting to point out that the oxidation states of these clusters do not appear to correlate to the activity of the two DHADs; the more oxygen-exposed cluster (aerobically activated) is more active in SaDHAD, whereas the less oxygen-exposed cluster (anaerobically activated) is more active in CjDHAD (Figure 5). The enhanced lability of the Fe–S clusters when activated aerobically, but the lack of a correlation between the mode of activation (aerobic vs. anaerobic) and activity in the two enzymes suggest that the oxidation state of the clusters affects their stability significantly but is of lesser relevance to their catalytic efficiency. This interpretation is also consistent with the rapid inactivation that was also observed for anaerobically purified EcDHAD (~ 50% loss of activity after 30 minutes at 25 °C in an aerobic environment), an enzyme with a confirmed oxygen-sensitive [4Fe–4S] cluster.[11,40]

Table 1. Fe content and catalytic rates (s⁻¹) of SaDHAD and CjDHAD as initially purified and after activation under aerobic or anaerobic conditions. Catalytic rates were determined with 100 nM SaDHAD and CjDHAD using 5 mM DHIV at 37 °C.

| Enzyme | Activation method | Fe atoms per subunit | Rate [kcat, s⁻¹] |
|--------|-------------------|----------------------|------------------|
| SaDHAD | None              | 0.30 ± 0.02          | 7.6 × 10⁻⁴       |
|        | Aerobic           | 4.66 ± 0.57          | 1.8 ± 0.1        |
|        | Anaerobic         | 5.45 ± 0.31          | 1.0 ± 0.0        |
|        | None              | 0.22 ± 0.07          | 3.4 × 10⁻⁵       |
|        | Aerobic           | 4.71 ± 0.01          | 3.8 ± 0.6        |
|        | Anaerobic         | 5.94 ± 0.19          | 6.7 ± 0.4        |
| CjDHAD | None              | 0.22 ± 0.07          | 3.4 × 10⁻⁵       |
|        | Aerobic           | 4.71 ± 0.01          | 3.8 ± 0.6        |
|        | Anaerobic         | 5.94 ± 0.19          | 6.7 ± 0.4        |

Figure 5. Stability of SaDHAD (A) and CjDHAD (B) after activation in an aerobic environment at 25 °C. Enzyme and substrate (DHIV) concentrations were 100 nM and 5 mM, respectively.
Characterisation of the catalytic properties of activated SaDHAD and CjDHAD

Both the aerobic and anaerobic activation procedures led to large increases in activity and Fe content for both enzymes (Figure 4 and Table 1). For SaDHAD the corresponding values are a ~15 and ~18-fold increase in iron loading, and ~2,400- and ~1,300-fold enhancement of the rate. For CjDHAD, the aerobic and anaerobic treatments led to a ~21 and ~27-fold increase in the iron loading, respectively, and a concomitant ~11,100- and ~19,700-fold enhancement of the catalytic rate. Thus, while there is a correlation between the iron loading and the activity, the two enzymes respond again differently to aerobic and anaerobic treatments (compare to previous paragraph); while the anaerobic incubation leads to higher loading in both enzymes, only in CjDHAD does this also lead to an increased rate (nearly two-fold). In SaDHAD, the rate after aerobic treatment reaches only ~60% of that after aerobic treatment despite a ~20% increase in iron loading. Since both enzymes were treated identically, the observed difference after anaerobic incubation supports the hypothesis that these enzymes may indeed prefer different oxidation state(s) for optimal activity.

In order to assess if the activation of the two enzymes affects their interaction with the substrate (DHIV), reaction rates were measured at various concentrations of substrate and analysed using Michaelis-Menten kinetics (Figure 4; data were only measured for the most active forms of the enzymes, i.e. the aerobically activated SaDHAD and the anaerobically activated CjDHAD). For both the enzymes as purified and activated, saturation-type behaviour for DHIV is observed. While the activation procedure greatly impacts on the $k_{cat}$ values, its effect on the $K_m$ values is minimal. For SaDHAD the $K_m$ value slightly increases from 0.93 mM to 1.1 mM upon activation whereas for CjDHAD virtually no difference is observed. Thus, substrate binding in both DHADs is largely unaffected by the activation procedure, indicating that minimal structural changes occurred during this process. Furthermore, the catalytic parameters of activated SaDHAD and CjDHAD are in good agreement with those of other known DHADs (Table 2).

**pH profile and thermostability**

Catalytic assays with the substrate DHIV were performed at different pH values ranging from 7 to 10 using three different buffers with activated SaDHAD and CjDHAD to determine the effect of pH on their activities (Figure 6). Both enzymes perform optimally at pH 9, slightly higher than the optimal pH of ~8 that has been observed in previously characterised DHADs and DHTs from organisms such as *S. solfataricus, M. tuberculosis*, *Methanococcus* spp. and *R. leguminosarum*.[28,41–43] The pH profiles of the two enzymes were fit to an equation derived for a diprotic system,[44–46] which provides estimates for the two relevant protonation equilibria ($pK_a^1$ and $pK_a^2$) that affect the catalytic rates of SaDHAD and CjDHAD. For both enzymes these $pK_a$ values are very similar (SaDHAD: 8.75 and 9.40; CjDHAD: 8.40 and 9.55). The assignment of these $pK_a$ values to respective residues in the active site is inherently difficult, and in the absence of structural and mutagenesis data it remains speculative. Nonetheless, the sharp increase in the activity as the pH is increased from 7.0 to 9.0 (corresponding to $pK_a^2$) is consistent with the deprotonation of Ser489, the residue proposed to initiate catalysis by abstracting a proton from the C2 position of the substrate (Figures 2 and 3). The sharp decline of the activity between pH 9.0 and 10.0 (corresponding to $pK_a^1$) may be due to the reduced ligand exchange rate of the hydroxide bound to the Fe-S cluster after the cleavage of the C3-OH (Figure 2). The $pK_a^1$ value of a water molecule bound to Fe$^{2+}$ is consistent with this assignment but it needs to be pointed out that no concrete evidence about the oxidation state(s) of the Fe–S clusters in SaDHAD and CjDHAD is currently available (see also below).

However, despite the uncertainties in the assignment of these $pK_a$ values the experimental data support the recently proposed

![Figure 6. The catalytic activities of SaDHAD (circles) and CjDHAD (squares) were determined at various pH values (at 37°C) using 50 mM HEPES (pH 7–7.5), Tris-HCl (pH 8–9) and Glycine buffers (pH 9.5–10). Enzyme and substrate concentrations were 100 nM SaDHAD, 50 nM CjDHAD and 5 mM DHIV. The data were fit to an equation derived for a diprotic system.](Image 314x649 to 539x775)

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**Table 2. Comparison of the kinetic parameters of several DHADs for their reaction with DHIV.**

| Organism (DHAD) | $K_m$ [mM] | $k_{cat}$ [s$^{-1}$] | $k_{cat}/K_m$ [M$^{-1}$·s$^{-1}$] | Nature of Fe–S cluster |
|-----------------|-------------|----------------------|-------------------------------|------------------------|
| *S. aureus* (SaDHAD) | 1.1         | 2.1                  | 1909                           | [Fe$^{3+}$]             |
| *C. jejuni* (CjDHAD) | 0.82        | 6.5                  | 7927                           | [Fe$^{3+}$]             |
| *E. coli* (EcDHAD)[28] | 1.5         | 70                   | 46,667                         | [Fe$^{2+}$]             |
| *M. tuberculosis* (MMDHAD)[29] | 2.0        | 1.87                 | 935                            | [Fe$^{2+}$]             |
| *A. thaliana* (AtDHAD)[28] | 5.7        | 1.2                  | 210                            | [Fe$^{2+}$]             |
| *S. oleracea* (SoDHAD)[30] | 1.5        | 25                   | 16,667                         | [Fe$^{2+}$]             |
| *Synechocystis* (SnDHAD)[30] | 0.14       | 0.47                 | 3,357                           | [Fe$^{2+}$]             |
| *S. solfataricus* (SsDHAD)[30] | 2.1        | 0.32                 | 152                            | [Fe$^{2+}$]             |
model for the reaction mechanism employed by DHADs (Figure 2).

The thermostability of SaDHAD and CjDHAD was also assessed with kinetic measurements after 15-minute incubation at temperatures ranging from 37°C to 60°C (Figure 7). After incubation, the enzymes were assayed at 37°C, the standard temperature used to assay DHAD activity. Activities sharply decrease for both enzymes, with virtually no activity measurable for samples incubated at 60°C. The ~30% drop in activity between the narrow temperature interval from 37°C to 40°C may indicate that the inactivation of CjDHAD and SaDHAD is likely to be due to the instability of the Fe–S cluster, not that of the protein itself, as also suggested for other DHADs in a previous study.\[12\]

Structure of the Fe–S cluster in SaDHAD and CjDHAD

The combined data presented above suggest that both SaDHAD and CjDHAD contain oxygen-sensitive [4Fe–4S] clusters in their active sites. Unfortunately, diverse attempts to crystallise these enzymes (as purified and after activation) failed, possibly a consequence of the instability of their Fe–S clusters (and the associated inhomogeneity of the protein samples). We thus performed a detailed sequence comparison between SaDHAD and CjDHAD and DHADs with well-established Fe–S clusters, i.e. EcDHAD with a [4Fe–4S] cluster and MtDHAD, AtDHAD and SoDHAD that have [2Fe–2S] clusters in their active sites (Figure 8). Two signature motifs from the lrv/EDD family are present in all DHADs. The first is located at the N-terminus and consists of 11 amino acids from sequence alignment position 121 to 131. Cysteine residue 121 is strictly conserved among all DHADs and provides a ligand for one of the iron ions in the Fe–S cluster (Figure 3; Cys121 in the alignment corresponds to Cys139 in the sequence of AtDHAD). This motif also contains two of the invariant ligands for the catalytically essential Mg\(^{2+}\) ion, i.e. Asp122 and the carboxylated Lys123 (Asp140 and Lys141 in AtDHAD; Figure 3).

The second signature motif in DHADs is located at the C-terminus and spans alignment residue positions 464 to 475 (Figure 8). Contained within this motif is an invariant serine residue at position 472 (Ser489 in AtDHAD; Figure 3). This residue has been proposed to act as the Lewis base during catalysis (Figure 2).\[10,25,48\] Other essential residues important for the function of DHADs are distributed across the protein sequence, including two additional cysteine ligands for the iron ions in the Fe–S clusters (Cys48 and Cys194, i.e. Cys66 and Cys211 in AtDHAD) and two ligands for the Mg\(^{2+}\) ion (Asp80 and Glu446, i.e. Asp98 and Glu463 in AtDHAD). However, it should be noted that the cysteine residue in alignment position 48 (Figure 8) is only conserved in DHADs with known [2Fe–2S] clusters, while in DHADs with a [4Fe–4S] cluster, notably EcDHAD,\[11\] a glycine residue resides in the corresponding location. Both SaDHAD and CjDHAD also have a glycine residue.
in this position (Figure 8) thus adding further evidence to the presence of a [4Fe–4S] cluster in these bacterial enzymes.

Since it has not yet been possible to visualise the three-dimensional structures of a [4Fe–4S] cluster in a DHAD, we employed electron paramagnetic resonance (EPR) and spectro-electrochemical measurements to gain further insight into the identity of the Fe–S clusters in SaDHAD and CjDHAD. Analysis of EPR spectra from Fe–S clusters frequently allow for distinction between two-iron, three-iron and four-ion clusters. Examples are illustrated in Figure 9A. Although the majority of Fe–S cluster containing enzymes play roles in electron transfer reactions, DHADs and DHTs are not the only examples of such enzymes that perform non-redox-type reactions. Aconitase, for instance, catalyses the stereo-specific isomerisation of citrate to isocitrate in the Krebs cycle. It can accommodate a [4Fe–4S] cluster, similar to EcDHAD and possibly SaDHAD and CjDHAD, but these clusters have proven to be mostly EPR silent; either they are in a diamagnetic spin state or a paramagnetic spin

Figure 9. (A) Simulated X-band CW EPR spectra of different Fe–S clusters. (B) X-band CW EPR spectra at 10 K of anaerobically activated SaDHAD and CjDHAD with and without sodium dithionite.
state with resonances beyond the field and frequency range of routine EPR spectrometers.$^{[26–52]}$ Indeed, aerobically and anaerobically activated CjDHAD is EPR silent (at 10 K, 9.38 GHz over the field range 0–1T), irrespective of the absence or presence of a reducing agent (Figure 9B). CW EPR spectra collected for SaDHAD under identical conditions contain only extremely weak signals consistent with the presence of a mixture of paramagnetic species at low concentration, probably indicating stages of cluster degradation (compare Figures 9A and 9B). The presence of a reducing agent produces no significant change to the EPR spectra. It thus appears that at least in SaDHAD, the Fe–S cluster quite rapidly disintegrates, consistent with the relatively rapid loss of catalytic activity (Figure 5).

Similar to the EPR spectra, the redox potentials of Fe–S clusters can vary significantly depending on the type of cluster that is present in the active site of an enzyme. For clusters of the [2Fe–2S] type potentials may vary between 0 and −0.4 V, while for [3Fe–4S] and [4Fe–4S] clusters potentials in the −0.2 V to −0.4 V and −0.2 V to −0.7 V range have been reported.$^{[54]}$ In order to assess the potential of Fe–S cluster of CjDHAD, spectrotelcochemical measurements (300 nm to 1000 nm) were performed with an anaerobically activated sample (Figure S6). Minimal changes in the UV-vis spectra were observed at potentials as low as −0.75 V vs. Ag/AgCl (i.e., −0.55 V vs. NHE). Overall, the spectral changes observed by sweeping from high to low potentials are insignificant. The peak at 410 nm, characteristic of an oxidized $[4Fe–4S]^{2+}$ cluster, persists throughout the experiments. During the course of the experiment, some loss of the Fe–S cluster via irreversible degradation was observed. Therefore, CjDHAD and SaDHAD, like aconitase, are not required to be reductive for catalysis. Indeed, reduction of the Fe–S cluster can lead to irreversible degradation of the active site.

Enzyme inhibition

DHAD, similar to other enzymes of the BCAA pathway (i.e., AHAS and KARI), is a suitable target for herbicides and antimicrobial agents.$^{[56–58]}$ N-isopropylxoyal hydroxamate (IpOHA; Figure 10), for instance, is a potent time-dependent inhibitor of KARI, the enzyme preceding DHAD in the BCAA pathway (Figure 1), with $K_i$ values in the nM range.$^{[6,21,59,60]}$ For both SaDHAD and CjDHAD, IpOHA also acts as a slow-binding inhibitor. In an assay containing 5 mM DHIV and 100 μM IpOHA, the inhibition increased from 2.1% to 91.9% upon pre-incubation for SaDHAD. The corresponding values for CjDHAD are 4.8% and 75.8%. Using pre-incubated samples, the $K_i$ values of SaDHAD and CjDHAD for IpOHA were determined to 7.8 μM and 32.9 μM, respectively (Figure 11). Thus, IpOHA is a good inhibitor for DHADs albeit not as effective as for KARI (with $K_i$ values ranging in the nM range$^{[6,21,59,60]}$). This is not surprising as IpOHA has been described as a transition state analog of KARI that resembles the substrate-bound rather than the product-bound state.$^{[26]}

Inhibition studies with aspterric acid (Figure 10), a known natural inhibitor of DHAD,$^{[26,29]}$ determined $K_i$ values of 51.6 μM and 35.1 μM for SaDHAD and CjDHAD, respectively (Figure 12). Aspterric acid has been tested against DHADs from plant and bacteria in recent studies$^{[26,28,29]}$ and was shown to be most potent for cyanobacterial DHAD (SnDHAD; $K_i = 9$ nM).$^{[26]}$ It was also shown to significantly inhibit both AtDHAD ($K_i = 0.3$ μM) and MdDHAD ($K_i = 10.1$ μM).$^{[26,29]}$ It is, however, currently unknown why the inhibitory potency of aspterric acid varies by three orders of magnitude between different DHADs. The

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**Figure 10.** Chemical structures of the known KARI inhibitor, IpOHA, and DHAD inhibitor, aspterric acid.

**Figure 11.** Inhibition of SaDHAD and CjDHAD by IpOHA (after 30-min pre-incubation).
nature of the Fe–S cluster alone may not account for this variation since SnDHAD, AtDHAD and MtDHAD contain [2Fe–2S] clusters, and SaDHAD and CjDHAD contain [4Fe–4S] clusters (although the enzymes with the less complex [2Fe–2S] clusters are generally stronger affected, possibly because their active sites are either more open or more flexible). In summary, both IpOHA and aspterric acid are suitable lead compounds for the development of novel anti-microbial drugs that target bacterial DHADs. IpOHA derivatives that may bypass the time-dependence of their interactions with KARI are currently being discovered. DHAD is still a relatively poorly characterised enzyme, and thus the characterisation of SaDHAD and CjDHAD provides significant functional and catalytic insight into these DHADs. A multiple sequence alignment of SaDHAD and CjDHAD with previously characterised DHADs suggests that they are likely to contain a [4Fe–4S] cluster like the DHAD from E. coli (EcDHAD). The Fe–S cluster-coordinating residues Cys-121 and Cys-194 and the Mg$^{2+}$ ion coordinating residues Asp-80, Asp-122, Lys-123 and Glu-446 are fully conserved in all DHADs, but the DHADs with a [4Fe–4S] cluster lack a cysteine in alignment position 48 (Figure 8). Recombinant SaDHAD and CjDHAD have been successfully expressed and purified from E. coli BL21 cells. Initial kinetic studies showed both enzymes to be virtually inactive, which was partially due to the lack of Fe–S cluster loading as indicated by ICP-OES analysis (Table 1). Reconstitution of the cluster by incubation with both Fe$^{2+}$ and reducing agents did not only increase the Fe–S loading of these enzymes, but also led to a dramatic increase in their catalytic rates (Figure 4). Stability experiments showed, however, that the activated enzymes lost their activity after 24 h under aerobic conditions at room temperature (Figure 5) but that activity can be preserved long term by storing the enzymes at −80 °C. The pH profiles for SaDHAD and CjDHAD indicate that optimal activity is reached at a pH of ~9 (Figure 6), slightly higher than the optima reported for other DHADs, but the catalytic mechanism is likely to be conserved. Furthermore, the thermostability of both SaDHAD and CjDHAD sharply decline at temperatures above 37 °C; at 60 °C the enzymes are essentially inactive (Figure 7). Thus, the sensitivity to inactivation by oxygen and elevated temperatures render both SaDHAD and CjDHAD as unsuitable for biotechnological processes that aim to exploit the catalytic potential of enzymes of the BCAA pathway to manufacture diverse platform chemicals (e.g. isobutanol$^{[12,13,31,32]}$). However, both SaDHAD and CjDHAD (and other bacterial DHADs) are suitable targets for the development of much needed new chemotherapeutics to treat infectious diseases. Catalytic measurements (Figures 11 and 12) have shown that IpOHA, a KARI inhibitor, and aspterric acid, a recently described DHAD inhibitor (Figure 10), inhibit both SaDHAD and CjDHAD potently with $K_i$ values in the low micromolar range. These compounds thus provide excellent starting points to further develop leads for novel treatments for infectious diseases. For instance, variants of IpOHA have recently been reported to be significantly more potent towards KARI from M. tuberculosis and similar modifications may thus also enhance the inhibitory potency of this compound for DHADs.$^{[22]}$

**Conclusions**

DHAD, along with other enzymes from the BCAA pathway (Figure 1), is an attractive target for novel anti-microbial drug discovery. DHAD is still a relatively poorly characterised enzyme, and thus the characterisation of SaDHAD and CjDHAD provides significant functional and catalytic insight into these DHADs. A multiple sequence alignment of SaDHAD and CjDHAD with previously characterised DHADs suggests that they are likely to contain a [4Fe–4S] cluster like the DHAD from E. coli (EcDHAD). The Fe–S cluster-coordinating residues Cys-121 and Cys-194 and the Mg$^{2+}$ ion coordinating residues Asp-80, Asp-122, Lys-123 and Glu-446 are fully conserved in all DHADs, but the DHADs with a [4Fe–4S] cluster lack a cysteine in alignment position 48 (Figure 8). Recombinant SaDHAD and CjDHAD have been successfully expressed and purified from E. coli BL21 cells. Initial kinetic studies showed both enzymes to be virtually inactive, which was partially due to the lack of Fe–S cluster loading as indicated by ICP-OES analysis (Table 1). Reconstitution of the cluster by incubation with both Fe$^{2+}$ and reducing agents did not only increase the Fe–S loading of these enzymes, but also led to a dramatic increase in their catalytic rates (Figure 4). Stability experiments showed, however, that the activated enzymes lost their activity after 24 h under aerobic conditions at room temperature (Figure 5) but that activity can be preserved long term by storing the enzymes at −80 °C. The pH profiles for SaDHAD and CjDHAD indicate that optimal activity is reached at a pH of ~9 (Figure 6), slightly higher than the optima reported for other DHADs, but the catalytic mechanism is likely to be conserved. Furthermore, the thermostability of both SaDHAD and CjDHAD sharply decline at temperatures above 37 °C; at 60 °C the enzymes are essentially inactive (Figure 7). Thus, the sensitivity to inactivation by oxygen and elevated temperatures render both SaDHAD and CjDHAD as unsuitable for biotechnological processes that aim to exploit the catalytic potential of enzymes of the BCAA pathway to manufacture diverse platform chemicals (e.g. isobutanol$^{[12,13,31,32]}$). However, both SaDHAD and CjDHAD (and other bacterial DHADs) are suitable targets for the development of much needed new chemotherapeutics to treat infectious diseases. Catalytic measurements (Figures 11 and 12) have shown that IpOHA, a KARI inhibitor, and aspterric acid, a recently described DHAD inhibitor (Figure 10), inhibit both SaDHAD and CjDHAD potently with $K_i$ values in the low micromolar range. These compounds thus provide excellent starting points to further develop leads for novel treatments for infectious diseases. For instance, variants of IpOHA have recently been reported to be significantly more potent towards KARI from M. tuberculosis and similar modifications may thus also enhance the inhibitory potency of this compound for DHADs.$^{[22]}$

**Materials and Methods**

**Transformation and expression trial**

The DHAD genes (ilvD) of Sa and Cj with hexa-histidine tag at the N-terminal of the polypeptide were cloned into pET-28a vector using NdeI and XhoI restriction sites and ordered from Gene Universal. The construct was transformed into E. coli BL21 cells using 100 μg/mL kanamycin plates. Single colonies were taken and inoculated in media for starter cultures and glycerol stock. For the expression trials, the protein was expressed in
1 mL autoinduction media. The cells were lysed and centrifuged at 14,500 rpm. The supernatant and the pellet samples were run on an SDS-PAGE gel to observe the presence of the proteins of interest.

**Protein expression and purification**

The inoculated cell culture was grown at 37°C in autoinduction media (ZYP-5052) containing 100 μg/mL kanamycin for three hours after which the temperature was decreased to 18°C and the protein was expressed overnight. The cells were harvested by centrifugation at 4000 rpm at 4°C for 30 minutes and was stored at –80°C until purification. The cell paste was resuspended in binding buffer with protease inhibitor cocktail, DNAse and lysozyme and was lysed by sonication. The binding buffer contained 300 mM NaCl, 20 mM imidazole, and 50 mM HEPES pH 8. The cell lysate was centrifuged at 4°C for one hour at 18000 RPM and the supernatant was loaded onto AKTA FPLC system. The protein was purified through immobilized metal affinity chromatography (IMAC) using 20 mM HisPrep Fast Flow 16/10 Ni-NTA-agarose resin column. The elution buffer used to elute the protein was same as the binding buffer except for the concentration of imidazole, which was increased to 300 mM. Next, the protein was further purified by size exclusion chromatography using a HiPrep 26/60 Sephacryl S-300 HR column with a buffer containing 10% glycerol and 50 mM HEPES pH 8. The protein concentration was quantified from its absorbance at 280 nm using its theoretical extinction coefficient calculated with Prot-param. TGX FastCast Acrylamide Kit from Bio-Rad was used to prepare all polyacrylamide gels. Precision Plus Protein Unstained Ladder was used to establish the standards. The gels were run with 1x running buffer at 110 V for 75 minutes. The gels were stained using Coomassie Instant Blue and destained with distilled water.

**Synthesis of DHIV**

Various syntheses have been reported for the formation of the DHAD substrates 2,3-dihydroxy-3-methylpentanoic acid (DHIV) or its derivative 2,3-dihydroxy-3-methylpentanoic acid (DHMV). Here, DHIV was synthesised in 25% yield across four steps from 3-methylbut-2-enoic acid (Scheme 1), similarly to the method reported by Chunduru et al. Details for the full synthesis are described in the Supporting Information.

![Scheme 1](image)

**Enzyme assay and characterisation**

For the enzyme assay, a modified version of the dinitrophenyl-hydrazide (DNPH) colorimetric endpoint assay described by Kiritani and Wagner and Katsuki et al. was used. DNPH was purchased from Sigma and 1.6 mg/mL was prepared in 2 N HCl. All assays were done at 37°C unless stated otherwise. A typical master reaction had a total volume of 2 mL and the reaction was started by adding dihydroxy-isovalerate (DHIV). Aliquots (380 μL) were taken from master reaction at each time point. The aliquot reaction was stopped by adding 0.1 volume (38 μL) of trichloroacetic acid (TCA) and was centrifuged at 14,500 rpm for 5 minutes to remove precipitation. Aliquot (360 μL) from the supernatant was taken and incubated with 120 μL of 1.6 mg/mL DNPH for 15 minutes at room temperature until 540 μL of 2 M NaOH was added. After 5 minutes, the formation of the hydrazone complex formed by the reaction of keto-acid with the hydrazine group of DNPH was measured spectrophotometrically at 430 nm. The amount of keto product was calculated from a standard curve that was plotted using KIV, which was purchased from Sigma. Michaelis-Menten plots were fitted using GraphPad Prism.

**Activation and stability of DHAD**

In order to activate DHAD, a modified protocol from Carsten et al. was used. 50 mM sodium dithionite, 200 mM 2-mercaptoethanol (2-ME) and 10 mM Fe²⁺ from ammonium ferrous sulfate were added to 500 μL of enzyme solution to make up a total volume of 3 mL. The solution was incubated at 37°C for 1 h before being buffer exchanged into 10% glycerol, 50 mM HEPES pH 8 solution using a 100G Econo desalting columns from Bio-Rad and concentrated for further experiments. The activated enzymes stored at –80°C retained stable activity for at least seven days (Figure S7). For the stability of activated CJ and SoDHADs, the activity was measured immediately after activation (t = 0), then after 30 minutes, one, two, four and 24 h. The activity was measured using the DNPH as described previously with 200 nM enzyme, 5 mM substrate, 5 mM MgCl₂ and 50 mM Tris-HCl, pH 8.5.

**pH profile and thermostability**

SoDHAD and CJDHAD were assayed with pH ranging from 7.0 to 10.0 to determine its pH profile. For pH 7.0 to 8.0 HEPES, for 8.0 to 9.0 Tris-HCl and for 9.5 to 10 glycine buffers were used. For the thermostability experiment, the enzyme was preincubated in the assay solution at said temperature for 15 minutes. Then the substrate was added and the reaction was monitored at 37°C with timepoints taken every five minutes. The rate was determined using the DNPH assay with 100 nM SoDHAD (50 nM CJDHAD), 5 mM substrate, 5 mM MgCl₂ and 50 mM of Tris-HCl pH 8.5.
EPR and spectrophotocatalystrochemistry of SaDHAD and CjDHAD

X-band CW EPR measurements were carried out at 10 K (Bruker E500 spectrometer equipped with a He closed cycle cryostat) on ~50 μM/100 μL of activated Sa and CjDHADs. Sodium dithionite was added to the reduced samples to a concentration of 200 μM. Spectroelectrochemistry was carried out (under nitrogen) with ~50 μM of anaerobically activated CjDHAD at 10°C. Spectra were measured down to -0.75 V vs. Ag/AgCl (−0.55 V vs. NHE) with a Pine Instruments honeycomb Au spectroelectrochemical electrode and thin layer cell. Mediators for electron transfer were the complexes [Co(AMMEN,Ssan)]³⁺, [Co(sep)]³⁺, [Co(AMMESan)]³⁺, [Co(cis-diammac)]³⁺ and [Co(trans-diammac)]³⁺ all at a concentration of 20 μM. The experiments were carried out under a constant blanket of nitrogen over the top of the cell.

Inhibition of DHAD

The inhibition of SaDHAD and CjDHAD by IpOHA and aspterric acid was evaluated using the DNPH assay. The reaction mixture contained 600 nM SaDHAD (100 nM CjDHAD), 5 mM DHIV, 5 mM MgCl₂ in 50 mM Tris-HCl pH 8.5. Inhibition of IpOHA was determined with 30-minute preincubation before starting the reaction with the addition of the substrate. Concentrations between 0 to 200 μM of IpOHA were used to determine the Kᵢ value. For assays with aspterric acid, no pre-incubation was applied and the inhibitor concentration ranged from 0 to 600 μM. The inhibition constants for IpOHA and aspterric acid were calculated using the simple inhibition equation (Equation [1]).

\[
V_i = \frac{V_u}{1 + \frac{[I]}{K_i}}
\]

where \(V_i\) is the rate at the inhibitor concentration \([I]\), \(V_u\) is the uninhibited rate and \(K_i\) is the inhibition constant.

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Conflict of Interest

The authors declare no conflict of interest.

Data Availability Statement

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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