Synthesis of 2-oxoglutarate derivatives and their evaluation as cosubstrates and inhibitors of human aspartate/asparagine-β-hydroxylase†

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2-Oxoglutarate (2OG) is involved in biological processes including oxidations catalyzed by 2OG oxygenases for which it is a cosubstrate. Eukaryotic 2OG oxygenases have roles in collagen biosynthesis, lipid metabolism, DNA/RNA modification, transcriptional regulation, and the hypoxic response. Aspartate/asparagine-β-hydroxylase (AspH) is a human 2OG oxygenase catalyzing post-translational hydroxylation of Asp/Asn-residues in epidermal growth factor-like domains (EGFDs) in the endoplasmic reticulum. AspH is of chemical interest, because its Fe(II) cofactor is complexed by two rather than the typical three residues. AspH is upregulated in hypoxia and is a prognostic marker on the surface of cancer cells. We describe studies on how derivatives of its natural 2OG cosubstrate modulate AspH activity. An efficient synthesis of C3- and/or C4-substituted 2OG derivatives, proceeding via cyanosulfur ylid intermediates, is reported. Mass spectrometry-based AspH assays with >30 2OG derivatives reveal that some efficiently inhibit AspH via competing with 2OG as evidenced by crystallographic and solution analyses. Other 2OG derivatives can substitute for 2OG enabling substrate hydroxylation. The results show that subtle changes, e.g. methyl- to ethyl-substitution, can significantly alter the balance between catalysis and inhibition. 3-Methyl-2OG, a natural product present in human nutrition, was the most efficient alternative cosubstrate identified; crystallographic analyses reveal the binding mode of (R)-3-methyl-2OG and other 2OG derivatives to AspH and inform on the balance between turnover and inhibition. The results will enable the use of 2OG derivatives as mechanistic probes for other 2OG utilizing enzymes and suggest 2-oxoacids other than 2OG may be employed by some 2OG oxygenases in vivo.

Introduction

2-Oxoglutarate (2OG, α-ketoglutarate; 1, Fig. 1a) is an integral metabolite in most of biology including prokaryotes, archaea, and eukaryotes; 2OG is crucially involved in cellular energy homeostasis and small-molecule metabolism and can act as a signaling molecule linking nitrogen and carbon metabolism. 2OG is an intermediate of the tricarboxylic acid (TCA) cycle where it is produced by isocitrate dehydrogenase (IDH)-catalyzed decarboxylation of β-isocitrate; 2OG is converted to succinyl-CoA and CO₂ by the 2OG dehydrogenase complex. Reductions in cellular 2OG coupled with the formation of (R)-2-hydroxyglutarate, which occur as a result of IDH mutations, are associated with changes in epigenetic regulation (e.g. DNA and histone methylation status) and certain types of cancer. These effects are proposed to be mediated, at least in part, by modulation of the activities of enzymes that rely on 2OG as a (co-)substrate, other than the 2OG dehydrogenase complex. Such enzymes include aminotransferases (e.g. branched-chain aminotransferases, BCATs), which convert 2OG to glutamate, and 2OG dependent oxygenases. The latter couple the conversion of 2OG and O₂ to succinate and CO₂ with substrate hydroxylation or demethylation via hydroxylation. There are approximately 60–70 assigned human 2OG oxygenases, all studied members of which likely employ Fe(II) as a cofactor and which have diverse roles, ranging from DNA/RNA modification and damage repair, histone/chromatin modification, lipid metabolism, post-translational modification of proteins with important functions in the extracellular matrix, to hypoxia sensing.

The human 2OG oxygenase aspartate/asparagine-β-hydroxylase (AspH, BAH, HAAH) is highly unusual amongst human 2OG dependent hydroxylases, because its Fe(II) cofactor is complexed by only two residues (His679 and His725) rather than by the typical triad of ligands (HisX/E/H) found in other human 2OG oxygenases (Fig. 1b). AspH catalyzes the post-translational hydroxylation of specific Asp- and Asn-residues in epidermal growth factor-like domains (EGFDs) of its substrate proteins in the endoplasmic
AspH is of significant interest from a cancer research perspective, because its levels are upregulated in invasive cancers (e.g. hepatocellular carcinoma\textsuperscript{16} and pancreatic cancer\textsuperscript{17}) and it is translocated to the cell surface where it can be used as a diagnostic and prognostic marker.\textsuperscript{18} Mouse models\textsuperscript{19} and heritable genetic diseases associated with mutations likely effecting AspH catalysis (e.g. Traboulsi syndrome)\textsuperscript{20} suggest that the Notch signaling pathway may be involved in transmitting the effect of AspH on cancer invasiveness. AspH levels are regulated by hypoxia which is a characteristic of many tumor cells.\textsuperscript{21} Thus, AspH appears to be an attractive medicinal chemistry and diagnostic target for certain types of cancer.

Recent studies have provided crystallographic and solution-based evidence that AspH accepts EGFD substrates with an unusual non-canonical disulfide connectivity (i.e. Cys 1–2, 3–4, 5–6) rather than the well-characterized canonical disulfide connectivity (i.e. Cys 1–3, 2–4, 5–6; ESI Fig. S1\textsuperscript{†}). High-throughput MS assays were established to monitor the catalytic activity of AspH using stable thioether-linked cyclic peptide substrate analogues mimicking the central non-canonical macrocyclic Cys 3–4 EGFD disulfide (ESI Fig. S1\textsuperscript{†}). Kinetic studies have revealed that AspH is sensitive towards subtle changes in oxygen availability and thus it is a candidate oxygenase for involvement in hypoxia sensing.\textsuperscript{23}

Previous studies have revealed that differences in the cosubstrate binding sites of 2OG oxygenases can be exploited for stereoselective selective inhibition employing 2OG analogues, e.g. N-oxalylamino acids.\textsuperscript{24} Analysis of reported AspH crystal structures wherein 2OG is replaced by a close 2OG analogue, i.e. N-oxalylglycine (NOG, 3; Fig. 1c),\textsuperscript{22} suggest that the AspH active site is sufficiently spacious to

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**Fig. 1** The AspH active site and stoichiometry of its reaction. (a) AspH catalyzes the post-translational hydroxylation of Asn- and Asp-residues in epidermal growth factor (EGF)-like domains; (b) analysis of an AspH-substrate (human Factor X, hFX) crystal structure (PDB ID: 5JQY)\textsuperscript{22} reveals that two AspH residues (His679 and His725) coordinate the active site metal rather than the typical three residues found in other human 2OG dependent hydroxylases. In the crystallographic analyses, Mn substitutes for Fe and NOG (3) for 2OG (1); (c) N-oxalylglycine (NOG, 3).

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**Fig. 2** Selected reported strategies for the synthesis of C3- and/or C4-substituted 2OG derivatives. (a) Alkylation reactions,\textsuperscript{25} (b) Michael reactions,\textsuperscript{27} and (b and c) oxidation reactions\textsuperscript{28a–c,29} have been employed in syntheses of C3/C4-substituted 2OG derivatives (6).
accompany substitutes at the 2OG C3- and/or C4-position, in part due to its unusual Fe([ii])-binding geometry (Fig. 1b). We were therefore interested to test this by exploring how a diverse set of 2OG derivatives interact with AspH.

To our knowledge, only a limited number of studies describing how 2OG derivatives bearing substituents at the C3- and/or C4-position modulate the activities of 2OG oxygenases (i.e., human JMjC histone N'-methyl lysine demethylase 4A, KDM4A,15 human factor inhibiting the hypoxia-inducible transcription factor HIF-α, FIH;24 and rat γ-butyrobetaine dioxygenase, BBOX25) are reported. In part, this likely reflects a lack of a simple synthetic method to access these types of 2OG derivatives. Some prior syntheses of C3/C4-substituted 2OG derivatives have relied inter alia on alkylation (Fig. 2a)25 and Michael reactions (Fig. 2b)27 to access key synthetic intermediates; the corresponding 2OG derivatives (6) were obtained after saponification. Other approaches rely on oxidation reactions using ozone28 or sodium periodate29 as oxidants to convert Michael acceptors (10) into 2OG derivatives (2c). The described syntheses are frequently associated with limited scalability, low overall chemical yields, and/or narrow substrate scopes due to harsh reaction conditions requiring, for example, the use of strong bases and acids,5,28 high pressure,27 or strong oxidants.28–30

Cyanophosphorous ylids are reported as valuable intermediates for the synthesis of ω-keto acids,13 however, only one example of a cyanophosphorous ylid (i.e., 11) being converted into a 2OG derivative is reported (Fig. 2c),27 possibly reflecting limitations associated with the conversion of cyanophosphorous ylids into ω-keto acids which requires strong oxidants.28–31 The use of cyanophosphorous ylids has been largely superseded by the corresponding cyanosulfur ylids, which can be oxidized under milder conditions.31,32 We thus envisaged that cyanosulfur ylids could be used for the synthesis of C3/C4-substituted 2OG derivatives.

Here we report the use of cyanosulfur ylids as intermediates that enable the facile synthesis of multiple 2OG derivatives bearing a diverse set of substituents at the C3- and/or C4-positions. The synthetic 2OG derivatives were used to modulate the activity of recombinant human AspH. Kinetic and crystallographic studies were employed to elucidate the mechanisms by which the 2OG derivatives modulate AspH activity and to garner information of the active site requirements of AspH. The results reveal an unexpectedly diverse set of 2OG derivatives can bind at the AspH active site and that subtle differences in the 2OG substitution pattern can cause significant disturbances in the balance between productive catalysis and inhibition.

**Results**

**Synthesis of 2OG derivatives**

A versatile and scalable synthetic route to access 2OG derivatives was developed employing cyanosulfur ylids as key intermediates (Scheme 1). The route employs mono-methyl dicarboxylic acid half-esters 13 as starting materials which were either commercially available or synthesized by established reactions, i.e. nuclophilic openings of the requisite symmetric cyclic anhydrides, formylation reactions of aryl iodides,35 Heck couplings36 of aryl iodides with orthogonally protected itaconates,37 or Horner–Wadsworth–Emmons (HWE)38 reactions (ESI Fig. S2†). In our work, racemic mixtures of mono-methyl dicarboxylic acid half-esters 13 bearing stereogenic carbon atoms at the 2OG C3- or C4-equivalent position were employed.

Cyanosulfur ylids 15, which are the key intermediates in our strategy, were obtained by reaction of mono-methyl dicarboxylic acid half-esters 13 with the reported tetrahydrothiophene bromide salt 1414 in yields ranging from 11 to 95%. T3P29 was chosen as the coupling reagent because it is suited for use with sterically hindered carboxylic acids, including those bearing substituents at the carboxylate ω-position. For some substrates, T3P-derived byproducts interfered with the purification process; however, these were completely removed after the subsequent reaction by chromatography.

The cyanosulfur ylids 15 were converted into the corresponding dimethyl dicarboxylic acid esters 12 using oxone38 as a mild oxidation reagent in methanol, in part to avoid ester exchange (Scheme 1). The dimethyl esters 12 were obtained in high purity after column chromatography in yields ranging from 39 to 98%. Lithium hydroxide-mediated saponification of dimethyl dicarboxylic acid esters 12 afforded the desired 2OG derivatives 6 (Scheme 1). The 2OG derivatives were obtained in sufficient purity after removal of excess base by acidic ion exchange chromatography yielding salt-free dicarboxylic acids 6 (ESI†), which are suitable for performing in vitro biochemical experiments with AspH. The 2OG derivatives and their synthetic precursors were stable when stored at −20 °C for more than six months.

![Scheme 1](https://example.com/scheme1.png)

**Scheme 1** Route and substrate scope for the synthesis of 2OG derivatives. Reagents and conditions: (a) T3P, 1,2,2,2-tetramethylimidazoline-1-oxyl-3-carboxylate (TEMPO), CH2Cl2, 0 °C to rt, 11–95%; (b) oxone, MeOH/H2O, rt, 39–98%; (c) LiOH, MeOH/H2O, 0 °C to rt, then: purification by ion exchange chromatography (Dowex® 50X8), 67% to apparent quantitative.
Scope of the synthesis
Following its development, the synthetic route was used to synthesize a diverse set of 2OG derivatives bearing aliphatic substituents at the 2OG C3- and/or C4-positions (Table 1, entries 1–21). The aliphatic substituents varied in both length and steric bulk of the carbon chain. Furthermore, 2OG derivatives were synthesized in which the C3/C4 ethylene unit of 2OG was replaced with different groups (Table 1, entries 22–32).

Table 1  Inhibition of AspH by 2OG derivatives

| 2OG derivative<sup>a</sup> | IC<sub>50</sub><sup>b,c</sup> [μM] | 2OG derivative<sup>a</sup> | IC<sub>50</sub><sup>b,c</sup> [μM] | 2OG derivative<sup>a</sup> | IC<sub>50</sub><sup>b,c</sup> [μM] |
|------------------------|-----------------|------------------------|-----------------|------------------------|-----------------|
| 1                      | Inactive        | 12                     | 0.61 ± 0.09     | 23                     | 12.9 ± 1.3      |
| 2                      | 1.2 ± 0.5       | 13                     | 0.47 ± 0.08     | 24                     | Inactive        |
| 3                      | 5.7 ± 1.1       | 14                     | 0.51 ± 0.12     | 25                     | Inactive        |
| 4                      | 48.2 ± 13.1     | 15                     | 0.70 ± 0.11     | 26                     | Inactive        |
| 5                      | 6.8 ± 0.9       | 16                     | 0.25 ± 0.05     | 27                     | Inactive        |
| 6                      | 1.6 ± 0.3       | 17                     | 0.43 ± 0.05     | 28                     | Inactive        |
| 7                      | 2.6 ± 0.8       | 18                     | 0.17 ± 0.03     | 29                     | Inactive        |
| 8                      | 6.3 ± 2.6       | 19                     | 0.3 ± 0.1       | 30                     | Inactive        |
| 9                      | 3.6 ± 1.4       | 20<sup>d</sup>         | 5.2 ± 1.7       | 31                     | Inactive        |
| 10                     | 4.7 ± 0.2       | 21<sup>e</sup>         | 19.3 ± 1.6      | 32                     | Inactive        |
| 11                     | Inactive        | 22                     | Inactive        | 33                     | 3.3 ± 1.0       |

<sup>a</sup> All chiral 2OG derivatives were prepared as racemic mixtures. <sup>b</sup> Mean of three independent runs (n = 3; mean ± SD). AspH inhibition assays were performed as described in the ESI using 50 nM His<sub>6</sub>-AspH<sub>315–758</sub> and 1.0 μM hFX-CP<sub>101–119</sub> (ESI Fig. S1d) as a substrate. <sup>c</sup> 2OG derivatives were termed inactive when the IC<sub>50</sub>-values were >50 μM. The AspH inhibition assays were of good quality which high S/N ratios and Z<sub>f</sub>-factors<sup>40</sup> (>0.5 for each plate) indicate (ESI Fig. S3). <sup>d</sup> Mixture of racemic diastereomers, dr (cis : trans) = 2.5 : 1. <sup>e</sup> (±)-(2-Exo,3-endo)-diastereomer.
was replaced by rings including heteroaromatic rings (Table 1, entries 22 and 23), aromatic rings (Table 1, entries 24–31), and aliphatic bicyclic rings (Table 1, entry 32). A cyclopropane-containing 2-oxocid that was not based on the glutarate C5 skeleton but on the succinate C4 skeleton (48; Table 1, entry 33), was synthesized.

The synthesis of 2OG derivatives bearing acid-labile or some oxidation-prone moieties was challenging. For example, during the cyano sulfur ylid oxidation reaction, both ketal and silyl ether alcohol protecting groups were cleaved and nitrogen-containing heteroaromatic rings (e.g., pyridines) formed N-oxides. Nonetheless, the oxidation conditions were sufficiently mild to tolerate alkenes (36; Table 1, entry 21) and substituted thiophenes (37 and 38; Table 1, entries 22 and 23), which constitutes an advantage compared to many prior syntheses of 2OG derivatives (Fig. 2).

AspH inhibition studies

We then evaluated the potential of the synthesized racemic 2OG derivatives to inhibit AspH by measuring AspH substrate depletion and product formation (i.e., by monitoring a +16 Da mass shift) using an established solid phase extraction coupled to mass spectrometry (SPE-MS) AspH inhibition assay. Half maximum inhibitory concentrations (IC50 values) for all the synthetic 2OG derivatives prepared were determined (Table 1).

3-Methyl-2OG (16) did not inhibit AspH (i.e., IC50 > 50 μM), while 2OG derivatives bearing longer carbon chains at the C3-position, including 3-ethyl-2OG (17), were efficient inhibitors, under the assay conditions. The inhibitory potency decreased on increasing the length or steric bulk of the carbon chain of the C3-substituent beyond that of an ethyl group, i.e., C3-Et (17): IC50 ~ 1.2 μM vs. C3-Pr (18): IC50 ~ 5.7 μM, C3-CH2Ph (21): IC50 ~ 1.6 μM vs. C3-CH2CH2CH2CH2Ph (20): IC50 ~ 6.8 μM, and C3-CH2CH2CH2Ph (20): IC50 ~ 6.8 μM vs. C3-CH2CH2CH2CH2CH2 (19): IC50 ~ 18.2 μM. For the C3 benzy l substituted 2OG derivatives 21–25 with differently substituted phenyl rings, the substitution pattern on the phenyl rings appears to not affect the inhibitor potency within experimental error (Table 1, entries 6–10). With the exception of 4-methyl-2OG (26), which did not inhibit AspH (Table 1, entry 11), 2OG derivatives bearing substituents at the 2OG C4-position (Table 1, entries 12–18) were substantially more potent in inhibiting AspH than those bearing substituents at the C3-position: their relative potencies increased by a factor of 2 (for C3/4-Et to ~70 for C3/4-CH2CH2CH2 (13)). The IC50 values of the C4-substituted 2OG derivatives range between 0.2 and 0.7 μM (Table 1, entries 12–18) and did not appear to depend on length or bulk of the tested C4-substituents. The C4-substituted 2OG derivatives inhibit AspH with comparable efficiency as the broad-spectrum 2OG oxygenase inhibitor NOG (3; IC50 ~ 1.0 μM; ESI Fig. S4†), but less efficiently than pyridine-2,4-dicarboxylate (2,4-PDCA; IC50 ~ 0.03 μM), which is another broad-spectrum 2OG oxygenase inhibitor.

4,4-Dimethyl-2OG (34) was an efficient AspH inhibitor (IC50 ~ 0.3 μM; Table 1, entry 19), whilst the more bulky bicyclo[2.2.2] octane-bearing 2OG derivative 47 did not inhibit (Table 1, entry 32). The dimethylcyclopropane-bearing 2OG derivative 35 (2.5:1 mixture of racemic cis/trans-diastereomers) inhibits (IC50 ~ 5.2 μM; Table 1, entry 20), albeit less efficiently than 34. This might reflect the reduced rotational flexibility of 35 around the C3–C4 bond due to the presence of the cyclopropane ring. 2OG derivative 36, which is also C3/C4-disubstituted, but which is significantly more bulky than 35, inhibits AspH (IC50 ~ 19.3 μM), but substantially less efficiently than does 35 (Table 1, entry 21). The thiophene-based 2OG derivative 37 did not inhibit AspH (Table 1, entry 22). By contrast, the regioisomeric thiophene-based 2OG derivative 38 inhibited AspH with moderate efficiency (IC50 ~ 12.9 μM; Table 1, entry 23). This observation might reflect the ability of 38, but likely not of 37 to better chelate Fe(n), including that at the AspH active site. Several Fe(n)-chelators have been previously identified to inhibit AspH in inhibitor screens.41a

The phenyl ring regioisomers 39, 40, and 41, which are 2OG derivatives bearing an aromatic core, did not inhibit AspH (Table 1, entries 24–26), as it was the case for the other tested phenyl ring containing 2OG derivatives 42–46 (Table 1, entries 27–31). The 2OG derivative 48 whose carbon scaffold was not based on glutarate, but derived from succinate, inhibited AspH with moderate efficiency (IC50 ~ 3.3 μM; Table 1, entry 33).

Interestingly, 4-benzyl-2OG (32) inhibits AspH significantly more efficiently (IC50 ~ 0.4 μM; Table 1, entry 17) than its N-oxalyl analogue N-oxalyl-a-phenylalanine (NOFD; IC50 ~ 15.5 μM),44 which is a reported inhibitor of human FIH.44 An opposite trend was observed for FIH, for which NOFD was a substantially more efficient inhibitor than 32,44 revealing the context dependent effect of the same 2OG substitutions. The structures of these two inhibitors are very similar: the C3 methylene-unit of 4-benzyl-2OG (32) is substituted for an NH-group in NOG and NOFD, which are C3/C4-disubstituted, but which have the cyanosulfur ylid oxidation reaction, both ketal and silyl ether alcohol protecting groups were cleaved and nitrogen-containing heteroaromatic rings (e.g., pyridines) formed N-oxides. Nonetheless, the oxidation conditions were sufficiently mild to tolerate alkenes (36; Table 1, entry 21) and substituted thiophenes (37 and 38; Table 1, entries 22 and 23), which constitutes an advantage compared to many prior syntheses of 2OG derivatives (Fig. 2).

2OG derivatives compete with 2OG for binding the AspH active site

To define whether the mechanism by which the 2OG derivatives inhibit AspH involves competition with 2OG for binding to the AspH active site, the effect of altered 2OG concentrations on the IC50 values of AspH was investigated. The IC50 values of four potent AspH inhibitors (i.e., 17, 29, 33, and 34) were determined at 2OG assay concentrations of 3, 200, 400, and 600 μM (Fig. 3a; ESI Table S1†). The results reveal an ascending linear dependence of the IC50 values on the 2OG assay concentration suggesting that the 2OG derivatives inhibit AspH by competing
with 2OG for binding to the active site. This is in agreement with a Hill coefficient analysis of the AspH inhibition curves which indicates that the 2OG derivatives did not inhibit AspH by forming colloidal aggregates; the Hill coefficients are in the range of the expected ‘ideal’ value –1 (Fig. 3b).44

Neither the position nor the size of the C3/C4-substituent of the 2OG derivatives had a detrimental effect on the linear dependence of their AspH IC50-values on the 2OG concentration (Fig. 3a). Efficient inhibition of AspH at higher 2OG assay concentrations was observed for 2OG derivatives 33 and 34 (IC50 ~ 5.6 and 18.7 μM at 0.6 mM 2OG assay concentration, respectively; ESI Table S1,† entries 3 and 4).

Some 2OG derivatives can replace 2OG as an AspH cosubstrate

During the assessment of the 2OG inhibition data, we observed that the presence of two 2OG derivatives, i.e. 3-methyl-2OG (16) and 4-carboxyphenylglyoxylic acid (41), the extent of AspH-catalyzed substrate hydroxylation appeared to increase. We proposed that these 2OG derivatives could replace 2OG and function as alternative substrates for AspH. AspH substrate hydroxylation was thus investigated in the absence of 2OG: high levels of AspH substrate hydroxylation were observed at elevated concentrations of the 2OG derivatives 16 (~95%) and, somewhat unexpectedly, the phenyl ring derivative 41 (~80%) in the absence of 2OG, demonstrating that these two 2OG derivatives can replace 2OG as an AspH substrate (Fig. 4a–c). NMR studies revealed that AspH converts the 2OG derivatives 16 and 41 into 2-methylsuccinate (50) and terephthalate (51), respectively, in an analogous manner to which it converts 2OG into succinate (Fig. 4a–c; ESI Fig. S5†).

To investigate whether the other synthesized 2OG derivatives can substitute for 2OG as a cosubstrate for AspH-catalyzed hydroxylation, assays were performed in the absence of 2OG, but in the presence of high concentrations (330 μM) of all the synthetic 2OG derivatives (ESI Table S2†). In addition to the 2OG derivatives 16 and 41, AspH substrate hydroxylation was observed for the 2OG derivatives 26 (~10%), 38 (~15%), 42 (~45%), 43 (~8%), 44 (~20%), 45 (~8%), 46 (~5%), and 47 (~8%) after 15 minutes. These results reveal that 3-methyl-2OG (16) is a substantially more efficient alternative AspH cosubstrate than its isomer 4-methyl-2OG (26), an observation in accord with the observation that C3-substituted 2OG derivatives are generally less efficient AspH inhibitors than their corresponding C4-substituted isomers (Table 1).

Increasing the steric bulk of the 2OG derivative, while maintaining the relative arrangement of the two carboxylate groups, decreases the catalytic efficiency of the cosubstrate analogue, as revealed by the comparison of the phenyl-ring 2OG derivative 41 (~80%) with its bridged bicyclo[2.2.2]octane analogue 47 (~8%). Derivatives of 41 bearing substituents ortho to the ketone (e.g. as in 42, ~45%) seemed to be more efficient AspH cosubstrates than those bearing substituents meta to the ketone (e.g. as in 43, ~8%). Increasing the size of the substituents ortho to the ketone of 2OG derivative 41 (i.e. ortho-F, 42; ortho-Br, 44; ortho methyl 46) results in a noticeable decrease in the efficiency to replace 2OG as an AspH cosubstrate (i.e. 42: ~45%; 44: ~20%; 46: ~5%); all derivatives of 41 were significantly less efficient with respect to the parent compound (~80%).

Kinetic analyses of the two most efficient 2OG substitute AspH cosubstrates identified (i.e. 16 and 41) were performed. Maximum velocities (v_max) and Michaelis constants (K_M) were determined employing SPE-MS turnover assays, albeit under modified conditions than previously reported for 2OG as a cosubstrate (Fig. 4d–f),23 as L-ascorbic acid (LAA), which is commonly added to 2OG oxygenase assays, was included in the assay buffer. The presence of LAA affected the kinetic parameters for 2OG when compared to the previous parameters (Table 2, entry 1). The v_max (2OG) did not change significantly, being ~16.8 × 10⁻³ μM s⁻¹ in the absence of LAA and ~15.0 × 10⁻³ μM s⁻¹ in the presence of LAA. The K_M (2OG)-value in the presence of LAA was approximately double (~1.3 μM) compared to that in the absence of LAA (~0.6 μM).23 However, this K_M (2OG)-value is still in the range of those values reported for most other human 2OG oxygenases including for the HIF-α prolyl hydroxylases and FIH (1–25 μM)44 and bovine AspH (~5 μM).45

**Fig. 3** Inhibition of AspH by 2OG derivatives. (a) The AspH IC50-values for the 2OG derivatives 17 (black diamonds), 29 (red squares), 33 (orange circles), and 34 (green triangles) depend on the 2OG concentration. AspH inhibition assays were performed in triplicate as described in the ESI† using 3, 200, 400, and 600 μM 2OG. IC50-values are summarized in ESI Table S1;† (b) representative dose–response curves used to determine IC50-values for the 2OG derivatives 17 (black diamonds), 29 (red squares), 33 (orange circles), and 34 (green triangles) at a 2OG assay concentration of 3 μM 2OG. Three dose–response curves each composed of technical duplicates were independently determined using SPE-MS AspH inhibition assays, performed as described in the ESI† and manifest high Z*-factors46 and signal-to-noise ratios (ESI Fig. S3†).
Compared to the $K_{\text{app}}$-value of AspH for 2OG ($1.3 \text{ mM}$; Table 2, entry 1), the $K_{\text{app}}$-value for 3-methyl-2OG (16) was about five times lower ($0.3 \text{ mM}$; Table 2, entry 2), indicating a higher affinity of AspH for 16 compared to 2OG. By contrast, the $K_{\text{app}}$-value of AspH for the 2OG derivative 41 was ~47 times higher (~62 $\mu$M; Table 2, entry 3) than that for 2OG, indicating much

**Table 2** Steady-state kinetic parameters for AspH. Maximum velocities ($v_{\text{app}}^\text{max}$), Michaelis constants ($K_{\text{app}}$), turnover numbers ($k_{\text{cat}}$), and specificity constants ($k_{\text{cat}}/K_m$) of His$_6$-AspH$_{315-758}$ for 2OG and the 2OG derivatives 16 and 41.$^{a,b}$

| AspH co-substrate                  | $v_{\text{app}}^\text{max}$ [$\mu$M s$^{-1}$] | $K_{\text{app}}$ [$\mu$M] | $k_{\text{cat}}$ [s$^{-1}$] | $k_{\text{cat}}/K_m$ [M$^{-1}$ s$^{-1}$] |
|-------------------------------------|-----------------------------------------------|----------------------------|----------------------------|-----------------------------------------|
| 1 2OG                               | $15.0 \times 10^{-3} \pm 0.9 \times 10^{-3}$ | $1.3 \pm 0.3$             | $0.17 \pm 0.03$            | $0.13 \pm 0.03$                        |
| 2 3-Methyl-2OG (16)                 | $6.9 \times 10^{-3} \pm 0.2 \times 10^{-3}$ | $0.27 \pm 0.04$           | $0.08 \pm 0.01$            | $0.30 \pm 0.06$                        |
| 3 4-Carboxyphenylglyoxylic acid (41) | $4.0 \times 10^{-3} \pm 0.3 \times 10^{-3}$ | $61.9 \pm 11.0$           | $0.04 \pm 0.01$            | $6.5 \times 10^{-3} \pm 0.2 \times 10^{-3}$ |

$^{a}$ Mean of three independent runs ($n = 3$; mean ± SD). $^{b}$ AspH assays were performed as described in the ESI using 0.1 $\mu$M His$_6$-AspH$_{315-758}$ and 2.0 $\mu$M hFX-CP$_{101-119}$ (ESI Fig. S1d) as a substrate.
less efficient binding. All three $k_{\text{cat}}^{\text{PP}}$-values range significantly below reported 2OG concentrations in healthy cells (up to $>1$ mM), which however vary substantially, but are in the approximate range of reported physiological 2OG levels in human plasma (9–12 μM 2OG).

Based on the determined concentration of active AspH (90.8 ± 13.7 nM for an original estimated AspH assay concentration of 100 nM AspH), turnover numbers (catalytic constants, $k_{\text{cat}}$) and specificity constants ($k_{\text{cat}}/K_m$) were calculated for the 2OG derivatives (Table 2). Comparison of the $k_{\text{cat}}$-values reveals that the impact of the 2OG derivatives on $k_{\text{cat}}$-values is notably smaller than on $k_{\text{cat}}^{\text{PP}}$-values which indicates that efficient AspH-catalyzed substrate hydroxylation is still feasible with the 2OG derivatives (Table 2): for AspH, the $k_{\text{cat}}$-value for 16 (~0.08 s⁻¹; Table 2, entry 2) was about half the $k_{\text{cat}}$ for 2OG (~0.17 s⁻¹; Table 2, entry 1), whereas the $k_{\text{cat}}$-value for 41 (~0.04 s⁻¹; Table 2, entry 3) was about a quarter of that for 2OG. Comparison of the $k_{\text{cat}}/K_m$-values clearly reveals the potential of 2OG derivatives to substitute for 2OG itself, with racemic 16 being of similar efficiency to 2OG. Although 41 is a much less efficient substrate, its conversion reveals the potential for unexpected cosubstrate utilization by 2OG oxygenases.

Crystallography

The AspH turnover assays indicated that the synthetic 2OG derivatives compete with 2OG for binding to the AspH active site. To investigate the divergent effects of C3/C4-substituted 2OG derivatives on AspH catalysis, i.e. AspH inhibition or promoting AspH activity, crystallographic studies were initiated. In the reported AspH crystal structures, the natural AspH cosubstrate 2OG was substituted for a 2OG competing inhibitor (e.g. NOG, 2,4-PDCA or l-malate), but an AspH crystal structure complexed with 2OG has not previously been reported.

To enable comparisons of how 2OG and the 2OG derivatives bind AspH, AspH was first crystallized in the presence of 2OG with the natural AspH cofactor Fe(n) being replaced by Mn(n). AspH crystallized in the absence of substrate in the P2₁2₁2₁ space group (AspH:2OG; 2.1 Å resolution), the structure, as were the subsequently described structures, was solved by molecular replacement using a reported AspH structure (PDB ID: 5JTC) as a search model (ESI Fig. S7†). Clear electron density corresponding to 2OG was observed (Fig. 5a); the C5-carboxylate of 2OG being positioned to form a salt bridge with the side chain of Arg735 (2.4 and 3.0 Å) and to interact with Ser668 (2.6 Å), which is part of an ‘RXS motif’ present in some other 2OG oxygenases. The C1-carboxylate of 2OG is positioned to interact with Arg688 (2.7 and 2.9 Å) and His690 (3.3 Å). The C1-carboxylate of 2OG and the 2OG C2-carbonyl group complex the Mn ion in a bidentate manner (1.6 and 2.5 Å; Fig. 5a). Two water molecules also coordinate the Mn ion (2.1 and 2.4 Å) along with the two anticipated residues His679 and His725 (2.2 and 2.1 Å; Fig. 5a). Thus, the AspH:2OG structure supports the proposal (based on the AspH structures in complex with the 2OG analogue NOG) that the active site metal, when bound to the natural AspH cosubstrate 2OG, is complexed by only two AspH residues rather than by the typical triad of ligands (HXD/E...H) found in other human 2OG hydroxylases. Superimposition of the AspH:2OG structure with the reported AspH:NOG and AspH:malate structures reveals that AspH adopts similar conformations in all structures and that 2OG binds the AspH active site in a similar manner to NOG (Cz RMSD = 0.21 and 0.21 Å, ESI Fig. S7†).

To investigate the effects of substrate binding in the presence of 2OG, AspH was crystallized in the presence of 2OG, Mn(n), and the synthetic hFX-EGFD186-124Ser substrate (ESI Fig. S1†), which mimics the EGFD1 of the reported AspH substrate human coagulation factor X (hFX). The structure was solved by molecular replacement using a reported structure (PDB ID: 5JTC) as a search model (P2₁2₁2₁, space group; 2.3 Å resolution). The active site region manifested electron density for both 2OG (Fig. 5b) and for the hFX-EGFD186-124Ser peptide (ESI Fig. S8†). Substrate binding to AspH affects the relative alignment of the oxygenase and TPR domains, i.e. the distance between the Cz atoms of Leu433 on TPR repeat z6 and Pro756 in the AspH C terminal region decreases from ~20 Å to ~14 Å upon substrate binding (ESI Fig. S9†). Evidence for an induced fit substrate binding mechanism involving this conformational change has been described when the cosubstrate 2OG was substituted for NOG. The significant conformational changes in the AspH oxygenase domain triggered by substrate binding (ESI Fig. S10†) do not affect the observed mode of 2OG binding in the active site (Fig. 5c); the binding modes of both 2OG and NOG are very similar, both in the presence or absence of substrate (ESI Fig. S9†). Interestingly, whilst in the 2OG complex structure the substrate residue Asp103hFX was observed in a single conformation whereas in the analogous NOG structure it was observed in two conformations (ESI Fig. S9†).

High resolution crystal structures of AspH complexed with 3-methyl-2OG (16) and Mn(n), both with and without the hFX-EGFD186-124Ser substrate bound, were obtained (AspH:16:hFX-EGFD186-124Ser, ESI Fig. S11†; AspH:16, ESI Fig. S12; 1.5 and 1.8 Å resolution, respectively). Substitution of 2OG for 16 did not trigger significant changes in the AspH conformations (superimposition of the AspH:2OG and the AspH:16 structures: Cz RMSD = 0.38 Å; superimposition of the AspH:2OG:hFX-EGFD186-124Ser and the AspH:16:hFX-EGFD186-124Ser structures: Cz RMSD = 0.23 Å; ESI Fig. S12 and S13†). A similar change in the relative alignment of the AspH oxygenase and TPR domains upon substrate binding was observed when 2OG was substituted for 16 (ESI Fig. S13†).

Although 3-methyl-2OG (16) was prepared and used as a racemic mixture, the electron density map corresponded to the, at least predominant, presence of the (R)-enantiomers at the active site in both the AspH:16 and AspH:16:hFX-EGFD186-124Ser structures (Fig. 5d and e; ESI Fig. S11 and S12†). The carboxylate groups of the (R)-enantiomer of 16 interact with the same AspH residues as 2OG in both the AspH:16 and AspH:16:hFX-EGFD186-124Ser structures. It is positioned to salt bridge with Arg733 (2.6/2.8 and 2.7/2.7 Å, respectively) with its C5-carboxylate and is positioned to interact with Ser668 (2.5 and 2.6 Å, respectively) through its C5-carboxylate, with His690 (2.8 and 2.8 Å, respectively) and Arg688 (2.8 and 2.9 Å,
respectively) through its C1-carboxylate, and with Mn(ii) through its C1-carboxylate (2.2 and 2.1 Å, respectively) and C2-carbonyl groups (2.2 and 2.1 Å, respectively). Notably, superimposition of the AspH:16 structure (beige: His6-AspH315–758; lavender: Mn ion; lemon: carbon-backbone of 2OG; green: carbon-backbone of hFX-EGFD186–124-4Ser peptide (ESI Fig. S1c†); red: oxygen; blue: nitrogen; w: water. (a) Representative OMIT electron density map (mFo-DFc) contoured to 5σ around 2OG of the AspH:2OG structure; (b) representative OMIT electron density map (mFo-DFc) contoured to 5σ around 2OG of the AspH:2OG:hFX-EGFD186–124-4Ser structure. The 2OG C5-carboxylate is positioned to salt bridge with the Arg735 side chain (2.5 and 2.9 Å) and to interact with Ser668 (2.5 Å). 2OG is positioned to interact with His690 (2.9 Å) and Arg688 (2.8 Å) through its C1-carboxylate, and with the Mn ion via its C1-carboxylate (2.2 Å) and C2-carbonyl (2.3 Å). The Mn ion is also complexed by His679 (2.2 Å), His725 (2.1 Å), a water molecule (2.1 Å), and the C4-carboxylate of the Asp103hFX peptide substrate (2.4 Å); (c) superimposition of views from the AspH:2OG:hFX-EGFD186–124-4Ser structure with one from the AspH:2OG structure (beige: His6-AspH315–758; lavender: Mn ion; lemon: carbon-backbone of 2OG); (d) representative OMIT electron density map (mFo-DFc) contoured to 5σ around (R)-16 of the AspH:16 structure; (e) representative OMIT electron density map (mFo-DFc) contoured to 5σ around (R)-16 of the AspH:16:hFX-EGFD186–124-4Ser structure; (f) superimposition of views from the AspH:16:hFX-EGFD186–124-4Ser structure with one from the AspH:2OG:hFX-EGFD186–124-4Ser structure (pale green: His6-AspH315–758; green: carbon-backbone of 2OG; lavender: Mn ion; aquamarine: carbon-backbone of hFX-EGFD186–124-4Ser).
also be noted that the (S)- and (R)-enantiomers of 16 will interconvert in aqueous media and in the presence of metal ions (ESI† Sections 4 and 5) and that, as (R)-16 is consumed in assays, this interconversion may become rate-limiting.

Next, AspH was crystallized in the presence of the inhibitor 3-ethyl-2OG (17); a crystal structure of AspH complexed with 17, Mn(II), and the hFX-EGFD186-124-4Ser substrate was obtained to 1.8 Å resolution (AspH:17:hFX-EGFD186-124-4Ser, ESI Fig. S14†). As with 16, the results suggest that 17 competes with 2OG for binding the AspH active site, in agreement with inhibition assays with varied 2OG concentrations (Fig. 3a and ESI Table S1†). As for 16, the observed electron density corresponded to the presence of the (R)-enantiomer of 17 (Fig. 6a and ESI Fig. S14†), even though 17 was used as a racemic mixture during crystallization. The (R)-enantiomers of 17 and 3-methyl-2OG (16) occupy similar conformations when bound to the AspH active site (Fig. 6c), with their C3-substituents facing towards the side chains of Val676 and Val727. Superimposition of the AspH:17:hFX-EGFD186-124-4Ser, the AspH:2OG:hFX-EGFD186-124-4Ser, and the AspH:16:hFX-EGFD186-124-4Ser structures reveals no significant changes in the AspH and the hFX-EGFD186-124-4Ser substrate conformations (ESI Fig. S15†), and the presence of the different 2OG analogues does not appear to affect the conformations of AspH active site side-chain residues (Fig. 6b and c).

![Diagram of binding of 3-ethyl-2OG (17) and 4,4-dimethyl-2OG (34) at the AspH active site.](image)
We also crystallized AspH in the presence of 4,4-dimethyl-2OG (34), Mn(ii), and the hFX-EGFD166-124Ser substrate (2.3 Å resolution; ESI Fig. S16†). In this AspH:34:hFX-EGFD166-124Ser structure, clear electron density for 34 was observed in the AspH active site (Fig. 6d). The pro-R methyl group of 34 projects towards one face of the hydrophobic pocket (Val626, Val727, Met670), whereas the pro-S methyl group of 34 is in proximity of Met670 and faces towards the indole ring of Trp625 (Fig. 6d and ESI Fig. S16c†).

While the AspH and the hFX-EGFD166-124Ser substrate conformations of the AspH:34:hFX-EGFD166-124Ser structure do not differ from those of previous structures (ESI Fig. S17†), the conformations of 34, of (R)-3-ethyl-2OG (17), and of N-oxallylaz-methylalanine (19), which is a derivative of 34, differ substantially in the crystalline state (Fig. 6f, ESI Fig. S18 and S19†). By contrast, the conformations of 2OG and 34 are very similar (Fig. 6e); we therefore investigated by NMR if 34 is converted into 2,2-dimethylsuccinate, in the presence of substantially higher AspH concentrations compared to the original SPE-MS assay conditions (~100 times more AspH). The results reveal that 34 indeed undergoes slow AspH-catalyzed oxidative decarboxylation, whereas another AspH inhibitor, 4-benzyl-2OG (32), was apparently not turned over under the same reaction conditions (ESI Fig. S20†).

Thus, the combined crystallographic analyses reveal that the C3/C4-substituted 2OG derivatives bind AspH in the same general manner as 2OG, with near identical binding modes for the oxalyl-groups and the C5-carboxylates. The different conformations for the C3- and C4-methylenes observed, however, do not correlate with catalysis versus inhibition (assuming the crystallographic binding modes reflect those in solution). Thus, whilst the C3- and C4-methylenes of (R)-16 and (R)-17 adopt a very similar conformation, which is distinct from that of 2OG and 34, 16 is a cosubstrate, whereas 17 is an inhibitor.

Hydrophobic interactions made by the C3-methyl and C3-ethyl groups of (R)-16 and (R)-17, respectively, and of one of the methyl groups of 34 with AspH, are consistent with the tighter binding of 16 as compared with 2OG and judged by the kcat values (Table 2). However, the kcat for 16 is approximately half that of 2OG whereas 17 and 34 are inhibitors and/or poor substrates, respectively (Table 2); the reason for these differences is uncertain, but it may reflect slower binding of O2, or a slower subsequent step during catalysis, e.g. reaction of the ferryl intermediate proposed to be present in the 2OG oxygenase catalytic cycle or release of the succinate coproduct (ESI Fig. S21†). Given the reduced kcat for 16, it is reasonable to propose that the kcat for the C3-ethyl substituted inhibitor 17 will be reduced even further, potentially approaching zero, potentially as a consequence of particular strong interaction with the hydrophobic pocket.

**Discussion**

The use of C3/C4-substituted 2OG derivatives has been a productive strategy to investigate *inter alia* the function, mechanism, and inhibition of aminotransferases and dehydrogenases that employ 2OG as a substrate or cosubstrate, but has been employed to a much lesser extent with 2OG oxygenases. By contrast with 2OG derivatives bearing major structural modifications of the 2-oxo-1,5-dicarboxylic acid scaffold, the synthetic accessibility of C3/C4-substituted 2OG derivatives has been hitherto limited, a factor which might have hampered detailed biochemical studies on their effects on 2OG oxygenases. To address this need, we developed an efficient synthesis of C3/C4-substituted 2OG derivatives relying on the use of cyanosulfur ylids as key intermediates (Scheme 1). Our synthesis compares favorably to reported syntheses of 2OG derivatives (Fig. 2); it is scalable, affords the 2OG derivatives and their synthetic precursors in high purity suitable for biochemical applications, and avoids the use of strong bases, acids, and oxidants. The broad substrate scope of the synthesis reflects the mild reaction conditions; for example, 2OG derivatives bearing oxidation-prone olefins and thiophenes were readily synthesized (ESI Table S1†).

Many of the C3/C4-substituted 2OG derivatives synthesized inhibited human AspH by a mechanism involving competition with 2OG for binding the active site as revealed by inhibition assays performed at variable 2OG concentrations and crystallographic studies (Fig. 3 and 6). In general, the C4-substituted 2OG derivatives were more efficient AspH inhibitors than the C3-substituted 2OG derivatives (Table 1). The C4-substituted 2OG derivatives were also more potent AspH inhibitors than the corresponding C4-substituted NOG derivatives (ESI Fig. S4†). NOG and other N-oxalyl amino acids are plant natural products and it is proposed that they may act as enzyme inhibitors in *vivo*. Our observations thus raise the possibility that naturally occurring 2-oxoacids may be biologically relevant modulators of the activities of 2OG oxygenases and related enzymes.

Our results indicate that some 2OG derivatives efficiently inhibit AspH at physiologically relevant 2OG levels which range from 9–12 μM 2OG in human plasma to >1 mM 2OG in cells. For example, 2OG derivative 33 inhibits AspH with 0.6 mM 2OG in the assay (IC50 ~ 5.6 μM; ESI Table S1†). 2OG competitive AspH inhibitors, such as 33, 34, or optimized variants of them, might be useful from a therapeutic perspective, because their inhibitory effect is unlikely to be compromised by mutations. Indeed an AspH mutation associated with Traboulsi syndrome
occurs in the 2OG binding site (i.e. R735W) and is likely
inactivating.28

At high enzyme concentrations, AspH converts some of the
inhibitors (e.g. 34) slowly into the corresponding succinate
derivatives (ESI Fig. S20†), indicating that the hydrophobic
interactions of the 2OG C3/C4-substituents with AspH stabilize
the AspH:2OG derivative complexes and/or reduce the rotary
flexibility of the cosubstrate necessary to enable its
oxidative decarboxylation. This observation is interesting
because in some contexts, e.g. the inhibition of HIF-α prolyl
hydroxylases, compounds that do not completely block activity
even when present in excess, including poor cosubstrates, may
actually be desirable, as they may help avoid overdose.27

C3/C4-substituted 2OG derivatives bear the potential to be
used as small-molecule probes26 in cells or in vivo to modulate
the catalytic activity of 2OG oxygenases, provided selective
interaction can be achieved. In this regard, the 2OG derivatives
might, at least partially, be selective AspH inhibitors consid-
ering that C4-substituted 2OG derivatives were reported to not
inhibit wild-type KDM4A,25 which is a 2OG dependent histone
demethylase.26 Note that, NOG derivatives have been used in vitro
and in cellular experiments to modulate the catalytic activities of FIH24 as well as of the KDM4 29 and ten-eleven-
transelocation (TET) enzymes28 with some selectivity. Our
results indicate that C4-substituted 2OG derivatives are more
efficient in inhibiting AspH than the corresponding C4-
substituted NOG derivatives (ESI Fig. S4†), as opposed to human FIH which also catalyzes the hydroxylation of Asp-
and Asn-residues and for which NOG derivatives were more potent
inhibitors.24 With regard to enzymes other than 2OG oxy-
genases, the AspH inhibitor 4,4-dimethyl-2OG (34), for example,
does not efficiently substitute for 2OG in glutamic oxaloacetic
aminotransferase catalyzed transamination reactions suggesting
that it might not interfere with the catalytic activities of human aminotransferases.41

The lysine metabolite 2-oxoadipate, which is based on a C6
rather than a C5 carbon skeleton as in 2OG, is capable of acting
as a relatively poor cosubstrate for procollagen prolyl hydroxy-
lases,42 phytanoyl-CoA 2-hydroxylase (PAHX),43 and for a bacte-
rial ethylene-forming 2OG dependent enzyme.44 However, to
our knowledge, C3/C4-substituted 2OG derivatives have so far
not been reported to substitute for 2OG as a cosubstrate for
wild-type 2OG oxygenases. The observation that several of our
synthetic 2OG derivatives efficiently substituted for 2OG as an
AspH cosubstrate, with 2OG derivatives 16 and 41 being the most
efficient (ESI Table S2†), is therefore of more general
interest.

The observation that 4-carboxyphenylglyoxylic acid (41) can
promote turnover of a 2OG oxygenase is remarkable because of
its cyclic aromatic scaffold. 41 is thus a promising candidate to
modulate AspH activity in vivo, as it might display selectivity for
AspH over other 2OG oxygenases because of its distinctive
structure. The oxidative decarboxylation of 41 by AspH is
reminiscent of the reaction catalyzed by 4-hydroxyphenyl pyru-
vate dioxygenase (HPPD),45 which is from a different class of Fe(n)-dependent oxygenases. A regioisomer of 41, 3-carbox-
yphenylglyoxylic acid (40), is a plant metabolite;46 thus, it is
possible that 41 or related aromatic compounds may modulate the catalytic activity of 2OG oxygenases in vivo.

The k_{cat}/K_m-value of AspH for 3-methyl-2OG (16) is approxi-
mately three times higher than for 2OG, suggesting the feasi-
bility of 2OG derivatives to selectively enhance AspH (or indeed other 2OG oxygenase) catalysis in cells or on the cell surface of
cancer cells in the presence of 2OG (Table 2). 16 is of particular
interest because it is a reported ingredient of human nutrition
(i.e. it is present in honey);47 thus, 16 might modulate the
activity of AspH and potentially other 2OG oxygenases in
humans. 2-Methylsuccinate, which is formed by the AspH-
catalyzed oxidative decarboxylation from 16 (ESI Fig. S5†), has
been detected in human urine48 and is inter alia used as a
biomarker for metabolic diseases as it is a product of other
metabolic pathways (i.e. isoleucine catabolism);49 this, however,
does not rule out the possibility that some 2-methylsuccinate
might originate from the oxidative decarboxylation of 16 cata-
yzed by 2OG oxygenases. 16 is also a proposed precursor of 3-
methyl glutamate, which is incorporated in natural products
such as polyethanamide A and B50 and daptomycin.51 16 is likely
biosynthesized in microorganisms through the direct reaction
of 2OG and SAM,52 suggesting that it might also be bio-
synthesized by animals in a similar manner. The results thus
raise the possibility that 16 or other C3/C4-substituted 2OG
derivatives, including their corresponding glutamate derivatives,
are human/animal metabolites and/or bioavailable through
nutrition or the gut microbiome.

In comparison with 16, its isomer 4-methyl-2OG (26), which is
a reported metabolite in plants53 and ingredient of wine,54 is
substantially less efficient in substituting for 2OG as a cosub-
strate (ESI Table S2†), highlighting that the position of the 2OG
substituent determines the ability of the 2OG derivatives to
serve as an alternative AspH cosubstrate.

AspH was co-crystallized in the presence of 3-methyl-2OG
(16) affording structures with the highest resolution reported
for AspH so far (1.5 and 1.8 Å; ESI Fig. S11 and S12†). The
crystallographic studies revealed that 16 substitutes for, and
binds similarly to, 2OG in the AspH active site (Fig. 5). However, the C3- and C4-methylene of 16 occupy different conforma-
tions orienting the 2OG C3-methyl-substituent towards Val676
and Val727, so enabling hydrophobic interactions. Superim-
position of 16 and 3-ethyl-2OG (17), which is a potent AspH
inhibitor, reveals that both 2OG derivatives adopt a similar
conformation when bound to AspH (Fig. 6c). Despite the use of
racemic 16 and 17 for crystallizations, in both cases the (3R)-
 enantiomers were observed by crystallography, with the C3-alkyl
substituents interacting with one face of a hydrophobic pocket
(ESI Fig. S16c†). Thus, subtle changes in the structure of the
2OG derivative can have a pronounced effect on AspH catalysis,
resulting in either efficient substrate hydroxylation or efficient
AspH inhibition. The exact factors that determine whether
a particular 2OG derivative inhibits or enables AspH catalysis
are unknown, but might relate to differences in oxygen binding,
modulation of steps after oxygen binding or to the stability of
the AspH:succinate derivative complexes (ESI Fig. S21†).

The potential of compensating catalytically inactivating
(with 2OG as a cosubstrate) mutants of 2OG oxygenases with 2-
oxoacids has been exemplified in the case of PAHX.⁵⁴ Our results thus raise to possibility that naturally occurring 2OG derivatives/analogues may modulate the activity of 2OG oxygenases and related enzymes (e.g. HPPD)⁵⁵, either by acting as inhibitors or by replacing 2OG in catalysis. Indeed, it is possible that some of the 2OG oxygenases without assigned biochemical functions (e.g. Jumonji C domain-containing protein 1c, JMJD1C)⁵⁶ or with unusual Fe(II) binding site geometries (e.g. PHD finger protein 2 (PHF2)⁵⁷ and hairless)⁵⁸ use 2-oxoacids cosubstrates other than 2OG. Future work will focus on exploring these possibilities using the 2OG derivatives described here and others prepared by the cyanosulfur ylid methodology.

Conclusions

A user-friendly efficient synthesis of C3/C4-substituted 2OG derivatives based on the use of cyanosulfur ylids was developed and employed to afford a diverse set of 2OG derivatives for detailed biochemical and structural investigations on the cosubstrate selectivity of the human 2OG oxygenase AspH. The overall results reveal that C3/C4-substituted 2OG derivatives can have profound effects on AspH catalysis and, by implication, likely other 2OG oxygenases. Simple alkyl substituents, e.g. methyl at the 2OG C-3 or C-4 position, enables retention of productive catalysis, likely in a stereoselective manner. By contrast, e.g. 2OG C4-dimethylation leads to inhibition/low levels of cosubstrate activity. The use of 2OG derivatives may thus inform on both the catalytic mechanisms and biological roles of AspH and other 2OG oxygenases, and aid in the development of new types of small-molecules that modulate 2OG oxygenase activity.

Data availability

Crystal structure data for His₆-AspH₁₁₅–₇₅₆ complexed to Mn, 2OG or a 2OG derivative (3-methyl-2OG, 16; 3-ethyl-2OG, 17; 4,4-dimethyl-2OG, 34; N-oxyl-α-methylalanine, 49), and, in some cases, substrate peptide (hFX-EGFD₁₈₆–₁₂₄-4Ser) are deposited in the protein data bank with PDB accession codes: 6YYU (AspH:2OG), 6YYW (AspH:2OG:hFX-EGFD₁₈₆–₁₂₄-4Ser), 6YYV (AspH:16), 6YYX (AspH:16:hFX-EGFD₁₈₆–₁₂₄-4Ser), 6YYY (AspH:34:hFX-EGFD₁₈₆–₁₂₄-4Ser), 6Z6Q (AspH:17:hFX-EGFD₁₈₆–₁₂₄-4Ser), and 6Z6R (AspH:49:hFX-EGFD₁₈₆–₁₂₄-4Ser).

Author contributions

L. B. synthesized the 2OG derivatives and the AspH substrate, produced recombinant human AspH, and performed AspH assays and crystallizations. Y. N. solved and refined the AspH crystal structures. All authors analyzed data. L. B. and C. J. S. wrote the manuscript with help from Y. N.

Conflicts of interest

There are no conflicts to declare.

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