



Cite this: *Org. Biomol. Chem.*, 2023, **21**, 4382

Received 24th April 2023,

Accepted 10th May 2023

DOI: 10.1039/d3ob00641g

rsc.li/obc

Total synthesis and biological evaluation of histone deacetylase inhibitor WF-3161†

Michael Kohr, ^a Niklas Papenhardt, ^b Manfred Jung ^b and Uli Kazmaier ^{*,a}

A novel synthesis of the naturally occurring HDAC inhibitor WF-3161 is described. Key steps include the Matteson homologation to generate the stereogenic centres in the side chain, and Pd-catalysed C–H functionalisation to connect the side chain to the peptide backbone. WF-3161 was found to be highly selective for HDAC1, whereas no activity was observed towards HDAC6. High activity was also found against the cancer cell line HL-60.

Introduction

Pharmaceuticals that target histone deacetylase (HDAC) enzymes have gained much attention in recent decades.¹ The overexpression of HDACs correlates with numerous prevalent diseases, such as Alzheimer's disease² and cancer,³ which makes these enzymes potential targets for drug development. Several acyclic HDAC inhibitors (HDACi), such as the naturally occurring trichostatin A (TSA)⁴ or the synthetic compound vorinostat (SAHA), have already been evaluated in clinical trials, with vorinostat the first HDACi to be approved by the FDA in the United States.⁵ More complex HDACi have been isolated from fungi, including chlamydocin,⁶ Cyl-1 and Cyl-2 (PMP: *p*-methoxyphenyl),⁷ the trapoxins,⁸ and WF-3161,⁹ among others (Fig. 1). All of these compounds have three major functional groups in common: (a) a zinc-binding motif, *e.g.* a hydroxamic acid or an epoxyketone; (b) a linker (spacer), which simulates the lysine side chain of the natural substrate; and (c) a cap region, which interacts with the protein surface surrounding the active site.¹⁰

Prototypical HDAC inhibitors, such as the natural product trichostatin A (TSA) or the approved drug vorinostat (SAHA),

lack HDAC isoform selectivity, which leads to undesirable interactions with the 'wrong' isoforms (Tables 1 and 2).¹¹ In contrast, macrocyclic HDACi contain a more complex cap region and are capable of differentiating among HDAC isoforms, enabling the investigation of their individual functions and association with cancer progression.¹² Macrocyclic HDACi containing (2*S*,9*S*)-2-amino-9,10-epoxy-8-oxodecanoic acid (Aeo), an amino acid bearing an epoxyketone group in the side chain, highly differentiate between HDAC1 and HDAC6, for example, which belong to different HDAC classes (Table 1).^{13a} The presence of the 8-ketogroup of Aeo has been proven to be essential for the interaction of HDACi with the protein's active site. Replacements of the keto-group or its reduction lead to a significantly lower bioactivity, as shown by Kim *et al.*^{13b}

While these results provide deep insights into the behaviour of the HDACi investigated, to the best of our knowledge, other important representatives of this class of compounds,

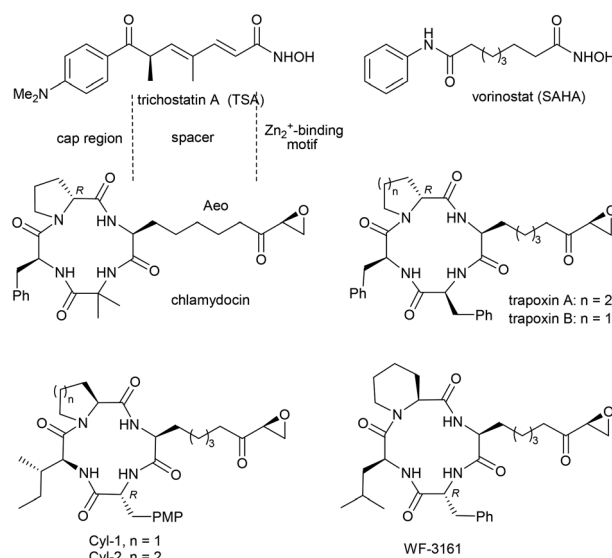


Fig. 1 Natural and synthetic HDAC inhibitors.

^aOrganic Chemistry, Saarland University, D-66123 Saarbrücken, Germany.

E-mail: u.kazmaier@mx.uni-saarland.de

^bInstitute of Pharmaceutical Sciences, University of Freiburg, Albertstr. 25, 79104 Freiburg, Germany

†Electronic supplementary information (ESI) available: Experimental details, compound characterization, copies of ¹H and ¹³C NMR spectra, HPLC chromatograms and MS² analysis. See DOI: <https://doi.org/10.1039/d3ob00641g>



Table 1 IC₅₀ of natural HDACi against HDAC1 and HDAC6^{11a}

HDACi	IC ₅₀ , [nM] HDAC1	IC ₅₀ , [nM] HDAC6	HDAC6/HDAC1
TSA	6.0 ± 2.5	8.6 ± 1.4	1.4
Trapoxin A	0.82 ± 0.29	524 ± 240	640
Trapoxin B	0.11 ± 0.01	360 ± 160	3300
Chlamydocin	0.15 ± 0.03	1100 ± 430	7300
Cyl-2	0.70 ± 0.45	40 000 ± 11 000	57 000

such as WF-3161 and Cyl-1, have not been thoroughly investigated to date.

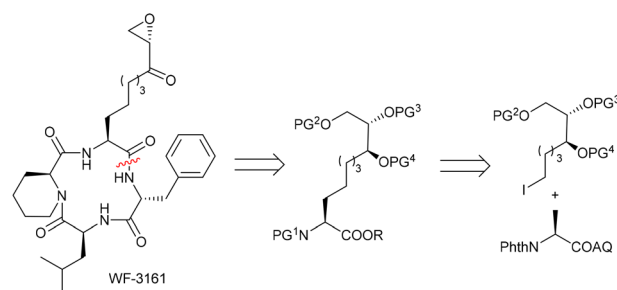
The interesting biological activities of these tetrapeptide-based HDACi and their derivatives have initiated wide-ranging efforts towards a synthesis of these compounds.¹⁴ A key challenge in this approach is the stereoselective synthesis of the unusual amino acid Aeo, which contains an epoxyketone as a warhead and a zinc-binding motif. Several synthetic approaches have been developed in recent years, including alkylation of imino glycinate,¹⁵ phosphonate condensation of glycine derivatives,¹⁶ and the use of enantiomerically pure amino acid precursors,¹⁷ among others. Since our group is also involved in the synthesis of unusual amino acids¹⁸ and peptidic natural products,¹⁹ we became interested in the synthesis of this type of HDACi and began to develop several independent routes towards protected Aeo-precursors. An asymmetric chelate-Claisen rearrangement²⁰ was the key step in the synthesis of chlamydocin²¹ and Cyl-1,²² while a Pd-catalysed peptide allylation²³ gave access to trapoxin A.²⁴ We subsequently became interested in developing a third independent route based on C–H functionalisation, with the goal of obtaining access to WF-3161. This natural product was first isolated from a strain of fungus, *Petriella guttulata*, by Umehara *et al.*, who amassed a large body of structural and biological data on its activity, including anti-cancer activity against blood cancer cells in mice and antifungal activity against *Trichophyton asteroides*.^{9a} In 2008, Proksch *et al.* obtained similar results with lymphoma cells from mice.²⁵ However, no investigations on the HDAC activity of WF-3161 have been carried out to date, although the first synthesis of this compound was already reported in 1989 by Schmidt *et al.*²⁶

Transition metal-catalysed C–H functionalisation has become an extremely popular tool in modern organic synthesis,²⁷ and its widespread applications have been described in a series of recent reviews.²⁸ Pd-catalysed functionalisation of amino acids and peptides, in particular, should offer an ideal method of synthesising and modifying peptidic natural products.²⁹ Suitable directing groups (DGs) are required to address a specific C_{sp³}–H bond in an amino acid in the presence of others.³⁰ This protocol is also suitable for the modification of peptides, as long as no acidic N–H-bond is present in the reacting amino acid.³¹ To date, the best results have been obtained with amides of 8-aminoquinoline (AQ)³² or 2-(methylthio)aniline (MTA),³³ with both directing groups developed by Daugulis *et al.*³⁴ In general, (het)aryliodides or -bro-

mides are employed for the functionalisation of cyclic or *N*-methylated amino acids at the β-position, whereas the introduction of alkyl groups is relatively rare. We were therefore inspired to determine if we could introduce a protected, highly functionalised side chain in a single step into a suitably protected alanine derivative, *e.g.* an *N*-phthaloylated aminoquinoline ester (Scheme 1). This particular approach is highly attractive, as it should enable the introduction of various side chains as potential zinc-binding motifs, regardless of the stereogenic centre at the α-position, as long as it does not epimerise under the reaction conditions used.

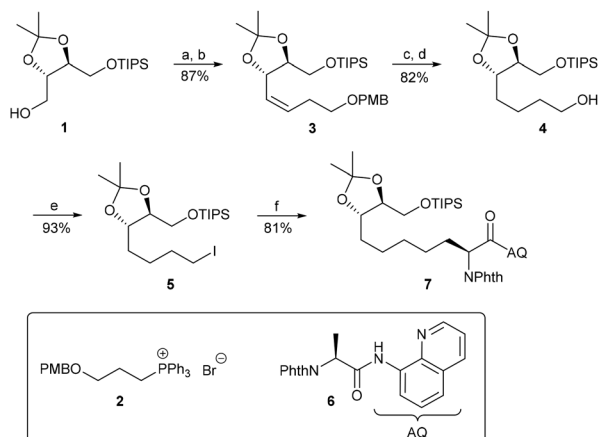
Results and discussion

Based on the total synthesis of WF-3161 by Schmidt *et al.*, who carefully investigated the four different macrocyclisation positions, we chose to focus on the position between Aeo and *D*-phenylalanine as the most promising. Before developing an entirely new route, we initially decided to verify the applicability of C–H functionalisation in this presumably challenging approach. Therefore, we synthesised iodide **5** in a similar manner as described by Schreiber *et al.*³⁵ In this approach, we started with threitol-derived triisopropylsilyl (TIPS)-ether **1** (Scheme 2), which we previously used as a starting material in Pd-catalysed allylic alkylations.³⁶ Swern oxidation and subsequent Wittig olefination using the *p*-methoxybenzyl (PMB)-protected phosphonium salt **2**³⁷ provided (*Z*)-alkene **3** as a single stereoisomer. The olefin geometry does not play a role in this case, because the double bond was hydrogenated before the PMB was removed. It should be noted, however, that the order of the reaction steps is important, because hydrogenation after PMB-cleavage preferentially affords the deoxygenated product. The primary alcohol **4** was converted into iodide **5** *via* the Appel reaction.³⁸ With the required building block in hand, we subsequently investigated the key step of our synthesis: the C–H functionalisation of *N*-phthaloyl-protected alanine 8-aminoquinoline ester **6**.³⁹ The first experiments were carried out in *t*-butanol/dichloroethane (1:1), which afforded the desired product **7** in an acceptable yield of 44%. As a major side product, the direct substitution of the iodine by *t*-butanol was observed. Switching to the more steri-



Scheme 1 Planned synthesis of protected Aeo *via* C–H-functionalization.

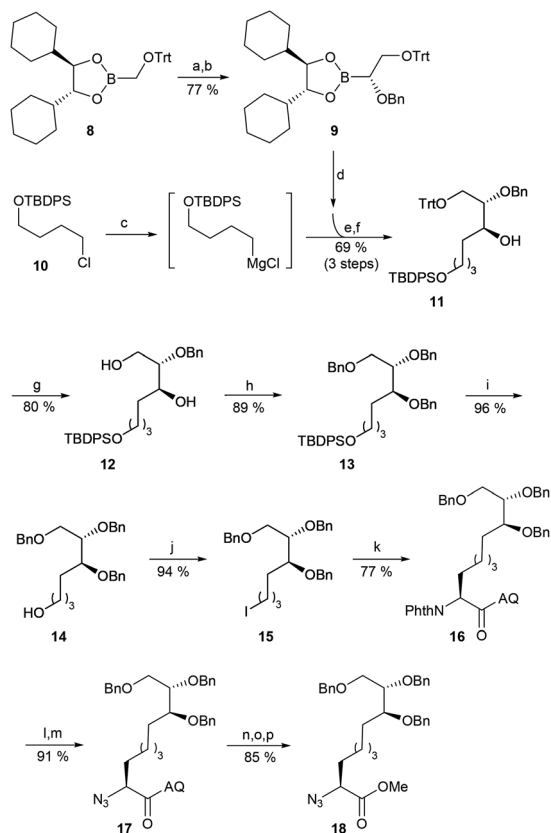




Scheme 2 Synthesis of an Ae-precursor **7** from threitol derivative **1** via C–H functionalization. Reagents and conditions: (a) $(\text{COCl})_2$ (1.6 equiv.), DMSO (3.0 equiv.), NEt_3 (5.0 equiv.), CH_2Cl_2 , 0 °C, 1 h. (b) KHMDS (1.1 equiv.), Wittig salt **2** (1.2 equiv.), THF, 0 °C, 75 min (c) H_2 (1 atm), Pd/C, MeOH, rt, 16 h. (d) DDQ (1.1 equiv.), CH_2Cl_2 : H_2O (20 : 1), rt, 2 h. (e) PPh_3 (1.25 equiv.), imidazole (1.25 equiv.), I_2 (1.24 equiv.), CH_2Cl_2 , 1. 0 °C, 25 min; 2. rt, 3.5 h. (f) **6**, Ag_2CO_3 (0.8 equiv.), $\text{Pd}(\text{OAc})_2$ (10 mol-%), dibenzyl phosphate (30 mol%), *tert*-amyl-OH : $\text{C}_2\text{H}_4\text{Cl}_2$ (1 : 1), 60 °C, 24 h.

cally hindered *t*-amylalcohol solved this problem and afforded **7** in excellent yield for an alkyl coupling.

Following the successful optimisation of the C–H functionalisation step, we developed a new synthetic approach by employing the Matteson homologation for the generation of the stereogenic centres in the side chain (Scheme 3).⁴⁰ We recently utilised this highly flexible concept during the total synthesis of several natural products.⁴¹ Because we were unable to homologate benzyloxy-substituted methyl boronic esters, we started our synthesis with the known *O*-trityl-protected boronic ester **8**,⁴² which had already been used by Matteson for a similar purpose (Scheme 3).⁴³ Deprotonated benzyl alcohol was chosen as a nucleophile under standard reaction conditions, which afforded boronic ester **9** in good yield and in enantiomerically pure form. The second stereogenic centre and the linear spacer unit were generated by converting the *O*-silylated alkyl halide **10** into the corresponding Grignard reagent, which was subsequently used as a nucleophile in the second homologation step. The prolonged boronic ester formed was directly oxidised to the corresponding alcohol **11**, which was obtained as a single stereoisomer. In the next step, we aimed to remove the trityl protecting group and protect the two OH-functionalities as benzyl esters, which would allow us to remove all protecting groups at once as the final step in the synthesis. Although, we had already replaced the previously used TIPS group with the more acid-stable *tert*-butyldiphenylsilyl (TBDPS) group, the selective cleavage of the trityl group was far from trivial. With HCl in dioxane or Amberlyst 15, partial or complete cleavage of the silyl protecting group was observed. With acetic acid, we were able to cleave the trityl group selectively, but by far the best



Scheme 3 Synthesis of an Ae-precursor **18** via Matteson homologation and C–H functionalization. Reagents and conditions: (a) (1) CH_2Cl_2 (3.0 equiv.), LDA (1.35 equiv.), –40 °C; (2) ZnCl_2 (3.0 equiv.), THF, rt. (b) BnOH (2.3 equiv.), NaH (2.0 equiv.), THF/DMSO, 0 °C to rt, 16 h. (c) Mg (2.7 equiv.), 1,2-dibromoethane (13 mol%), THF, Δ , 2 h. (d) (1) CH_2Cl_2 (3.0 equiv.), LDA (1.35 equiv.), –40 °C; (2) ZnCl_2 (4.0 equiv.), THF, rt. (e) 0 °C to rt, 16 h. (f) NaOH (2.25 equiv.), H_2O_2 (2.25 equiv.), THF, rt, 90 min (g) ZnBr_2 (4.0 equiv.), H_2O (4.0 equiv.), CH_2Cl_2 , rt, 3.5 h. (h) KOtBu (3.0 equiv.), BnBr (4.0 equiv.), THF, 0 °C, 100 min (i) TBAF (1.25 equiv.), THF, rt, 16 h. (j) PPh_3 (1.25 equiv.), imidazole (1.25 equiv.), I_2 (1.24 equiv.), CH_2Cl_2 , rt, 3.5 h. (k) **6**, Ag_2CO_3 (0.8 equiv.), $\text{Pd}(\text{OAc})_2$ (10 mol-%), dibenzyl phosphate (30 mol-%), *tert*-amyl-OH : $\text{C}_2\text{H}_4\text{Cl}_2$ (1 : 1), 60 °C, 22 h. (l) hydrazine (1.2 equiv.), DIPEA (1.0 equiv.), EtOH, 80 °C, 20 h. (m) TfN_3 (3.1 equiv.) in toluene, NaHCO_3 (4.0 equiv.), $\text{CSO}_4 \cdot 5 \text{H}_2\text{O}$ (4.3 mol%), MeOH/ H_2O , rt, 2 h. (n) Boc_2O (4.0 equiv.), DMAP (60 mol%), MeCN, rt, 16 h. (o) LiOH (2.0 equiv.), H_2O_2 (5.0 equiv.), THF, 0 °C, 1 h. (p) K_2CO_3 (2.5 equiv.), MeI (3.0 equiv.), DMF, 0 °C to rt, 3 h.

results were obtained using anhydrous ZnBr_2 in the presence of exactly one equivalent of H_2O per equivalent ZnBr_2 . Under these conditions, only traces of the silyl cleavage product were observed and 7% of **11** could be recovered. The resulting diol **12** was double benzyl-protected (**13**) and following cleavage of the silyl protecting group, the terminal primary alcohol **14** was converted into the desired iodide **15** in an analogous manner to **5**.

The subsequent C–H functionalisation under the previously optimised conditions afforded comparable results as previously. Protected amino acid **16** was obtained in high yield and as a single stereoisomer. The phthaloyl protecting group was removed under standard conditions in excellent yield, and



the free amine was converted into the corresponding azide **17**. Subsequent cleavage of the aminoquinoline, followed by *O*-methylation, provided methylester **18**, which was used in the subsequent peptide couplings.

In our first attempt, we tried to couple our new amino acid with a preformed tripeptide; unfortunately, however, significant epimerisation in the peptide fragment was observed. Therefore, we decided to build up the peptide step by step (Scheme 4). Staudinger reduction of the azide of **18** afforded the free amine, which was coupled with Boc-protected pipecolic acid to **19**. Boc-cleavage under standard conditions, using HCl in dioxane, and coupling with Boc-Leu afforded the desired tripeptide **20** in an overall high yield. Interestingly, a partial peptide bond cleavage was observed when reacting tripeptide **20** with HCl in dioxane, as in the previous step. After some optimisations, however, high yields were achieved with TFA/triisopropylsilane (TIPS-H),⁴⁴ in which no decomposition was observed. Following peptide coupling to tetrapeptide **21** and saponification, the free carboxylic acid was obtained, and was first activated with pentafluorophenol in analogy to the Schmidt synthesis. Unfortunately, no synthetically useful yield was obtained from our attempts to cyclise this active ester. Instead, macrocyclisation with PyAOP and HOAt afforded the desired product **22** in high yields and without any epimerisation. The *O*-benzyl protecting groups were easily removed at ambient pressure in the presence of small amounts of HCl. Under these conditions, however, the formation of peptide cleavage products was an issue, as WF-3161 has a relatively high ring tension compared to other tetrapeptidic HDACi.⁴⁵ This problem was solved by carrying out the hydrogenation reaction under neutral conditions and at higher pressure. The triol obtained was subsequently converted into the corresponding epoxide, which was not a trivial step. Attempts to tosylate the primary alcohol led to low conversions and poor selectivity. In contrast, a selective mesylation at low tempera-

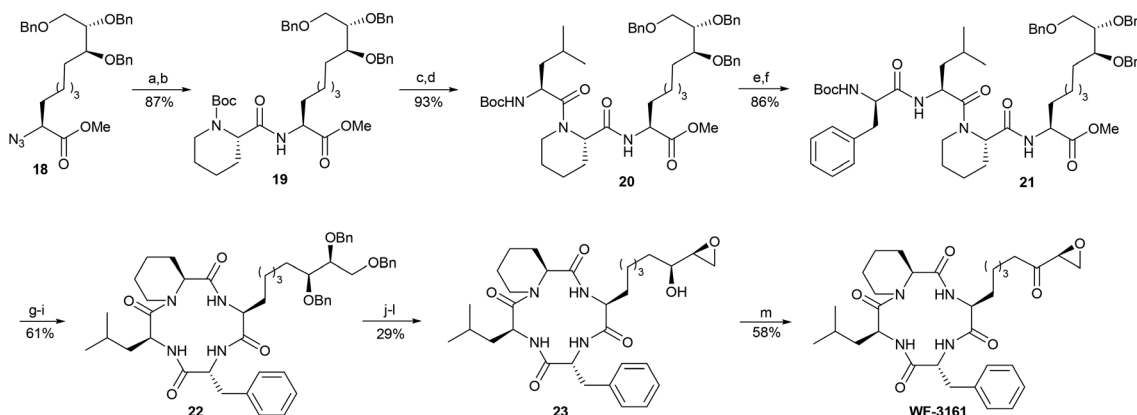
Table 2 IC₅₀ values (mean ± SD) detected by a biochemical *in vitro* assay for HDAC1 and HDAC6. n.i. = no inhibition. GI₅₀ values (mean ± SD) detected by the cell viability assay

HDACi	IC ₅₀ , [nM] HDAC1	IC ₅₀ , [nM] HDAC6	GI ₅₀ , [nM] HL-60
WF-3161	0.7 ± 0.1	n.i. @ 25 000	13 ± 1.1
Vorinostat	126 ± 7	58 ± 8	846 ± 48
Mocetinostat	110 ± 8	n.i.	592 ± 63

ture was possible. Full conversion was still not achieved, though it was possible to suppress double mesylation using 1.2 equiv. of mesyl chloride. Fortunately, unreacted triol could be recovered from the reaction mixture. The best results in the epoxide formation step were obtained using the non-nucleophilic base DBU to afford epoxy alcohol **23**, which was subsequently oxidised with Dess–Martin periodinane (DMP) to afford the desired WF-3161.

Next, WF-3161 was initially screened *in vitro* for HDAC inhibitory activity, followed by an assessment of its ability to reduce cancer cell viability. A fluorescence-based assay was used to determine the activity of WF-3161 on the class I nuclear isoforms HDAC1, as well as the class II cytoplasmatic isoform HDAC6. The data are presented as IC₅₀ values in Table 2. The already approved HDAC inhibitor vorinostat and the HDAC1 selective inhibitor mocetinostat were included as positive controls. WF-3161 inhibited HDAC activity in a dose-dependent manner, with IC₅₀ values of 0.7 nM ± 0.1 nM for HDAC1. Interestingly, no inhibition of HDAC6 was observed at 25 μM, suggesting a strong selectivity towards class I HDACs, similar to that observed with Cyl-2, but different from that of the trapoxins.

To further measure the activity of the synthesised WF-3161, we examined its growth-inhibiting effects on an acute myeloid leukaemia cell line, based on the MTS assay (Promega), with



Scheme 4 Synthesis of WF-3161. Reagents and conditions: (a) PPh₃ (3.0 equiv.), THF/H₂O, 50 °C, 3 h. (b) Boc-L-Pip-OH (1.1 equiv.), TBUTU (1.1 equiv.), DIPEA (1.2 equiv.), MeCN, 0 °C to rt, 16 h. (c) HCl/1,4-dioxane (10 equiv.), rt, 60 min (d) Boc-L-Leu-OH (1.2 equiv.), PyAOP (1.2 equiv.), DIPEA (2.5 equiv.), DMF, 0 °C to rt, 16 h. (e) TFA/TIPS-H/H₂O (92.5 : 5 : 2.5), CH₂Cl₂, 0 °C, 60 min (f) Boc-D-Phe-OH (1.1 equiv.), PyAOP (1.1 equiv.), DIPEA (2.5 equiv.), DMF, 0 °C to rt, 16 h. (g) LiOH (1.2 equiv.), 1,4-dioxane/H₂O, 0 °C to rt, 16 h. (h) TFA/TIPS-H/H₂O (92.5 : 5 : 2.5), CH₂Cl₂, 0 °C, 40 min (i) PyAOP (10 equiv.), HOAt (10 equiv.), DIPEA (10 equiv.), DMF, high dilution (1 mM), 0 °C to rt, 16 h. (j) H₂ (20 bar), Pd/C, MeOH, 16 h. (k) MsCl (1.2 equiv.), 2,4,6-collidine (10 equiv.), CH₂Cl₂, 0 °C, 22 h. (l) DBU (5.0 equiv.), MeOH, 0 °C, 5.5 h. (m) DMP (2.0 equiv.), CH₂Cl₂, rt, 2 h.

vorinostat and mocetinostat as reference controls. The results are shown in Table 2. As indicated by the data, the macrocyclic WF-3161, which contains an epoxyketone, inhibits the cell viability of HL-60 in the low nanomolar range, with a GI_{50} value of 13 ± 1.1 nM. The low GI_{50} value obtained is also in accordance with the low IC_{50} value observed in the *in vitro* tests.

The increased potency of WF-3161 compared to that of vorinostat or mocetinostat *in vitro* and in the MTS assay could be related to the epoxyketone as a zinc-binding group. Interestingly, the mechanism of epoxyketone inhibition of HDACs is not entirely clear. In 1993, M. Kijima and colleagues hypothesised that trapoxin A, which also contains the epoxyketone, might act as either a tight noncovalent binder or covalently bind to the HDAC.⁴⁶ This observation suggests that WF-3161 inhibits class I HDACs in a similar fashion and leads to increased potency compared to the hydroxamic acid and benzamide zinc-binding groups.

Conclusions

We have developed a novel synthesis for the HDAC inhibitor WF-3161, based on Matteson homologations and C–H functionalisation. The Matteson homologation was used to generate the stereogenic centre in the side chain and should be a suitable method for introducing other comparable zinc-binding motifs. The side chain was subsequently coupled to an alanine derivative *via* palladium-catalysed C–H-functionalisation. The WF-3161 obtained was screened *in vitro* for HDAC inhibitory activity and for its ability to reduce cancer-cell viability. Highly selective inhibition of HDAC1 compared to HDAC6 was observed, in addition to low nM activity towards the myeloid leukaemia cell line HL-60.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Financial support from Saarland University and the Deutsche Forschungsgemeinschaft (DFG, Bruker Neo 500-447298507 to U. K.; SFB 992 (Project-ID: 192904750) to M. J. is gratefully acknowledged.

References

- (a) H. Rajak, A. Singh, P. K. Dewangan, V. Patel, D. K. Jain, S. K. Tiwari, R. Veerasamy and P. C. Sharma, *Curr. Med. Chem.*, 2013, **20**, 1887–1903; (b) S. Tan and Z.-P. Liu, *ChemMedChem*, 2015, **10**, 441–450; (c) X. Peng, Z. Sun, P. Kuang and J. Chen, *Eur. J. Med. Chem.*, 2020, **208**, 112831; (d) T. C. S. Ho, A. H. Y. Chan and A. Ganesan, *J. Med. Chem.*, 2020, **63**, 12460–12484; (e) M. Daško, B. de Pascual-Teresa, I. Ortín and A. Ramos, *Molecules*, 2022, **27**, 715.
- M. Kilgore, C. A. Miller, D. M. Fass, K. M. Hennig, S. J. Haggarty, J. D. Sweatt and G. Rumbaugh, *Neuropsychopharmacology*, 2010, **35**, 870–880.
- (a) D. Marchion and P. Münster, *Expert Rev. Anticancer Ther.*, 2007, **7**, 583–598; (b) J. M. Wagner, B. Hackanson, M. Lübbert and M. Jung, *Clin. Epigenet.*, 2010, **1**, 117–136; (c) G. Shanmugam, S. Rakshit and K. Sarka, *Transl. Oncol.*, 2022, **16**, 101312.
- (a) M. Yoshida, M. Kijima, M. Akita and T. Beppu, *J. Biol. Chem.*, 1990, **265**, 17174–17179; (b) A. G. Kazantsev and L. M. Thompson, *Nat. Rev. Drug Discovery*, 2008, **7**, 854–868; (c) M. Jakovcevski and S. Akbarian, *Nat. Med.*, 2012, **18**, 1194–1204; (d) F. F. Wagner, M. Weiwer, M. C. Lewis and E. B. Holson, *Neurotherapeutics*, 2013, **10**, 589–604; (e) A. C. West and R. W. Johnstone, *J. Clin. Invest.*, 2014, **124**, 30–39.
- P. A. Marks and R. Breslow, *Nat. Biotechnol.*, 2007, **25**, 84–90.
- A. Closse and R. Huguenin, *Helv. Chim. Acta*, 1974, **57**, 533–545.
- (a) A. Hirota, A. Suzuki, H. Suzuki and S. Tamura, *Agric. Biol. Chem.*, 1973, **37**, 643–647; (b) S. Takayama, A. Isogai, M. Nakata, H. Suzuki and A. Suzuki, *Agric. Biol. Chem.*, 1984, **48**, 839–842.
- H. Itazaki, K. Nagashima, K. Sugita, H. Yoshida, Y. Kawamura, Y. Yasuda, K. Matsumoto, K. Ishii, N. Uotani, H. Nakai, A. Terui, S. Yoshimatsu, Y. Ikenishi and Y. Nakagawa, *J. Antibiot.*, 1990, **43**, 1524–1532.
- (a) K. Umehara, K. Nakahara, S. Kiyoto, M. Iwami, M. Okamoto, H. Tanaka, M. Kohsaka, H. Aoki and H. Imanaka, *J. Antibiot.*, 1983, **36**, 478–483; (b) M. Kawai, R. Pottorf and D. H. Rich, *J. Med. Chem.*, 1986, **29**, 2409–2411.
- M. S. Finnin, J. R. Donigian, A. Cohen, V. M. Richon, R. A. Rifkind, P. A. Marks, R. Breslow and N. P. Pavletich, *Nature*, 1999, **401**, 188–193.
- (a) R. Furumai, Y. Komatsu, N. Nishino, S. Khochbin, M. Yoshida and S. Horinouchi, *Proc. Natl. Acad. Sci. U. S. A.*, 2001, **98**, 87–92; (b) P. Gupta, R. C. Reid, A. Iyer, M. J. Sweet and D. P. Fairlie, *Curr. Top. Med. Chem.*, 2012, **12**, 1479–1499.
- K. Cheng, S. Li and C. Liao, *Curr. Med. Chem.*, 2017, **24**, 4166–4179.
- (a) A. V. Bieliauskas and M. K. H. Pflum, *Chem. Soc. Rev.*, 2008, **37**, 1402–1413; (b) S.-D. Kim, H. W. Knoche and L. D. Dunkle, *Physiol. Mol. Plant Pathol.*, 1987, **30**, 433–440.
- (a) T. L. Newkirk, A. A. Bowers and R. M. Williams, *Nat. Prod. Rep.*, 2009, **26**, 1293–1320; (b) A. R. Maolanon, H. M. E. Kristensen, L. J. Leman, M. R. Ghadiri and C. A. Olsen, *ChemBioChem*, 2017, **18**, 5–49.
- D. H. Rich, J. Singh and J. H. Gardner, *J. Org. Chem.*, 1983, **48**, 432–434.
- U. Schmidt, A. Lieberknecht, H. Griesser and F. Bartkowiak, *Angew. Chem., Int. Ed. Engl.*, 1984, **23**, 318–320.



- 17 (a) J. E. Baldwin, R. M. Adlington, C. R. A. Godfrey and V. K. Patel, *J. Chem. Soc., Chem. Commun.*, 1991, 1277; (b) J. E. Baldwin, R. M. Adlington, C. R. A. Godfrey and V. K. Patel, *Tetrahedron*, 1993, **49**, 7837–7856.
- 18 (a) U. Schmidt, A. Lieberknecht, U. Kazmaier, H. Griesser, G. Jung and J. Metzger, *Synthesis*, 1991, 49–55; (b) A. Krebs and U. Kazmaier, *Tetrahedron Lett.*, 1996, **37**, 7945–7946; (c) M. Pohlman and U. Kazmaier, *Org. Lett.*, 2003, **5**, 2631–2633.
- 19 (a) L. Karmann, K. Schulz, J. Herrmann, R. Müller and U. Kazmaier, *Angew. Chem., Int. Ed.*, 2015, **54**, 4502–4507; (b) P. Barbie and U. Kazmaier, *Org. Lett.*, 2016, **18**, 204–207; (c) J. Gorges and U. Kazmaier, *Org. Lett.*, 2018, **20**, 2033–2036; (d) M. Kohr, C. Walt, J. Dastbaz, R. Müller and U. Kazmaier, *Org. Biomol. Chem.*, 2022, **20**, 9609–9612.
- 20 U. Kazmaier, *Liebigs Ann./Recl.*, 1997, 285–295.
- 21 C. Quirin and U. Kazmaier, *Eur. J. Org. Chem.*, 2009, 371–377.
- 22 P. Servatius and U. Kazmaier, *Org. Biomol. Chem.*, 2018, **16**, 3464–3472.
- 23 (a) U. Kazmaier, J. Deska and A. Watzke, *Angew. Chem., Int. Ed.*, 2006, **45**, 4855–4858; (b) U. Kazmaier, *Org. Chem. Front.*, 2016, **3**, 1541–1560.
- 24 P. Servatius and U. Kazmaier, *J. Org. Chem.*, 2018, **83**, 11341–11349.
- 25 P. Proksch, R. Ebel, R. Edrada, F. Riebe, H. Liu, A. Diesel, M. Bayer, X. Li, W. H. Lin, V. Grebenyuk, W. E. G. Müller, S. Draeger, A. Zuccaro and B. Schulz, *Bot. Mar.*, 2008, **51**, 209–218.
- 26 U. Schmidt, U. Beutler and A. Lieberknecht, *Angew. Chem., Int. Ed. Engl.*, 1989, **28**, 333–335.
- 27 (a) A. R. Dick, K. L. Hull and M. S. Sanford, *J. Am. Chem. Soc.*, 2004, **126**, 2300–2301; (b) D. Shabashov and O. Daugulis, *Org. Lett.*, 2005, **7**, 3657–3659; (c) R. Giri, J. Liang, J. Lei, J. Li, D. Wang, X. Chen, I. C. Naggar, C. Guo, B. M. Yu and J. Foxman, *Angew. Chem., Int. Ed.*, 2005, **44**, 7420–7424; (d) S. Aspin, A. S. Goutierre, P. Larini, R. Jazzar and O. Baudoin, *Angew. Chem., Int. Ed.*, 2012, **51**, 10808–10811; (e) E. T. Nades, G. I. F. Santos, D. Shabashov and O. Daugulis, *J. Org. Chem.*, 2013, **78**, 9689–9714.
- 28 (a) J. Yamaguchi, A. D. Yamaguchi and K. Itami, *Angew. Chem., Int. Ed.*, 2012, **51**, 8960–9009; (b) A. F. M. Noisier and M. A. Brimble, *Chem. Rev.*, 2014, **114**, 8775–8806; (c) X. Lu, B. Xiao, R. Shang and L. Liu, *Chin. Chem. Lett.*, 2016, **27**, 305–311; (d) G. He, B. Wang, W. A. Nack and G. Chen, *Acc. Chem. Res.*, 2016, **49**, 635–645; (e) J. He, M. Wasa, K. S. L. Chan, Q. Shao and J. Yu, *Chem. Rev.*, 2017, **117**, 8754–8786; (f) S. Sengupta and G. Mehta, *Org. Biomol. Chem.*, 2020, **18**, 1851–1876.
- 29 (a) W. R. Gutekunst and P. S. Baran, *Chem. Soc. Rev.*, 2011, **40**, 1976–1991; (b) J. Yamaguchi, A. D. Yamaguchi and K. Itami, *Angew. Chem., Int. Ed.*, 2012, **51**, 8960–9009; (c) T. Cernak, K. D. Dykstra, S. Tyagarajan, P. Vachal and S. W. Krska, *Chem. Soc. Rev.*, 2016, **45**, 546–576; (d) R. R. Karimov and J. F. Hartwig, *Angew. Chem., Int. Ed.*, 2018, **57**, 4234–4241.
- 30 C. Sambaglio, D. Schönbauer, R. Blicke, T. Dao-Huy, G. Pototschnig, P. Schaaf, T. Wiesinger, M. F. Zia, J. Wencel-Delord, T. Besset, B. U. W. Maes and M. Schnürch, *Chem. Soc. Rev.*, 2018, **47**, 6603–6743.
- 31 (a) B. Mondal, B. Roy and U. Kazmaier, *J. Org. Chem.*, 2016, **81**, 11646–11655; (b) T. Kinsinger and U. Kazmaier, *Org. Lett.*, 2018, **20**, 7726–7730; (c) T. Kinsinger and U. Kazmaier, *Org. Biomol. Chem.*, 2019, **17**, 5595–5600; (d) M. Kohr and U. Kazmaier, *Org. Lett.*, 2021, **23**, 5947–5951.
- 32 (a) V. G. Zaitsev, D. Shabashov and O. Daugulis, *J. Am. Chem. Soc.*, 2005, **127**, 13154–13155; (b) L. D. Tran and O. Daugulis, *Angew. Chem., Int. Ed.*, 2012, **51**, 5188–5191.
- 33 D. Shabashov and O. Daugulis, *J. Am. Chem. Soc.*, 2010, **132**, 3965–3972.
- 34 (a) A. A. Kulkarni and O. Daugulis, *Synthesis*, 2009, 4087–4109; (b) J. Roane and L. D. Tran, *Acc. Chem. Res.*, 2015, **48**, 1053–1064.
- 35 J. Taunton, J. L. Collins and S. L. Schreiber, *J. Am. Chem. Soc.*, 1996, **118**, 10412–10422.
- 36 A. Horn and U. Kazmaier, *Org. Lett.*, 2019, **21**, 4595–4599.
- 37 M. Brovetto and G. Seoane, *J. Org. Chem.*, 2008, **73**, 5776–5785.
- 38 V. S. C. de Andrade and M. C. S. de Mattos, *Curr. Org. Synth.*, 2015, **12**, 309–327.
- 39 G. Liao, X. S. Yin, K. Chen, Q. Zhang, S. Q. Zhang and B. F. Shi, *Nat. Commun.*, 2016, **7**, 12901.
- 40 (a) D. S. Matteson, *Tetrahedron*, 1989, **45**, 1859–1885; (b) D. S. Matteson, *J. Org. Chem.*, 2013, **78**, 10009–10023; (c) S. Kirupakaran, H.-G. Korth and C. Hirschhäuser, *Synthesis*, 2018, 2307–2322; (d) D. S. Matteson, B. S. L. Collins, V. K. Aggarwal and E. Ciganik, *Org. React.*, 2021, **105**, 427–860.
- 41 (a) J. Gorges and U. Kazmaier, *Org. Lett.*, 2018, **20**, 2033–2036; (b) O. Andler and U. Kazmaier, *Chem. – Eur. J.*, 2021, **27**, 949–953; (c) O. Andler and U. Kazmaier, *Org. Biomol. Chem.*, 2021, **19**, 4866–4870; (d) O. Andler and U. Kazmaier, *Org. Lett.*, 2022, **24**, 2541–2545.
- 42 (a) O. C. Ho, R. Soundararajan, J. Lu, D. S. Matteson, Z. Wang, X. Chen, M. Wei and R. D. Willett, *Organometallics*, 1995, **14**, 2855–2860; (b) M. Tost, O. Andler and U. Kazmaier, *Eur. J. Org. Chem.*, 2021, 6459–6471; (c) T. Kinsinger and U. Kazmaier, *Org. Lett.*, 2022, **24**, 3599–3603.
- 43 (a) D. S. Matteson and J.-J. Yang, *Tetrahedron: Asymmetry*, 1997, **8**, 3855–3861; (b) D. S. Matteson, R. Soundararajan, O. C. Ho and W. Gatzweiler, *Organometallics*, 1996, **15**, 152–163.
- 44 A. Chan, W. M. Hussein, K. A. Ghaffar, N. Marasini, A. Mostafa, S. Eskandari, M. R. Batzloff, M. F. Good, M. Skwarczynski and I. Toth, *Bioorg. Med. Chem.*, 2016, **24**, 3095–3101.
- 45 M. Kawai, R. S. Pottorf and D. H. Rich, *J. Med. Chem.*, 1986, **29**, 2409–2411.
- 46 M. Kijima, M. Yoshida, K. Sugita, S. Horinouchi and T. Beppu, *J. Biol. Chem.*, 1993, **268**, 22429–22435.

