

REVIEW

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Lysine methyltransferase inhibitors: where we are now

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Protein lysine methyltransferases constitute a large family of epigenetic writers that catalyse the transfer of a methyl group from the cofactor *S*-adenosyl-L-methionine to histone- and non-histone-specific substrates. Alterations in the expression and activity of these proteins have been linked to the genesis and progress of several diseases, including cancer, neurological disorders, and growing defects, hence they represent interesting targets for new therapeutic approaches. Over the past two decades, the identification of modulators of lysine methyltransferases has increased tremendously, clarifying the role of these proteins in different physio-pathological states. The aim of this review is to furnish an updated outlook about the protein lysine methyltransferases disclosed modulators, reporting their potency, their mechanism of action and their eventual use in clinical and preclinical studies.

Introduction

Despite evidence of histone methylation being reported in the 1960s,¹ the first enzyme that is able to catalyse this modification was only identified forty years later.^{2,3}

From the chemical point of view, histone methyltransferases (HMTs) catalyse, using *S*-adenosyl-L-methionine (SAM, AdoMet) as a cofactor, the introduction of a methyl mark to the arginine or lysine residues of target proteins. Lysine methyltransferases (KMTs) are responsible for the mono-, di- and trimethylation of N ϵ -lysine in histone and non-histone substrates (Fig. 1).

The introduction of this methyl mark is a dynamic process that controls transcriptional regulation and gene expression and it has a pivotal role in cell fate determination. As a result, these proteins have been related to several disease conditions such as cancer and neurodegeneration.^{4–8} These findings make KMTs an appealing target for therapeutic intervention

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Monica Viviano

been an assistant professor at the Department of Pharmacy of the same university, focusing on advanced procedures for the synthesis and biological evaluation of small molecules of biological interest.

Monica Viviano received her Pharmaceutical Chemistry degree in 2009 at the University of Salerno. During her PhD, she spent 6 months in the Kappe Laboratory at Karl Franz University (Austria) to expand her knowledge in MW and continuous-flow techniques. In 2013, she obtained her PhD in Pharmaceutical Sciences at the University of Salerno, where she was a post-doc working on small-molecule modulators of epigenetic targets. Since October 2021, she has

and have encouraged the identification of modulators of these enzymes.

The first group of KMTs includes around 55 members, which share a common evolutionarily conserved SET [Su(var),⁹ enhancer of zeste,¹⁰ trithorax¹¹] domain that is responsible

for the catalytic activity. The SET domain is approximately 130 amino acids long and the central region (i-SET) is preceded and followed by n-SET and c-SET domains, which interact with each other and are required for methyltransferase activity.^{12,13}



Alessandra Cipriano

Alessandra Cipriano received her Master of Science degree (MSc) in Pharmaceutical Chemistry and Technologies in 2016 at the University of Salerno. During her PhD program, she spent 6 months in the laboratory of Prof. Ciulli (University of Dundee, Scotland, UK) to expand her knowledge in the PROTACTM field. In 2020, she obtained her PhD in Drug Discovery and Development at the University of Salerno, where she is currently a post-doctoral researcher working on the design and synthesis of modulators of epigenetic targets. Her main research interests are the development of advanced procedures for the synthesis of small molecules that targeting epigenetic and non-epigenetic targets.



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Ciro Milite received his Master of Science degree (MSc) in Pharmaceutical Chemistry in 2007 at the University of Salerno where he obtained his PhD in Medicinal Chemistry in 2011. During his PhD, he spent a period at the "SCRIPPS Research Institute" at the Barbas Laboratory to expand his knowledge in the field of organocatalyzed enantioselective chemical reactions. In 2018 he gained a position as assistant professor at the Department of Pharmacy, University of Salerno and, since 2021, he has been an associate professor. His research activity is focused on epigenetic drug discovery, spanning synthetic strategies, chemical biology, and biophysical techniques.



Sabrina Castellano

Sabrina Castellano obtained her Medicinal Chemistry degree in 1995 from the University of Trieste (IT) and her PhD at the University of Milan (IT). In 1999, she got a Postdoctoral Fellowship from the University of Trieste and in 2001 a position as an assistant professor. She has spent two years as a visiting scientist in Oyun Kwon's group at UCLA (USA). Since 2014 she has been an associate Professor at the University of Salerno (IT). In 2018, she obtained the National Scientific Habilitation to become full professor in medicinal chemistry. She has published more than 70 scientific peer-reviewed papers and holds one patent. Her main research interests are the broad area of medicinal chemistry, covering the design of small-molecule modulators of biological targets, the development of advanced procedures for the synthesis of compounds to be used in drug-discovery projects, and the application of modern biophysical approaches to study the different aspects of protein-ligand interactions.



Gianluca Sbardella

Gianluca Sbardella graduated in Pharmaceutical Chemistry and Technology at the University of Rome "La Sapienza" (Italy) in 1993 and received his PhD from the same university in 1997. In 2001 he joined the Department of Pharmacy of the University of Salerno, where since 2016 he has been Full Professor in Medicinal Chemistry. During his career, Gianluca was a Research Fellow at the Drug Chemistry Center of CNR (Italian National Research Center) in 1997, a Research Fellow at the Istituto Superiore di Sanità (ISS) – the Italian National Institute of Health – in 1998–1999, a postdoctoral fellow at the University of Rome "La Sapienza" in 2000, and a research associate at the University of Siena in 2000–2001. In 2004 he spent a sabbatical period as a visiting professor at the Department of Chemistry & Biochemistry of UCLA. After early interest in the discovery of antiviral drugs, since 2000 his research has regarded most aspects of epigenetic drug discovery, spanning synthetic strategies, medicinal chemistry, chemical biology, and biophysical techniques. He is currently the Past President of the Medicinal Chemistry Division of the Italian Chemical Society (Divisione di Chimica Farmaceutica della Società Chimica Italiana).



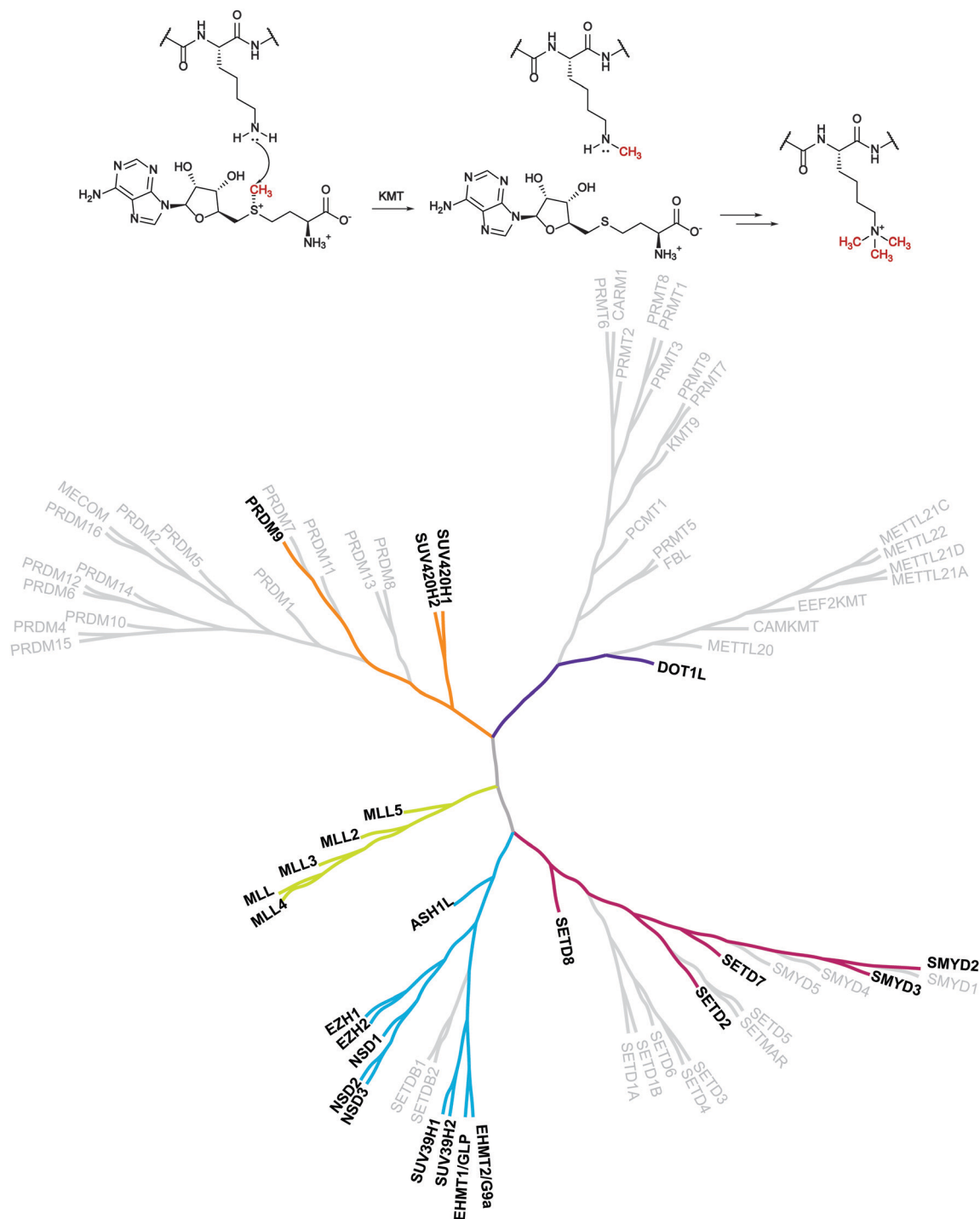


Fig. 1 Lysine methylation catalysed by lysine methyltransferases and the KMT phylogenetic tree. In bold are depicted the proteins that are the object of this review for which inhibitors are known.

The second group of KMTs is composed of non-SET-containing proteins that transfer the methyl group, exploiting a catalytic portion which looks like that of DNA and protein arginine methyltransferases (DNMTs and PRMTs).¹⁴

In the past decade, several reviews regarding these proteins have been published, some of them focusing on structural features of the proteins, some on their involvement in different

pathologies, and others reporting the currently available inhibitors for some of the proteins.^{15–23}

In this review, we report an updated overview of the current literature regarding selected KMTs for which high-quality chemical probes have been reported. For each of these proteins, we provide a summary of their biological functions, including the implications in pathological states, and describe the

corresponding inhibitors on the basis of their mechanism of action. For selected inhibitors (at least one for each protein), structural insights into the binding mode are also provided.

DOT1L

The protein DOT1L was identified for the first time in a yeast screen (*Saccharomyces cerevisiae*) performed for searching genes that, when overexpressed, disrupt telomeric silencing.²⁴ Homolog genes of DOT1 (DOT1-like or DOT1L) present 88% homology in their amino acid sequence and they have been found in different species such as mammals, drosophila, and protozoa.²⁵ DOT1L (disruptor of telomeric silencing 1-like) is the most studied non-SET-containing methyltransferase that is responsible for the mono-, di- and trimethylation of lysine 79 of histone H3 (H3K79) in yeasts and mammals, where it is associated with transcriptionally active genes. DOT1L represents the sole epigenetic protein responsible for this modification; indeed, the knockout of its gene in different species causes the complete loss of H3K79 methylation.^{26–28} DOT1L catalyses the transfer of a methyl group from the cofactor SAM to the H3 substrate thanks to the presence of an AdoMet binding motif, which resembles that of DNMTs and PRMT1.¹⁴

DOT1L can methylate histone H3 only in the presence of DNA, while no activity has been detected on free histone H3, meaning that only parental histones will be methylated by this protein.²⁶

DOT1L is involved in different processes, including physiological as well as pathological ones. From the physiological point of view, this protein has a key role in transcriptional activation, promoting the formation of heterochromatin. In addition, it is involved in embryonic development, in the regulation of haematopoiesis, in cell cycle progression and in DNA repair after damage.²⁷

Moreover, DOT1L together with other proteins such as AF10, AF17, AF9, ENL, Skp1, TRRAP and β -catenin, forms the multi-metric complex “DotCom”, which has a specific enzymatic activity as it can catalyse H3K79 di- and trimethylation.²⁹ Considering that three of these proteins (AF9, ENL and AF10) are fusion partners with the mixed-lineage leukaemia (MLL) genes, there is a strict link between their chromatin-reading effect and DOT1L activity in leukemogenesis.³⁰ Indeed, it has been demonstrated that aberrant transcriptional activation *via* H3K79 methylation by DOT1L is linked to the insurgence of MLL, a subtype of childhood leukaemia that derives from oncogenic chromosomal rearrangements of the MLL genes.³¹ Thus, although the expression of DOT1L is not modified in leukaemia, its altered methyltransferase activity is a consequence of the chromosomal translocations observed in MLL. However, the precise mechanism by which DOT1L contributes to the gene-activation process has not been fully elucidated and it requires further investigation.

DOT1L inhibitors

Because of the importance of the protein DOT1L in many physiological and pathological processes, different inhibitors of this protein have been developed over the years and they can

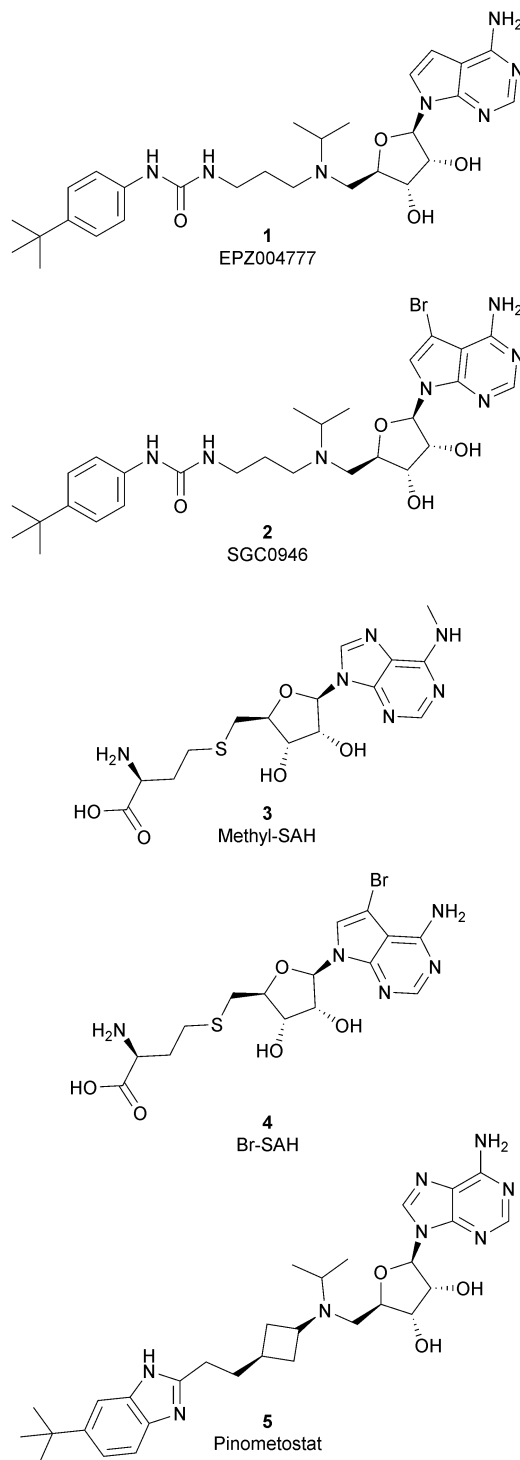


Fig. 2 Nucleoside inhibitors of DOT1L.

be classified, based on their structure, into two main classes: nucleoside inhibitors (Fig. 2) and non-nucleoside inhibitors (Fig. 4).

Nucleoside inhibitors

As the DOT1L mechanism of action is based on the presence of an AdoMet binding motif, early efforts to find inhibitors were



focused on the design of small molecules that are able to occupy the same binding pocket of the cofactor SAM. Therefore, the first class of DOT1L inhibitors is characterized by structural features that mimic the SAM molecule. **1** (EPZ004777) was the first SAM competitor able to strongly inhibit DOT1L (IC_{50} of 0.4 nM), showing *in vitro* and *in vivo* antiproliferative activity in MLL-re-arranged leukemic cells (MV4-11).

In addition, selectivity assays against a panel of 9 other protein methyltransferases (PMTs) showed no inhibition at concentrations up to 50 μ M.³² Despite being one of the strongest DOT1L inhibitors, **1** cannot be used in therapy due to its low half-life, poor pharmacological properties, and side effects. Moreover, even if the etiology of the effect is not clear, a significant inhibition of this enzyme by **1** was observed to be responsible, in a 14 day treatment, for the increasing number of white blood cells, neutrophils, monocytes and lymphocytes in mice.^{32,33}

Starting from **1**, different analogues were synthesized, such as **2** (SGC0946), characterized by a bromide on the adenosine ring, which was more active than the parent compound, showing an IC_{50} of 0.3 nM in a cell-free assay, and an IC_{50} of 8.8 nM and 2.6 nM in MCF10a and A431 cells, respectively.³⁴ Several compounds based on the SAH (S-adenosylhomocysteine) structure also belong to the first class of DOT1L inhibitors: some of them were identified by alkylation of the primary amine of the adenine ring of SAH (methyl-SAHA, **3**)³⁵ or by adding a bromide on the adenosine ring (BrSAHA, **4**), but these modifications led to a deep decrease in DOT1L inhibition and selectivity with respect to the parent compound.³⁶

In 2013, **5** (pinometostat) was identified as a potent ($K_i \leq 80$ pM) and selective DOT1L inhibitor, showing a 37 000-fold selectivity over

other PMTs. It exhibits strong inhibitory activity towards DOT1L in cells and is able to inhibit MV4-11 proliferation with an IC_{50} value of 3.5 nM.³⁷ However, this molecule has low oral bioavailability in mouse, rat, and dog;³⁸ therefore, it is necessary to carry out intraperitoneal administration. A representation of the binding mode of compound **5** obtained using the LigPlot+ program³⁹ is depicted in Fig. 3. The X-ray crystal structure of the inhibitor in complex with DOT1L (PDB 4HRA) shows that **5** is able to occupy the SAM binding site and to induce the opening of a hydrophobic pocket beyond the amino acid portion of SAM. Specifically, the adenosine moiety binds in a similar manner as in SAM structures. The positively charged 5'-amino group is responsible for a hydrogen bond with the carbonyl moiety of Gly163. On the other hand, the benzimidazole nitrogen engages with the carboxylate side chain of Asp161.

Similar to other SAM-mimicking inhibitors, **5** has a low half-life and the effects are visible only after more than 7 days of treatment. However, preclinical experiments evidenced clear tumour regression and little or no re-growth for more than 30 days after cessation of treatment with the compound. **5** was subjected to phase I clinical trials against relapsed or refractory acute myeloid leukaemia in children (NCT02141828)⁴⁰ and adults (NCT01684150),⁴¹ and two different clinical trials are now evaluating the combination of **5** with azacytidine, cytarabine and daunorubicin (NCT03724084;⁴² NCT03701295⁴³) to improve cell killing and tolerability *in vivo*.

Non-nucleoside inhibitors

In recent years, several approaches to discover new DOT1L inhibitors that overcome the pharmacokinetic (PK) limitations of SAM-related compounds have been developed. Luo and co-workers identified the first non-nucleoside DOT1L inhibitor through structure-based virtual screening (SBVS):⁴⁴ the bisaminoquinoline **6** showed high inhibitory activity towards DOT1L, but its poor cell permeability was responsible for the weak activity against MV4-11 proliferation. In order to improve the permeability, the same research group developed compound **7**, which showed strong DOT1L inhibition (IC_{50} of 3.55 μ M) and could effectively down-regulate both the level of H3K79 methylation and the overexpression of leukemogenic genes in MV4-11, *via* qRT-PCR and western blot studies.⁴⁵

In 2016, a fragment-based screening (FBS) approach led to the identification of the weak inhibitor **8** (IC_{50} of 320 μ M), whose co-crystallization with DOT1L and structure-based ligand optimization yielded **9** with nanomolar activity (IC_{50} of 14 nM). X-ray crystallography studies confirmed that these new SAM-competitive compounds are not able to interact within the SAM binding site, but with an induced binding pocket adjacent to the binding site of the cofactor: upon the binding of compound **9**, a conformational change of the enzyme did not allow anymore the binding of the SAM cofactor that is fundamental for the activity of DOT1L.⁴⁶ Exploiting again the FBS approach, the same research group identified compounds **10** and **11** as potent and selective DOT1L inhibitors, where both interact with the same binding pocket and had IC_{50} values of 1.4 and 0.4 nM, respectively.⁴⁷ Cellular assays showed that these compounds are able to

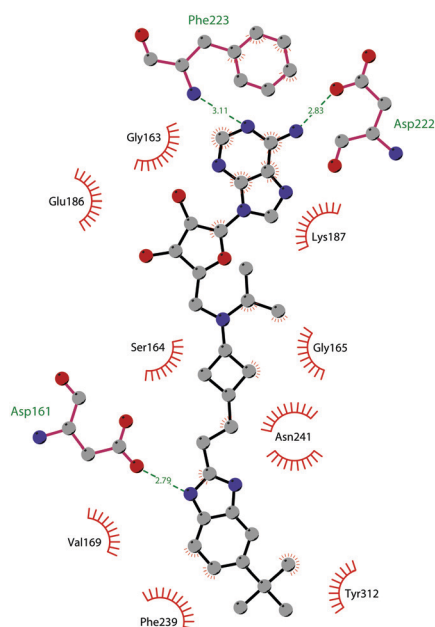


Fig. 3 Flat representation (generated using LigPlot+ from PDB ID 4HRA) showing contacts between compound **5** and active-site residues in DOT1L. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with the compound.



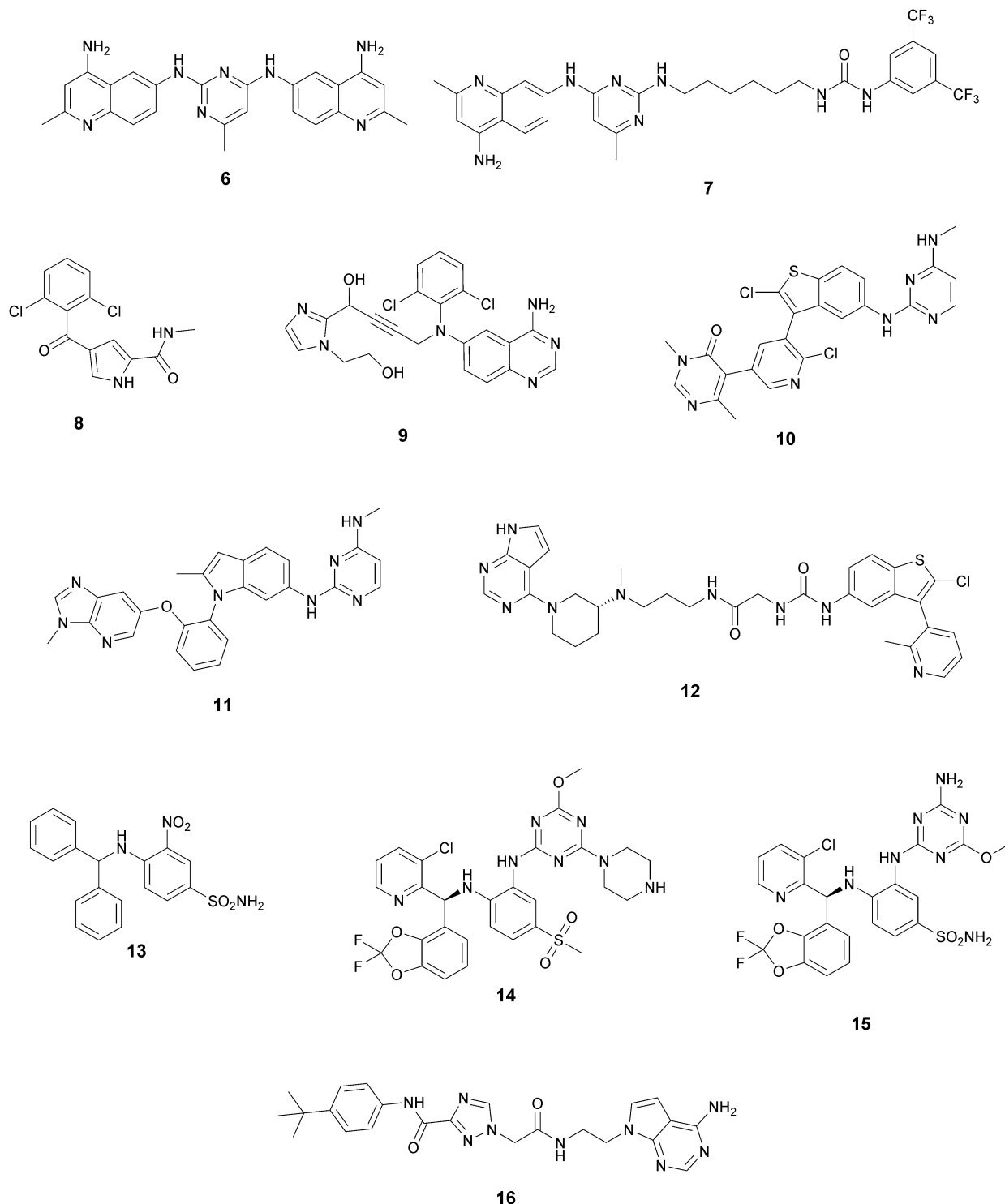


Fig. 4 Non-nucleoside inhibitors of DOT1L.

significantly reduce dimethylated H3K79 levels (IC_{50} of 23 and 16 nM, respectively) and can inhibit the proliferation of MV4-11 leukaemia cells in the nanomolar range, whilst maintaining good PK properties, a moderate half-life and good oral bioavailability.

In 2017, Mobits and co-workers used a combination of FBS, virtual screening (VS) and fragment-linking approaches, which led to the discovery of compound 12. This derivative was

obtained by connecting two different fragments: a non-nucleoside fragment, which mimics the key interactions of SAM bound to DOT1L, and an induced back pocket binder. This new inhibitor showed a K_i in the very low picomolar range and a favourable selectivity profile against a panel of 22 KMTs and PRMTs with no inhibitory activity up to 50 μ M. Moreover, in a head-to-head comparison in cellular assays, 12 worked



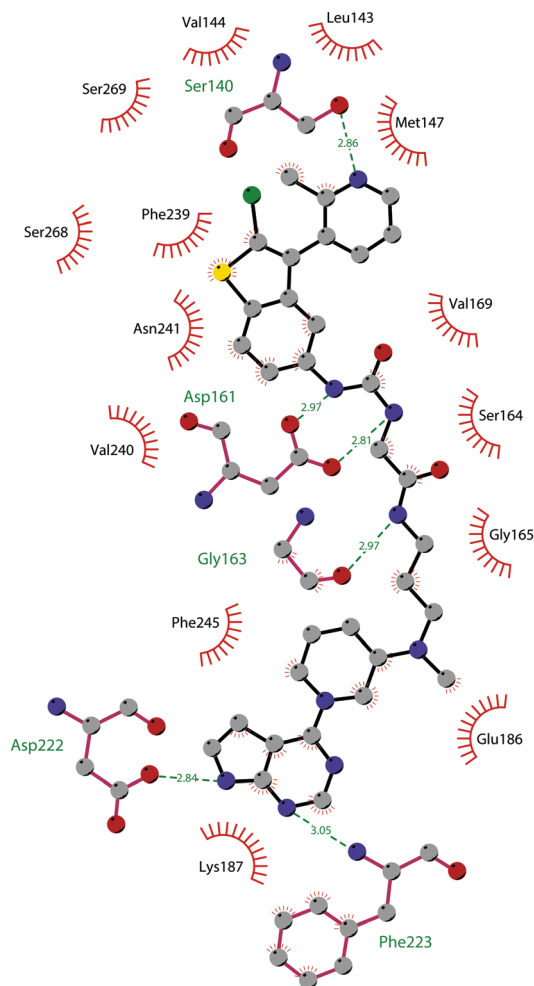


Fig. 5 Flat representation (generated using LigPlot+ from PDB ID 5MW4) showing contacts between compound **12** and active-site residues in DOT1L. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with the compound.

equal to or better than **5**, suppressing H3K79 dimethylation (IC_{50} of 3 nM) both in HeLa and Molm-13 cells and inhibiting proliferation of MV4-11 cells.⁴⁸

The binding interactions of compound **12** with the enzyme are represented in Fig. 5. The X-ray crystal structure of the inhibitor bound to the enzyme (PDB 5MW4) confirmed the expected binding mode: the NH of the pyrrolopyrimidine core serves as a hydrogen-bond donor, interacting with the side chain of Asp222, while the adjacent nitrogen acts as a hydrogen-bond acceptor for the backbone NH of Phe223. An interaction of the pyridine nitrogen with Ser140 is also observed.

Two years later, Stauffer and co-workers exploited the structure-guided optimization of a high-throughput screening (HTS) hit (**13**, IC_{50} of 8.5 μ M) to obtain new DOT1L inhibitors (**14** and **15**) with sub-nanomolar potency (IC_{50} of 0.11 nM and 0.17 nM, respectively). Moreover, selectivity testing for these compounds against a selected panel of histone methyltransferases showed no inhibition below 5 μ M. Despite the good results, a limited efficacy in a mouse tumour xenograft model

was obtained, illustrating the difficulty of reaching the good level of DOT1L inhibition *in vivo* required to effectively suppress MLL-fusion tumour growth.⁴⁹

More recently, Gibbons and co-workers applied a VS strategy on a nucleoside-focused library of 1200 compounds, adenosine-based as well as small molecules with 'nucleoside-like' scaffolds, which is different from the DOT1L clinical candidates and other reported inhibitors. Using structure-based design and molecular modelling studies, compound **16** was identified, showing an IC_{50} of 1.0 μ M and very modest activity against DNMT1, PRMT3, PRMT5 and PRMT8, with selectivity for DOT1L of 250, 130, 175, and >250-fold, respectively. This compound exhibited dose-dependent cellular activity against the proliferation of murine MLL-AF9 transformed cells, accompanied by a dose-dependent decrease of H3K79 dimethylation and induction of cell differentiation. Although this new inhibitor is only modestly potent, it represents a new chemical probe with a unique non-nucleoside scaffold that is able to bind and compete with the SAM-binding site of DOT1L, which could be the starting point for further optimizations.⁵⁰

G9A/GLP

G9a and the related G9a-like protein (GLP) are two proteins with a homology of around 80% in the sequence of their catalytic domain. G9a and GLP are reported also as euchromatin histone lysine methyltransferases (EHMTs), a family of proteins that regulate gene transcription. G9a is also known as lysine methyltransferase 1C or euchromatic histone lysine *N*-methyltransferase 2 (EHMT2) while GLP is reported as lysine methyltransferase 1D or euchromatic histone methyltransferase 1 (EHMT1).^{51,52} G9a and GLP are mainly responsible for the mono- and dimethylation of H3K9, but they are also able to catalyse the methylation of other histone substrates (H3K27, H3K56 and H1),^{53–57} as well as non-histone proteins, including p53,⁵⁸ sirtuin⁵⁹ and G9a itself. In particular, autocatalytic G9a methylation represents a mechanism of gene silencing.⁶⁰

Dimethylated H3K9 is associated with repressed euchromatic regions of the genome, meaning that G9a and GLP repress euchromatic gene transcription through H3K9 dimethylation of specific target genes. In addition, both G9a and GLP have been found to interact with several transcriptionally repressive protein complexes.⁶¹

From the structural point of view, G9a and GLP are characterized by the presence of a C-terminal SET domain, a large N-terminal region for a nuclear localization signal (with a glutamate-rich region and a cysteine-rich ring finger-like domain) and a middle domain characterized by 6 ankyrin repeats as reader domains.^{62,63}

G9a and GLP can exist as homomeric and heteromeric complexes. The heteromeric complex is functional *in vivo* so it was initially believed that this complex was necessary for the catalytic activity. Subsequent studies suggested that G9a and GLP proteins have independent physiological functions.⁶⁴

G9a and GLP are widely expressed in several organs and tissues, as they have multiple biological roles. Mainly, they are linked with both facultative and constitutive heterochromatin



formation and transcriptional repression. They are also involved in the development of the brain, as G9a knockout led to embryonic lethality caused by growth defects.^{52,65}

On the other hand, aberrant activity and expression of G9a and GLP is linked to the development of diseases, mainly cancer.⁶⁶ Several pieces of experimental evidence correlate G9a and GLP up-regulation with metastatic disease and poor prognosis in various types of human cancer. Consistently, knockdown of G9a reduces tumour cell growth in *in vitro* experiments and the invasiveness of cancer.⁶⁷ In addition, dysregulation of G9a and GLP is also associated with inflammatory and neurodegenerative disorders.^{68,69}

G9a/GLP inhibitors

Considering the crucial role of G9a and GLP in many biological processes, over the past decade several G9a and GLP inhibitors have been developed. Based on their mode of action, these compounds can be categorized into two different groups: SAM-competitive inhibitors (Fig. 6) and substrate-competitive inhibitors (Fig. 7).

SAM-competitive inhibitors

SAM-competitive inhibitors bind the cofactor SAM binding pocket of G9a, preventing histone methylation and, consequently, G9a activity. Among these compounds, **17** (**sinefungin**), a natural product structurally correlated to SAM, inhibits GLP/G9a with an IC_{50} of 28.4 and 30.1 μ M, respectively, although it displays no selectivity over a wide range of methyltransferases.

Modifications at the amino acid moiety led to the identification of a series of cycloalkyl-substituted analogues such as **18a,b**. These compounds showed a comparable or better potency than sinefungin (IC_{50} of 21.8 and 39.6 μ M for **18a** and IC_{50} of 1.5 and 1.6 μ M for **18b** on GLP and G9a, respectively). Although the selectivity was evaluated on a limited number of methyltransferases, these compounds appear to be more selective than sinefungin, *i.e.*, **18a** showed no inhibition on PRMT1 and **18b** showed a weak inhibition of DNMT1 and no activity on PRMT1 and SET7/9.^{70,71}

Starting from compound **19** (**BIX01338**), identified *via* HTS as a good G9a inhibitor (IC_{50} of 4.7 μ M) but with poor selectivity and cellular activity, other SAM-mimetic benzimidazoles were designed, leading to the identification of **20** (**BRD9539**).^{72,73} This compound has an IC_{50} value of 6.3 μ M against G9a but is inactive in cell-based assays, probably due to its low cell permeability. Conversely, the methyl ester analogue **21** (**BRD4770**) displayed a higher cellular activity, although its reduced biochemical potency compared to **20**. It was hypothesized that **20** is the potentially active form of **21** in cells, considering that the methyl ester group could be easily hydrolysed in cells acting as a “prodrug”. **21** reduced cellular levels of di- and trimethylated H3K9 with low cellular toxicity and inhibited both anchorage-dependent and -independent proliferation in the pancreatic cancer cell line PANC-1. Despite their interesting profiles, these compounds suffer from low selectivity, considering that **20** also inhibits PRC2 activity, with a similar potency to G9a inhibition as well as NSD2, even if the effects were observed only at 40 μ M.⁷³

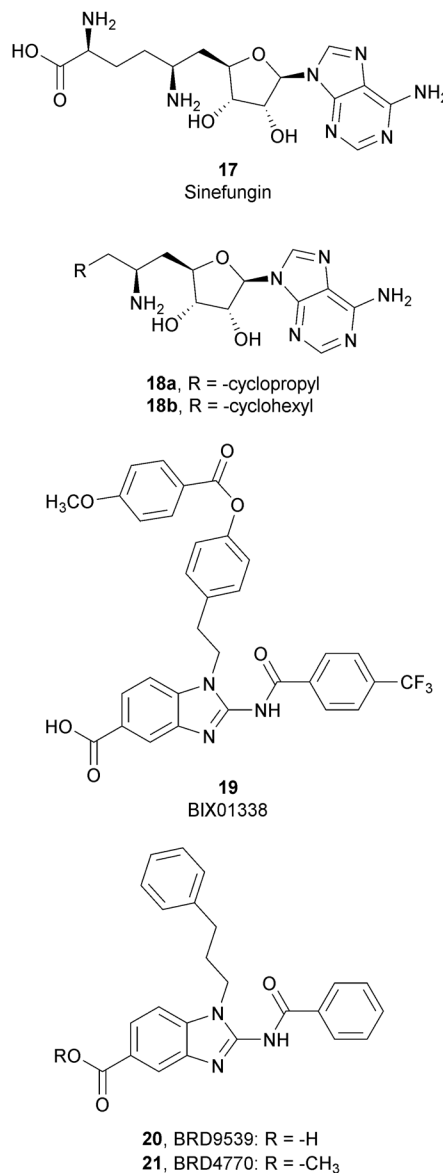


Fig. 6 G9a SAM-competitive inhibitors.

The low selectivity is one of the major drawbacks of this class of inhibitors; consequently, SAM-competitive inhibitors have not been further investigated.

Substrate-competitive inhibitors

The most potent, selective, and widely used chemical probes of G9a/GLP are substrate-competitive inhibitors. These compounds specifically occupy the substrate site in G9a, gaining a better selectivity against the protein target over SAM-competitive inhibitors. In 2007, *via* the HTS of approximately 125 000 compounds, Kubicek and co-workers identified the quinazoline **22** (**BIX-01294**) as a G9a/GLP inhibitor. **22** inhibits G9a and GLP (IC_{50} of 1.9 μ M and 0.7 μ M, respectively) in an uncompetitive manner with SAM. **22** reduces dimethylated H3K9 levels in different G9a target genes, promoting autophagy-dependent cell death and reducing cell proliferation in different types of cancer cell lines,



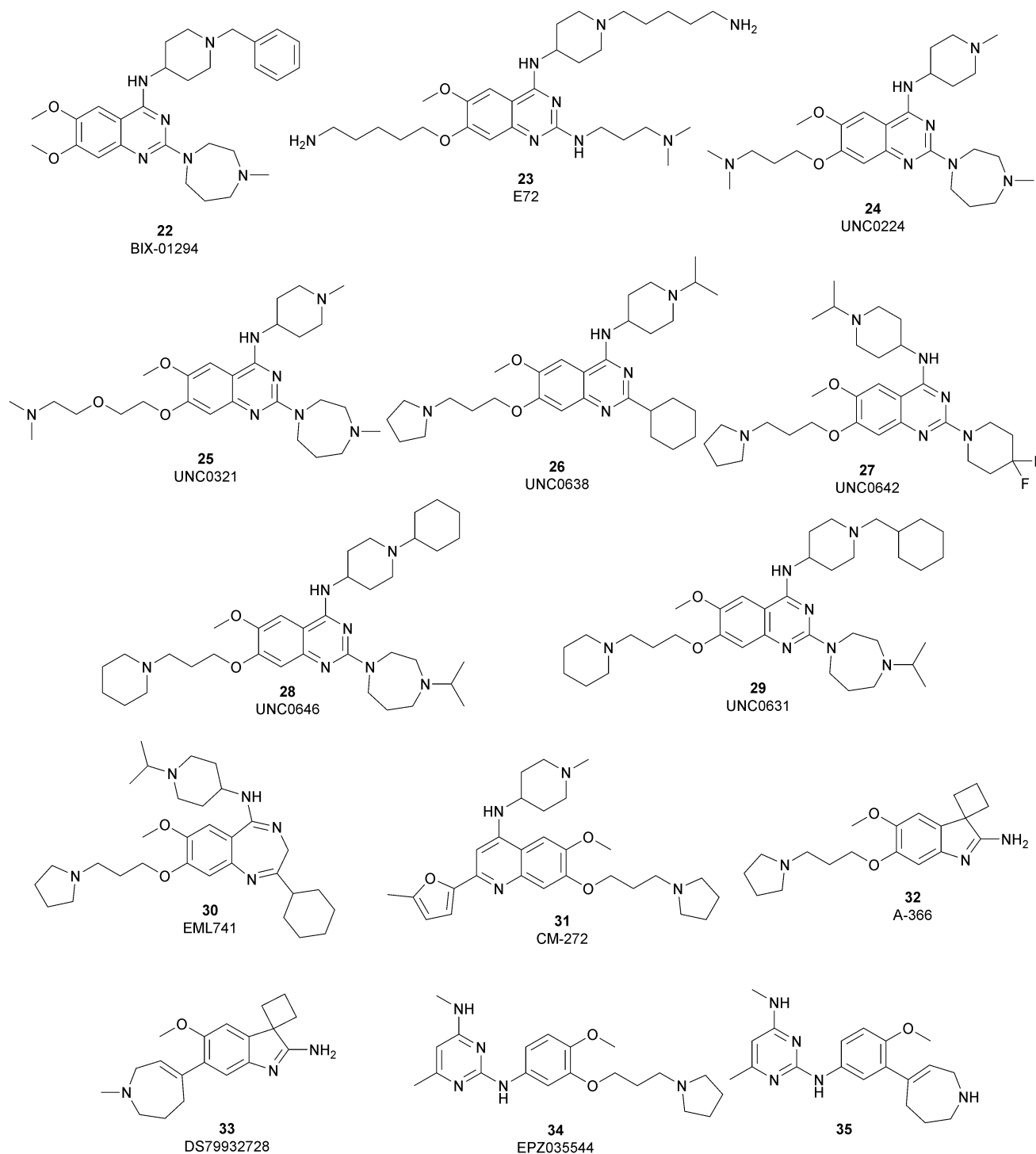


Fig. 7 G9a substrate-competitive inhibitors.

such as colorectal, breast, bladder cancer and glioma cells.^{74–76} Insertion of the 5-aminopentyloxy moiety on **22** to mimic the lysine side chain, yielded **23** (E72), which retains a similar activity to **22** but displays a very low cytotoxicity.⁷⁷ The 2,4-diamino-6,7-dimethoxyquinazoline core was identified as the key pharmacophore, and structure–activity relationship (SAR) investigations allowed the identification of several derivatives. **24** (UNC0224) is an efficient and selective G9a/GLP inhibitor

(IC₅₀ of 15 nM and 20 nM, respectively), whose X-ray crystal structure in complex with G9a showed that the 7-dimethylaminopropoxy group is responsible for important interactions with the lysine-binding channel of this enzyme.⁷⁸

In the quinazoline scaffold, replacement of 7-dimethylaminopropoxy group with an ethoxyethyl chain resulted in **25** (UNC0321) (*K_i* of 63 pM), which is the first reported G9a inhibitor with picomolar potency, although this had a lower



cellular activity compared with **22**.⁷⁹ Despite their high *in vitro* potency, the cellular activity of the above mentioned quinazoline-based compounds is limited, considering that the concentration required to produce functional effects is close to the toxic concentration and that they are characterized by poor druggability properties.

In order to overcome these drawbacks, further optimization of the quinazoline class of inhibitors led to the discovery of **26** (**UNC0638**), a cell-permeable G9a/GLP inhibitor ($IC_{50} < 15$ nM and 19 nM, respectively) with high selectivity over a wide range of other lysine and arginine methyltransferases, as well as over non-epigenetic targets. The X-ray crystal structure of **26** bound to the protein and the SAH (PDB 3RJW) provided structural information about the binding mode, confirming its substrate-competitive mode of action. As depicted in Fig. 8A, the compound engages the substrate binding pocket, inserting the 7-(3-pyrrolidin-1-yl)propoxy side chain into the lysine-binding groove. Moreover, the hydrogen of the amino group at the 4-position of the quinazoline ring establishes an important hydrogen bond with Asp1083, as well as the nitrogen of the quinazoline being hydrogen bonded with Asp1088.

26 potently reduces levels of dimethylated H3K9 in a variety of cell lines and with a lower cellular toxicity than the previously reported compounds. However, the poor PK properties limited its employment in animal studies.⁸⁰ The hypothesis that the cyclohexyl group at position 2 of the quinazoline core could be related to the poor metabolic stability, and supported its replacement with a 4,4-difluoropiperidine moiety, yielding **27** (**UNC0642**). This new derivative maintains the high *in vitro* potency ($IC_{50} < 2.5$ nM) and cellular activity and with a lower

toxicity than the parent compounds but also exhibits improved *in vivo* PK properties.⁸¹ Other related compounds are **28** (**UNC0646**) and **29** (**UNC0631**), potent G9a inhibitors (IC_{50} of 6 and 4 nM, respectively) with excellent potency in a variety of cell lines.⁸²

Apart from the quinazoline derivatives, a limited number of G9a/GLP inhibitors have been identified. Exploiting the scaffold-hopping approach, the quinazoline moiety of **25** was replaced with a benzodiazepine scaffold, resulting in the identification of **30** (**EML741**). This derivative showed high *in vitro* (IC_{50} of 23 nM) and cellular potency, improved selectivity against other methyltransferases and low cellular toxicity. Notably, **30** is also able to inhibit DNMT1, synergizing with G9a inhibition in reducing cellular proliferation. The major interactions of **30** with the target protein are depicted in Fig. 8B. The co-crystal structure of the compound in complex with the catalytic SET domain of GLP in the presence of SAH (PDB 6MBP) showed that the inhibitor and the cofactor are located in two opposite binding sites. Specifically, **30** is bound to the acidic surface of the substrate-binding pocket, inserting the pyrrolidine moiety into the lysine-binding channel. The benzodiazepine scaffold is lodged with Arg1214, Ile1218, and Leu1143 forming van der Waals interactions. Also, the amino group at the 4-position and the protonated nitrogen of the *N*-isopropyl-piperidine group are hydrogen bonded with two negatively charged residues, Asp1140 and Asp1145.⁸³

Starting from the quinazoline scaffold, its bioisosteric replacement with the quinoline one led to **31** (**CM-272**), a dual small-molecule inhibitor that is able to target simultaneously the methyltransferase activity of G9a and DNMTs. In fact, G9a

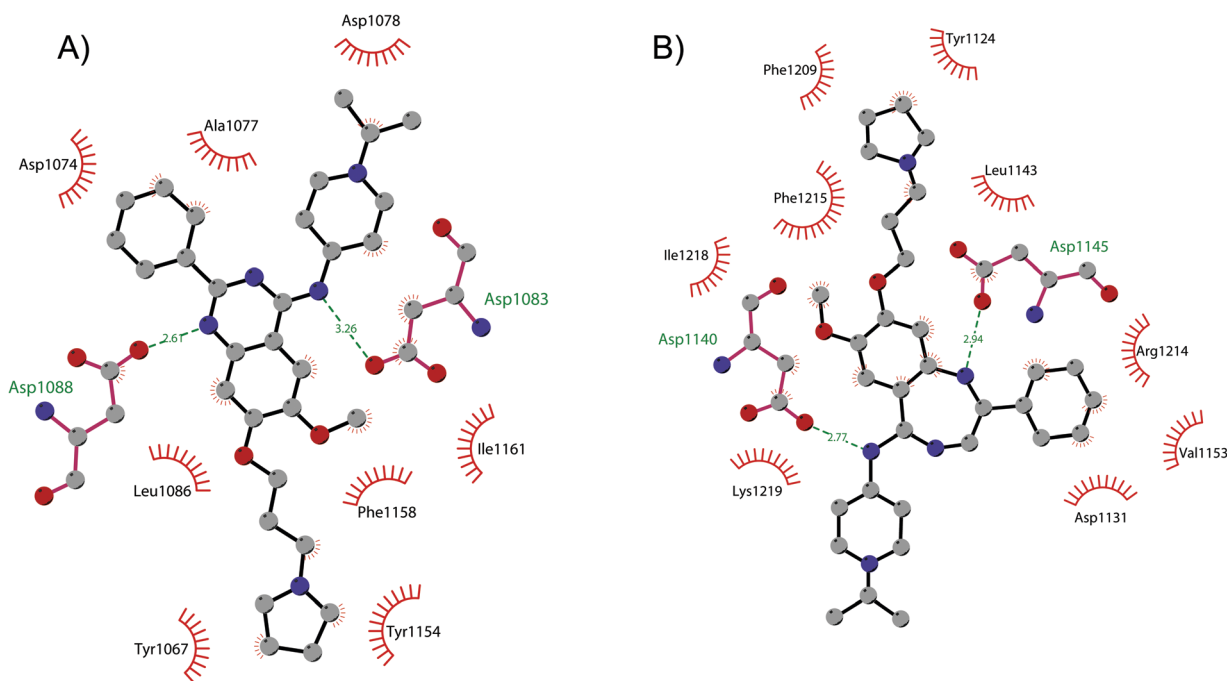
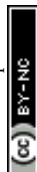


Fig. 8 Flat representation (generated using LigPlot+) showing contacts between (A) compound **26** and active-site residues in G9a (from PDB ID 3RJW) and (B) compound **30** and active-site residues in GLP (from PDB ID 6MBP). Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with compounds.



interacts with DNMT1 to coordinate DNA and histone methylation, promoting transcriptional silencing of the target genes. The dual small-molecule **31** inhibits cellular proliferation, promotes apoptosis, and prolongs the survival of haematological neoplasia xenogeneic models, offering a novel approach for treating several human cancers with poor prognoses.⁸⁴ Through a library screen of a subset of compounds, the spiroindole **32** (**A-366**) was identified as a potent and selective inhibitor (IC₅₀ of 3.3 nM and 38 nM for G9a and GLP, respectively).⁸⁵ The compound occupies the lysine-binding site of G9a and is peptide-competitive and SAM-uncompetitive. Therefore, it is not very surprising that, two years after its identification as a G9a/GLP inhibitor, compound **32** was also reported as a sub-micromolar inhibitor of the epigenetic methyl reader protein Spindlin1 (IC₅₀ of 186.3 nM), which is also able to engage the target in cells (although only at 100 µM).^{86,87} In fact, the capability of the compound to bind to the same pocket of the lysine substrate of the methyltransferase makes it also able to bind to the reader protein that recognizes the methylation reaction product.

The introduction of a tetrahydroazepine in **32** led to the identification of **33** (**DS79932728**), which retained the potency on G9a (IC₅₀ of 4.50 nM and 33.9 nM for G9a and GLP, respectively), showed good metabolic stability, and the capability to induce γ-globin in monkeys, with interesting implications in the study of β-thalassemia.⁸⁸

Beyond the well-characterized quinazoline structures, interest in the identification of new valuable chemical scaffolds as G9a inhibitors led to consideration of the diaminopyrimidine core of **34** (**EP2035544**), a reported G9a/GLP inhibitor (IC₅₀ of 16.9 and 185 nM, respectively) with low PK profile, as an optimizable novel template. Insertion of the tetrahydroazepine group as a side chain, used successfully in the design of **33**, allowed the identification of **35**, with increased potency on G9a/GLP (IC₅₀ of 4.3 and 32.3 nM, respectively) compared with **34** and an improved PK profile in mice (the oral bioavailability increased up to 40%).⁸⁹

SUV39H1 and SUV39H2

The suppressors of variegation 3–9 homolog 1 (SUV39H1, also known as KMT1A) and 2 (SUV39H2, also known as KMT1B) are members of the SUV39 subfamily of KMTs and are localized in the nucleus. They contain a highly conserved C-terminal SET domain that is responsible for the di- and trimethylation of H3K9, which is associated with gene silencing and heterochromatin formation, while in the N-terminal region there is a chromodomain that is fundamental for the catalytic activity.^{90,91} SUV39H2 has a homology sequence of 59% with SUV39H1; however, it differs from this last protein by the presence of a basic histone H1-like N-terminus.⁹²

SUV39H1 preferentially methylates the monomethylated H3K9, exploiting the binding of the chromodomain to the methylated residue, although it remains unclear if SUV39H2 prefers the methylated or unmethylated substrate H3.^{93,94} In addition, SUV39H2 methylates histone H2AX on Lys134, which is a critical modification for the production of phosphorylated H2AX (γ-H2AX) in cancer cells.⁹⁵

SUV39H1 and SUV39H2 play an essential role in several biological mechanisms, such as transcriptional repression and epigenetic silencing in euchromatin, heterochromatin organization, the regulation of telomere length in mammalian cells, and embryonic development.^{96,97}

Knockout of both SUV39H1 and SUV39H2 can lead to abnormally long telomeres, due to the lack of their enzymatic activities of targeting heterochromatin,⁹⁷ and it causes prenatal lethality, accompanied by a global reduction in trimethylated H3K9,⁹⁸ as well as a reduction of hematopoietic stem cell function.⁹⁹

Despite sharing the same enzymatic activity, SUV39H1 and SUV39H2 might play different roles in cancer, probably due to their different substrate preference. In fact, SUV39H1 is considered as a tumour-suppressor gene because of its cell-proliferation-suppressing activity, while for SUV39H2 a tumour-promoting activity has been reported.¹⁰⁰ In particular, alterations as well as an overexpression of SUV39H2 can be found in different cancers, indicating that SUV39H2 is involved in carcinogenesis, invasion, and metastasis of different kinds of solid tumours and haematological malignancies.^{85,101} Dysregulation of SUV39H2 has also been reported in several other diseases, such as hereditary nasal parakeratosis, which is caused by a mutation of SUV39H2 in dogs.¹⁰² Moreover, it has been reported that alterations in the regulation of SUV39H2 are also associated with autoimmune diabetes¹⁰³ and steatohepatitis in mice.¹⁰⁴

SUV39H1 and SUV39H2 inhibitors

As discussed above, SUV39H1 and SUV39H2 have emerged as critical proteins involved in the insurgence of several pathological states, mainly cancer. In spite of great efforts having been spent toward the development of modulators of both these proteins, only three inhibitors have been reported so far (Fig. 9).

Until recently, the fungal epipolythiodioxopiperazine metabolites chaetocin¹⁰⁵ and verticillin A¹⁰⁶ were the sole SUV39H1 inhibitors described; however, although potent (an IC₅₀ value of 0.6 µM was reported for chaetocin), both compounds also inhibit several other histone methyltransferases and are non-specific redox and covalently active compounds.^{105–107}

In 2019 a virtual screening campaign performed on the SET domain of human SUV39H1, followed by an *in vitro* validation, led to the identification of a few hits, including a 4,5-dioxo-6H-pyrrolo(3,2-*e*)indole-2,8-dicarboxylate derivative. Subsequent structural optimization studies yielded compound **36** (**F5446**), which was found to inhibit SUV39H1 with an IC₅₀ of 0.496 µM and decreased the deposition level of the repressive H3K9me3 mark at the promoter regions of the genes coding for the major proteins used by CD8⁺ cytotoxic T lymphocytes (CTLs) to induce tumour cell apoptosis, thus increasing their expression.¹⁰⁸ Compound **36** also showed low toxicity in mice models, and *in vivo* studies showed that it suppresses established tumour growth in a CD8⁺ CTL-dependent manner. Subsequent studies also showed that the inhibition of SUV39H1 by **36** effectively suppresses human colon carcinoma xenograft growth *in vivo*.¹⁰⁹ In 2018 the screening of a panel of commercially available



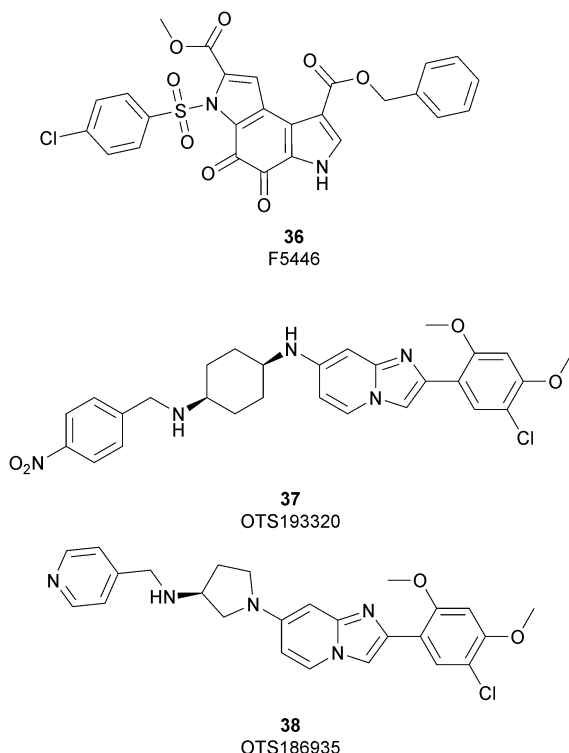


Fig. 9 SUV39H1 and SUV39H2 inhibitors.

compounds enabled the identification of some imidazo[1,2-*a*]pyridine compounds with weak SUV39H2 methyltransferase inhibitory activity, with IC_{50} values of $<10 \mu M$. A subsequent SAR-based drug optimization study yielded **37** (**OTS193320**), a potent SUV39H2 inhibitor (IC_{50} of 22.2 nM) that displays a growth-suppressive outcome of SUV39H2-positive A549 lung cancer cells (IC_{50} of 0.38 μM). In *in vitro* studies, this compound decreased global H3K9 trimethylation levels in breast cancer cells and activated apoptotic cell death. Furthermore, when combined with doxorubicin, **37** reduced γ -H2AX levels as well as cancer cell viability compared with a single agent administration. Further optimization allowed the identification of **38** (**OTS186935**) (IC_{50} of 6.49 nM), which is able to significantly inhibit tumour growth without any detectable toxicity in mouse xenograft models, exploiting MDA-MB-231 breast cancer cells and A549 lung cancer cells.¹¹⁰ In order to show the binding interactions between **38** and the active-site residues of SUV39H2, a flat representation using LigPlot+ has been prepared (Fig. 10). The deposited X-ray crystal structure of the methyltransferase domain of the enzyme bound to **38** (PDB 6P0R) revealed the importance of the nitrogen of the imidazo[1,2-*a*]pyridine scaffold for the interaction with Thr303, responsible of the correct allocation of the molecule inside the binding site.

However, it should be noted that these compounds were not well characterized by the authors in terms of their selectivity against other PMTs and, consequently, caution should be exercised when considering their use as chemical probes. Nevertheless, to the best of our knowledge they are the most potent SUV39H2 inhibitors reported so far.

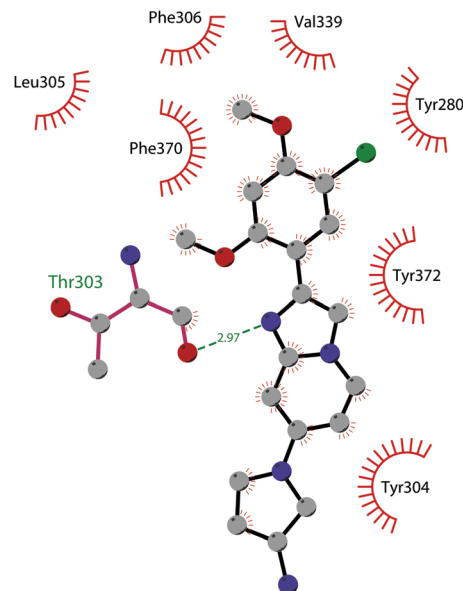


Fig. 10 Flat representation (generated using LigPlot+ from PDB ID 6P0R) showing contacts between compound **38** and active-site residues in SUV39H2. Hydrogen bonds are shown as dashed lines. The spoke arcs represent protein residues making non-bonded (hydrophobic) contacts with the compound.

EZH1 and EZH2

E(Z) is one of the founding members of the SET domain family of methyltransferase proteins.¹¹¹ Enhancer of zeste homologs 1 and 2 (EZH1 and EZH2) are two homolog proteins belonging to Polycomb Repressive Complex 2 (PRC2), a family of proteins involved in the maintenance of gene silencing thanks to the introduction of post-translational modifications on histones and chromatin condensation.¹¹² PRC2 is composed of four core components: EZH1 and EZH2, suppressor of zeste 12 (SUZ12), retinoblastoma suppressor associated protein 46/48 (RBAP46/48) and embryonic ectoderm development (EED). EZH1 and EZH2 represent the catalytic subunit and they are responsible for lysine mono-, di- and trimethylation of H3K27 thanks to their SET domain,¹¹³ while the other core subunits behave as regulators of the methyltransferase activity of EZH1 and EZH2. However, it remains unclear if they possess other functions independently from EZH1 and EZH2.^{114–116} Among all of these, of particular importance is the EED subunit, a WD40-repeat-containing protein that recognizes, thanks to the presence of an aromatic cage, the methyl marks deposited by EZH proteins, and this interaction causes a conformational change in the structure of these methyltransferases, which increases their catalytic activity. This mechanism allows the propagation of H3K27 trimethylation on nucleosomes, contributing to the maintenance of the chromatin in a repressive form.¹¹⁶

From the functional point of view, the PRC2 complex is recruited to non-methylated CpG islands (CGIs) and trimethylates H3K27, with the subsequent recruitment of canonical PRC1 *via* chromodomain proteins (CBX), a subunit of PRC1 that binds to these trimethylated residues. In turn, PRC1 mono-ubiquitylates histone H2A at K119, inducing the condensation of chromatin.^{111,117,118}

Despite sharing the same target genes, EZH1 and EZH2 are differently expressed in tissues and have distinct functions. While EZH1 is ubiquitously expressed, EZH2 expression is found in proliferating tissues. In addition, EZH1 exhibits low levels of histone lysine methyltransferase activity while EZH2 performs this modification robustly. Regarding the condensation of chromatin, EZH1 represses transcription and compacts chromatin more efficiently than EZH2; nevertheless, EZH1 and EZH2 have a complementary role, as EZH2 knockdown resulted in EZH1 upregulation.¹¹²

EZH1 and EZH2 are responsible for the regulation of gene expression during development and haematopoiesis, therefore mutations and abnormal expression of these proteins are reported in various types of cancer, mainly haematological malignancies, breast, and prostate cancer. Specifically, over-expression of both EZH1 and EZH2 is associated with tumour aggressiveness, poor prognosis, and relapses.^{119–122}

EZH1 and EZH2 inhibitors

The development of small-molecule inhibitors of EZH1/EZH2 has been an intense area of research. So far, several EZH1/EZH2 inhibitors have been developed and some of them have been entered in clinical trials alone or in combination with other therapies for a variety of tumours. Considering that EZH1/EZH2 are the catalytic subunit of the PRC2 complex, different drug-design strategies have been explored, targeting both the catalytic domain as well as other complex components such as the EED subunit. Consequently, inhibitors can be categorized into two different classes: inhibitors of the EZH1/EZH2 catalytic domain (Fig. 11) and inhibitors of the EED subunit (Fig. 13). Moreover, the development of degraders (Fig. 18) that are able to induce protein degradation of the PRC2 complex will be discussed.

Inhibitors of the EZH1/EZH2 catalytic domain

The first class of inhibitors developed comprises small molecules targeting the catalytic SET domain EZH1/EZH2. It is worth noting that EZH1 and EZH2 share 96% sequence identity within their catalytic SET domain and 76% sequence identity overall, making the identification of selective ligands challenging. Despite these difficulties, most of the compounds identified have been shown to be more active towards EZH2 over EZH1, while some molecules targeted both proteins, acting as dual inhibitors. The benefits of selective or dual EZH1/EZH2 inhibition are still under discussion. Notwithstanding the good effects of compounds with a greater activity on EZH2 *versus* EZH1, good results can also be achieved with dual EZH1/EZH2 inhibitors considering that EZH1, despite the lower methyltransferases activity compared with EZH2, plays a minor role in balancing the deletion of EZH2.^{122,123}

From a structural point of view, the pyridone-benzamide core is a highly optimized scaffold for the binding to EZH2 in a SAM-competitive mode. In fact, in 2012 Epizyme Inc. identified **39** (EPZ005687) as an EZH2 inhibitor (K_i of 24 nM), displaying 500-fold selectivity over 15 other PMTs and 50-fold selectivity over the closely related enzyme EZH1. This compound selectively inhibits H3K27 methylation in EZH2 wild type and Y641- or

A677-mutant lymphoma cells as well as other tumour cell lines, such as breast and prostate cancers.¹²⁴ The optimization process of **39** led to the identification of **40** (EPZ6438 or tazemetostat), with an increased potency (K_i of 2.5 nM) and a better PK profile compared with the parent compound. **40** is more than 35-fold more selective towards EZH2 over EZH1¹²⁵ and it selectively killed wild type as well as human lymphoma cell lines harbouring point mutations in the EZH2 catalytic domain, inhibiting H3K27 trimethylation.¹²⁶ **40** is orally available and, in 2020, the US FDA approved it, and it was marketed as Tazverik[®], the first EZH2 inhibitor for the treatment of metastatic or locally advanced epithelioid sarcoma in adults and paediatric patients older than 16 years. In the same year, the FDA conferred accelerated approval to **40** for patients with relapsed or refractory follicular lymphoma, who have received at least two preceding systemic therapies and whose tumours have an EZH2 mutation or have no satisfactory alternative treatment options. In addition, **40** was also evaluated in other clinical trials in combination with other therapeutic agents (NCT04179864; NCT01897571; and NCT03854474).^{127–129}

In 2015, Epizyme Inc. developed **41** (EPZ011989), a selective and orally available inhibitor of EZH2, structurally related to **40**. In particular, the central aromatic core was differently decorated: besides the pyridone scaffold, the pyran substituent of **40** was replaced with a *N*-(2-methoxyethyl)-*N*-methylcyclohexanamine and the morpholine was connected through an acetylene linker instead of a benzene moiety. **41** maintains its activity both on mutant and wild-type EZH2 (K_i of <3 nM) with more than 3000-fold selectivity over the other tested HMTs and more than 15-fold selectivity for EZH2 over EZH1. **41** significantly reduced tumour growth in a mouse xenograft model of human B cell lymphoma.¹³⁰

Maintaining the pyridone-amidic moiety, which is crucial for the EZH2 binding, Lu and co-workers exploited a scaffold-hopping approach on the benzene rings of compound **40**, obtaining a series of benzofuran-based EZH2 inhibitors. Among them, **42** (EBI-2511) displayed good enzymatic (IC_{50} of 4.0 nM) as well as cellular activity, showing a reduction of trimethylated H3K27 levels in a dose-dependent manner both in Pfeiffer cells (IC_{50} of 6.0 nM) and WSU-DLCL2 cells (IC_{50} of 55 nM). In a Pfeiffer xenograft mouse model, **42** showed a dose-dependent inhibition of the tumour growth, resulting in a good reduction in tumour size with no changes in body weight. Notably, **42** showed a superior antitumor efficacy compared with **40** at the same dose, suggesting that it might accomplish a similar efficacy but at lower dosages in clinical studies.¹³¹

Through HTS, in 2012 Glaxo Smith Kline USA (GSK) disclosed **43** (GSK126) a selective inhibitor of both wild-type and mutant EZH2 (K_i^{app} of 0.5–3 nM).¹³² Suppressing EZH2 activity, **43** significantly decreased the growth of EZH2-mutant diffuse large B-cell lymphoma (DLBCL) in xenograft mice upon intravenous administration.¹³³ **43** entered in a phase I clinical trial for different types of solid tumours and lymphoma but showed insufficient clinical activity (NCT02082977).¹³⁴ It was suggested that the activity of EZH2 inhibitors is drastically affected by the tumour microenvironment, as confirmed by the antitumor efficacy of **43** when administered to immunodeficient but not



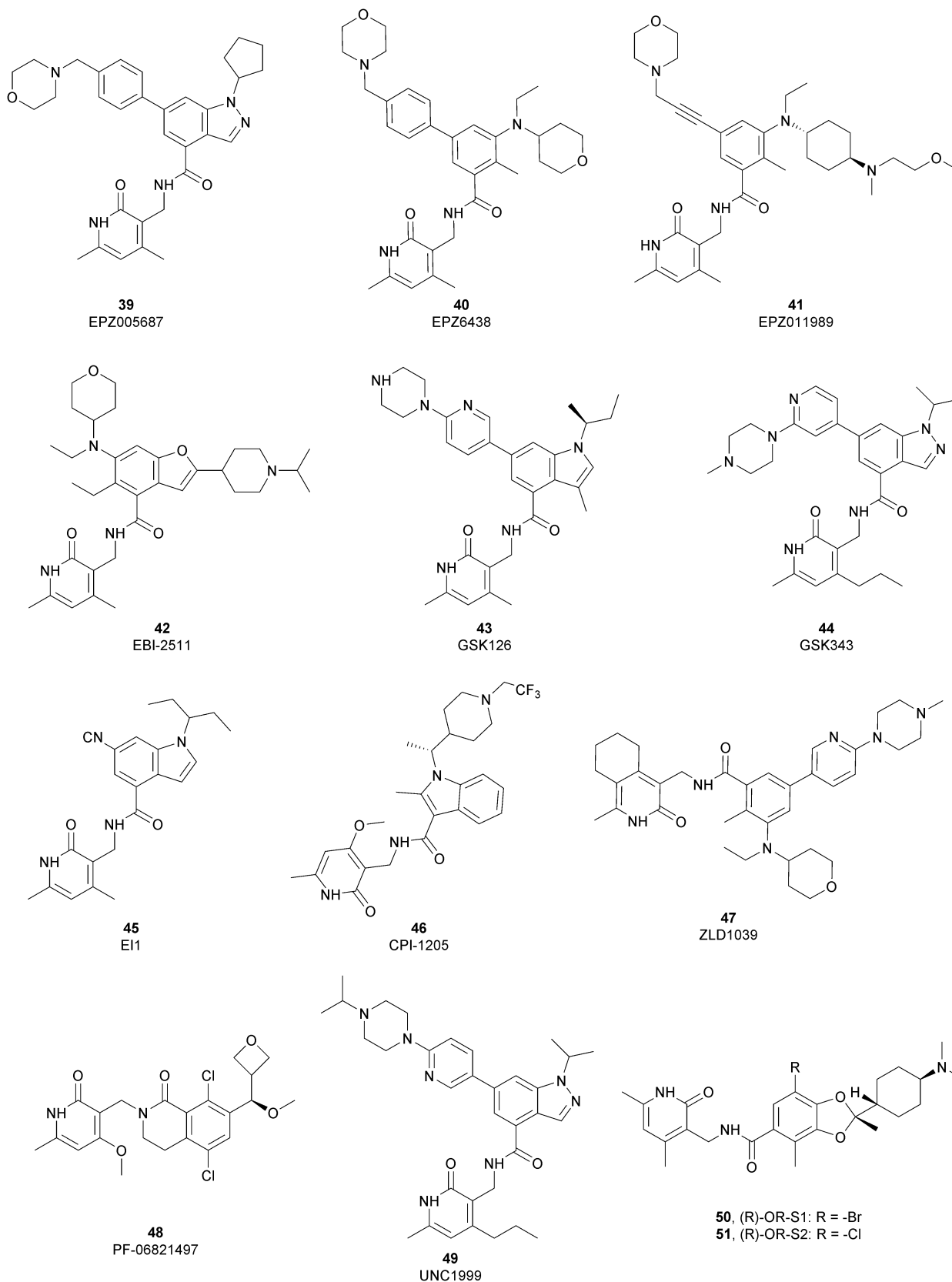


Fig. 11 Inhibitors of the EZH1/2 catalytic domain.



to immunocompetent hosts.¹³⁵ Liu and co-workers were able to solve the crystal structure of human PRC2 bound to **43** (PDB 5WG6).¹³⁶ The main interactions are depicted in Fig. 12. The pyridone moiety is embedded in a binding pocket formed by residues of Trp624, Arg685, and Phe686. In particular, the interactions of the pyridone group confirmed the SAM-competitive mode of action of this compound. In fact, the amide nitrogen and the carbonyl oxygen establish distorted hydrogen bonds with both the carbonyl oxygen and the amide nitrogen of Trp624 of EZH2, thus preventing binding to the cofactor SAM. In addition, the Tyr111 is hydrogen bonded with the carbonyl oxygen between the indole group and the pyridone ring. Interestingly, the linked pyridine and piperazine rings of **43** protrude above the gate to the solvent.

In 2012, compounds **44** (GSK343) and **45** (EI1) were also identified. Specifically, GSK developed **44** as an efficient (EZH2 K_i^{app} of 1.2 nM), selective, and cell-active EZH2 inhibitor. Despite its great potential as an *in vitro* tool to explore EZH2 biology, **44** is not suitable for *in vivo* studies, considering the high clearance in rat PK studies.¹³⁷ At the same time, Novartis exploited HTS, followed by structural optimization, and obtained derivative **45**, which is able to inhibit the methyltransferases activity of EZH2 with more than 10 000-fold selectivity over other methyltransferases and 90-fold selectivity over EZH1. **45** showed good cellular activity, decreasing H3K27 methylation without disrupting the chromatin binding of PRC2. **45** is equally active against both wild-type and Y641-mutant DLBCL cells, reducing proliferation, causing cell cycle arrest, and apoptosis.¹³⁸

In 2016, Constellation Pharmaceuticals published a series of inhibitors characterized by the presence of an indole ring, identifying **46** (CPI-1205) as a compound with a modest selectivity for EZH2 over EZH1 (IC_{50} of 2 nM and 52 nM, respectively) compared with the high selectivity shown against 30 other histone or DNA methyltransferases.¹³⁹ In a KARPAS-422 lymphoma xenograft model, **46** demonstrated significant tumour growth reduction

upon 25 days of administration. The compound completed a phase I study in patients with B-cell lymphomas, (NCT02395601)¹⁴⁰ showing anti-cancer activity and no significant toxicity. In a phase Ib study, **46** also produced positive results for patients with metastatic castration-resistant prostate cancer (mCRPC), demonstrating good tolerability, minimal safety issues and encouraging clinical activity. Currently, **46** is under investigation in a phase Ib/II trial to evaluate its effects on mCRPC in combination with enzalutamide or abiraterone/prednisone (NCT03480646).¹⁴¹ Another interesting aspect is that EZH2 inhibition in human T cells by **46** produced phenotypic and functional alterations of the regulatory T-cells and enhanced the cytotoxic activity of effector T-cells. It was speculated that modulation of EZH2 expression in T-cells can improved anti-tumor responses *via* anti-CTLA-4 therapy. Indeed, ipilimumab

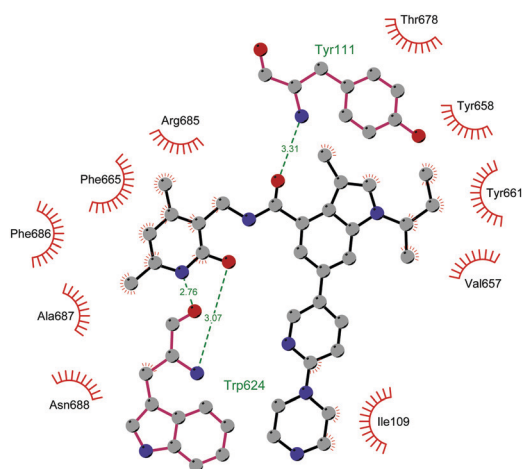
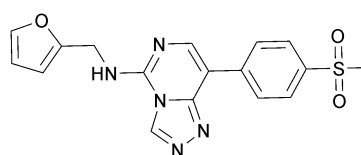
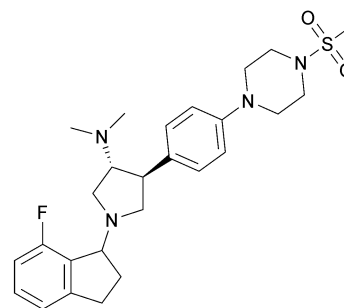


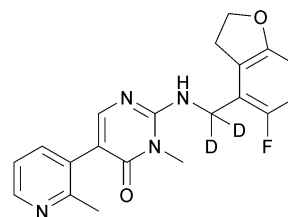
Fig. 12 Flat representation (generated using LigPlot+ from PDB ID 5WG6) showing contacts between compound **43** and human PRC2. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with the compound.



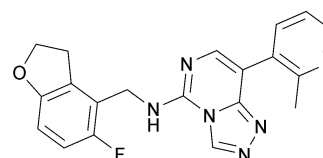
52
EED226



53
A-395



54
BR-001



55
MAK683

Fig. 13 Inhibitors of the EED subunit.



increased EZH2 expression in peripheral T-cells, suggesting a strong rationale for its combination trial with **46**.¹⁴²

Compound **47** (**ZLD1039**), featuring a cyclohexane group fused with the pharmacophoric pyridone moiety, inhibited EZH2 wild type as well as Y641F and A677G mutant (IC_{50} of 5.6, 15, and 4 nM, respectively). **47** reduced the global H3K27 methylation levels and re-established the expression tumour suppressors that are responsible for the higher survival of patients with breast cancer. In three different breast tumour xenograft models (MCF-7, MDA-MB-231, and 4T1 cell lines), **47** decreased tumour growth and metastasis, endorsing the role of EZH2 in breast cancer progression.¹⁴³

In 2018, the Pfizer Global Research group optimized a previous series of lactam-derived EZH2 inhibitors, developing the oxetane-containing compound **48** (**PF-06821497**), which exhibited remarkable potency (K_i less than 0.1 nM) and pharmaceutical properties as well as robust tumour growth inhibition *in vivo*.¹⁴⁴ **48** entered a phase I clinical trial for the treatment of adults with relapsed/refractory small cell lung cancer, castration-resistant prostate cancer and follicular lymphoma (NCT03460977).¹⁴⁵

Regarding dual inhibitors, **49** (**UNC1999**) was reported by researchers at the University of North Carolina as dual EZH1 and EZH2 inhibitor (IC_{50} of 45 and <10 nM, respectively). This compound reduced trimethylated H3K27 levels in cells and selectively killed DLBCL cells carrying the EZH2-Y641 mutations together with inhibition in growth of MV4-11 cells.^{146,147}

Other orally bioavailable EZH1/EZH2 dual inhibitors are **50** (**(R)-OR-S1**) and **51** (**(R)-OR-S2**), which strongly and selectively inhibited methyltransferase activity of both EZH2 and EZH1 (IC_{50} of 8.4 and 7.4 nM and 2.5 and 10 nM for EZH2/EZH1, respectively). Compared with EZH2 selective inhibitors, these compounds showed a remarkable antitumor efficacy *in vitro* as well as *in vivo* against DLBCL cells with EZH2 mutation. These dual inhibitors displayed a strong efficacy both in haematological and solid cancer panel assays, suggesting that both EZH2 and EZH1 are implicated in some types of cancer and their simultaneous inhibition would be effective for targeting them. Accordingly, **51** inhibited the *in vitro* growth of MV4-11 cells more effectively than **49**, which is probably due to the weaker enzymatic activity of **49** on EZH1.¹⁴⁸

Inhibitors of the EED subunit

Besides inhibitors of the EZH1/2 catalytic subunit, other components of the PRC2 complex have been targeted to inhibit the methyltransferase activity. This strategy could prevent some recently emerged issues, such as the gain-of-function mutations of EZH2 resulting in EZH2-inhibitor-resistant cancer cells lines. Considering the role of EED in enhancing the catalytic efficiency of the PRC2 complex, inhibition of the EED interaction with the trimethylated histone substrate was explored as a strategy to modulate the enzymatic activity of the complex. So far, several inhibitors targeting the EED subunit have been reported.

In 2017, **52** (**EED226**) and **53** (**A-395**) were identified as nanomolar inhibitors of EED and studies helped to clarify the effects on EED inhibition. Specifically, Novartis identified **52** as

a potent allosteric binder of the PRC2 complex (IC_{50} of 23.4 nM). The high-resolution crystal structure of **52** bound to EED and to an EED binding domain (EBD) (PDB 5GSA) revealed that the compound binds the pocket responsible for the recognition of the trimethylated H3K27 substrate. Specifically, the compound is inserted into a deeper pocket that is created by a conformational change of the four aromatic residues which form the 'aromatic cage' of the binding site (Phe97, Tyr148, Trp364, and Tyr365), positioning the furan group on the inside and the methylsulfonic group towards the solvent. Fig. 14 depicts the major interactions of **52**: the triazolopyrimidine core forms π - π interactions with Tyr148, Tyr365, and Phe97 residues while the furan group is stabilized by van der Waals and cation- π interactions with other residues of the binding pocket. Moreover, two hydrogen bonds stabilize the interactions of **52** inside the pocket: the amino group between the bicyclic and the furan moieties is hydrogen bonded with the side chain carbonyl of Asn194 while the bottom nitrogen of the triazolopyrimidine core establishes a hydrogen bond with the side chain of Lys211. Despite the moderate cell permeability, **52** led to a dose-dependent reduction of both di- and trimethylated H3K27 markers in G401 cells in addition to a strong proliferation inhibition in lymphoma cells carrying EZH2 mutations as well as cell lines with acquired resistance to SAM-competitive EZH2 inhibitors. Administration of **52** in mice using a subcutaneous xenograft model of Karpas422 showed a clear dose-dependent reduction of trimethylated H3K27 levels with a good tolerability. To better evaluate the effect of PRC2 inhibition *in vivo*, **52** was administered orally to mice for a longer period and a complete tumour regression was observed after 32 days. Furthermore, a

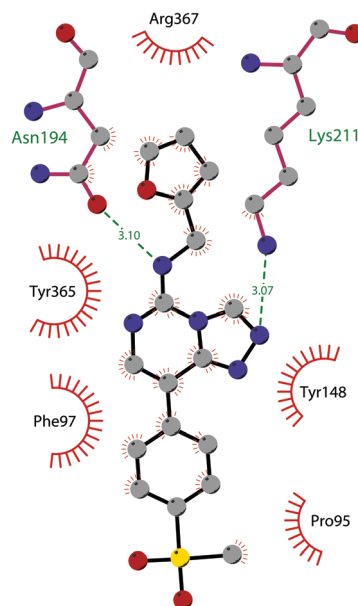


Fig. 14 Flat representation (generated using LigPlot+ from PDB ID 5GSA) showing contacts between compound **52**, EED, and an EZH2 peptide referred to as the EED binding domain (EBD). Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with compound.



combination of **52** and the EZH2 inhibitor **45** exhibited synergistic effects in suppressing cancer cell growth, suggesting a potential combination therapy.¹⁴⁹

In the same year, AbbVie reported the identification of **53** as a selective EED inhibitor with high *in vitro* potency (K_D of 1.5 nM from surface plasmon resonance (SPR) assay and IC_{50} of 18 nM in a radioactivity-based assay). Structurally, **53** is a mixture of two diastereomers, epimeric at the indane stereocenter, and is used for biological evaluation without separation. The crystal structure of **53** with the EED WD40 domain showed that the compound competes with the histone substrate for binding to the aromatic cage within the WD40 repeat domain. Cellular studies showed that **53** is effective against PRC2 in human cancer cell lines that are sensitive to EZH2 inhibition, leading to phenotypic effects on H3K27 methylation that are comparable to inhibitors of the catalytic EZH1/EZH2 domain.

In a DLBCL Pfeiffer xenograft model, administration of **53** two times weekly for 5 weeks, showed a reduction of tumour growth of 84%. Moreover, **53** retains its effects on EZH2-inhibitor-resistant cell lines. Unfortunately, oral administration and systemic administration using an osmotic pump resulted in insufficient exposures and the only route to administer the compound was subcutaneously.¹⁵⁰

In 2019, the pyrimidone-based compound **54** (**BR-001**) was identified, which is able to selectively bind to EED, competing with trimethylated H3K27 binding (IC_{50} of 4.5 nM). The compound exhibited a strong antiproliferative effect on Karpas 422 cells and anti-tumour efficacy in the CT26 syngeneic model. It has been reported that **54** acts both directly, leading to inactivation of PRC2, as well as modulating the tumour microenvironment promoting CD8⁺ T-cell tumour infiltration.¹⁵¹ Currently, **55** (**MAK683**), patented by Novartis as a new triazolo pyrimidine EED inhibitor, is in a phase I/II study in patients with advanced malignancies such as DLBCL, nasopharyngeal carcinoma or other advanced solid tumours for whom no further effective standard treatment is available (NCT02900651).¹⁵²

EZH2 and EED degraders

Besides canonical single-site inhibitors, novel approaches have been exploited to target the PRC2 complex, developing chemical agents that are able to selectively induce protein degradation. Specifically, PROteolysis TArgeting Chimeras (PROTACTM) have been developed, linking a binder for the protein-of-interest and an E3 ligase ligand to induce ubiquitination, thereby achieving protein degradation *via* the proteasome. At the same time, hydrophobic tagging was exploited, attaching a hydrophobic group to a selected protein binder, leading the misfolding or unfolding of the target protein and its subsequent proteasome-degradation. Following the same strategies developed to obtain single-site inhibitors, EZH2 and EED degraders were developed, using selective warheads for both proteins (Fig. 15a and b, respectively).

In 2018, the discovery of a first-in-class EZH2-selective degrader **56** (**MS1943**) was reported. This compound was designed by merging a non-covalent and selective inhibitor of EZH2 (IC_{50} of 12 nM and 2.5 μ M for EZH2 and EZH1, respectively) that is structurally related to **44** to a bulky adamantyl group as a

hydrophobic tag. Compound **56** preserved the potency (IC_{50} of 120 nM) and selectivity in inhibiting EZH2 activity (IC_{50} more than 10 μ M for EZH1). In cancer and non-cancerous cell lines, **56** significantly reduced the EZH2 protein levels while it had a minimum or no effect in reducing the EZH1 protein levels. **56**-induced EZH2 degradation resulted in significant cell growth inhibition and cytotoxic effects in multiple triple-negative breast cancer cell lines but not in normal cells. Moreover, **56** is able to completely suppress tumour growth *in vivo* with good tolerability, suggesting an alternative treatment for EZH2-dependent cancer.¹⁵³

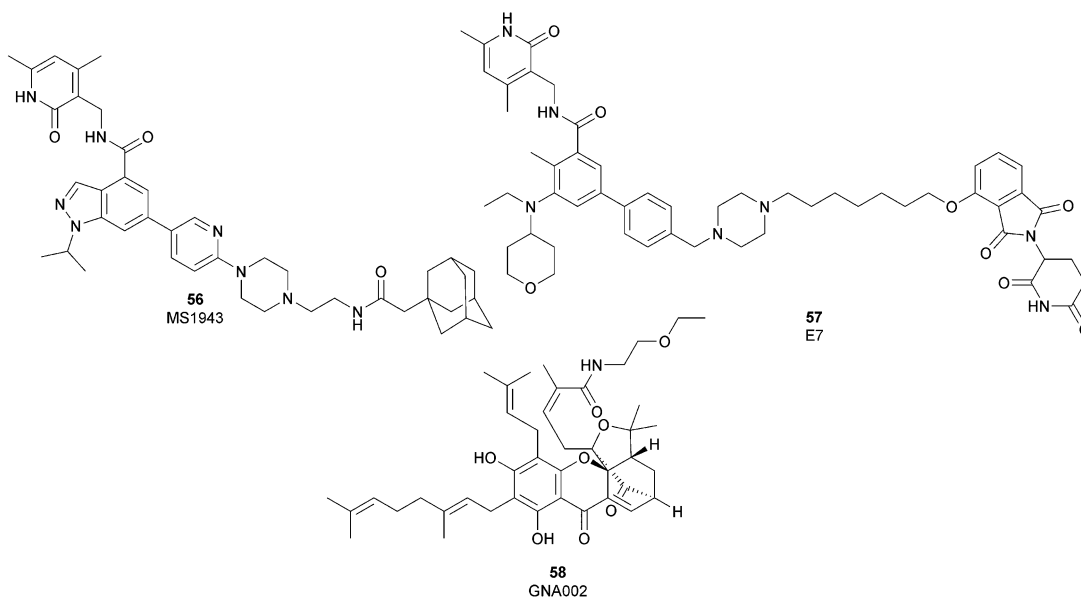
The PROTACTM **57** (**E7**), formally obtained by connecting the protein binder **43** to 4-hydroxythalidomide, a well-studied ligand of the cereblon (CRBN) E3 ligase, through an alkyl linker, has been also reported. **57** showed high potency (IC_{50} of 2.7 nM) in reducing the EZH2 activity and good selectivity over EZH1 (IC_{50} of 179 nM), fully inhibiting the oncogenic activity of EZH2 with antiproliferative effects on EZH2-dependent tumours. Notably, the recruitment of E3 ubiquitin ligase resulted in the indiscriminate ubiquitination and degradation of all PRC2 components, including EZH2, EED, SUZ12, and RbAp48.¹⁵⁴

Besides bifunctional degraders, the compound **58** (**GNA002**) is able to induce EZH2 degradation without featuring a hydrophobic tail or an E3 ligase binder. Specifically, the compound covalently binds the Cys668 residue located within the EZH2-SET-domain, leading to its ubiquitination and consequently its degradation by the protein quality control E3 ligase, CHIP. Moreover, as **58** covalently bound a specific cysteine of EZH2, it did not affect the activity of EZH1 or other examined SET domain-containing PMTs. The ability to significantly suppress tumour growth was investigated using both *in vitro* and *in vivo* assays in wild-type EZH2 solid tumours as well as in lymphomas harbouring EZH2 mutations. These encouraging results suggested an alternative approach for EZH2 degradation as an anticancer strategy.¹⁵⁵

On the other hand, three bifunctional degraders were developed using a selective EED warhead. AstraZeneca researchers reported the identification of two EED-targeted PROTACsTM (**59** and **60**) obtained by merging an EED binder, structurally related to compound **55**, and a well-characterized ligand of von Hippel-Lindau ligase (VHL) with two different types of linker. Both compounds displayed a comparable binding affinity for EED *via* an *in vitro* assay (pIC_{50} of 8.11 and 8.17 for **59** and **60**, respectively). In cellular assays, these PROTACsTM induced rapid EED degradation in cells, inhibiting proliferation of the EZH2 mutant DLBCL cell line as well as the EZH2 wild-type rhabdoid cancer cell line. Unfortunately, no growth inhibition was observed in a limited number of tested EZH2-dependent cell lines, probably due to a cellular fraction of EZH2 that is resistant to PROTACTM-mediated degradation. Interestingly, these degraders also induce the degradation of EED associated proteins in the PRC2 complex, such as EZH2 and SUZ12. Considering that EED degradation is faster than for EZH2 and SUZ12, it was suggested that EED degradation destabilizes the PRC2 complex, triggering other components to the proteasome. Otherwise, it was speculated that formation of the EED-PROTACsTM-E3 ligase ternary complex could expose the lysine residues of EZH2 or



a EZH2 Degraders



b EED Degraders

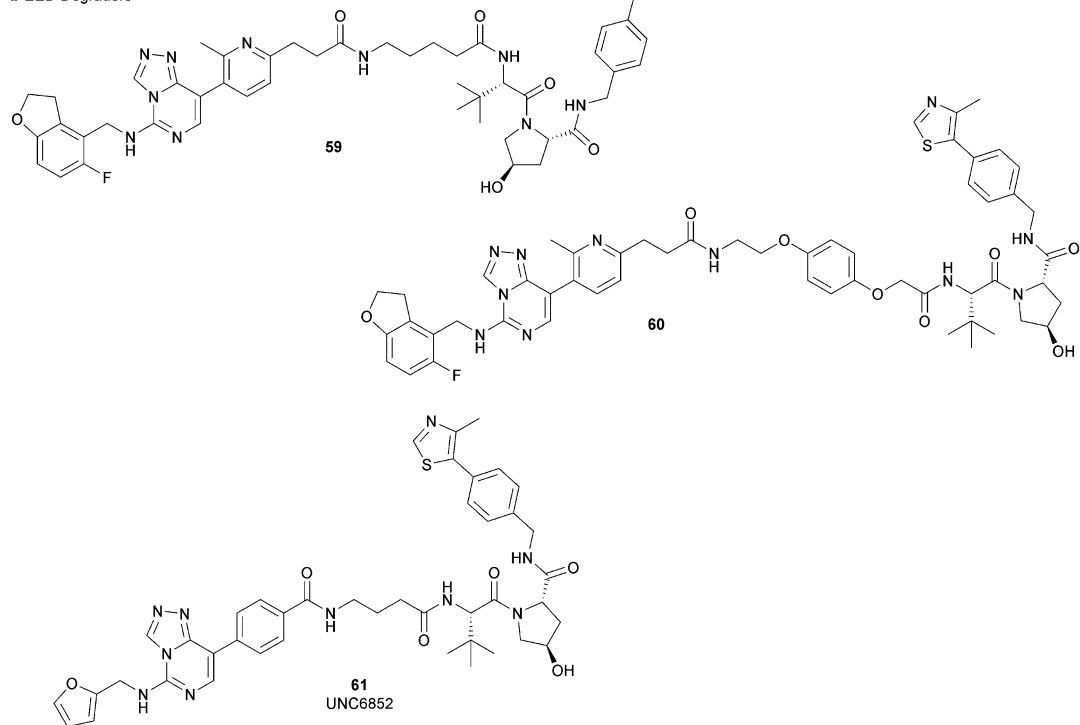


Fig. 15 (a) EZH2 and (b) EED degraders.

SUZ12, which are consequently ubiquitinated.¹⁵⁶ In the same year, **61** (UNC6852) was disclosed as an EED degrader, featuring a 52-derived ligand and a VHL binder. Even if the degrader showed a slight decrease in potency in the binding of WD40 of the EED protein compared with **52** (IC_{50} of 247 nM and 45 nM for **61** and **52** in the same assay, respectively), it is still able to bind the target protein. Like compounds **59** and **60**, **61** also induces proteasomal degradation of all PRC2 components, probably due to the aforementioned mechanisms. **61** is able to prevent the histone

methyltransferase activity of EZH2 in both HeLa cells and DLBCL cells that contain EZH2 gain-of-function mutations. Furthermore, **61** has anti-proliferative effects that are comparable to EZH2 and EED inhibitors in DB and Pfeiffer cells but it is substantially less toxic.¹⁵⁷

SETD7

The SET domain containing lysine methyltransferase 7 (SETD7, also called SET7/9 or KMT7) was the first lysine methyltransferase identified to specifically monomethylate H3K4.^{158–160}



This mark furnishes a site for further modifications, but several studies reported a limited involvement of SETD7 in the di- and trimethylation of H3K4.^{161,162}

SETD7 can also methylate many non-histone proteins (including p53, TAF10, ER, P65, STAT3, SOX2, pRb, SIRT1, DNMT1, SUV39H1 and FOXO34-14) and probably these are the main SETD7 substrates.¹⁴³ Overall, the monomethylation of K372 of p53 represents a central regulatory event as it has a key role for p53 activation in human cell lines. SETD7 activity is triggered by DNA damage and induces the acetylation of lysine residues in the regulatory domain of p53 by different acetyltransferases (p300, CBP and TIP60). The acetylation prevents the ubiquitin-mediated degradation of p53 by the proteasome and, as a final outcome, K372 methylation extends the p53 life.¹⁶³

On the other hand, SETD7 methylation of other substrates can be associated with protein degradation. For instance, human DNMT1 methylation at K1094 by SETD7 increases the turnover of the protein.¹⁶⁴ In addition, DNMT1 can also be monomethylated at K142, leading to its ubiquitination and degradation.¹⁶⁵

Considering the abundance of its substrates in cells, SETD7 is involved in the regulation of various pathways. It has been demonstrated that SETD7, in atherosclerosis and diabetes, acts as a coactivator of inflammatory genes that interact with NF- κ B.¹⁶⁶ Increasing evidence suggests that SETD7 also has a key role in the regulation of processes such as cell differentiation and proliferation. Indeed, many genes required for normal embryonic development frequently harbour the monomethylated H3K4 modification. SETD7 is also overexpressed in a variety of cancers, such as colorectal cancer¹⁶⁷ and hepatocellular carcinoma,¹⁶⁸ making this protein an interesting target for drug discovery.

SETD7 inhibitors

To better explore the regulatory role of SETD7, several attempts have been made to develop potent and selective SETD7 inhibitors. However, the number of selective compounds reported so far is still limited, and the majority of these have a weak inhibitory activity (Fig. 16). As for other methyltransferases, different derivatives have been designed, exploiting as a starting point the structure of the cofactor SAM. The replacement of the sulphur atom with a nitrogen and the introduction of different alkylamino chains that mimic the side chain of lysine yielded different derivatives.¹⁶⁹ Among these, in an ELISA assay, **62** (DAAM-3) showed approximately 50% inhibition against SETD7 at 10 μ M. The co-crystal structure of the SET domain in complex with **62** revealed that the compound binds the cofactor binding site, inserting the *n*-hexyl group within the lysine access channel.¹⁷⁰

An HTS campaign led to the identification of **63** (cyproheptadine), an antagonist of the histamine and serotonin receptor clinically approved as an antiallergy drug, as a SETD7 inhibitor (IC₅₀ of 1 μ M). The X-ray structure of SETD7 in complex with **63** revealed that the compound binds to the substrate-binding site of the enzyme, inducing conformational changes in the C-terminal region and in the amino acid residues involved in

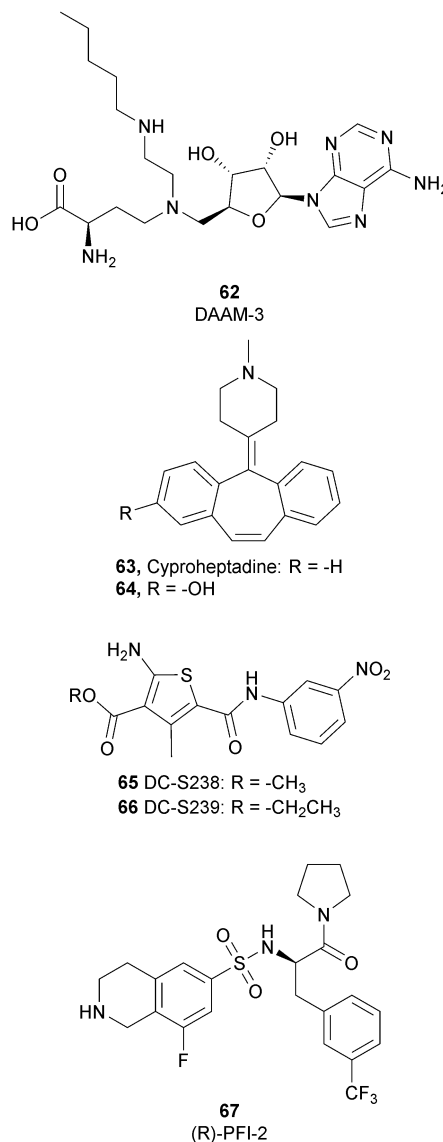


Fig. 16 SETD7 inhibitors.

the formation of the channel for lysine access. According to SETD7 knockdown, **63** reduced the amount of ER α in a dose- and time-dependent manner, enhancing its degradation by the proteasome. This evidence highlights the critical role of SETD7 in the estrogen-dependent transactivation of ER target genes and its potential therapeutic use in diseases associated with upregulation of ER α such as breast cancer.¹⁷¹ Structural optimization of **63** *via* the insertion of various functional groups on the aromatic rings of the dibenzocycloheptene moiety led to the identification of derivative **64**, with an improved potency (IC₅₀ of 0.41 μ M).¹⁷²

SAM competitive and selective inhibitors of SETD7 are **65** (DC-S238) and **66** (DC-S239) (IC₅₀ of 4.88 μ M and 4.59 μ M, respectively). Compound **66** exhibited anti-proliferative effects on breast cancer and leukaemia cell lines MCF7, HL60 and MV4-11 cells but had no cellular activity against HCT116 (human colon cancer) and DHL4 (lymphoma) cells, suggesting



that this derivative is more active towards breast cancer and leukaemia cells.¹⁷³

So far, the most potent and widely used SETD7 inhibitor is **67** ((*R*)-PFI-2), which was identified *via* the HTS of approximately 150 000 compounds of Pfizer's collection.¹⁷⁴ This compound showed a high potency (IC₅₀ of 2.0 nM) and is more than 1000-fold selective over a panel of 18 other methyltransferases and DNMT1. The X-ray crystal structure of the SETD7 catalytic domain bound to **67** revealed an uncommon mode of action, suggesting a substrate-competitive and SAM-dependent binding mode. This evidence was supported by the X-ray crystal structure of the compound bound to the target protein (PDB 4JLG). As depicted in Fig. 17, the pyrrolidine amide engages the pocket that is normally occupied by the target lysine residue, also making direct hydrophobic interactions with the leaving methyl group of SAM, preventing useful interactions of the protein with lysine substrate. Upon the binding of **67** with the protein, the post-SET loop (residues 336–349) adopts a distinctive conformation, making hydrophobic interactions with the trifluoromethyl group. Moreover, the ligand anchors the peptide-binding site, making several hydrogen bonds with different residues, including Gly336 of the post-SET loop Ser268 on the opposite side of the substrate-binding channel.

The cellular activity of **67** highlighted the role of SETD7-dependent methylation in the control, subcellular localization, and function of the transcriptional coactivator Yes-associated protein (YAP). Treatment of **67** in high-density MCF7 cells resulted in a dose-dependent increase of the nuclear YAP and intensified the expression of its target genes, such as AREG and CYR61, indicating that YAP localization in this cell line requires persistent SETD7 activity. It is worthy of note that only **67** is active whereas its (*S*)-enantiomer is 500-fold less active. Molecular dynamics simulations helped to discriminate their different efficacy based on different bioactive energy barriers. This study provides helpful information for future structure-

based drug development, revealing the impact of chirality on SETD7 inhibition.¹⁷⁵

SMYD2

SMYD2 (SET and MYND Domain-Containing Protein 2, also called KMT3C) is a member of a subfamily of SET-domain-containing proteins with a particular domain architecture. This family of proteins is composed of a SET domain that is divided into two parts by a MYND domain/zinc finger motif, followed by a cysteine-rich post-SET domain.¹⁷⁶ SMYD2 has been shown to methylate H3K36 even if it lacks specificity on the free histones. This protein is a relatively promiscuous enzyme as it is responsible for the methylation of many substrates (at least six non-histone substrates have been reported in cells),¹⁷⁷ the best characterized of which is p53, where it shows a highly selective activity. SMYD2 methylates p53 at K370, a repressing modification to p53-mediated transcriptional regulation, and reduces p53-mediated apoptosis. It is of note that the methylation of K370 in p53 is inhibited by SET9-mediated methylation of K372, corroborating the regulatory crosstalk between these post-translational modifications.¹⁷⁸

Physiologically, SMYD2 is involved in the differentiation of embryonic stem cells and, differently from the other SMYD proteins that are equally expressed in somatic and pluripotent cells, it is expressed at high levels in somatic rather than in pluripotent cells.¹⁷⁹ Moreover, it has been reported that SMYD2 is important for cardiac and skeletal function, although it is not essential for the development of the heart.¹⁸⁰

From the pathological point of view, SMYD2 overexpression promotes oncogenic transformation, and high protein levels have been reported in various cancers such as gastric, breast, cervical, pancreatic and lymphoma.^{181–184}

SMYD2 inhibitors

Since the structure description of SMYD2 that was specified in 2011,¹⁸⁵ several pharmaceutical companies have focused research interest on this enzyme. Despite great efforts so far, no SMYD2 inhibitor has been subjected to a clinical-evaluation process, and all compounds reported (Fig. 18) are still at the research level.

In 2011, within a large screening campaign, Ferguson and co-workers from AstraZeneca identified the first substrate-competitive and selective SMYD2 inhibitor **68** (AZ-505). This compound showed an inhibitory potency in the sub-micromolar range (IC₅₀ of 0.12 μM) and the crystallographic structure of SMYD2 in the complex of the cofactor SAM and **68** confirmed the mechanism of action of the inhibitor, giving interesting guidelines for further optimization.¹⁸⁵ Starting from **68**, the formal shift of the hydroxyl group from position 5 of the benzoxazinone to the amino ethyl linker at position 8 resulted in an increased potency, as compound **69** (A-893) is 80-fold more active than the parent compound (IC₅₀ of 2.8 nM). The co-crystal structure of SMYD2 with the inhibitor (PDB 4YND) highlighted the binding mode of this compound. As depicted in Fig. 19, the ether oxygen of the benzoxazinone accepts a hydrogen bond from the side-chain hydroxyl of Tyr258 of the

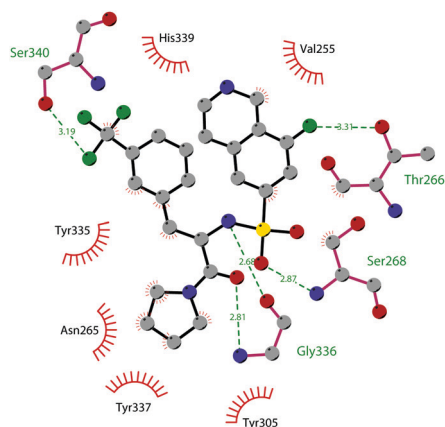


Fig. 17 Flat representation (generated using LigPlot+ from PDB ID 4JLG) showing contacts between compound **67** and active-site residues in SETD7. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with compound.



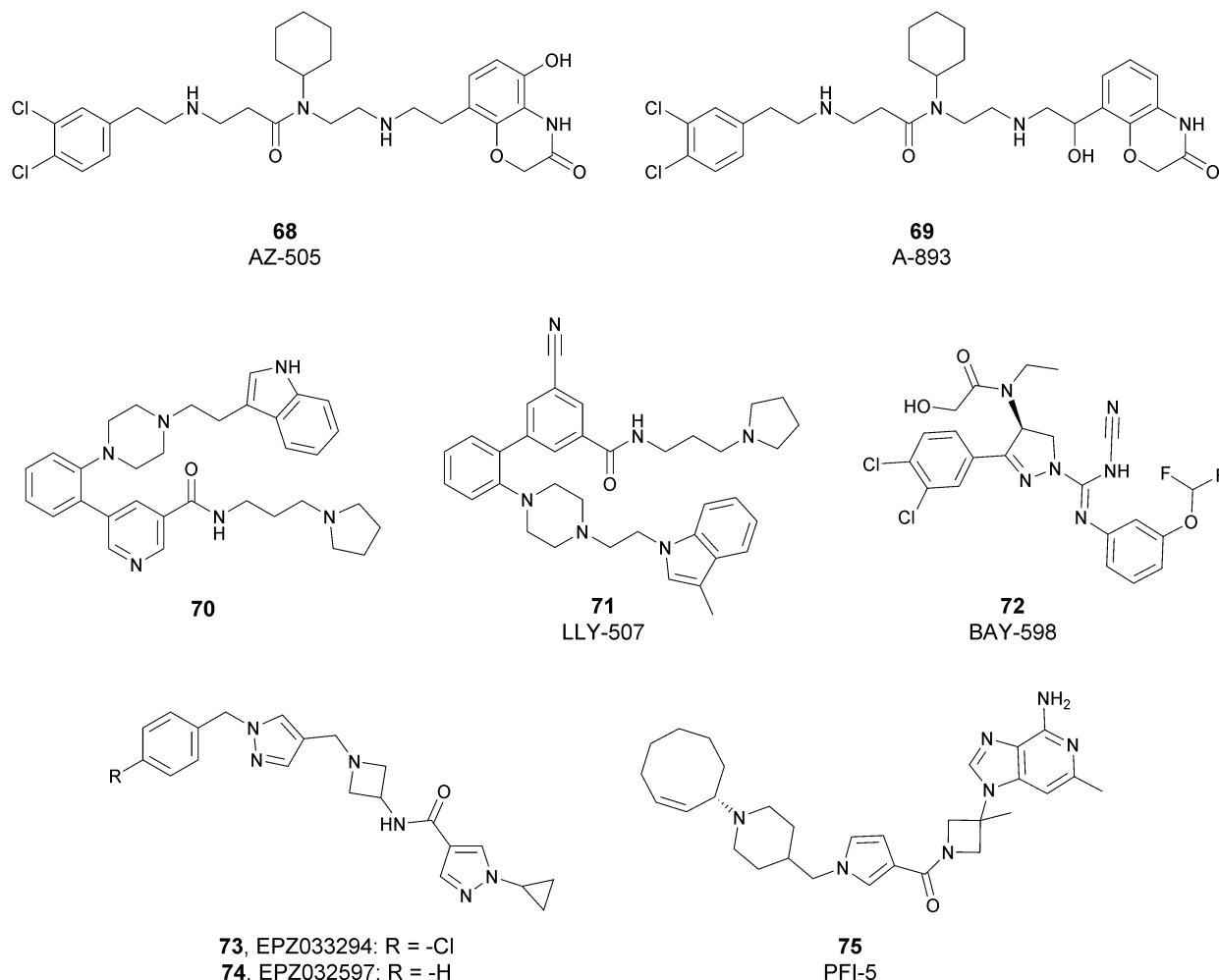


Fig. 18 SMYD2 inhibitors.

lysine channel, and the ring amine maintains a water-bridged interaction. The configuration of the chiral carbon has only a marginal influence on the *in vitro* inhibitory activity (IC_{50} of 2.8 vs. 5.6 nM for (*R*)- and (*S*)-**69**, respectively). **69**, as the parent compound, is a substrate-competitive inhibitor with a good selectivity for SMYD2 over a panel of 31 PMTs and can reduce p53 methylation levels by 42% in A549 lung cancer cells.¹⁸⁶

In 2013, the AstraZeneca group performed an internal hit identification campaign using a combination of HTS and high-concentration fragment screening, which led to the discovery of different substrate-competitive chemical series of SMYD2 inhibitors.^{187,188} In order to improve the biochemical and cellular potency as well as the PK properties, scaffold hybridization techniques were used and compound **70** was developed. This new pyridyl inhibitor showed good selectivity towards SMYD2 over a focused panel of KMTs (*i.e.*, EZH2, NSD1, G9a, and DOT1L), as well as an interesting potency both *in vitro* (IC_{50} of 17 nM) than in the U2OS cell line (EC_{50} of 1.02 μ M).¹⁸⁹

Again, the lead generation efforts by the AstraZeneca group paved the way for further studies that led to the discovery of novel SMYD2 inhibitors. In 2015, Nguyen and co-workers from Eli Lilly and Company, in collaboration with the SGC, designed

the potent ($IC_{50} < 15$ nM) SMYD2 inhibitor **71** (LLY-507), which shows activity in biochemical as well as cell-based assays. This compound showed high selectivity towards SMYD2 (more than two-orders of magnitude on a panel of 25 different methyltransferases, including SMYD3, and other non-epigenetic targets), which is strictly related to the presence of the pyrrolidine group that is able to bind the lysine pocket of the peptide substrate channel of the enzyme. This substrate-competitive compound reduced p53 methylation at sub-micromolar concentrations (IC_{50} of 0.6 μ M) in osteosarcoma (U2OS) and other cell lines and inhibited proliferation in cancer cells characterized by SMYD2 overexpression.¹⁹⁰

In 2016, an optimization of HTS hits by Bayer Pharma resulted in the identification of **72** (BAY-598) as a potent (IC_{50} of 27 nM) substrate-competitive (K_i of 8 nM) SMYD2 inhibitor endowed with suitable PK properties. In this case, stereo-selectivity played a significant role in binding to SMYD2, as the *S*-enantiomer is 50-fold more potent than the *R*-enantiomer (IC_{50} of 27 nM vs. 1.7 μ M for (*S*)-**72** and (*R*)-**72**, respectively). **72** exhibited more than 100-fold selectivity toward SMYD2 on a panel of 32 PMTs, including the related SMYD3 and SUV420H1/H2. The high inhibitory activity of **72** was confirmed in

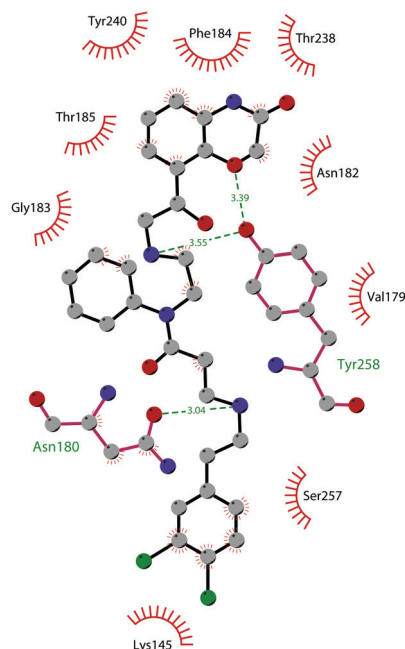


Fig. 19 Flat representation (generated using LigPlot+ from PDB ID 4YND) showing contacts between compound **69** and active-site residues in SMYD2. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with the compound.

HEK293T (embryonic kidney) cells in which it decreased p53 methylation in a dose-dependent manner (IC_{50} of 58 nM) without altering the protein levels. Nevertheless, when tested on a variety of 240 cancer cell lines of different origin, the antiproliferative effect of this inhibitor was limited. *In vivo* studies on mice bearing subcutaneous tumour xenografts derived from the KYSE-150 cell line overexpressing SMYD2 confirmed the capability of this compound to reduce methylation upon oral administration. On the other hand, only mice groups treated with **72** in combination with doxorubicin exhibited a considerable decrease of tumour area with respect to monotherapy or control groups.¹⁹¹

In 2018, Thomenius and co-workers (Epizyme Inc.) described two novel SMYD2 inhibitors, **73** (**EPZ033294**) and **74** (**EPZ032597**), characterized by a very good potency (IC_{50} of 3.9 nM and 16 nM, respectively) against SMYD2 methyltransferase activity. Both compounds showed a 3-fold selectivity ratio toward SMYD2 over a panel of 15 PMTs, including SMYD3. The crystal structure of the ternary SAM-SMYD2-**73** complex underlined the unique binding mode of **73**, positioning the cyclopropyl pyrazole portion in the lysine groove and the hydrophobic tail into a new channel generated by the presence of the inhibitor itself.¹⁹² It is noteworthy that, due to its nanomolar inhibitory activity, **73** is the most potent and the first non-competitive SMYD2 inhibitor known so far.

In 2019, Taylor and co-workers, within a HTS campaign of a large number of SAM-mimicking compounds, identified the first SAM-competitive SMYD2 inhibitors. The structural optimization of a promising hit resulted in the development

of **75** (**PFI-5**), characterized by the presence of a 3-deaza-adenine ring. Moreover, the amino acid side chain and the ribose of SAM were replaced with a linear string of three rings and an azetidine group, respectively. **75** showed a good capability to inhibit the methylation of p53 in MCF7 cells (IC_{50} of 1.3 μ M). In addition, this inhibitor showed high selectivity towards SMYD2 (less than 20% inhibition against a panel of 33 PKMTs, PRMTs and DNMTs) when tested at 50 μ M.¹⁹³

SMYD3

SMYD3 (SET and MYND Domain-Containing 3, also known as KMT3E) is a protein that catalyses the methylation of H3K4.¹⁹⁴ Structural analysis of the SMYD3 protein reveals the presence of a tetratricopeptide repeat (TPR) domain, which is responsible for the peculiar structure of the substrate-binding region and a deep and narrow pocket, which constitutes the C-terminal domain of SMYD3. Both the TPR and post-SET domains are of main importance for the methyltransferase activity.¹⁹⁵

SMYD3 is not able to methylate H3K4 on free histones or on nucleosomes; however, HSP90A can assist SMYD3 in the protein folding, and methylation can occur at H3K4. This interaction of HSP90A with SMYD3 takes place *via* the MYND domain and the TPR motif.¹⁹⁶

Among the non-histone substrates, SMYD3 methylates K831 in VEGFR1, resulting in an enhancement of the kinase activity of the enzyme, which is strictly connected to the migration and invasion of cancer cells.¹⁹⁷ In addition, SMYD3 plays a pivotal role in the regulation of Ras signalling in oncogenic processes through the methylation of MAP3K2.¹⁹⁸

SMYD3 expression has been reported in different tissues such as the brain, kidney, and ovary with the highest expression in skeletal muscle and the thymus. Like SMYD2, SMYD3 is involved in embryonic development, as it has a key role in early embryonic lineage commitment and peri-implantation development in mice, thanks to the activation of specific genes.¹⁹⁹

From the pathological side, SMYD3 is responsible for the transcriptional regulation of carcinogenesis and the development of human cancer.^{200,201} This protein activates the transcription of a set of downstream genes (oncogenes, cell adhesion genes, and signal transduction genes) and it is overexpressed in different human cancers (especially hepatocellular and colorectal carcinomas).²⁰² However, the molecular mechanism by which SMYD3 regulates cancer pathways and its relationship to tumorigenesis *in vivo* are largely unknown.

SMYD3 inhibitors

As described above for SMYD2, an intense involvement of several pharmaceutical companies in the discovery of SMYD3-modulators resulted in the identification of a good number of validated SMYD3-selective inhibitors (Fig. 20), which can be employed as tools to investigate the effects of SMYD3 inhibition.

As the result of *in silico* structure-based HTS, in 2015 Peserico and co-workers described **76** (**BCI-121**) as the first-in-class SMYD3 inhibitor that is able to significantly reduce selected methylation marks at 100 μ M. This compound demonstrated a good potency both in biochemical (IC_{50} of 3 nM) and



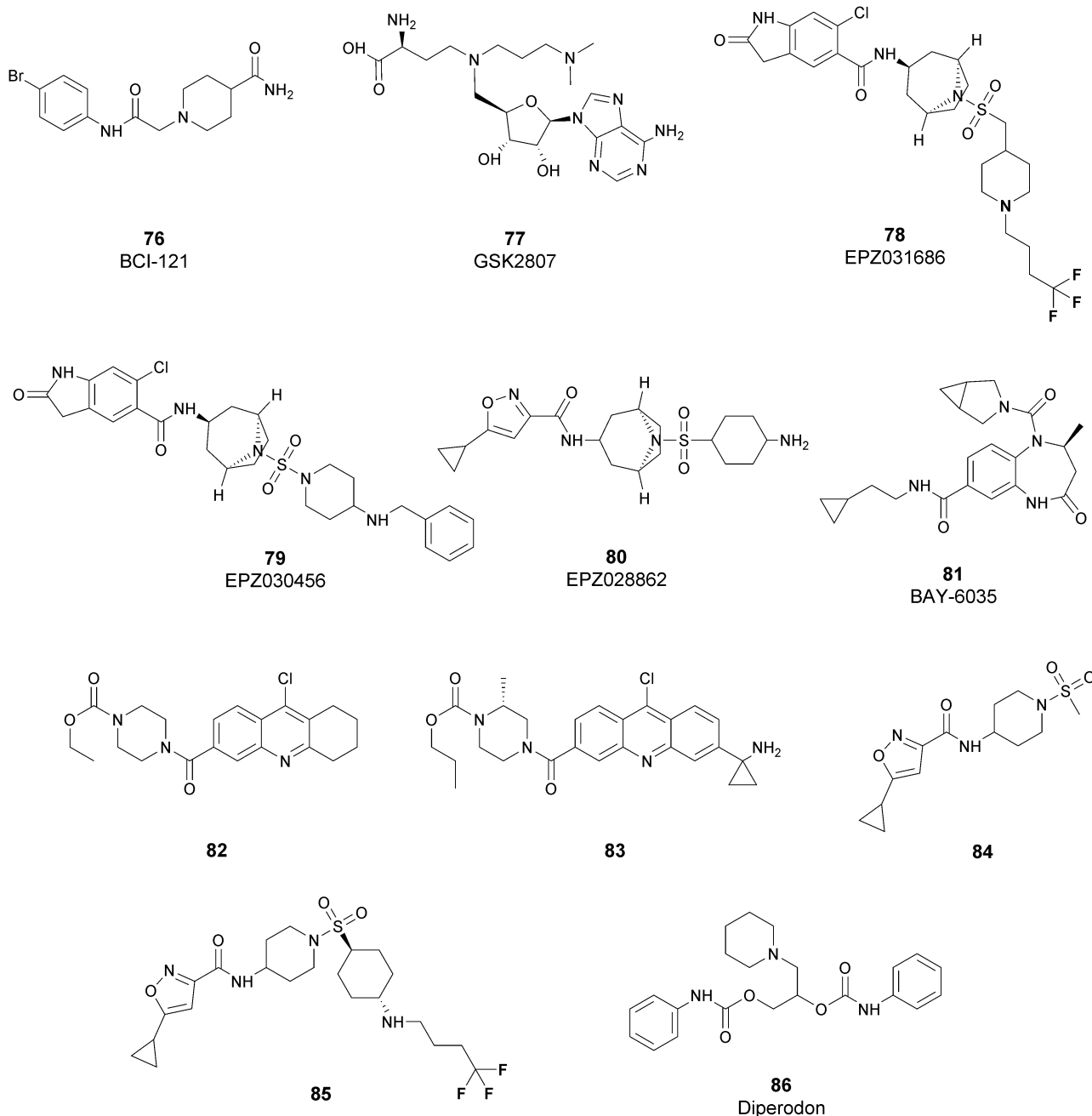


Fig. 20 SMYD3 inhibitors.

cellular (IC_{50} of 36 nM) assays, which also confirmed it to be orally bioavailable and suitable for *in vivo* analyses. The treatment of HCT116 and HT29 (Caucasian colon adenocarcinoma) cell lines with **76** at 100 μ M caused a substantial antiproliferative effect after 72 h (54 and 46% inhibition of cell growth, respectively).²⁰³

In 2016, starting from visual inspection of the ternary SMYD3-MEKK2 peptide-SAH complex, Van Aller and co-workers designed bi-substrate inhibitors. A series of aza-SAH analogues was synthesized and tested as SMYD3 inhibitors, focusing on the identification of a better side-chain length for good inhibition. The most potent (IC_{50} of 130 nM) derivative

was **77** (GSK2807), which is characterized by a propyl dimethylamine side chain, and is able to fit into the lysine channel anchoring the inhibitor in the pocket with a high degree of similarity with SAH-binding mode. The inhibition of methyltransferase activity by **77** was selective for SMYD3 toward a panel of 8 methyltransferases, and was 24-fold less active toward SMYD2. Despite these interesting data, the poor PK properties of this compound did not permit *in vivo* assays.²⁰⁴

In the same year, screening of the proprietary histone methyltransferase-biased library and subsequent hit optimization phases enabled Epizyme Inc. to develop **78** (EPZ031686) and **79** (EPZ030456). These derivatives showed a common oxindole



scaffold, capable of making several interactions with amino acid residues in the lysine channel of SMYD3. **78**, whose 4,4,4-trifluorobutyl chain is responsible for increasing of the lipophilicity of the molecule, showed strong activity both in biochemical (IC_{50} of 3 nM) and cellular (IC_{50} of 36 nM) assays. In the same way, the insertion of a sulphonamide portion allowed the identification of **79**, which was characterized by a nanomolar inhibitory activity in biochemical (IC_{50} of 4 nM) and cellular (IC_{50} of 48 nM) assays. Both derivatives were selective SMYD3 inhibitors, even if the low solubility of **79** troubled its *in vivo* testing.²⁰⁵

Two years later, after a screening campaign on isoxazole sulphonamides, the same company identified **80** (EPZ028862), which was active both in biochemical (IC_{50} of 1.3 nM) and cellular (IC_{50} of 32 nM) assays with satisfactory PK properties for further *in vivo* studies using different animal models. This derivative was a selective SMYD3 inhibitor that is able to reduce methylation marks on MEKK2 in 293T cells co-transfected with SMYD3 and MEKK2. The crystal structure of **80** in complex with the protein (PDB 5V37) disclosed a binding mode similar to **79**, which is probably due to the fact that both compounds are characterized by a common azabicyclo-octanyl moiety, as well as a sulphonamide group, and is able to involve different protein moieties. Moreover, as shown in Fig. 21, the acetamide motif is responsible for the interactions with both Thr184 and Tyr239 residues of the binding pocket. Despite the good activity profile, no antiproliferative activity was observed when tested in lung cancer cell lines, and no effect on cell growth was detected in a panel of other cancer cell lines.¹⁹²

In 2018, a collaboration between SGC and Bayer allowed the identification of **81** (BAY-6035), a substrate-competitive SMYD3

inhibitor with nanomolar IC_{50} values both *in vitro* (88 nM) and in cell-based (70 nM) assays toward the methylation of MEKK2. **81** is commercially available as a chemical probe for SMYD3, even if the corresponding publication has not been yet released.

Starting from a HTS campaign of 503 954 compounds, in 2019 Huang and co-workers identified the tetrahydroacridine derivative **82** (IC_{50} of 4.78 μ M) as a reasonable chemical starting point for the development of new SMYD3 inhibitors. Subsequent SAR studies and further optimization resulted in the discovery of a new class of 4-chloroacridines that are able to covalently modify SMYD3 *via* a nucleophilic aromatic substitution reaction at position 4. Among them, compound **83** showed a high inhibitory potency towards the enzymatic activity of SMYD3 (IC_{50} of 11.7 nM), as well as antiproliferative activity against the HepG2 human liver cancer cell line in 3D cell culture.²⁰⁶

Recently, a new HTS within the GSK proprietary compounds collection resulted in the identification of compound **84**, an isoxazole amide related to **80**, which is characterized by a lower biochemical potency (IC_{50} of 5 μ M *vs.* 1.3 nM for **84** and **80**, respectively) but has an interesting ligand efficiency (LE of 0.35) and PK properties. Modifications of the HTS hit on the basis of the structure-guided SAR study resulted in the identification of compound **85**, a potent (IC_{50} of 6.3 and 44 nM in biochemical and cellular MEKK2me assays, respectively), selective and orally bioavailable small-molecule inhibitor of SMYD3, and is considered a useful tool to explore the SMYD3 function in cancer as well as other disorders.²⁰⁷

Very recently, Talibov and co-workers identified **86** (diperodon) as an allosteric SMYD3 ligand. The *R*- and *S*-enantiomers showed a K_D of 42 and 84 μ M, respectively, and co-crystallization studies disclosed that both compounds were able to bind to an allosteric site in the C-terminal protein binding domain, representing a novel starting point for the design of new compounds that interact with a new druggable allosteric site.²⁰⁸

SETD2

SETD2 (also known as KMT3A and HYPB) was first isolated from human hemopoietic stem cells, and it represents a unique protein since it is the sole methyltransferase that is responsible for the trimethylation of H3K36.²⁰⁹ SETD2 has a high homology with the yeast protein SET2, which was previously identified in *S. cerevisiae* and currently SET2 is the sole nucleosomal H3K36 methyltransferase reported in yeast (responsible for all the methylation states of H3K36), where it acts as a transcriptional repressor.²¹⁰ Besides H3K36, two non-histone substrates of SETD2 are mainly reported, which are α -tubulin (methylated at K40)²¹¹ and STAT1 (methylated at K525).²¹² Nevertheless, in a recent study, other additional non-histone peptide substrates were identified and nine of them showed strong methylation at the target lysine.²¹³

From the structural point of view, SETD2 is characterized by the presence of different functional domains: the methyltransferase activity domains (AWS, SET and post-SET domains), the protein-binding domains (WW (tryptophan-tryptophan), CC (coiled-coiled), SRI (Set2-Rpb1 interacting)) and other domains with a function that is still unknown.^{214,215}

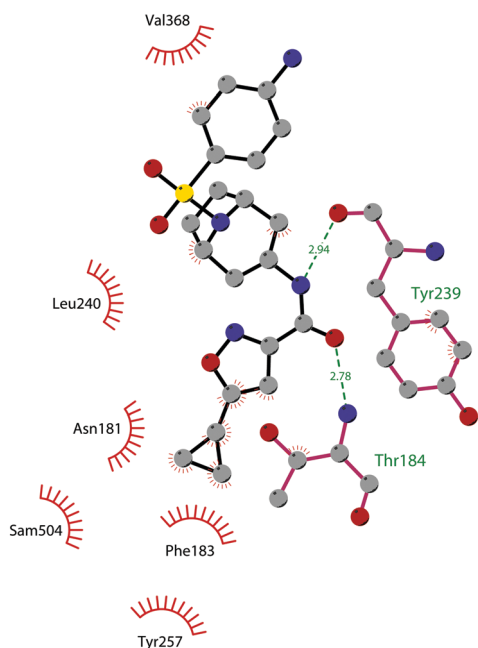


Fig. 21 Flat representation (generated using LigPlot+ from PDB ID 5V37) showing contacts between compound **80** and active-site residues in SMYD3. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with the compound.



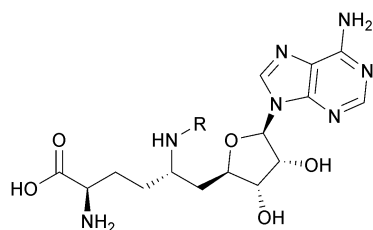
SETD2 is involved in different important cellular processes, such as transcriptional regulation, DNA damage repair and cytoskeletal remodelling. Regarding transcriptional regulation, SETD2, exploiting its SRI domain, binds to the hyperphosphorylated C terminal domain of RNA polymerase II, and it trimethylates H3K36 allowing the recruitment of reader proteins (MRG15, PTB) during transcriptional processes.²¹⁶ SETD2 contributes in maintaining the microtubule stability *via* the methylation of α -tubulin, highlighting the cross-talk between the epigenome and the cytoskeleton.²¹¹

From the pathological point of view, it has been reported that mutations in the sequence or aberrations in the SETD2 activity are linked to renal cancer,²¹⁷ glioma²¹⁸ and leukaemia.²¹⁹ In particular, these alterations can lead to tumorigenesis, the progression of cancer and unfavourable prognosis, even if the exact mechanism is still to be clarified.^{220–222}

SETD2 inhibitors

So far, the only known inhibitors of SETD2 are **sinefungin** derivatives (Fig. 22). With the aim of identifying target-specific methyltransferase inhibitors, in 2012 Zheng and co-workers carried out screening against a panel of methyltransferases of *N*-alkyl **sinefungin** analogues as privileged small-molecule scaffolds. *N*-Propyl **sinefungin** (**87**, Pr-SNF) and *N*-benzyl **sinefungin** (**88**, Bn-SNF) were identified as SAM-competitive inhibitors of human SETD2-specific inhibitors with definite potency and selectivity. **87** and **88** exhibited an IC₅₀ of 0.80 and 0.48 μ M, respectively, a value that is 10-fold higher than that of **sinefungin** itself and displayed 2/200-fold and 10/100-fold preferences, respectively, for SETD2 over other examined methyltransferases. Specifically, **87** was shown to preferentially interact with SETD2 by matching the distinct transition-state features of SETD2's catalytically active conformer, locating its secondary amine at the substrate-cofactor interface and the *N*-alkyl chain in the lysine-binding pocket.²²³

In fact, as depicted in Fig. 23, the SETD2–**87** co-crystal structure (PDB 4FMU) shows that the *N*-propyl moiety of the inhibitor is buried in a hydrophobic binding pocket formed by Tyr1579, Tyr1605, Phe1664, and Tyr1666. On the other hand, a large number of hydrogen bonds are also detected, engaging the hydroxy groups of the ribose portion and His1603, Tyr1604 and Gln1676. Despite their specificity, the activity of **87** and **88**



87, Pr-SNF: R = *n*-Pr

88, Bn-SNF: R = -Bn

Fig. 22 SETD2 inhibitors.

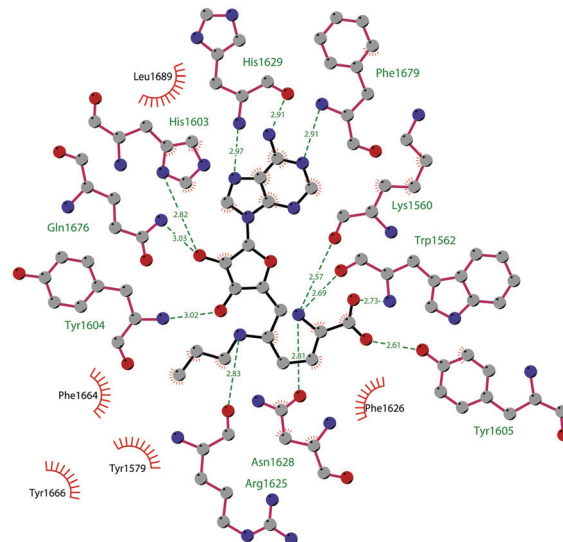


Fig. 23 Flat representation (generated using LigPlot+ from PDB ID 4FMU) showing contacts between compound **87** and active-site residues in SETD2. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with the compound.

is influenced by the peptide substrate, and their poor membrane permeability prevented in cell or *in vivo* assays.

SETD8

SET domain-containing lysine methyltransferase 8 (also called PR-SET7 or KMT5A) is a member of the SET domain-containing family and is characterized by a unique activity as it is the only reported mammalian protein responsible for the monomethylation of H4K20 *in vivo*. SETD8 preferentially monomethylates nucleosomes instead of histones, suggesting this protein is able to interact with other regions of the nucleosome as well as the H4 N-terminal region surrounding the residue H4K20.²³ These multivalent interactions of SETD8 involve both the SET domain and a basic region N-terminal to the SET domain.^{224,225}

As with other lysine methyltransferases, SETD8 methylates not only histones but also other proteins, including p53, Numb and PCNA. Monomethylation of p53 K382 by SETD8 triggers the suppression of p53-dependent transcription activation in cancer cells.²²⁶ Numb is methylated by SETD8 on the K158 and K163 residues, and this methylation disrupts the complex Numb–p53–MDM2, which is responsible for the promotion of apoptosis. Hence, the methylation of Numb prevents apoptosis and finally results in the ubiquitination and degradation of p53.²²⁷ The monomethylation of K248 of proliferating cell nuclear antigen (PCNA) (PCNAK248me1) enhances its interaction with the flap endonuclease FEN1, promoting carcinogenesis by deregulating the PCNA expression.²²⁸

Several studies have provided evidence that SETD8 may be involved in many cellular processes, such as DNA replication, DNA damage response, transcription modulation, and cell cycle regulation as well as in carcinogenesis and cancer invasiveness.^{229–231} SETD8 is overexpressed in different types of cancer tissues including bladder cancer, non-small-cell and



small-cell lung carcinoma, chronic myelogenous leukaemia, hepatocellular carcinoma, pancreatic and breast cancer.^{232,233}

SETD8 inhibitors

To better elucidate the SETD8 biology as well as its effects in both normal and diseased cells, some inhibitors have been developed (Fig. 24), even if just a few of them displayed a certain degree of selectivity and/or cellular activity.²²

The screening of a library of marine natural products allowed the identification of the polyketide **89a** (nahuic acid A), a decalin polyketide isolated from cultures of *Streptomyces* sp. obtained from a marine sediment collected in Papua New Guinea. **89a** inhibited SETD8 activity with an IC_{50} value of 6.5 μ M, and with no significant effect on other tested enzymes. Cellular treatment with **89a** led to a decrease in the levels of both mono- and trimethylated H4K20 without affecting other methylated histone marks.^{234,235} Exposure of U2OS cells to **89a** and its prodrug version (nahuic acid A pentaacetate, **89b**), prepared to improve the cellular permeability, resulted in a modest decrease of cellular proliferation (IC_{50} of 65 μ M and 39 μ M, respectively). The same results were obtained when treating breast cancer cell lines such as SUM159 and MDA-MB-436.

In 2014, Luo and co-workers reported three small molecules (**90–92**) as SETD8 inhibitors, which were identified *via* the HTS of a library containing more than 5000 commercially available compounds. These derivatives differ in their mechanisms for SETD8 inhibition: **90** (SPS8I1, IC_{50} of 0.21 μ M) turned out to be a substrate-dependent inhibitor, **91** (SPS8I2, IC_{50} of 0.5 μ M) exhibited no substrate or SAM dependence, while **92** (SPS8I3, IC_{50} of 0.5 μ M) was found to be both a substrate- and SAM-dependent inhibitor. In all cases, the quinonic motif, common to all inhibitors, interacts with active cysteine residues (Cys270) in SETD8 through an irreversible slow-onset process.

According to SETD8 knockdown, these compounds produced a cell cycle arrest phenotype and reduced the monomethylation of H4K20. Unfortunately, their low selectivity against other protein methyltransferases and other cellular targets prevents their further development.²³⁶

In the same year, Jin and co-workers reported the identification of **93** (UNC0379) *via* a cross-screening of a library containing more than 150 quinazoline-based compounds, the same scaffold previously explored for G9a ligands. In a radioactive methyl transfer assay, **93** showed an IC_{50} value of 7.3 μ M. This compound showed no appreciable activity on 14 other methyltransferases tested, with IC_{50} values above 100 μ M also for G9a and GLP. By contrast, compound **93** inhibited PRC2 only at the two highest concentrations (50 and 100 μ M), showing an IC_{50} value of more than 50 μ M. A SAR study was undertaken on the 4-amino group of the quinazoline scaffold, showing that reducing the length of the alkyl linker between the two amino groups resulted in a lower potency. Moreover, it was shown that the basicity of the pyrrolidinyl group was critical for the SETD8 inhibitory activity. Regarding the mode of action, **93** acts as a substrate-competitive inhibitor, non-competing with the cofactor SAM, showing a good selectivity over other methyltransferases.²³⁷ Recently, it has been shown that the pharmacological inhibition of SETD8 by **93** led to cell-cycle defects and apoptosis in a p53-dependent or -independent manner. In particular, SETD8 inhibition in multiple myeloma (MM) cell lines induced by **93** led to cell death activating the p53 canonical pathway. However, **93**-induced cytotoxicity in MM cells is not necessarily dependent on p53 activation but is mainly caused by cellular stress during G1 progression. Moreover, a low dose of **93** co-administered with the cytotoxic agent melphalan strongly increases DNA damage and overcomes drug resistance in both p53 wild-type and deficient MM cells.

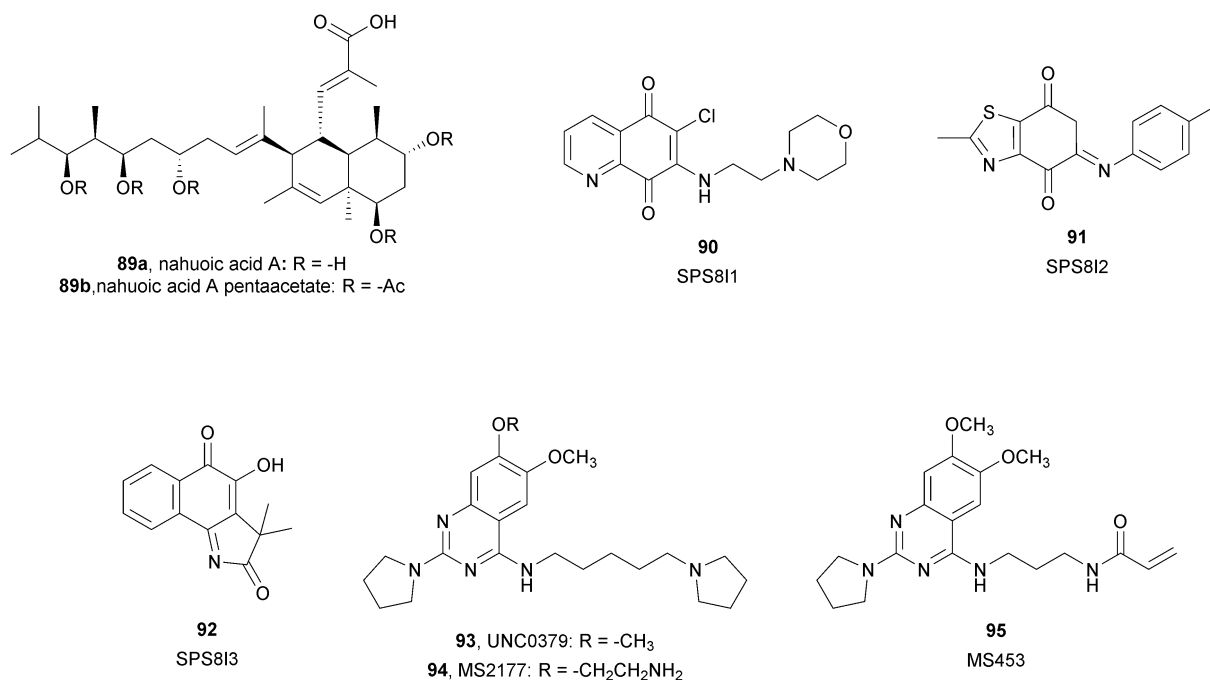


Fig. 24 SETD8 inhibitors.



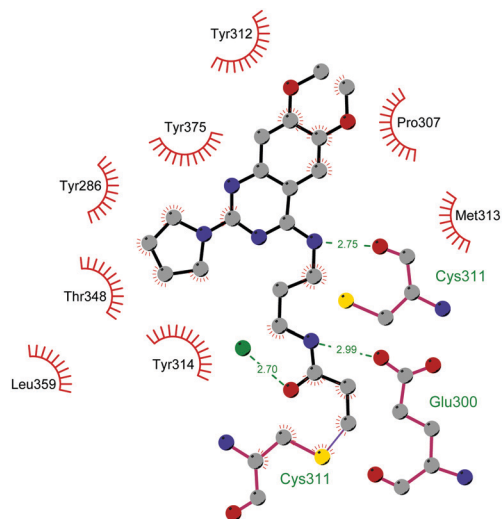


Fig. 25 Flat representation (generated using LigPlot+ from PDB ID 5TH7) showing contacts between compound **95** with active-site residues in SETD8. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with compound.

This evidence suggested that targeting SETD8 could be an alternative therapeutic strategy in high-risk patients with multiple myeloma independently of their mutational status for p53.²³⁸

In 2016, a higher potent inhibitor, **94** (**MS2177**), was identified by inserting an aminoethyl group as a side chain on **93**. Indeed, the resulting compound **94** had an *in vitro* IC₅₀ of 1.9 μ M and a much lower affinity in isothermal titration calorimetry (ITC) experiments compared with **93** (K_D of 1.3 μ M vs. 18 μ M). The compound was found to be competitive with the substrate but not with the co-factor SAM.²³⁹ In the same study, a potent covalent SETD8 inhibitor, **95** (**MS453**), was identified by inserting into compound **93** an acrylamide group as the electrophilic warhead, and which is responsible for the reaction with a surface-exposed Cys311 in the active site. The crystal structure of **95** in complex with SETD8 (PDB 5TH7) showed the formation of a homodimer in which each covalently modified Cys311 residue locates the quinazoline moiety to its binding site of the other complementary subunit. As illustrated in Fig. 25, the compound establishes a hydrogen bond between the secondary amino group in the 4-position and the carbonyl oxygen of Cys311 on the peptide backbone. Moreover, **95** makes solvent-mediated hydrogen bonds between the quinazoline N1 nitrogen and the carbonyl groups on the protein backbone, inserting the 2-pyrrolidine motif into a hydrophobic pocket induced by the binding of the inhibitor to the protein.

95 showed an IC₅₀ value of 804 nM with a good selectivity over 28 other methyltransferases. Unfortunately, no significant reduction in the methylated H4K20 levels was observed from treating HeLa cells with the inhibitor (10 μ M), probably due to its poor cellular permeability.²³⁹

SUV420H1 and SUV420H2

SUV420H1 (KMT5B) and SUV420H2 (KMT5C) are two highly homologous enzymes in mammals (65% sequence identity in the

catalytic domain). Although both enzymes can use as a substrate the unmodified H4K20, they prefer the monomethylated H4K20 as a substrate *in vivo* and exploit the SETD8-generated monomethylated H4K20 to produce di- and trimethylated H4K20.²⁴⁰ In particular, it has been reported that, in mouse embryonic fibroblasts, SUV420H1 catalyses the formation of most of the dimethylated H4K20 while SUV420H2 is responsible for trimethylated H4K20. The catalytic domains of SUV420H1 and SUV420H2 consist of three subdomains: an N-terminal helical domain, a SET domain and a Zn-binding post-SET domain. SUV420H1 has a bigger N-domain than SUV420H2 due to a 22-residue insertion between the first and second helices.²⁴¹

SUV420H1 and SUV420H2 proteins play critical roles in the chromatin structure, genomic integrity and in the regulation of the cell cycle. Indeed, both these proteins are linked to transcriptional repression, DNA replication and repair.^{242,243} Therefore, the loss of both of them is accompanied with chromosomal aberrations, deficiencies in DNA double-strand break repair and reduction in telomere elongation, together with a decrease in proliferation rates and in cell cycle progression.^{230,243–245} SUV420H1 and SUV420H2 are pivotal regulators of normal embryonic development and in self-renewal or lineage-commitment of stem cell populations. Indeed, the loss of SUV420H1/2 results in perinatal lethality. Alterations in the expression and mutations of SUV420 enzymes are related to the insurgence of cancer. Regarding SUV420H2, it has been reported that in breast cancer and in osteosarcoma there is a reduction in the levels of trimethylated H4K20, together with a decrease in the expression of SUV420H2, which is associated with invasiveness and poor prognosis.²⁴⁶ However, in pancreatic cancer, SUV420H2 is overexpressed and it is associated with invasive cancer. In particular, it has been reported that this protein behaves as a regulator of epithelial/mesenchymal state control and its high levels promote the epithelial-to-mesenchymal transition, which is related to metastasis.²⁴⁷ Regarding SUV420H1, it has been reported that this protein is overexpressed and mutated in different cancers. On the basis of the type of mutation, different outcomes in cancer have been observed, where the protein can assume a tumour suppressor or a tumorigenic role.²⁴⁸ Hence, we can conclude that both SUV420 enzymes have multiple and divergent roles in cancer that need to be fully elucidated.

SUV420H1 and SUV420H2 inhibitors

Although the identification of modulators of these enzymes may be essential for understanding their function, so far only three inhibitors have been reported (Fig. 26). In 2017, Bromberg and co-workers developed a high-throughput homogeneous activity-based scintillation proximity assay with the aim of identifying SUV420 inhibitors. A library of approximately 470 000 compounds was screened, and among them **96** (**A-196**) was described as a first-in-class chemical probe of both SUV420 enzymes and a good starting point for the development of new inhibitors. Co-crystallization and biochemical analyses confirmed that **96** is a substrate-competitive inhibitor of SUV420H1 and SUV420H2 with IC₅₀ values of 25 and 144 nM, respectively.



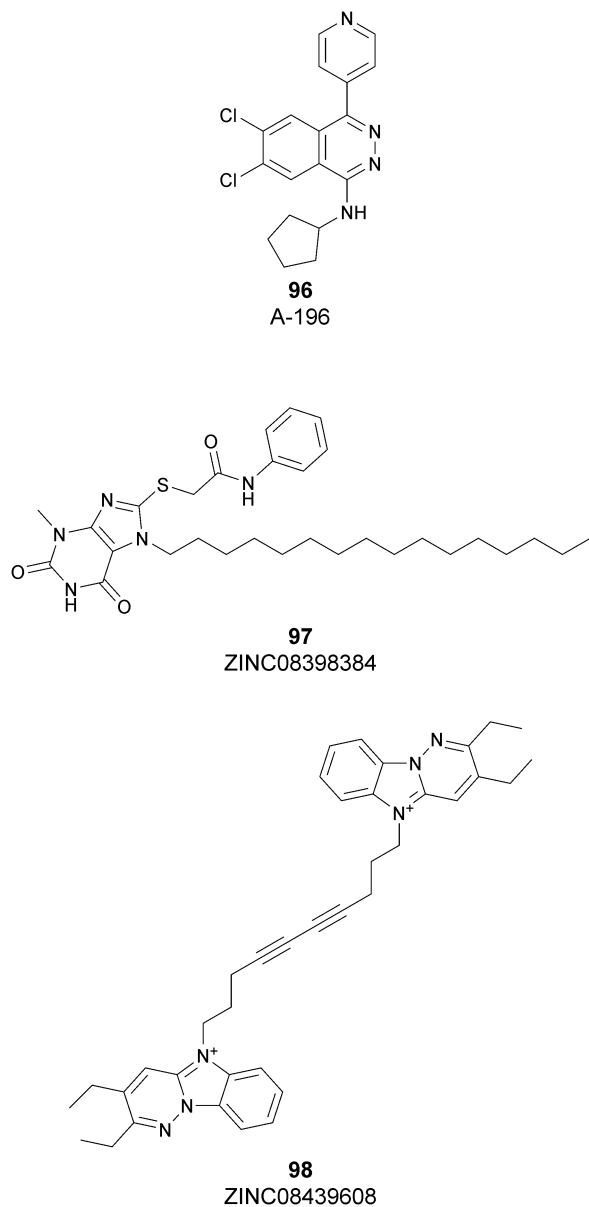


Fig. 26 SUV420H1 and SUV420H2 inhibitors.

Examination of the crystal structure of **96** bound to SUV420H1 (PDB 5CPR) revealed that the cyclopentyl moiety is able to occupy a hydrophobic pocket made by Ser251, Val252, Tyr307 and Met253, which could be further explored. On the other hand, the pyridyl unit is located in a solvent-exposed region (Fig. 27). A further global decrease in di- and trimethylated H4K20 and a simultaneous increase in monomethylated H4K20 were found after treatment of the prostate adenocarcinoma cell line PC-3 or U2OS osteosarcoma cells with **96**, without any toxic effects.

Furthermore, this inhibitor does not induce resistance/compensation effects, as di- and trimethylated H4K20 remained low after 20 population doublings.²⁴⁹

Recently, a HT and SBVS combined with molecular dynamics simulation has been used by Zheng and co-workers in order to find novel potential SUV420H1 inhibitors. Starting

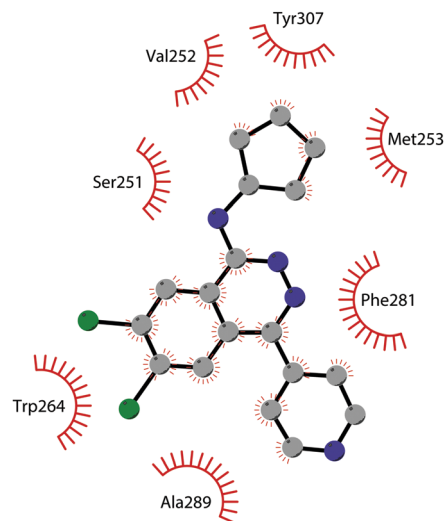


Fig. 27 Flat representation (generated using LigPlot+ from PDB ID 5CPR) showing contacts between compound **96** and active-site residues in SUV420H1. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with the compound.

from the Specs subset of the ZINC database, containing over 200 000 compounds, **97** (ZINC08398384) and **98** (ZINC08439608) were selected and showed an inhibitory effect similar to the known inhibitor **96**, effectively reducing the expression level of dimethylated H4K20 in U2OS (at 1.25 and 2.5 μ M, respectively) and attenuating the migration of osteosarcoma cells *in vitro*.²⁵⁰

NSD1, NSD2, NSD3

The NSD (Nuclear receptor SET Domain-containing) family is a phylogenetically distinct family of methyltransferases comprised of NSD1, NSD2 (MMSET/WHSC1) and NSD3 (WHSC1L1), three proteins with pivotal roles in chromatin regulation. These proteins share the presence of a catalytic SET domain (with a high homology with the SETD2 protein) and multiple conserved domains, such as proline-tryptophan-tryptophan-proline (PWWP) and the plant homeodomain (PHD) zinc finger domains, which are involved in the interaction with DNA and other methylated histones.²⁵¹

Regarding their specific substrates, NSD proteins catalyse the methylation of several lysine marks on histones H3 and H4.²⁵² In particular, NSD1 dimethylates H3K36 and H4K20, whereas NSD2 methylates H4K20, H3K4, H3K27 and H3K36, and NSD3 methylates H3K4 and H3K27.^{253,254}

NSD proteins play essential roles in embryonic development, and it has been shown that the deletion of NSD1 protein causes problems in the development of post-implantation embryos.²⁵⁵ Moreover, the dysregulation of NSD proteins in human cells affects cell growth and differentiation and has been related to several diseases, such as mental retardation and developmental defects that are common symptoms of Sotos syndrome^{256,257} and Wolf-Hirschhorn syndrome.^{258,259} In addition, aberrant activity of NSD proteins, as well as their overexpression, has been linked to the insurgence of neoplastic diseases such as



breast carcinoma, acute myeloid leukaemia, and multiple myeloma.^{260–262}

NSD1, NSD2, NSD3 inhibitors

Although NSD proteins are appealing therapeutic targets, a limited number of small-molecule inhibitors have been described so far (Fig. 28). Based on the homology of the catalytic SET domain of the NSDs to that of G9a-like protein, in 2017 the G9a/GLP inhibitor **22** (Fig. 8) was tested on NSD proteins, showing an *in vitro* inhibitory activity on all these

targets and lowering monomethylated H3K36 levels. Specifically, **22** revealed the highest inhibition on NSD2 (IC₅₀ of 41 μM), followed by NSD3 (IC₅₀ of 95 μM), and finally NSD1 (IC₅₀ of 112 μM). This different activity has been associated with the diverse binding modes of the compound towards the catalytic portion of the three proteins. Specifically, **22** is able to shallowly bind into the lysine channel of NSD1; conversely, in NSD2, the diazepane side of the molecule fits into the lysine channel; in NSD3 the dimethoxy quinazoline group spreads deeply into the lysine channel, like in GLP. Furthermore, the better IC₅₀ value observed against NSD2 is probably due to a tighter hydrogen-bond network in the NSD2 active site compared with NSD1 and NSD3.²⁶³

Recently, an NMR fragment-based screening approach, followed by chemical optimization, resulted in the identification of **99** (**BT5**), the first-in-class irreversible small-molecule inhibitor of NSD1, whose electrophilic methyl-aziridine moiety is responsible for the covalent binding to the C2062 residue of the NSD1 SET domain. The crystal structure of NSD1 in complex with **99** disclosed a conformational modification in the autoinhibitory loop of the SET domain, leading to the formation of a unique, channel-like site that is suitable for targeting with small molecules. **99** inhibits NSD1 at low micromolar concentrations (IC₅₀ of 5.8 μM upon 4 h incubation, which improved to IC₅₀ of 1.4 μM after 16 h due to irreversible binding). NSD2 and NSD3 inhibition was also observed, although non-covalent and at higher concentrations. The cellular thermal shift assay confirmed that **99** is able to bind the NSD1 SET domain in cells, generating a global reduction of dimethylated H3K36 levels without perturbing trimethylated H3K36.²⁶⁴

The natural product **sinfungin 17** (Fig. 6) and the *N*-alkyl **sinfungin** derivative **87** (Fig. 22) have been identified as NSD2 inhibitors (IC₅₀ of 26 μM and 3.3 μM, respectively), but inhibition data on NSD1 and NSD3 have not been reported.²⁶⁵ In 2019, de Luccio and co-workers by VS identified **100** (**LEM-14**). This inhibitor showed an *in vitro* IC₅₀ of 132 μM against H3K36 methylation by NSD2, a very weak activity against NSD1, and no activity against NSD3, representing an interesting tool for studying the biology of NSD2.²⁶⁶

In 2021, a VS followed by a ligand-based scaffold-hopping approach, enabled the identification of **101**, a small-molecule antagonist that blocks the protein-protein interaction between the N-terminal PWWP domain of NSD2 (PWWP1) and dimethylated H3K36 with a *K_D* of 3.4 μM.²⁶⁷ The crystal structure of **101** in complex with NSD2 and subsequent molecular docking simulations enabled the development of the more potent **102** (**MRT866**), which is capable of binding NSD2–PWWP1 with a *K_D* of 349 nM. Further structure-based optimization resulted in **103** (**UNC6934**), which showed potent binding to PWWP1 (*K_D* of 91 nM).²⁶⁸ In order to better understand the binding mode of **103**, a representation of the main interaction is shown in Fig. 29. The crystal structure of the inhibitor in complex with NSD2–PWWP1 (PDB 6XCG) shows how the cyclopropyl moiety is inserted in a hydrophobic pocket made of Val230 and Phe266. Moreover, the amidic NH is involved in a hydrogen bond with Gln321, while the carbonyl oxygen establishes a strong interaction with Ala270.

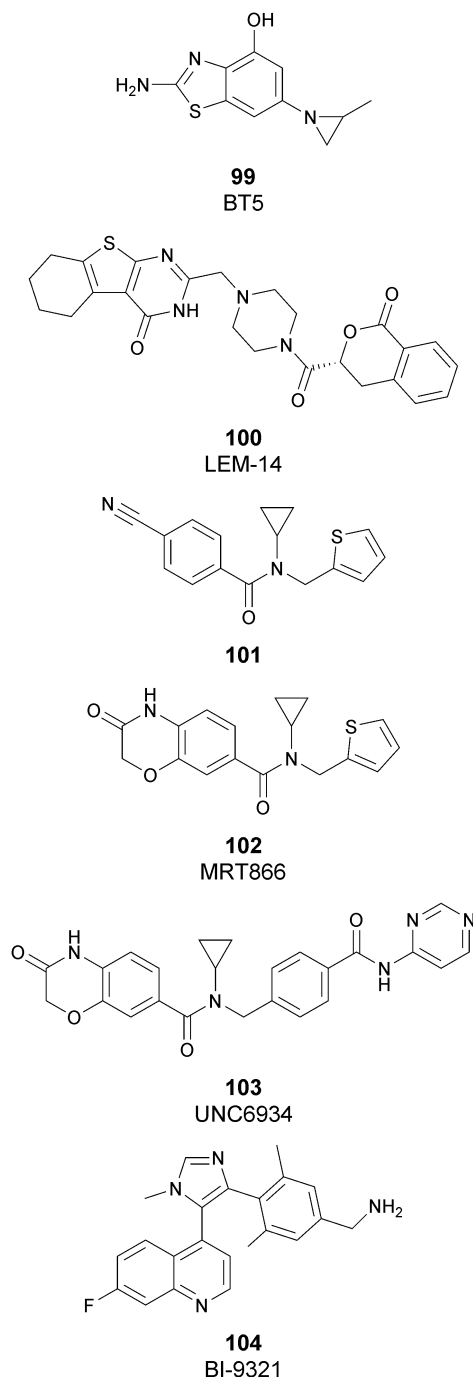


Fig. 28 NSD1–3 inhibitors.



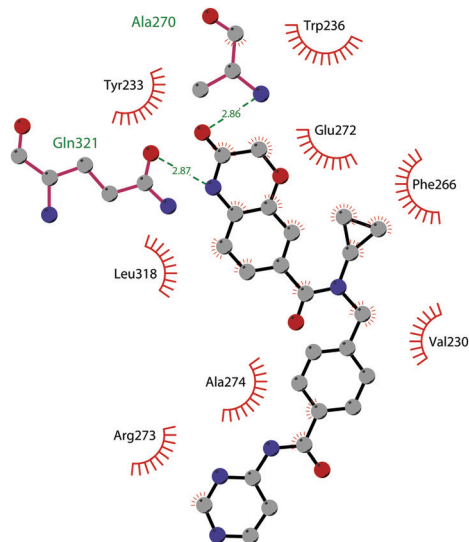


Fig. 29 Flat representation (generated using LigPlot+ from PDB ID 6XCG) showing contacts between compound **103** and active-site residues in NSD2–PWWP1. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with the compound.

In 2019 Mc Connell and co-workers reported the fragment-based virtual screening discovery of **104** (**BI-9321**), a strong and selective antagonist of the NSD3–PWWP1 domain. This compound showed the capability of targeting the methyl-lysine binding site of the PWWP1 domain with an IC_{50} of 203 nM and a K_D of 166 nM. Moreover, **104** was able to downregulate Myc messenger RNA expression, as well as reduce cellular proliferation in the MOLM-13 human leukaemia cell line.²⁶⁹ The co-crystal structure of NSD3–PWWP1 in complex with **104** (PDB 6G2O) showed the importance of the aminic group in the binding mode of the inhibitor. In fact, as depicted in Fig. 30,

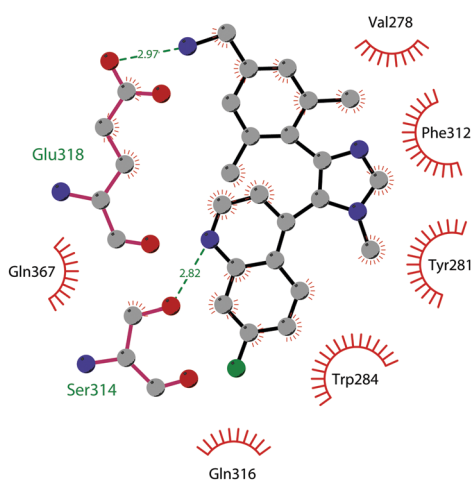


Fig. 30 Flat representation (generated using LigPlot+ from PDB ID 6G2O) showing contacts between compound **104** and active-site residues in NSD3–PWWP1. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with the compound.

this motif is responsible for a hydrogen bond with Glu318, while the pyridinic nitrogen is involved in an interaction with the Ser314 residue of the binding pocket.

MLL proteins

The family of lysine methyltransferase KMT2 is composed of SETD1A, SETD1B and four mixed-lineage leukaemia proteins (MLL 1,2,3, and 4) that play pivotal roles in the regulation of gene expression during embryogenesis and development. These proteins catalyse the methylation of H3K4,²⁷⁰ and they exist in nuclear complexes (human COMPASS), which include ASH2L, RbBP5, Cfp1, WDR5 and Dpy30. The components of the complex greatly enhance the enzymatic activity of MLL methyltransferases.^{271,272}

Structurally, MLL proteins are characterized by a peculiar domain architecture: in addition to the C-terminal SET domain, all the proteins of the family feature a different number of DNA-binding and protein–protein interaction domains, which are exploited for binding to specific interaction partners. Despite the structural homology in the organization of the domains, each MLL protein differs from the others on the basis of the type, number and position of each specific domain. In particular, there is high variability in the number of PHD finger domains: four for MLL1 and 2, eight for MLL3, and seven for MLL4.²⁷³ In addition, MLL1 and MLL2 display the CXXC zinc-finger domain for the interaction with unmethylated CpG islands, and this domain is absent in the other members of the family.²⁷⁴ The structural differences in MLL proteins are reflected in their enzymatic activity: for instance, while MLL1 and 2 are responsible for both the mono- and dimethylation of H3K4, proteins MLL3 and MLL4 have been reported as monomethyltransferases.^{275,276}

Many studies have shown that alterations in the expression of MLL genes are linked to the insurgence of diseases. Beside the involvement of MLL proteins in the pathogenesis of leukaemia, for instance through the interaction with menin (which lead to myeloid transformation),²⁷⁷ a huge number of somatic mutations for MLL proteins have been identified in specific solid cancer types, mainly of the lung, endometrium, and large intestine.^{278–280}

MLL protein inhibitors

A deeper understanding of the biological role and function of MLL proteins could be triggered by the discovery of potent chemical probes that selectively target them. So far, only a small SAM-based focused chemical library has been designed with the aim of directly targeting the active site of MLL proteins. Among them, compound **105** (**TC-5115**, Fig. 31) has emerged as the best candidate (IC_{50} of 16 nM). Its co-crystal structure revealed that the compound is positioned in the cofactor binding site, while the azetidine group induces the “open” conformation of the substrate binding pocket, thus allowing additional interactions between the inhibitor and the MLL SET domain.²⁸¹

On the other hand, efforts made to target protein–protein interactions in the MLL core complex have been more



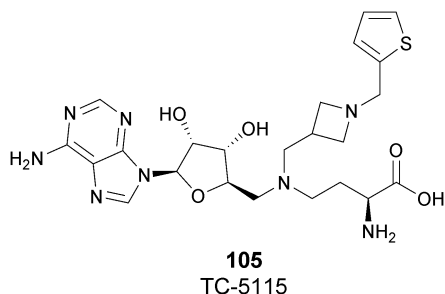


Fig. 31 Structure of SAM-based MLLs inhibitor.

successful. Specifically, disruptors of WDR5–MLL (Fig. 32) as well as menin–MLL interactions (Fig. 34) have been developed.

WDR5–MLL interaction disruptors

The central role of WDR5 in the maintenance of integrity and the methyltransferase activity of MLL complexes has suggested that disrupting the WDR5–MLL interaction is a possible strategy to develop inhibitors of MLL methyltransferases. The identification of a conserved arginine-containing motif within MLL, *i.e.*, the WIN (WDR5-interacting) sequence, is responsible for the association with WDR5 and, consequently, is essential for the enzymatic activity of MLL1²⁸² and represents a good starting point from which to design disruptors of WDR5–MLL interactions.

Karatas and co-workers reported the design of peptidomimetics to disrupt WDR5–MLL interactions. Specifically, the identification of the minimum motif responsible for the high-affinity binding of MLL1 to WDR5 (–CO-ARA-NH–), led to the design of two 3-mer peptides (Ac-ARA-NH₂ and Ac-ART-NH₂) as efficient WDR5 binders with K_i values of 120 and 20 nM, respectively.²⁸³ Further extensive modifications enabled the identification of **106** (**MM-102**), a potent peptidomimetic compound (K_i of <1 nM) that mimics the WIN peptide for binding to WDR5. However, **106** exhibits a very modest cellular potency and poor metabolic stability in human microsomes.²⁸⁴ It was anticipated that the reduction of conformational flexibility could overcome these issues. Therefore, the linear structure of the parent compound was replaced with a macrocyclic equivalent, obtaining diverse compounds.^{285,286} Among them, derivative **107** (**MM-589**) binds to WDR5 with a very high affinity (K_i less than 1 nM), significantly inhibiting the MLL1 methyltransferase activity (IC_{50} of 12.7 nM). Moreover, **107** inhibits cell growth in the MOLM-13 and MV4-11 human AML cell lines that carry the MLL translocation with more than 30-fold selectivity over the HL-60 leukaemia cell line that lacks the MLL translocation.²⁸⁶ In a screening of commercially available compounds, **108** (**WDR5-0103**) was identified as a modest WDR5 binder (K_{dis} of 3.0 μ M in a peptide-displacement assay, and K_D of 0.45 μ M). The compound efficiently antagonizes the WDR5–WIN peptide interaction, negatively affecting the catalytic activity of MLL complexes *in vitro*.²⁸⁷ The X-ray crystal structure supported the optimization of **108**-based nitro analogues, facilitating the identification of **109** (**WDR5-47**), an improved WDR5 binder (K_{dis} of 0.3 μ M).²⁸⁸ Replacement of the nitro group with the 4-pyridinyl group led to **110** (**W-23**), which efficiently disrupted the MLL1–WDR5 interaction with an IC_{50} value of 104 nM.²⁸⁹

Further optimization yielded derivative **111** (**WL-15**), which better disrupted the interaction of MLL1–WDR5 (IC_{50} of 26.4 nM) and inhibiting the catalytic activity of the MLL1 complex (IC_{50} of 5.4 μ M). Unfortunately, the poor cellular permeability limited its cellular activity in inhibiting the proliferation of leukaemia cell lines with or without the MLL1 fusion protein.²⁹⁰

Based on the *N*-(2-(4-methylpiperazin-1-yl)phenyl)acetamide scaffold, other derivatives were designed as disruptors of MLL1–WDR5 interactions with increased cellular activity. Specifically, **112** (**DDO-2084**) inhibited MLL1 methyltransferase activity both *in vitro* (IC_{50} of 1.65 μ M) and in the MV4-11 cell line (IC_{50} of 17.7 μ M), showing a down-regulation of the MLL-target genes expression level. Insertion of the 4-aminobutanamido group on **112** improved the inhibitory activity: compound **113** (**DDO-2117**) exhibited an IC_{50} of 0.19 μ M and strongly inhibited the MLL1–WDR5 interaction (IC_{50} of 7.6 nM and K_D of 13.6 nM).²⁹¹

Additional optimization of the substituents in the above-mentioned scaffold led to the identification of **114** (**OICR-9429**), a compound that competitively disrupted the WDR5 interaction with the WIN motif of MLL (K_{dis} of 64 nM) with good WDR5 affinity (K_D of 30 and 93 nM in SPR and ITC assay, respectively). **114** is highly selective for WDR5 with no inhibition of 22 other human methyltransferases, 9 different WD40 domains, and other non-epigenetic targets. The crystal structure of WDR5 in complex with **114** helped to clarify its detailed binding mode (PDB 4QL1). The compound is inserted in an arginine-binding pocket of WDR5, precluding the interaction of the WIN peptide residue with MLL. Upon **114** binding, the protein undergoes conformational rearrangements to better accommodate the compound (Fig. 33). In particular, the Phe263 moved to lodge the *N*-methylpiperazine moiety in position of the WIN peptide guanidinium group, while the Phe133 and Phe149 residues moved to accommodate the opposite side of the molecule. Moreover, the piperazine, the amide linker and the pyridone group interact (directly or by water-mediated hydrogen bonds) with the surrounding residues of Cys261, Ser91 and Asp107, whereas the other side of the molecule interacts with a hydrophobic pocket in WDR5 formed by the Phe133, Tyr191 and Pro173 residues. Critical for compound binding is the interaction with Ser91. Notably, the morpholino ring is solvated, lacking a clear electron density, suggesting that it can assume different conformations. The efficacy and target selectivity of **114** were also confirmed in cells in which the oncogenic transcription factor p-30 drives cell growth in a WDR5–MLL-dependent manner. These findings offer an attractive strategy for targeting cancer depending on the oncogenic transcription factor mutations.^{292,293}

In 2018, exploiting fragment-based methods and structure-based design, dihydro-pyrroloimidazole was disclosed as an effective WDR5 binder. Crystal structure analysis depicted the binding of this core to WDR5 in a pocket that normally accommodates the MLL1–peptide-R3765–guanidine side chain. The most potent inhibitor derivative, **115**, displayed a high potency (K_i of 0.9 nM) causing the dose-dependent inhibition of H3K4-methylation activity and a moderate growth reduction in the MV4-11 cell line.²⁹⁴



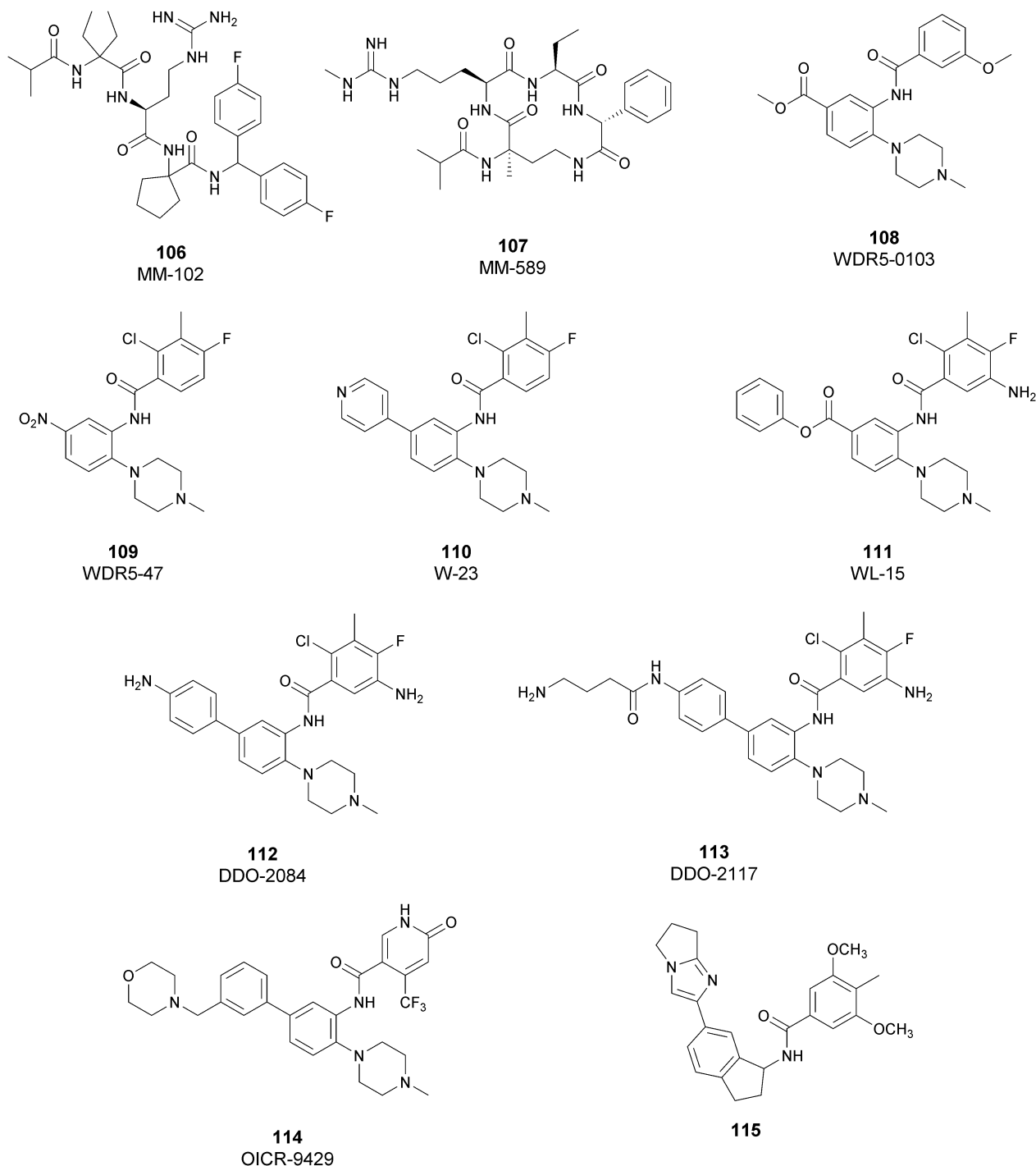


Fig. 32 WDR5–MLL interaction disruptors.

Menin–MLL interaction disruptors

Another therapeutic strategy explored to develop new drugs for MLL leukaemia patients is *via* disruption of the protein–protein interactions between MLL and the oncogenic cofactor menin, which are required for the initiation of MLL-mediated leukemogenesis as well as for regulating the expression of target genes, such as HOXA9 and MEIS1 genes, essential for the leukemogenic activity of MLL fusion proteins. Menin binds with two different fragments of MLL: the high-affinity menin binding motif (MBM1

with a K_D of 56 nM) and the low-affinity menin binding motif (MBM2 with a K_D of 1 μ M). Even if MBM2 contributes to binding to menin, deletion of the MBM1 motif is sufficient to abolish the activity of MLL oncoproteins.²⁹⁵ Therefore, small-molecule inhibitors have been developed that mainly target the MBM1 motif.

From a structural point of view, most of the MLL–menin inhibitors developed feature a thienopyrimidine core structure.

HTS and further medicinal chemistry optimization guided the identification of the parent compound of this class, **116** (**MI-2**).



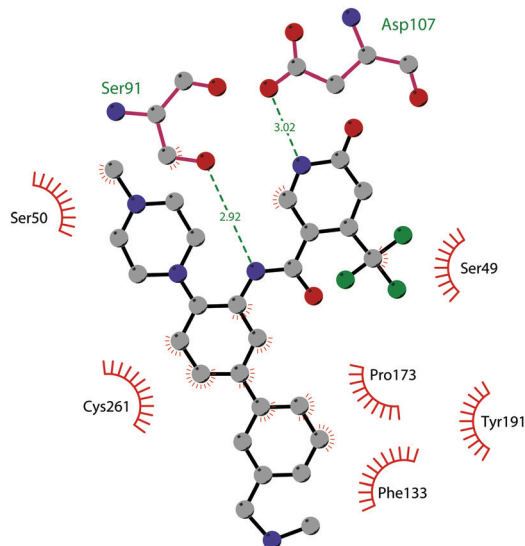


Fig. 33 Flat representation (generated using LigPlot+ from PDB ID 4QL1) showing contacts between compound **114** and active-site residues in WDR5. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with compound.

It inhibited the the menin–MLL interaction with an IC_{50} of 446 nM, disrupting the menin–MLL interaction both *in vitro* and in human cells. The treatment of different MLL leukaemia cell lines with **116** resulted in different effects, such as the downregulation of HOXA9 and MEIS1 expression, inhibition of proliferation and induction of apoptosis and differentiation.²⁹⁶ These pioneering results highlighted the potentiality of targeting menin–MLL interactions in acute leukaemia associated with MLL translocations and prompted the synthesis of several derivatives featuring the thienopyrimidine scaffold.

117 (MI-2-2) showed an 8-fold improved binding affinity toward menin (K_D of 22 nM) disrupting the MBM1–menin interaction with high affinity (IC_{50} of 46 nM) comparable to the fragment including both MBM1 and MBM2. In this way, the compound is able to fully dissociate the entire menin–MLL fusion protein complex.²⁹⁷ Despite the good cellular permeability, **117** showed modest activity in human MLL leukaemia cell lines and poor metabolic stability, limiting *in vivo* studies. Insertion of a cyano-indole ring connected to the thienopyrimidine scaffold *via* a piperidine linker furnished the compound **118 (MI-136)**, which maintained a good inhibitory activity and a strong binding affinity for menin (IC_{50} of 31 nM and K_D of 24 nM, respectively).²⁹⁸ The introduction of a methyl and/or a methyl pyrazole substituent on the indole nitrogen yielded compounds **119 (MI-463)** and **120 (MI-503)** as efficient menin binders with low-nanomolar binding affinities (IC_{50} of 15.3 nM and 14.7 nM and K_D of 9.9 nM and 9.3 nM, respectively). Treatment of human leukaemia cell lines, such as MV4-11 and MOLM-13 (which express the MLL–AF9 oncogene), with **120** resulted in a strong downregulation expression of homeobox genes (HOXA9, HOXA10, HOXA7, PBX3, and MEIS1) and other downstream targets of MLL fusion proteins. **119** and **120** displayed good metabolic stability and a PK profile in mice, and their *in vivo* oral administration

resulted in a longer survival time in a mouse model of MLL leukaemia.²⁹⁸ Further SAR studies led to the identification of **121 (MI-538)**, which displayed a strong *in vitro* inhibition of the menin–MLL interaction and a high binding affinity for menin (IC_{50} of 21 nM and K_D of 6.5 nM, respectively). Compared with **119** and **120**, this compound revealed a significant improvement regarding the cellular activity, selectivity for MLL leukaemia cells, and PK properties.²⁹⁹ Additional optimization of the substituent of the indole nitrogen led to an improvement in inhibitory activity over previously identified compounds. **122 (MI-1481)** emerged as potent inhibitor of the menin–MLL interaction (IC_{50} of 3.6 nM). The improved inhibitory activity of **122** could be associated with a stronger interference, not only with the MBM1 motif but also with the MBM2 portion of MLL, resulting in a better displacement of the bivalent MLL peptide from its binding to menin. On the other hand, the increased polarity of **122** negatively affected the oral bioavailability of this compound.³⁰⁰ The introduction of an amino group in the thienopyrimidine scaffold, combined with optimization of the pattern of substituents on the indole ring, furnished **123 (MI-3454)**. This compound exhibited a sub-nanomolar inhibitory activity (IC_{50} of 0.51 nM), efficiently disrupting the interaction of menin with an MLL fragment. **123** has a high oral bioavailability and its administration in mice significantly induces a complete remission or regression of leukaemia in different models, including patient-derived xenografts (PDXs).³⁰¹

Iterative structure-based drug design of a piperazinyl pyrimidine fragment initially identified subsequently yielded **124 (VTP50469)**, a potent, highly selective, and orally bioavailable inhibitor. Structurally, the compound maintains the pyrimidine core, while the piperazine was replaced with a spirocyclic moiety decorated with a cyclohexyl sulfonamide group. The binding of **124** into the menin–MLL binding pocket was highlighted *via* the co-crystal structure (PDB 6PKC). In Fig. 35 are summarized the key binding features, including the hydrogen bond with Tyr276 and Trp341 and cation– π interactions with Tyr319 and Tyr323 side chains. In cellular studies, **124** inhibited proliferation and produced apoptosis of leukaemia cell lines harbouring MLL rearrangements, with no effect on the cell lines lacking them. Moreover, the treatment of PDX models developed from patients with different types of leukaemia with **124** resulted in impressive reductions of leukaemia. Interestingly, **124** is able to displace menin from chromatin genome-wide also decreasing the DOT1L chromatin occupancy, probably causing its destabilization.³⁰²

Aiming to identify new menin–MLL inhibitors with diverse chemical scaffolds, the Grembecka group identified a novel class of compounds starting from the HTS of approximately 288 000 small molecules. **125 (MIV-6R)**, an aminomethylpiperidine-based compound, displayed a high binding affinity for menin and high potency in disrupting menin–MLL interactions (K_D of 85 nM and IC_{50} of 56 nM). The compound efficiently induced the hematopoietic differentiation in MLL leukaemia cells and inhibited their proliferation, reducing the expression level of key MLL–fusion protein target genes.³⁰³ However, the primary amino group could affect the metabolic stability of **125**. In order to improve this issue, some modifications were applied, leading to the identification of



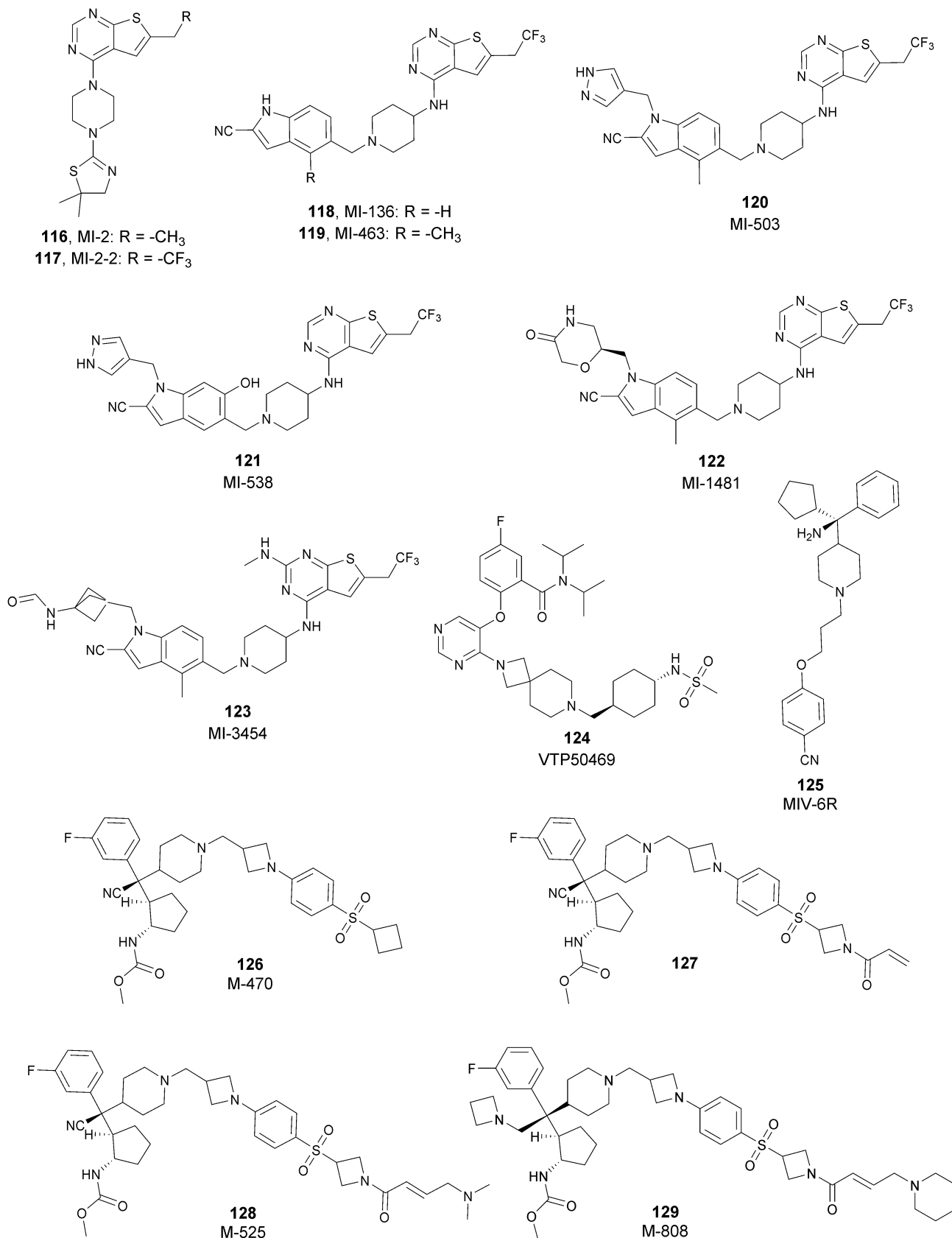


Fig. 34 MLL-menin interaction disruptors.



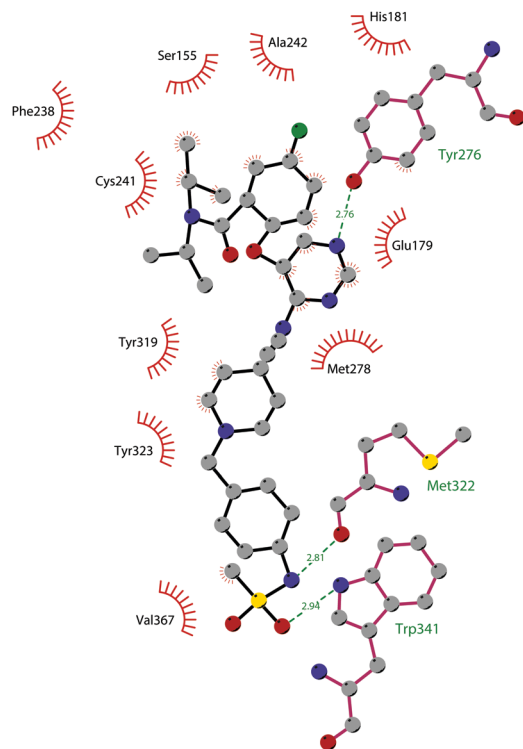


Fig. 35 Flat representation (generated using LigPlot+ from PDB ID 6PKC) showing contacts between compound **124** and the active-site residues in menin. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with compounds.

126 (M-470). Specifically, the primary amino group was replaced with a metabolically stable nitrile group and the cyclopentyl ring was functionalized with a reverse carbamate, inserting a fluorine into the phenyl ring to improve the binding affinity. Moreover, the linker was constrained into a 3-methylazetidine group and the nitrile group in position 6 was replaced with a cyclobutylsulfonyl group.³⁰⁴ These modifications improved the inhibitory activity (IC_{50} of 7 nM), the metabolic stability and, above all, offered the chance to design a covalent inhibitor. Indeed, the insertion of an acrylamide group as a Michael acceptor on the 4-membered ring allowed the identification of **127**, a covalent binder of menin (IC_{50} of 3 nM), which strongly inhibited cell growth in MV4-11 cells. To improve the solubility, a (dimethylamino)-methyl group was attached to the acrylamide group, obtaining **128 (M-525)**, which showed a potent antiproliferative effect in a variety of human acute leukaemia cell lines, mostly harbouring MLL fusions. The co-crystal structure of **128** in complex with menin revealed that the compound forms a covalent bond with the sulphur atom of the C329 residue through its acrylamide group.³⁰⁴

Further optimization of **128** led to the identification of **129 (M-808)**, a covalent menin inhibitor that is able to potently inhibit cell growth in MV4-11 and MOLM-13 cell lines harbouring MLL fusion (IC_{50} of 1 and 4 nM, respectively). Notably, the compound demonstrated strong antitumor effects in the MV4-11 xenograft tumour model in mice with severe combined immunodeficiency, and its intravenous administration (25 mg kg⁻¹ every day, three times per week) resulted in 97% tumour growth inhibition.³⁰⁵

PRDM9

PRDM9 is a member of the PRDM (PRDI-BF1 (positive regulatory domain I-binding factor 1) and RIZ1 (retinoblastoma protein-interacting zinc finger gene 1)) family, a group of 19 methyltransferases that are involved in the epigenetic regulation of gene expression.³⁰⁶ A common feature of all these proteins is the presence of several protein–protein and protein–DNA binding domains and an N-terminal PR domain, which is structurally and functionally related to the SET domain.³⁰⁷

However, despite the high homology with the SET domain, only a few PRDM proteins (PRDM2, PRDM7 and PRDM9) have been characterized by their methyltransferase activity, while the others are mainly involved in recruiting other writers through protein–protein interactions.³⁰⁸

PRDM9 catalyses the trimethylation of H3K4 and H3K36 on nucleosome substrates; however, thanks to the presence of its specific protein–protein domains, it interacts with other enzymes, leading to genetic homologous recombination. In addition, PRDM9 is considered an important enzyme in the meiotic prophase.³⁰⁹ Physiologically, PRDM9 is expressed only in germ cells in the gonads and testes and the absence of PRDM9 in mice is associated with sterility for the male as well as the female phenotype. Alterations in the expression of the protein and also the presence of PRDM9 mutations has been related to the insurgence of tumours such as head and neck squamous cell carcinoma, renal cell cancer and leukaemia.^{310–313}

PRDM9 inhibitors

Despite the increasing interest in the PRDM subfamily of enzymes, so far only one PRDM9 inhibitor has been reported (Fig. 36). In 2019, a radioactivity-based methyltransferase assay was used to screen a library of 7500 compounds, enabling the identification of 39 hits that are able to inhibit PRDM9 with IC_{50} values ranging from 4 to 30 μ M. After an extensive SAR campaign, **130 (MRK-740)** emerged as a potent (IC_{50} of 80 nM) and selective PRDM9 inhibitor. The crystal structure of the PRDM9 catalytic domain–SAM–**130** ternary complex (PDB 6NM4) enabled the confirmation that this compound acts as a SAM-dependent substrate-competitive inhibitor.

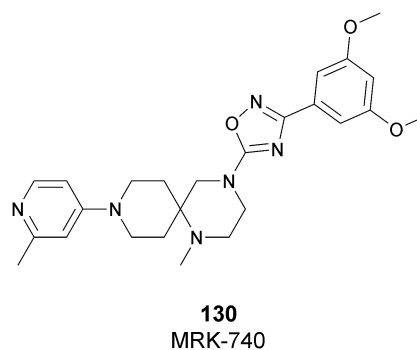


Fig. 36 PRDM9 inhibitor.

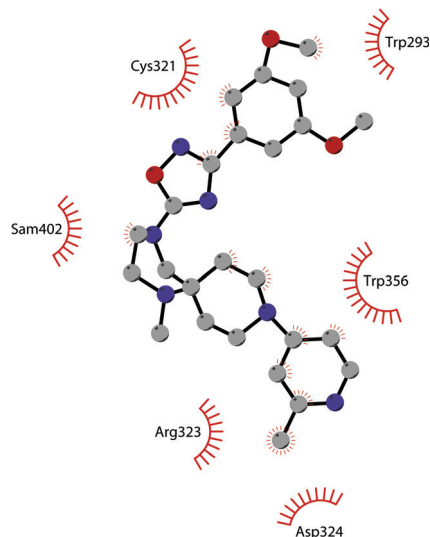


Fig. 37 Flat representation (generated using LigPlot+ from PDB ID 6NM4) showing contacts between compound **130** and active-site residues in PRDM9. Hydrogen bonds are shown as dashed lines. The spoked arcs represent protein residues making non-bonded (hydrophobic) contacts with the compound.

A representation of the intermolecular contacts is reported in Fig. 37, revealing a large number of hydrophobic interactions between the PRDM9 side chains and **130**. In addition, the pyridine ring is responsible for the π -stacking interactions with Trp356. On the other hand, the pyridine nitrogen is assumed to be protonated and forms a long-range electrostatic interaction with the side chain of Asp324. In cellular assays, **130** showed the direct inhibition of H3K4 methylation at endogenous PRDM9 target loci, representing the first-in-class PRDM sub-family chemical probe.³¹⁴

ASH1L

The protein Ash1 (absent, small, or homeotic disc 1) was first identified in *Drosophila* as a member of the trithorax group of proteins,³¹⁵ which act as major regulators in embryonic development and whose function counteracts the polycomb group of proteins that are responsible for gene silencing. The mammalian homolog of Ash1, named ASH1L (ASH1-like, also known as KMT2H), is a methyltransferase that catalyses the dimethylation of H3K36. This protein is localized in the promoter-proximal coding regions of active genes, suggesting a role as an epigenetic activator.³¹⁶

From the structural point of view, ASH1L contains a SET domain that is located in the middle of the sequence, followed by a post-SET domain, a bromodomain, a bromo-adjacent homology (BAH) domain and a PHD zinc finger domain.³¹⁷

Functionally, ASH1L interacts with proteins MRG15 and NURF55, constituting the ASH1 complex. In particular, the interaction with MRG15 significantly promotes the catalytic activity of ASH1L, probably by inducing a conformational change that abolishes the block of the auto-inhibitory loop of ASH1L.³¹⁸

ASH1L is widely expressed mostly in the brain as it plays an important role in the development and in the function of the

nervous system. As a matter of fact, alterations of the ASH1L gene are related to the development of neurological disorders such as Tourette syndrome, autism spectrum disorder, and intellectual disability during development.³¹⁹ In addition, ASH1L is overexpressed in breast cancer,¹⁸⁴ in hepatocellular carcinoma³²⁰ and in anaplastic thyroid cancer,³²¹ although the role in the biology of these tumours has still not been fully disclosed.

ASH1L inhibitors

The involvement of ASH1L in several cancer types makes this protein an interesting therapeutic target. Unfortunately, the identification of selective inhibitors has been challenging, mainly due to the limited access to the active site of the protein, considering that the catalytic SET domain adopts an inactive conformation with an autoinhibitory loop. Very recently, Grembecka and co-workers reported the identification of the first-in-class ASH1L inhibitors. The NMR-based screening of approximately 1600 fragments from an internal library, followed by medicinal chemistry optimization and structure-based design, enabled the identification of thioamide-containing compounds, of which derivative **131** (AS-99, Fig. 38) was the best candidate with an IC_{50} of 0.79 μ M and a K_D of 0.89 μ M.

This inhibitor was found to be highly selective towards ASH1L, with no inhibition observed at 50 μ M on a panel of 20 HMTs as well as on a panel of representative kinases. Interestingly, structural studies revealed that the thioamide-containing derivatives allosterically inhibited the enzymatic activity of the protein, inserting into a pocket that flanks the autoinhibitory loop in the SET domain and stabilizing this conformation. Compound **131** is able to reduce dimethylated levels of H3K36 in MLL leukaemia cells, inhibiting their proliferation and inducing their differentiation and apoptosis. Moreover, this compound downregulates the expression of MLL fusion target genes and reduces the leukaemia burden in a systemic model of MLL leukaemia. This evidence endorses **131** as a valuable chemical probe to further investigation of ASH1L biology.³²²

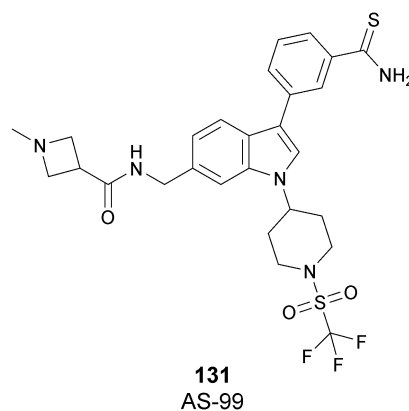


Fig. 38 ASH1L inhibitor.

Conclusions

In the past 20 years, a great abundance of studies has shown the importance of KMTs in a huge number of physiological processes and in the development of several diseases. These findings have led to the discovery of a plethora of modulators with the aim of fully elucidating the biological role of these proteins as well as establishing them as targets for new therapeutic approaches. In this review, we have reported the so-far identified modulators of KMTs. Among them, one compound has been recently approved by the FDA, some of them are in advanced phases of preclinical studies, and others have only recently been identified and for them a thorough characterization is still needed before they can enter the clinic. Nevertheless, there is a huge number of proteins, endowed with a clear role in several pathological conditions, for which no compounds have been identified yet. Hence, despite the recent advancements in this field, there is still an urgent need to identify new potent and selective modulators of these epigenetic targets.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

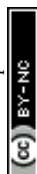
- 1 K. Murray, *Biochemistry*, 1964, **3**, 10–15.
- 2 D. Chen, H. Ma, H. Hong, S. S. Koh, S.-M. Huang, B. T. Schurter, D. W. Aswad and M. R. Stallcup, *Science*, 1999, **284**, 2174–2177.
- 3 S. Rea, F. Eisenhaber, D. O'Carroll, B. D. Strahl, Z.-W. Sun, M. Schmid, S. Opravil, K. Mechtler, C. P. Ponting, C. D. Allis and T. Jenuwein, *Nature*, 2000, **406**, 593–599.
- 4 R. Hamamoto, V. Saloura and Y. Nakamura, *Nat. Rev. Cancer*, 2015, **15**, 110–124.
- 5 T. Vougiouklakis, B. J. Bernard, N. Nigam, K. Burkitt, Y. Nakamura and V. Saloura, *Clin. Epigenet.*, 2020, **12**, 146.
- 6 M. Urulangodi and A. Mohanty, *J. Cell Commun. Signal*, 2020, **14**, 31–45.
- 7 X. Wei, X. Yi, X.-H. Zhu and D.-S. Jiang, *Clin. Epigenet.*, 2020, **12**, 30.
- 8 E. M. Michalak, M. L. Burr, A. J. Bannister and M. A. Dawson, *Nat. Rev. Mol. Cell Biol.*, 2019, **20**, 573–589.
- 9 B. Tschiersch, A. Hofmann, V. Krauss, R. Dorn, G. Korge and G. Reuter, *EMBO J.*, 1994, **13**, 3822–3831.
- 10 R. S. Jones and W. M. Gelbart, *Mol. Cell. Biol.*, 1993, **13**, 6357–6366.
- 11 M. J. Stassen, D. Bailey, S. Nelson, V. Chinwalla and P. J. Harte, *Mech. Dev.*, 1995, **52**, 209–223.
- 12 R. C. Trievel, B. M. Beach, L. M. Dirk, R. L. Houtz and J. H. Hurley, *Cell*, 2002, **111**, 91–103.
- 13 M. Schapira, *Curr. Chem. Genomics*, 2011, **5**, 85–94.
- 14 K. Sawada, Z. Yang, J. R. Horton, R. E. Collins, X. Zhang and X. Cheng, *J. Biol. Chem.*, 2004, **279**, 43296–43306.
- 15 P. A. Del Rizzo and R. C. Trievel, *Epigenetics*, 2011, **6**, 1059–1067.
- 16 D. Husmann and O. Gozani, *Nat. Struct. Mol. Biol.*, 2019, **26**, 880–889.
- 17 E. M. Rowe, V. Xing and K. K. Biggar, *Brain Res.*, 2019, **1707**, 164–171.
- 18 K. P. Bhat, H. Ümit Kaniskan, J. Jin and O. Gozani, *Nat. Rev. Drug Discovery*, 2021, **20**, 265–286.
- 19 C. Zagni, U. Chiacchio and A. Rescifina, *Curr. Med. Chem.*, 2013, **20**, 167–185.
- 20 R. A. Copeland, *Philos. Trans. R. Soc., B*, 2018, **373**, 20170080.
- 21 H. U. Kaniskan and J. Jin, *ACS Chem. Biol.*, 2015, **10**, 40–50.
- 22 C. Milite, A. Feoli, M. Viviano, D. Rescigno, A. Mai, S. Castellano and G. Sbardella, *ChemMedChem*, 2016, **11**, 1680–1685.
- 23 C. Milite, A. Feoli, M. Viviano, D. Rescigno, A. Cianciulli, A. L. Balzano, A. Mai, S. Castellano and G. Sbardella, *Clin. Epigenet.*, 2016, **8**, 102.
- 24 M. S. Singer, A. Kahana, A. J. Wolf, L. L. Meisinger, S. E. Peterson, C. Goggin, M. Mahowald and D. E. Gottschling, *Genetics*, 1998, **150**, 613–632.
- 25 J. Min, Q. Feng, Z. Li, Y. Zhang and R. M. Xu, *Cell*, 2003, **112**, 711–723.
- 26 F. van Leeuwen, P. R. Gafken and D. E. Gottschling, *Cell*, 2002, **109**, 745–756.
- 27 B. Jones, H. Su, A. Bhat, H. Lei, J. Bajko, S. Hevi, G. A. Baltus, S. Kadam, H. Zhai, R. Valdez, S. Gonzalo, Y. Zhang, E. Li and T. Chen, *PLoS Genet.*, 2008, **4**, e1000190.
- 28 G. A. Shanower, M. Muller, J. L. Blanton, V. Honti, H. Gyurkovics and P. Schedl, *Genetics*, 2005, **169**, 173–184.
- 29 M. Mohan, H. M. Herz, Y. H. Takahashi, C. Lin, K. C. Lai, Y. Zhang, M. P. Washburn, L. Florens and A. Shilatifard, *Genes Dev.*, 2010, **24**, 574–589.
- 30 Y. Okada, Q. Feng, Y. Lin, Q. Jiang, Y. Li, V. M. Coffield, L. Su, G. Xu and Y. Zhang, *Cell*, 2005, **121**, 167–178.
- 31 K. M. Bernt, N. Zhu, A. U. Sinha, S. Vempati, J. Faber, A. V. Krivtsov, Z. Feng, N. Punt, A. Daigle, L. Bullinger, R. M. Pollock, V. M. Richon, A. L. Kung and S. A. Armstrong, *Cancer Cell*, 2011, **20**, 66–78.
- 32 S. R. Daigle, E. J. Olhava, C. A. Theriksen, C. R. Majer, C. J. Sneeringer, J. Song, L. D. Johnston, M. P. Scott, J. J. Smith, Y. Xiao, L. Jin, K. W. Kuntz, R. Chesworth, M. P. Moyer, K. M. Bernt, J. C. Tseng, A. L. Kung,



- S. A. Armstrong, R. A. Copeland, V. M. Richon and R. M. Pollock, *Cancer Cell*, 2011, **20**, 53–65.
- 33 A. T. Nguyen, J. He, O. Taranova and Y. Zhang, *Cell Res.*, 2011, **21**, 1370–1373.
- 34 S. S. Spurr, E. D. Bayle, W. Yu, F. Li, W. Tempel, M. Vedadi, M. Schapira and P. V. Fish, *Bioorg. Med. Chem. Lett.*, 2016, **26**, 4518–4522.
- 35 J. L. Anglin, L. Deng, Y. Yao, G. Cai, Z. Liu, H. Jiang, G. Cheng, P. Chen, S. Dong and Y. Song, *J. Med. Chem.*, 2012, **55**, 8066–8074.
- 36 W. Yu, D. Smil, F. Li, W. Tempel, O. Fedorov, K. T. Nguyen, Y. Bolshan, R. Al-Awar, S. Knapp, C. H. Arrowsmith, M. Vedadi, P. J. Brown and M. Schapira, *Bioorg. Med. Chem.*, 2013, **21**, 1787–1794.
- 37 S. R. Daigle, E. J. Olhava, C. A. Therkelsen, A. Basavapathruni, L. Jin, P. A. Boriack-Sjodin, C. J. Allain, C. R. Klaus, A. Raimondi, M. P. Scott, N. J. Waters, R. Chesworth, M. P. Moyer, R. A. Copeland, V. M. Richon and R. M. Pollock, *Blood*, 2013, **122**, 1017–1025.
- 38 A. Basavapathruni, E. J. Olhava, S. R. Daigle, C. A. Therkelsen, L. Jin, P. A. Boriack-Sjodin, C. J. Allain, C. R. Klaus, A. Raimondi, M. P. Scott, A. Dovletoglou, V. M. Richon, R. M. Pollock, R. A. Copeland, M. P. Moyer, R. Chesworth, P. G. Pearson and N. J. Waters, *Biopharm. Drug Dispos.*, 2014, **35**, 237–252.
- 39 R. A. Laskowski and M. B. Swindells, *J. Chem. Inf. Model.*, 2011, **51**, 2778–2786.
- 40 I. Epizyme and C. Corporation, A Phase 1 Dose Escalation and Expanded Cohort Study of EPZ-5676 in the Treatment of Pediatric Patients With Relapsed/Refractory Leukemias Bearing a Rearrangement of the MLL Gene, <https://ClinicalTrials.gov/show/NCT02141828>, 2014.
- 41 I. Epizyme and Celgene, A Phase 1, Open-Label, Dose-Escalation & Expanded Cohort, Continuous IV Infusion, Multi-center Study of the Safety, Tolerability, PK & PD of EPZ-5676 in Treatment Relapsed/Refractory Patients With Leukemias Involving, <https://ClinicalTrials.gov/show/NCT01684150>, 2012.
- 42 N. C. Institute, Pinometostat With Standard Chemotherapy in Treating Patients With Newly Diagnosed Acute Myeloid Leukemia and MLL Gene Rearrangement, <https://ClinicalTrials.gov/show/NCT03724084>, 2019.
- 43 N. C. Institute, Pinometostat and Azacitidine in Treating Patients With Relapsed, Refractory, or Newly Diagnosed Acute Myeloid Leukemia With 11q23 Rearrangement, <https://ClinicalTrials.gov/show/NCT03701295>, 2019.
- 44 S. Chen, L. Li, Y. Chen, J. Hu, J. Liu, Y.-C. Liu, R. Liu, Y. Zhang, F. Meng, K. Zhu, J. Lu, M. Zheng, K. Chen, J. Zhang, H. Jiang, Z. Yao and C. Luo, *J. Chem. Inf. Model.*, 2016, **56**, 527–534.
- 45 Y. Zhang, Y. Zhang, J. Liu, L. Chen, L. Zhao, B. Li and W. Wang, *Bioorg. Med. Chem. Lett.*, 2017, **27**, 1584–1587.
- 46 C. Scheufler, H. Möbitz, C. Gaul, C. Ragot, C. Be, C. Fernández, K. S. Beyer, R. Tiedt and F. Stauffer, *ACS Med. Chem. Lett.*, 2016, **7**, 730–734.
- 47 C. Chen, H. Zhu, F. Stauffer, G. Caravatti, S. Vollmer, R. Machauer, P. Holzer, H. Mobitz, C. Scheufler, M. Klumpp, R. Tiedt, K. S. Beyer, K. Calkins, D. Guthy, M. Kiffe, J. Zhang and C. Gaul, *ACS Med. Chem. Lett.*, 2016, **7**, 735–740.
- 48 H. Möbitz, R. Machauer, P. Holzer, A. Vaupel, F. Stauffer, C. Ragot, G. Caravatti, C. Scheufler, C. Fernandez, U. Hommel, R. Tiedt, K. S. Beyer, C. Chen, H. Zhu and C. Gaul, *ACS Med. Chem. Lett.*, 2017, **8**, 338–343.
- 49 F. Stauffer, A. Weiss, C. Scheufler, H. Mobitz, C. Ragot, K. S. Beyer, K. Calkins, D. Guthy, M. Kiffe, B. Van Eerdenbrugh, R. Tiedt and C. Gaul, *ACS Med. Chem. Lett.*, 2019, **10**, 1655–1660.
- 50 G. S. Gibbons, A. Chakraborty, S. M. Grigsby, A. C. Umeano, C. Liao, O. Moukha-Chafiq, V. Pathak, B. Mathew, Y. T. Lee, Y. Dou, S. C. Schurer, R. C. Reynolds, T. S. Snowden and Z. Nikolovska-Coleska, *Eur. J. Med. Chem.*, 2020, **189**, 112023.
- 51 M. Tachibana, K. Sugimoto, T. Fukushima and Y. Shinkai, *J. Biol. Chem.*, 2001, **276**, 25309–25317.
- 52 M. Tachibana, J. Ueda, M. Fukuda, N. Takeda, T. Ohta, H. Iwanari, T. Sakihama, T. Kodama, T. Hamakubo and Y. Shinkai, *Genes Dev.*, 2005, **19**, 815–826.
- 53 D. Patnaik, H. G. Chin, P. O. Estève, J. Benner, S. E. Jacobsen and S. Pradhan, *J. Biol. Chem.*, 2004, **279**, 53248–53258.
- 54 H. Wu, X. Chen, J. Xiong, Y. Li, H. Li, X. Ding, S. Liu, S. Chen, S. Gao and B. Zhu, *Cell Res.*, 2011, **21**, 365–367.
- 55 Y. Yu, C. Song, Q. Zhang, P. A. DiMaggio, B. A. Garcia, A. York, M. F. Carey and M. Grunstein, *Mol. Cell*, 2012, **46**, 7–17.
- 56 P. Trojer, J. Zhang, M. Yonezawa, A. Schmidt, H. Zheng, T. Jenuwein and D. Reinberg, *J. Biol. Chem.*, 2009, **284**, 8395–8405.
- 57 T. Weiss, S. Hergeth, U. Zeissler, A. Izzo, P. Tropberger, B. M. Zee, M. Dunder, B. A. Garcia, S. Daujat and R. Schneider, *Epigenet. Chromatin*, 2010, **3**, 7.
- 58 J. Huang, J. Dorsey, S. Chuikov, L. Pérez-Burgos, X. Zhang, T. Jenuwein, D. Reinberg and S. L. Berger, *J. Biol. Chem.*, 2010, **285**, 9636–9641.
- 59 K. E. Moore, S. M. Carlson, N. D. Camp, P. Cheung, R. G. James, K. F. Chua, A. Wolf-Yadlin and O. Gozani, *Mol. Cell*, 2013, **50**, 444–456.
- 60 S. C. Sampath, I. Marazzi, K. L. Yap, S. C. Sampath, A. N. Krutchinsky, I. Mecklenbräuer, A. Viale, E. Rudensky, M. M. Zhou, B. T. Chait and A. Tarakhovsky, *Mol. Cell*, 2007, **27**, 596–608.
- 61 C. Mozzetta, J. Pontis, L. Fritsch, P. Robin, M. Portoso, C. Proux, R. Margueron and S. Ait-Si-Ali, *Mol. Cell*, 2014, **53**, 277–289.
- 62 R. E. Collins, J. P. Northrop, J. R. Horton, D. Y. Lee, X. Zhang, M. R. Stallcup and X. Cheng, *Nat. Struct. Mol. Biol.*, 2008, **15**, 245–250.
- 63 H. Wu, J. Min, V. V. Lunin, T. Antoshenko, L. Dombrowski, H. Zeng, A. Allali-Hassani, V. Campagna-Slater, M. Vedadi, C. H. Arrowsmith, A. N. Plotnikov and M. Schapira, *PLoS One*, 2010, **5**, e8570.



- 64 N. Liu, Z. Zhang, H. Wu, Y. Jiang, L. Meng, J. Xiong, Z. Zhao, X. Zhou, J. Li, H. Li, Y. Zheng, S. Chen, T. Cai, S. Gao and B. Zhu, *Genes Dev.*, 2015, **29**, 379–393.
- 65 M. Tachibana, K. Sugimoto, M. Nozaki, J. Ueda, T. Ohta, M. Ohki, M. Fukuda, N. Takeda, H. Niida, H. Kato and Y. Shinkai, *Genes Dev.*, 2002, **16**, 1779–1791.
- 66 T. H. W. Dobson, R. J. Hatcher, J. Swaminathan, C. M. Das, S. Shaik, R.-H. Tao, C. Milite, S. Castellano, P. H. Taylor, G. Sbardella and V. Gopalakrishnan, *Mol. Cancer Res.*, 2017, **15**, 1073–1084.
- 67 T. Huang, P. Zhang, W. Li, T. Zhao, Z. Zhang, S. Chen, Y. Yang, Y. Feng, F. Li, X. Shirley Liu, L. Zhang, G. Jiang and F. Zhang, *Cell Death Dis.*, 2017, **8**, e2726.
- 68 T. Kleefstra, M. Smidt, M. J. Banning, A. R. Oudakker, H. Van Esch, A. P. de Brouwer, W. Nillesen, E. A. Sistermans, B. C. Hamel, D. de Bruijn, J. P. Fryns, H. G. Yntema, H. G. Brunner, B. B. de Vries and H. van Bokhoven, *J. Med. Genet.*, 2005, **42**, 299–306.
- 69 F. Antignano, K. Burrows, M. R. Hughes, J. M. Han, K. J. Kron, N. M. Penrod, M. J. Oudhoff, S. K. Wang, P. H. Min, M. J. Gold, A. L. Chenery, M. J. Braam, T. C. Fung, F. M. Rossi, K. M. McNagny, C. H. Arrowsmith, M. Lupien, M. K. Levings and C. Zaph, *J. Clin. Invest.*, 2014, **124**, 1945–1955.
- 70 Q. Liu, X. Cai, D. Yang, Y. Chen, Y. Wang, L. Shao and M. W. Wang, *Bioorg. Med. Chem.*, 2017, **25**, 4579–4594.
- 71 K. Devkota, B. Lohse, Q. Liu, M. W. Wang, D. Stærk, J. Berthelsen and R. P. Clausen, *ACS Med. Chem. Lett.*, 2014, **5**, 293–297.
- 72 S. Kubicek, R. J. O'Sullivan, E. M. August, E. R. Hickey, Q. Zhang, M. L. Teodoro, S. Rea, K. Mechtler, J. A. Kowalski, C. A. Homon, T. A. Kelly and T. Jenuwein, *Mol. Cell*, 2007, **25**, 473–481.
- 73 Y. Yuan, Q. Wang, J. Paulk, S. Kubicek, M. M. Kemp, D. J. Adams, A. F. Shamji, B. K. Wagner and S. L. Schreiber, *ACS Chem. Biol.*, 2012, **7**, 1152–1157.
- 74 Y. Chang, X. Zhang, J. R. Horton, A. K. Upadhyay, A. Spannhoff, J. Liu, J. P. Snyder, M. T. Bedford and X. Cheng, *Nat. Struct. Mol. Biol.*, 2009, **16**, 312–317.
- 75 I. A. Ciechomska, P. Przanowski, J. Jackl, B. Wojtas and B. Kaminska, *Sci. Rep.*, 2016, **6**, 38723.
- 76 Z. Rahman, M. R. Bazaz, G. Devabattula, M. A. Khan and C. Godugu, *J. Biochem. Mol. Toxicol.*, 2020, e22674, DOI: 10.1002/jbt.22674.
- 77 Y. Chang, T. Ganesh, J. R. Horton, A. Spannhoff, J. Liu, A. Sun, X. Zhang, M. T. Bedford, Y. Shinkai, J. P. Snyder and X. Cheng, *J. Mol. Biol.*, 2010, **400**, 1–7.
- 78 F. Liu, X. Chen, A. Allali-Hassani, A. M. Quinn, G. A. Wasney, A. Dong, D. Barsyte, I. Kozieradzki, G. Senisterra, I. Chau, A. Siarheyeva, D. B. Kireev, A. Jadhav, J. M. Herold, S. V. Frye, C. H. Arrowsmith, P. J. Brown, A. Simeonov, M. Vedadi and J. Jin, *J. Med. Chem.*, 2009, **52**, 7950–7953.
- 79 F. Liu, X. Chen, A. Allali-Hassani, A. M. Quinn, T. J. Wigle, G. A. Wasney, A. Dong, G. Senisterra, I. Chau, A. Siarheyeva, J. L. Norris, D. B. Kireev, A. Jadhav, J. M. Herold, W. P. Janzen, C. H. Arrowsmith, S. V. Frye, P. J. Brown, A. Simeonov, M. Vedadi and J. Jin, *J. Med. Chem.*, 2010, **53**, 5844–5857.
- 80 M. Vedadi, D. Barsyte-Lovejoy, F. Liu, S. Rival-Gervier, A. Allali-Hassani, V. Labrie, T. J. Wigle, P. A. Dimaggio, G. A. Wasney, A. Siarheyeva, A. Dong, W. Tempel, S. C. Wang, X. Chen, I. Chau, T. J. Mangano, X. P. Huang, C. D. Simpson, S. G. Pattenden, J. L. Norris, D. B. Kireev, A. Tripathy, A. Edwards, B. L. Roth, W. P. Janzen, B. A. Garcia, A. Petronis, J. Ellis, P. J. Brown, S. V. Frye, C. H. Arrowsmith and J. Jin, *Nat. Chem. Biol.*, 2011, **7**, 566–574.
- 81 F. Liu, D. Barsyte-Lovejoy, F. Li, Y. Xiong, V. Korboukh, X. P. Huang, A. Allali-Hassani, W. P. Janzen, B. L. Roth, S. V. Frye, C. H. Arrowsmith, P. J. Brown, M. Vedadi and J. Jin, *J. Med. Chem.*, 2013, **56**, 8931–8942.
- 82 F. Liu, D. Barsyte-Lovejoy, A. Allali-Hassani, Y. He, J. M. Herold, X. Chen, C. M. Yates, S. V. Frye, P. J. Brown, J. Huang, M. Vedadi, C. H. Arrowsmith and J. Jin, *J. Med. Chem.*, 2011, **54**, 6139–6150.
- 83 C. Milite, A. Feoli, J. R. Horton, D. Rescigno, A. Cipriano, V. Pisapia, M. Viviano, G. Pepe, G. Amendola, E. Novellino, S. Cosconati, X. Cheng, S. Castellano and G. Sbardella, *J. Med. Chem.*, 2019, **62**, 2666–2689.
- 84 E. San Jose-Eneriz, X. Agirre, O. Rabal, A. Vilas-Zornoza, J. A. Sanchez-Arias, E. Miranda, A. Ugarte, S. Roa, B. Paiva, A. Estella-Hermoso de Mendoza, R. M. Alvarez, N. Casares, V. Segura, J. I. Martin-Subero, F. X. Ogi, P. Soule, C. M. Santiveri, R. Campos-Olivas, G. Castellano, M. G. F. de Barrena, J. R. Rodriguez-Madoz, M. J. Garcia-Barchino, J. J. Lasarte, M. A. Avila, J. A. Martinez-Climent, J. Oyarzabal and F. Prosper, *Nat. Commun.*, 2017, **8**, 15424.
- 85 R. F. Sweis, M. Pliushchev, P. J. Brown, J. Guo, F. Li, D. Maag, A. M. Petros, N. B. Soni, C. Tse, M. Vedadi, M. R. Michaelides, G. G. Chiang and W. N. Pappano, *ACS Med. Chem. Lett.*, 2014, **5**, 205–209.
- 86 T. Wagner, H. Greschik, T. Burgahn, K. Schmidtkunz, A. K. Schott, J. McMillan, L. Baranauskiene, Y. Xiong, O. Fedorov, J. Jin, U. Oppermann, D. Matulis, R. Schule and M. Jung, *Nucleic Acids Res.*, 2016, **44**, e88.
- 87 V. Fagan, C. Johansson, C. Gileadi, O. Monteiro, J. E. Dunford, R. Nibhani, M. Philpott, J. Malzahn, G. Wells, R. Faram, A. P. Cribbs, N. Halidi, F. Li, I. Chau, H. Greschik, S. Velupillai, A. Allali-Hassani, J. Bennett, T. Christott, C. Giroud, A. M. Lewis, K. V. M. Huber, N. Athanasou, C. Bountra, M. Jung, R. Schule, M. Vedadi, C. Arrowsmith, Y. Xiong, J. Jin, O. Fedorov, G. Farnie, P. E. Brennan and U. Oppermann, *J. Med. Chem.*, 2019, **62**, 9008–9025.
- 88 K. Katayama, K. Ishii, H. Terashima, E. Tsuda, M. Suzuki, K. Yotsumoto, K. Hiramoto, I. Yasumatsu, M. Torihata, T. Ishiyama, T. Muto and T. Katagiri, *ACS Med. Chem. Lett.*, 2021, **12**, 121–128.
- 89 K. Katayama, K. Ishii, E. Tsuda, K. Yotsumoto, K. Hiramoto, M. Suzuki, I. Yasumatsu, W. Igarashi, M. Torihata, T. Ishiyama and T. Katagiri, *Bioorg. Med. Chem. Lett.*, 2020, **30**, 127475.



- 90 H. G. Chin, D. Patnaik, P. O. Esteve, S. E. Jacobsen and S. Pradhan, *Biochemistry*, 2006, **45**, 3272–3284.
- 91 T. Wang, C. Xu, Y. Liu, K. Fan, Z. Li, X. Sun, H. Ouyang, X. Zhang, J. Zhang, Y. Li, F. Mackenzie, J. Min and X. Tu, *PLoS One*, 2012, **7**, e52977.
- 92 D. O'Carroll, H. Scherthan, A. H. Peters, S. Opravil, A. R. Haynes, G. Laible, S. Rea, M. Schmid, A. Lebersorger, M. Jerratsch, L. Sattler, M. G. Mattei, P. Denny, S. D. Brown, D. Schweizer and T. Jenuwein, *Mol. Cell. Biol.*, 2000, **20**, 9423–9433.
- 93 M. K. Schuhmacher, S. Kudithipudi, D. Kusevic, S. Weirich and A. Jeltsch, *Biochim. Biophys. Acta*, 2015, **1849**, 55–63.
- 94 A. Allali-Hassani, G. A. Wasney, A. Siarheyeva, T. Hajian, C. H. Arrowsmith and M. Vedadi, *J. Biomol. Screen*, 2012, **17**, 71–84.
- 95 K. Sone, L. Piao, M. Nakakido, K. Ueda, T. Jenuwein, Y. Nakamura and R. Hamamoto, *Nat. Commun.*, 2014, **5**, 5691.
- 96 M. Puschendorf, R. Terranova, E. Boutsma, X. Mao, K. Isono, U. Brykczynska, C. Kolb, A. P. Otte, H. Koseki, S. H. Orkin, M. van Lohuizen and A. H. Peters, *Nat. Genet.*, 2008, **40**, 411–420.
- 97 M. García-Cao, R. O'Sullivan, A. H. Peters, T. Jenuwein and M. A. Blasco, *Nat. Genet.*, 2004, **36**, 94–99.
- 98 A. H. Peters, D. O'Carroll, H. Scherthan, K. Mechtler, S. Sauer, C. Schöfer, K. Weipoltshammer, M. Pagani, M. Lachner, A. Kohlmaier, S. Opravil, M. Doyle, M. Sibilia and T. Jenuwein, *Cell*, 2001, **107**, 323–337.
- 99 C. R. Keenan, N. Iannarella, G. Naselli, N. G. Bediaga, T. M. Johanson, L. C. Harrison and R. S. Allan, *Blood*, 2020, **135**, 2049–2058.
- 100 Y. Zheng, B. Li, J. Wang, Y. Xiong, K. Wang, Y. Qi, H. Sun, L. Wu and L. Yang, *Clin. Epigenet.*, 2018, **10**, 129.
- 101 W. Shuai, J. Wu, S. Chen, R. Liu, Z. Ye, C. Kuang, X. Fu, G. Wang, Y. Li, Q. Peng, W. Shi, Y. Li, Q. Zhou and W. Huang, *Cancer Lett.*, 2018, **422**, 56–69.
- 102 V. Jagannathan, J. Bannoehr, P. Plattet, R. Hauswirth, C. Drögemüller, M. Drögemüller, D. J. Wiener, M. Doherr, M. Owczarek-Lipska, A. Galichet, M. M. Welle, K. Tengvall, K. Bergvall, H. Lohi, S. Rüfenacht, M. Linek, M. Paradis, E. J. Müller, P. Roosje and T. Leeb, *PLoS Genet.*, 2013, **9**, e1003848.
- 103 X. Y. Liu and H. Li, *J. Diabetes Res.*, 2017, **2017**, 8365762.
- 104 Z. Fan, L. Li, M. Li, X. Zhang, C. Hao, L. Yu, S. Zeng, H. Xu, M. Fang, A. Shen, T. Jenuwein and Y. Xu, *Hepatology*, 2017, **65**, 1904–1919.
- 105 D. Greiner, T. Bonaldi, R. Eskeland, E. Roemer and A. Imhof, *Nat. Chem. Biol.*, 2005, **1**, 143–145.
- 106 A. V. Paschall, D. Yang, C. Lu, J.-H. Choi, X. Li, F. Liu, M. Figueroa, N. H. Oberlies, C. Pearce, W. B. Bollag, A. Nayak-Kapoor and K. Liu, *J. Immunology*, 2015, **195**, 1868–1882.
- 107 C. H. Arrowsmith, J. E. Audia, C. Austin, J. Baell, J. Bennett, J. Blagg, C. Bountra, P. E. Brennan, P. J. Brown, M. E. Bunnage, C. Buser-Doepner, R. M. Campbell, A. J. Carter, P. Cohen, R. A. Copeland, B. Cravatt, J. L. Dahlin, D. Dhanak, A. M. Edwards, M. Frederiksen, S. V. Frye, N. Gray, C. E. Grimshaw, D. Hepworth, T. Howe, K. V. M. Huber, J. Jin, S. Knapp, J. D. Kotz, R. G. Kruger, D. Lowe, M. M. Mader, B. Marsden, A. Mueller-Fahrnow, S. Muller, R. C. O'Hagan, J. P. Overington, D. R. Owen, S. H. Rosenberg, R. Ross, B. Roth, M. Schapira, S. L. Schreiber, B. Shoichet, M. Sundstrom, G. Superti-Furga, J. Taunton, L. Toledo-Sherman, C. Walpole, M. A. Walters, T. M. Willson, P. Workman, R. N. Young and W. J. Zuercher, *Nat. Chem. Biol.*, 2015, **11**, 536–541.
- 108 C. Lu, D. Yang, J. D. Klement, I. K. Oh, N. M. Savage, J. L. Waller, A. H. Colby, M. W. Grinstaff, N. H. Oberlies, C. J. Pearce, Z. Xie, S. K. Kulp, C. C. Coss, M. A. Phelps, T. Albers, I. O. Lebedeva and K. Liu, *Cancer Immunol. Res.*, 2019, **7**, 414–427.
- 109 C. Lu, J. D. Klement, D. Yang, T. Albers, I. O. Lebedeva, J. L. Waller and K. Liu, *Cancer Lett.*, 2020, **476**, 87–96.
- 110 T. Vougiouklakis, V. Saloura, J. H. Park, N. Takamatsu, T. Miyamoto, Y. Nakamura and Y. Matsuo, *Oncotarget*, 2018, **9**, 31820–31831.
- 111 J. Müller, C. M. Hart, N. J. Francis, M. L. Vargas, A. Sengupta, B. Wild, E. L. Miller, M. B. O'Connor, R. E. Kingston and J. A. Simon, *Cell*, 2002, **111**, 197–208.
- 112 R. Margueron, G. Li, K. Sarma, A. Blais, J. Zavadil, C. L. Woodcock, B. D. Dynlacht and D. Reinberg, *Mol. Cell*, 2008, **32**, 503–518.
- 113 A. Kuzmichev, K. Nishioka, H. Erdjument-Bromage, P. Tempst and D. Reinberg, *Genes Dev.*, 2002, **16**, 2893–2905.
- 114 R. Cao, L. Wang, H. Wang, L. Xia, H. Erdjument-Bromage, P. Tempst, R. S. Jones and Y. Zhang, *Science*, 2002, **298**, 1039–1043.
- 115 D. Pasini, A. P. Bracken, M. R. Jensen, E. Lazzerini Denchi and K. Helin, *EMBO J.*, 2004, **23**, 4061–4071.
- 116 R. Margueron, N. Justin, K. Ohno, M. L. Sharpe, J. Son, W. J. Drury, 3rd, P. Voigt, S. R. Martin, W. R. Taylor, V. De Marco, V. Pirrotta, D. Reinberg and S. J. Gambelin, *Nature*, 2009, **461**, 762–767.
- 117 B. Czermin, R. Melfi, D. McCabe, V. Seitz, A. Imhof and V. Pirrotta, *Cell*, 2002, **111**, 185–196.
- 118 H. Wang, L. Wang, H. Erdjument-Bromage, M. Vidal, P. Tempst, R. S. Jones and Y. Zhang, *Nature*, 2004, **431**, 873–878.
- 119 S. Varambally, S. M. Dhanasekaran, M. Zhou, T. R. Barrette, C. Kumar-Sinha, M. G. Sanda, D. Ghosh, K. J. Pienta, R. G. Sewalt, A. P. Otte, M. A. Rubin and A. M. Chinnaiyan, *Nature*, 2002, **419**, 624–629.
- 120 C. G. Kleer, Q. Cao, S. Varambally, R. Shen, I. Ota, S. A. Tomlins, D. Ghosh, R. G. Sewalt, A. P. Otte, D. F. Hayes, M. S. Sabel, D. Livant, S. J. Weiss, M. A. Rubin and A. M. Chinnaiyan, *Proc. Natl. Acad. Sci. U. S. A.*, 2003, **100**, 11606–11611.
- 121 B. D. Karanikolas, M. L. Figueiredo and L. Wu, *Mol. Cancer Res.*, 2009, **7**, 1456–1465.
- 122 M. Yamagishi, M. Hori, D. Fujikawa, T. Ohsugi, D. Honma, N. Adachi, H. Katano, T. Hishima, S. Kobayashi, K. Nakano, M. Nakashima, M. Iwanaga, A. Utsunomiya,



- Y. Tanaka, S. Okada, K. Tsukasaki, K. Tobinai, K. Araki, T. Watanabe and K. Uchimar, *Cell Rep.*, 2019, **29**, 2321–2337 e2327.
- 123 S. Fujita, D. Honma, N. Adachi, K. Araki, E. Takamatsu, T. Katsumoto, K. Yamagata, K. Akashi, K. Aoyama, A. Iwama and I. Kitabayashi, *Leukemia*, 2018, **32**, 855–864.
- 124 S. K. Knutson, T. J. Wigle, N. M. Warholc, C. J. Sneeringer, C. J. Allain, C. R. Klaus, J. D. Sacks, A. Raimondi, C. R. Majer, J. Song, M. P. Scott, L. Jin, J. J. Smith, E. J. Olhava, R. Chesworth, M. P. Moyer, V. M. Richon, R. A. Copeland, H. Keilhack, R. M. Pollock and K. W. Kuntz, *Nat. Chem. Biol.*, 2012, **8**, 890–896.
- 125 S. K. Knutson, N. M. Warholc, T. J. Wigle, C. R. Klaus, C. J. Allain, A. Raimondi, M. Porter Scott, R. Chesworth, M. P. Moyer, R. A. Copeland, V. M. Richon, R. M. Pollock, K. W. Kuntz and H. Keilhack, *Proc. Natl. Acad. Sci. U. S. A.*, 2013, **110**, 7922–7927.
- 126 S. K. Knutson, S. Kawano, Y. Minoshima, N. M. Warholc, K. C. Huang, Y. Xiao, T. Kadowaki, M. Uesugi, G. Kuznetsov, N. Kumar, T. J. Wigle, C. R. Klaus, C. J. Allain, A. Raimondi, N. J. Waters, J. J. Smith, M. Porter-Scott, R. Chesworth, M. P. Moyer, R. A. Copeland, V. M. Richon, T. Uenaka, R. M. Pollock, K. W. Kuntz, A. Yokoi and H. Keilhack, *Mol. Cancer Ther.*, 2014, **13**, 842–854.
- 127 I. Epizyme, Study of Tazemetostat With Enzalutamide or Abiraterone/Prednisone in Subjects With Castration Resistant Prostate Cancer Who Have Not Received Chemotherapy, <https://ClinicalTrials.gov/show/NCT04179864>, 2019.
- 128 I. Epizyme, Open-Label, Multicenter, Phase 1/2 Study of Tazemetostat (EZH2 Histone Methyl Transferase HMT Inhibitor) as a Single Agent in Subjects With Adv. Solid Tumors or With B-cell Lymphomas and Tazemetostat in Combination With Prednisolone in Subjects With DLBCL, <https://ClinicalTrials.gov/show/NCT01897571>, 2013.
- 129 N. C. Institute, Tazemetostat and Pembrolizumab in Treating Patients With Locally Advanced or Metastatic Urothelial Carcinoma, <https://ClinicalTrials.gov/show/NCT03854474>, 2019.
- 130 J. E. Campbell, K. W. Kuntz, S. K. Knutson, N. M. Warholc, H. Keilhack, T. J. Wigle, A. Raimondi, C. R. Klaus, N. Rioux, A. Yokoi, S. Kawano, Y. Minoshima, H. W. Choi, M. Porter Scott, N. J. Waters, J. J. Smith, R. Chesworth, M. P. Moyer and R. A. Copeland, *ACS Med. Chem. Lett.*, 2015, **6**, 491–495.
- 131 B. Lu, X. Shen, L. Zhang, D. Liu, C. Zhang, J. Cao, R. Shen, J. Zhang, D. Wang, H. Wan, Z. Xu, M. H. Ho, M. Zhang, L. Zhang, F. He and W. Tao, *ACS Med. Chem. Lett.*, 2018, **9**, 98–102.
- 132 M. T. McCabe, H. M. Ott, G. Ganji, S. Korenchuk, C. Thompson, G. S. Van Aller, Y. Liu, A. P. Graves, A. Della Pietra, 3rd, E. Diaz, L. V. LaFrance, M. Mellinger, C. Duquenne, X. Tian, R. G. Kruger, C. F. McHugh, M. Brandt, W. H. Miller, D. Dhanak, S. K. Verma, P. J. Tummino and C. L. Creasy, *Nature*, 2012, **492**, 108–112.
- 133 D. Zeng, M. Liu and J. Pan, *Oncotarget*, 2017, **8**, 3396–3411.
- 134 GlaxoSmithKline, A Study to Investigate the Safety, Pharmacokinetics, Pharmacodynamics and Clinical Activity of GSK2816126 in Subjects With Relapsed/Refractory Diffuse Large B Cell Lymphoma, Transformed Follicular Lymphoma, Other Non-Hodgkin's Lymphomas, Solid Tumors and Multiple Myeloma, <https://ClinicalTrials.gov/show/NCT02082977>, 2014.
- 135 S. Huang, Z. Wang, J. Zhou, J. Huang, L. Zhou, J. Luo, Y. Y. Wan, H. Long and B. Zhu, *Cancer Res.*, 2019, **79**, 2009–2020.
- 136 M. Bratkowski, X. Yang and X. Liu, *Sci. Rep.*, 2018, **8**, 9092.
- 137 S. K. Verma, X. Tian, L. V. LaFrance, C. Duquenne, D. P. Suarez, K. A. Newlander, S. P. Romeril, J. L. Burgess, S. W. Grant, J. A. Brackley, A. P. Graves, D. A. Scherzer, A. Shu, C. Thompson, H. M. Ott, G. S. Aller, C. A. Machutta, E. Diaz, Y. Jiang, N. W. Johnson, S. D. Knight, R. G. Kruger, M. T. McCabe, D. Dhanak, P. J. Tummino, C. L. Creasy and W. H. Miller, *ACS Med. Chem. Lett.*, 2012, **3**, 1091–1096.
- 138 W. Qi, H. Chan, L. Teng, L. Li, S. Chuai, R. Zhang, J. Zeng, M. Li, H. Fan, Y. Lin, J. Gu, O. Ardayfio, J. H. Zhang, X. Yan, J. Fang, Y. Mi, M. Zhang, T. Zhou, G. Feng, Z. Chen, G. Li, T. Yang, K. Zhao, X. Liu, Z. Yu, C. X. Lu, P. Atadja and E. Li, *Proc. Natl. Acad. Sci. U. S. A.*, 2012, **109**, 21360–21365.
- 139 R. G. Vaswani, V. S. Gehling, L. A. Dakin, A. S. Cook, C. G. Nasveschuk, M. Duplessis, P. Iyer, S. Balasubramanian, F. Zhao, A. C. Good, R. Campbell, C. Lee, N. Cantone, R. T. Cummings, E. Normant, S. F. Bellon, B. K. Albrecht, J. C. Harmange, P. Trojer, J. E. Audia, Y. Zhang, N. Justin, S. Chen, J. R. Wilson and S. J. Gamblin, *J. Med. Chem.*, 2016, **59**, 9928–9941.
- 140 C. Pharmaceuticals, A Study Evaluating CPI-1205 in Patients With B-Cell Lymphomas, <https://ClinicalTrials.gov/show/NCT02395601>, 2015.
- 141 C. Pharmaceuticals, ProSTAR: A Study Evaluating CPI-1205 in Patients With Metastatic Castration Resistant Prostate Cancer, <https://ClinicalTrials.gov/show/NCT03480646>, 2017.
- 142 S. Goswami, I. Apostolou, J. Zhang, J. Skepner, S. Anandhan, X. Zhang, L. Xiong, P. Trojer, A. Aparicio, S. K. Subudhi, J. P. Allison, H. Zhao and P. Sharma, *J. Clin. Invest.*, 2018, **128**, 3813–3818.
- 143 X. Song, T. Gao, N. Wang, Q. Feng, X. You, T. Ye, Q. Lei, Y. Zhu, M. Xiong, Y. Xia, F. Yang, Y. Shi, Y. Wei, L. Zhang and L. Yu, *Sci. Rep.*, 2016, **6**, 20864.
- 144 P. P. Kung, P. Bingham, A. Brooun, M. Collins, Y. L. Deng, D. Dinh, C. Fan, K. S. Gajiwala, R. Grantner, H. J. Gukasyan, W. Hu, B. Huang, R. Kania, S. E. Kephart, C. Krivacic, R. A. Kumpf, P. Khamphavong, M. Kraus, W. Liu, K. A. Maegley, L. Nguyen, S. Ren, D. Richter, R. A. Rollins, N. Sach, S. Sharma, J. Sherrill, J. Spangler, A. E. Stewart, S. Sutton, S. Uryu, D. Verhelle, H. Wang, S. Wang, M. Wythes, S. Xin, S. Yamazaki, H. Zhu, J. Zhu, L. Zehnder and M. Edwards, *J. Med. Chem.*, 2018, **61**, 650–665.
- 145 Pfizer, PF-06821497 Treatment Of Relapsed/Refractory SCLC, Castration Resistant Prostate Cancer, and Follicular Lymphoma, <https://ClinicalTrials.gov/show/NCT03460977>, 2018.



- 146 K. D. Konze, A. Ma, F. Li, D. Barsyte-Lovejoy, T. Parton, C. J. Macnevin, F. Liu, C. Gao, X. P. Huang, E. Kuznetsova, M. Rougie, A. Jiang, S. G. Pattenden, J. L. Norris, L. I. James, B. L. Roth, P. J. Brown, S. V. Frye, C. H. Arrowsmith, K. M. Hahn, G. G. Wang, M. Vedadi and J. Jin, *ACS Chem. Biol.*, 2013, **8**, 1324–1334.
- 147 B. Xu, D. M. On, A. Ma, T. Parton, K. D. Konze, S. G. Pattenden, D. F. Allison, L. Cai, S. Rockowitz, S. Liu, Y. Liu, F. Li, M. Vedadi, S. V. Frye, B. A. Garcia, D. Zheng, J. Jin and G. G. Wang, *Blood*, 2015, **125**, 346–357.
- 148 D. Honma, O. Kanno, J. Watanabe, J. Kinoshita, M. Hirasawa, E. Nosaka, M. Shiroishi, T. Takizawa, I. Yasumatsu, T. Horiuchi, A. Nakao, K. Suzuki, T. Yamasaki, K. Nakajima, M. Hayakawa, T. Yamazaki, A. S. Yadav and N. Adachi, *Cancer Sci.*, 2017, **108**, 2069–2078.
- 149 W. Qi, K. Zhao, J. Gu, Y. Huang, Y. Wang, H. Zhang, M. Zhang, J. Zhang, Z. Yu, L. Li, L. Teng, S. Chuai, C. Zhang, M. Zhao, H. Chan, Z. Chen, D. Fang, Q. Fei, L. Feng, L. Feng, Y. Gao, H. Ge, X. Ge, G. Li, A. Lingel, Y. Lin, Y. Liu, F. Luo, M. Shi, L. Wang, Z. Wang, Y. Yu, J. Zeng, C. Zeng, L. Zhang, Q. Zhang, S. Zhou, C. Oyang, P. Atadja and E. Li, *Nat. Chem. Biol.*, 2017, **13**, 381–388.
- 150 Y. He, S. Selvaraju, M. L. Curtin, C. G. Jakob, H. Zhu, K. M. Comess, B. Shaw, J. The, E. Lima-Fernandes, M. M. Szewczyk, D. Cheng, K. L. Klinge, H. Q. Li, M. Pliushchev, M. A. Algire, D. Maag, J. Guo, J. Dietrich, S. C. Panchal, A. M. Petros, R. F. Sweis, M. Torrent, L. J. Bigelow, G. Senisterra, F. Li, S. Kennedy, Q. Wu, D. J. Osterling, D. J. Lindley, W. Gao, S. Galasinski, D. Barsyte-Lovejoy, M. Vedadi, F. G. Buchanan, C. H. Arrowsmith, G. G. Chiang, C. Sun and W. N. Pappano, *Nat. Chem. Biol.*, 2017, **13**, 389–395.
- 151 H. Dong, S. Liu, X. Zhang, S. Chen, L. Kang, Y. Chen, S. Ma, X. Fu, Y. Liu, H. Zhang and B. Zou, *Cancer Res.*, 2019, **79**, 5587–5596.
- 152 N. Pharmaceuticals and Novartis, Safety and Efficacy of MAK683 in Adult Patients With Advanced Malignancies, <https://ClinicalTrials.gov/show/NCT02900651>, 2016.
- 153 A. Ma, E. Stratikopoulos, K. S. Park, J. Wei, T. C. Martin, X. Yang, M. Schwarz, V. Leshchenko, A. Rialdi, B. Dale, A. Lagana, E. Guccione, S. Parekh, R. Parsons and J. Jin, *Nat. Chem. Biol.*, 2020, **16**, 214–222.
- 154 Z. Liu, X. Hu, Q. Wang, X. Wu, Q. Zhang, W. Wei, X. Su, H. He, S. Zhou, R. Hu, T. Ye, Y. Zhu, N. Wang and L. Yu, *J. Med. Chem.*, 2021, **64**, 2829–2848.
- 155 X. Wang, W. Cao, J. Zhang, M. Yan, Q. Xu, X. Wu, L. Wan, Z. Zhang, C. Zhang, X. Qin, M. Xiao, D. Ye, Y. Liu, Z. Han, S. Wang, L. Mao, W. Wei and W. Chen, *EMBO J.*, 2017, **36**, 1243–1260.
- 156 J. H. Hsu, T. Rasmusson, J. Robinson, F. Pahl, J. Read, S. Kawatkar, O. D. Dh, S. Bagal, E. Code, P. Rawlins, A. Argyrou, R. Tomlinson, N. Gao, X. Zhu, E. Chiarparin, K. Jacques, M. Shen, H. Woods, E. Bednarski, D. M. Wilson, L. Drew, M. P. Castaldi, S. Fawell and A. Bloecher, *Cell Chem. Biol.*, 2020, **27**, 41–46 e17.
- 157 F. Potjewyd, A. W. Turner, J. Beri, J. M. Rectenwald, J. L. Norris-Drouin, S. H. Cholensky, D. M. Margolis, K. H. Pearce, L. E. Herring and L. I. James, *Cell Chem. Biol.*, 2020, **27**, 47–56 e15.
- 158 K. Nishioka, S. Chuikov, K. Sarma, H. Erdjument-Bromage, C. D. Allis, P. Tempst and D. Reinberg, *Genes Dev.*, 2002, **16**, 479–489.
- 159 H. Wang, R. Cao, L. Xia, H. Erdjument-Bromage, C. Borchers, P. Tempst and Y. Zhang, *Mol. Cell*, 2001, **8**, 1207–1217.
- 160 T. Kwon, J. H. Chang, E. Kwak, C. W. Lee, A. Joachimiak, Y. C. Kim, J. Lee and Y. Cho, *EMBO J.*, 2003, **22**, 292–303.
- 161 J. R. Wilson, C. Jing, P. A. Walker, S. R. Martin, S. A. Howell, G. M. Blackburn, S. J. Gamblin and B. Xiao, *Cell*, 2002, **111**, 105–115.
- 162 P. Hu, S. Wang and Y. Zhang, *J. Am. Chem. Soc.*, 2008, **130**, 3806–3813.
- 163 S. Chuikov, J. K. Kurash, J. R. Wilson, B. Xiao, N. Justin, G. S. Ivanov, K. McKinney, P. Tempst, C. Prives, S. J. Gamblin, N. A. Barlev and D. Reinberg, *Nature*, 2004, **432**, 353–360.
- 164 J. Wang, S. Hevi, J. K. Kurash, H. Lei, F. Gay, J. Bajko, H. Su, W. Sun, H. Chang, G. Xu, F. Gaudet, E. Li and T. Chen, *Nat. Genet.*, 2009, **41**, 125–129.
- 165 P.-O. Estève, H. G. Chin, J. Benner, G. R. Feehery, M. Samaranayake, G. A. Horwitz, S. E. Jacobsen and S. Pradhan, *Proc. Natl. Acad. Sci. U. S. A.*, 2009, **106**, 5076–5081.
- 166 Y. Li, M. A. Reddy, F. Miao, N. Shanmugam, J. K. Yee, D. Hawkins, B. Ren and R. Natarajan, *J. Biol. Chem.*, 2008, **283**, 26771–26781.
- 167 B. Duan, J. Bai, J. Qiu, J. Wang, C. Tong, X. Wang, J. Miao, Z. Li, W. Li, J. Yang and C. Huang, *EBioMedicine*, 2018, **37**, 134–143.
- 168 Y. Gu, X. Wang, H. Liu, G. Li, W. Yu and Q. Ma, *Oncol. Rep.*, 2018, **40**, 1863–1874.
- 169 S. Mori, K. Iwase, N. Iwanami, Y. Tanaka, H. Kagechika and T. Hirano, *Bioorg. Med. Chem.*, 2010, **18**, 8158–8166.
- 170 H. Niwa, N. Handa, Y. Tomabechi, K. Honda, M. Toyama, N. Ohsawa, M. Shirouzu, H. Kagechika, T. Hirano, T. Umehara and S. Yokoyama, *Acta Crystallogr., Sect. D: Biol. Crystallogr.*, 2013, **69**, 595–602.
- 171 Y. Takemoto, A. Ito, H. Niwa, M. Okamura, T. Fujiwara, T. Hirano, N. Handa, T. Umehara, T. Sonoda, K. Ogawa, M. Tariq, N. Nishino, S. Dan, H. Kagechika, T. Yamori, S. Yokoyama and M. Yoshida, *J. Med. Chem.*, 2016, **59**, 3650–3660.
- 172 T. Hirano, T. Fujiwara, H. Niwa, M. Hirano, K. Ohira, Y. Okazaki, S. Sato, T. Umehara, Y. Maemoto, A. Ito, M. Yoshida and H. Kagechika, *ChemMedChem*, 2018, **13**, 1530–1540.
- 173 F. Meng, S. Cheng, H. Ding, S. Liu, Y. Liu, K. Zhu, S. Chen, J. Lu, Y. Xie, L. Li, R. Liu, Z. Shi, Y. Zhou, Y. C. Liu, M. Zheng, H. Jiang, W. Lu, H. Liu and C. Luo, *J. Med. Chem.*, 2015, **58**, 8166–8181.
- 174 D. Barsyte-Lovejoy, F. Li, M. J. Oudhoff, J. H. Tatlock, A. Dong, H. Zeng, H. Wu, S. A. Freeman, M. Schapira, G. A. Senisterra,



- E. Kuznetsova, R. Marcellus, A. Allali-Hassani, S. Kennedy, J. P. Lambert, A. L. Couzens, A. Aman, A. C. Gingras, R. Al-Awar, P. V. Fish, B. S. Gerstenberger, L. Roberts, C. L. Benn, R. L. Grimley, M. J. Braam, F. M. Rossi, M. Sudol, P. J. Brown, M. E. Bunnage, D. R. Owen, C. Zaph, M. Vedadi and C. H. Arrowsmith, *Proc. Natl. Acad. Sci. U. S. A.*, 2014, **111**, 12853–12858.
- 175 B. Tang, B. Li, B. Li, J. Qin, J. Zhao, J. Xu, Y. Qiu, Z. Wu and M. Fang, *RSC Adv.*, 2019, **9**, 9218–9227.
- 176 M. A. Brown, R. J. Sims, 3rd, P. D. Gottlieb and P. W. Tucker, *Mol. Cancer*, 2006, **5**, 26.
- 177 S. Weirich, M. K. Schuhmacher, S. Kudithipudi, C. Lungu, A. D. Ferguson and A. Jeltsch, *ChemBioChem*, 2020, **21**, 256–264.
- 178 J. Huang, L. Perez-Burgos, B. J. Placek, R. Sengupta, M. Richter, J. A. Dorsey, S. Kubicek, S. Opravil, T. Jenuwein and S. L. Berger, *Nature*, 2006, **444**, 629–632.
- 179 B. Sesé, M. J. Barrero, M. C. Fabregat, V. Sander and J. C. Izpisua Belmonte, *Int. J. Dev. Biol.*, 2013, **57**, 357–364.
- 180 F. Diehl, M. A. Brown, M. J. van Amerongen, T. Novoyatleva, A. Wietelmann, J. Harriss, F. Ferrazzi, T. Böttger, R. P. Harvey, P. W. Tucker and F. B. Engel, *PLoS One*, 2010, **5**, e9748.
- 181 S. Komatsu, D. Ichikawa, S. Hirajima, H. Nagata, Y. Nishimura, T. Kawaguchi, M. Miyamae, W. Okajima, T. Ohashi, H. Konishi, A. Shiozaki, H. Fujiwara, K. Okamoto, H. Tsuda, I. Imoto, J. Inazawa and E. Otsuji, *Br. J. Cancer*, 2015, **112**, 357–364.
- 182 N. Reynoird, P. K. Mazur, T. Stellfeld, N. M. Flores, S. M. Lofgren, S. M. Carlson, E. Brambilla, P. Hainaut, E. B. Kaznowska, C. H. Arrowsmith, P. Khatri, C. Stresemann, O. Gozani and J. Sage, *Genes Dev.*, 2016, **30**, 772–785.
- 183 S. Komatsu, I. Imoto, H. Tsuda, K. I. Kozaki, T. Muramatsu, Y. Shimada, S. Aiko, Y. Yoshizumi, D. Ichikawa, E. Otsuji and J. Inazawa, *Carcinogenesis*, 2009, **30**, 1139–1146.
- 184 L. Liu, S. Kimball, H. Liu, A. Holowatyj and Z. Q. Yang, *Oncotarget*, 2015, **6**, 2466–2482.
- 185 A. D. Ferguson, N. A. Larsen, T. Howard, H. Pollard, I. Green, C. Grande, T. Cheung, R. Garcia-Arenas, S. Cowen, J. Wu, R. Godin, H. Chen and N. Keen, *Structure*, 2011, **19**, 1262–1273.
- 186 R. F. Sweis, Z. Wang, M. Algire, C. H. Arrowsmith, P. J. Brown, G. G. Chiang, J. Guo, C. G. Jakob, S. Kennedy, F. Li, D. Maag, B. Shaw, N. B. Soni, M. Vedadi and W. N. Pappano, *ACS Med. Chem. Lett.*, 2015, **6**, 695–700.
- 187 S. Cowen, EpiCongress 2013, Boston, MA, July 23–24, 2013, 2013.
- 188 S. Throner, S. Cowen, D. Russell, L. Dakin, H. W. Chen, N. Larsen, R. Godin, X. L. Zheng, A. Molina, J. Q. Wu, T. Cheung, T. Howard, R. Garcia-Arenas, N. Keen and A. Ferguson, *Abstr. Pap. Am. Chem. Soc.*, 2015, **250**, 1.
- 189 S. D. Cowen, D. Russell, L. A. Dakin, H. Chen, N. A. Larsen, R. Godin, S. Throner, X. Zheng, A. Molina, J. Wu, T. Cheung, T. Howard, R. Garcia-Arenas, N. Keen, C. S. Pendleton, J. A. Pietenpol and A. D. Ferguson, *J. Med. Chem.*, 2016, **59**, 11079–11097.
- 190 H. Nguyen, A. Allali-Hassani, S. Antonysamy, S. Chang, L. H. Chen, C. Curtis, S. Emtage, L. Fan, T. Gheyi, F. Li, S. Liu, J. R. Martin, D. Mendel, J. B. Olsen, L. Pelletier, T. Shatseva, S. Wu, F. F. Zhang, C. H. Arrowsmith, P. J. Brown, R. M. Campbell, B. A. Garcia, D. Barsyte-Lovejoy, M. Mader and M. Vedadi, *J. Biol. Chem.*, 2015, **290**, 13641–13653.
- 191 E. Eggert, R. C. Hillig, S. Koehr, D. Stockigt, J. Weiske, N. Barak, J. Mowat, T. Brumby, C. D. Christ, A. Ter Laak, T. Lang, A. E. Fernandez-Montalvan, V. Badock, H. Weinmann, I. V. Hartung, D. Barsyte-Lovejoy, M. Szewczyk, S. Kennedy, F. Li, M. Vedadi, P. J. Brown, V. Santhakumar, C. H. Arrowsmith, T. Stellfeld and C. Stresemann, *J. Med. Chem.*, 2016, **59**, 4578–4600.
- 192 M. J. Thomenius, J. Totman, D. Harvey, L. H. Mitchell, T. V. Riera, K. Cosmopoulos, A. R. Grassian, C. Klaus, M. Foley, E. A. Admirand, H. Jahic, C. Majer, T. Wigle, S. L. Jacques, J. Gureasko, D. Brach, T. Lingaraj, K. West, S. Smith, N. Rioux, N. J. Waters, C. Tang, A. Raimondi, M. Munchhof, J. E. Mills, S. Ribich, M. Porter Scott, K. W. Kuntz, W. P. Janzen, M. Moyer, J. J. Smith, R. Chesworth, R. A. Copeland and P. A. Boriack-Sjodin, *PLoS One*, 2018, **13**, e0197372.
- 193 A. P. Taylor, M. Szewczyk, S. Kennedy, V. V. Trush, H. Wu, H. Zeng, A. Dong, R. Ferreira de Freitas, J. Tatlock, R. A. Kumpf, M. Wythes, A. Casimiro-Garcia, R. A. Denny, M. D. Parikh, F. Li, D. Barsyte-Lovejoy, M. Schapira, M. Vedadi, P. J. Brown, C. H. Arrowsmith and D. R. Owen, *J. Med. Chem.*, 2019, **62**, 7669–7683.
- 194 G. S. Van Aller, N. Reynoird, O. Barbash, M. Huddleston, S. Liu, A. F. Zmoos, P. McDevitt, R. Sinnamon, B. Le, G. Mas, R. Annan, J. Sage, B. A. Garcia, P. J. Tummino, O. Gozani and R. G. Kruger, *Epigenetics*, 2012, **7**, 340–343.
- 195 S. Xu, J. Wu, B. Sun, C. Zhong and J. Ding, *Nucleic Acids Res.*, 2011, **39**, 4438–4449.
- 196 M. A. Brown, K. Foreman, J. Harriss, C. Das, L. Zhu, M. Edwards, S. Shaaban and H. Tucker, *Oncotarget*, 2015, **6**, 4005–4019.
- 197 M. Kunizaki, R. Hamamoto, F. P. Silva, K. Yamaguchi, T. Nagayasu, M. Shibuya, Y. Nakamura and Y. Furukawa, *Cancer Res.*, 2007, **67**, 10759–10765.
- 198 P. K. Mazur, N. Reynoird, P. Khatri, P. W. Jansen, A. W. Wilkinson, S. Liu, O. Barbash, G. S. Van Aller, M. Huddleston, D. Dhanak, P. J. Tummino, R. G. Kruger, B. A. Garcia, A. J. Butte, M. Vermeulen, J. Sage and O. Gozani, *Nature*, 2014, **510**, 283–287.
- 199 S. Suzuki, Y. Nozawa, S. Tsukamoto, T. Kaneko, H. Imai and N. Minami, *Reproduction*, 2015, **150**, 21–30.
- 200 X. G. Luo, T. Xi, S. Guo, Z. P. Liu, N. Wang, Y. Jiang and T. C. Zhang, *IUBMB Life*, 2009, **61**, 679–684.
- 201 A. M. Cock-Rada, S. Medjkane, N. Janski, N. Yousfi, M. Perichon, M. Chaussepied, J. Chluba, G. Langsley and J. B. Weitzman, *Cancer Res.*, 2012, **72**, 810–820.
- 202 R. Hamamoto, Y. Furukawa, M. Morita, Y. Iimura, F. P. Silva, M. Li, R. Yagyu and Y. Nakamura, *Nat. Cell Biol.*, 2004, **6**, 731–740.



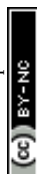
- 203 A. Peserico, A. Germani, P. Sanese, A. J. Barbosa, V. Di Virgilio, R. Fittipaldi, E. Fabini, C. Bertucci, G. Varchi, M. P. Moyer, G. Caretti, A. Del Rio and C. Simone, *J. Cell. Physiol.*, 2015, **230**, 2447–2460.
- 204 G. S. Van Aller, A. P. Graves, P. A. Elkins, W. G. Bonnette, P. J. McDevitt, F. Zappacosta, R. S. Annan, T. W. Dean, D. S. Su, C. L. Carpenter, H. P. Mohammad and R. G. Kruger, *Structure*, 2016, **24**, 774–781.
- 205 L. H. Mitchell, P. A. Boriack-Sjodin, S. Smith, M. Thomenius, N. Rioux, M. Munchhof, J. E. Mills, C. Klaus, J. Totman, T. V. Riera, A. Raimondi, S. L. Jacques, K. West, M. Foley, N. J. Waters, K. W. Kuntz, T. J. Wigle, M. P. Scott, R. A. Copeland, J. J. Smith and R. Chesworth, *ACS Med. Chem. Lett.*, 2016, **7**, 134–138.
- 206 C. Huang, S. S. Liew, G. R. Lin, A. Poulsen, M. J. Y. Ang, B. C. S. Chia, S. Y. Chew, Z. P. Kwek, J. L. K. Wee, E. H. Ong, P. Retna, N. Baburajendran, R. Li, W. Yu, X. Koh-Stenta, A. Ngo, S. Manesh, J. Fulwood, Z. Ke, H. H. Chung, S. Sepramaniam, X. H. Chew, N. Dinie, M. A. Lee, Y. S. Chew, C. B. Low, V. Pendharkar, V. Manoharan, S. Vuddagiri, K. Sangthongpitag, J. Joy, A. Matter, J. Hill, T. H. Keller and K. Foo, *ACS Med. Chem. Lett.*, 2019, **10**, 978–984.
- 207 D. S. Su, J. Qu, M. Schulz, C. W. Blackledge, H. Yu, J. Zeng, J. Burgess, A. Reif, M. Stern, R. Nagarajan, M. B. Pappalardi, K. Wong, A. P. Graves, W. Bonnette, L. Wang, P. Elkins, B. Knapp-Reed, J. D. Carson, C. McHugh, H. Mohammad, R. Kruger, J. Luengo, D. A. Heerding and C. L. Creasy, *ACS Med. Chem. Lett.*, 2020, **11**, 133–140.
- 208 V. O. Talibov, E. Fabini, E. FitzGerald, D. Tedesco, D. Eriksson, M. J. Talu, M. M. Rachman, F. Mihalic, E. Manoni, M. Naldi, P. Sanese, G. Forte, M. L. Signorile, X. Barril, C. Simone, M. Bartolini, D. Dobritzsch, A. Del Rio and U. H. Danielson, *ChemBioChem*, 2021, **22**, 1597–1608.
- 209 J. W. Edmunds, L. C. Mahadevan and A. L. Clayton, *EMBO J.*, 2008, **27**, 406–420.
- 210 B. D. Strahl, P. A. Grant, S. D. Briggs, Z. W. Sun, J. R. Bone, J. A. Caldwell, S. Mollah, R. G. Cook, J. Shabanowitz, D. F. Hunt and C. D. Allis, *Mol. Cell. Biol.*, 2002, **22**, 1298–1306.
- 211 I. Y. Park, R. T. Powell, D. N. Tripathi, R. Dere, T. H. Ho, T. L. Blasius, Y. C. Chiang, I. J. Davis, C. C. Fahey, K. E. Hacker, K. J. Verhey, M. T. Bedford, E. Jonasch, W. K. Rathmell and C. L. Walker, *Cell*, 2016, **166**, 950–962.
- 212 K. Chen, J. Liu, S. Liu, M. Xia, X. Zhang, D. Han, Y. Jiang, C. Wang and X. Cao, *Cell*, 2017, **170**, 492–506 e414.
- 213 M. K. Schuhmacher, S. Beldar, M. S. Khella, A. Brohm, J. Ludwig, W. Tempel, S. Weirich, J. Min and A. Jeltsch, *Commun. Biol.*, 2020, **3**, 511.
- 214 M. Li, H. P. Phatnani, Z. Guan, H. Sage, A. L. Greenleaf and P. Zhou, *Proc. Natl. Acad. Sci. U. S. A.*, 2005, **102**, 17636–17641.
- 215 X. J. Sun, J. Wei, X. Y. Wu, M. Hu, L. Wang, H. H. Wang, Q. H. Zhang, S. J. Chen, Q. H. Huang and Z. Chen, *J. Biol. Chem.*, 2005, **280**, 35261–35271.
- 216 I. Sanidas, C. Polyarchou, M. Hatzia Apostolou, S. A. Ezell, F. Kottakis, L. Hu, A. Guo, J. Xie, M. J. Comb, D. Iliopoulos and P. N. Tsiachlis, *Mol. Cell*, 2014, **53**, 577–590.
- 217 N. Cancer Genome Atlas Research, W. M. Linehan, P. T. Spellman, C. J. Ricketts, C. J. Creighton, S. S. Fei, C. Davis, D. A. Wheeler, B. A. Murray, L. Schmidt, C. D. Vocke, M. Peto, A. A. Al Mamun, E. Shinbrot, A. Sethi, S. Brooks, W. K. Rathmell, A. N. Brooks, K. A. Hoadley, A. G. Robertson, D. Brooks, R. Bowlby, S. Sadeghi, H. Shen, D. J. Weisenberger, M. Bootwalla, S. B. Baylin, P. W. Laird, A. D. Cherniack, G. Saksena, S. Haake, J. Li, H. Liang, Y. Lu, G. B. Mills, R. Akbani, M. D. Leiserson, B. J. Raphael, P. Anur, D. Bottaro, L. Albiges, N. Barnabas, T. K. Choueiri, B. Czerniak, A. K. Godwin, A. A. Hakimi, T. H. Ho, J. Hsieh, M. Ittmann, W. Y. Kim, B. Krishnan, M. J. Merino, K. R. Mills Shaw, V. E. Reuter, E. Reznik, C. S. Shelley, B. Shuch, S. Signoretti, R. Srinivasan, P. Tamboli, G. Thomas, S. Tickoo, K. Burnett, D. Crain, J. Gardner, K. Lau, D. Mallory, S. Morris, J. D. Paulauskis, R. J. Penny, C. Shelton, W. T. Shelton, M. Sherman, E. Thompson, P. Yena, M. T. Avedon, J. Bowen, J. M. Gastier-Foster, M. Gerken, K. M. Leraas, T. M. Lichtenberg, N. C. Ramirez, T. Santos, L. Wise, E. Zmuda, J. A. Demchok, I. Felau, C. M. Hutter, M. Sheth, H. J. Sofia, R. Tarnuzzer, Z. Wang, L. Yang, J. C. Zenklusen, J. Zhang, B. Ayala, J. Baboud, S. Chudamani, J. Liu, L. Lolla, R. Nares, T. Pihl, Q. Sun, Y. Wan, Y. Wu, A. Ally, M. Balasundaram, S. Balu, R. Beroukhi, T. Bodenheimer, C. Buhay, Y. S. Butterfield, R. Carlsen, S. L. Carter, H. Chao, E. Chuah, A. Clarke, K. R. Covington, M. Dahdouli, N. Dewal, N. Dhalla, H. V. Doddapaneni, J. A. Drummond, S. B. Gabriel, R. A. Gibbs, R. Guin, W. Hale, A. Hawes, D. N. Hayes, R. A. Holt, A. P. Hoyle, S. R. Jefferys, S. J. Jones, C. D. Jones, D. Kalra, C. Kovar, L. Lewis, J. Li, Y. Ma, M. A. Marra, M. Mayo, S. Meng, M. Meyerson, P. A. Mieczkowski, R. A. Moore, D. Morton, L. E. Mose, A. J. Mungall, D. Muzny, J. S. Parker, C. M. Perou, J. Roach, J. E. Schein, S. E. Schumacher, Y. Shi, J. V. Simons, P. Sipahimalani, T. Skelly, M. G. Soloway, C. Sougnez, A. Tam, D. Tan, N. Thiessen, U. Veluvolu, M. Wang, M. D. Wilkerson, T. Wong, J. Wu, L. Xi, J. Zhou, J. Bedford, F. Chen, Y. Fu, M. Gerstein, D. Haussler, K. Kasaian, P. Lai, S. Ling, A. Radenbaugh, D. Van Den Berg, J. N. Weinstein, J. Zhu, M. Albert, I. Alexopoulou, J. J. Andersen, J. T. Auman, J. Bartlett, S. Bastacky, J. Bergsten, M. L. Blute, L. Boice, R. J. Bollag, J. Boyd, E. Castle, Y. B. Chen, J. C. Cheville, E. Curley, B. Davies, A. DeVolk, R. Dhir, L. Dike, J. Eckman, J. Engel, J. Harr, R. Hrebinko, M. Huang, L. Huelsenbeck-Dill, M. Iacocca, B. Jacobs, M. Lobis, J. K. Maranchie, S. McMeekin, J. Myers, J. Nelson, J. Parfitt, A. Parwani, N. Petrelli, B. Rabeno, S. Roy, A. L. Salner, J. Slaton, M. Stanton, R. H. Thompson, L. Thorne, K. Tucker, P. M. Weinberger, C. Winemiller, L. A. Zach and R. Zuna, *N. Engl. J. Med.*, 2016, **374**, 135–145.



- 218 A. M. Fontebasso, J. Schwartzentruber, D. A. Khuong-Quang, X. Y. Liu, D. Sturm, A. Korshunov, D. T. Jones, H. Witt, M. Kool, S. Albrecht, A. Fleming, D. Hadjadj, S. Busche, P. Lepage, A. Montpetit, A. Staffa, N. Gerges, M. Zakrzewska, K. Zakrzewski, P. P. Liberski, P. Hauser, M. Garami, A. Klekner, L. Bogner, G. Zadeh, D. Faury, S. M. Pfister, N. Jabado and J. Majewski, *Acta Neuropathol.*, 2013, **125**, 659–669.
- 219 X. Zhu, F. He, H. Zeng, S. Ling, A. Chen, Y. Wang, X. Yan, W. Wei, Y. Pang, H. Cheng, C. Hua, Y. Zhang, X. Yang, X. Lu, L. Cao, L. Hao, L. Dong, W. Zou, J. Wu, X. Li, S. Zheng, J. Yan, J. Zhou, L. Zhang, S. Mi, X. Wang, L. Zhang, Y. Zou, Y. Chen, Z. Geng, J. Wang, J. Zhou, X. Liu, J. Wang, W. Yuan, G. Huang, T. Cheng and Q. F. Wang, *Nat. Genet.*, 2014, **46**, 287–293.
- 220 C. Kandath, M. D. McLellan, F. Vandin, K. Ye, B. Niu, C. Lu, M. Xie, Q. Zhang, J. F. McMichael, M. A. Wyczalkowski, M. D. M. Leiserson, C. A. Miller, J. S. Welch, M. J. Walter, M. C. Wendl, T. J. Ley, R. K. Wilson, B. J. Raphael and L. Ding, *Nature*, 2013, **502**, 333–339.
- 221 A. A. Hakimi, I. Ostrovskaya, B. Reva, N. Schultz, Y. B. Chen, M. Gonen, H. Liu, S. Takeda, M. H. Voss, S. K. Tickoo, V. E. Reuter, P. Russo, E. H. Cheng, C. Sander, R. J. Motzer, J. J. Hsieh and R. C. C. G. A. R. N. I. cc, *Clin. Cancer Res.*, 2013, **19**, 3259–3267.
- 222 N. Kanu, E. Gronroos, P. Martinez, R. A. Burrell, X. Yi Goh, J. Bartkova, A. Maya-Mendoza, M. Mistrik, A. J. Rowan, H. Patel, A. Rabinowitz, P. East, G. Wilson, C. R. Santos, N. McGranahan, S. Gulati, M. Gerlinger, N. J. Birkbak, T. Joshi, L. B. Alexandrov, M. R. Stratton, T. Powles, N. Matthews, P. A. Bates, A. Stewart, Z. Szallasi, J. Larkin, J. Bartek and C. Swanton, *Oncogene*, 2015, **34**, 5699–5708.
- 223 W. Zheng, G. Ibanez, H. Wu, G. Blum, H. Zeng, A. Dong, F. Li, T. Hajian, A. Allali-Hassani, M. F. Amaya, A. Siarheyeva, W. Yu, P. J. Brown, M. Schapira, M. Vedadi, J. Min and M. Luo, *J. Am. Chem. Soc.*, 2012, **134**, 18004–18014.
- 224 K. Nishioka, J. C. Rice, K. Sarma, H. Erdjument-Bromage, J. Werner, Y. Wang, S. Chuikov, P. Valenzuela, P. Tempst, R. Steward, J. T. Lis, C. D. Allis and D. Reinberg, *Mol. Cell*, 2002, **9**, 1201–1213.
- 225 T. S. Girish, R. K. McGinty and S. Tan, *J. Mol. Biol.*, 2016, **428**, 1531–1543.
- 226 X. Shi, I. Kachirskaya, H. Yamaguchi, L. E. West, H. Wen, E. W. Wang, S. Dutta, E. Appella and O. Gozani, *Mol. Cell*, 2007, **27**, 636–646.
- 227 G. K. Dhami, H. Liu, M. Galka, C. Voss, R. Wei, K. Muranko, T. Kaneko, S. P. Cregan, L. Li and S. S. Li, *Mol. Cell*, 2013, **50**, 565–576.
- 228 M. Takawa, H. S. Cho, S. Hayami, G. Toyokawa, M. Kogure, Y. Yamane, Y. Iwai, K. Maejima, K. Ueda, A. Masuda, N. Dohmae, H. I. Field, T. Tsunoda, T. Kobayashi, T. Akasu, M. Sugiyama, S. Ohnuma, Y. Atomi, B. A. Ponder, Y. Nakamura and R. Hamamoto, *Cancer Res.*, 2012, **72**, 3217–3227.
- 229 S. Jorgensen, I. Elvers, M. B. Trelle, T. Menzel, M. Eskildsen, O. N. Jensen, T. Helleday, K. Helin and C. S. Sorensen, *J. Cell Biol.*, 2007, **179**, 1337–1345.
- 230 G. Schotta, R. Sengupta, S. Kubicek, S. Malin, M. Kauer, E. Callen, A. Celeste, M. Pagani, S. Opravil, I. A. De La Rosa-Velazquez, A. Espejo, M. T. Bedford, A. Nussenzweig, M. Busslinger and T. Jenuwein, *Genes Dev.*, 2008, **22**, 2048–2061.
- 231 H. Oda, M. R. Hubner, D. B. Beck, M. Vermeulen, J. Hurwitz, D. L. Spector and D. Reinberg, *Mol. Cell*, 2010, **40**, 364–376.
- 232 F. Yang, L. Sun, Q. Li, X. Han, L. Lei, H. Zhang and Y. Shang, *EMBO J.*, 2012, **31**, 110–123.
- 233 R. Huang, Y. Yu, X. Zong, X. Li, L. Ma and Q. Zheng, *Cancer Lett.*, 2017, **390**, 1–10.
- 234 D. E. Williams, F. Izard, S. Arnould, D. S. Dalisay, C. Tantapakul, W. Maneerat, T. Matainaho, E. Julien and R. J. Andersen, *J. Org. Chem.*, 2016, **81**, 1324–1332.
- 235 D. E. Williams, D. S. Dalisay, F. Li, J. Amphlett, W. Maneerat, M. A. Chavez, Y. A. Wang, T. Matainaho, W. Yu, P. J. Brown, C. H. Arrowsmith, M. Vedadi and R. J. Andersen, *Org. Lett.*, 2013, **15**, 414–417.
- 236 G. Blum, G. Ibanez, X. Rao, D. Shum, C. Radu, H. Djaballah, J. C. Rice and M. Luo, *ACS Chem. Biol.*, 2014, **9**, 2471–2478.
- 237 A. Ma, W. Yu, F. Li, R. M. Bleich, J. M. Herold, K. V. Butler, J. L. Norris, V. Korboukh, A. Tripathy, W. P. Janzen, C. H. Arrowsmith, S. V. Frye, M. Vedadi, P. J. Brown and J. Jin, *J. Med. Chem.*, 2014, **57**, 6822–6833.
- 238 L. Herviou, F. Izard, O. Karmous-Gadacha, C. Gourzones, C. Bellanger, E. Desmedt, A. Ma, L. Vincent, G. Cartron, K. Vanderkerken, J. Jin, E. De Bruyne, C. Grimaud, E. Julien and J. Moreaux, *bioRxiv*, 2019, 776930, DOI: 10.1101/776930.
- 239 K. V. Butler, A. Ma, W. Yu, F. Li, W. Tempel, N. Babault, F. Pittella-Silva, J. Shao, J. Wang, M. Luo, M. Vedadi, P. J. Brown, C. H. Arrowsmith and J. Jin, *J. Med. Chem.*, 2016, **59**, 9881–9889.
- 240 H. Yang, J. J. Pesavento, T. W. Starnes, D. E. Cryderman, L. L. Wallrath, N. L. Kelleher and C. A. Mizzen, *J. Biol. Chem.*, 2008, **283**, 12085–12092.
- 241 H. Wu, A. Siarheyeva, H. Zeng, R. Lam, A. Dong, X. H. Wu, Y. Li, M. Schapira, M. Vedadi and J. Min, *FEBS Lett.*, 2013, **587**, 3859–3868.
- 242 X. Lu, M. D. Simon, J. V. Chodaparambil, J. C. Hansen, K. M. Shokat and K. Luger, *Nat. Struct. Mol. Biol.*, 2008, **15**, 1122–1124.
- 243 G. Schotta, M. Lachner, K. Sarma, A. Ebert, R. Sengupta, G. Reuter, D. Reinberg and T. Jenuwein, *Genes Dev.*, 2004, **18**, 1251–1262.
- 244 R. Benetti, S. Gonzalo, I. Jaco, G. Schotta, P. Klatt, T. Jenuwein and M. A. Blasco, *J. Cell Biol.*, 2007, **178**, 925–936.
- 245 R. M. Marion, G. Schotta, S. Ortega and M. A. Blasco, *PLoS One*, 2011, **6**, e25680.
- 246 L. Piao, X. Yuan, L. Wang, X. Xu, M. Zhuang, J. Li, R. Kong and Z. Liu, *Oncol. Lett.*, 2020, **20**, 26.



- 247 M. Viotti, C. Wilson, M. McClelland, H. Koeppen, B. Haley, S. Jhunjhunwala, C. Klijn, Z. Modrusan, D. Arnott, M. Classon, J. P. Stephan and I. Mellman, *J. Cell Biol.*, 2018, **217**, 763–777.
- 248 A. Brohm, H. Elsayy, P. Rathert, S. Kudithipudi, T. Schoch, M. K. Schuhmacher, S. Weirich and A. Jeltsch, *J. Mol. Biol.*, 2019, **431**, 3068–3080.
- 249 K. D. Bromberg, T. R. Mitchell, A. K. Upadhyay, C. G. Jakob, M. A. Jhala, K. M. Comess, L. M. Lasko, C. Li, C. T. Tuzon, Y. Dai, F. Li, M. S. Eram, A. Nuber, N. B. Soni, V. Manaves, M. A. Algire, R. F. Sweis, M. Torrent, G. Schotta, C. Sun, M. R. Michaelides, A. R. Shoemaker, C. H. Arrowsmith, P. J. Brown, V. Santhakumar, A. Martin, J. C. Rice, G. G. Chiang, M. Vedadi, D. Barsyte-Lovejoy and W. N. Pappano, *Nat. Chem. Biol.*, 2017, **13**, 317–324.
- 250 R. Zheng, D. Liang, N. Jiang, J. Zhou, X. Long, X. Wang, M. Wu, C. Wu and J. Bao, *J. Biomol. Struct. Dyn.*, 2021, **39**, 526–537.
- 251 W. Li, W. Tian, G. Yuan, P. Deng, D. Sengupta, Z. Cheng, Y. Cao, J. Ren, Y. Qin, Y. Zhou, Y. Jia, O. Gozani, D. J. Patel and Z. Wang, *Nature*, 2021, **590**, 498–503.
- 252 M. Morishita, D. Mevius and E. di Luccio, *BMC Struct. Biol.*, 2014, **14**, 25.
- 253 J. Marango, M. Shimoyama, H. Nishio, J. A. Meyer, D. J. Min, A. Sirulnik, Y. Martinez-Martinez, M. Chesi, P. L. Bergsagel, M. M. Zhou, S. Waxman, B. A. Leibovitch, M. J. Walsh and J. D. Licht, *Blood*, 2008, **111**, 3145–3154.
- 254 J. Y. Kim, H. J. Kee, N. W. Choe, S. M. Kim, G. H. Eom, H. J. Baek, H. Kook, H. Kook and S. B. Seo, *Mol. Cell. Biol.*, 2008, **28**, 2023–2034.
- 255 G. V. Rayasam, O. Wendling, P. O. Angrand, M. Mark, K. Niederreither, L. Song, T. Lerouge, G. L. Hager, P. Chambon and R. Losson, *EMBO J.*, 2003, **22**, 3153–3163.
- 256 J. Douglas, S. Hanks, I. K. Temple, S. Davies, A. Murray, M. Upadhyaya, S. Tomkins, H. E. Hughes, T. R. Cole and N. Rahman, *Am. J. Hum. Genet.*, 2003, **72**, 132–143.
- 257 N. Kurotaki, K. Imaizumi, N. Harada, M. Masuno, T. Kondoh, T. Nagai, H. Ohashi, K. Naritomi, M. Tsukahara, Y. Makita, T. Sugimoto, T. Sonoda, T. Hasegawa, Y. Chinen, H. A. Tomita Ha, A. Kinoshita, T. Mizuguchi, K. Yoshiura Ki, T. Ohta, T. Kishino, Y. Fukushima, N. Niikawa and N. Matsumoto, *Nat. Genet.*, 2002, **30**, 365–366.
- 258 K. Nimura, K. Ura, H. Shiratori, M. Ikawa, M. Okabe, R. J. Schwartz and Y. Kaneda, *Nature*, 2009, **460**, 287–291.
- 259 N. L. Shannon, E. L. Maltby, A. S. Rigby and O. W. Quarrell, *J. Med. Genet.*, 2001, **38**, 674–679.
- 260 P. O. Angrand, F. Apiou, A. F. Stewart, B. Dutrillaux, R. Losson and P. Chambon, *Genomics*, 2001, **74**, 79–88.
- 261 M. Chesi, E. Nardini, R. S. C. Lim, K. D. Smith, W. M. Kuehl and P. L. Bergsagel, *Blood*, 1998, **92**, 3025–3034.
- 262 J. L. Brito, B. Walker, M. Jenner, N. J. Dickens, N. J. Brown, F. M. Ross, A. Avramidou, J. A. Irving, D. Gonzalez, F. E. Davies and G. J. Morgan, *Haematologica*, 2009, **94**, 78–86.
- 263 M. Morishita, D. E. H. F. Mevius, Y. Shen, S. Zhao and E. di Luccio, *Med. Chem. Res.*, 2017, **26**, 2038–2047.
- 264 H. Huang, C. A. Howard, S. Zari, H. J. Cho, S. Shukla, H. Li, J. Ndoj, P. Gonzalez-Alonso, C. Nikolaidis, J. Abbott, D. S. Rogawski, M. A. Potopnyk, K. Kempinska, H. Miao, T. Purohit, A. Henderson, A. Mapp, M. L. Sulis, A. Ferrando, J. Grembecka and T. Cierpicki, *Nat. Chem. Biol.*, 2020, **16**, 1403–1410.
- 265 D. Tisi, E. Chiarparin, E. Tamanini, P. Pathuri, J. E. Coyle, A. Hold, F. P. Holding, N. Amin, A. C. Martin, S. J. Rich, V. Berdini, J. Yon, P. Acklam, R. Burke, L. Drouin, J. E. Harmer, F. Jeganathan, R. L. van Montfort, Y. Newbatt, M. Tortorici, M. Westlake, A. Wood, S. Hoelder and T. D. Heightman, *ACS Chem. Biol.*, 2016, **11**, 3093–3105.
- 266 N. P. Coussens, S. C. Kales, M. J. Henderson, O. W. Lee, K. Y. Horiuchi, Y. Wang, Q. Chen, E. Kuznetsova, J. Wu, S. Chakka, D. M. Cheff, K. C. Cheng, P. Shinn, K. R. Brimacombe, M. Shen, A. Simeonov, M. Lal-Nag, H. Ma, A. Jadhav and M. D. Hall, *J. Biol. Chem.*, 2018, **293**, 13750–13765.
- 267 R. Ferreira de Freitas, Y. Liu, M. M. Szewczyk, N. Mehta, F. Li, D. McLeod, C. Zepeda-Velazquez, D. Dilworth, R. P. Hanley, E. Gibson, P. J. Brown, R. Al-Awar, L. I. James, C. H. Arrowsmith, D. Barsyte-Lovejoy, J. Min, M. Vedadi, M. Schapira and A. Allali-Hassani, *J. Med. Chem.*, 2021, **64**, 1584–1592.
- 268 D. Dilworth, R. P. Hanley, R. Ferreira de Freitas, A. Allali-Hassani, M. Zhou, N. Mehta, M. R. Marunde, S. Ackloo, R. A. Carvalho Machado, A. Khalili Yazdi, D. D. G. Owens, V. Vu, D. Y. Nie, M. Alqazzaz, E. Marcon, F. Li, I. Chau, A. Bolotokova, S. Qin, M. Lei, Y. Liu, M. M. Szewczyk, A. Dong, S. Kazemzadeh, T. Abramyan, I. K. Popova, N. W. Hall, M. J. Meiners, M. A. Cheek, E. Gibson, D. Kireev, J. F. Greenblatt, M.-C. Keogh, J. Min, P. J. Brown, M. Vedadi, C. H. Arrowsmith, D. Barsyte-Lovejoy, L. I. James and M. Schapira, *Nat. Chem. Biol.*, 2022, **18**, 56–63.
- 269 J. Bottcher, D. Dilworth, U. Reiser, R. A. Neumuller, M. Schleicher, M. Petronczki, M. Zeeb, N. Mischerikow, A. Allali-Hassani, M. M. Szewczyk, F. Li, S. Kennedy, M. Vedadi, D. Barsyte-Lovejoy, P. J. Brown, K. V. M. Huber, C. M. Rogers, C. I. Wells, O. Fedorov, K. Rumpel, A. Zoephel, M. Mayer, T. Wunberg, D. Bose, S. Zahn, H. Arnhof, H. Berger, C. Reiser, A. Hormann, T. Krammer, M. Corcokovic, B. Sharps, S. Winkler, D. Haring, X. L. Cockcroft, J. E. Fuchs, B. Mullauer, A. Weiss-Puxbaum, T. Gerstberger, G. Boehmelt, C. R. Vakoc, C. H. Arrowsmith, M. Pearson and D. B. McConnell, *Nat. Chem. Biol.*, 2019, **15**, 822–829.
- 270 T. A. Milne, S. D. Briggs, H. W. Brock, M. E. Martin, D. Gibbs, C. D. Allis and J. L. Hess, *Mol. Cell*, 2002, **10**, 1107–1117.
- 271 S. A. Shinsky, K. E. Monteith, S. Viggiano and M. S. Cosgrove, *J. Biol. Chem.*, 2015, **290**, 6361–6375.



- 272 Y. H. Takahashi, G. H. Westfield, A. N. Oleskie, R. C. Trievel, A. Shilatifard and G. Skiniotis, *Proc. Natl. Acad. Sci. U. S. A.*, 2011, **108**, 20526–20531.
- 273 N. T. Crump and T. A. Milne, *Cell. Mol. Life Sci.*, 2019, **76**, 2885–2898.
- 274 P. M. Ayton, E. H. Chen and M. L. Cleary, *Mol. Cell. Biol.*, 2004, **24**, 10470–10478.
- 275 L. Wu, S. Y. Lee, B. Zhou, U. T. Nguyen, T. W. Muir, S. Tan and Y. Dou, *Mol. Cell*, 2013, **49**, 1108–1120.
- 276 D. Hu, X. Gao, M. A. Morgan, H. M. Herz, E. R. Smith and A. Shilatifard, *Mol. Cell. Biol.*, 2013, **33**, 4745–4754.
- 277 C. Caslini, Z. Yang, M. El-Osta, T. A. Milne, R. K. Slany and J. L. Hess, *Cancer Res.*, 2007, **67**, 7275–7283.
- 278 S. Yin, J. Yang, B. Lin, W. Deng, Y. Zhang, X. Yi, Y. Shi, Y. Tao, J. Cai, C. I. Wu, G. Zhao, L. D. Hurst, J. Zhang, L. Hu and X. Kong, *Sci. Rep.*, 2014, **4**, 6036.
- 279 M. Le Gallo, M. L. Rudd, M. E. Urick, N. F. Hansen, S. Zhang, N. C. S. Program, F. Lozy, D. C. Sgroi, A. Vidal Bel, X. Matias-Guiu, R. R. Broaddus, K. H. Lu, D. A. Levine, D. G. Mutch, P. J. Goodfellow, H. B. Salvesen, J. C. Mullikin and D. W. Bell, *Cancer*, 2017, **123**, 3261–3268.
- 280 T. Sjoblom, S. Jones, L. D. Wood, D. W. Parsons, J. Lin, T. D. Barber, D. Mandelker, R. J. Leary, J. Ptak, N. Silliman, S. Szabo, P. Buckhaults, C. Farrell, P. Meeh, S. D. Markowitz, J. Willis, D. Dawson, J. K. Willson, A. F. Gazdar, J. Hartigan, L. Wu, C. Liu, G. Parmigiani, B. H. Park, K. E. Bachman, N. Papadopoulos, B. Vogelstein, K. W. Kinzler and V. E. Velculescu, *Science*, 2006, **314**, 268–274.
- 281 T. R. Chern, L. Liu, E. Petrunak, J. A. Stuckey, M. Wang, D. Bernard, H. Zhou, S. Lee, Y. Dou and S. Wang, *ACS Med. Chem. Lett.*, 2020, **11**, 1348–1352.
- 282 A. Patel, V. E. Vought, V. Dharmarajan and M. S. Cosgrove, *J. Biol. Chem.*, 2008, **283**, 32162–32175.
- 283 H. Karatas, E. C. Townsend, D. Bernard, Y. Dou and S. Wang, *J. Med. Chem.*, 2010, **53**, 5179–5185.
- 284 H. Karatas, E. C. Townsend, F. Cao, Y. Chen, D. Bernard, L. Liu, M. Lei, Y. Dou and S. Wang, *J. Am. Chem. Soc.*, 2013, **135**, 669–682.
- 285 F. Cao, E. C. Townsend, H. Karatas, J. Xu, L. Li, S. Lee, L. Liu, Y. Chen, P. Ouillette, J. Zhu, J. L. Hess, P. Atadja, M. Lei, Z. S. Qin, S. Malek, S. Wang and Y. Dou, *Mol. Cell*, 2014, **53**, 247–261.
- 286 H. Karatas, Y. Li, L. Liu, J. Ji, S. Lee, Y. Chen, J. Yang, L. Huang, D. Bernard, J. Xu, E. C. Townsend, F. Cao, X. Ran, X. Li, B. Wen, D. Sun, J. A. Stuckey, M. Lei, Y. Dou and S. Wang, *J. Med. Chem.*, 2017, **60**, 4818–4839.
- 287 G. Senisterra, H. Wu, A. Allali-Hassani, G. A. Wasney, D. Barsyte-Lovejoy, L. Dombrovski, A. Dong, K. T. Nguyen, D. Smil, Y. Bolshan, T. Hajian, H. He, A. Seitova, I. Chau, F. Li, G. Poda, J. F. Couture, P. J. Brown, R. Al-Awar, M. Schapira, C. H. Arrowsmith and M. Vedadi, *Biochem. J.*, 2013, **449**, 151–159.
- 288 Y. Bolshan, M. Getlik, E. Kuznetsova, G. A. Wasney, T. Hajian, G. Poda, K. T. Nguyen, H. Wu, L. Dombrovski, A. Dong, G. Senisterra, M. Schapira, C. H. Arrowsmith, P. J. Brown, R. Al-Awar, M. Vedadi and D. Smil, *ACS Med. Chem. Lett.*, 2013, **4**, 353–357.
- 289 D. D. Li, W. L. Chen, X. L. Xu, F. Jiang, L. Wang, Y. Y. Xie, X. J. Zhang, X. K. Guo, Q. D. You and H. P. Sun, *Eur. J. Med. Chem.*, 2016, **118**, 1–8.
- 290 D. D. Li, Z. H. Wang, W. L. Chen, Y. Y. Xie, Q. D. You and X. K. Guo, *Bioorg. Med. Chem.*, 2016, **24**, 6109–6118.
- 291 D. D. Li, W. L. Chen, Z. H. Wang, Y. Y. Xie, X. L. Xu, Z. Y. Jiang, X. J. Zhang, Q. D. You and X. K. Guo, *Eur. J. Med. Chem.*, 2016, **124**, 480–489.
- 292 M. Getlik, D. Smil, C. Zepeda-Velazquez, Y. Bolshan, G. Poda, H. Wu, A. Dong, E. Kuznetsova, R. Marcellus, G. Senisterra, L. Dombrovski, T. Hajian, T. Kiyota, M. Schapira, C. H. Arrowsmith, P. J. Brown, M. Vedadi and R. Al-Awar, *J. Med. Chem.*, 2016, **59**, 2478–2496.
- 293 F. Grebien, M. Vedadi, M. Getlik, R. Giamb Bruno, A. Grover, R. Avellino, A. Skucha, S. Vittori, E. Kuznetsova, D. Smil, D. Barsyte-Lovejoy, F. Li, G. Poda, M. Schapira, H. Wu, A. Dong, G. Senisterra, A. Stukalov, K. V. M. Huber, A. Schonegger, R. Marcellus, M. Bilban, C. Bock, P. J. Brown, J. Zuber, K. L. Bennett, R. Al-Awar, R. Delwel, C. Nerlov, C. H. Arrowsmith and G. Superti-Furga, *Nat. Chem. Biol.*, 2015, **11**, 571–578.
- 294 F. Wang, K. O. Jeon, J. M. Salovich, J. D. Macdonald, J. Alvarado, R. D. Gogliotti, J. Phan, E. T. Olejniczak, Q. Sun, S. Wang, D. Camper, J. P. Yuh, J. G. Shaw, J. Sai, O. W. Rossanese, W. P. Tansey, S. R. Stauffer and S. W. Fesik, *J. Med. Chem.*, 2018, **61**, 5623–5642.
- 295 A. Yokoyama, T. C. Somervaille, K. S. Smith, O. Rozenblatt-Rosen, M. Meyerson and M. L. Cleary, *Cell*, 2005, **123**, 207–218.
- 296 J. Grembecka, S. He, A. Shi, T. Purohit, A. G. Muntean, R. J. Sorenson, H. D. Showalter, M. J. Murai, A. M. Belcher, T. Hartley, J. L. Hess and T. Cierpicki, *Nat. Chem. Biol.*, 2012, **8**, 277–284.
- 297 A. Shi, M. J. Murai, S. He, G. Lund, T. Hartley, T. Purohit, G. Reddy, M. Chruszcz, J. Grembecka and T. Cierpicki, *Blood*, 2012, **120**, 4461–4469.
- 298 D. Borkin, S. He, H. Miao, K. Kempinska, J. Pollock, J. Chase, T. Purohit, B. Malik, T. Zhao, J. Wang, B. Wen, H. Zong, M. Jones, G. Danet-Desnoyers, M. L. Guzman, M. Talpaz, D. L. Bixby, D. Sun, J. L. Hess, A. G. Muntean, I. Maillard, T. Cierpicki and J. Grembecka, *Cancer Cell*, 2015, **27**, 589–602.
- 299 D. Borkin, J. Pollock, K. Kempinska, T. Purohit, X. Li, B. Wen, T. Zhao, H. Miao, S. Shukla, M. He, D. Sun, T. Cierpicki and J. Grembecka, *J. Med. Chem.*, 2016, **59**, 892–913.
- 300 D. Borkin, S. Klossowski, J. Pollock, H. Miao, B. M. Linhares, K. Kempinska, Z. Jin, T. Purohit, B. Wen, M. He, D. Sun, T. Cierpicki and J. Grembecka, *J. Med. Chem.*, 2018, **61**, 4832–4850.
- 301 S. Klossowski, H. Miao, K. Kempinska, T. Wu, T. Purohit, E. Kim, B. M. Linhares, D. Chen, G. Jih, E. Perkey, H. Huang, M. He, B. Wen, Y. Wang, K. Yu, S. C. Lee, G. Danet-Desnoyers, W. Trotman, M. Kandarpa, A. Cotton,



- O. Abdel-Wahab, H. Lei, Y. Dou, M. Guzman, L. Peterson, T. Gruber, S. Choi, D. Sun, P. Ren, L. S. Li, Y. Liu, F. Burrows, I. Maillard, T. Cierpicki and J. Grembecka, *J. Clin. Invest.*, 2020, **130**, 981–997.
- 302 A. V. Krivtsov, K. Evans, J. Y. Gadrey, B. K. Eschle, C. Hatton, H. J. Uckelmann, K. N. Ross, F. Perner, S. N. Olsen, T. Pritchard, L. McDermott, C. D. Jones, D. Jing, A. Braytee, D. Chacon, E. Earley, B. M. McKeever, D. Claremon, A. J. Gifford, H. J. Lee, B. A. Teicher, J. E. Pimanda, D. Beck, J. A. Perry, M. A. Smith, G. M. McGeehan, R. B. Lock and S. A. Armstrong, *Cancer Cell*, 2019, **36**, 660–673 e611.
- 303 S. He, T. J. Senter, J. Pollock, C. Han, S. K. Upadhyay, T. Purohit, R. D. Gogliotti, C. W. Lindsley, T. Cierpicki, S. R. Stauffer and J. Grembecka, *J. Med. Chem.*, 2014, **57**, 1543–1556.
- 304 S. Xu, A. Aguilar, T. Xu, K. Zheng, L. Huang, J. Stuckey, K. Chinnaswamy, D. Bernard, E. Fernandez-Salas, L. Liu, M. Wang, D. McEachern, S. Przybranowski, C. Foster and S. Wang, *Angew. Chem., Int. Ed.*, 2018, **57**, 1601–1605.
- 305 S. Xu, A. Aguilar, L. Huang, T. Xu, K. Zheng, D. McEachern, S. Przybranowski, C. Foster, K. Zawacki, Z. Liu, K. Chinnaswamy, J. Stuckey and S. Wang, *J. Med. Chem.*, 2020, **63**, 4997–5010.
- 306 K. Hayashi, K. Yoshida and Y. Matsui, *Nature*, 2005, **438**, 374–378.
- 307 H. Wu, N. Mathioudakis, B. Diagouraga, A. Dong, L. Dombrowski, F. Baudat, S. Cusack, B. de Massy and J. Kadlec, *Cell Rep.*, 2013, **5**, 13–20.
- 308 I. Fumasoni, N. Meani, D. Rambaldi, G. Scafetta, M. Alcalay and F. D. Ciccarelli, *BMC Evol. Biol.*, 2007, **7**, 187.
- 309 S. Myers, R. Bowden, A. Tumian, R. E. Bontrop, C. Freeman, T. S. MacFie, G. McVean and P. Donnelly, *Science*, 2010, **327**, 876–879.
- 310 A. Sorrentino, A. Federico, M. Rienzo, P. Gazzarro, M. Bifulco, A. Ciccodicola, A. Casamassimi and C. Abbondanza, *Int. J. Mol. Sci.*, 2018, **19**, 3250.
- 311 A. E. Zou, H. Zheng, M. A. Saad, M. Rahimy, J. Ku, S. Z. Kuo, T. K. Honda, J. Wang-Rodriguez, Y. Xuan, A. Korrapati, V. Yu, P. Singh, J. R. Grandis, C. C. King, S. M. Lippman, X. Q. Wang, A. Hinton and W. M. Ongkeko, *Oncotarget*, 2016, **7**, 51211–51222.
- 312 J. Hussin, D. Sinnett, F. Casals, Y. Idaghdour, V. Bruat, V. Saillour, J. Healy, J. C. Grenier, T. de Malliard, S. Busche, J. F. Spinella, M. Lariviere, G. Gibson, A. Andersson, L. Holmfeldt, J. Ma, L. Wei, J. Zhang, G. Andelfinger, J. R. Downing, C. G. Mullighan and P. Awadalla, *Genome Res.*, 2013, **23**, 419–430.
- 313 A. A. Houle, H. Gibling, F. C. Lamaze, H. A. Edgington, D. Soave, M. J. Fave, M. Agbessi, V. Bruat, L. D. Stein and P. Awadalla, *Genome Res.*, 2018, **28**, 1611–1620.
- 314 A. Allali-Hassani, M. M. Szewczyk, D. Ivanochko, S. L. Organ, J. Bok, J. S. Y. Ho, F. P. H. Gay, F. Li, L. Blazer, M. S. Eram, L. Halabelian, D. Dilworth, G. M. Luciani, E. Lima-Fernandes, Q. Wu, P. Loppnau, N. Palmer, S. Z. A. Talib, P. J. Brown, M. Schapira, P. Kaldis, R. C. O'Hagan, E. Guccione, D. Barsyte-Lovejoy, C. H. Arrowsmith, J. M. Sanders, S. D. Kattar, D. J. Bennett, B. Nicholson and M. Vedadi, *Nat. Commun.*, 2019, **10**, 5759.
- 315 N. Tripoulas, D. LaJeunesse, J. Gildea and A. Shearn, *Genetics*, 1996, **143**, 913–928.
- 316 H. Miyazaki, K. Higashimoto, Y. Yada, T. A. Endo, J. Sharif, T. Komori, M. Matsuda, Y. Koseki, M. Nakayama, H. Soejima, H. Handa, H. Koseki, S. Hirose and K. Nishioka, *PLoS Genet.*, 2013, **9**, e1003897.
- 317 S. An, K. J. Yeo, Y. H. Jeon and J. J. Song, *J. Biol. Chem.*, 2011, **286**, 8369–8374.
- 318 C. Huang, F. Yang, Z. Zhang, J. Zhang, G. Cai, L. Li, Y. Zheng, S. Chen, R. Xi and B. Zhu, *Nat. Commun.*, 2017, **8**, 1649.
- 319 W. Shen, P. Krautscheid, A. M. Rutz, P. Bayrak-Toydemir and S. L. Dugan, *Eur. J. Med. Genet.*, 2019, **62**, 55–60.
- 320 B. Skawran, D. Steinemann, A. Weigmann, P. Flemming, T. Becker, J. Flik, H. Kreipe, B. Schlegelberger and L. Wilkens, *Mod. Pathol.*, 2008, **21**, 505–516.
- 321 B. Xu, T. Qin, J. Yu, T. J. Giordano, M. A. Sartor and R. J. Koenig, *J. Biol. Chem.*, 2020, **295**, 8834–8845.
- 322 D. S. Rogawski, J. Deng, H. Li, H. Miao, D. Borkin, T. Purohit, J. Song, J. Chase, S. Li, J. Ndoj, S. Klossowski, E. Kim, F. Mao, B. Zhou, J. Ropa, M. Z. Krotoska, Z. Jin, P. Ernst, X. Feng, G. Huang, K. Nishioka, S. Kelly, M. He, B. Wen, D. Sun, A. Muntean, Y. Dou, I. Maillard, T. Cierpicki and J. Grembecka, *Nat. Commun.*, 2021, **12**, 2792.

