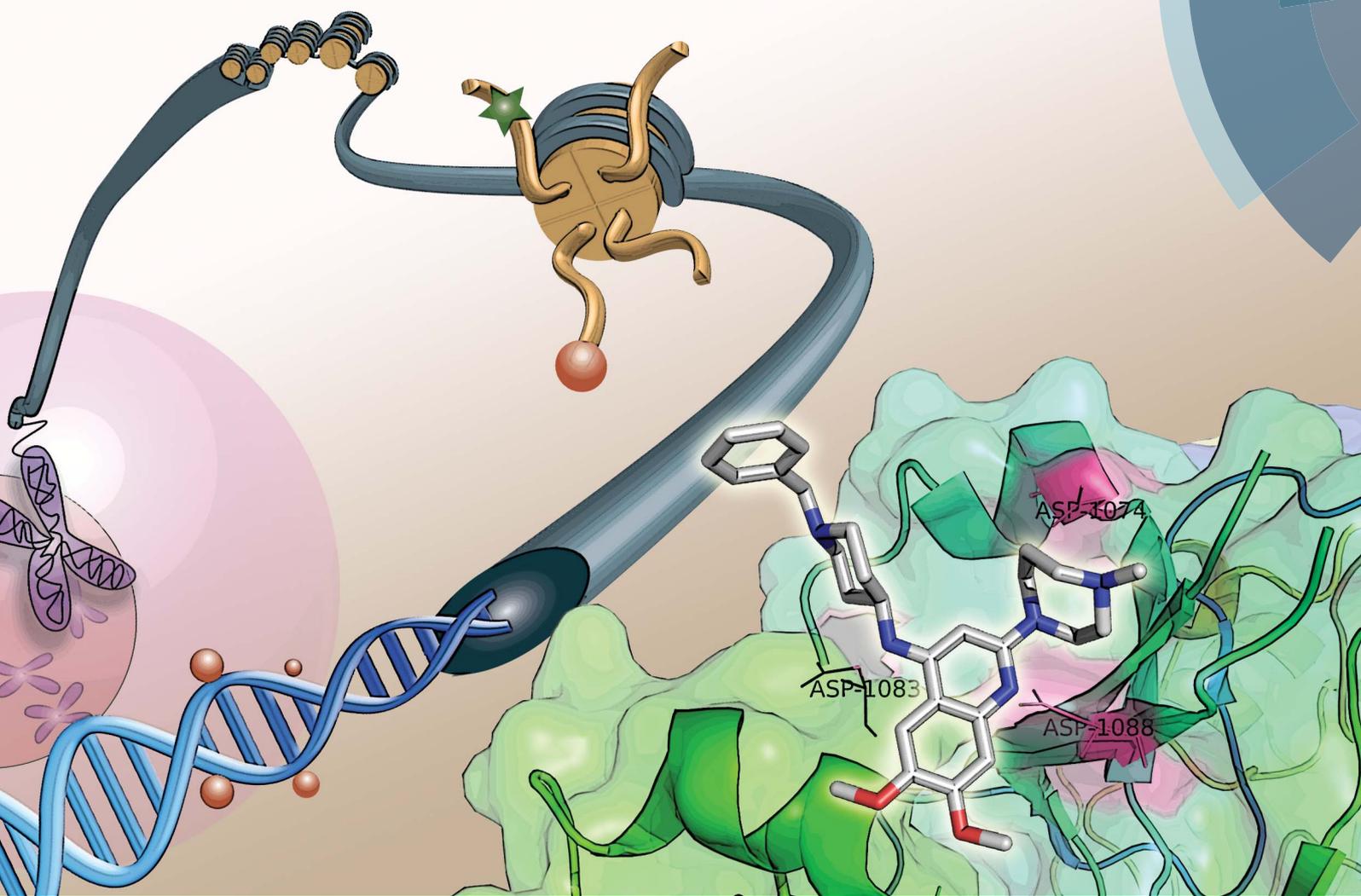


MedChemComm

Broadening the field of opportunity for medicinal chemists

www.rsc.org/medchemcomm



Themed issue: Epigenetics

ISSN 2040-2503



CONCISE ARTICLE
Matthew J. Fuchter *et al.*
Identification of 2,4-diamino-6,7-dimethoxyquinoline
derivatives as G9a inhibitors





Cite this: *Med. Chem. Commun.*, 2014, 5, 1821

Identification of 2,4-diamino-6,7-dimethoxyquinoline derivatives as G9a inhibitors†

Nitipol Srimongkolpithak,^{ab} Sandeep Sundriyal,^a Fengling Li,^c Masoud Vedadi^c and Matthew J. Fuchter^{*a}

G9a is a histone lysine methyltransferase (HKMT) involved in epigenetic regulation via the installation of histone methylation marks. 6,7-Dimethoxyquinazoline analogues, such as BIX-01294, are established as potent, substrate competitive inhibitors of G9a. With an objective to identify novel chemotypes for substrate competitive inhibitors of G9a, we have designed and synthesised a range of heterocyclic scaffolds, and investigated their ability to inhibit G9a. These studies have led to improved understanding of the key pharmacophoric features of BIX-01294 and the identification of a new core quinoline inhibitory scaffold, which retains excellent potency and high selectivity. Molecular docking was carried out to explain the observed *in vitro* data.

Received 24th June 2014
Accepted 25th August 2014

DOI: 10.1039/c4md00274a

www.rsc.org/medchemcomm

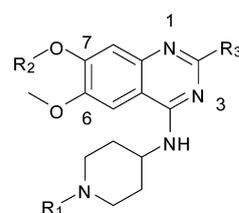
Introduction

G9a (also known as EHMT2) is a histone-lysine *N*-methyltransferase (HKMT), which catalyses the addition of one or two methyl groups to lysine 9 of histone H3 (H3K9me1 and H3K9me2) within a chromatin environment.¹ Due to its central role in epigenetic control, the methylation of H3K9 is associated with many biological pathways and is aberrantly regulated in several diseases including cancer² and AIDS.³ According to a genome-wide analysis of histone modifications,⁴ mono-methylation of histone H3 (H3K9me1) is associated with permissive chromatin, while di- and tri-methylation (H3K9me2/3) label a repressed chromatin state. Overexpression of G9a has been found in many types of cancer and is associated with a poor prognosis.^{5,6} Furthermore, G9a has been reported to methylate a variety of non-histone targets including the chromatin-modifying factor Tip49a⁷ and CDYL1.⁸

Like the majority of HKMTs, the SET (Su(var), E(z) and Trithorax) domain of G9a mediates catalysis and is comprised of two binding pockets; one for the cofactor *S*-adenosylmethionine (SAM), the other for the protein substrate. It is increasingly being realised that both pockets are 'druggable' and occupancy of either with a small molecule inhibitor is an effective strategy to block HKMT mediated methylation.⁹ The first substrate-competitive inhibitor of G9a, BIX-01294, was discovered using high throughput screening (Fig. 1).¹⁰ Focused medicinal

chemistry efforts from a number of groups has led to a series of optimised diaminoquinazoline analogues including UNC0224,¹¹ UNC0321,¹² UNC0631,¹³ UNC0638,¹⁴ UNC0646,¹³ UNC0965,¹⁵ E72,¹⁶ and UNC0642,¹⁷ which is suitable for *in vivo* work. Recently, 2-amino-indole derivatives have also been reported as substrate competitive G9a inhibitors.¹⁸ There is little doubt that the provision of such high quality inhibitors has dramatically facilitated the study of G9a biology,^{14,15,19,20} and related targets, especially in a disease context.^{21–24}

BIX-01294 and its optimised analogues are composed of a quinazoline heterocyclic core, substituted at positions 2, 4 and 7 (Fig. 1). The co-crystallized structure of UNC0224 with G9a (PDB



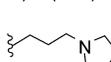
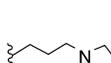
	R ₁	R ₂	R ₃
BIX-01294	-CH ₂ Ph	-CH ₃	
UNC0224	-CH ₃	-(CH ₂) ₃ N(CH ₃) ₂	
UNC0321	-CH ₃	-(CH ₂) ₂ O(CH ₂) ₂ N(CH ₃) ₂	
UNC0638	-CH(CH ₃) ₂		
UNC0642	-CH(CH ₃) ₂		

Fig. 1 Representative examples of substrate-competitive G9a inhibitors.

^aDepartment of Chemistry, Imperial College London, London SW7 2AZ, UK. E-mail: m.fuchter@imperial.ac.uk; Fax: +44 (0)2075945805; Tel: +44 (0)2075945815

^bInstitute of Chemical Biology, Imperial College London, London SW7 2AZ, UK

^cStructural Genomics Consortium, University of Toronto, Toronto, Ontario M5G 1L7, Canada

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c4md00274a



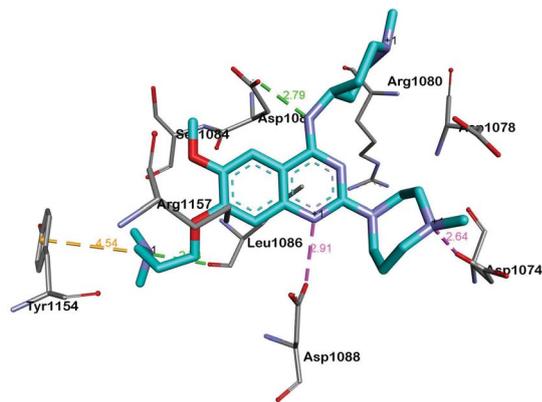


Fig. 2 A summary of the interactions between UNC0224 and G9a (PDB code 3K5K). Salt bridges, cation- π interactions and hydrogen bonds are depicted as pink, golden and green dashed lines, respectively.

code 3K5K)¹¹ reveals important interactions between the inhibitor and the substrate pocket of G9a. Key binding interactions include (Fig. 2): (1) a salt bridge between N-1 of the quinazoline core, which is expected to be protonated at physiological pH,^{25,26} and Asp1088; (2) a hydrogen bond between the C-4 NH functionality and Asp1083; (3) a hydrogen bond between the protonated 'lysine mimic' amine at C-7 with the backbone of Leu1086, as well as a cation- π interaction between the same protonated amine and Tyr1154.

Whilst the prior studies, particularly those of Jin and co-workers,^{11–17} have established important G9a structure-activity relationships (SAR) with respect to the side chains of the quinazoline core, it was apparent to us that the pharmacophoric features of the central heterocycle were yet to be determined. Thus we set out to better define the important features of the central inhibitor scaffold, while attempting to maintain the aforementioned interactions within the substrate pocket of the enzyme.

Chemistry

BIX-01294 (**1**) and other quinazoline derivatives (**2–4**, Table 1), were synthesised following the established two step synthesis.^{11,27}

Initially the importance of the dimethoxy-benzenoid ring of BIX-01294 was explored, while retaining the substituted pyrimidine ring. Hence, compounds **12** to **25** were synthesised (Table 1) containing a pyrimidine ring fused to a furan, thiophene, imidazole, or cyclopentane ring.

Amino furan **8a** was synthesised using a slight modification of a previously reported method (Scheme 1).²⁸ The carboxyl group of furan **5** was converted to a Boc protected amine **6** in a single step *via* a Curtius rearrangement. Treatment of amino furan **6** with *n*-butyllithium, followed by a methyl chloroformate quench, resulted in the isolation of product **7**. Boc deprotection of **7** yielded the desired intermediate **8a** with a free amino group at position 3. Amino furan **8a** and its commercially available analogues **8b** and **8c** were then converted to the corresponding

fused-pyrimidinediones **9a–9c** following reported reaction conditions.^{28,29} The desired dichloropyrimidines **10a–10c** were obtained by heating **9a–9c** to reflux in the presence of phosphoryl chloride, while the commercially available fused pyrimidinediones **9d** and **9e** were converted to **10d** and **10e** using pyrophosphoryl chloride³⁰ and phosphoryl chloride, respectively. Key intermediates **10a–10e** were then substituted with 4-amino-1-benzylpiperidine at C-4 to yield **11a–11e** which were further heated with a variety of secondary amines to obtain the fully substituted heterocyclic compounds **12–25** (Table 1).

As an alternative strategy to completely replacing the benzenoid ring of the original quinazoline analogues, a number of derivatives were prepared where the dimethoxy functionality had been altered or removed. Thus, the commercially available benzonitrile **26** was nitrated, followed by reduction of the nitro-group with sodium dithionite to give amine **27** (Scheme 2). Acylation of **27** using methyl chloroformate, followed by cyclisation under basic conditions, resulted in quinazoline-dione **28a**. The synthesised quinazoline-dione **28a** and commercially available desmethoxyquinazoline-dione **28b** were both chlorinated and substituted with amines as described previously^{11,27} to provide **31–37** (Table 1).

Finally, the pyrimidine ring portion of the quinazoline scaffold was varied. In light of the ligand-G9a interaction analysis above (Fig. 2), we decided to remove the N-3 nitrogen, since obvious interactions with G9a were not apparent. Thus, dimethoxyquinoline derivatives **41** and **42** were synthesised (Scheme 3). Unlike the synthesis of the related quinazoline derivatives, the regioselective synthesis of our target 2,4-diaminoquinolines was challenging. We found very few examples of such compounds in the literature, one example being from Campbell and co-workers, who have previously reported the synthesis of quinoline analogues of the antihypertensive diaminoquinazoline drug, prazosin.^{26,31} With a slight modification of their methodology, we were able to prepare free-amino quinoline derivatives **40a** and **40b**. Thus, the commercially available **38** was converted to an intermediate amino-benzonitrile and it was then treated with triethylorthoacetate under slightly reduced pressure to yield an imidate, which was heated with either methylpiperazine or methylhomopiperazine to yield corresponding derivatives **39a** and **39b**, respectively (Scheme 3). The ring closure of **39a** and **39b** was carried out by treatment with zinc chloride, which occurs presumably *via* the enamine tautomer.^{32,33} Finally, the free-amino quinoline analogues **40a** and **40b** were converted to target compounds **41** and **42** *via* reductive amination with 1-benzyl-4-piperidone.

Results and discussion

All the compounds synthesised were evaluated for G9a inhibitory activity using a previously reported SPA assay.¹⁷ The preliminary screen was conducted at three concentrations (50, 10, 1 μ M), with full dose response data derived for compounds with >20% inhibition at the 1 μ M dose. UNC0638 (Fig. 1) was used as a positive control and gave a comparable activity to that observed previously (IC₅₀ \sim 2.5 nM). Expectedly, BIX-01294 and the other quinazolines (**2–4**) exhibited IC₅₀ values in the



Table 1 SAR, biological and computational results of the BIX-01294 derivatives^a

Compound ID	Scaffold	R ₂	Activity% compound ^b (μM)			IC ₅₀ ^c (μM)	c log P ^d	N-1 pK _a ^e	Docking score (kcal mol ⁻¹)	
			50	10	1				SP	XP
1 (BIX-01294)			6	6	11	0.067 ± 0.003	3.9	8.13 (±2.22)	-7.859	na
2 (HKMTI-1-005)			5	6	19	0.101 ± 0.010	3.5	5.60 (±2.22)	-6.995	-6.380
3 (HKMTI-1-022)			7	9	35	0.472 ± 0.017	4.7	7.47 (±1.47)	-5.789	-4.244
4 (HKMTI-1-011)			15	31	79	3.190 ± 0.080	4.4	8.10 (±1.47)	na	na
12			99	102	99	—	2.9	4.57 (±0.70)	na	na
13			98	100	107	—	2.5	4.66 (±0.70)	na	na
14			97	96	101	—	3.7	5.31 (±0.70)	na	na
15			102	98	107	—	3.4	5.77 (±0.70)	na	na
16			47	82	91	—	3.9	5.68 (±0.70)	na	na
17			62	83	105	—	3.5	5.77 (±0.70)	na	na
18			91	102	114	—	4.8	6.42 (±0.70)	na	na
19			104	105	98	—	4.5	6.88 (±0.70)	na	na
20			81	96	91	—	3.9	3.28 (±2.22)	na	na
21			88	92	105	—	3.5	3.35 (±2.22)	na	na
22			93	105	105	—	4.8	3.86 (±2.22)	na	na
23			67	90	101	—	2.6	3.15 (±2.22)	na	na
24			101	90	97	—	3.4	3.73 (±2.22)	na	na
25			55	84	96	—	3.2	6.62 (±0.70)	na	na
31			63	91	98	—	3.2	5.58 (±2.22)	na	na
32			78	95	96	—	4.4	7.45 (±1.47)	na	na
33			109	98	96	—	4.2	8.08 (±1.47)	na	na
34			18	46	93	—	3.8	8.06 (±2.22)	na	na
35			27	59	94	—	3.5	5.53 (±2.22)	na	na
36			72	82	94	—	4.7	7.40 (±2.22)	na	na
37			97	108	104	—	4.4	8.03 (±2.22)	na	na



Table 1 (Contd.)

Compound ID	Scaffold	R ₂	Activity% compound ^b (μM)			IC ₅₀ ^c (μM)	c log P ^d	N-1 pK _a ^e	Docking score (kcal mol ⁻¹)	
			50	10	1				SP	XP
41 (HKMTI-1-248)			5	4	6	0.013 ± 0.001	4.5	10.57 (±2.22)	-7.522	-6.904
42 (HKMTI-1-247)			5	6	7	0.031 ± 0.003	4.1	9.84 (±2.22)	na	-6.476

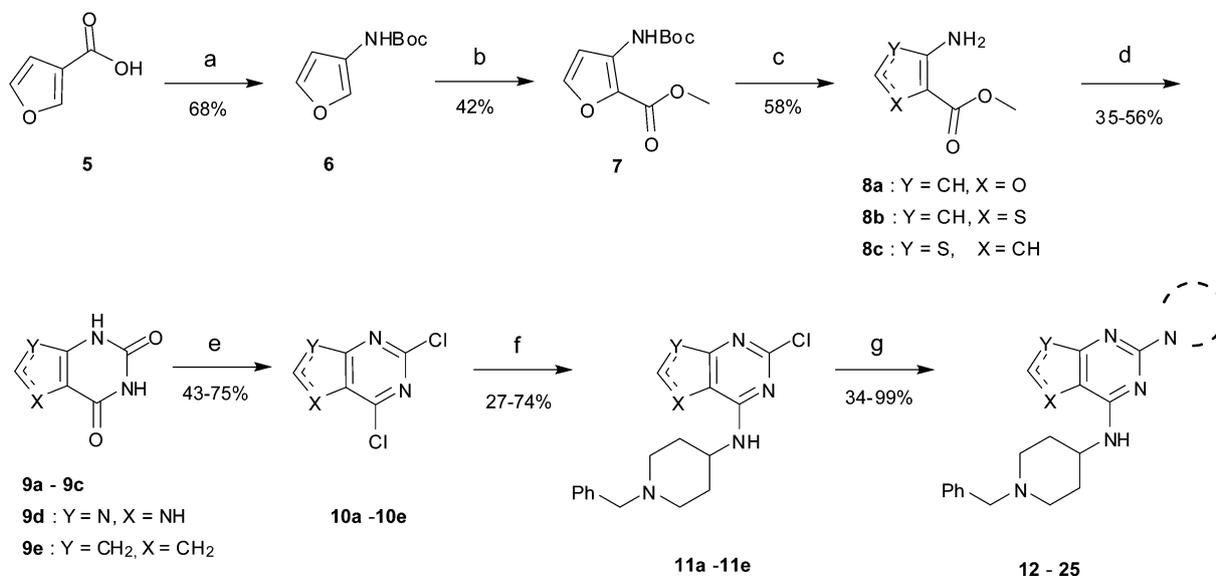
^a R₁ is the same for all compounds: R₁ = , SP = standard precision mode, XP = extra precision mode, na = no desired pose found.

^b The experiment was conducted in duplicate. ^c The assay was conducted in triplicate at K_m of both substrates (0.8 μM peptide [H3 1–25] and 8 μM SAM) for G9a (5 nM).¹⁷ ^d c log P values were calculated using the freely available program RDKit.³⁴ ^e 'Sequential' pK_a values were calculated at pH 7.0, with water as the solvent model, using the Epik 2.7 program implemented in Schrodinger (see ESI).

nanomolar to low micromolar range. For the derivatives where the benzenoid ring had been replaced with a furan, thiophene, imidazole or a cyclopentane ring (12–25), no significant G9a inhibitory activity was observed. While obvious interactions are not apparent for this region of the quinazoline G9a inhibitors (*vide supra*), except when a 'lysine mimic' is present at C-7, it would seem the benzenoid ring does play an important role in binding. Interestingly, the recently reported G9a inhibitor A-366 shares a similar structural feature with BIX-01294 in this regard.¹⁸

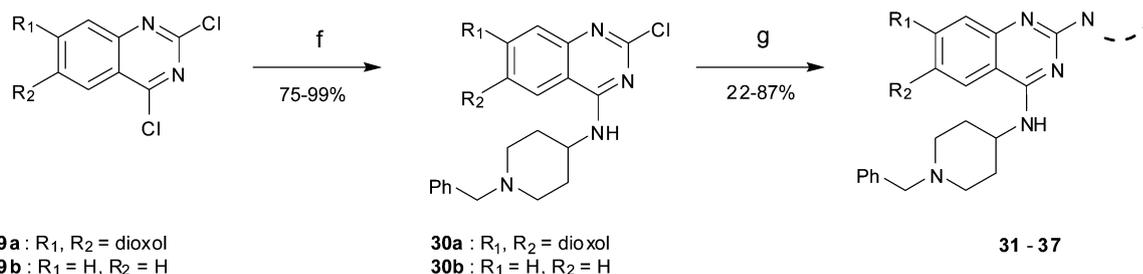
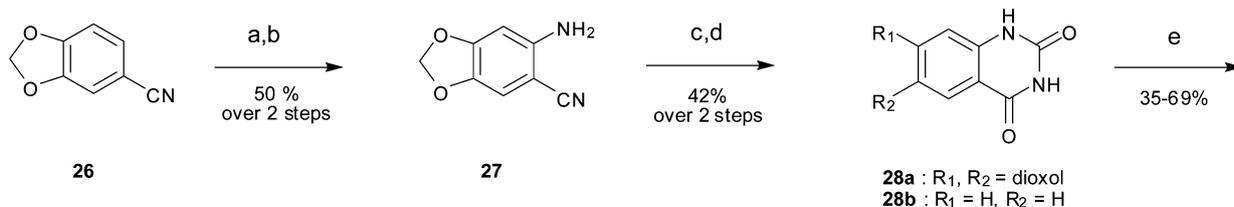
Since the benzenoid ring is important for inhibitory activity, the role of the dimethoxy substituents on this ring should be questioned. Desmethoxyquinazoline derivatives (34–37) were found to be far less active against G9a, which is in agreement

with a recent report on related analogues,^{35–42} and demonstrates that these substituents are also of importance. A number of theoretical and experimental studies have shown that *ortho*-dimethoxybenzenes prefer to adopt a co-planar conformation, with the sp² oxygen lone pairs projecting towards each other and the corresponding methyl groups angled away from each other. Indeed, such a conformation is apparent in the inhibitor-G9a/GLP crystal structures (such as PDB 3FPD,⁴³ 3K5K (Fig. 2),¹¹ and 3RJW¹⁴). Thus, analogues 31–33 (Table 1) were prepared where the methoxy groups were replaced by a dioxalane ring; forcing the oxygen lone pairs to point in the opposite direction. Interestingly, these analogues were also found to be largely devoid of G9a inhibitory activity, suggesting the conformational preference of the dimethoxy functionality to be important for G9a activity.



Scheme 1 The synthesis of fused-pyrimidine derivatives. Reagents and conditions: (a) DPPA, TEA, *t*-BuOH, reflux; (b) CH₃OCOCl, TMEDA, *n*-BuLi, THF; (c) TFA, DCM, rt, (d) ClSO₂NCO, DCM, aq. NaOH, aq. HCl or urea, 190 °C (neat), microwave; (e) POCl₃, DMF, reflux or O(POCl₂)₂, 145 °C, sealed tube, 2 day; (f) 4-amino-1-benzylpiperidine, DIEA, THF, rt, 24 h or 4-amino-1-benzylpiperidine, TEA, *n*-butanol, sealed tube, overnight; (g) cyclic amine, 185 °C (neat), microwave, 30 min.



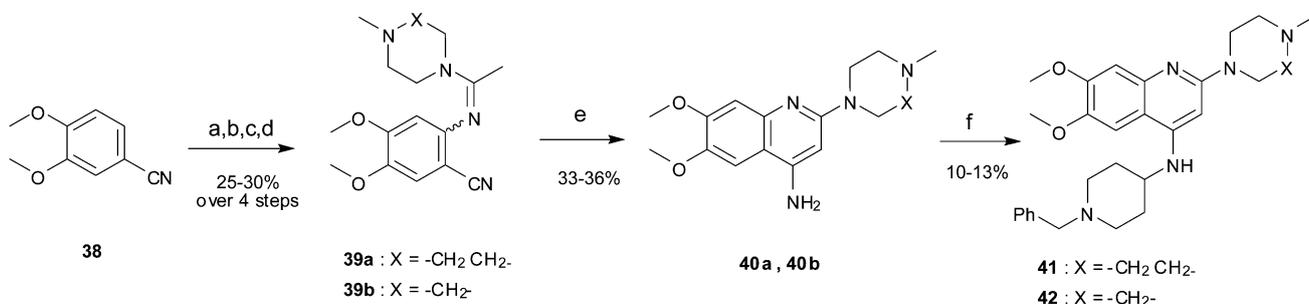


Scheme 2 The synthesis of desmethoxy and dioxoloquinazoline derivatives. Reagents and conditions: (a) HNO₃, Ac₂O, rt; (b) Na₂S₂O₄, TBAB, DCM/H₂O, rt; (c) ClCOOCH₃, DIEA, DMA/DCM; (d) H₂O, NaOH, EtOH, reflux; (e) POCl₃, PhNEt₂, MeCN, reflux; (f) 4-amino-1-benzylpiperidine, DIEA, THF, rt; (g) cyclic amine, 185 °C (neat), microwave, 30 min or amine, toluene, reflux, overnight.

In inhibitor designs where the dimethoxybenzenoid moiety was retained, but N-3 (Fig. 1) replaced by carbon atom (**41** and **42**), significant activity was observed. Indeed, to our delight, **41** and **42** were found to be ~5-fold more potent than the parental inhibitor BIX-01294. This result is clearly in agreement with our initial inhibitor-G9a crystal structure analysis (*vide supra*), which suggested that the N-3 nitrogen of the quinazoline scaffold might not be essential for G9a binding (Fig. 2). Comparison of the calculated pK_as revealed that N-1 of the quinolines **41** and **42** is more basic than the corresponding nitrogen of quinazolines **1** and **2** (Table 1). This computational prediction is in agreement with experimental data which demonstrates diaminoquinolines to be more basic than the analogous quinazolines.^{25,26} N-1 of the quinoline derivatives were predicted to have the highest pK_as (10.57 and 9.84 for **41** and **42**, respectively) of all the analogues prepared (Table 1). Interestingly, the N-1 atom in all inactive molecules (**12**–**25**, Table 1) were predicted to have lower pK_a values. A similar analysis between quinazoline and

quinoline activity based on pK_a dependent interactions has been described earlier for the design of α₁-adrenoreceptor antagonists.^{25,26}

In light of this trend, we measured the experimental pK_a values for compounds **1** (BIX-01294) and **41** in the pH range 2–12 using UV-vis spectroscopy. As expected, both compounds were found to be tribasic due to the presence of two aliphatic basic centres (the benzylpiperidine nitrogen and the homo piperazine nitrogen) in addition to N-1. The experimental pK_a values of BIX-01294 (**1**) were 6.94 ± 0.01, 8.24 ± 0.05 and 9.22 ± 0.04. The experimental pK_a values of quinoline **41** were 7.14 ± 0.01, 8.05 ± 0.02 and 9.50 ± 0.01. Since the largest spectral change for **1** and **41** would be expected upon ring protonation, the UV-vis spectra observed upon each protonation event can be used to assign the pK_a of N-1 (see Fig. S5 and S6, ESI⁺). We therefore assign the pK_a of N-1 of quinazoline **1** to be 6.94 ± 0.01 and N-1 of quinoline **41** to be 9.50 ± 0.01.



Scheme 3 The synthesis of quinoline derivatives. Reagents and conditions: (a) HNO₃, Ac₂O, rt; (b) Na₂S₂O₄, TBAB, DCM/H₂O, rt; (c) CH₃C(OCH₂CH₂)₃, 150 °C; (d) 1-methylhomopiperazine or 1-methylpiperazine, pTsoH, 100 °C; (e) ZnCl₂, DMA, reflux; (f) 1-benzyl-4-piperidone, AcOH, toluene, reflux, Dean–Stark, then a reducing agent (NaBH₄ or NaBH(OAc)₃), AcOH, THF, reflux.



This data is highly suggestive that a protonated N-1 position is required for good G9a binding, through a strong electrostatic interaction with Asp1088. Indeed, using the Henderson-Hasselbalch equation, under the given assay conditions (pH 8), 96.9% of **41** will be in the N-1 protonated form, whereas for BIX-01294 (**1**), 8.0% of the molecule will be N-1 protonated. We note however that there is an alternative explanation of the excellent potency of our quinoline hit compounds: the enthalpy loss due to breakage of a hydrogen bond between N-3 and water before ligand binding is not been compensated for upon ligand binding to G9a.⁴⁴ Despite this trend in N-1 pK_a, inactive quinazolines **31–37** are predicted to have N-1 pK_a values that are similar to active derivatives **1–4**. Therefore clearly both the dimethoxybenzenoid ring system and basic N-1 functionality are important features for G9a binding and hence inhibitory activity.

Our initial design of the new fused heterocyclic scaffolds was based on the hypothesis that they could bind to the substrate binding pocket of G9a with a pose comparable to that of the quinazoline co-crystallized ligand (Fig. 2). In light of the *in vitro* data obtained, all molecules prepared were docked without constraints into the substrate pocket of G9a to see if active and inactive molecules could be differentiated computationally. The G9a X-ray structure co-crystallised with UNC0224 (PDB code 3K5K)¹¹ was used, employing both standard precision (SP) and extra precision (XP) modes of the Glide program (Schrodinger, see ESI†).

Interestingly, analysis of the top scoring poses revealed that none of the inactive molecules could reproduce the expected pose in either SP or XP mode (Table 1). On the other hand all active derivatives were predicted to bind to G9a in a comparable manner to UNC0224, in at least one of the precision modes. For example, the poses of quinolines **41** (Fig. 3) and **42** (see ESI†)

overlaid perfectly with UNC0224, with the protonated N-1 functionality interacting with Asp1088. Similarly, other active compounds **1–3** exhibited a similar binding mode (see ESI, Fig. S1–S4†). Overall, the docking scores gave a qualitative correlation with the IC₅₀ data; compounds **1** and **41** giving higher scores than **2** and **3**.

Interestingly, the docking study reinforced the importance of dimethoxy structural feature in acquiring the correct pose. For example, quinazoline derivatives either lacking dimethoxy groups (**34–37**) or with the bridged methoxy groups (**31–33**) did not display the desired pose which is in agreement to their lack of *in vitro* activity. Also, moderately active derivative **4** could not reproduce the expected pose in spite of possessing dimethoxy groups, plausibly due to the large pyridylpiperazine substituent at position 2.

In light of the excellent potency of our quinoline inhibitors **41** and **42** against G9a, the selectivity of these compounds was examined in a methyltransferase enzyme panel, and compared to BIX-01294. This panel consisted of twenty three additional methyltransferases including sixteen HKMTs, six protein arginine methyltransferases (PRMTs), and one DNA methyltransferase (DNMT) (see ESI, Table S1†).¹⁷ Compound **41** and **42** were found to be equipotent against G9a and GLP (G9a-like protein, EHMT1), which is not surprising given the high degree of homology between the G9a and GLP SET domains. Such dual G9a/GLP activity is a common feature of the diaminoquinazoline inhibitors¹⁷ and was shared by BIX-01294. However to our delight, quinoline inhibitors **41** and **42** were found to be inactive against all other methyltransferases in the panel, with the exception of SETD2 and EZH2 for which a moderate inhibitory activity was at the highest concentrations surveyed (50 μM).

Conclusions

In summary, we have designed and synthesised a variety of heterocyclic derivatives in order to identify novel G9a inhibitors and better define the pharmacophoric features associated with the core heterocycle. These efforts resulted in the identification of potent and selective G9a/GLP inhibitors **41** and **42** based on a quinoline scaffold. Activity and computational data highlight the importance of the dimethoxy groups on the benzenoid ring of this scaffold and a basic nitrogen at position 1 for potent G9a activity. The ability of molecular docking to predict inhibitor activity based on the UNC0224 binding pose, once further validated, may be helpful in the future for predicting and prioritising novel G9a/GLP inhibitors.

Acknowledgements

N.S. was supported by a Royal Thai Government Scholarship and the EPSRC-funded Institute of Chemical Biology Doctoral Training Centre. S.S. acknowledges European Commission for awarding Marie Curie International Incoming Fellowship (Agreement no. 299857). The SGC is a registered charity (number 1097737) that receives funds from AbbVie, Boehringer Ingelheim, the Canada Foundation for Innovation, the Canadian Institutes for Health Research, Genome Canada through

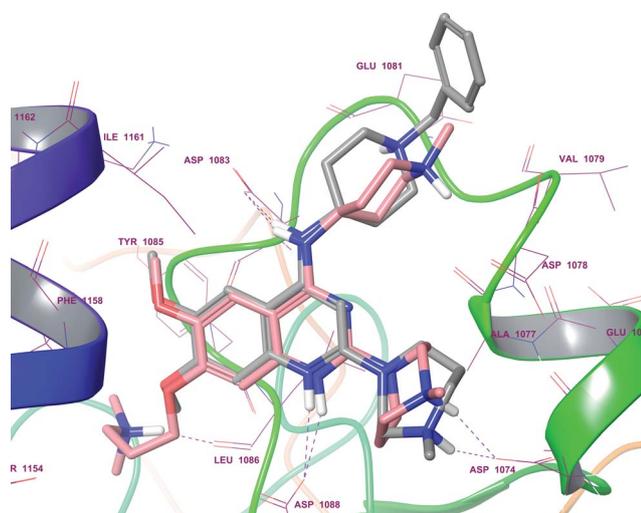


Fig. 3 Docking pose of quinoline analogue **41** (grey sticks) overlaid with the co-crystallized quinazoline derivative, UNC0224 (pink sticks; PDB 3K5K), in the G9a substrate binding pocket. Purple dashed lines display H-bonds. As expected, protonated the N-1 and 4-amino moiety of **41** were shown to interact with Asp1088 and Asp1083, respectively.



the Ontario Genomics Institute [OGI-055], GlaxoSmithKline, Janssen, Lilly Canada, the Novartis Research Foundation, the Ontario Ministry of Economic Development and Innovation, Pfizer, Takeda, and the Wellcome Trust [092809/Z/10/Z].

Notes and references

- M. Tachibana, K. Sugimoto, T. Fukushima and Y. Shinkai, *J. Biol. Chem.*, 2001, **276**, 25309–25317.
- H. Tao, H. Li, Y. Su, D. Feng, X. Wang, C. Zhang, H. Ma and Q. Hu, *Mol. Cell. Biochem.*, 2014, **394**, 23–30.
- S. Bouchat, J. S. Gatot, K. Kabeya, C. Cardona, L. Colin, G. Herbein, S. De Wit, N. Clumeck, O. Lambotte, C. Rouzioux, O. Rohr and C. Van Lint, *AIDS*, 2012, **26**, 1473–1482.
- A. Barski, S. Cuddapah, K. Cui, T. Y. Roh, D. E. Schones, Z. Wang, G. Wei, I. Chepelev and K. Zhao, *Cell*, 2007, **129**, 823–837.
- J. Ding, T. Li, X. Wang, E. Zhao, J. H. Choi, L. Yang, Y. Zha, Z. Dong, S. Huang, J. M. Asara, H. Cui and H. F. Ding, *Cell Metabolism*, 2013, **18**, 896–907.
- X. Zhong, X. Chen, X. Guan, H. Zhang, Y. Ma, S. Zhang, E. Wang, L. Zhang and Y. Han, *Histopathology*, 2014, DOI: 10.1111/his.12456.
- J. S. Lee, Y. Kim, I. S. Kim, B. Kim, H. J. Choi, J. M. Lee, H. J. Shin, J. H. Kim, J. Y. Kim, S. B. Seo, H. Lee, O. Binda, O. Gozani, G. L. Semenza, M. Kim, K. I. Kim, D. Hwang and S. H. Baek, *Mol. Cell*, 2010, **39**, 71–85.
- P. Rathert, A. Dhayalan, M. Murakami, X. Zhang, R. Tamas, R. Jurkowska, Y. Komatsu, Y. Shinkai, X. Cheng and A. Jeltsch, *Nat. Chem. Biol.*, 2008, **4**, 344–346.
- K. T. Nguyen, F. Li, G. Poda, D. Smil, M. Vedadi and M. Schapira, *J. Chem. Inf. Model.*, 2013, **53**, 681–691.
- 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.
- 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.
- 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.
- 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.
- 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.
- K. D. Konze, S. G. Pattenden, F. Liu, D. Barsyte-Lovejoy, F. Li, J. M. Simon, I. J. Davis, M. Vedadi and J. Jin, *ChemMedChem*, 2014, **9**, 549–553.
- 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.
- 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.
- R. F. Sweis, M. Pliushchev, P. J. Brown, J. Guo, F. L. 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.
- Y. Shi, C. Despons, J. T. Do, H. S. Hahm, H. R. Scholer and S. Ding, *Cell Stem Cell*, 2008, **3**, 568–574.
- K. Imai, H. Togami and T. Okamoto, *J. Biol. Chem.*, 2010, **285**, 16538–16545.
- L. Dembele, J. F. Franetich, A. Lorthiois, A. Gego, A. M. Zeeman, C. H. Kocken, R. Le Grand, N. Dereuddre-Bosquet, G. J. van Gemert, R. Sauerwein, J. C. Vaillant, L. Hannoun, M. J. Fuchter, T. T. Diagana, N. A. Malmquist, A. Scherf, G. Snounou and D. Mazier, *Nat. Med.*, 2014, **20**, 307–312.
- A. K. Upadhyay, D. Rotili, J. W. Han, R. Hu, Y. Chang, D. Labella, X. Zhang, Y. S. Yoon, A. Mai and X. Cheng, *J. Mol. Biol.*, 2012, **416**, 319–327.
- N. A. Malmquist, T. A. Moss, S. Mecheri, A. Scherf and M. J. Fuchter, *Proc. Natl. Acad. Sci. U. S. A.*, 2012, **109**, 16708–16713.
- S. Sundriyal, N. A. Malmquist, J. Caron, S. Blundell, F. Liu, X. Chen, N. Srimongkolpithak, J. Jin, S. Charman, A. Scherf and M. Fuchter, *ChemMedChem*, 2014, DOI: 10.1002/cmde.201402098.
- J. Bordner, S. F. Campbell, M. J. Palmer and M. S. Tute, *J. Med. Chem.*, 1988, **31**, 1036–1039.
- S. F. Campbell, J. D. Hardstone and M. J. Palmer, *J. Med. Chem.*, 1988, **31**, 1031–1035.
- R. Brown, M. Fuchter, N. Chapman-Rothe, N. Srimongkolpithak, J. Caron, J. Synder, T. Ganesh, J. Liu and A. Sun, *EU Pat.*, WO/2013/140148, 2013.
- T. P. Heffron, B. Wei, A. Olivero, S. T. Staben, V. Tsui, S. Do, J. Dotson, A. J. Folkes, P. Goldsmith, R. Goldsmith, J. Gunzner, J. Lesnick, C. Lewis, S. Mathieu, J. Nonomiya, S. Shuttleworth, D. P. Sutherlin, N. C. Wan, S. Wang, C. Wiesmann and B. Y. Zhu, *J. Med. Chem.*, 2011, **54**, 7815–7833.
- W. Zhu, Y. Liu, X. Zhai, X. Wang, Y. Zhu, D. Wu, H. Zhou, P. Gong and Y. Zhao, *Eur. J. Med. Chem.*, 2012, **57**, 162–175.
- M. Chiriac, D. Axente, N. Palibroda and C. T. Craescu, *J. Labelled Compd. Radiopharm.*, 1999, **42**, 377–385.
- S. F. Campbell, J. D. Hardstone and M. J. Palmer, *Tetrahedron Lett.*, 1984, **25**, 4813–4816.



- 32 H. Schafer, K. Sattler and K. Gewald, *J. Prakt. Chem.*, 1979, **321**, 695–698.
- 33 J. A. Moore and L. D. Kornreich, *Tetrahedron Lett.*, 1963, **4**, 1277–1281.
- 34 <http://www.rdkit.org/>, accessed December, 2013.
- 35 D. Rotili, D. Tarantino, B. Marrocco, C. Gros, V. Masson, V. Poughon, F. Ausseil, Y. Chang, D. Labella, S. Cosconati, S. Di Maro, E. Novellino, M. Schnekenburger, C. Grandjenette, C. Bouvy, M. Diederich, X. Cheng, P. B. Arimondo and A. Mai, *PLoS One*, 2014, **9**, e96941.
- 36 G. M. Anderson, P. A. Kollman, L. N. Domelsmith and K. N. Houk, *J. Am. Chem. Soc.*, 1979, **101**, 2344–2352.
- 37 J. Caillet and J. Caillet, *Acta Crystallogr., Sect. B: Struct. Crystallogr. Cryst. Chem.*, 1982, **38**, 1786–1791.
- 38 T. Schaefer, R. Sebastian, A. Lemire and G. H. Penner, *Can. J. Chem.*, 1990, **68**, 1393–1398.
- 39 M. Gerzain, G. W. Buchanan, A. B. Driega, G. A. Facey, G. Enright and R. A. Kirby, *J. Chem. Soc., Perkin Trans. 2*, 1996, 2687–2693.
- 40 J. W. Emsley, E. K. Foord and J. C. Lindon, *J. Chem. Soc., Perkin Trans. 2*, 1998, 1211–1218.
- 41 S. Tsuzuki, H. Houjou, Y. Nagawa and K. Hiratani, *J. Chem. Soc., Perkin Trans. 2*, 2002, 1271–1273.
- 42 C. V. Velde, E. Bultinck, K. Tersago, C. V. Alsenoy and F. Blockhuys, *Int. J. Quantum Chem.*, 2007, **107**, 670–679.
- 43 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.
- 44 H. Zhao and D. Huang, *PLoS One*, 2011, **6**, e19923.

