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inhibitor of BRD4 as a potent anticancer agent

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protein–protein interaction *in vitro*, and exhibited potent
anticancer activity in an *in vivo* model of melanoma.

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An iridium(III)-based irreversible protein–protein interaction inhibitor of BRD4 as a potent anticancer agent†

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Bromodomain-containing protein 4 (BRD4) has recently emerged as an attractive epigenetic target for anticancer therapy. In this study, an iridium(III) complex is reported as the first metal-based, irreversible inhibitor of BRD4. Complex **1a** is able to antagonize the BRD4-acetylated histone protein–protein interaction (PPI) *in vitro*, and to bind BRD4 and down-regulate *c-myc* oncogenic expression *in cellulo*. Chromatin immunoprecipitation (ChIP) analysis revealed that **1a** could modulate the interaction between BRD4 and chromatin in melanoma cells, particular at the MYC promoter. Finally, the complex showed potent activity against melanoma xenografts in an *in vivo* mouse model. To our knowledge, this is the first report of a Group 9 metal complex inhibiting the PPI of a member of the bromodomain and extraterminal domain (BET) family. We envision that complex **1a** may serve as a useful scaffold for the development of more potent epigenetic agents against cancers such as melanoma.

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Introduction

Gene transcription is a dynamic process tightly regulated by chromatin, which is a complex structure comprised of DNA and histone proteins.¹ The function of gene regulation is controlled by post-translational modification states of DNA-packing histones in the chromatin complex.² For example, the N-terminal lysine residues of histone proteins can be acetylated and deacetylated to control gene expression *via* the interplay of

a range of enzymes such as histone acetyltransferase (HAT), histone deacetylase (HDAC) and methyltransferase (MT).¹ Hence, these enzymes have become the targets of drug discovery efforts.^{3,4} However, the reader domains that interrogate post-translational modification states have been less intensively pursued as epigenetic targets.^{5,6}

Acetylated histones are recognized by small protein pockets called bromodomains.⁷ The bromodomain and extraterminal domain (BET) family of bromodomain-containing proteins (BRD2, BRD3, BRD4 and BRDT) are a class of transcriptional regulators containing tandem bromodomains and a carboxyl-terminal recruitment domain.^{8,9} In particular, BRD4 plays a significant role in cell cycle progression and viability *via* its effects on growth-related genes at the M/G1 boundary.^{10,11} Recently, BRD4 has been shown to play an important role in sustaining the proliferation of metastatic melanoma, a mostly incurable disease, thus rendering it as a possible target for epigenetic therapy.¹²

The selective inhibition of the bromodomain 4 (BRD4)/histone interaction has been demonstrated by several small molecule inhibitors such as (+)-JQ1, which is capable of occupying the ϵ -N-acetylated lysine residue (Kac) binding site of BRD4 and act as a Kac-competitive inhibitor.¹³ Subsequent reports have shown that (+)-JQ1 can directly regulate transcription mediated by the *c-myc* gene and reduce the expression of oncogenic *c-myc* protein.^{14,15}

The success of the anti-cancer compound cisplatin and its analogues has inspired the investigation of metal-based

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compounds as therapeutic agents over the past few decades.^{16–24} While classical metal-based chemotherapeutic agents typically target double-helical DNA, increasing knowledge in molecular biology has uncovered the possibility of specifically targeting therapeutically relevant proteins or enzymes using transition-metal complexes.^{25–32} Metal-based compounds can offer distinct opportunities in targeting proteins or enzymes compared to organic small molecules due to their interesting structural diversity and electronic properties. Moreover, metal complexes can undergo ligand exchange reactions with biomolecules, and such irreversible inhibitors may show enhanced potency and potentially allow for less frequent and lower dosages *in vivo*.³³ Examples of approved drugs that act *via* a covalent mechanism include EGFR inhibitors Neratinib (Pfizer), Afatinib/BIBW-2992 (Boehringer Ingelheim) and PF-00299804 (Pfizer), and anti-HCV agents Telaprevir (Vertex Pharmaceuticals and Johnson & Johnson) and Boceprevir/Victrelis (Merck) (Fig. S1†).³³ Neratinib, Afatinib/BIBW-2992 and PF-00299804 target cysteine in EGFR, and Carfilzomib/Kyprolis, a selective proteasome inhibitor, targets threonine, while Telaprevir, used for the treatment of HCV, targets serine. Boceprevir/Victrelis also targets serine of HCV protease, and is used for the treatment of hepatitis caused by HCV.

Metal complexes can adopt a wide range of geometrical shapes defined by the oxidation state of the metal center and the nature of the co-ligands, while organic compounds are mainly restricted to linear, trigonal-planar and tetrahedral geometries. Therefore, metal complexes may be able to sample additional chemical space within the active site of enzymes or proteins. In addition, the steric and electronic properties of metal complexes can be easily tuned without lengthy synthetic protocols due to the modular nature of inorganic synthesis. We and others have previously demonstrated that certain Ir(III),^{34–36} Rh(III)^{37,38} and Ru(II)^{39–42} complexes can be developed as inhibitors of enzymes or protein–protein interactions (PPI). Continuing in our quest to explore the therapeutic applications of Group 9 complexes, we report herein the first metal-based epigenetic inhibitor of BRD4 and of any BET protein in general.

Results and discussion

Screening of Group 9 metal complexes as BRD4 inhibitors

The Ir(III)/Rh(III) metal complexes 1–27 were synthesized as racemates according to literature methods (Fig. 1).^{25,43} As lysine-acetylated histone peptides are known substrates for bromodomain-containing proteins, we initially screened the complexes at a concentration of 100 μ M for their ability to modulate the protein–protein interaction between tetra-acetylated lysine histone 4 peptide (H4AcK4) and BRD4 using a time-resolved-fluorescent resonance energy transfer (TR-FRET) assay (Cayman Chemical, Ann Arbor, MI, USA). Among the 27 compounds tested, complex 1 emerged as a top candidate, with slightly lower activity compared to the positive control compound (+)-JQ-1 (Fig. S1 and S2†). We analyzed the structure of complexes 1–27 to identify favorable substructures for the



Fig. 1 Chemical structures of the cyclometallated Ir(III) and Rh(III) complexes (racemates) used for preliminary screening. Only one enantiomer is shown for clarity; complexes 1 and 17 are OTf salts, and others are PF₆ salts.

development of the next round of complexes. We observed that complex 1 possessed two 2-phenylpyridine C^N ligands and two acetonitrile ligands, whereas the other complexes tested generally possessed bidentate N^N donor ligands. Complex 5, which possessed the dipyrido[3,2-*a*:2',3'-*c*]phenazine (dppz) N^N ligand, showed the weakest potency out of the complexes tested, suggesting that this ligand detracted from biological activity (Fig. 2).

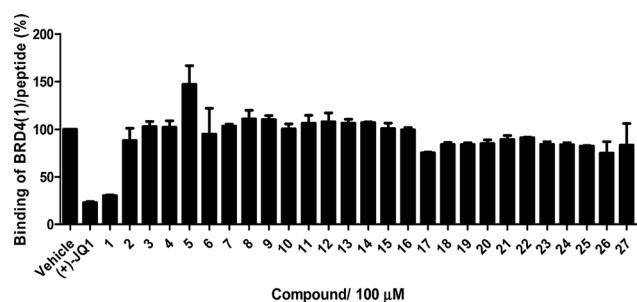


Fig. 2 Displacement of a tetra-acetylated H4 peptide from BRD4 by a selection of Ir(III)/Rh(III) complexes at 100 μ M in a TR-FRET assay. Error bars represent the standard deviations of the results from three independent experiments.



Synthesis and structure-based optimization of novel BRD4 inhibitors

Based on the structure of **1**, a focused library of 10 cyclometalated rhodium(III) and iridium(III) complexes containing different C[^]N or N-donor ligands (**1a–1j**) (Fig. 3a) was designed and synthesized. The spectroscopic data of the complexes are presented in Table S1.† This library was enriched in acetonitrile or acetonitrile-based N-donor ligands that were identified in the first round of screening to be favorable substructures for biological potency. In the second round of screening, the iridium(III) complex **1a** showed the greatest inhibition of BRD4-H4AcK4 binding and was comparably potent to (+)-JQ1 (Fig. 3b). Notably, complex **1a**, which contains two 2-phenyl-6-methylpyridine C[^]N ligands and two acetonitrile ligands, was more potent than the parent complex **1**. Based on these results, preliminary structure-activity relationships (SAR) could be deduced. Varying the N-donor ligands from acetonitrile to other nitrile-based ligands (such as in **1i**) did not result in improved activity against BRD4, as complex **1i** was one of the least active compounds in the TR-FRET assay. Additionally, substituting the 2-phenyl-6-methylpyridine (as in **1a**) or 2-phenylpyridine (as in **1**) C[^]N co-ligands with more extended aromatic systems such as 2-phenylquinoline (as in **1d**) or 1-phenylisoquinoline (as in **1f**) also decreased the potency of the complex. Moreover, the presence of fluorine substituents on the C[^]N ligands appeared to be highly detrimental for activity, as complex **1b** showed the weakest activity out of this series. Finally, replacing the iridium(III) center of complex **1** with rhodium(III) (as in congener **1h**) resulted in greatly decreased activity against BRD4. Taken together, these results suggest that the binding between **1a** and BRD4 is highly sensitive to the steric and/or electronic properties of the metal complexes.

As complex **1** has been previously reported to bind covalently to histidine and generate a luminescence signal,²⁵ we were interested to investigate whether complex **1a** would also show a luminescent response to histidine or various other natural amino acids. In the presence of histidine, **1a** exhibits an intense emission enhancement at $I_{\text{max}} = 497$ nm, whereas no significant changes in its emission were observed upon the addition of other natural amino acids (Fig. S2†). Competition experiments were carried out by incubating **1a** with histidine and 10 equivalents of another natural amino acid. No significant difference was observed between the luminescence intensity detected in the competition experiments compared to that in the presence of histidine alone (data not shown). We further analyzed the binding of **1a** to histidine by electrospray ionization mass spectrometry. After incubation of **1a** (m/z 529.1) with histidine for 30 min at 20 °C, a new peak at m/z 684.2 was observed (Fig. S3b†). This peak corresponds to the covalent attachment of one histidine molecule (m/z 155) to **1a**. No mass change was recorded for **1a** upon incubation with other natural amino acids under same reaction conditions (Fig. S3c†). This data demonstrates that **1a** selectively and covalently binds to histidine, leading to a luminescence response.

To assess the selectivity of complex **1a**, we tested its activity against two unrelated proteins, caspase-6 and STAT3, which



Fig. 3 (a) Chemical structures of the cyclometalated Ir(III) and Rh(III) complexes (racemates) used for structure-activity analysis (SAR). (b) Displacement of a tetra-acetylated H4 peptide from BRD4 by complex **1** and analogues **1a–1j** at 10 μM in a TR-FRET assay. Error bars represent the standard deviations of the results from three independent experiments.

contain 12 and 13 histidine residues, respectively. The results showed that **1a** exhibited no significant effect on caspase-6 activity, and only slightly inhibited STAT3 DNA-binding activity *in vitro* (Fig. S4†). This data therefore demonstrates that **1a** does not bind equally well to all histidine-containing proteins, and suggests that there exists further criteria that determine the activity of **1a** against BRD4.

BRD4 inhibitor antagonizes the BRD4-acetylated histone PPI *in vitro*

To further verify the BRD4 inhibitory activity of iridium(III) complex **1a**, the complex was subjected to a dose-response experiment in the TR-FRET assay. The results showed that **1a** inhibited the peptide-binding activity of BRD4 with an IC_{50} value (dose required to inhibit 50% TR-FRET ratio) of 0.07 μM (Fig. 4a). The inhibitory activity of complex **1a** against the interaction between BRD4 and H4AcK4 was further confirmed using an AlphaScreen assay (Fig. S5†).

BRD4 contains two conserved N-terminal bromodomains BRD4(1) and BRD4(2). The activity of complex **1a** against the binding of BRD4(2) to H4AcK4 was also investigated using the TR-FRET assay (Fig. 4b). The results revealed that **1a** exhibited





Fig. 4 Ability of **1a** to displace H4AcK4 peptide from (a) BRD4(1) and (b) BRD4(2) in a time-resolved-fluorescent resonance energy transfer (TR-FRET) assay. Binding of H4AcK4 to BRD4(1) was strongly inhibited by **1a**, with half-maximum inhibitory concentration (IC_{50}) value of $0.07 \mu\text{M}$. Error bars represent the standard deviations of the results from three independent experiments. LC-MS/MS analysis of (c) BRD4(1), (d) BRD4(1) with **1a** and (e) **1a** only. BRD4(1) and **1a** complex were buffered in 10 mM Tris-HCl, pH = 7.5, 500 mM NaCl and incubated at 25°C for 2 h. The sample was analyzed by positive ion mass spectra.

no significant inhibition of the interaction between BRD4(2) and H4AcK4. Therefore, complex **1a** selectively inhibits the BRD4(1) domain. This result was further corroborated by a fluorescence polarization assay (Fig. S6[†]).

The interaction between complex **1a** with the BRD4 proteins was also monitored by luminescence spectroscopy, since complex **1a** exhibits a high luminescence response in the presence of BRD4(1) and BRD4(2) (Fig. S7[†]). A time-course experiment revealed that the luminescence signal of **1a** reached steady-state within 5 and 8 min upon the addition of BRD4(1) or BRD4(2) at 25°C , respectively (Fig. S8[†]). These data suggest that complex **1a** may react slightly faster with the BRD4(1) protein.

LC-MS/MS further demonstrated the binding of **1a** to BRD4(1). Mass spectrometer data was pooled and analyzed for the BRD4(1)-**1a** complex of 17 036.5 Da or intact BRD4(1) corresponding to 16 472.3 Da (Fig. 4c and d). As shown in Fig. 4e, MS/MS fragmentation of the singly-charged ion (m/z 529.1) matched the molecular weight of **1a** with cleavage of two ACN ligands. After 2 h of incubation with **1a**, a complex was observed corresponding to BRD4(1) binding to $\text{C}_{24}\text{H}_{20}\text{N}_2\text{Ir}$ (529.1) with an additional buffer adduct NH_4^+OH^- (35.0). The MS data therefore suggests that **1a** loses two ACN ligands upon binding to BRD4(1).

To further investigate the mechanism of action of **1a**, we incubated the complex in DMSO solution. The results showed that **1a** exchanges its acetonitrile ligands for DMSO ligands from the solution (Fig. S9[†]). This is similar to previous complexes,⁴⁴ as well as NAMI-A and KP1019. Moreover, since DMSO ligands are also labile, this should not affect the ability of **1a** to bind covalently to the protein target, as is the case for the previously described complexes. This makes the mechanism of **1a** likely to be similar to that of NAMI-A/KP1019,^{45–49} which also interact covalently with their biomolecular targets *via* ligand exchange.^{50,51} Furthermore, after ligand exchange with DMSO, the complex was stable for at least 24 h in DMSO solution and in plasma under our test conditions, as revealed by the absence of significant changes in the absorbance.

BRD4 inhibition suppresses MYC and Bcl-2 *in cellulo* and represses cancer cell growth

Given by the promising activity of complex **1a** at antagonizing the BRD4-H4AcK4 interaction *in vitro*, the complex was further examined for its biological activity in cells. We first performed a chromatin immunoprecipitation (ChIP) assay to investigate whether **1a** can modulate the binding of BRD4 to chromatin in human malignant melanoma A375 and human caucasian metastatic melanoma A2058 cell lines (Fig. 5). ChIP analysis at the MYC promoter showed that 10 nM of complex **1a** decreased the recruitment of BRD4 after 6 h. A similar pattern was observed at Bcl-2 and CDK6 loci, but not at the housekeeping B2M gene. These results suggest that **1a** is able to modulate the interaction between chromatin and BRD4 in A375 and A2058 cells, particularly at the MYC promoter.

Furthermore, the impact of complex **1a** on *c-myc* and Bcl-2 expression in A375 and A2058 cells was investigated. Immunoblotting analysis of lysates from treated cells revealed that the expression of *c-myc* and Bcl-2 proteins was reduced by **1a** in a dose-dependent manner (Fig. 6 and Fig. S10[†]). These observations are also consistent with the result of the ChIP assay described above, which showed that complex **1** could disrupt the binding of BRD4 to the MYC and Bcl-2 promoters. These results therefore suggest that **1a** may act as a transcriptional modulator of *c-myc* and Bcl-2 expression.

c-myc and Bcl-2 have been intensely studied as anti-cancer targets due to their roles in cell cycle progression, cellular transformation and apoptosis.^{52,53} Therefore, we were interested to investigate whether or not complex **1a** could exhibit anti-proliferative effects against cancer cells. In *in vitro* evaluation, complex **1a** exhibited potent cytotoxicity against the A375 ($IC_{50} = 12.5 \mu\text{M}$) and A2058 ($IC_{50} = 3 \mu\text{M}$) cell lines (Fig. 6c and d). The anti-proliferative activity of complex **1a** towards A375 and A2058 cells was further determined using the colony formation assay. The results showed that **1a** was cytotoxic against A375 and A2058 melanoma cells with estimated IC_{50} values (dose required to inhibit 50% cellular growth after 24 h exposure to **1a**) of 5 nM and 1 nM, respectively (Fig. 6e and f). We reason that the cytotoxicity imparted by **1a** could be attributed, at least in part, to the suppression of *c-myc* and Bcl-2 transcription *via* BRD4 inhibition. Additionally, ERK 1/2, p-ERK 1/2 and PARP





Fig. 5 Chromatin immunoprecipitation (ChIP) analysis showed that **1a** selectively decreased the binding of BRD4 to MYC, Bcl-2 and CDK6, but not housekeeping genes (B2M) in (a) A375 and (b) A2058 cells. Bar graphs represented the mean enrichment relative to input and error bars reflect standard deviation of results derived from biological triplicate experiments. Significantly different at *** $p < 0.01$. Error bars represent the standard deviations of the results from three independent experiments.

expression were also down-regulated in A375 and A2058 cells after treatment with **1a** (Fig. 6a and b). Aberrant ERK 1/2 activation is implicated in numerous tumors, while PARP promotes cell survival due to its role in DNA repair. Therefore, the inhibition of ERK 1/2 and/or PARP activity by complex **1a** may represent an alternative mechanism by which the complex exerts anti-proliferative activity.

BRD4 inhibitor displays anti-proliferative activity in an animal model of melanoma

To further explore the relationship between BRD4 inhibition and cytotoxic activity, we also tested the metal complexes **1**, **1a–1j** from the second round of screening and **9–11** with chemical structures similar to **1a** from the first round of screening for their *in vitro* anticancer activity against A375 cells. Plotting the antiproliferative IC_{50} values of the complexes against the IC_{50} values for the inhibition of BRD4(1)-H4AcK4 binding revealed a positive relationship ($r = 0.8207$, $n = 14$) (Fig. 6g), suggesting that the anticancer activity of the metal complexes may be mediated by their inhibition of BRD4 binding activity.

To further investigate the antitumor activity exerted by BRD4 inhibition, we explored the biological activity of **1a** in a mouse xenograft tumor model. Six to seven week-old male CB.17 SCID mice were injected subcutaneously with human malignant melanoma A375 cells, and after the establishment of palpable tumors, were administered with **1a** (100 mg kg^{-1}) or vehicle (13% DMSO in normal saline) intraperitoneally (i.p.) once daily

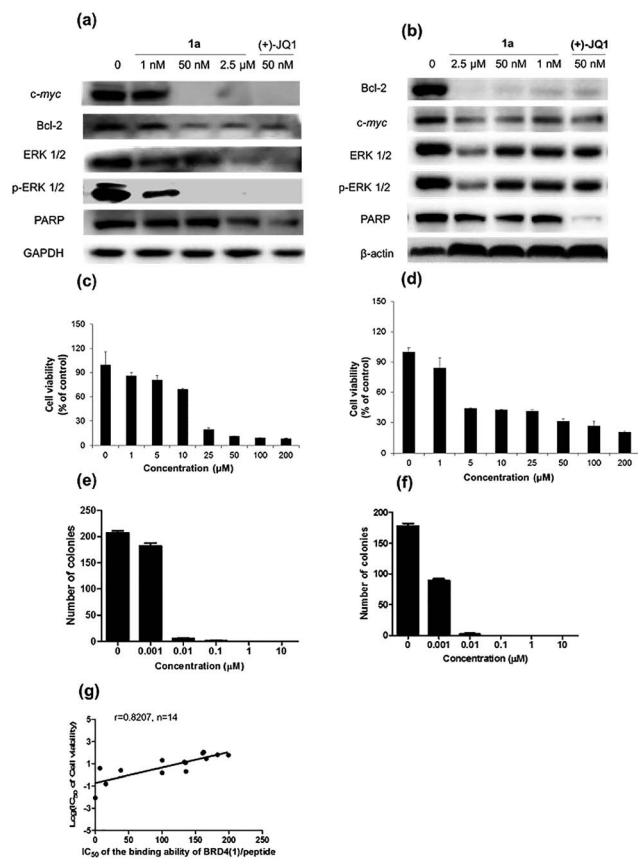


Fig. 6 Immunoblotting analysis of the effect of **1a** and (+)-JQ1 treatment in (a) A375 and (b) A2058 cells. Densitometry analysis revealed that **1a** inhibited c-myc, Bcl-2, ERK 1/2, p-ERK 1/2 and PARP expression. Dose response analysis of cell viability of complex **1a** against (c) A375 cells and (d) A2058 cells. Error bars represent the standard deviations of the results from three independent experiments. Normalized proliferation curves in the colony formation assay for (e) A375 and (f) A2058 cells treated with vehicle or **1a** (0.001–10 μM) measured by crystal violet staining. Error bars represent the standard deviations of the results from three independent experiments. (g) The relationship between the IC_{50} of the binding ability of BRD4(1)/peptide and the log of IC_{50} of A375 cell viability, and a trend of positive correlation was observed ($r = 0.8207$, $n = 14$).

for 16 days. Encouragingly, the treated tumors were *ca.* 40% smaller than the control tumors over the course of the treatment, with a significant difference in estimated tumor volume being observed after 16 days (Fig. 7a–c). Tumor weight measurements after sacrifice confirmed a reduction in tumor growth in mice administered with **1a** (Fig. 7d). We also observed that the treated mice exhibited no signs of weight loss over the course of the experiment (Fig. 7e). Taken together, these results indicate that complex **1a** significantly inhibited the growth of melanoma tissue in an *in vivo* xenograft model, without causing overt toxicity to the mice.

Microarray analysis was performed on the excised tumor tissues to identify signaling pathways that were up-regulated or down-regulated by complex **1a**. The results showed that treatment with **1a** resulted in MYC down-regulation as well as a significant decrease in expression of the c-myc target gene set in



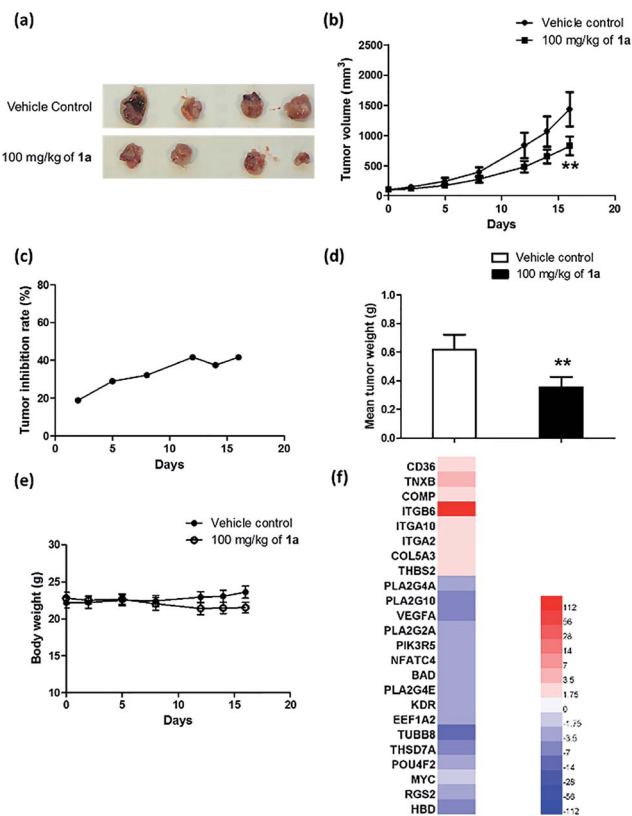


Fig. 7 Anti-proliferative activity of **1a** in an *in vivo* xenograft model of melanoma. (a) Photographs of dissected tumors from the control (vehicle) and treatment (**1a**, 100 mg kg⁻¹). (b) Average A375 tumour volume in the control group and treatment group (**1a**, 100 mg kg⁻¹). Each group contained six mice and results are reported as the values of the mean \pm SEM. (c) Tumour inhibition of A375 xenografts in the treatment group (**1a**, 100 mg kg⁻¹) expressed as percentage reduction in tumor volume compared to the control group. The results were analyzed using the Student's *t*-test. Significantly different at 0.01 < *p* < 0.05. (d) Average tumor weight of the vehicle control group versus the treatment group (**1a**, 100 mg kg⁻¹). (e) Average body weight of the two groups. Each group contained six mice, and results were reported as the values of the mean \pm SEM. The results were analyzed using the Student's *t*-test. Significantly different at 0.01 < *p* < 0.05. (f) Heat map of regulated genes of the ECM pathway and VEGF signaling pathway following treatment with **1a**. The color scale in the inset represents the log-fold change of expression compared with untreated control.

the tumor tissues. Moreover, complex **1a** up-regulated genes in the extracellular matrix (ECM) pathway while down-regulating genes in the VEGF signaling pathway (Table S2† and Fig. 7f). Differential ECM gene expression is an important marker of metastatic activity, and BRD4 modulates the expression of many ECM genes that are dysregulated in tumors.⁵⁴ Moreover, increased MYC activity can also suppress ECM gene expression.^{55–57} Hence, the up-regulation of the ECM pathway in tumor tissues could potentially be attributed to the inhibition of BRD4-directed transcription by **1a** *in vivo*. Furthermore, MYC promotes angiogenesis through the up-regulation of VEGF.^{58,59} Therefore, the down-regulation of genes in the VEGF signaling pathway could also be attributed to the effects of **1a** on BRD4-mediated transcription *in vivo*.

Conclusions

In conclusion, an iridium(III) complex **1a** was found to be a potential modulator of the epigenetic reader protein BRD4. Complex **1a** inhibited the PPI between BRD4 and an acetylated histone peptide as revealed by multiple biochemical assays, including FRET, AlphaScreen and FP assays. Although mass spectrometry data suggested that **1a** binds to histidine residues with the loss of ACN ligands, **1a** was found not to significantly interact with other histidine-containing proteins such as caspase-6 and STAT3. Additionally, complex **1a** displaced BRD4 from chromatin and hence inhibited *c-myc* expression in melanoma cells through blocking the binding of BRD4 to the *c-myc* promoter. Cytotoxicity and colony formation experiments suggested **1a** is capable of anti-proliferative activity in melanoma cells, possibly through down-regulation of *c-myc* protein expression. Finally, complex **1a** significantly repressed A375 melanoma xenograft growth in an *in vivo* mouse model without causing visible toxicity to the mice. Preliminary structure-activity analysis indicated that the nature of the metal ion and the C^N and N^N co-ligands were important for the biological activity of **1a**. To our knowledge, complex **1a** represents the first metal-based inhibitor of BRD4 and of any BET bromodomain-containing protein in general. We envision that complex **1a** may serve as a useful scaffold for the development of more potent epigenetic agents against cancers such as melanoma. Additionally, the irreversible and selective nature of BRD4 inhibition by **1a** may allow the complex to be used at lower dosages in potential *in vivo* applications.

Experimental

Materials and cell lines

All the chemicals were purchased from Sigma-Aldrich and were used as received. BRD4 bromodomain 1 TR-FRET assay kit and BRD4 bromodomain 2 TR-FRET assay kitware purchased from Cayman Chemical (Ann Arbor, MI, USA). Fetal bovine serum (FBS) and Dulbecco's modified Eagle's medium (DMEM) were obtained from Gibco BRL (Gaithersburg, MD, USA). All other reagents and chemicals were obtained from commercial sources.

General experimental

Mass spectrometry was performed at the Mass Spectroscopy Unit at the Department of Chemistry, Hong Kong Baptist University, Hong Kong (China). Deuterated solvents for NMR purposes were obtained from Armar and used as received.¹H and ¹³C NMR were recorded on a Bruker Avance 400 spectrometer operating at 400 MHz (¹H) and 100 MHz (¹³C). ¹H and ¹³C chemical shifts were referenced internally to solvent shift (acetonitrile-*d*₃: ¹H, δ 1.94, ¹³C δ 118.7; acetone-*d*₆: ¹H δ 2.05, ¹³C δ 29.7). Chemical shifts (δ) are quoted in ppm, the downfield direction being defined as positive. Uncertainties in chemical shifts are typically \pm 0.01 ppm for ¹H and \pm 0.05 for ¹³C. Coupling constants are typically \pm 0.1 Hz for ¹H-¹H and \pm 0.5 Hz



for ^1H – ^{13}C couplings. The following abbreviations are used for convenience in reporting the multiplicity of NMR resonances: s, singlet; d, doublet; t, triplet; q, quartet; m, multiplet. All NMR data was acquired and processed using standard Bruker software (Topspin).

Cell cultures

The human melanoma A375, A2058 cell lines were purchased from the Bioresource Collection and Research Center (Hsinchu, Taiwan, ROC) and the American Type Culture Collection (Manassas, VA, USA).

Time-resolved-fluorescence resonance energy transfer (TR-FRET) assay

The disruption of BRD4/tetra-acetylated lysine histone 4 peptide (H4AcK4) binding by the tested complexes was evaluated according to the manufacturer's instructions (Cayman Chemical, Carlsbad, CA). Briefly, the tested compounds, BRD4 bromodomain 1 or BRD4 bromodomain 2 europium chelate and BRD4 bromodomain 1 or BRD4 bromodomain 2 ligand/APC acceptor mixture were dissolved in $1 \times$ TR-FRET buffer provided by the manufacturer and stored in -80°C before use. To each test well, $10\ \mu\text{L}$ of the diluted BRD4 bromodomain 1 europium chelate was added. Indicated concentrations of the tested compounds and (+)-JQ1 were then added to the well and incubated for 15 min at room temperature and dark condition to allow pre-equilibration of the compounds with the BRD4 bromodomain 1 or BRD4 bromodomain 2 europium chelate. Subsequently, $5\ \mu\text{L}$ of the BRD4 bromodomain 1 or BRD4 bromodomain 2 ligand/APC acceptor mixture was added to each tested well. The microtitre plate was sealed and incubated at room temperature in the dark for 1 h. The experimental results were qualified using a spectrophotometer by measuring the dual emissions at 620 nm and 670 nm using an excitation at 340 nm. Data analysis is performed using the TR-FRET ratio (670 nm emission/620 nm emission).

Mass spectrometry analysis

$1\ \mu\text{g}$ BRD4(1) and $25\ \mu\text{M}$ complex **1a** were incubated in reaction buffer (10 mM Tris-HCl, pH = 7.5, 500 mM NaCl) at 25°C for 2 h. Positive ion mass spectra were acquired as accurate mass centroid data using a Thermo Q Exactive with a nano spray source, connected to a thermo Easy-nLC 1000 HPLC system. Survey full-scan MS spectra (from m/z 310–1800) were acquired in the Orbitrap analyzer with resolution $r = 70\ 000$ at m/z 400. The elemental composition was calculated using Xcalibur 2.1 for the $[\text{M} + \text{H}]^+$.

For amino acid mass spectra analysis, histidine were incubated with complex **1a** in Tris-HCl buffer (10 mM Tris-HCl, pH = 7.5, 500 mM NaCl) 30 min, then were directly injected into an ESI time-of-flight mass spectrometer at a flow rate of $5\ \mu\text{L}\ \text{min}^{-1}$. The capillary voltage was set at $-4500\ \text{V}$, and the dry N_2 gas flow was $4.0\ \text{L}\ \text{min}^{-1}$ at 180°C . Data were analyzed by the software Bruker Daltonics Data Analysis.

Chromatin immunoprecipitation

Approximately 1×10^8 A375 and A2058 cells were treated with 500 nM complex **1a** or DMSO for 24 h and cross-linked with $1 \times$ crosslink solution (1.1% formaldehyde, 5 mM 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES), pH 7.3, 10 mM NaCl, 100 μM ethylenediaminetetraacetic acid (EDTA), 50 μM ethylene glycol tetraacetic acid (EGTA)) followed by two washes using phosphate buffered saline (PBS). Cells were scraped and frozen in liquid nitrogen. ChIP-PCR analysis was done following a published protocol.⁶⁰

Immunoblotting

Cells were harvested to obtain whole-cell extracts by the addition of one volume of 250 mM Tris-HCl (pH 6.8), 20% glycerol, 2% sodium dodecyl sulfate (SDS), 5% 2-mercaptoethanol, and 0.2% bromophenol blue to cells in one volume of PBS followed by boiling for 5 min. Samples were resolved on sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS/PAGE) gels and transferred to polyvinylidene fluoride (PVDF) membranes. Blots were probed with antibodies to *c-myc*, Bcl-2, ERK 1/2, p-ERK 1/2, PARP, GAPDH and beta-actin (cell signaling). After incubation with secondary antibodies (Santa Cruz Biotechnology), blots were developed with ECL reagent (ThermoFisher).

Animal materials

In this study, the use of animals complied with the Guiding Principles in the Care and Use of Animals of the American Physiology Society and was approved by the Animal Care and Use Committee at the National Kaohsiung Medical University. Six to seven weeks old male CB.17 SCID mice were purchased from BioLascoTaiwan Co., LTD and quarantined for one week. During the experiment period, 5 mice were housed in one cage. All animals were hosted in the Da-Hu animal facility in a 12 h light/12 h dark cycle at 19 – 25°C . Animals had free access to rodent pellet food and water *ad libitum*. The experimental protocol of animal study was reviewed and approved by the Institutional Animal Care and Use Committee, DCB.

Xenograft tumor assay

The human A375 melanoma cells used for implantation were harvested during log phase growth and re-suspended in phosphate buffered saline to a concentration containing 1×10^7 cells per mL. Each six to seven weeks' old male CB.17 SCID mice was injected subcutaneously (s.c.) in the flank with 1.5×10^7 cells in 0.15 mL of a 50% matrigel solution (BD Biosciences, MA, USA). When the average tumor volume had reached $100\ \text{mm}^3$, the mice were randomly divided into 2 groups and were administrated with **1a** (50 or 100 $\text{mg}\ \text{kg}^{-1}$) or vehicle (13% DMSO in normal saline) intraperitoneally (i.p.) once daily for 16 days. The diameters of xenograft tumors were measured with Digimatic caliper (Series no. 500, Mitutoyo Corp., Japan) and the tumor volume (in mm^3) was calculated using the formula:

$$\text{Tumor volume} = (w^2 \times l)/2$$



where w = width and l = length in diameter (mm) of the tumor. The treatment and control groups contained five mice each.

Tumors were measured twice per week using calipers. The percentage of tumor growth inhibition (TGI) was calculated using the following formula:

$$\%TGI = [1 - (T/C)] \times 100\%$$

where T and C represent the mean tumor volumes of the treatment group and the control group, respectively.

Animals were weighed twice weekly until the completion of the study. The body weight change was calculated as the percentage increase in body weight compared to the initial body weight.

Microarray analysis

Xenografts in the mouse models were removed 16 days after the administration of A375 cells. Total RNA from tumours from three independent vehicle and treatment mice were prepared using TRIzol reagent (Invitrogen). Gene expression profiles were examined using Mouse One Array DNA Microarray chips (Phalanx Biotech Group, Inc., Hsinchu, Taiwan). The criterion for affected genes was a significant difference in tumor RNA levels between the treatment and vehicle groups ($p < 0.05$), with at least a 2-fold difference. Gene ontology analysis was performed using PANTHER.

Statistical analysis

Data were analyzed using the Prism software version 6 (Graph Pad software Inc., San Diego, CA, USA). The Student's t -test was used for comparisons between the two groups. The correlation between BRD4 inhibition and cytotoxic activity was calculated using the Pearson's correlation coefficient equation. Quantitative data are reported as mean \pm SEM from at least three independent experiments. Differences were considered statistically significant at $p < 0.05$. All statistical tests were two-sided.

All synthetic methods, characterization and biological assays details are given in the ESI.†

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Notes and references

- 1 B. Li, M. Carey and J. L. Workman, *Cell*, 2007, **128**, 707–719.
- 2 G. E. Zentner and S. Henikoff, *Nat. Struct. Mol. Biol.*, 2013, **20**, 259–266.
- 3 A. J. Ruthenburg, H. Li, D. J. Patel and C. D. Allis, *Nat. Rev. Mol. Cell Biol.*, 2007, **8**, 983–994.
- 4 C. H. Arrowsmith, C. Bountra, P. V. Fish, K. Lee and M. Schapira, *Nat. Rev. Drug Discovery*, 2012, **11**, 384–400.
- 5 S. D. Taverna, H. Li, A. J. Ruthenburg, C. D. Allis and D. J. Patel, *Nat. Struct. Mol. Biol.*, 2007, **14**, 1025–1040.
- 6 C.-w. Chung, A. W. Dean, J. M. Woolven and P. Bamborough, *J. Med. Chem.*, 2012, **55**, 576–586.
- 7 L. Zeng and M.-M. Zhou, *FEBS Lett.*, 2002, **513**, 124–128.
- 8 B. Florence and D. V. Faller, *Front. Biosci.*, 2001, **6**, d1008–d1018.
- 9 C.-w. Chung and D. F. Tough, *Drug Discovery Today: Ther. Strategies*, 2012, **9**, e111–e120.
- 10 Z. Yang, N. He and Q. Zhou, *Mol. Cell Biol.*, 2008, **28**, 967–976.
- 11 A. Dey, A. Nishiyama, T. Karpova, J. McNally and K. Ozato, *Mol. Biol. Cell*, 2009, **20**, 4899–4909.
- 12 M. F. Segura, B. Fontanals-Cirera, A. Gaziol-Sovran, M. V. Guijarro, D. Hanniford, G. Zhang, P. Gonzalez-Gomez, M. Morante, L. Jubierre, W. Zhang, F. Darvishian, M. Ohlmeyer, I. Osman, M. M. Zhou and E. Hernando, *Cancer Res.*, 2013, **73**, 6264–6276.
- 13 P. Filippakopoulos, J. Qi, S. Picaud, Y. Shen, W. B. Smith, O. Fedorov, E. M. Morse, T. Keates, T. T. Hickman, I. Felletar, M. Philpott, S. Munro, M. R. McKeown, Y. Wang, A. L. Christie, N. West, M. J. Cameron, B. Schwartz, T. D. Heightman, N. la Thangue, C. A. French, O. Wiest, A. L. Kung, S. Knapp and J. E. Bradner, *Nature*, 2010, **468**, 1067–1073.
- 14 J. E. Delmore, G. C. Issa, M. E. Lemieux, P. B. Rahl, J. Shi, H. M. Jacobs, E. Kastiris, T. Gilpatrick, R. M. Paranal, J. Qi, M. Chesi, A. C. Schinzel, M. R. McKeown, T. P. Heffernan, C. R. Vakoc, P. L. Bergsagel, I. M. Ghobrial, P. G. Richardson, R. A. Young, W. C. Hahn, K. C. Anderson, A. L. Kung, J. E. Bradner and C. S. Mitsiades, *Cell*, 2011, **146**, 904–917.
- 15 J. A. Mertz, A. R. Conery, B. M. Bryant, P. Sandy, S. Balasubramanian, D. A. Mele, L. Bergeron and R. J. Sims, *Proc. Natl. Acad. Sci. U. S. A.*, 2011, **108**, 16669–16674.
- 16 C. G. Hartinger and P. J. Dyson, *Chem. Soc. Rev.*, 2009, **38**, 391–401.
- 17 A. V. Klein and T. W. Hambley, *Chem. Rev.*, 2009, **109**, 4911–4920.
- 18 G. Gasser, I. Ott and N. Metzler-Nolte, *J. Med. Chem.*, 2011, **54**, 3–25.



- 19 A. L. Noffke, A. Habtemariam, A. M. Pizarro and P. J. Sadler, *Chem. Commun.*, 2012, **48**, 5219–5246.
- 20 A. Meyer, C. P. Bagowski, M. Kokoschka, M. Stefanopoulou, H. Alborzina, S. Can, D. H. Vlecken, W. S. Sheldrick, S. Wolf and I. Ott, *Angew. Chem., Int. Ed.*, 2012, **51**, 8895–8899.
- 21 A. de Almeida, B. L. Oliveira, J. D. G. Correia, G. Soveral and A. Casini, *Coord. Chem. Rev.*, 2013, **257**, 2689–2704.
- 22 C. G. Hartinger, M. Groessel, S. M. Meier, A. Casini and P. J. Dyson, *Chem. Soc. Rev.*, 2013, **42**, 6186–6199.
- 23 H. Huang, P. Zhang, B. Yu, Y. Chen, J. Wang, L. Ji and H. Chao, *J. Med. Chem.*, 2014, **57**, 8971–8983.
- 24 A. Leonidova, V. Pierroz, L. A. Adams, N. Barlow, S. Ferrari, B. Graham and G. Gasser, *ACS Med. Chem. Lett.*, 2014, **5**, 809–814.
- 25 D. L. Ma, W. L. Wong, W. H. Chung, F. Y. Chan, P. K. So, T. S. Lai, Z. Y. Zhou, Y. C. Leung and K. Y. Wong, *Angew. Chem., Int. Ed.*, 2008, **47**, 3735–3739.
- 26 E. Meggers, *Chem. Commun.*, 2009, 1001–1010.
- 27 X. Meng, M. L. Leyva, M. Jenny, I. Gross, S. Benosman, B. Fricker, S. Harlepp, P. Hébraud, A. Boos, P. Wlosik, P. Bischoff, C. Sirlin, M. Pfeffer, J.-P. Loeffler and C. Gaidon, *Cancer Res.*, 2009, **69**, 5458–5466.
- 28 C. M. Che and F. M. Siu, *Curr. Opin. Chem. Biol.*, 2010, **14**, 255–261.
- 29 K. J. Kilpin and P. J. Dyson, *Chem. Sci.*, 2013, **4**, 1410–1419.
- 30 D. L. Ma, H. Z. He, K. H. Leung, D. S. H. Chan and C. H. Leung, *Angew. Chem., Int. Ed.*, 2013, **52**, 7666–7682.
- 31 D. L. Ma, L. J. Liu, K. H. Leung, Y. T. Chen, H. J. Zhong, D. S. H. Chan, H. M. D. Wang and C. H. Leung, *Angew. Chem., Int. Ed.*, 2014, **53**, 9178–9182.
- 32 C. H. Leung, L. J. Liu, L. Lu, B. He, D. W. J. Kwong, C. Y. Wong and D. L. Ma, *Chem. Commun.*, 2015, **51**, 3973–3976.
- 33 J. Singh, R. C. Petter, T. A. Baillie and A. Whitty, *Nat. Rev. Drug Discovery*, 2011, **10**, 307–317.
- 34 L. Feng, Y. Geisselbrecht, S. Blanck, A. Wilbuer, G. E. Atilla-Gokcumen, P. Filippakopoulos, K. Kräling, M. A. Celik, K. Harms, J. Maksimoska, R. Marmorstein, G. Frenking, S. Knapp, L.-O. Essen and E. Meggers, *J. Am. Chem. Soc.*, 2011, **133**, 5976–5986.
- 35 S. Blanck, Y. Geisselbrecht, K. Kraling, S. Middel, T. Mietke, K. Harms, L.-O. Essen and E. Meggers, *Dalton Trans.*, 2012, **41**, 9337–9348.
- 36 C. H. Leung, H. J. Zhong, H. Yang, Z. Cheng, D. S. H. Chan, V. P. Y. Ma, R. Abagyan, C. Y. Wong and D. L. Ma, *Angew. Chem., Int. Ed.*, 2012, **51**, 9010–9014.
- 37 S. Dieckmann, R. Riedel, K. Harms and E. Meggers, *Eur. J. Inorg. Chem.*, 2012, 813–821.
- 38 S. Mollin, S. Blanck, K. Harms and E. Meggers, *Inorg. Chim. Acta*, 2012, **393**, 261–268.
- 39 R. Anand, J. Maksimoska, N. Pagano, E. Y. Wong, P. A. Gimotty, S. L. Diamond, E. Meggers and R. Marmorstein, *J. Med. Chem.*, 2009, **52**, 1602–1611.
- 40 G. E. Atilla-Gokcumen, L. Costanzo and E. Meggers, *JBIC, J. Biol. Inorg. Chem.*, 2011, **16**, 45–50.
- 41 S. Blanck, T. Cruchter, A. Vultur, R. Riedel, K. Harms, M. Herlyn and E. Meggers, *Organometallics*, 2011, **30**, 4598–4606.
- 42 C. Streu, L. Feng, P. J. Carroll, J. Maksimoska, R. Marmorstein and E. Meggers, *Inorg. Chim. Acta*, 2011, **377**, 34–41.
- 43 M. S. Lowry, W. R. Hudson, R. A. Pascal and S. Bernhard, *J. Am. Chem. Soc.*, 2004, **126**, 14129–14135.
- 44 C. Li, M. Yu, Y. Sun, Y. Wu, C. Huang and F. Li, *J. Am. Chem. Soc.*, 2011, **133**, 11231–11239.
- 45 F. Piccioli, S. Sabatini, L. Messori, P. Orioli, G. H. Ch and B. K. Keppler, *J. Inorg. Biochem.*, 2004, **98**, 1135–1142.
- 46 A. R. Timerbaev, L. S. Foteeva, A. V. Rudnev, J. K. Abramski, K. Polec-Pawlak, C. G. Hartinger, M. Jarosz and B. K. Keppler, *Electrophoresis*, 2007, **28**, 2235–2240.
- 47 O. Domotor, C. G. Hartinger, A. K. Bytzeck, T. Kiss, B. K. Keppler and E. A. Enyedy, *JBIC, J. Biol. Inorg. Chem.*, 2013, **18**, 9–17.
- 48 R. Trondl, P. Heffeter, C. R. Kowol, M. A. Jakupec, W. Berger and B. K. Keppler, *Chem. Sci.*, 2014, **5**, 2925–2932.
- 49 M. I. Webb and C. J. Walsby, *Dalton Trans.*, 2015, DOI: 10.1039/c5dt02021b.
- 50 F. Kratz, M. Hartmann, B. Keppler and L. Messori, *J. Biol. Chem.*, 1994, **269**, 2581–2588.
- 51 L. Messori, P. Orioli, D. Vullo, E. Alessio and E. Iengo, *Eur. J. Biochem.*, 2000, **267**, 1206–1213.
- 52 M. A. Stoff-Khalili, P. Dall and D. T. Curiel, *Cancer Gene Ther.*, 2006, **13**, 633–647.
- 53 K. J. Falkenberg and R. W. Johnstone, *Nat. Rev. Drug Discovery*, 2014, **13**, 673–691.
- 54 N. P. S. Crawford, J. Alsarraj, L. Lukes, R. C. Walker, J. S. Officewala, H. H. Yang, M. P. Lee, K. Ozato and K. W. Hunter, *Proc. Natl. Acad. Sci. U. S. A.*, 2008, **105**, 6380–6385.
- 55 H. A. Coller, C. Grandori, P. Tamayo, T. Colbert, E. S. Lander, R. N. Eisenman and T. R. Golub, *Proc. Natl. Acad. Sci. U. S. A.*, 2000, **97**, 3260–3265.
- 56 J. D. Watson, S. K. Oster, M. Shago, F. Khosravi and L. Z. Penn, *J. Biol. Chem.*, 2002, **277**, 36921–36930.
- 57 E. R. Lawlor, L. Soucek, L. Brown-Swigart, K. Shchors, C. U. Bialucha and G. I. Evan, *Cancer Res.*, 2006, **66**, 4591–4601.
- 58 R. Chen, J. H. Yik, Q. J. Lew and S. H. Chao, *BioMed Res. Int.*, 2014, **2014**, 232870.
- 59 S. W. J. Ember, J. Y. Zhu, S. H. Olesen, M. P. Martin, A. Becker, N. Berndt, G. I. Georg and E. Schönbrunn, *ACS Chem. Biol.*, 2014, **9**, 1160–1171.
- 60 P. B. Rahl, C. Y. Lin, A. C. Seila, R. A. Flynn, S. McCuine, C. B. Burge, P. A. Sharp and R. A. Young, *Cell*, 2010, **141**, 432–445.

