

RESEARCH ARTICLE

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View Journal | View IssueCite this: *RSC Med. Chem.*, 2025, 16, 5419Tailored SirReal-type inhibitors enhance SIRT2 inhibition through ligand stabilization and disruption of NAD⁺ co-factor bindingRicky Wirawan, ^{†a} Matthias Frei, ^{†a} Anna Heider, ^b Niklas Papenkordt, ^c Florian Friedrich, ^c Thomas Wein, ^a Manfred Jung, ^c Michael Groll, ^b Eva M. Huber ^b and Franz Bracher ^{*a}

Human sirtuin 2 (SIRT2) is an NAD⁺ dependant enzyme that has been linked to the pathogenesis of various diseases, making it a promising target for pharmaceutical intervention. This study presents a systematic investigation on the inhibitory effects of SIRT2 inhibitors functionalized with diverse electrophilic functional groups. Guided by initial docking studies, we designed and synthesised 14 derivatives of two published potent lead structures **24a** and **SirReal2**. The most potent and subtype selective SIRT2 inhibitor **29** (RW-78) exhibits an IC₅₀ of 26 nM, which outperforms its lead structure **24a** (IC₅₀ = 79 nM) by a factor of 3. The increased potency of **29** is explained by halogen- π interactions with SIRT2 residues as visualized by X-ray crystallography. Furthermore, **29** interferes with NAD⁺ binding, highlighting co-factor displacement as a valid strategy to inhibit SIRT2. Additionally, we showed cellular target engagement *via* NanoBRET assays in HEK293T cells (EC₅₀ = 15 nM). Altogether our findings provide a deeper insight into the structure-activity relationships of these SirReal-type inhibitors and offer new avenues for optimisation of SIRT2 inhibitors.

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Introduction

Sirtuins, belonging to class III histone deacetylases, constitute a family of highly conservative NAD⁺-dependant proteins that are involved in the regulation of various biological processes such as metabolism, aging, inflammation and oxidative stress.¹ Although sirtuins were initially recognised solely for their deacetylation activity,² recent studies unveiled a broader catalytic scope that includes desuccinylation,³ demalonylation,³ demyristoylation⁴ and ADP-ribosylation.⁵ Furthermore, sirtuin substrates extend beyond histones to include other proteins, such as α -tubulin,⁶ p53 (ref. 7) and NF- κ B.⁸ Of the seven sirtuin subtypes, SIRT2 emerged as a key target, particularly due to its involvement in the pathogenesis of cancer,⁹ viral infections¹⁰ and neurodegenerative diseases.^{11,12} Despite numerous efforts over the past few years, the development of potent and subtype selective SIRT2 inhibitors with satisfactory physicochemical properties still

poses a significant challenge. Published SIRT2 inhibitors with drug-like properties have been shown to possess a wide range of structural diversity with various binding mechanisms (Fig. 1). SIRT2 inhibitors with greater potency tend to incorporate thioamide (**KPM-2** (ref. 13) and **TM**⁹) and thiourea (**AF8**)¹⁴ scaffolds that can form stable covalent intermediates with the essential co-factor NAD⁺. Additionally, SIRT2 inhibitors with other structural motifs such as oxadiazole (**Moniot_39**)¹⁵ and chroman-4-one (**6f**)¹⁶ were identified with IC₅₀ values covering low micromolar ranges. The alkaloid cytosine derived compound **NPD11033** was shown by Kudo *et al.* to likewise be a potent SIRT2 inhibitor.¹⁷ Further potent SIRT2 inhibitors such as **AGK2** were detected by means of high-throughput screening.¹⁸ Similarly, a series of aminothiazoles were discovered through library screening, in which the term sirtuin rearranging ligands (SirReals) was coined.^{19,20} Among these aminothiazoles, **SirReal2** displayed high inhibitory potency with an IC₅₀ value of 0.44 μ M.¹⁹ X-ray crystallographic studies on a SIRT2–**SirReal2**–NAD⁺ complex unveiled a ligand-induced structural rearrangement of the active site that generates the emergence of a selectivity pocket that is occupied by the 4,6-dimethyl-2-mercaptopyrimidine motif. Further optimisations employing triazole motifs led to SIRT2 inhibitors with improved potency (**Vogelmann_12** (**SH10**)²¹ and **Schiedel_9** and **Schiedel_10**²²). Extending efforts for the development of alternative SIRT2 inhibitors exploiting this selectivity pocket, Yang *et al.* synthesised a library of

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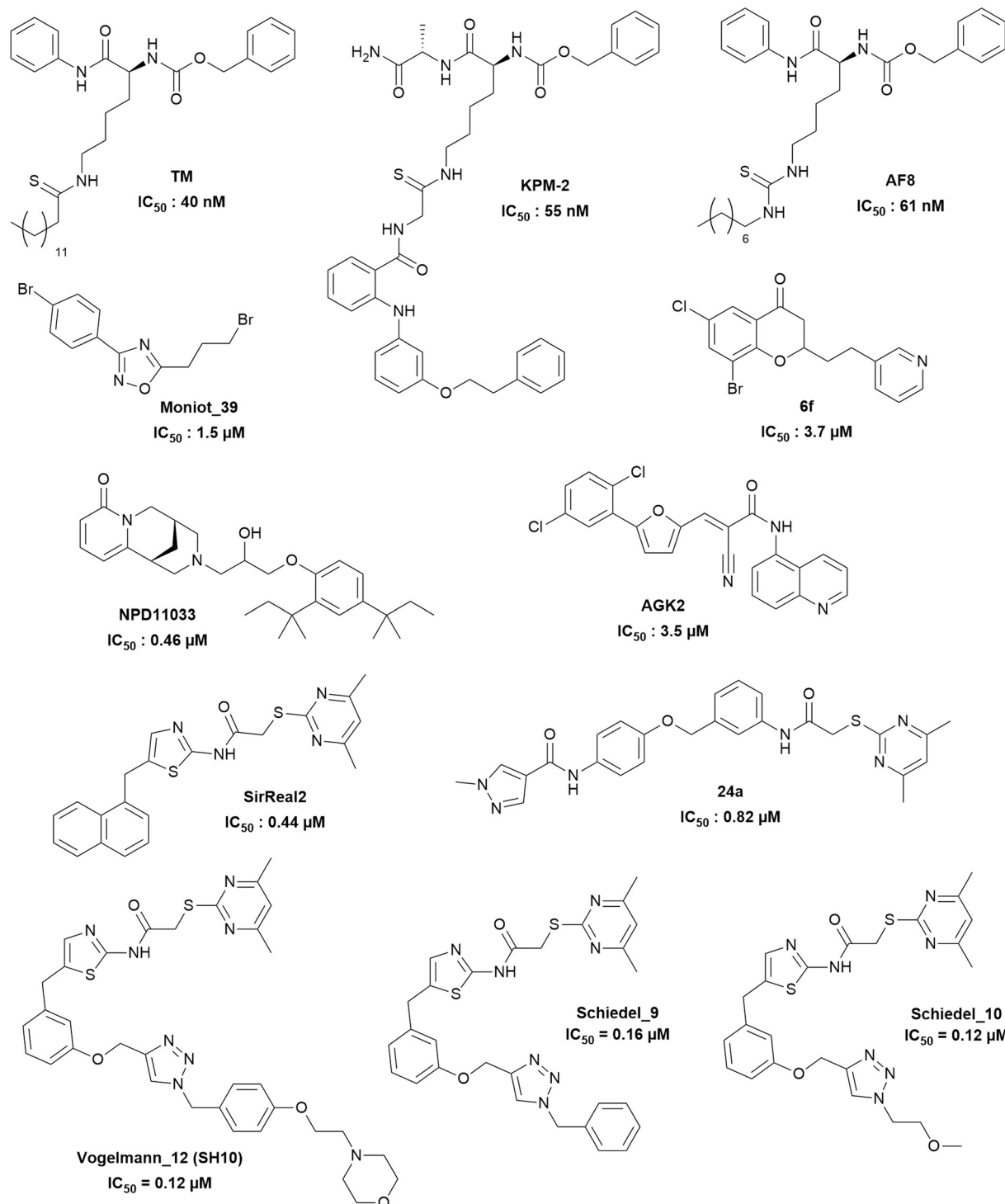


Fig. 1 Examples of published SIRT2 inhibitors.

N-acylaniline derivatives that identified **24a** as a potent and selective SIRT2 inhibitor with an IC_{50} value of 0.82 μ M.^{23,24}

Design rationale

Based on the published co-crystal structures of SIRT2 with either **24a** (PDB ID: 5YQO)²³ or **SirReal2** and NAD^+ (PDB ID:

4RMG),¹⁹ we hypothesised that significant increase in the potency of these lead structures can be achieved through strong, targeted interactions with the vicinal hydroxy groups of the ribose unit of NAD^+ by utilizing appropriate functional groups that enable reversible covalent binding or halogen bonding. Although reactive warheads such as Michael acceptors, β -lactones and β -lactams, epoxides and sulfonyl



fluorides have seen application in drug development and present themselves as attractive irreversible covalent binders, the potential for immunogenic reactions and off-target toxicity is almost inevitable.^{25–27} In contrast, reversible covalent inhibitors can dissociate from off-targets, reducing the potential of unwanted side effects while retaining extended binding at the intended target protein.²⁸ Several reversible covalent inhibitors have been successfully approved as therapeutic drugs, such as the boronic acid bortezomib that inhibits the 20S proteasome for the treatment of multiple myeloma,^{29,30} the nitrile-based dipeptidylpeptidase 4 (DPP4) inhibitor saxagliptin for the treatment of type 2 diabetes mellitus,³¹ and the aldehyde-bearing voxelotor for the treatment of sickle cell anaemia.³² Functional groups such as boronic acids, nitriles and aldehydes do not only possess the capability to undergo reversible covalent bonding, but can also form strong non-covalent interactions in the form of hydrogen bonding, *e.g.* with the ribose hydroxy groups of the co-factor NAD⁺. Comparably, non-covalent interactions *via* halogen bonding, which is sometimes referred to as the hydrophobic equivalent of hydrogen bonding,³³ represent an area of interest in current rational drug design approaches as demonstrated by the optimisation of PDE5 (ref. 34) and HIV-1 reverse transcriptase inhibitors^{35,36} and offer, as such, a promising drug optimisation technique. This non-covalent interaction arises from the anisotropic electron distribution in halogen residues, creating a σ -hole with depleted electron density that enables strong bonding with diverse nucleophiles.^{37–40} By tailoring lead structures **24a** and **SirReal2** with such functional groups, we aimed to achieve, in a (to our knowledge) unprecedented manner in the sirtuin field, reversible covalent binding and strong halogen bonding with the co-factor NAD⁺ of SIRT2 that strives for significant enhancement in potency (Fig. 2).

Encouraged by initial docking studies of the envisaged boronic acid derivatives of **24a** and **SirReal2** that showed poses of the boronic acid moiety in proximity to the vicinal

diol unit of NAD⁺ (Fig. 3), we continued our efforts⁴¹ in developing highly potent and sub-type selective SIRT2 inhibitors through systematic investigation of the effects of such functional groups on these two selected lead structures.

Results and discussion

Chemistry

The lead structure **24a** (here **10**) was synthesised according to literature with slight modifications in the chronology.^{23,24} Syntheses of functionalised **24a** derivatives **11–14** were initiated by the amide coupling of 4-amino-2-bromophenol (**2**) and 1-methyl-1*H*-pyrazole-4-carboxylic acid to give amide **4** (Scheme 1). Williamson ether synthesis of **4** with 3-nitrobenzyl bromide gave bromobenzene intermediate **6**, which afforded nitrile **9** with zinc(II) cyanide following published procedures,⁴² and boronic acid **8** following Miyaura borylation protocols⁴³ and subsequent oxidative cleavage of boronic acid pinacol ester **7**. In the final step, reduction of the nitrobenzene derivatives **6**, **8** and **9** followed by amide coupling with 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid gave bromobenzene derivative **11**, boronic acid **13** and nitrile **14**, respectively. In contrast, the introduction of the aldehyde functional group was performed in the final step due to its instability. Here, bromobenzene derivative **11** was formylated to aldehyde **12** with *in situ* generated carbon monoxide from *N*-formylsaccharin following protocols from Ueda *et al.*⁴⁴ A liquid–liquid extraction protocol⁴⁵ utilising sodium bisulfite to form a charged bisulfite adduct with subsequent regeneration of the aldehyde *via* basification was necessary, since purification with flash column chromatography alone did not afford the desired aldehyde **12** with sufficient purity. However, the obtained 3% yield was rather unsatisfactory, owing to stability issues. Nevertheless, the amount of product **12** obtained sufficed for chemical analysis and biological testing. Analogously, the preparation of the isomeric functionalised **24a** derivatives **29–34** followed a

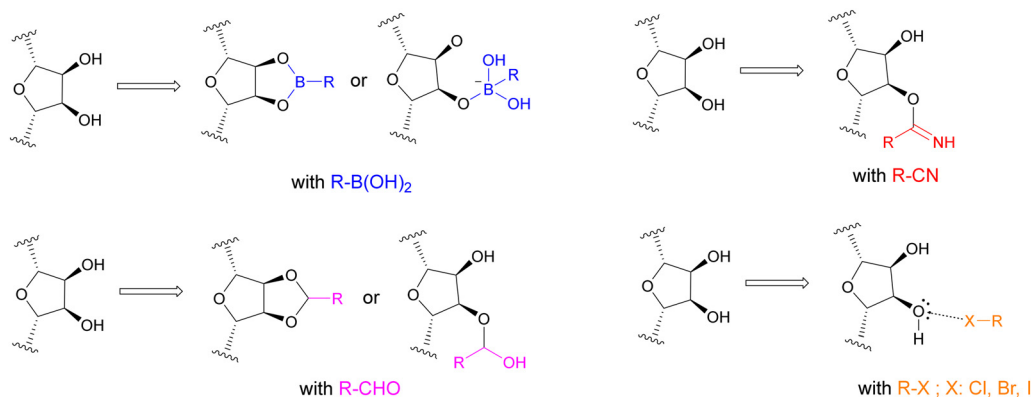


Fig. 2 Proposed binding mechanisms and interactions of the selected functional groups with the ribose unit of the co-factor NAD⁺. Boronic acids can undergo reversible covalent bonding with one or both hydroxy groups to form (cyclic) boronates. Nitriles can form iminoethers, and aldehydes can form hemiacetals or cyclic acetals. Halogen residues such as chlorine, bromine and iodine can form halogen bonds with the ribose hydroxy groups.



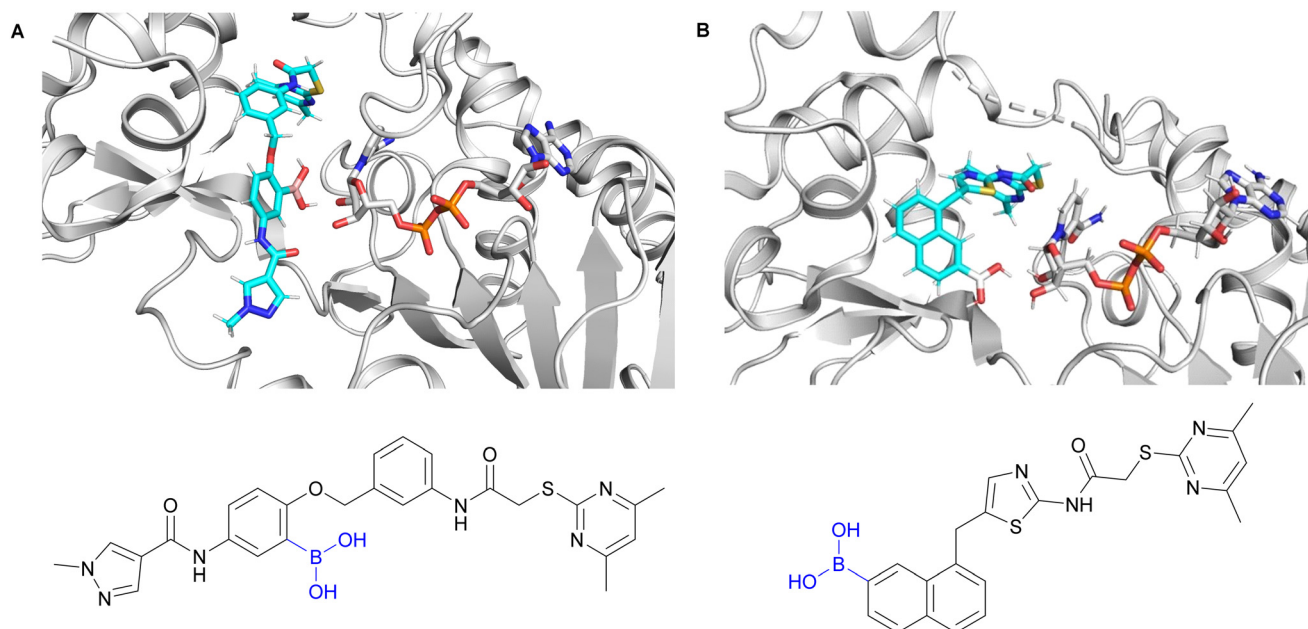


Fig. 3 (A) Docking poses of the envisaged **24a** boronic acid derivative (cyan) based on PDB ID: 5YQO²³ (NAD⁺ shown was extracted separately from PDB ID: 4RMG¹⁹) and (B) **SirReal2** boronic acid derivative (cyan) based on PDB ID: 4RMG¹⁹ in the presence of NAD⁺ showing the boronic acids in proximity to the vicinal diol unit of ribose of NAD⁺ for reversible covalent bonding.

similar synthetic route. Building blocks 4-amino-3-bromophenol (**18**) and 4-amino-3-iodophenol (**19**) were obtained from the reduction of 3-bromo-4-nitrophenol (**15**) and 3-iodo-4-nitrophenol (**16**), respectively. These phenols and the commercially available 4-amino-3-chlorophenol (**17**) were then subjected to Williamson ether synthesis prior to amide coupling with 1-methyl-1*H*-pyrazole-4-carboxylic acid as we noticed a significantly higher reactivity of the phenols compared to the aromatic amines. Similar to the previous synthetic route, bromobenzene intermediate **24** served as a key intermediate for the preparation of boronic acid **27** and nitrile **28**. Reduction of the nitrobenzene derivatives **23–25** and **28** and subsequent amide coupling with 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid in the final step afforded halogenated derivatives **29–31** and nitrile **34**, respectively. It is noteworthy that the amide coupling with 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid after the reduction of the boronic acid-bearing nitrobenzene **27** failed with various amide coupling reagents. To overcome this problem, the highly reactive bromoacetyl bromide was instead implemented, thereby generating a α -bromoacetamide intermediate that underwent nucleophilic substitution with 4,6-dimethylpyrimidine-2-thiol to give boronic acid **33**. Aldehyde **32** was obtained from bromobenzene derivative **30** with the same formylation⁴⁴ and bisulfite-mediated purification⁴⁵ protocol that was developed for the synthesis of aldehyde **12**.

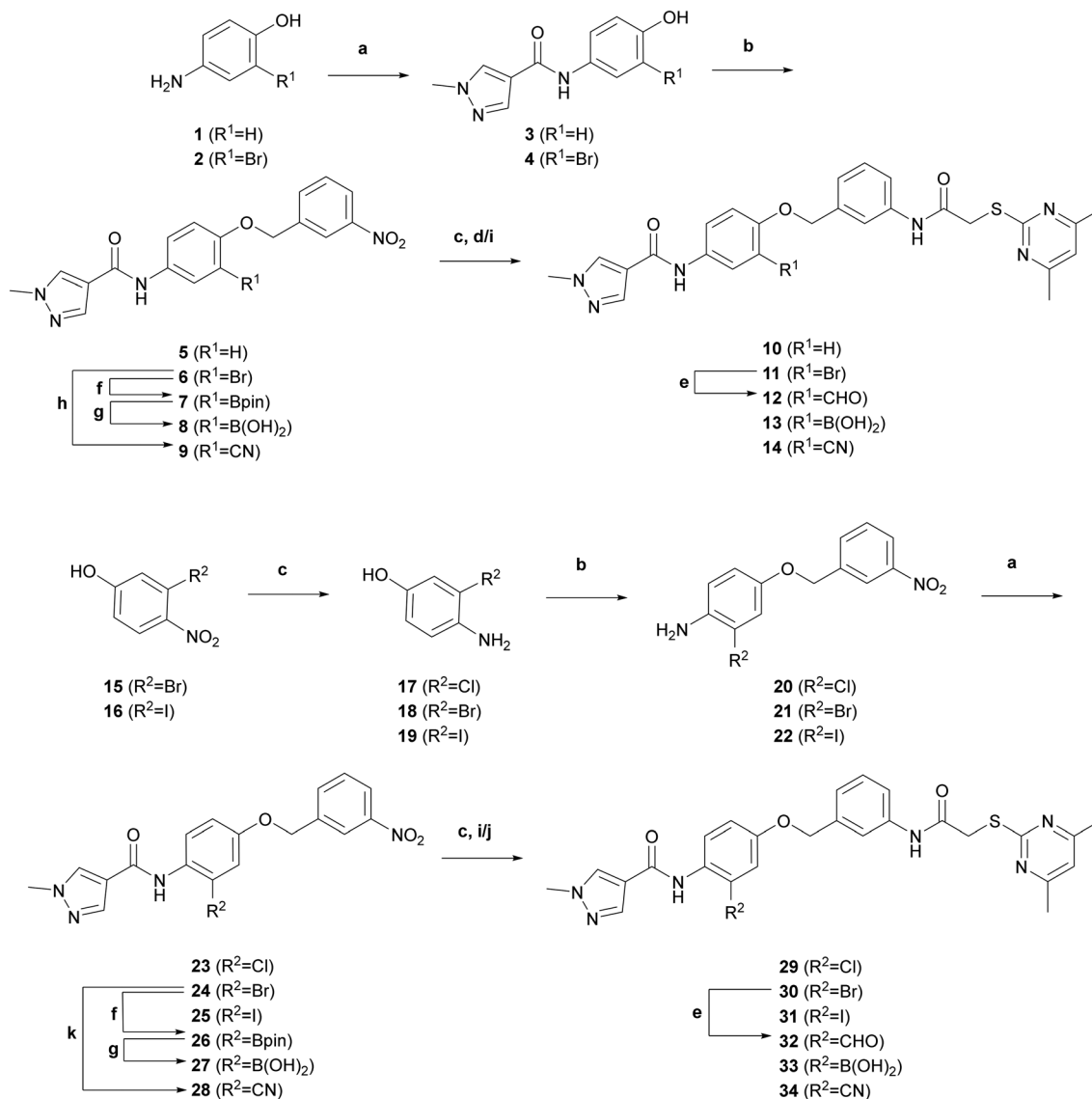
The synthesis of functionalised **SirReal2** derivatives was initiated by the preparation of aminothiazole **38** mainly according to literature (Scheme 2).²⁰ In deviation therefrom, intermediate 7-bromonaphthalene-1-amine (**37**) was prepared in a two-step process according to a

patented method.⁴⁶ Starting from 7-bromo-1-tetralone (**35**), oxime **36** was prepared and then converted to 7-bromonaphthalene-1-amine (**37**) *via* Semmler-Wolff aromatisation. Subsequently, a modified Meerwein arylation afforded aminothiazole **38** that served as a key intermediate for further functional group modifications, such as the palladium-catalysed synthesis^{42,47} of nitrile **39** and cyanomethyl derivative **40**. Notably, the introduction of the boronic acid necessitated the protection of the amine. *N*-Boc protection of aminothiazole **38** gave bromobenzene intermediate **41**, which underwent Miyaura borylation⁴³ to give boronic acid pinacol ester **42**. Oxidative cleavage of **42** and subsequent *N*-Boc cleavage with TFA gave boronic acid **44**. The functionalised **SirReal2** derivatives **45–48** were then obtained in the final step from **38–40** and **44** *via* amide coupling with 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid. Formylation of **45** was again performed in the final step of the synthesis, in this case using a protocol from Konishi *et al.*⁴⁸ that utilises a different phosphine ligand, which generated a better yield for the preparation of aldehyde **49**. Laborious purification of aldehyde **49** with the aforementioned liquid–liquid extraction protocol⁴⁵ *via* a bisulfite adduct was not necessary in this case.

In vitro sirtuin inhibitory activities

The determination of SIRT2 inhibitory activity was performed by Reaction Biology Corporation (Malvern, USA) with a fluorescence-based assay utilizing the fluorogenic peptide of p53 residues 379–382 (RHKK(Ac)-Fl). In general, the determined IC₅₀ values of literature-known lead



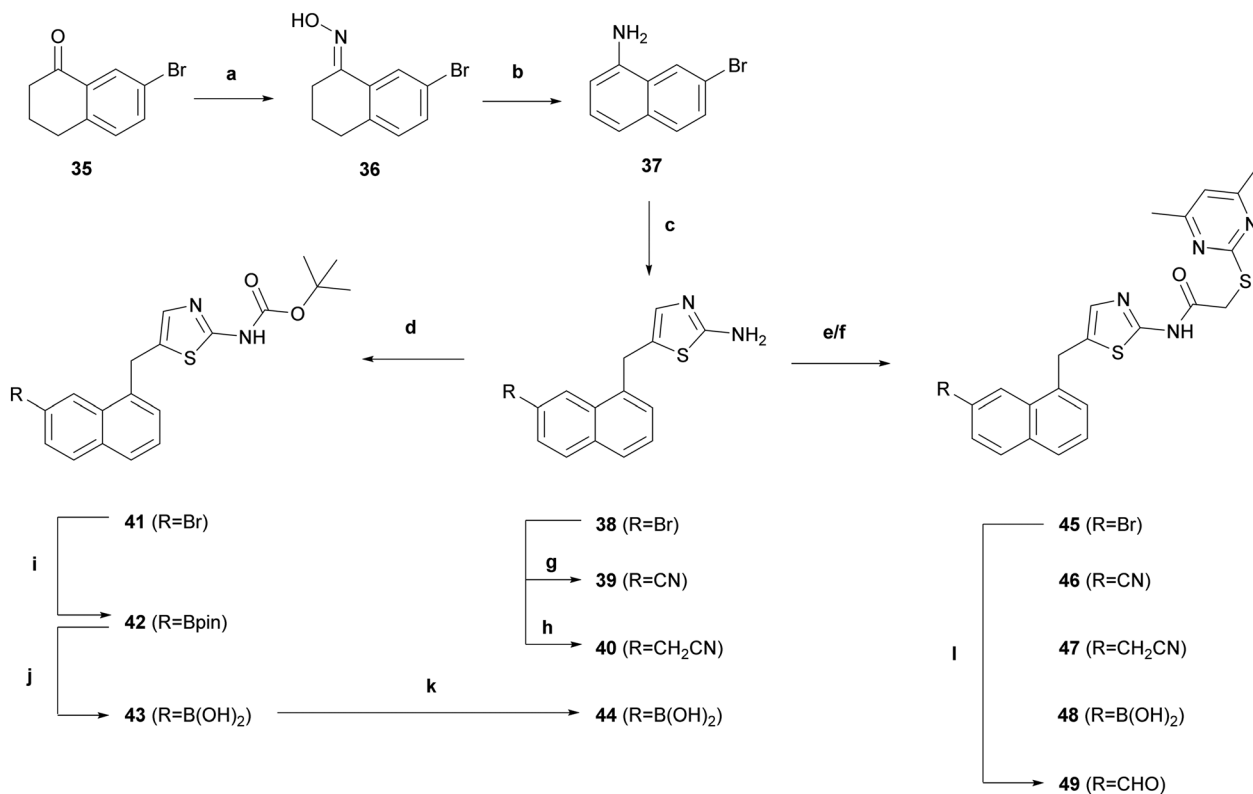


Scheme 1 Reagents and conditions: (a) 1-methyl-1*H*-pyrazole-4-carboxylic acid, HATU, DIPEA, THF, rt–65 °C, 3 h–6 d, 21–71%; (b) 3-nitrobenzyl bromide, K₂CO₃, DMF, 0 °C–rt, 5 h–16 h, 38% quant.; (c) Fe, NH₄Cl, EtOH, 90 °C, 2 h, 86–89%; (d) 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid, EDC, HOBT, DIPEA, DCM, rt, 18 h, 62% (over 2 steps); (e) Pd(OAc)₂, dppb, *N*-formylsaccharin, Na₂CO₃, Et₃SiH, DMF, 75 °C, 19 h, 3%; (f) B₂pin₂, PdCl₂(dppf)-DCM, KOAc, 1,4-dioxane, 95 °C, 19 h, 26–45%; (g) NaIO₄, NH₄OAc or HCl, acetone/H₂O or THF/H₂O, rt, 2 h–15 h; 45% quant. (h) Zn(CN)₂, Pd(PPh₃)₄, DMF, 150 °C, 22 h, 7%; (i) 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid, EDC-HCl, 4-DMAP, DMF, rt, 18 h–7 d, 40–69% (over 2 steps); (j) bromoacetyl bromide, DMF, rt, 30 min; then 4,6-dimethylpyrimidine-2-thiol, *t*-BuOK, DMF, rt, 19 h, 26% (over 3 steps); (k) CuCN, DMF, 150 °C, 22 h, 46%.

structures **24a** (here **10**)²³ and **SirReal2** (ref. 20) in this assay are in accordance with their published values. Furthermore, all **24a** derivatives showed potent and subtype selective inhibition of SIRT2 in the nanomolar range (Table 1). However, the introduction of the boronic acid, nitrile and aldehyde moieties in both positions R¹ and R² (**12–14**, **32–34**) showed no improvement, and to some extent a decrease in the potency compared to the lead structure **24a** (**10**). Similarly, halogen modifications with bromine at the R¹ position (**11**) showed no significant improvement in potency. On the contrary, we observed a significant increase in potency by the introduction of

halogens at the R² position (compounds **29–31**), highlighting the importance of the position of the functional group modifications. In particular, chlorobenzene derivative **29** displayed a 3-fold increase in potency, exhibiting an IC₅₀ value of 26 nM, but also the iodine (compound **31**, IC₅₀ = 29 nM) and bromine derivatives (compound **30**, IC₅₀ = 54 nM) were highly potent. These results lead to the following ranking in descending order of potency: Cl > I > Br. Although in theory the chlorine atom should form the weakest halogen bond with the ribose hydroxy groups of NAD⁺,³⁷ the observation that the chlorobenzene derivative **29**





Scheme 2 Reagents and conditions: (a) $\text{NH}_2\text{OH}\cdot\text{HCl}$, pyridine, $\text{EtOH}/\text{H}_2\text{O}$, rt, 24 h, quant.; (b) Ac_2O , H_2SO_4 , AcOH , $120\text{ }^\circ\text{C}$, 24 h, 25%. (c) NaNO_2 , HCl ; then $\text{FeCl}_3\cdot 6\text{H}_2\text{O}$, HCl , H_2O , $0\text{ }^\circ\text{C}$, 10 min; then $\text{CuCl}_2\cdot 2\text{H}_2\text{O}$, HCl , acetone/ EtOH , $0\text{ }^\circ\text{C}$, 10 min; then acrolein, acetone/ H_2O , rt, 4 h; then thiourea, EtOH , $80\text{ }^\circ\text{C}$, 30 h, 41% (over 4 steps); (d) Boc_2O , toluene, $100\text{ }^\circ\text{C}$, 4.5 h, 85%; (e) 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid, HATU, DIPEA, DMF, rt, 18 h, 29–40%; (f) 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid, EDC-HCl, 4-DMAP, DMF, rt, 16 h, 17–52%; (g) $\text{Zn}(\text{CN})_2$, $\text{Pd}(\text{PPh}_3)_4$, DMF, $80\text{ }^\circ\text{C}$, 18 h, 61%; (h) 4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)isoxazole, $\text{PdCl}_2(\text{dppf})\cdot\text{DCM}$, DMF, $90\text{ }^\circ\text{C}$, 20 h, 26% (i) B_2pin_2 , $\text{PdCl}_2(\text{dppf})\cdot\text{DCM}$, KOAc, 1,4-dioxane, $80\text{ }^\circ\text{C}$, 1 h; then rt, 2 h, 51%; (j) NaIO_4 , HCl , $\text{THF}/\text{H}_2\text{O}$, rt, 4 h, 84%; (k) TFA, CHCl_3 , rt, 17 h, quant.; (l) $\text{Pd}(\text{OAc})_2$, dppf, *N*-formylsaccharin, Na_2CO_3 , Et_3SiH , DMF, $75\text{ }^\circ\text{C}$, 19 h, 11%.

outperformed both bromobenzene and iodobenzene derivatives 30 and 31 suggests a much more complex mechanism than mere halogen bonding for the underlying increase in potency. Additional factors such as the atomic radius and thus steric requirements of these halogens may also play a significant role in the increase in potency, where the smaller chlorine atom may experience less steric effects upon binding at the active site compared to the larger bromine and iodine atoms. Halogen modification of **SirReal2** with bromine (compound 45) pointed similarly to an increase in potency. This is consistent with the published IC_{50} values of **SirReal2** and 7-bromo-**SirReal2** (45).²⁰ Although the introduction of boronic acid and aldehyde groups (compounds 48, 49) did not display any significant improvement in potency, the **SirReal2** derivative 46 bearing a nitrile group showed a 2-fold increase in SIRT2 inhibition with an IC_{50} value of 122 nM. In comparison, the homologous cyanomethyl derivative 47 only had similar potency to **SirReal2** (Table 1).

The determination of subtype selectivity against SIRT1, 3 and 5 was performed by measuring the residual enzyme activity after treatment with the corresponding inhibitor at a fixed concentration of 50 μM and subsequent

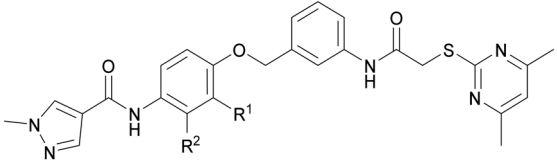
calculation of the percentage inhibition. All target compounds tested showed selectivity towards SIRT2, as corresponding inhibition values of SIRT1, 3 and 5 indicate IC_{50} values of over 50 μM with an exception for compound 33 that showed 71% inhibition at 50 μM . Nevertheless, considering the low IC_{50} value of 202 nM for the desired target SIRT2, selectivity is considered as sufficient.

Thermal stability of SIRT2–inhibitor complexes

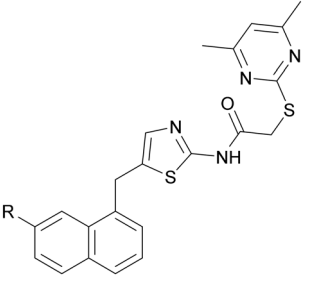
To investigate the potential role of the co-factor NAD^+ in the potency enhancement of the chloro-derivate 29 (**RW-78**) and nitrile 46 (**FM295**) compared to their corresponding lead structures 24a and **SirReal2**, we performed fluorescence thermal shift assays (Fig. 4). The thermal stability of SIRT2 was determined in the presence of 10 μM and 30 μM inhibitor and in the presence and absence of 2.5 mM co-factor NAD^+ . The presence of 30 μM of compound 29 resulted in a significant increase in the melting temperature of SIRT2 ($\Delta T = 6.5\text{ }^\circ\text{C}$) compared to its lead structure 24a ($\Delta T = 3\text{ }^\circ\text{C}$). However, results showed that the inhibitor-induced SIRT2 stabilization was



Table 1 *In vitro* inhibition of human SIRT1, 2, 3 and 5 by functionalised derivatives of **24a** and **SirReal2**. IC₅₀ values against SIRT2 are given as mean with standard deviations (*n* = 3). Inhibition percentages at 50 μM for SIRT1, 3 and 5 are given as mean without standard deviations (*n* = 2)



Compound ID	R ¹	R ²	IC ₅₀ (nM)		Inhibition of SIRT1/3/5 in %@50 μM		
			SIRT2	SIRT1	SIRT3	SIRT5	
10 (24a ^a)	H	H	79 ± 4	n.d.	n.d.	n.d.	
11	Br	H	81 ± 10	n.i.	24	13	
12	CHO	H	147 ± 13	2	24	16	
13	B(OH) ₂	H	294 ± 17	29	5	10	
14	CN	H	125 ± 18	6	35	2	
29	H	Cl	26 ± 2	12	26	7	
30	H	Br	54 ± 7	4	43	1	
31	H	I	29 ± 2	5	22	12	
32	H	CHO	184 ± 9	n.i.	7	6	
33	H	B(OH) ₂	202 ± 6	71	24	14	
34	H	CN	91 ± 7	23	31	4	



Compound ID	R	IC ₅₀ (nM)		Inhibition of SIRT1/3/5 in %@50 μM		
		SIRT2	SIRT1	SIRT3	SIRT5	
SirReal2 ^b	H	235 ± 10	n.d.	n.d.	n.d.	
45 ^c	Br	196 ± 17	n.d.	n.d.	n.d.	
46	CN	122 ± 8	6	9	8	
47	CH ₂ CN	235 ± 19	n.i.	15	2	
48	B(OH) ₂	235 ± 36	19	12	23	
49	CHO	356 ± 42	11	13	21	

^a Literature-known compound **24a** by Yang *et al.* with a published IC₅₀ of 0.82 μM for SIRT2.²³ ^b Literature-known compound **SirReal2** by Rumpf *et al.* with a published IC₅₀ of 0.44 μM for SIRT2.¹⁹ ^c Literature-known compound **45** by Schiedel *et al.* with a published IC₅₀ of 0.21 μM for SIRT2 (ref. 20).

independent from NAD⁺, contradicting our initial proposed binding mode. Similar results were obtained with the nitrile **46**.

Co-crystal structure of SIRT2 with compounds **29** (RW-78) and **31** (RW-80)

To clarify the binding mode of the most potent inhibitors we aimed for co-crystal structures with human SIRT2. To this end, we expressed human SIRT2 56-356 as a N-terminal His₆-SUMO fusion in *Escherichia coli* and purified the protein *via* affinity and size exclusion chromatography with tag removal in between (Fig. S1). Crystallization trials finally yielded an apo structure of SIRT2 (2.15 Å resolution, Table S1, PDB ID: 9S44) and two

complex structures with **29** (RW-78) (1.45 Å, Table S2, Fig. 5, PDB ID: 9S46) and **31** (RW-80, Fig. S2A) (1.45 Å, Table S3, PDB ID: 9S48) in the same space group.

Both **29** and **31** occupy a pocket located slightly below the NAD⁺-binding cleft, previously exploited by other potent and selective SIRT2 inhibitors like **SirReal2** and **24a** (Fig. 5A and S2A and B).^{19,23} Compared to our SIRT2 apo structure, the ligand bound structures adopt the “open-locked-state” reported first in the SIRT2:**SirReal2** co-crystal structure (PDB ID: 4RMG; root mean square deviation ≤ 0.223 Å over 218 C^α atoms).^{19,49} Both **29** and **31** are coordinated by the same protein residues and only differ by their halogen atom (Fig. 5B and S2B). In the binding pocket, the inhibitor is stabilized by multiple interactions previously described for the lead structure **24a** (PDB: 5YQO, Fig. S2B).²³ Most



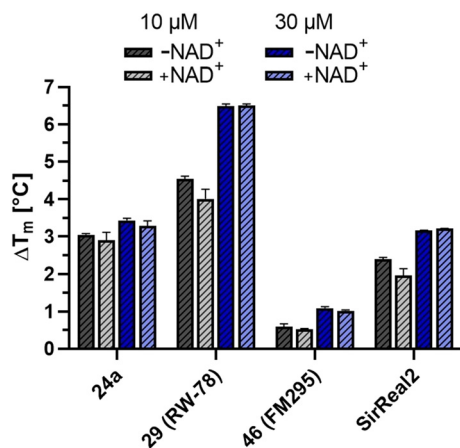


Fig. 4 Melting point shifts of SIRT2 with inhibitors at 10 μM and 30 μM in the presence and absence of 2.5 mM co-factor NAD^+ with no dependency on the co-factor observed. All melting temperature shifts measured were referenced to the melting temperature of SIRT2 without inhibitor, also in the presence and absence of the co-factor. Under both these conditions, the reference melting temperatures showed no significant change, each yielding a melting temperature of 53 $^{\circ}\text{C}$.

importantly, the amide next to the methyl pyrazole hydrogen bonds to the carbonyl oxygen of V^{233} and R^{97} . Additionally, the chlorine (**29**) or iodine (**31**) atom engages in halogen- π -interactions with F^{235} (Fig. 5B and S2A and B). The spacing between the chlorine atom of **29** and the centroid of the π -system of F^{235} (3.6 \AA) matches the average distance of $\text{C}-\text{Cl}\cdots\pi$ -bonds (3.854 \AA),⁵⁰ explaining the high potency of **29**. Contrary to our design, **29** and **31** do not engage in interactions with the SIRT2 co-factor NAD^+ but stabilize the enzyme in an inactive state. This state is established by rearrangements of residues near the active site (Fig. 5B and S2C). For instance, the catalytic histidine H^{187} is slightly shifted compared to SIRT2 apo structures (PDB ID: 1J8F,⁴⁹

PDB ID: 9S44). In addition, by interacting with the inhibitor, F^{96} and R^{97} move into the NAD^+ -binding pocket, displacing the co-factor and prohibiting NAD^+ -binding. Therefore, the determined structures do not show electron densities for the co-factor, although 5 mM NAD^+ was included in the crystallization screens. Instead of the co-factor, the NAD^+ -binding site contains many water molecules. Considering the high affinity of **29** and **31** (Table 1), the entropic barrier that is associated with the displacement of NAD^+ and the binding of water molecules must be counteracted by an enormous enthalpic stabilization of the inhibitors in the SIRT2 pocket. Notably, the NAD^+ displacement is supported by thermal shift assays (Fig. 4), which revealed that **29** increases the melting temperature of SIRT2 independently of NAD^+ .

Cellular target engagement in HEK293T cells via NanoBRET assay

Cellular SIRT2 target engagement of chloro-derivative **29** was assessed in HEK293T cells via NanoBRET assay that was developed by Vogelmann *et al.*²¹ The NanoBRET tracer pre-treated cells were incubated with increasing concentrations of chloro-derivative **29**, reflecting its intracellular binding to SIRT2. Compound **29** showed high target engagement with an EC_{50} value of 15 nM (Fig. 6), validating our data obtained from *in vitro* inhibition studies. For comparison, the highly potent triazole-based SirReal inhibitor Vogelmann_12 (SH10) was used as a reference and showed an EC_{50} value of 99 nM.

Conclusions

Based on the rationale to target the essential co-factor NAD^+ of SIRT2 via reversible covalent binding or halogen bonding with appropriate functional groups, a total of 14 potent and subtype selective SIRT2 inhibitors were synthesised. While lead structure modifications of SirReal2 and **24a** with polar,

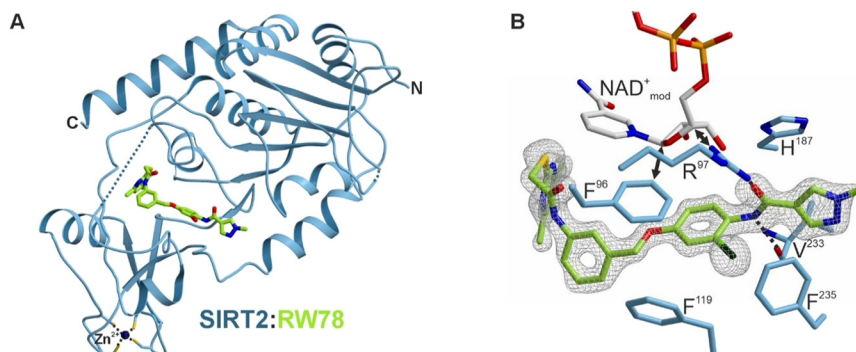


Fig. 5 (A) Co-crystal structure of human SIRT2 (blue) with its inhibitor **29** (RW-78, green) (PDB ID: 9S46). Residues 104–106 and 303 are disordered and indicated by blue dotted lines. The zinc ion bound to SIRT2 is shown as a dark blue sphere and coordinating cysteines are depicted as sticks. (B) Close-up view of **29** (green) bound to SIRT2. The experimental F_o-F_c omit electron density is shown as a gray mesh contoured to 3σ . Coordinating amino acid side chains (blue) are shown as sticks and labelled by the one-letter code. **29** is stabilized by hydrogen bonds to R^{97} and the V^{233} carbonyl oxygen (black dotted lines). The chlorine atom of **29** interacts with F^{235} via halogen- π -bonding. Binding of **29** induces movements of F^{96} and R^{97} into the NAD^+ -binding site (see also Fig. S2C). Modelling of the NAD^+ co-factor (extracted from the SIRT2:SirReal2 coordinates (PDB ID: 4RMG))¹⁹ into the SIRT2:RW-78 structure by superposition illustrates that F^{96} and R^{97} clash with NAD^+ (black double arrows) and hence displace the co-factor.



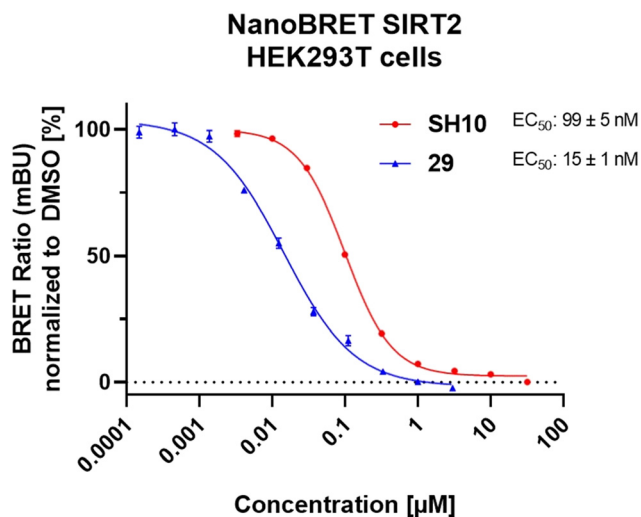


Fig. 6 NanoBRET-based cellular target engagement of the SIRT2 inhibitor **29**. Cellular binding of **29** was evaluated using a NanoBRET target engagement assay in live HEK293T cells. Cells were incubated with 2 μM NanoBRET tracer and treated with increasing concentrations **29** for 2 hours at 37 $^{\circ}\text{C}$. Binding for **SH10** is plotted as a reference. For compound **29**, a stable HEK293T cell line expressing NanoLuc-SIRT2 fusion construct was used, whereas for **SH10** HEK293T cells were transiently transfected with the NanoLuc-SIRT2 fusion construct.

potential reversible covalent binding warheads showed ambiguous results, nitrile **46** (**FM295**; $\text{IC}_{50} = 122 \text{ nM}$) demonstrated significant potency enhancement compared to its lead structure **SirReal2** ($\text{IC}_{50} = 235 \text{ nM}$). Among the halogenated derivatives of **24a** (**10**), compounds **30** ($\text{IC}_{50} = 54 \text{ nM}$) and **31** ($\text{IC}_{50} = 29 \text{ nM}$) outperformed the lead structure **24a** ($\text{IC}_{50} = 79 \text{ nM}$). In addition, ligand **29** (**RW-78**) emerged as the best member of the whole series and as one of the most potent SIRT2 inhibitors known to date ($\text{IC}_{50} = 26 \text{ nM}$) while maintaining high subtype selectivity. X-ray crystallographic data visualized that compound **29** (**RW-78**) undergoes halogen- π interactions with SIRT2 and induces structural changes that interfere with co-factor binding. These results are in agreement with thermal shift assays indicating that the inhibition of compound **29** (**RW-78**) is NAD^{+} independent. In addition, the utility of compound **29** (**RW-78**) in the cellular context was validated *via* NanoBRET assay in HEK293T cells that showed high target engagement to SIRT2 ($\text{EC}_{50} = 15 \text{ nM}$). In conclusion, our findings provide a valuable contribution to a deeper understanding of the structure-activity relationships of SIRT2 inhibitors and a foundation for further optimisation of SIRT2-selective inhibitors.

Experimental

Chemistry

Materials and instruments. All solvents and reagents were purchased from commercial sources and used without further purification. Standard vacuum line techniques were

applied. Reactions were monitored *via* thin layer silica gel chromatography (TLC) using polyester sheets POLYGRAM SIL G/UV254 coated with 0.2 mm silica gel (Macherey-Nagel). Plates were visualised using UV light (254 nm or 365 nm) or staining with KMnO_4 , CAM (ceric ammonium molybdate) or DNPH (dinitrophenylhydrazine). Products were purified by flash column chromatography (normal-phase silica gel chromatography) using SiO_2 60 (0.040–0.063 mm, 230–400 mesh ASTM) from Merck. NMR spectra were recorded with Avance III HD 400 MHz Bruker BioSpin and Avance III HD 500 MHz Bruker BioSpin ($^1\text{H-NMR}$: 400 MHz and 500 MHz, $^{13}\text{C-NMR}$: 101 MHz and 126 MHz) using the deuterated solvent stated. Chemical shifts (δ) are quoted in parts per million (ppm) and referenced to the residual solvent peak. Multiplicities are denoted as s-singlet, d-doublet, t-triplet, q-quartet and quin-quintet. Coupling constants J are given in Hz and round to the nearest 0.1 Hz. Infrared spectra were recorded from 4000 to 650 cm^{-1} on a PERKIN ELMER Spectrum BX-59343 FT-IR instrument. A Smiths Detection DuraSamp IR II Diamond ATR sensor was used for detection. The absorption bands are reported in wavenumbers [cm^{-1}]. High resolution mass spectra (HR-MS) were recorded using a Jeol Mstation 700 or JMS GCmate II Jeol instrument for electron impact ionisation (EI). Thermo Finnigan LTQ was used for electrospray ionisation (ESI). Melting points were measured with a Büchi Schmelzpunktapparat B-540. HPLC analytical measurements at 210 nm and 254 nm for purities determination was performed with the following methods:

Method 1:

- Zorbax SB C18 3.5 μm (4.6 \times 100 mm), injection volume 5 μL .
 a) MeCN/water (35 : 65); flow rate 1.0 mL min^{-1} ; temp. 35 $^{\circ}\text{C}$.
 b) MeCN/water (50 : 50); flow rate 1.2 mL min^{-1} ; temp. 35 $^{\circ}\text{C}$.
 c) MeCN/water (50 : 50); flow rate 1.2 mL min^{-1} ; temp. 50 $^{\circ}\text{C}$.

Method 2:

- Raptor C18 5 μm (4.6 \times 150 mm), injection volume 5 μL .
 a) MeCN/water (35 : 65); flow rate 0.7 mL min^{-1} ; temp. 35 $^{\circ}\text{C}$.

Method 3:

- Zorbax Eclipse Plus $^{\circ}$ C18 5 μm (4.6 \times 150 mm), injection volume 5 μL .
 a) MeCN/water (70 : 30); flow rate 1.0–1.5 mL min^{-1} ; temp. 30–50 $^{\circ}\text{C}$.
 b) MeCN/water (50 : 50); flow rate 1.0–1.5 mL min^{-1} ; temp. 30–50 $^{\circ}\text{C}$.
 c) MeCN/phosphate buffer pH = 5 (70 : 30); flow rate 1.0–1.2 mL min^{-1} ; temp. 30–50 $^{\circ}\text{C}$.

Synthetic procedures

General procedure A – amide coupling (I) with HATU and DIPEA. To a stirred solution of the appropriate carboxylic acid (1.0 equivalent) in THF with a concentration of 0.20 M were added DIPEA (3.0 equivalents) and HATU (1.5 equivalents). The reaction mixture was stirred at room temperature for 1 h. The appropriate amine (1.0 equivalent) was then added and unless stated otherwise, the reaction mixture was further stirred at room temperature for 3 h.



DCM and water were then added, the resulting two phases were separated, and the aqueous phase was extracted with DCM (3×). The combined organic phases were dried using a phase separation paper and the solvent was removed *in vacuo*. Unless stated otherwise, the crude product was then purified by FCC using the indicated eluent.

General procedure B – amide coupling (II) with EDC, HOBT and DIPEA. To a stirred solution of the appropriate amine (1.0 equivalent) in DCM with a concentration of 0.10 M were added HOBT (1.5 equivalents), EDC (2.0 equivalents), DIPEA (2.0 equivalents) and the appropriate carboxylic acid (2.0 equivalents). The reaction mixture was stirred at room temperature for 18 h. DCM and water were then added, the resulting two phases were separated, and the aqueous phase was extracted with DCM (3×). The combined organic phases were dried using a phase separation paper and the solvent was removed *in vacuo*. The crude product was then purified by FCC using the indicated eluent.

General procedure C – amide coupling (III) with EDC·HCl and 4-DMAP. To a stirred solution of the appropriate carboxylic acid (2.0 equivalents) in DMF with a concentration of 0.40 M were added EDC·HCl (2.0 equivalents) and 4-DMAP (2.0 equivalents). The reaction mixture was stirred at room temperature for 15 minutes. Afterwards, the appropriate amine (1.0 equivalent) was added and the solution was stirred at the stated temperature for the stated time. The solution was then diluted with EtOAc and the organic phase was washed with brine (3×). The organic phase was dried using a phase separation paper and the solvent was removed *in vacuo*. The crude product was then purified by FCC using the indicated eluent.

General procedure D – Williamson ether synthesis. To a stirred solution of the appropriate phenol (1.0 equivalent) in DMF with a concentration of 0.10 M was added K₂CO₃ (3.0 equivalents). The reaction mixture was stirred at room temperature for 30 minutes. Afterwards, the appropriate alkyl halide (1.2 equivalents) was added and the mixture was stirred at room temperature for 16 h. The solution was then diluted with EtOAc and the organic phase was washed with brine (3×). The organic phase was dried using a phase separation paper and the solvent was removed *in vacuo*. The crude product was then purified by the stated procedure.

General procedure E – nitrobenzene reduction. To a stirred solution of the appropriate nitrobenzene derivative (1.0 equivalent) in EtOH with a concentration of 0.010 M were added iron powder (5.0 equivalents) and 0.30 M aq. NH₄Cl (5.0 equivalents) at 50 °C. The reaction mixture was then refluxed at 90 °C for 2 h. Afterwards, the iron powder was filtered off from the still hot mixture, and the filtrate was concentrated *in vacuo*. Unless stated otherwise, the crude product was used for the next step without further purification.

N-(4-Hydroxyphenyl)-1-methyl-1H-pyrazole-4-carboxamide (3). Prepared according to General procedure A from 1-methyl-1H-pyrazole-4-carboxylic acid (500 mg, 3.96 mmol) and aminophenol 1 (433 mg, 3.96 mmol). The crude product

was purified by FCC (DCM/MeOH 96:4) to give 3 (179 mg, 3.96 mmol, 21%) as a white solid; m.p. 225 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3347, 3020, 1641, 1602, 1558, 1530, 1510, 1430, 1351, 1202, 1099, 1006, 860, 813, 749; δ_{H} (400 MHz; (CD₃)₂SO) 9.57 (s, 1H, CONH), 9.18 (s, 1H, OH), 8.24 (s, 1H, 5-H), 7.96 (s, 1H, 3-H), 7.47–7.41 (m, 2H, 2'-H and 6'-H), 6.74–6.68 (m, 2H, 3'-H and 5'-H), 3.87 (s, 3H, CH₃). δ_{C} (101 MHz; (CD₃)₂SO) 160.00 (CONH), 153.40 (C-4'), 138.62 (C-3), 132.34 (C-5), 130.63 (C-1'), 121.92 (C-2' and C-6'), 118.72 (C-4), 114.96 (C-3' and C-5'), 38.79 (CH₃); HRMS (ESI): calcd.: 218.0924; found: $\{m/z: [M + H]^+\}$ 218.0928.

N-(3-Bromo-4-hydroxyphenyl)-1-methyl-1H-pyrazole-4-carboxamide (4). Prepared according to General procedure A from 1-methyl-1H-pyrazole-4-carboxylic acid (500 mg, 3.96 mmol) and aminophenol 2 (761 mg, 3.96 mmol). The crude product was purified by FCC (DCM/MeOH 98:2) to give amide 4 (649 mg, 2.19 mmol, 55%) as a light pink solid; m.p. 218 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3108, 1637, 1596, 1558, 1410, 1268, 1227, 1197, 1008, 852, 801, 753, 657; δ_{H} (400 MHz; (CD₃)₂SO) 9.98 (s, 1H, OH), 9.69 (s, 1H, CONH), 8.25 (s, 1H, 5-H), 7.96 (s, 1H, 3-H), 7.89 (d, *J* = 2.5 Hz, 1H, 2'-H), 7.45 (dd, *J* = 8.8, 2.5 Hz, 1H, 6'-H), 6.90 (d, 1H, 5'-H), 3.88 (s, 3H, CH₃). δ_{C} (101 MHz; (CD₃)₂SO) 160.14 (CONH), 150.02 (C-4'), 138.65 (C-3), 132.50 (C-5), 131.83 (C-1'), 124.45 (C-2'), 120.73 (C-6'), 118.37 (C-4), 116.01 (C-5'), 108.52 (C-3'), 38.83 (CH₃); HRMS (ESI): calcd.: 296.0029; found: $\{m/z: [M + H]^+\}$ 296.0033.

1-Methyl-N-(4-((3-nitrobenzyl)oxy)phenyl)-1H-pyrazole-4-carboxamide (5). Prepared according to General procedure D from phenol 3 (165 mg, 0.760 mmol) and 1-(bromomethyl)-3-nitrobenzene (197 mg, 0.911 mmol). The crude product was purified by FCC (DCM/MeOH 95:5) to give ether 5 (221 mg, 0.626 mmol, 82%) as a yellow solid. Analytical data are in alignment with literature.²³

N-(3-Bromo-4-((3-nitrobenzyl)oxy)phenyl)-1-methyl-1H-pyrazole-4-carboxamide (6). Prepared according to General procedure D from phenol 4 (3.10 g, 10.5 mmol) and 1-(bromomethyl)-3-nitrobenzene (2.71 g, 12.6 mmol). The crude product was resuspended in hexanes, vacuum filtered, and the residue washed sequentially with hexanes (2×), EtOAc (2×) and DCM (1×). The filtrate was discarded and the solid residue collected to give ether 6 (4.52 g, 10.5 mmol, quant.) as a beige solid; m.p. 226 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3394, 3114, 1649, 1593, 1550, 1520, 1499, 1397, 1357, 1275, 1229, 1057, 1001, 881, 853, 790, 728.; δ_{H} (400 MHz; (CD₃)₂SO) 9.82 (s, 1H, CONH), 8.38–8.36 (m, 1H, 2''-H), 8.27 (s, 1H, 5-H), 8.22–8.19 (m, 1H, 4''-H), 8.04 (d, *J* = 2.5 Hz, 1H, 2'-H), 8.00–7.97 (m, 1H, 3-H), 7.96–7.91 (m, 1H, 6''-H), 7.73 (t, *J* = 7.9 Hz, 1H, 5''-H), 7.65 (dd, *J* = 8.9, 2.6 Hz, 1H, 6'-H), 7.22 (d, *J* = 9.0 Hz, 1H, 5'-H), 5.34 (s, 2H, CH₂), 3.89 (s, 3H, CH₃). δ_{C} (101 MHz; (CD₃)₂SO) 160.31 (CONH), 149.96 (C-4'), 147.86 (C-3''), 139.16 (C-1''), 138.71 (C-3), 133.87 (C-1'), 133.74 (C-6''), 132.63 (C-5), 130.10 (C-5''), 124.42 (C-2'), 122.78 (C-4''), 121.78 (C-2''), 120.29 (C-6'), 118.20 (C-4), 114.43 (C-5'), 110.77 (C-3'), 69.05 (CH₂), 38.85 (CH₃); HRMS (ESI): calcd.: 431.0349; found: $\{m/z: [M + H]^+\}$ 431.0355.



1-Methyl-N-(4-((3-nitrobenzyl)oxy)-3-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)phenyl)-1H-pyrazole-4-carboxamide (7). PdCl₂(dppf)·DCM (93.3 mg, 0.128 mmol), bis(pinacolato)diboron (648 mg, 2.55 mmol), bromo derivative **6** (550 mg, 1.28 mmol) and KOAc (501 mg, 5.10 mmol) were dissolved in degassed 1,4-dioxane (40 mL) under N₂ atmosphere and stirred at 95 °C for 19 h. The reaction mixture was cooled to room temperature and then filtered. The filtrate was then diluted with EtOAc (50 mL). Water (50 mL) was added, the resulting two phases were separated, and the aqueous phase was extracted with EtOAc (3 × 60 mL). The combined organic phases were then dried using a phase separation paper and concentrated *in vacuo*. The crude product was purified by FCC (hexanes/EtOAc 20:80) to give boronic acid pinacol ester **7** (277 mg, 0.579 mmol, 45%) as a yellow solid; m.p. 254 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3339, 2982, 1640, 1557, 1530, 1495, 1344, 1318, 1239, 1140, 1068, 1008, 812, 731; δ_{H} (400 MHz; (CD₃)₂SO) 9.75 (s, 1H, CONH), 8.55 (s, 1H, 2''-H), 8.28 (s, 1H, 5-H), 8.21–8.16 (m, 1H, 4''-H), 8.03–7.99 (m, 2H, 3-H and 6''-H), 7.97 (dd, *J* = 9.0, 2.7 Hz, 1H, 6'-H), 7.83 (d, *J* = 2.8 Hz, 1H, 2'-H), 7.71 (t, *J* = 8.0 Hz, 1H, 5''-H), 7.09 (d, *J* = 9.0 Hz, 1H, 5'-H), 5.25 (s, 2H, CH₂), 3.89 (s, 3H, NCH₃), 1.33 (s, 12H, (CH₃)₄). δ_{C} (101 MHz; (CD₃)₂SO) 160.14 (CONH), 158.47 (C-4'), 147.97 (C-3''), 140.25 (C-1''), 138.72 (C-3), 133.04 (C-6''), 132.48 (C-5 and C-1'), 129.59 (C-5''), 128.41 (C-2'), 124.72 (C-6'), 122.23 (C-4''), 120.94 (C-2''), 118.52 (C-4), 117.15 (C-3'), 112.48 (C-5'), 83.30 ((C(CH₃)₂)₂), 68.07 (CH₂), 38.83 (NCH₃), 24.63 ((CH₃)₄); HRMS (ESI): calcd.: 479.2096; found: $\{m/z: [M + H]^+\}$ 479.2114.

(5-(1-Methyl-1H-pyrazole-4-carboxamido)-2-((3-nitrobenzyl)oxy)phenyl)boronic acid (8). To a stirred solution of boronic acid pinacol ester **7** (80.0 mg, 0.167 mmol) in acetone (20 mL) were added sodium periodate (107 mg, 0.502 mmol), ammonium acetate (38.7 mg, 0.502 mmol) and water (7 mL). The reaction mixture was stirred vigorously at room temperature for 2 h. Afterwards, acetone was removed *in vacuo* and the reaction mixture was diluted with DCM (20 mL). Water (25 mL) was added, the resulting two phases were separated, and the aqueous phase was extracted with DCM (3 × 25 mL). The combined organic phases were then dried with Na₂SO₄ and then concentrated *in vacuo*. The crude product was then purified by FCC (DCM/MeOH 97:3) to give boronic acid **8** (30.0 mg, 75.7 μmol , 45%) as an off-white solid; m.p. 190 °C (decomposition); IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3292, 2919, 1638, 1557, 1528, 1492, 1417, 1351, 1316, 1227, 1157, 1008, 861, 808, 759, 733, 673; δ_{H} (400 MHz; (CD₃)₂SO) 9.68 (s, 1H, CONH), 8.40–8.38 (m, 1H, 2'-H), 8.27 (s, 1H, 5''-H), 8.21–8.17 (m, 1H, 4'-H), 7.99 (s, 1H, 3''-H), 7.95 (d, *J* = 7.4 Hz, 1H, 6'-H), 7.86 (s, 2H, B(OH)₂), 7.81–7.76 (m, 1H, 4-H), 7.74–7.70 (m, 2H, 6-H and 5'-H), 7.01 (d, *J* = 8.9 Hz, 1H, 3-H), 5.30 (s, 2H, CH₂), 3.88 (s, 3H, CH₃). δ_{C} (101 MHz; (CD₃)₂SO) 160.11 (CONH), 157.64 (C-2), 147.88 (C-3'), 139.80 (C-1'), 138.70 (C-3''), 133.96 (C-6'), 132.44 (C-5 and C-5''), 129.99 (C-5'), 127.17 (C-6), 123.02 (C-4), 122.65 (C-4'), 122.04 (C-2'), 118.59 (C-4''), 111.88 (C-3), 68.30 (CH₂), 38.81 (CH₃); HRMS (ESI): calcd.: 419.1133; found: $\{m/z: [M + Na]^+\}$ 419.1140.

N-(3-Cyano-4-((3-nitrobenzyl)oxy)phenyl)-1-methyl-1H-pyrazole-4-carboxamide (9). Bromo derivative **6** (200 mg, 0.464 mmol), Pd(PPh₃)₄ (53.6 mg, 46.4 μmol) and Zn(CN)₂ (32.7 mg, 0.278 mmol) were dissolved in dry DMF (1 mL) under N₂ atmosphere and stirred at 150 °C for 22 h. The reaction mixture was cooled to room temperature and then diluted with EtOAc (25 mL). The organic phase was washed with brine (4 × 30 mL) and then dried with Na₂SO₄ and concentrated *in vacuo*. The crude product was purified by FCC (DCM/MeOH 98:2) to give nitrile **9** (11.6 mg, 30.7 μmol , 7%) as a beige solid; m.p. 251 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3397, 2226, 1668, 1593, 1553, 1528, 1510, 1351, 1319, 1287, 1247, 1207, 1011, 870, 861, 814, 748, 731; δ_{H} (400 MHz; (CD₃)₂SO) 9.97 (s, 1H, CONH), 8.37 (s, 1H, 2''-H), 8.29 (s, 1H, 5-H), 8.23 (d, *J* = 8.3 Hz, 1H, 4''-H), 8.08 (s, 1H, 2'-H), 7.99 (s, 1H, 3-H), 7.95–7.88 (m, 2H, 6'-H and 6''-H), 7.75 (t, *J* = 7.9 Hz, 1H, 5''-H), 7.37 (d, *J* = 9.2 Hz, 1H, 5'-H), 5.43 (s, 2H, CH₂), 3.89 (s, 3H, CH₃). δ_{C} (101 MHz; (CD₃)₂SO) 160.50 (CONH), 155.32 (C-4'), 147.89 (C-3''), 138.75 (C-3), 138.51 (C-1''), 133.96 (C-6''), 133.11 (C-1'), 132.75 (C-5), 130.24 (C-5''), 126.68 (C-6'), 124.31 (C-2'), 123.04 (C-4''), 122.06 (C-2''), 117.98 (C-4), 116.15 (CN), 114.17 (C-5'), 100.54 (C-3'), 69.01 (CH₂), 38.87 (CH₃); HRMS (ESI): calcd.: 378.1197; found: $\{m/z: [M + H]^+\}$ 378.1195.

N-(4-((3-(2-((4,6-Dimethylpyrimidin-2-yl)thio)acetamido)benzyl)oxy)phenyl)-1-methyl-1H-pyrazole-4-carboxamide (10). Prepared according to literature.²³

N-(3-Bromo-4-((3-(2-((4,6-dimethylpyrimidin-2-yl)thio)acetamido)benzyl)oxy)phenyl)-1-methyl-1H-pyrazole-4-carboxamide (11). Prepared according to General procedure E from nitrobenzene **6** (1.00 g, 2.32 mmol). The obtained crude amine was reacted with 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid (924 mg, 4.66 mmol) according to General procedure C. The reaction mixture was stirred at room temperature for 4 days. The crude product was purified by FCC (DCM/MeOH 98:2) to give **11** (610 mg, 1.05 mmol, 45% over two steps) as a pale-yellow solid; m.p. 96 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3284, 1646, 1582, 1551, 1526, 1490, 1441, 1339, 1264, 1223, 1048, 1005, 873, 755, 690; δ_{H} (400 MHz; (CD₃)₂SO) 10.29 (s, 1H, 3''-NHCO), 9.79 (s, 1H, 4-CONH), 8.27 (s, 1H, 5-H), 8.02 (d, *J* = 2.5 Hz, 1H, 2'-H), 7.98 (s, 1H, 3-H), 7.69 (t, *J* = 1.9 Hz, 1H, 2''-H), 7.60 (dd, *J* = 9.0, 2.5 Hz, 1H, 6'-H), 7.57–7.53 (m, 1H, 4''-H), 7.34 (t, *J* = 7.8 Hz, 1H, 5''-H), 7.18–7.17 (m, 1H, 5'-H or 6''-H), 7.16–7.14 (m, 1H, 5'-H or 6''-H), 6.96 (s, 1H, 5''-H), 5.15 (s, 2H, OCH₂), 4.04 (s, 2H, SCH₂), 3.89 (s, 3H, NCH₃), 2.32 (s, 6H, 4''-CH₃ and 6''-CH₃). δ_{C} (101 MHz; (CD₃)₂SO) 169.32 (C-2'''), 166.95 (C-4''' and C-6'''), 166.62 (3''-NHCO), 160.27 (4-CONH), 150.34 (C-4'), 139.19 (C-3''), 138.71 (C-3), 137.39 (C-1''), 133.53 (C-1'), 132.61 (C-5), 128.89 (C-5''), 124.41 (C-2'), 122.26 (C-6''), 120.22 (C-6'), 118.65 (C-4''), 118.24 (C-4), 117.95 (C-2''), 116.05 (C-5''), 114.29 (C-5'), 110.68 (C-3'), 70.27 (OCH₂), 38.84 (NCH₃), 35.47 (SCH₂), 23.33 (4''-CH₃ and 6''-CH₃); HRMS (ESI): calcd.: 581.0965; found: $\{m/z: [M + H]^+\}$ 581.0973; purity (HPLC): 210 nm: >95%; 254 nm: >95% (method 1a).



***N*-(4-((3-(2-((4,6-Dimethylpyrimidin-2-yl)thio)acetamido)benzyl)oxy)-3-formylphenyl)-1-methyl-1*H*-pyrazole-4-carboxamide (12).** Bromoarene **11** (250 mg, 0.430 mmol), Pd(OAc)₂ (2.90 mg, 12.9 μmol, 3 mol%), dppb (8.25 mg, 19.3 μmol, 4.5 mol%), *N*-formylsaccharin (287 mg, 1.29 mmol) and Na₂CO₃ (137 mg, 1.29 mmol) were added to a 30 mL glass tube, which was then evacuated and backfilled three times with N₂. A degassed solution of Et₃SiH (90.3 μL, 0.560 mmol) in DMF (1 mL) was added to the glass tube under N₂ atmosphere. The mixture was stirred for 15 min at room temperature and subsequently warmed to 75 °C and stirred for another 19 h. The reaction mixture was cooled to room temperature, then diluted with EtOAc (15 mL) and washed with brine (3 × 15 mL). The organic phase was dried with a phase separation paper and concentrated *in vacuo*. The crude product was redissolved in MeOH (5 mL), sat. aq. NaHSO₃ (25 mL) was added, stirred for approximately 30 s, diluted with H₂O (25 mL), and then extracted with EtOAc (3 × 25 mL). The collected aqueous phase was basified with 50% NaOH (10 mL) and extracted with DCM (3 × 25 mL). The combined organic layers were dried with Na₂SO₄ and concentrated *in vacuo*. The obtained crude residue was purified by FCC (DCM/MeOH 98:2) to give **13** (7.0 mg, 13 μmol, 3%) as an off-white solid; m.p. 92 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3284, 2921, 2852, 1667, 1582, 1553, 1532, 1442, 1308, 1264, 1223, 1168, 1005, 873, 755, 690; δ_{H} (400 MHz; (CD₃)₂SO) 10.48 (s, 1H, CHO), 9.59 (s, 1H, 3''-NHCO), 7.97 (dd, *J* = 9.0, 2.9 Hz, 1H, 6''-H), 7.90 (s, 1H, 5''-H), 7.83–7.80 (m, 2H, 3-H and 2''-H), 7.63 (t, 1H, 2''-H), 7.59 (s, 1H, 4-CONH), 7.38 (dt, 1H, 4''-H), 7.33 (t, *J* = 7.7 Hz, 1H, 5''-H), 7.17 (dt, *J* = 7.5 Hz, 1H, 6''-H), 7.05 (d, *J* = 9.0 Hz, 1H, 5''-H), 6.85 (s, 1H, 5'''-H), 5.18 (s, 2H, OCH₂), 3.92 (s, 3H, NCH₃), 3.85 (s, 2H, SCH₂), 2.46 (s, 6H, 4'''-CH₃ and 6'''-CH₃). δ_{C} (101 MHz; (CD₃)₂SO) 189.29 (CHO), 170.40 (C-2'''), 168.40 (3''-NHCO), 168.11 (C-4''' and C-6'''), 160.89 (4-CONH), 157.95 (C-4'), 139.17 (C-3'''), 138.30 (C-3), 137.70 (C-1''), 132.25 (C-1' or C-3'), 132.23 (C-5), 129.66 (C-5''), 128.70 (C-6'), 125.49 (C-1' or C-3'), 123.13 (C-6''), 119.83 (C-2'), 119.61 (C-4''), 119.04 (C-4), 118.68 (C-2''), 117.09 (C-5'''), 114.35 (C-5'), 70.98 (OCH₂), 39.71 (NCH₃), 35.90 (SCH₂), 24.09 (4'''-CH₃ and 6'''-CH₃); HRMS (ESI): calcd.: 529.1658; found: {*m/z*: [M-H]⁻ 529.1665}; purity (HPLC): 210 nm: >95%; 254 nm: >95% (method 1a).

2-((3-(2-((4,6-Dimethylpyrimidin-2-yl)thio)acetamido)benzyl)oxy)-5-(1-methyl-1*H*-pyrazole-4-carboxamido)phenylboronic acid (13). Prepared according to General procedure E from nitrobenzene **8** (15.0 mg, 37.9 μmol). The obtained crude amine was reacted with 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid (15.2 mg, 76.5 μmol) according to General procedure B. The crude product was purified by FCC (DCM/MeOH 96:4) to give **13** (13.0 mg, 23.8 μmol, 62% over two steps) as a white solid; m.p. 193 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3435, 3325, 1683, 1625, 1581, 1550, 1530, 1495, 1443, 1374, 1322, 1265, 1225, 1166, 1043, 1017, 885, 815, 750; δ_{H} (400 MHz; (CD₃)₂SO) 10.30 (s, 1H, 3''-NHCO), 9.69 (s, 1H, 4'-CONH), 8.27 (s, 1H, 5'-H), 7.99 (s, 1H, 3'-H), 7.83–7.76 (m, 2H, 4-H and 6-H), 7.74 (s, 2H, B(OH)₂), 7.67 (s, 1H, 2''-H), 7.55 (d, *J* = 8.1

Hz, 1H, 4''-H), 7.34 (t, *J* = 7.8 Hz, 1H, 5''-H), 7.18 (d, *J* = 7.6 Hz, 1H, 6''-H), 7.01 (d, *J* = 8.9 Hz, 1H, 3-H), 6.95 (s, 1H, 5'''-H), 5.13 (s, 2H, OCH₂), 4.04 (s, 2H, SCH₂), 3.88 (s, 3H, NCH₃), 2.32 (s, 6H, 4'''-CH₃ and 6'''-CH₃). δ_{C} (101 MHz; (CD₃)₂SO) 169.31 (C-2'''), 166.96 (C-4''' and C-6'''), 166.62 (3''-NHCO), 160.12 (4'-CONH), 158.53 (C-2), 139.23 (C-3''), 138.71 (C-3'), 137.76 (C-1''), 132.44 (C-5), 132.33 (C-5'), 129.01 (C-5''), 127.56 (C-4 or C-6), 123.46 (C-4 or C-6), 122.53 (C-6''), 121.82 (C-1), 118.68 (C-4'), 118.61 (C-4''), 118.13 (C-2''), 116.06 (C-5'''), 111.89 (C-5''), 69.79 (OCH₂), 38.82 (NCH₃), 35.47 (SCH₂), 23.33 (4'''-CH₃ and 6'''-CH₃); HRMS (ESI): calcd.: 545.1784; found: {*m/z*: [M-H]⁻ 545.1788}; purity (HPLC): 210 nm: 92%; 254 nm: >95% (method 1a).

***N*-(3-Cyano-4-((3-(2-((4,6-dimethylpyrimidin-2-yl)thio)acetamido)benzyl)oxy)phenyl)-1-methyl-1*H*-pyrazole-4-carboxamide (14).** Prepared according to General procedure E from nitrobenzene **9** (90.0 mg, 0.238 mmol). The obtained crude amine was reacted with 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid (100 mg, 0.507 mmol) according to General procedure B. The crude product was purified by FCC (DCM/MeOH 96:4) to give **14** (82.5 mg, 0.156 mmol, 62% over two steps) as a yellow solid; m.p. 165 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3353, 2227, 1661, 1587, 1555, 1535, 1503, 1446, 1410, 1263, 1232, 1004, 860, 817, 752, 688; δ_{H} (400 MHz; (CD₃)₂SO) 10.31 (s, 1H, 3''-NHCO), 9.94 (s, 1H, 4-CONH), 8.29 (s, 1H, 5-H), 8.07 (s, 1H, 2'-H), 7.99 (s, 1H, 3-H), 7.85 (d, *J* = 9.1 Hz, 1H, 6'-H), 7.67 (s, 1H, 2''-H), 7.57 (d, *J* = 8.1 Hz, 1H, 4''-H), 7.39–7.33 (m, 1H, 5''-H), 7.33–7.28 (m, 1H, 5'-H), 7.17 (d, *J* = 7.6 Hz, 1H, 6''-H), 6.94 (s, 1H, 5'''-H), 5.25 (s, 2H, OCH₂), 4.04 (s, 2H, SCH₂), 3.89 (s, 3H, NCH₃), 2.31 (s, 6H, 4'''-CH₃ and 6'''-CH₃). δ_{C} (101 MHz; (CD₃)₂SO) 169.31 (C-2'''), 166.95 (C-4''' and C-6'''), 166.66 (3''-NHCO), 160.47 (4-CONH), 155.68 (C-4'), 139.28 (C-3''), 138.74 (C-3), 136.78 (C-1''), 132.78 (C-1'), 132.72 (C-5), 129.03 (C-5''), 126.61 (C-6'), 124.30 (C-2'), 122.43 (C-6''), 118.87 (C-4''), 118.02 (C-2'), 116.28 (CN), 116.04 (C-5'''), 114.08 (C-5'), 100.36 (C-3'), 70.20 (OCH₂), 38.86 (NCH₃), 35.48 (SCH₂), 23.32 (4'''-CH₃ and 6'''-CH₃); HRMS (ESI): calcd.: 550.1637; found: {*m/z*: [M + Na]⁺ 550.1633}; purity (HPLC): 210 nm: >95%; 254 nm: >95% (method 1a).

4-Amino-3-bromophenol (18). Prepared according to General procedure E from nitroarene **15** (5.00 g, 22.9 mmol). The crude product was purified by FCC (DCM/MeOH 97:3) to give 4-amino-3-bromophenol (**18**) (3.85 g, 20.5 mmol, 89%) as a light pink solid. Analytical data are in alignment with literature.⁵¹

4-Amino-3-iodophenol (19). Prepared according to General procedure E from nitroarene **16** (1.00 g, 3.66 mmol). The crude product was purified by FCC (DCM/MeOH 97:3) to give aniline **19** (738 mg, 3.14 mmol, 86%) as a grey-white solid. Analytical data are in alignment with literature.⁵¹

2-Chloro-4-((3-nitrobenzyl)oxy)aniline (20). To a stirred solution of phenol **17** (500 mg, 3.41 mmol) in DMF (25 mL) was added K₂CO₃ (1.42 g, 10.2 mmol) and the reaction mixture was stirred for 30 minutes. 1-(Bromomethyl)-3-nitrobenzene (885 mg, 4.10 mmol) was then added and the reaction mixture was stirred at 0 °C for another 5 h. The



reaction mixture was diluted with EtOAc and washed with brine (3 × 200 mL). The organic phase was dried using a phase separation paper and concentrated *in vacuo*. The crude product was purified by FCC (DCM) to give ether **20** (475 mg, 1.70 mmol, 50%) as an orange solid; m.p. 102 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3443, 3361, 3090, 2866, 1607, 1572, 1527, 1500, 1480, 1467, 1387, 1350, 1231, 1093, 1041, 901, 887, 823, 808, 796, 732; δ_{H} (400 MHz; $(\text{CD}_3)_2\text{SO}$) 8.27 (dt, $J = 2.3, 0.6$ Hz, 1H; 2'-H), 8.18 (ddd, $J = 8.2, 2.4, 1.0$ Hz, 1H, 4'-H), 7.87 (ddd, $J = 7.7, 1.7, 1.0$ Hz, 1H, 6'-H), 7.69 (t, $J = 7.9$ Hz, 1H, 5'-H), 6.97 (d, $J = 2.7$ Hz, 1H, 3-H), 6.81 (dd, $J = 8.8, 2.8$ Hz, 1H, 5-H), 6.75 (d, $J = 8.8$ Hz, 1H, 6-H), 5.15 (d, $J = 0.8$ Hz, 2H, CH_2), 4.91 (s, 2H, NH_2). δ_{C} (101 MHz; $(\text{CD}_3)_2\text{SO}$) 149.24 (C-4), 147.81 (C-3'), 139.75 (C-1'), 139.16 (C-1), 133.99 (C-6'), 130.00 (C-5'), 122.62 (C-4'), 121.88 (C-2'), 117.28 (C-2), 116.21 (C-6), 115.63 (C-3 or C-5), 115.53 (C-3 or C-5), 68.72 (CH_2); HRMS (ESI): calcd.: 279.0531; found: $\{m/z: [\text{M} + \text{H}]^+\}$ 279.0532}.

2-Bromo-4-((3-nitrobenzyl)oxy)aniline (21). Prepared according to General procedure D from phenol **18** (3.85 g, 20.5 mmol) and 1-(bromomethyl)-3-nitrobenzene (5.31 g, 24.6 mmol). The crude product was purified by FCC (DCM + 0.5% MeOH) to give ether **21** (3.15 g, 9.75 mmol, 48%) as an orange solid; m.p. 84 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3417, 3340, 3073, 1707, 1601, 1581, 1525, 1499, 1387, 1343, 1318, 1228, 1094, 1050, 1030, 811, 729, 667; δ_{H} (400 MHz; $(\text{CD}_3)_2\text{SO}$) 8.28–8.25 (m, 1H, 2'-H), 8.18 (ddd, $J = 8.2, 2.4, 1.0$ Hz, 1H, 4'-H), 7.87 (ddd, 1H, 6'-H), 7.69 (t, $J = 7.9$ Hz, 1H, 5'-H), 7.11 (d, $J = 2.8$ Hz, 1H, 3-H), 6.85 (dd, $J = 8.8, 2.8$ Hz, 1H, 5-H), 6.76 (d, $J = 8.8$ Hz, 1H, 6-H), 5.15 (s, 2H, CH_2), 4.88 (s, 2H, NH_2). δ_{C} (101 MHz; $(\text{CD}_3)_2\text{SO}$) 149.41 (C-4), 147.81 (C-3'), 140.34 (C-1), 139.74 (C-1'), 133.99 (C-6'), 130.00 (C-5'), 122.62 (C-4'), 121.88 (C-2'), 118.37 (C-3), 116.30 (C-5 or C-6), 116.10 (C-5 or C-6), 107.43 (C-2), 68.75 (CH_2); HRMS (EI): calcd.: 321.9953; found: $\{m/z: [\text{M}]^+\}$ 321.9940}.

2-Iodo-4-((3-nitrobenzyl)oxy)aniline (22). Prepared according to General procedure D from phenol **19** (682 mg, 2.90 mmol) and 1-(bromomethyl)-3-nitrobenzene (752 mg, 3.48 mmol). The crude product was purified by FCC (DCM/hexanes 90 : 10) to give ether **22** (405 mg, 1.09 mmol, 38%) as a brown solid; m.p. 109 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3420, 3345, 3050, 2856, 1615, 1599, 1569, 1519, 1493, 1452, 1346, 1317, 1230, 1091, 1043, 930, 869, 802, 729, 665; δ_{H} (500 MHz; $(\text{CD}_3)_2\text{SO}$) 8.27 (td, $J = 1.7, 0.8$ Hz, 1H, 2'-H), 8.18 (ddd, $J = 8.3, 2.4, 1.1$ Hz, 1H, 4'-H), 7.87 (ddd, $J = 7.6, 1.7, 1.0$ Hz, 1H, 6'-H), 7.69 (t, $J = 7.9$ Hz, 1H, 5'-H), 7.28 (d, $J = 2.8$ Hz, 1H, 3-H), 6.87 (dd, $J = 8.8, 2.9$ Hz, 1H, 5-H), 6.72 (d, $J = 8.8$ Hz, 1H, 6-H), 5.13 (s, 2H, CH_2), 4.81 (s, 2H, NH_2). δ_{C} (126 MHz; $(\text{CD}_3)_2\text{SO}$) 149.63 (C-4), 147.81 (C-3'), 143.22 (C-1), 139.80 (C-1'), 133.99 (C-6'), 129.99 (C-5'), 124.28 (C-3), 122.61 (C-4'), 121.88 (C-2'), 117.04 (C-5), 114.88 (C-6), 83.11 (C-2), 68.73 (CH_2); HRMS (ESI): calcd.: 370.9887; found: $\{m/z: [\text{M} + \text{H}]^+\}$ 370.9886}.

N-(2-Chloro-4-((3-nitrobenzyl)oxy)phenyl)-1-methyl-1H-pyrazole-4-carboxamide (23). Prepared according to General procedure A from 1-methyl-1H-pyrazole-4-carboxylic acid (195 mg, 1.55 mmol) and aniline **20** (440 mg, 1.55 mmol). The reaction mixture was stirred at 65 °C for 3 days. The crude

product was purified by recrystallisation from DCM to give amide **23** (363 mg, 0.938 mmol, 61%) as an off-white solid; m.p. 191 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3413, 3119, 1671, 1584, 1552, 1527, 1477, 1349, 1279, 1248, 1216, 1093, 1050, 1003, 876, 850, 807, 800, 750, 731, 708; δ_{H} (400 MHz; $(\text{CD}_3)_2\text{SO}$) 9.53 (s, 1H, CONH), 8.33 (t, $J = 2.0$ Hz, 1H, 2''-H), 8.27 (s, 1H, 5-H), 8.21 (ddd, $J = 8.2, 2.4, 1.1$ Hz, 1H, 4''-H), 7.97 (s, 1H, 3-H), 7.93 (ddd, $J = 7.7, 1.7, 1.0$ Hz, 1H, 6''-H), 7.72 (t, $J = 7.9$ Hz, 1H, 5''-H), 7.41 (d, $J = 8.8$ Hz, 1H, 6'-H), 7.26 (d, $J = 2.8$ Hz, 1H, 3'-H), 7.06 (dd, $J = 8.8, 2.9$ Hz, 1H, 5'-H), 5.32 (s, 2H, CH_2), 3.88 (s, 3H, CH_3). δ_{C} (101 MHz; $(\text{CD}_3)_2\text{SO}$) 160.71 (CONH), 156.35 (C-4'), 147.87 (C-3''), 139.04 (C-1''), 138.83 (C-3), 134.14 (C-6''), 132.61 (C-5), 130.53 (C-2'), 130.13 (C-5''), 129.70 (C-6'), 128.13 (C-1'), 122.88 (C-4''), 122.11 (C-2''), 117.82 (C-4), 115.47 (C-3'), 114.23 (C-5'), 68.41 (CH_2), 38.83 (CH_3); HRMS (ESI): calcd.: 387.0855; found: $\{m/z: [\text{M} + \text{H}]^+\}$ 387.0854}.

N-(2-Bromo-4-((3-nitrobenzyl)oxy)phenyl)-1-methyl-1H-pyrazole-4-carboxamide (24). Prepared according to General procedure A from 1-methyl-1H-pyrazole-4-carboxylic acid (1.23 g, 9.75 mmol) and aniline **21** (3.15 g, 9.75 mmol). The reaction mixture was stirred at 65 °C for 6 days. The crude product was purified by recrystallisation from DCM to give amide **24** (2.97 g, 6.89 mmol, 71%) as a white solid; m.p. 192 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3399, 3116, 2992, 1674, 1605, 1587, 1523, 1475, 1410, 1385, 1342, 1279, 1217, 1046, 1028, 852, 811, 749, 732.; δ_{H} (400 MHz; $(\text{CD}_3)_2\text{SO}$) 9.52 (s, 1H, CONH), 8.33 (t, $J = 2.0$ Hz, 1H, 2''-H), 8.26 (s, 1H, 5-H), 8.21 (dt, $J = 8.2, 1.6$ Hz, 1H, 4''-H), 7.97 (s, 1H, 3-H), 7.93 (d, $J = 7.7$ Hz, 1H, 6''-H), 7.72 (t, $J = 7.9$ Hz, 1H, 5''-H), 7.41 (d, $J = 2.8$ Hz, 1H, 3'-H), 7.38 (d, $J = 8.8$ Hz, 1H, 6'-H), 7.10 (dd, $J = 8.8, 2.8$ Hz, 1H, 5'-H), 5.32 (s, 2H, CH_2), 3.88 (s, 3H, CH_3). δ_{C} (101 MHz; $(\text{CD}_3)_2\text{SO}$) 160.45 (CONH), 156.24 (C-4'), 147.61 (C-3''), 138.79 (C-1''), 138.55 (C-3), 133.88 (C-6''), 132.31 (C-5), 129.87 (C-6' or C-5''), 129.70 (C-6' or C-5''), 129.37 (C-1'), 122.61 (C-4''), 121.84 (C-2''), 121.24 (C-2'), 118.14 (C-3'), 117.62 (C-4), 114.54 (C-5'), 68.16 (CH_2), 38.57 (CH_3); HRMS (ESI): calcd.: 429.0204; found: $\{m/z: [\text{M}-\text{H}]^-\}$ 429.0203}.

N-(2-Iodo-4-((3-nitrobenzyl)oxy)phenyl)-1-methyl-1H-pyrazole-4-carboxamide (25). Prepared according to General procedure A from 1-methyl-1H-pyrazole-4-carboxylic acid (122 mg, 0.964 mmol) and aniline **22** (357 mg, 0.964 mmol). The reaction mixture was stirred at 65 °C for 16 h. The crude product was purified by FCC (DCM/MeOH 98 : 2) to give amide **25** (216 mg, 0.452 mmol, 47%) as a brown solid; m.p. 213 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3378, 3125, 1657, 1579, 1553, 1520, 1440, 1399, 1347, 1299, 1273, 1218, 1095, 1057, 998, 866, 846, 808, 752, 727, 697; δ_{H} (500 MHz; $(\text{CD}_3)_2\text{SO}$) 9.51 (s, 1H, CONH), 8.33 (t, $J = 2.0$ Hz, 1H, 2''-H), 8.25 (s, 1H, 6-H), 8.21 (ddd, $J = 8.3, 2.4, 1.1$ Hz, 1H, 4''-H), 7.97 (s, 1H, 4-H), 7.94–7.91 (m, 1H, 6''-H), 7.72 (t, $J = 7.9$ Hz, 1H, 5''-H), 7.58 (d, $J = 2.8$ Hz, 1H, 3'-H), 7.28 (d, $J = 8.7$ Hz, 1H, 6'-H), 7.11 (dd, $J = 8.7, 2.8$ Hz, 1H, 5'-H), 5.30 (s, 2H, CH_2), 3.89 (s, 3H, CH_3). δ_{C} (126 MHz; $(\text{CD}_3)_2\text{SO}$) 160.69 (CONH), 156.46 (C-4'), 147.86 (C-3''), 139.13 (C-1''), 138.78 (C-4), 134.12 (C-6''), 133.10 (C-1'), 132.52 (C-6), 130.12 (C-5''), 129.26 (C-6'), 124.29 (C-3'), 122.84



(C-4''), 122.07 (C-2''), 118.08 (C-5), 115.41 (C-5'), 99.74 (C-2'), 68.33 (CH₂), 38.24 (CH₃); HRMS (ESI): calcd.: 479.0211; found: {*m/z*: [M + H]⁺ 479.0208}.

1-Methyl-*N*-(4-((3-nitrobenzyl)oxy)-2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)phenyl)-1*H*-pyrazole-4-carboxamide (26). PdCl₂(dppf)·DCM (20.4 mg, 27.8 μmol), bis(pinacolato)diboron (141 mg, 0.557 mmol), bromoarene **24** (120 mg, 0.278 mmol), and KOAc (109 mg, 1.11 mmol) were dissolved in degassed 1,4-dioxane (15 mL) under N₂ atmosphere and stirred at 95 °C for 20 h. The reaction mixture was cooled to room temperature and then filtered. The filtrate was diluted with EtOAc (50 mL). Water (50 mL) was added, the resulting two phases were separated, and the aqueous phase was extracted with EtOAc (3 × 60 mL). The combined organic phases were dried using a phase separation paper and concentrated *in vacuo*. The crude product was purified by FCC (hexanes/EtOAc 4:96 + 3% MeOH) to give boronic acid pinacol ester **26** (34.5 mg, 72.1 μmol, 26%) as a pale-yellow solid; m.p. 199 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 2967, 1641, 1606, 1530, 1482, 1349, 1315, 1266, 1157, 1129, 1013, 876, 802, 726; δ_{H} (400 MHz; (CD₃)₂SO) 11.67 (s, 1H, CONH), 8.42 (s, 1H, 5-H), 8.31 (t, *J* = 2.0 Hz, 1H, 2''-H), 8.19 (ddd, *J* = 8.2, 2.4, 1.0 Hz, 1H, 4''-H), 8.13 (s, 1H, 3-H), 7.93–7.90 (m, 1H, 6''-H), 7.70 (t, *J* = 7.9 Hz, 1H, 5''-H), 7.16 (d, 1H, 6'-H), 6.98–6.94 (m, 2H, 5'-H), 5.26 (s, 2H, CH₂), 3.95 (s, 3H, NCH₃), 1.16 (s, 12H, (CH₃)₄). δ_{C} (101 MHz; (CD₃)₂SO) 159.77 (CONH), 155.40 (C-4'), 147.83 (C-3''), 139.95 (C-1''), 139.05 (C-3), 133.89 (C-6''), 133.72 (C-5), 132.37 (C-1'), 130.07 (C-5''), 122.58 (C-4''), 121.85 (C-2''), 117.34 (C-3' and C-6'), 114.48 (C-5'), 113.92 (C-4), 79.44 ((C(CH₃)₂)₂), 67.98 (CH₂), 38.89 (NCH₃), 26.05 ((CH₃)₄); HRMS (ESI): calcd.: 479.2096; found: {*m/z*: [M + H]⁺ 479.2092}.

(2-(1-Methyl-1*H*-pyrazole-4-carboxamido)-5-((3-nitrobenzyl)oxy)phenyl)boronic acid (27). To a stirred solution of boronic acid pinacol ester **26** (330 mg, 0.690 mmol) in THF/water 4:1 v/v (25 mL) was added sodium periodate (738 mg, 3.45 mmol) and the reaction mixture was stirred vigorously at room temperature for 45 minutes. 1 M HCl (0.828 mmol, 0.828 mL) was then added, and the mixture stirred for 15 h. The milky suspension was diluted with DCM (15 mL), filtered and the residue washed with DCM (1×) and MeOH (1×). The filtrate was discarded and the solid residue collected to give boronic acid **27** (274 mg, 0.692 mmol, quant.) as an off-white solid; m.p. 260 °C (decomposition); IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3545, 3295, 3107, 2858, 1640, 1602, 1532, 1476, 1346, 1206, 1098, 1015, 882, 805, 739; δ_{H} (400 MHz; CF₃COOD) 9.00 (s, 1H, 3''-H), 8.65 (s, 1H, 5''-H), 8.46 (s, 1H, 2'-H), 8.34 (d, *J* = 8.8 Hz, 1H, 4'-H), 7.92 (d, *J* = 7.6 Hz, 1H, 6'-H), 7.69 (t, *J* = 8.0 Hz, 1H, 5'-H), 7.38–7.33 (m, 2H, 3-H and 6-H), 7.25 (dd, *J* = 8.7, 2.8 Hz, 1H, 4-H), 5.38 (s, 2H, CH₂), 4.29 (s, 3H, CH₃). δ_{C} (101 MHz; CF₃COOD) 160.53 (C-5), 159.44 (CONH), 149.88 (C-3'), 140.65 (C-1'), 139.87 (C-3''), 139.00 (C-5''), 136.12 (C-6'), 132.11 (C-2), 131.70 (C-5'), 125.19 (C-4'), 124.13 (C-2'), 120.57 (C-3), 119.89 (C-4), 117.07 (C-6), 114.82 (C-4''), 71.13 (CH₂), 40.45 (CH₃); HRMS (ESI): calcd.: 379.1208; found: {*m/z*: [M-H₂O + H]⁺ 379.1205}.

***N*-(2-Cyano-4-((3-nitrobenzyl)oxy)phenyl)-1-methyl-1*H*-pyrazole-4-carboxamide (28).** To a stirred solution of bromoarene **24** (350 mg, 0.812 mmol) in dry DMF (4 mL) was added CuCN (147 mg, 1.62 mmol). The reaction mixture was stirred at 150 °C for 22 h. Afterwards, the reaction mixture was concentrated *in vacuo* and the crude residue was resuspended in 20 mL aq. NH₄OH. The resulting suspension was vacuum filtered, and the residue was washed with aq. NH₄OH until the filtrate was no longer blue. The filtrate was discarded, and the filter residue was further washed sequentially with water (3 × 25 mL), DCM (4 × 25 mL), EtOAc (2 × 25 mL) and MeOH (2 × 25 mL). The filter residue was collected to give nitrile **28** (142 mg, 0.376 mmol, 46%) as a brownish-yellow solid; m.p. 197 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3293, 3117, 3074, 2922, 2229, 1651, 1615, 1587, 1518, 1483, 1419, 1387, 1348, 1286, 1227, 1161, 1101, 1048, 1003, 977, 886, 875, 823, 730; δ_{H} (400 MHz; (CD₃)₂SO) 10.06 (s, 1H, CONH), 8.34 (t, *J* = 2.0 Hz, 1H, 2''-H), 8.29 (s, 1H, 5-H), 8.24–8.20 (m, 1H, 4''-H), 7.99 (s, 1H, 3-H), 7.95–7.91 (m, 1H, 6''-H), 7.72 (t, *J* = 8.0 Hz, 1H, 5''-H), 7.57 (d, *J* = 2.6 Hz, 1H, 3'-H), 7.44–7.40 (m, 2H, 5'-H and 6'-H), 5.35 (s, 2H, OCH₂), 3.90 (s, 3H, CH₃). δ_{C} (101 MHz; (CD₃)₂SO) 160.77 (CONH), 155.43 (C-4'), 147.88 (C-3''), 138.90 (C-1''), 138.77 (C-3), 134.23 (C-6''), 133.61 (C-1'), 132.86 (C-5), 130.16 (C-5''), 128.68 (C-6'), 122.97 (C-4''), 122.22 (C-2''), 121.10 (C-5'), 117.98 (C-3'), 117.45 (C-4), 116.72 (CN), 110.38 (C-2'), 68.58 (OCH₂), 38.89 (CH₃); HRMS (ESI): calcd.: 376.1046; found: {*m/z*: [M-H]⁻ 376.1053}.

***N*-(2-Chloro-4-((3-(2-((4,6-dimethylpyrimidin-2-yl)thio)acetamido)benzyl)oxy)phenyl)-1-methyl-1*H*-pyrazole-4-carboxamide (29).** Prepared according to General procedure E from nitroarene **23** (150 mg, 0.388 mmol). The obtained crude amine was reacted with 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid (154 mg, 0.779 mmol) according to General procedure C. The reaction mixture was stirred at room temperature for 2 days. The crude product was purified by FCC (DCM/MeOH 98:2) to give **29** (145 mg, 0.269 mmol, 69% over two steps) as a pale-yellow solid; m.p. 86 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3270, 3081, 1651, 1580, 1552, 1514, 1490, 1440, 1264, 1203, 1045, 1004, 874, 784, 755, 690; δ_{H} (400 MHz; (CD₃)₂SO) 10.29 (s, 1H, 3''-NHCO), 9.51 (s, 1H, 4-CONH), 8.26 (s, 1H, 5-H), 7.97 (s, 1H, 3-H), 7.69 (t, *J* = 1.9 Hz, 1H, 2''-H), 7.54 (ddd, *J* = 8.2, 2.2, 1.1 Hz, 1H, 4''-H), 7.38 (d, *J* = 8.9 Hz, 1H, 6'-H), 7.33 (t, *J* = 7.8 Hz, 1H, 5''-H), 7.18 (d, *J* = 2.8 Hz, 1H, 3'-H), 7.16–7.12 (m, 1H, 6''-H), 6.99 (dd, *J* = 8.8, 2.8 Hz, 1H, 5'-H), 6.96 (s, 1H, 5'''-H), 5.13 (s, 2H, OCH₂), 4.04 (s, 2H, SCH₂), 3.88 (s, 3H, NCH₃), 2.32 (d, *J* = 0.5 Hz, 6H, 4'''-CH₃ and 6'''-CH₃). δ_{C} (101 MHz; (CD₃)₂SO) 169.31 (C-2'''), 166.96 (C-4''' and C-6'''), 166.63 (3''-NHCO), 160.71 (4-CONH), 156.70 (C-4'), 139.23 (C-3''), 138.82 (C-3), 137.33 (C-1''), 132.59 (C-5), 130.48 (C-2'), 129.64 (C-6'), 128.92 (C-5''), 127.81 (C-1'), 122.48 (C-6''), 118.65 (C-4''), 118.08 (C-2''), 117.85 (C-4), 116.06 (C-5'''), 115.32 (C-3'), 114.18 (C-5'), 69.65 (OCH₂), 38.82 (NCH₃), 35.47 (SCH₂), 23.32 (4'''-CH₃ and 6'''-CH₃); HRMS (ESI): calcd.: 537.1470; found: {*m/z*: [M + H]⁺ 537.1466}; purity (HPLC): 210 nm: >95%; 254 nm: 89% (method 1a).



***N*-(2-Bromo-4-((3-(2-((4,6-dimethylpyrimidin-2-yl)thio)acetamido)benzyl)oxy)phenyl)-1-methyl-1*H*-pyrazole-4-carboxamide (30).** Prepared according to General procedure E from nitroarene **24** (1.36 g, 3.14 mmol). The obtained crude amine was reacted with 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid (1.25 g, 6.31 mmol) according to General procedure C. The reaction mixture was stirred at room temperature for 4 days. The crude product was purified by FCC (DCM/MeOH 98:2) to give **30** (1.18 g, 2.03 mmol, 64% over two steps) as a yellow solid; m.p. 83 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3269, 2923, 1651, 1581, 1553, 1514, 1490, 1441, 1265, 1219, 1203, 1032, 1004, 872, 785, 757, 692; δ_{H} (500 MHz; $(\text{CD}_3)_2\text{SO}$) 10.29 (s, 1H, 3''-NHCO), 9.50 (s, 1H, 4-CONH), 8.26 (s, 1H, 5-H), 7.96 (s, 1H, 3-H), 7.68 (t, $J = 1.8$ Hz, 1H, 2''-H), 7.54 (dt, $J = 8.0, 1.7$ Hz, 1H, 4''-H), 7.37–7.31 (m, 3H, 3'-H, 6'-H and 5''-H), 7.14 (dt, $J = 7.7$ Hz, 1H, 6''-H), 7.04 (dd, $J = 8.8, 2.8$ Hz, 1H, 5'-H), 6.96 (s, 1H, 5'''-H), 5.13 (s, 2H, OCH₂), 4.04 (s, 2H, SCH₂), 3.88 (s, 3H, NCH₃), 2.32 (s, 6H, 4'''-CH₃ and 6'''-CH₃). δ_{C} (126 MHz; $(\text{CD}_3)_2\text{SO}$) 169.31 (C-2''), 166.96 (C-4'' and C-6''), 166.63 (3''-NHCO), 160.71 (4-CONH), 156.85 (C-4'), 139.23 (C-3''), 138.81 (C-3), 137.34 (C-1''), 132.57 (C-5), 129.90 (C-6'), 129.31 (C-1'), 128.93 (C-5''), 122.47 (C-6''), 121.47 (C-2'), 118.64 (C-4''), 118.26 (C-3'), 118.06 (C-2''), 117.91 (C-4), 116.06 (C-5''), 114.73 (C-5'), 69.65 (OCH₂), 38.83 (NCH₃), 35.47 (SCH₂), 23.33 (4'''-CH₃ and 6'''-CH₃); HRMS (ESI): calcd.: 581.0965; found: $\{m/z: [\text{M} + \text{H}]^+\}$ 581.0967; purity (HPLC): 210 nm: >95%; 254 nm: >95% (method 1a).

***N*-(4-((3-(2-((4,6-Dimethylpyrimidin-2-yl)thio)acetamido)benzyl)oxy)-2-iodophenyl)-1-methyl-1*H*-pyrazole-4-carboxamide (31).** Prepared according to General procedure E from nitroarene **25** (100 mg, 0.209 mmol). The obtained crude amine was reacted with 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid (83.1 mg, 0.419 mmol) according to General procedure C. The reaction mixture was stirred at room temperature for 18 h. The crude product was purified by FCC (DCM/MeOH 98:2) to give **31** (83.7 mg, 0.133 mmol, 64% over two steps) as a beige solid; m.p. 107 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3269, 2919, 1649, 1581, 1552, 1511, 1487, 1441, 1265, 1205, 1004, 978, 867, 784, 756, 691; δ_{H} (500 MHz; $(\text{CD}_3)_2\text{SO}$) 10.29 (s, 1H, 3''-NHCO), 9.49 (s, 1H, 4-CONH), 8.25 (s, 1H, 5-H), 7.97 (s, 1H, 3-H), 7.68 (t, $J = 1.9$ Hz, 1H, 2''-H), 7.54 (dt, $J = 8.3, 1.5$ Hz, 1H, 4''-H), 7.51 (d, $J = 2.8$ Hz, 1H, 3'-H), 7.33 (t, $J = 7.8$ Hz, 1H, 5''-H), 7.25 (d, $J = 8.7$ Hz, 1H, 6'-H), 7.13 (d, $J = 7.6$ Hz, 1H, 6''-H), 7.05 (dd, $J = 8.7, 2.9$ Hz, 1H, 5'-H), 6.97 (s, 1H, 5'''-H), 5.11 (s, 2H, OCH₂), 4.04 (s, 2H, SCH₂), 3.88 (s, 3H, NCH₃), 2.32 (s, 6H, 4'''-CH₃ and 6'''-CH₃). δ_{C} (126 MHz; $(\text{CD}_3)_2\text{SO}$) 169.31 (C-2''), 166.96 (C-4'' and C-6''), 166.62 (3''-NHCO), 160.69 (4-CONH), 156.81 (C-4'), 139.22 (C-3''), 138.78 (C-3), 137.42 (C-1''), 132.78 (C-1'), 132.51 (C-5), 129.20 (C-6'), 128.91 (C-5''), 124.20 (C-3'), 122.44 (C-6''), 118.62 (C-4''), 118.10 (C-4), 118.02 (C-2''), 116.06 (C-5''), 115.31 (C-5'), 99.71 (C-2'), 69.56 (OCH₂), 38.83 (NCH₃), 35.47 (SCH₂), 23.33 (4'''-CH₃ and 6'''-CH₃); HRMS (ESI): calcd.: 629.0826; found: $\{m/z: [\text{M} + \text{H}]^+\}$ 629.0822; purity (HPLC): 210 nm: >95%; 254 nm: >95% (method 1c).

***N*-(4-((3-(2-((4,6-Dimethylpyrimidin-2-yl)thio)acetamido)benzyl)oxy)-2-formylphenyl)-1-methyl-1*H*-pyrazole-4-carboxamide (32).** **30** (150 mg, 0.258 mmol), Pd(OAc)₂ (1.74 mg, 7.74 μmol , 3 mol%), dppb (4.95 mg, 11.6 μmol , 4.5 mol%), *N*-formylsaccharin (172 mg, 0.774 mmol), and Na₂CO₃ (82.0 mg, 0.774 mmol) were added to a 30 mL glass tube, which was then evacuated and backfilled three times with N₂. A degassed solution of Et₃SiH (54.2 μL , 0.335 mmol) in DMF (0.6 mL) was added to the glass tube under N₂ atmosphere. The mixture was stirred for 15 min at room temperature and subsequently warmed to 75 °C and stirred for another 19 h. The reaction mixture was cooled to room temperature, then diluted with EtOAc (15 mL) and washed with brine (3 \times 15 mL). The organic phase was dried with a phase separation paper and concentrated *in vacuo*. The crude product was redissolved in MeOH (5 mL), sat. aq. NaHSO₃ (25 mL) was added, stirred for approximately 30 seconds, diluted with H₂O (25 mL), and then extracted with EtOAc (3 \times 25 mL). The aqueous phase was basified with 50% NaOH (10 mL) and extracted with DCM (3 \times 25 mL). The combined organic layers were dried with Na₂SO₄ and concentrated *in vacuo*. The obtained crude residue was purified by FCC (DCM/MeOH 98:2) to give **32** (3.8 mg, 7.2 μmol , 3%) as a yellow solid; m.p. 87 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3232, 2921, 2852, 1727, 1654, 1582, 1553, 1529, 1487, 1435, 1286, 1264, 1220, 1154, 1031, 893, 785, 696; δ_{H} (500 MHz; $(\text{CD}_3)_2\text{SO}$) 11.45 (s, 1H, 4-CONH), 9.91 (s, 1H, CHO), 9.58 (s, 1H, 3''-NHCO), 8.76 (d, $J = 10.1$ Hz, 1H, 6'-H), 7.96 (s, 2H, 3-H and 5-H), 7.63 (t, 1H, 2''-H), 7.38 (dt, $J = 8.0, 1.6$ Hz, 1H, 4''-H), 7.32 (t, $J = 7.8$ Hz, 1H, 5''-H), 7.28–7.25 (m, 2H, 3'-H and 5'-H), 7.16 (d, $J = 7.4$ Hz, 1H, 6''-H), 6.85 (s, 1H, 5'''-H), 5.11 (s, 2H, OCH₂), 3.95 (s, 3H, NCH₃), 3.85 (s, 2H, SCH₂), 2.47 (s, 6H, 4'''-CH₃ and 6'''-CH₃). δ_{C} (126 MHz; $(\text{CD}_3)_2\text{SO}$) 195.98 (CHO), 170.43 (C-2''), 168.31 (3''-NHCO), 168.11 (4'''-CH₃ and 6'''-CH₃), 161.20 (4-CONH), 154.24 (C-4'), 139.16 (C-3''), 139.03 (C-3), 138.00 (C-1''), 135.85 (C-1'), 131.99 (C-5), 129.60 (C-5''), 123.70 (C-5'), 123.20 (C-6''), 122.70 (C-2'), 121.61 (C-6'), 121.03 (C-3'), 119.76 (C-4), 119.46 (C-4''), 118.76 (C-2''), 117.06 (C-5''), 70.71 (OCH₂), 39.73 (NCH₃), 35.90 (SCH₂), 30.09 (4'''-CH₃ and 6'''-CH₃); HRMS (ESI): calcd.: 553.1633; found: $\{m/z: [\text{M} + \text{Na}]^+\}$ 553.1633; purity (HPLC): 210 nm: >95%; 254 nm: >95% (method 1b).

5-((3-(2-((4,6-Dimethylpyrimidin-2-yl)thio)acetamido)benzyl)oxy)-2-(1-methyl-1*H*-pyrazole-4-carboxamido)phenyl-boronic acid (33). Prepared according to General procedure F from nitroarene **27** (90.0 mg, 0.227 mmol). The obtained crude amine was dissolved in dry DMF (3 mL) and bromoacetyl bromide (23.8 μL , 0.274 mmol) was added. The reaction mixture was stirred at room temperature for 30 minutes. The solution was then diluted with EtOAc (50 mL) and washed with brine (3 \times 20 mL). The organic phase was dried using a phase separation paper and concentrated *in vacuo*. The brown oily crude product was redissolved in dry DMF (8 mL), 4,6-dimethylpyrimidine-2-thiol (64.5 mg, 0.46 mmol) and *t*-BuOK (51.6 mg, 0.46 mmol) were added, and the reaction mixture was stirred at room temperature for 19 h. The solution was then diluted with EtOAc (150 mL) and



washed with brine (3 × 100 mL). The organic phase was dried using a phase separation paper and concentrated *in vacuo*. The crude product was purified by FCC (DCM/10% NH₃ in MeOH 88:12) to give product **33** (32.1 mg, 58.7 μmol, 26% over three steps) as a pale-yellow solid; m.p. 165 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3271, 2922, 2853, 1667, 1633, 1601, 1581, 1553, 1532, 1480, 1441, 1312, 1264, 1205, 1009, 874, 787, 749, 690; δ_{H} (400 MHz; (CD₃)₂SO) 11.78 (s, 1H, 4'-CONH), 10.20 (s, 1H, 3''-NHCO), 8.25 (s, 1H, 5'-H), 7.96 (s, 1H, 3'-H), 7.62 (d, *J* = 8.3 Hz, 1H, 3-H), 7.58 (s, 1H, 2''-H), 7.53 (d, *J* = 8.2 Hz, 1H, 4''-H), 7.30 (d, *J* = 3.0 Hz, 1H, 6-H), 7.21 (t, *J* = 7.8 Hz, 1H, 5''-H), 7.04 (d, *J* = 7.6 Hz, 1H, 6''-H), 6.94 (dd, *J* = 8.9, 3.0 Hz, 1H, 4-H), 6.86 (s, 1H, 5'''-H), 5.00 (s, 2H, OCH₂), 3.98 (s, 2H, SCH₂), 3.71 (s, 3H, NCH₃), 2.28 (s, 6H, 4'''-CH₃ and 6'''-CH₃). δ_{C} (101 MHz; (CD₃)₂SO) 169.29 (C-2'''), 166.90 (C-4''' and C-6'''), 166.57 (3''-NHCO), 159.32 (4'-CONH), 155.13 (C-5) 139.07 (C-3''), 138.65 (C-3'), 137.98 (C-1''), 133.14 (C-2 and C-5'), 128.68 (C-5''), 122.30 (C-6''), 119.01 (C-6), 118.40 (C-4''), 117.91 (C-2''), 117.72 (C-3), 115.98 (C-4' and C-5'''), 114.14 (C-4), 69.07 (OCH₂), 38.75 (NCH₃), 35.41 (SCH₂), 23.28 (4'''-CH₃ and 6'''-CH₃); HRMS (ESI): calcd.: 527.1673; found: {*m/z*: [M-H₂O-H]⁻ 527.1680}; purity (HPLC): 210 nm: >95%; 254 nm: >95% (method 2a).

N-(2-Cyano-4-((3-(2-((4,6-dimethylpyrimidin-2-yl)thio)acetamido)benzyl)oxy)phenyl)-1-methyl-1H-pyrazole-4-carboxamide (34). Prepared according to General procedure E from nitroarene **28** (130 mg, 0.344 mmol). The obtained crude amine was reacted with 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid (103 mg, 0.520 mmol) according to General procedure C. The reaction mixture was stirred at room temperature for 7 days. The crude product was purified by FCC (DCM/10% NH₃ in MeOH 99:1) to give **34** (73.4 mg, 0.139 mmol, 40% over two steps) as a yellow solid; m.p. 93 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3271, 2922, 2853, 2229, 1652, 1581, 1552, 1490, 1441, 1265, 1226, 1161, 1003, 873, 785, 756, 691; δ_{H} (500 MHz; (CD₃)₂SO) 10.30 (s, 1H, 3''-NHCO), 10.04 (s, 1H, 4-CONH), 8.29 (s, 1H, 5-H), 7.98 (s, 1H, 3-H), 7.70 (t, *J* = 2.0 Hz, 1H, 2''-H), 7.54 (dt, *J* = 8.1, 1.9 Hz, 1H, 4''-H), 7.50 (d, *J* = 2.9 Hz, 1H, 3'-H or 5'-H), 7.41 (d, *J* = 8.9 Hz, 1H, 6'-H), 7.36–7.35 (m, 1H, 5''-H), 7.34–7.32 (m, 1H, 3'-H or 5'-H), 7.14 (d, *J* = 7.5 Hz, 1H, 6''-H), 6.96 (s, 1H, 5'''-H), 5.16 (s, 2H, OCH₂), 4.04 (s, 2H, SCH₂), 3.90 (s, 3H, NCH₃), 2.32 (s, 6H, 4'''-CH₃ and 6'''-CH₃). δ_{C} (126 MHz; (CD₃)₂SO) 169.31 (C-2'''), 166.96 (C-4''' and C-6'''), 166.64 (3''-NHCO), 160.76 (4-CONH), 155.75 (C-4'), 139.25 (C-3''), 138.89 (C-3), 137.08 (C-1''), 133.30 (C-1'), 132.85 (C-5), 128.96 (C-3' or C-5'), 128.63 (C-6'), 122.59 (C-6''), 121.03 (C-5''), 118.73 (C-4''), 118.17 (C-2''), 117.82 (C-3' or C-5'), 117.47 (C-4), 116.77 (CN), 116.06 (C-5'''), 110.36 (C-2'), 69.80 (OCH₂), 38.87 (NCH₃), 35.48 (SCH₂), 23.32 (4'''-CH₃ and 6'''-CH₃); HRMS (ESI): calcd.: 528.1812; found: {*m/z*: [M + H]⁺ 528.1814}; purity (HPLC): 210 nm: >95%; 254 nm: >95% (method 1a).

8-((2-Aminothiazol-5-yl)methyl)-2-naphthonitrile (39). To a stirred solution of bromo derivative **38** (500 mg, 1.57 mmol) in dry DMF (5 mL) were added zinc cyanide (110 mg, 0.940 mmol) and Pd(PPh₃)₄ (181 mg, 0.157 mmol). The reaction

mixture was stirred at 80 °C for 18 h under N₂ atmosphere, then was diluted with water (50 mL) and extracted with EtOAc (3 × 100 mL). The combined organic layers were dried over sodium sulfate, the solvent was evaporated *in vacuo* and the crude product was purified by FCC (hexanes/EtOAc/NET₃ 30:70:1) to give **39** (254 mg, 0.957 mmol, 61%) as a light-yellow solid; m.p. 216 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3397, 3283, 3116, 2221, 1629, 1517, 1438, 1377, 1326, 1310, 1270, 1204, 1160, 1120, 1052, 889, 861, 835, 794, 749, 721, 688; δ_{H} (500 MHz; (CD₃)₂SO) 8.75 (d, *J* = 1.6 Hz 1H, 8-H), 8.13 (d, *J* = 8.4 Hz, 1H, 5-H), 7.95 (d, *J* = 8.2 Hz, 1H, 4-H), 7.80 (dd, *J* = 8.5, 1.5 Hz, 1H, 6-H), 7.67 (dd, *J* = 8.2, 7.0 Hz, 1H, 3-H), 7.57 (dd, *J* = 7.2, 1.2 Hz, 1H, 2-H), 6.77 (s, 1H, 4'-H), 6.69 (s, 2H, NH₂), 4.46 (s, 2H, CH₂). δ_{C} (126 MHz; (CD₃)₂SO) 167.80 (C-2'), 137.86 (C-1), 135.88 (C-4'), 134.94 (C-4a), 130.64 (C-8), 130.14 (C-8a), 130.13 (C-5), 129.10 (C-3), 127.99 (C-2), 127.25 (C-4), 126.23 (C-6), 124.03 (C-5'), 119.37 (CN), 108.50 (C-7), 29.46 (CH₂); HRMS (ESI): calcd.: 266.0746; found: {*m/z*: [M + H]⁺ 266.0745}.

2-(8-((2-Aminothiazol-5-yl)methyl)naphthalen-2-yl)acetonitrile (40). To a stirred solution of bromoarene **38** (323 mg, 1.01 mmol) in DMF (1.21 mL) was added 4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)isoxazole (237 mg, 1.22 mmol) and 1 M KF solution (513 μL, 513 μmol). The mixture was degassed in an ultrasonic bath under nitrogen atmosphere. PdCl₂(dppf)-DCM (12.5 mg, 0.0171 mmol) was then added under nitrogen counterflow and the reaction mixture was stirred at 90 °C for 20 h. The mixture was allowed to cool to room temperature, diluted with water (50 mL) and brine (50 mL), and extracted with EtOAc (3 × 50 mL). The combined organic layers were dried over sodium sulfate. After evaporating the solvent *in vacuo*, the crude product was purified by FCC (hexanes/EtOAc 50:50) to give cyanomethyl derivative **40** (74.4 mg, 0.266 mmol, 26%) as a light brown solid; m.p. 203 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 3463, 3112, 1613, 1519, 1403, 1354, 1301, 1195, 1035, 930, 844, 825, 794, 770, 754, 673; δ_{H} (400 MHz; (CD₃)₂SO) 8.14 (d, *J* = 1.6 Hz, 1H, 8-H), 7.98 (d, *J* = 8.4 Hz, 1H, 5-H), 7.84 (dd, *J* = 8.1, 1.6 Hz 1H, 4-H), 7.52–7.41 (m, 3H, 2-H, 3-H, 6-H), 6.77 (s, 1H, 4'-H), 6.66 (s, 2H, NH₂), 4.37 (s, 2H, CH₂), 4.23 (s, 2H, CH₂CN). δ_{C} (101 MHz; (CD₃)₂SO) 167.69 (C-2'), 136.37 (C-1), 135.67 (C-4'), 132.58 (C-4a), 131.01 (C-8a), 129.46 (C-5), 128.98 (C-7), 126.95 (C-4), 126.89 (C-2), 125.99 (C-3), 135.87 (C-6), 124.19 (C-5'), 122.85 (C-8), 119.21 (CN), 29.74 (CH₂), 22.89 (CH₂CN); HRMS (ESI): calcd.: 280.0903; found: {*m/z*: [M + H]⁺ 280.0902}.

tert-Butyl (5-((7-bromonaphthalen-1-yl)methyl)thiazol-2-yl) carbamate (41). To a stirred solution of aminothiazole **38** (0.738 g, 2.31 mmol) in toluene (75 mL) was added di-*tert*-butyl dicarbonate (1.98 mL, 9.24 mmol) dropwise at room temperature. The reaction mixture was stirred at 100 °C for 4.5 h and then the solvent was evaporated *in vacuo*. The crude product was purified by FCC (hexanes/EtOAc 80:20) to give **41** (828 mg, 1.98 mmol, 85%) as a beige solid; m.p. 296 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 2726, 1710, 1579, 1546, 1495, 1447, 1391, 1368, 1321, 1297, 1251, 1236, 1080, 1056, 1028, 880, 860, 844, 825, 809, 763, 750, 683; δ_{H} (400 MHz; (CD₃)₂SO) 11.25 (s, 1H, NHCO), 8.32 (d, *J* = 2.0 Hz, 1H, 8-H), 7.93 (d, *J* = 8.8 Hz, 1H,



5-H), 7.87 (m, 1H, 4-H), 7.65 (dd, $J = 8.8, 1.9$ Hz, 1H, 6-H), 7.56–7.48 (m, 2H, 2-H, 3-H), 7.19 (s, 1H, 4'-H), 4.52 (s, 2H, CH₂), 1.41 (s, 9H, C(CH₃)₃). δ_C (126 MHz; (CD₃)₂SO) 158.38 (C-2'), 152.70 (NHCO), 135.65 (C-1), 134.90 (C-4'), 132.28 (C-8a), 132.08 (C-4a), 130.91 (C-5), 130.39 (C-5'), 128.88 (C-6), 127.75 (C-2), 127.37 (C-4), 126.40 (C-3), 126.02 (C-8), 119.78 (C-7), 80.88 (C(CH₃)₃), 29.29 (CH₂), 27.84 (C(CH₃)₃); HRMS (ESI): calcd.: 417.0278; found: $\{m/z: [M-H]^- 417.0285\}$.

tert-Butyl (5-((7-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)naphthalen-1-yl)methyl)thiazol-2-yl)carbamate (42). PdCl₂(dppf)-DCM (281 mg, 0.383 mmol), bis(pinacolato)diboron (1.46 g, 5.75 mmol), bromoarene derivative **41** (804 mg, 1.92 mmol) and KOAc (753 mg, 7.67 mmol) were dissolved in degassed anhydrous 1,4-dioxane (15 mL) under N₂ atmosphere and stirred at 80 °C for 1 h and then at room temperature for another 2 h. Subsequently the mixture was diluted with water (100 mL) and brine (100 mL) and extracted with EtOAc (3 × 100 mL). The combined organic layers were dried over sodium sulfate. After evaporating the solvent *in vacuo*, the crude product was purified by FCC (hexanes/EtOAc 100:0 → 100:20) to give boronic acid pinacol ester **42** (456 mg, 0.978 mmol, 51%) as a beige solid; m.p. 203 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 2975, 1719, 1624, 1568, 1457, 1369, 1341, 1309, 1251, 1240, 1142, 1090, 1059, 1009, 982, 962, 836, 804, 768, 755, 690; δ_H (400 MHz; (CD₃)₂SO) 11.24 (s, 1H, NHCO), 8.49 (s, 1H, 8-H), 7.93 (d, $J = 8.2$ Hz, 1H, 5-H), 7.85 (d, $J = 8.1$ Hz, 1H, 4-H), 7.73 (dd, $J = 8.2, 1.0$ Hz, 1H, 6-H), 7.53 (dd, $J = 8.2, 7.0$ Hz, 1H, 3-H), 7.45 (d, $J = 7.0$ Hz, 1H, 2-H), 7.13 (s, 1H, 4'-H), 4.55 (s, 2H, CH₂), 1.41 (s, 9H, C(CH₃)₃), 1.33 (s, 12H, O₂C₂(CH₃)₄). δ_C (101 MHz; (CD₃)₂SO) 158.35 (C-2'), 152.69 (NHCO), 136.82 (C-1), 135.08 (C-4a), 134.86 (C-4), 131.27 (C-8), 130.34 (C-5'), 130.31 (C-8a), 130.00 (C-6), 127.95 (C-5), 127.23 (C-4), 126.76 (C-3), 126.70 (C-2), 125.41 (C-7), 83.83 (O₂C₂(CH₃)₄), 80.86 (C(CH₃)₃), 29.25 (CH₂), 27.82 (C(CH₃)₃), 24.71 (O₂C₂(CH₃)₄); HRMS (ESI): calcd.: 465.2025; found: $\{m/z: [M-H]^- 465.2033\}$.

(8-((2-((tert-Butoxycarbonyl)amino)thiazol-5-yl)methyl)naphthalen-2-yl)boronic acid (43). To a stirred solution of **42** (541 mg, 1.16 mmol) in THF/water 3:1 v/v (6 mL), was added sodium periodate (744 mg, 3.48 mmol) and the reaction mixture was stirred at room temperature for 1 h. Then, 1 M HCl (1.16 mL) was added and the mixture was stirred for another 3 h, then the mixture was extracted with EtOAc (3 × 50 mL). The combined organic layers were dried over sodium sulfate. After evaporating the solvent *in vacuo*, the crude product was purified by FCC (DCM/MeOH 100:2) to give boronic acid **43** (376 mg, 0.978 mmol, 84%) as a white solid; m.p. 194 °C (decomposition); IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 2976, 1716, 1624, 1558, 1458, 1368, 1346, 1306, 1250, 1151, 1094, 1061, 1024, 837, 755, 697; δ_H (400 MHz; (CD₃)₂SO) 11.21 (s, 1H, NHCO), 8.68 (s, 1H, 8-H), 8.23 (s, 2H, B(OH)₂), 7.88–7.86 (m, 2H, 5-H, 6-H), 7.80 (d, $J = 8.1$ Hz, 1H, 4-H), 7.48 (dd, $J = 8.1, 7.0$ Hz, 1H, 3-H), 7.42 (dd, $J = 7.0, 1.3$ Hz, 1H, 2-H), 7.19 (s, 1H, 4'-H), 4.54 (s, 2H, CH₂), 1.41 (s, 9H, C(CH₃)₃). δ_C (126 MHz; (CD₃)₂SO) 158.33 (C-2'), 152.73

(NHCO), 136.95 (C-1), 134.82 (C-4), 134.57 (C-4a), 131.72 (C-7), 130.74 (C-6, C-8a), 130.55 (C-8), 130.46 (C-5'), 127.28 (C-5), 127.02 (C-4), 126.41 (C-3), 126.32 (C-2), 80.90 (C(CH₃)₃), 29.23 (CH₂), 27.88 (C(CH₃)₃); HRMS (ESI): calcd.: 385.1388; found: $\{m/z: [M + H]^+ 358.1388\}$.

(8-((2-Aminothiazol-5-yl)methyl)naphthalen-2-yl)boronic acid (44). To a stirred solution of *N*-Boc-derivative **43** (407 mg, 1.06 mmol) in chloroform (10 mL) was added TFA (3.98 mL, 53.0 mmol) at room temperature. The reaction mixture was stirred at room temperature for 17 h. Subsequently brine (15 mL) was added, and the solution was basified with NaOH (2.33 g, 58.3 mmol). After collecting the organic phase, the aqueous phase was extracted with DCM/2-propanol (3 × 30 mL, 4:1 v/v) and then EtOAc/2-propanol (3 × 30 mL, 4:1). The combined organic layers were dried over sodium sulfate and the solvent was evaporated *in vacuo* to give **44** (302 mg, 1.06 mmol, quant.) as a light greyish-white solid; m.p. 281 °C (decomposition); IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 2929, 1599, 1554, 1515, 1456, 1379, 1315, 1256, 1157, 1051, 834, 754, 699; δ_H (500 MHz; (CD₃)₂SO) 8.69 (d, $J = 1.1$ Hz, 1H, 8-H), 8.21 (s, 2H, B(OH)₂), 7.90–7.82 (m, 2H, 5-H, 6-H), 7.77 (d, $J = 8.1$ Hz, 1H, 4-H), 7.46 (dd, $J = 8.2, 7.0$ Hz, 1H, 3-H), 7.37 (dd, $J = 7.0, 1.2$ Hz, 1H, 2-H), 6.74 (s, 1H, 4'-H), 6.64 (s, 2H, NH₂), 4.40 (s, 2H, CH₂). δ_C (126 MHz; (CD₃)₂SO) 167.70 (C-2'), 137.26 (C-1), 135.59 (C-4'), 134.46 (C-4a), 131.55 (C-7) 130.78 (C-8), 130.49 (C-8a), 130.44 (C-6), 127.16 (C-5), 126.74 (C-4), 126.30 (C-3), 125.94 (C-2), 124.49 (C-5'), 29.58 (CH₂); HRMS (ESI): calcd.: 285.0864; found: $\{m/z: [M + H]^+ 285.0863\}$.

***N*-(5-((7-Bromonaphthalen-1-yl)methyl)thiazol-2-yl)-2-((4,6-dimethylpyrimidin-2-yl)thio)acetamide (45).** To a stirred solution of aminothiazole **38** (627 mg, 1.96 mmol) in dry DMF (5 mL) were added DMAP (120 mg, 0.982 mmol), EDC·HCl (461 mg, 2.36 mmol) and a solution of 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid (389 mg, 1.96 mmol) in dry DMF (1 mL) at room temperature. The reaction mixture was stirred at room temperature for 16 h. Water (50 mL) and brine (50 mL) were added, and the mixture was extracted with DCM (3 × 100 mL). The combined organic layers were dried over sodium sulfate, the solvent was evaporated *in vacuo* and the crude product was purified by FCC (DCM/EtOAc 90:10) to give **45** (509 mg, 1.02 mmol, 52%) as a white solid. Analytical data are in alignment with literature.²⁰

***N*-(5-((7-Cyanonaphthalen-1-yl)methyl)thiazol-2-yl)-2-((4,6-dimethylpyrimidin-2-yl)thio)acetamide (46).** To a stirred solution of 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid (194 mg, 0.976 mmol) in dry DMF (3 mL) were added DIPEA (429 μ L, 2.44 mmol) and HATU (371 mg, 0.976 mmol) and the reaction mixture was stirred at room temperature for 1 h. Aminothiazole **39** (216 mg, 0.814 mmol) was added and the reaction mixture was stirred at room temperature for another 18 h. Then the mixture was diluted with water (150 mL) and extracted with EtOAc (4 × 100 mL). The combined organic layers were dried over sodium sulfate. After evaporating the solvent *in vacuo*, the crude product was purified by FCC (hexanes/EtOAc 40:60) to give amide **46**



(105 mg, 0.235 mmol, 29%) as a white solid; m.p. 218 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 2897, 2221, 1687, 1581, 1530, 1429, 1374, 1322, 1258, 1243, 1161, 966, 876, 841, 819, 755, 715; δ_{H} (500 MHz; $(\text{CD}_3)_2\text{SO}$) 12.20 (s, 1H, CONH), 8.75 (d, $J = 1.6$ Hz, 1H, 8-H), 8.13 (d, $J = 8.4$ Hz, 1H, 5-H), 7.97 (d, $J = 8.2$ Hz, 1H, 4-H), 7.80 (dd, $J = 8.5, 1.5$ Hz, 1H, 6-H), 7.68 (dd, $J = 8.2, 7.0$ Hz, 1H, 3-H), 7.61 (dd, $J = 7.1, 1.3$ Hz, 1H, 2-H), 7.33 (s, 1H, 4'-H), 6.92 (s, 1H, CH_2), 4.64 (s, 2H, CH_2S), 4.05 (s, 2H, CH_2S), 2.25 (s, 6H, 4- CH_3 , 6- CH_3). δ_{C} (126 MHz; $(\text{CD}_3)_2\text{SO}$) 168.86 (C-2''), 166.96 (C-4'', C-6''), 166.85 (NHCO), 156.77 (C-2'), 137.45 (C-1), 135.07 (C-4'), 134.99 (C-4a), 130.74 (C-5'), 130.58 (C-8), 130.19 (C-5), 130.09 (C-8a), 129.12 (C-3), 128.31 (C-2), 127.49 (C-4), 126.32 (C-6), 119.32 (CN), 116.08 (C-5''), 108.65 (C-7), 34.01 (CH_2S), 28.95 (CH_2), 23.19 (4- CH_3 , 6- CH_3); HRMS (ESI): calcd.: 444.0958; found: $\{m/z: [\text{M}-\text{H}]^- 444.0956\}$; purity (HPLC): 210 nm: >95%; 254 nm: >95% (method 3c).

N-(5-((7-(Cyanomethyl)naphthalen-1-yl)methyl)thiazol-2-yl)-2-((4,6-dimethylpyrimidin-2-yl)thio)acetamide (47). To a stirred solution of 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid (73.4 mg, 0.370 mmol) in dry DMF (0.5 mL) were added DIPEA (130 μL , 0.741 mmol) and HATU (141 mg, 0.370 mmol) and the reaction mixture was stirred at room temperature for 1 h. Aminothiazole **40** (69.0 mg, 0.247 mmol) was added and the reaction mixture was stirred at room temperature for another 18 h. Then the mixture was diluted with water (50 mL) and extracted with EtOAc (4 \times 50 mL). The combined organic layers were dried over sodium sulfate. After evaporating the solvent *in vacuo*, the crude product was purified by FCC (hexanes/EtOAc 50 : 50) to give amide **47** (45.5 mg, 0.0990 mmol, 40%) as a white solid; m.p. 179 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 2925, 1687, 1580, 1549, 1524, 1505, 1434, 1403, 1340, 1280, 1260, 1229, 1177, 1155, 1121, 1035, 951, 932, 891, 845, 823, 792, 757, 717; δ_{H} (500 MHz; $(\text{CD}_3)_2\text{SO}$) 12.18 (s, 1H, NHCO), 8.13 (s, 1H, 8-H), 7.98 (d, $J = 8.4$ Hz, 1H, 5-H), 7.85 (dd, $J = 6.5, 3.0$ Hz, 1H, 4-H), 7.51–7.45 (m, 3H, 2-H, 3-H, 6-H), 7.31 (s, 1H, 4'-H), 6.92 (s, 1H, 5''-H), 4.54 (s, 2H, CH_2), 4.21 (s, 2H, CH_2CN), 4.05 (s, 2H, CH_2S), 2.26 (s, 6H, 4- CH_3 , 6- CH_3). δ_{C} (126 MHz; $(\text{CD}_3)_2\text{SO}$) 168.85 (C-2''), 166.98 (C-4'', C-6''), 166.77 (NHCO), 156.62 (C-2'), 135.98 (C-1), 134.87 (C-4'), 132.64 (C-4a), 130.97 (C-8a), 130.91 (C-5'), 129.52 (C-5), 129.15 (C-7), 127.20 (C-2, C-4), 126.03 (C-3), 125.95 (C-6), 122.86 (C-8), 119.18 (CN), 116.09 (C-5''), 34.00 (CH_2S), 29.28 (CH_2), 23.21 (4- CH_3 , 6- CH_3), 22.89 (CH_2CN); HRMS (ESI): calcd.: 458.1115; found: $\{m/z: [\text{M}-\text{H}]^- 458.1113\}$; purity (HPLC): 210 nm: >95%; 254 nm: >95% (method 3c).

(8-((2-(2-((4,6-Dimethylpyrimidin-2-yl)thio)acetamido)-thiazol-5-yl)methyl)naphthalen-2-yl)boronic acid (48). To a stirred solution of amine **44** (304 mg, 1.07 mmol) in dry DMF (5 mL) were added DMAP (65.4 mg, 0.535 mmol), EDC $\cdot\cdot\cdot\text{HCl}$ (251 mg, 1.28 mmol) and a solution of 2-((4,6-dimethylpyrimidin-2-yl)thio)acetic acid (212 mg, 1.07 mmol) in dry DMF (1 mL) at room temperature. The reaction mixture was stirred at room temperature for 16 h. Water (100 mL) and brine (50 mL) were added, and the mixture was extracted with EtOAc (3 \times 100 mL). The combined organic layers were dried over sodium sulfate, the solvent was

evaporated *in vacuo*, and the crude product was purified by FCC (DCM/MeOH/ NH_3 (aq., 25%) 100 : 5 : 0.05) to give amide **48** (82.5 mg, 0.178 mmol, 17%) as a white solid; m.p. 153 °C (decomposition); IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 2928, 1691, 1623, 1583, 1533, 1438, 1385, 1313, 1265, 1163, 1030, 972, 893, 837, 758, 668; δ_{H} (500 MHz; $(\text{CD}_3)_2\text{SO}$) 12.17 (s, 1H, NHCO), 8.68 (s, 1H, 8-H), 8.20 (s, 2H, $\text{B}(\text{OH})_2$), 7.88–7.84 (m, 2H, 5-H, 6-H), 7.79 (d, $J = 8.1$ Hz, 1H, 4-H), 7.47 (dd, $J = 8.2, 7.0$ Hz, 1H, 3-H), 7.41 (dd, $J = 7.0, 1.3$ Hz, 1H, 2-H), 7.27 (s, 1H, 4'-H), 6.92 (s, 1H, 5''-H), 4.57 (s, 2H, CH_2), 4.05 (s, 2H, CH_2S), 2.26 (s, 6H, 4- CH_3 , 6- CH_3). δ_{C} (126 MHz; $(\text{CD}_3)_2\text{SO}$) 168.87 (C-2''), 166.99 (C-4'', C-6''), 166.76 (NHCO), 156.58 (C-2'), 136.80 (C-1), 134.80 (C-4'), 134.51 (C-4a), 131.73 (C-7), 131.21 (C-5'), 130.70 (C-8), 130.51 (C-6), 130.43 (C-8a), 127.23 (C-5), 127.00 (C-4), 126.33 (C-3), 126.27 (C-2), 116.11 (C-5''), 34.01 (CH_2S), 29.10 (CH_2), 23.23 (4- CH_3 , 6- CH_3); HRMS (ESI): calcd.: 465.1221; found: $\{m/z: [\text{M} + \text{H}]^+ 465.1220\}$; purity (HPLC): 210 nm: >95%; 254 nm: >95% (method 3a).

2-((4,6-Dimethylpyrimidin-2-yl)thio)-N-(5-((7-formylnaphthalen-1-yl)methyl)thiazol-2-yl)acetamide (49). Bromoarene **45** (200 mg, 0.400 mmol), $\text{Pd}(\text{OAc})_2$ (2.70 mg, 0.0120 mmol, 3 mol%), dppf (9.99 mg, 0.0180 mmol, 4.5 mol%), *N*-formylsaccharin (134 mg, 0.601 mmol) and Na_2CO_3 (63.7 mg, 0.601 mmol) were added to a 30 mL glass tube, which was then evacuated and backfilled three times with N_2 . A degassed solution of Et_3SiH (84.1 μL , 0.521 mmol) in DMF (2 mL) was added to the glass tube under N_2 atmosphere. The mixture was stirred for 10 min at room temperature and subsequently warmed to 80 °C and stirred for another 16 h. After cooling to room temperature, the mixture was diluted with water (100 mL) and extracted with DCM (3 \times 50 mL). After the combined organic layers were dried over sodium sulfate, the solvent was evaporated *in vacuo* and the crude product was purified using two consecutive PTLCS (hexanes/EtOAc/ NET_3 50 : 50 : 1 and DCM/EtOAc 70 : 30) to give **49** (18.8 mg, 0.0419 mmol, 11%) as a white solid; m.p. 219 °C; IR (ATR) $\tilde{\nu}/\text{cm}^{-1}$ 2961, 1921, 2852, 1691, 1580, 1533, 1434, 1368, 1341, 1297, 1259, 1231, 1168, 1187, 1168, 1134, 1014, 972, 886, 860, 794, 762, 751, 711; δ_{H} (500 MHz; CDCl_3) 11.43 (bs, 1H, NHCO), 10.02 (s, 1H, CHO), 8.45 (s, 1H, 8-H), 7.90 (d, $J = 8.5$ Hz, 1H, 5-H), 7.84 (dd, $J = 8.5, 1.5$ Hz, 1H, 6-H), 7.78 (d, $J = 8.2$ Hz, 1H, 4-H), 7.52 (dd, $J = 8.3, 7.0$ Hz, 1H, 3-H), 7.44 (d, $J = 7.1$ Hz, 1H, 2-H), 7.01 (s, 1H, 4'-H), 6.76 (s, 1H, 5''-H), 4.53 (s, 2H, CH_2), 3.76 (s, 2H, CH_2S), 2.40 (s, 6H, 4- CH_3 , 6- CH_3). δ_{C} (126 MHz; CDCl_3) 192.53 (CHO), 170.05 (C-2''), 168.49 (C-4'', C-6''), 167.65 (NHCO), 157.19 (C-2'), 137.92 (C-1), 137.47 (C-4a), 135.61 (C-4'), 134.64 (C-7), 131.92 (C-5'), 131.41 (C-8a), 130.54 (C-8), 130.35 (C-5), 129.29 (C-3), 128.39 (C-4), 128.17 (C-2), 123.02 (C-6), 117.24 (C-5''), 34.71 (CH_2S), 30.74 (CH_2), 23.94 (4- CH_3 , 6- CH_3); HRMS (ESI): calcd.: 417.0920; found: $\{m/z: [\text{M} + \text{Na}]^+ 417.0919\}$; purity (HPLC): 210 nm: >95%; 254 nm: >95% (method 3b).

Biology

Sirtuin assays. The determination of inhibitory activity on the corresponding sirtuin enzymes was carried out based on



a fluorescence-based assay commissioned by Reaction Biology Corporation (Malvern, USA) following an internal experimental protocol. Test substances were prepared in DMSO, delivered into the respective enzyme mixture (prepared with reaction buffer: Tris-HCl, pH = 8) and incubated for 10 minutes at 30 °C. Subsequently the substrate mixture (NAD⁺ and a 7-amino-4-methylcoumarin-based fluorogenic peptide substrate) was added to initiate the deacetylation reaction. After 2 hours incubation at 30 °C, 2 mM nicotinamide (universal sirtuin inhibitor, to stop the reaction) and protease-based developer (to generate fluorescence by the cleavage of 7-amino-4-methylcoumarin) was added. At 30 °C 1 hour later, the respective fluorescence was measured (extinction/emission = 360 nm/460 nm). The inhibitory effect of the test compounds is indirectly proportional to the amount of converted fluorescent substrate standardized as 100% activity of the control without inhibitor. The test substances were tested in 10-dose IC₅₀ triplicate mode, with 3-fold serial dilution starting at 50 μM final reaction concentration and if necessary, starting at 100 μM. For each serially diluted replicate of the triplicate, an IC₅₀ value was determined by sigmoidal curve fitting, resulting in three IC₅₀ values, from which the mean and corresponding standard deviation were subsequently calculated. Data processing was performed based on Prism 8.0.2 software (GraphPad Software, Boston, USA). Subtype selectivity on SIRT1, SIRT3 and SIRT5 was evaluated in a single dose duplicate mode at 50 μM final test compound concentration according to previous outlined internal protocol of Reaction Biology Corporation by determining the enzyme activity of the respective sirtuins in % (no inhibitor control as 100% activity).

Fluorescence thermal shift assays. Fluorescence thermal shift assays were performed in white 96-well plates (Hard-Shell PCR Plates, BioRad, USA) with a total volume of 20 μL per well and a final DMSO concentration of 5% (v/v), following a published protocol.^{19,21} Briefly, 10 μL SIRT2 (final concentration of 6.0 μM, purified according to published protocol²¹) and SYPRO Orange (5× final concentration, Sigma-Aldrich, Germany) in assay buffer (25 mM Tris-HCl, 150 mM NaCl, 1 mM DTT, pH 8.0) were mixed with 10 μL of compound (final concentration of 30 μM and 10 μM) and incubated at 25 °C and 350 rpm for 5 min. The compounds were prepared as 10 mM stock solutions in DMSO and diluted in the assay such that a final DMSO concentration of 5% (v/v) was achieved. If necessary, NAD⁺ was added to a final concentration of 2.5 mM. Fluorescence intensity was recorded during a temperature gradient of 1 °C per 20 s from 25 to 95 °C using a real-time PCR machine (C1000 Touch™ Thermal Cycler, CFX96™ Real-Time System, BioRad, USA). Melting temperatures were determined using GraphPad Prism, following a published procedure.⁵²

Cloning and expression of SIRT2 56-356. The human *SIRT2* 2-389 gene (Table S4) was synthesized and cloned in a pETDuet based vector encoding an N-terminal His₆-SUMO tag by Eurofins Genomics. This construct served for Q5

mutagenesis with primers A-D (Table S5) to yield His₆-SUMO-Ser-SIRT2 56-356. All plasmids were controlled by sequencing. *SIRT2* 56-356 was expressed as an N-terminal His₆-SUMO fusion in Rosetta(DE3) cells. Overnight cultures were grown in LB with ampicillin (180 mg L⁻¹) and chloramphenicol (50 mg L⁻¹) at 37 °C, 130 rpm. Expression cultures were inoculated with 25 mL overnight culture per liter autoinduction medium from Studier⁵³ (adapted according to Table 2) with the antibiotics ampicillin (90 mg L⁻¹) and chloramphenicol (25 mg L⁻¹) added. After 18 h at 37 °C, 130 rpm, the cells were cultured for 24 h at 16 °C, 130 rpm for protein expression. Cells were harvested by centrifugation for 20 min at 5000 × g, 20 °C, and washed with 0.9% (w/v) NaCl. The supernatant was removed by centrifugation for 20 min at 5000 × g, 4 °C, and the cell pellets were stored at -20 °C.

Protein purification of SIRT2 56-356. Cell pellets were resuspended on ice in 100 mM HEPES pH 7.0, 300 mM NaCl, 20 mM imidazole, 2 mM β-mercaptoethanol, 1 mg of DNase I, and Pefabloc® SC. After lysis by sonication, the cell debris was removed by centrifugation for 20 min at 41 000 × g, 4 °C. The lysate was loaded onto a NiSepharose column (HisTrap™ HP 5 mL, Cytiva) previously equilibrated with buffer A (100 mM HEPES pH 7.0, 300 mM NaCl, 20 mM imidazole, 2 mM β-mercaptoethanol) by an Äkta pure™ System (GE Healthcare) at 4 °C. The column was washed with 10 CV of buffer A, and SIRT2 was eluted by a continuous gradient from 0 to 100% buffer B (100 mM HEPES pH 7.0, 300 mM NaCl, 500 mM imidazole, 2 mM β-mercaptoethanol). All purification steps were monitored by UV absorption at 280 nm and SDS-PAGE. To remove the His₆-SUMO-tag, the fractions containing SIRT2 were dialyzed overnight at 4 °C with His₆-SUMO-protease against 2 L of 50 mM HEPES pH 7.0, 150 mM NaCl, 2 mM β-mercaptoethanol. Precipitated protein was removed by centrifugation, and the supernatant was subjected to a second Ni²⁺ affinity chromatography. The flow-through containing cleaved SIRT2 was collected and concentrated to a volume of 2 mL using an Amicon® centrifugal filter (Merck) with 10 000 MWCO. The protein solution was loaded on a HiLoad® 16/600 Superdex® 200 pg column (Cytiva) at 4 °C using 25 mM HEPES pH 7.0, 100 mM NaCl, 2 mM DTT as running buffer. Fractions containing SIRT2 were pooled and concentrated using Amicon® 10 000

Table 2 Composition of autoinduction medium

Content [g L ⁻¹]	Compound
5.0	g L ⁻¹ Yeast extract
10	g L ⁻¹ Peptone
0.5	g L ⁻¹ Glucose
1.9	g L ⁻¹ Lactose
5.0	g L ⁻¹ Glycerol
0.2	g L ⁻¹ Magnesium sulfate
3.4	g L ⁻¹ Potassium dihydrogen phosphate
3.5	g L ⁻¹ Disodium hydrogen phosphate
2.8	g L ⁻¹ Ammonium chloride
pH adjusted to	pH 6.8



MWCO centrifugal filters (Merck). The protein was frozen in liquid nitrogen and stored at $-80\text{ }^{\circ}\text{C}$ until further use.

Crystallisation and X-ray structure determination of SIRT2 56-356. SIRT2 56-356 (23.7 mg mL^{-1}) was crystallized without ligand by sitting drop vapor diffusion at $20\text{ }^{\circ}\text{C}$. A 2:1 ratio of protein and reservoir solution containing 0.2 M ammonium formate, 20% (w/v) PEG3350 led to crystals within two months. Crystals were cryoprotected with 35% (v/v) glycerol and flash frozen in liquid nitrogen.

Crystal structures of SIRT2 56-356 with ligands were obtained from hanging drop crystallization experiments at $20\text{ }^{\circ}\text{C}$. The SIRT2 56-356 protein was diluted to 7.3 mg mL^{-1} in 25 mM HEPES pH 7.0, 100 mM NaCl, 2 mM DTT, and mixed with 5 mM NAD^{+} and 500 μM of **29 (RW-78)/31 (RW-80)** dissolved in DMSO, respectively. The solution was incubated for 1 h on ice, and any precipitate was removed by centrifugation for 10 min at $16000 \times g$. Crystals grew after 5 days in 24-well, with 300 μL of 0.1 M sodium acetate, 17% (w/v) PEG3350 as reservoir, and a drop ratio of 1 μL protein to 1 μL reservoir solution. The crystals were soaked with 10 μL of 30% (v/v) ethylene glycol and vitrified in liquid nitrogen. Diffraction images of SIRT2 56-356:**RW-78**, and SIRT2 56-356:**RW-80** were recorded using synchrotron radiation of $\lambda = 1.060\text{ \AA}$ at the P13 beamline from PETRAIII at DESY (Deutsches Elektronen-Synchrotron, EMBL, Hamburg, Germany). Recorded reflections were processed using the XDS suite, and data reduction was performed with XSCALE.^{54,55} Phasing of the SIRT2 56-356 apo structure was performed by molecular replacement with PHASER⁵⁶ and the coordinates of SIRT2:**SirReal2** (PDB ID: 4RMG¹⁹). The solution was refined by restrained refinement with REFMAC5 (ref. 57) and iteratively rebuilt with COOT (v. 0.9).⁵⁸ The resulting model was used to phase the SIRT2 56-356:**RW-78** data set, followed by iterating model building in COOT⁵⁸ with restrained refinements in REFMAC5.⁵⁷ The ligand was prepared in AceDRG.⁵⁹ Water molecules were positioned with ARP/wARP 8.0.⁶⁰ The apo structure was completed by TLS refinement, whereas the SIRT2 56-356:**RW-78** structure was refined with anisotropic restraints to satisfactory R_{work} and R_{free} values. The SIRT2 56-356:**RW-80** structure was phased with the SIRT2:**RW-78** structure in REFMAC5 (ref. 57) and built as described for SIRT2 56-356:**RW-78**. The geometry of the final structures was analyzed by the MOLPROBITY⁶¹ online tool, and the structures were deposited in the RCSB Protein Data Bank (Tables S1–S3).

NanoBRET assay. The NanoBRET target engagement assay was performed as previously described by Vogelmann *et al.*,²¹ with the modification that stably transfected HEK293T cells expressing the NanoLuc-tagged SIRT2(50-356) fusion protein were used instead of transient transfection. HEK293T-NLuc-Sirt250-356 cells were cultured under standard conditions ($37\text{ }^{\circ}\text{C}$, 5% CO_2) in DMEM supplemented with 10% fetal bovine serum, 2 mM glutamine, and 300 $\mu\text{g mL}^{-1}$ hygromycin as a selection

marker. For the assay, cells were trypsinized, resuspended in Opti-MEM® reduced serum medium, and adjusted to a concentration of 2×10^5 cells per mL. To determine the affinities of the inhibitors, a final tracer concentration of 2 μM was used. Serially diluted inhibitor and tracer were added to the cell suspension, and 100 μL were seeded into 96-well white, sterile, nonbinding surface plates. Plates were incubated at $37\text{ }^{\circ}\text{C}$ with 5% CO_2 for 2 hours. For BRET measurements, NanoBRET NanoGlo Substrate (Promega cat. #N1571) was added to the wells according to the manufacturer's protocol. All measurements were performed using the 2102 EnVision™ Multilabel Reader (PerkinElmer), equipped with a 460 nm filter (donor) and a 615 nm filter (acceptor). The BRET ratio was calculated as the ratio of acceptor to donor signal, and data were normalized to vehicle-treated controls. Apparent intracellular binding affinities (EC_{50} values) were determined by fitting dose-response curves using nonlinear regression analysis in GraphPad Prism. All experiments were conducted in triplicate unless otherwise noted, and results are presented as mean \pm standard deviation.

Computational methods

Docking simulations. Docking simulations were carried out with Schrödinger software suite (Schrödinger Inc., New York City, USA, version 2020-3).⁶² Crystal structures of SIRT2 and respective lead structures were imported from the Protein Data Bank (PDB)⁶³ (**SirReal2**: PDB ID: 4RMG;¹⁹ **24a**: PDB ID: 5YQO²³) and prepared with the Protein Preparation Wizard (Schrödinger Inc. New York City, USA). All ligands were prepared with the Ligand Preparation Wizard using Epik for protonation and charge calculations.⁶⁴ Docking was performed using Glide in standard precision mode SP (all docking parameters left to their default values). Results were inspected and visualized with PyMOL 2.5.8 (Schrödinger Inc., New York city, USA). The top-ranking poses were analysed, considering the favourable spatial orientation in relation to the crystal structures of the corresponding lead compounds.

Author contributions

This study was conceptualised and supervised by F. B. Investigation, methodology, experimental execution and data curation were performed by R. W., M. F., A. H., N. P., F. F., T. W., M. J., M. G. and E. M. H. Original manuscript draft was written and edited by R. W. and M. F. with inputs from A. H., E. M. H. and F. B. All authors partook in the discussion of the findings and thoroughly reviewed the final manuscript for publication.

Conflicts of interest

There are no conflicts to declare.



Data availability

Supplementary information is available: ^1H and ^{13}C NMR spectra of compounds, HPLC chromatograms of tested compounds, Tables S1–S5: Crystallographic data, Fig. S1: Purification of human SIRT2 56-356 and Fig. S2: Close-up view of the SIRT2 ligand binding site can be found in the SI. See DOI: <https://doi.org/10.1039/D5MD00144G>.

The data supporting this article have been included as part of the SI. X-ray coordinates have been deposited in the RCSB Protein Data Bank under the accession codes 9S44 (SIRT2 56-356), 9S46 (SIRT2 56-356:RW-78) and 9S48 (SIRT2 56-356:RW-80).

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