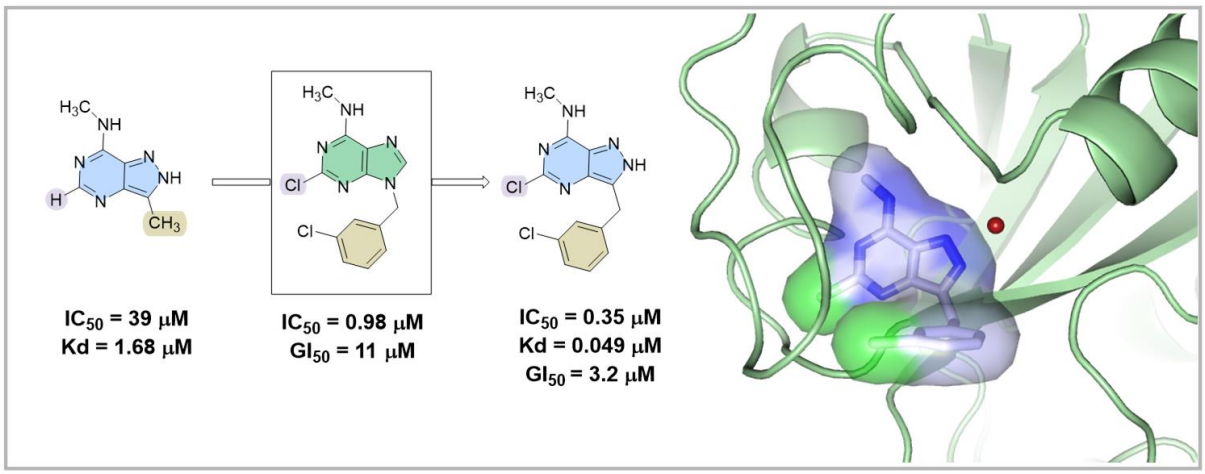
Structure-based design of a potent and selective YTHDC1 ligand

*František Zálešák, Francesco Nai, Rajiv K. Bedi, Yaozong Li, Marcin Herok, Elena Bochenkova, Francesco Errani, Amedeo Caflisch**

Department of Biochemistry, University of Zurich, Winterthurerstrasse 190, CH-8057, Zurich, Switzerland

N^6-Adenosine methylation (m6A) is a prevalent post-transcriptional modification of mRNA, with YTHDC1 being the reader protein responsible for recognizing this modification in the nucleus of the cell. Here we present a protein structure-based medicinal chemistry campaign that resulted in the YTHDC1 inhibitor **40** which shows an equilibrium dissociation constant (K_d) of 49 nM. The crystal structure of the complex (1.6-Å resolution) validated the design. Compound **40** is selective against the cytoplasmic m6A-RNA readers YTHDF1-3 and YTHDC2 and shows antiproliferative activity against the acute myeloid leukemia (AML) cell lines THP-1, MOLM-13, and NOMO1. For the series of compounds that culminated into ligand **40**, the good correlation between the affinity in the biochemical assay and antiproliferative activity in the cellular assay (THP-1) provides evidence of YTHDC1 target engagement in the cell. Thus compound **40** meets chemical probe properties for studying the role of YTHDC1 in AML.



Introduction. Post-transcriptional changes of eukaryotic messenger RNA (mRNA) play a pivotal role in cellular processes.[1] Before leaving the nucleus, mRNA undergoes a series of chemical alterations, such as methylation, acetylation, and splicing.[2,3] These modifications directly affect mRNA stability, processing, and translation efficiency.[4,5] Recent investigation of post-transcriptional events have given rise to a dynamic research field known as epitranscriptomics.[3] Among the diverse mRNA modifications in the human transcriptome, the most prevalent and extensively studied is the N^6 methylation of adenosine (m6A).[6] This reversible process is facilitated by proteins referred to as "writers", such as METTL3/14,[7,8] while the demethylation is achieved by "erasers" like FTO or ALKBH5.[9,10] Our research group has made recent contributions to this field through the development of small-molecule inhibitors targeting METTL3/14, demonstrating the modulation of interconnected cellular events.[11,12]

Recognition of the m6A modification is mediated by "readers" that subsequently impact downstream processes.[13] The YTH family, comprising five proteins (YTHDC1, YTHDC2, and YTHDF1-3), is the best characterized family of m6A readers.[14] Due to their crucial role in diverse biological processes, these proteins hold great promise as therapeutical targets.[15,16] Predominantly localized in the nucleus, YTHDC1 is responsible for the regulation of pre-mRNA splicing and

mRNA export from the nucleus.[17] A growing number of reports associate the activity of YTHDC1 with various biological functions, including embryonic development, neuronal development, and others.[18] Furthermore, its critical role has been identified in various types of cancer.[19,20]

Our group recently reported fragments selectively binding to YTHDC1 or YTHDF2, respectively.[21,22] Other studies in the literature have reported inhibition using rather promiscuous binders providing limited selectivity towards proteins of the YTH family,[23–25] except for a recent report which describes a selective YTHDC1 inhibitor identified by *in vitro* high-throughput screening followed by a structure-based optimization.[26]

In this study, we present a structure-based design campaign aimed at developing a potent and selective ligand of YTHDC1. Additionally, we provide a biochemical evaluation (HTRF assay, ITC, TSA, X-Ray) and report a direct connection between YTHDC1 inhibition and the effect on the cell proliferation of acute monocytic leukemia (THP-1, MOLM-13, NOMO1).

Results and discussion. The present medicinal chemistry campaign builds upon our initial *in silico* screening for YTHDC1.[21,27] The structure-based medicinal chemistry optimization started here with the ligand-efficient fragment **1** (Figure 1C) which had been characterized by biochemical assays and crystallography.[21] Structurally, this fragment contains a pyrazolo[4,3-d]pyrimidine core that mimics the natural ligand – m^6adenine (we use m^6adenine for nucleotide, in contrast to m^6A for nucleoside). The recognition of fragment **1** is achieved by the aromatic cage consisting of two tryptophan residues (Trp428, Trp377) and five hydrogen bonds between the pyrazolopyrimidine ring and the binding pocket (Asn367, Asn363, Asp476, backbone Ser378, and structural water - bridging sidechains of Trp377 and Asp476). Notably, unlike the natural ligand, the pyrazolopyrimidine **1** forms an additional hydrogen bond with the sidechain of Asp476 (Figure 1, PDB: 7P8F). This interaction significantly enhances the YTHDC1 affinity of fragment **1** by nearly

10-fold compared to m6-adenine (IC_{50} = 39 uM vs 306 uM).[21] Upon evaluating the binding pose of pyrazolopyrimidine **1** within the binding pocket, we identified two positions on the aromatic core that are conducive to ligand growth and optimization. The first position lies between N^6 and N^4 of the pyrimidine ring. The substituent in this position leads to a small pocket, suggesting that a suitable substituent could occupy the space and enhance binding. The second substituent leads from C^3 towards a shallow, positively charged pocket that binds the negatively charged part of RNA in the natural ligand.

During the initial stages of the campaign, we identified that a chloride substituent located between N^6 and N^4 fits into the aforementioned small pocket, notably improving the binding affinity. (Please note the difference in the numbering of purine and pyrazolopyrimidine core as shown in Figure 1C). This improvement was demonstrated by comparing the potency of non-chlorinated **2a** and chlorinated m^6adenine **2b** resulting in IC_{50} values of 306 μM and 37 μM, respectively. However, the adenine aromatic core cannot form a conventional hydrogen bond with Asp476, unlike pyrazolopyrimidine **1** moiety. Therefore, we decided to merge the two structural features and synthesize 5-chloropyrazolopyrimidine fragment **3**. This compound exhibited improved potency together with ligand efficiency (IC_{50} = 2 uM, LE = 0.60) and sub-micromolar equilibrium dissociation constant (K_D = 146 nM). As mentioned above, an alternative position for ligand growth involved extending or replacing the methyl substituent at position 3 of 5-chloropyrazolopyrimidine **3**. Considering the synthetic challenges, we decided to pursue the optimization using 2-chloropurine **2b** instead of 5-chloropyrazolopyrimidine **3**. This approach offered the advantage of convenient derivatization at N^9 of the purine moiety, as compared to C^3 in the equivalent position of the pyrazolopyrimidine core. The primary rationale behind this choice was to focus on extending the purine scaffold and synthesizing only the most promising examples

in combination with the 5-chloropyrazolopyrimidine **3**. This strategy was based on the observation of essentially identical binding poses among the fragments (See Figure 1B) and the hypothesis that interactions outside the aromatic cage would be preserved across different fragments.

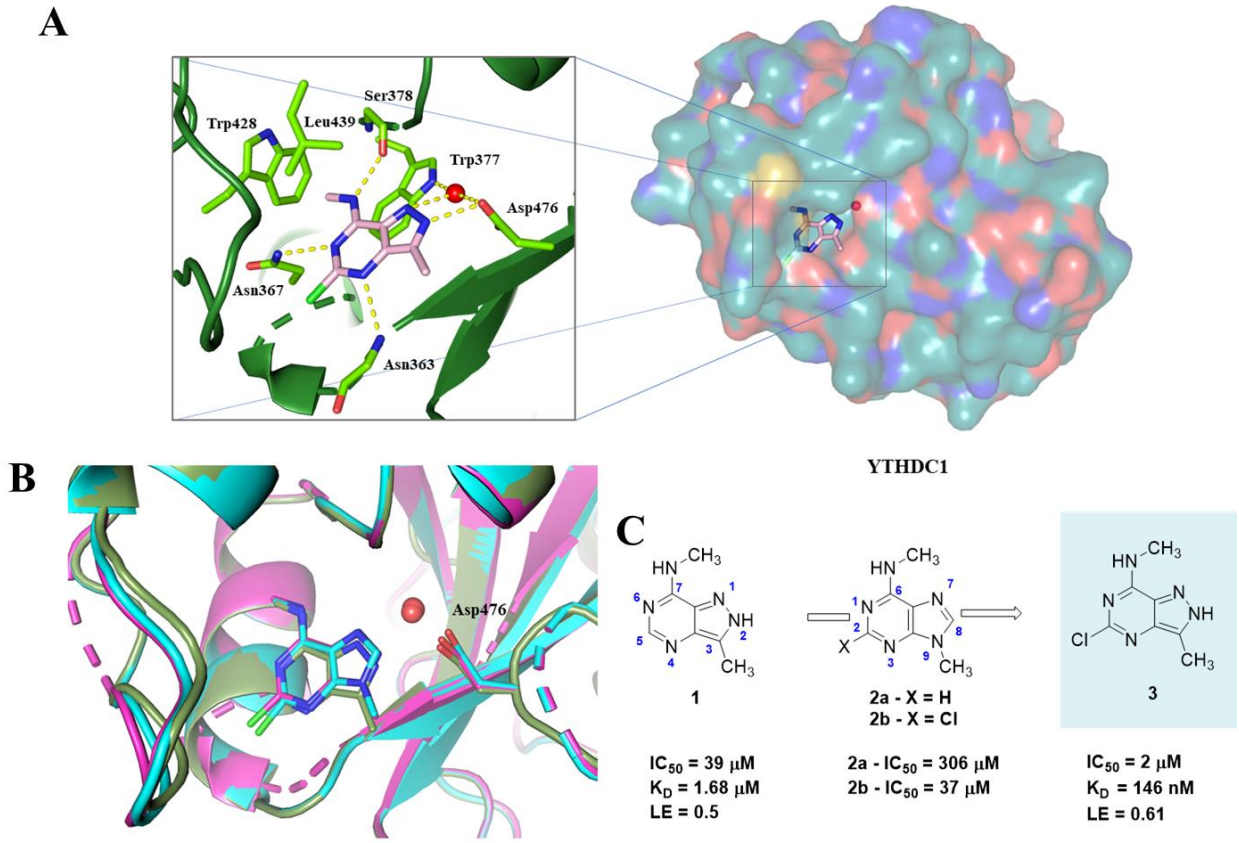
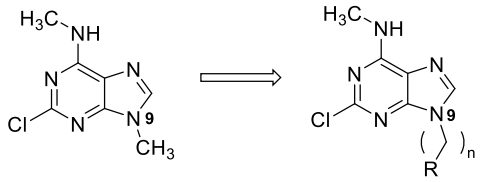


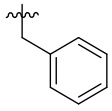
Figure 1. (A) Crystal structure of fragment **3** in the YTHDC1 binding site with relevant residues (light green). (B) Structural overlap of fragments **1-3** in the binding site of YTHDC1 (Compound **1** – smudge, Compound **2b** – cyan, Compound **3** – magenta). (C) 2D structures of fragments **1-3**. Fragment **1** is the starting point of the optimization.[21]

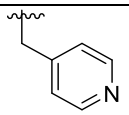
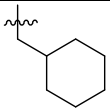
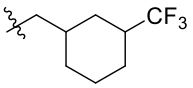
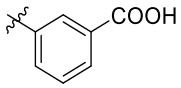
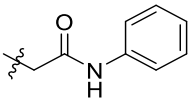
a) Ligand growing – First round of optimization. In the initial screening phase, we opted to replace/extend the methyl group at position 9 with various aliphatic and aromatic rings, including both -CH_2- bridged and non-bridged structures (Table 1). Notably, the substituents with a

methylene linker (**5**, **6**, **8**, **9,** and **11**) exhibited improved binding leading to low micromolar potency (IC_{50} = 3, 6, 8, 11, and 9 µM, respectively). These results suggest the presence of van der Waals interaction between the rings and lipophilic residues (Leu380, Pro431, Met434). The flexibility provided by the methylene allows the aromatic ring to achieve better interactions within the binding site residues, which is not the case for compounds **4** and **7** (Figures 2A and 2B). However, there was one exemption among the tested molecules. Compound **10**, which features a carboxylic acid moiety and displayed enhanced potency (IC_{50} = 11 µM). This observation indicates that the absence of hydrophobic interactions is compensated with ion-ion or dipole-ion interactions. Such interactions are likely to be formed between negatively charged carboxyl group and positively charged residues within the binding pocket. The significant impact of the carboxylic group is evident when comparing the potency of compound **4**, featuring a non-substituted phenyl ring to its carboxylic acid derivative **10**.

Table 1. 2-Chloropurine derivatives with different substituents on N^9



Compound number	R	IC_{50}[a] (µM)	LE[b]	LLE[c]	GI_{50}[d] (THP-1, µM)	PDB Code Resolution (Å)
4		>100	-	-	-	8Q2S (1.41)
5		3	0.40	3.2	31	8Q2T (1.40)

6		6	0.8	3.9	81	8Q2U (1.36)
7		47	0.33	2.2	-	8Q2V (1.71)
8		8	0.37	2.5	-	8Q2W (1.41)
9		11	0.30	1.6	-	-
10		11	0.32	3.2	-	8Q2X (1.60)
11		9	0.31	3.4	-	8Q2Y (1.71)

[a] Homogeneous time-resolved fluorescence (HTRF) [b] Ligand efficiency (kcal·mol^{-1}·heavy atom count^{-1})[28]. [c] Lipophilic ligand efficiency (pIC50 − clogP)[28].[d] Growth inhibition 50 (GI_{50}) values after 72h treatment (THP-1).

b) Second round of optimization - Benzyl rings SAR study. Due to its promising potency, the benzyl substituent was retained for further optimization. The easily accessible and commercially available nature of benzyl halides facilitated an efficient structure-activity relationship (SAR) study, enabling identifications of potent functionalization. The benzylic scaffolds we synthesized and tested could be divided into subgroups based on the position and number of substituents (Table 2). In the following, *ortho, meta,* and *para* positions refer to the substitution relative to the methylene connecting the purine heterocycle.

Among the *ortho*-substituted compounds, improved potency was exhibited by compound **12** with free amine moiety and its sulfonamide derivatives (**13, 14**) with IC_{50} values of 9, 0.46, and 2 μM, respectively. The potency of compound **13** could be potentially attributed to an additional hydrogen bond formed between the sulfonamide oxygen and the backbone N-H of Asp476 (Figure

2C). On the other hand, the presence of trifluoroacetic amide **15** and methoxy derivative **18** provided the IC_{50} value above 100 μM and 13 μM, respectively. This suggested that the carbonyl of **15** is unable to adopt the favorable geometry orientation for hydrogen bond formation observed between the sulfonamide oxygen of **13** and Asp476. Although perfluorinated alkyl substituents (**16**, **17**) exhibited lower potency in comparison with the most promising compound from this set (IC_{50} = 2 μM and 3 μM versus 0.46 μM), compound **16** provided the best results against THP-1 cell line (GI_{50} = 14 μM). Additionally, compounds **12, 13, 15, 16,** and **18** were soaked to YTHDC1 enabling X-ray confirmation of their binding poses with high resolution (< 1.5 Å).

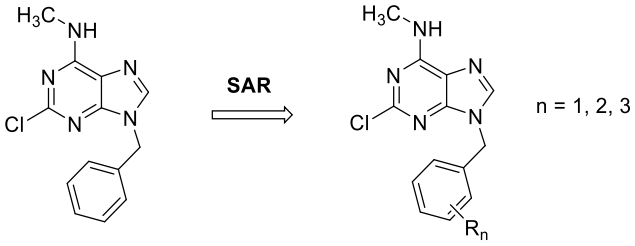
Among the *meta*-substituted derivatives, our focus primarily centered on carboxylic acid derivatives, as encouraged by the aforementioned significant potency increase between compounds **4** and **10**. To expand structural and functional diversity, we also evaluated purine analogs containing halogen and methoxy groups. As expected, *meta*-carboxylic acid analog **20** demonstrated the highest affinity with an IC_{50} value of 0.51 μM. In contrast, the tetrazole heterocycle **21,** serving as carboxylic acid bioisoster, displayed lower affinity (IC_{50} = 1 μM) compared to the free carboxylic acid compound **20**. Conversely, *meta*-substituted compounds with amide (**22, 23**) and methyl ester moiety **19** exhibited compared affinity (IC_{50} = 2, 1, 1 μM) compared to nitrile group **26** (IC_{50} = 2 μM**)** and did not surpass the potency of compound **20**. We hypothesized that the improved binding towards YTHDC1 might be a result of ion-ion or ion–dipole interaction involving the positively charged side chain of Arg475. However, X-Ray crystallography could not confirm this, due to the lack of density for both the carboxylic group and the Arg475 side chain. Regarding the antiproliferative activity against THP-1, the methyl ester derivative **19** exhibited low micromolar GI_{50} (6.2 μM) while carboxylic acid derivative **20**, which showed greater potency in the HTRF assay, had a minimal effect (GI_{50} > 100 μM), likely due to

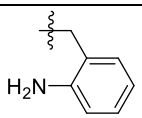
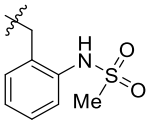
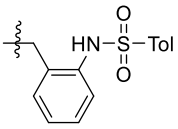
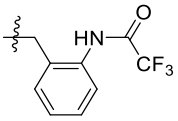
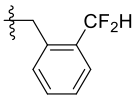
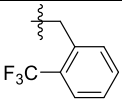
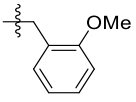
poor cell permeation of the charged carboxylic residue at physiological pH. Among the compounds tested, we also discovered that a chloride substituent in the *meta* position exhibited enhanced binding with an IC_{50} value of 0.96 μM (compound **25**) and displayed antiproliferative activity against THP-1 (GI_{50} = 11 μM). This compound also exhibited a very favorable LE value (0.41). The improved affinity could potentially arise from the formation of a halogen bond. X-ray crystallography indeed provided evidence for an interaction between the chlorine substituent and the backbone carbonyl of Val432 (Figure 2C). On the other hand, a bigger and more lipophilic bromide substituent present in compound **24** resulted in a 2-fold decrease in potency (IC_{50} = 2 μM) in comparison to compound **25**. Because of the positive effect observed with carboxylic acid functionality, we also prepared derivatives of carboxylic acid (**28**, **29**) and benzyl alcohol derivative **30** in the *para* position. However, these compounds did not exhibit improved binding compared to the ones with functional groups in *meta* position, presumably due to an unfavorable orientation of the substituents that lead towards solvent exposed area out of the pocket.

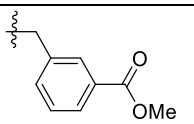
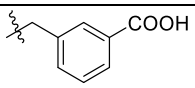
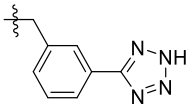
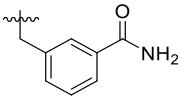
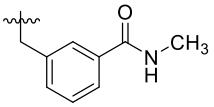
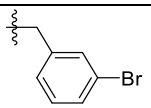
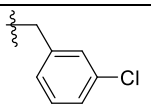
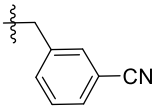
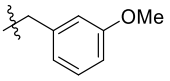
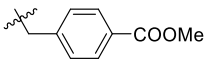
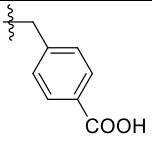
The combination of the most potent aromatic ring substitution, namely methylsulfonamide, in *ortho* (Compound **13**) and chloride in *meta* (Compound **31**) position did not substantially improve potency with respect to the monosubstituted analogs. Although Compound **31** showed an improved biochemical potency (IC_{50} = 0.45 μM), the presence of sulfonamide moiety once again deteriorated its activity against THP-1 growth (GI_{50} = 22 μM). On the other hand, compound **32** containing *meta*-chloro and *meta*-carboxyl substitution showed greater enhancement in binding resulting in the most potent compound (IC_{50} = 0.18 μM). Unfortunately, the biochemical potency increase was not transferred into the improved antiproliferative activity (GI_{50} > 100 μM) which is also true for its methyl ester analog **33** (GI_{50} = 9.6 μM). Furthermore, while the combination of *para*-methoxy and *meta*-chloro substitution provided sub-micromolar potency (Compound **34**, 0.59 μM), the

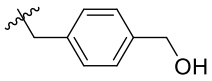
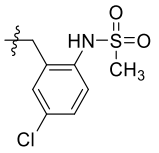
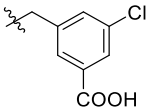
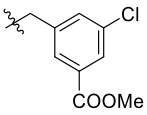
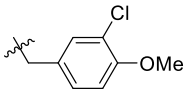
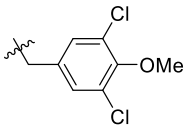
presence of two chloro substituents in meta position diminished the binding (Compound **35**, IC_{50} = 2 μM).

Table 2. Structure-activity relationship – benzyl ring substitutions



Compound number	R^1	IC_{50} (μM)	LE	LLE	GI_{50} (THP-1)	PDB Code (Resolution - Å)
Monosubstitution (*ortho*)						
12		9	0.35	3.4	-	8Q31 (1.32)
13		0.46	0.36	5.0	22	8Q32 (1.43)
14		2	0.26	2.7	-	-
15		> 100	-	-	-	8Q33 (1.43)
16		2	0.36	2.7	14	8Q35 (1.31)
17		3	0.33	2.4	-	-
18		13	0.32	2.6	-	8Q37 (1.28)

Monosubstitution (*meta*)						
19		1	0.36	3.8	6.2	8Q38 (1.42)[e]
20		0.51	0.39	4.7	>100	-
21		1	0.34	4.4	-	8Q39 (1.42)
22		2	0.36	4.3	-	8Q3A (1.43)
23		1	0.36	4.2	31	8Q3G (1.42)[e]
24		2	0.39	2.7	-	-
25		0.96	0.41	3.1	11	8Q4M (1.43)
26		2	0.37	3.5	-	8Q4N (1.42)[e]
27		4	0.35	3.2	-	8Q4P (1.43)
Monosubstitution (para)						
28		0.97	0.36	3.8	9.5	-
29		0.87	0.38	4.4	-	8Q4Q (1.40)

30		2	0.37	4.0	23	8Q4R (1.43)
Di- and Trisubstitution						
31		0.45	0.35	4.4	22	8Q4T (1.51)
32		0.18	0.4	4.5	> 100	-
33		1	0.34	3.2	9.6	-
34		0.59	0.39	3.4	9.5	-
35		2	0.34	2.2	-	-

Same as Table 1. [e] No density beyond the phenyl ring

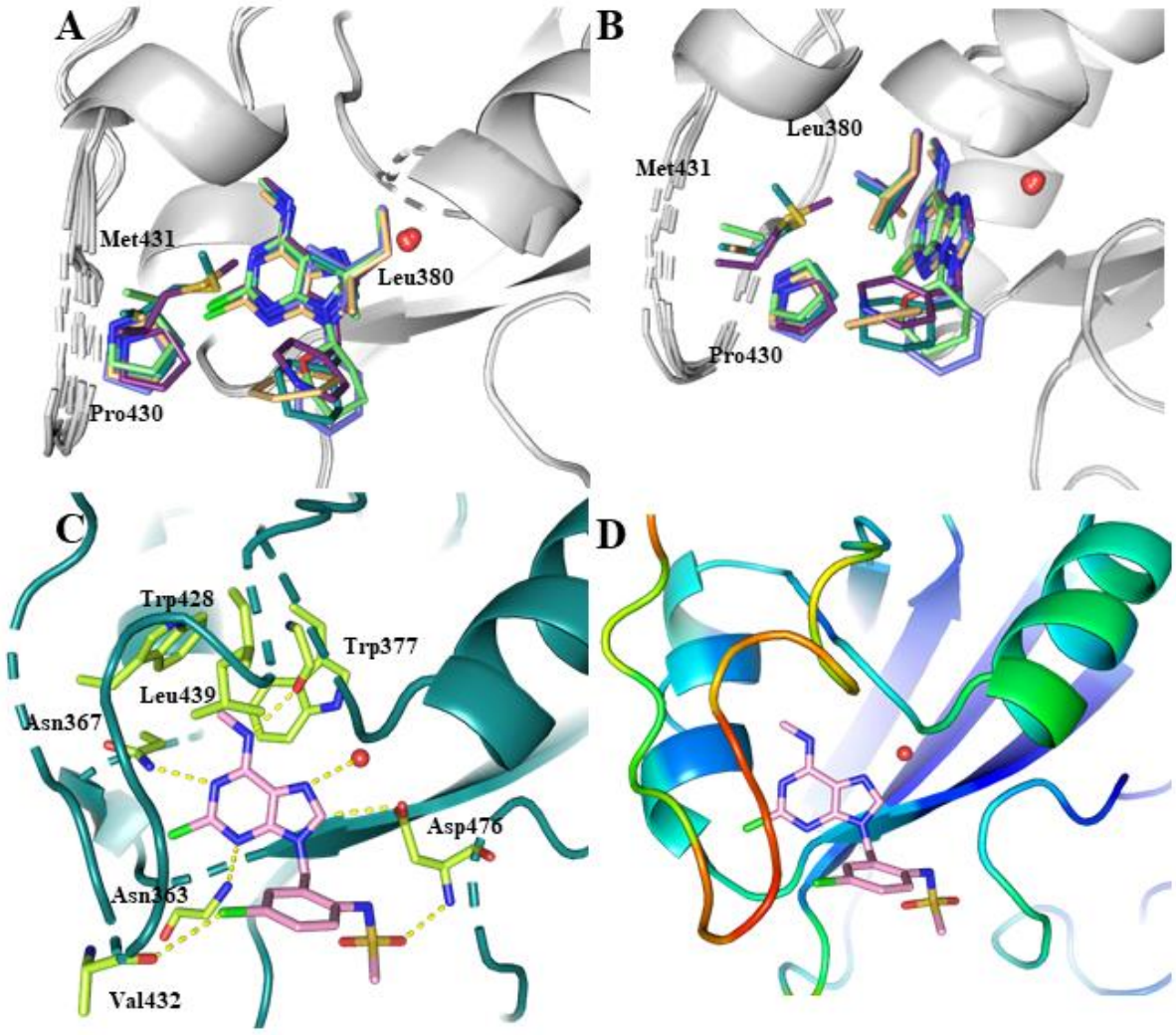


Figure 2. Interactions between ligands and YTHDC1 in crystal structures of the complexes**.** (A+B)**.** Structural overlap of the complexes with compounds **4** (slate), **5** (light orange), **6** (magenta), **7** (lime), **8** (deep teal). Compounds **5** and **6** have a methylene linker so that the phenyl and pyridine ring, respectively, form van der Waals interactions with the Pro430, Met431, and Leu380. (C) The binding pose of ligand **31** shows a halogen bond with the carbonyl oxygen of Val432 (which is also observed for **25**) and a hydrogen bond towards the backbone NH of Asp476 (also for **13**). (D) The protein backbone is colored according to the crystallographic B-factor (disorder increasing from blue to red).

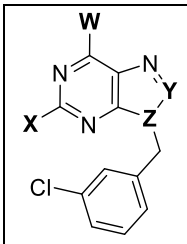
c) Headgroup optimization – After optimizing the benzylic sidechain and identifying suitable substituents, we shifted our focus back to the heterocyclic core optimization. For this reason, we explored further modifications on the purine fragment while maintaining the *meta*-chlorobenzyl substituent at N^9 (Table 3). Firstly, we confirmed that the presence of a hydrogen bond between the methylamino group and the backbone of Ser378 is essential for the binding. This was verified by testing compounds **38** and **39**, both containing a chlorine substituent at position 6 of the purine ring. The presence of chlorine atom instead of methylamino moiety resulted in a notable decrease in potency. Additionally, compound **36**, bearing fluoro substituent instead of chloro (position 2 of purine ring), pointing towards a small lipophilic pocket, also exhibited weakened binding (IC_{50} = 2 μM). Moreover, the substitution of the methylamino group with cyclopropyl amino moiety **37** also led to weakened binding, indicating that the aromatic cage of YTHDC1 is intolerant to increased bulkiness.

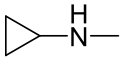
As outlined in our strategy, our primary objective was to combine the most promising (LE, GI_{50}) substitution pattern optimized on purine heterocycle with the 5-chloropyrazolopyrimidine core. Thus, we decided to obtain compound **40**, which combines 5-chloropyrazolopyrimidine and benzyl substituent in position 3. As anticipated, X-ray analysis confirmed the interactions between compound **40** and YTHDC1, which are the same as observed with the purine analog **25**, along with an additional hydrogen bond with Asp476 (Figure 3A). Furthermore, compound **40** exhibits an IC_{50} of 0.35 (Figure 3B), good LE (0.44), and antiproliferative activity against THP-1 (GI_{50} = 3.2 μM) MOLM-13 (5.6 μM) and NOMO1 (8.2 μM)(Figure 3, Figure S1). We further confirmed the binding of compound **40** to YTHDC1 by thermal shift assay (TSA), where the protein exhibited a thermal stabilization upon binding of pyrazolopyrimidine **40,** increasing the melting temperature by 12 °C at a compound concentration

of 100 µM (Figure 3C). Moreover, an equilibrium dissociation constant (K_D) of 49 nM was measured by isothermal titration calorimetry (ITC) for compound **40** and YTHDC1.

Next, we tested the selectivity of **40** against YTHDF1-3. Compound **40** displayed an IC_{50} value of 89 µM, 60 µM, and 83 µM against YTHDF1, YTHDF2, and YTHDF3, respectively (Figure S2) which is about 200-fold difference in binding preference towards YTHDC1. Finally, it is important to note that the correlation between biochemical IC_{50} values measured by the HTRF assay and the GI_{50} values (THP-1) provides evidence of engagement of the YTHDC1 target in the cell (Figure 3F).

Table 3. Additional optimization of the original fragment



Compound number	W	X	Y	Z	IC_{50}	LE	LLE	GI_{50}	PDB Code Resolution (Å)
36			C	N	2	0.39	3.2	-	8Q4U (1.37)
37			C	N	20	0.29	1.2	-	8Q4V (1.36)
38			C	N	>5[f]	-	-	-	-
39			C	N	>100	-	-	-	-
40[e]			N	C	0.35 K_d = 0.05	0.44	3.7	3.2	8Q4W (1.61)

Same as Table 1. For compound **40** the K_D determined by ITC is reported below the IC_{50} value. [e] Compound **40** was tested in the form of HCl salt [f] tested only in single dose at 5 micromolar (103%)

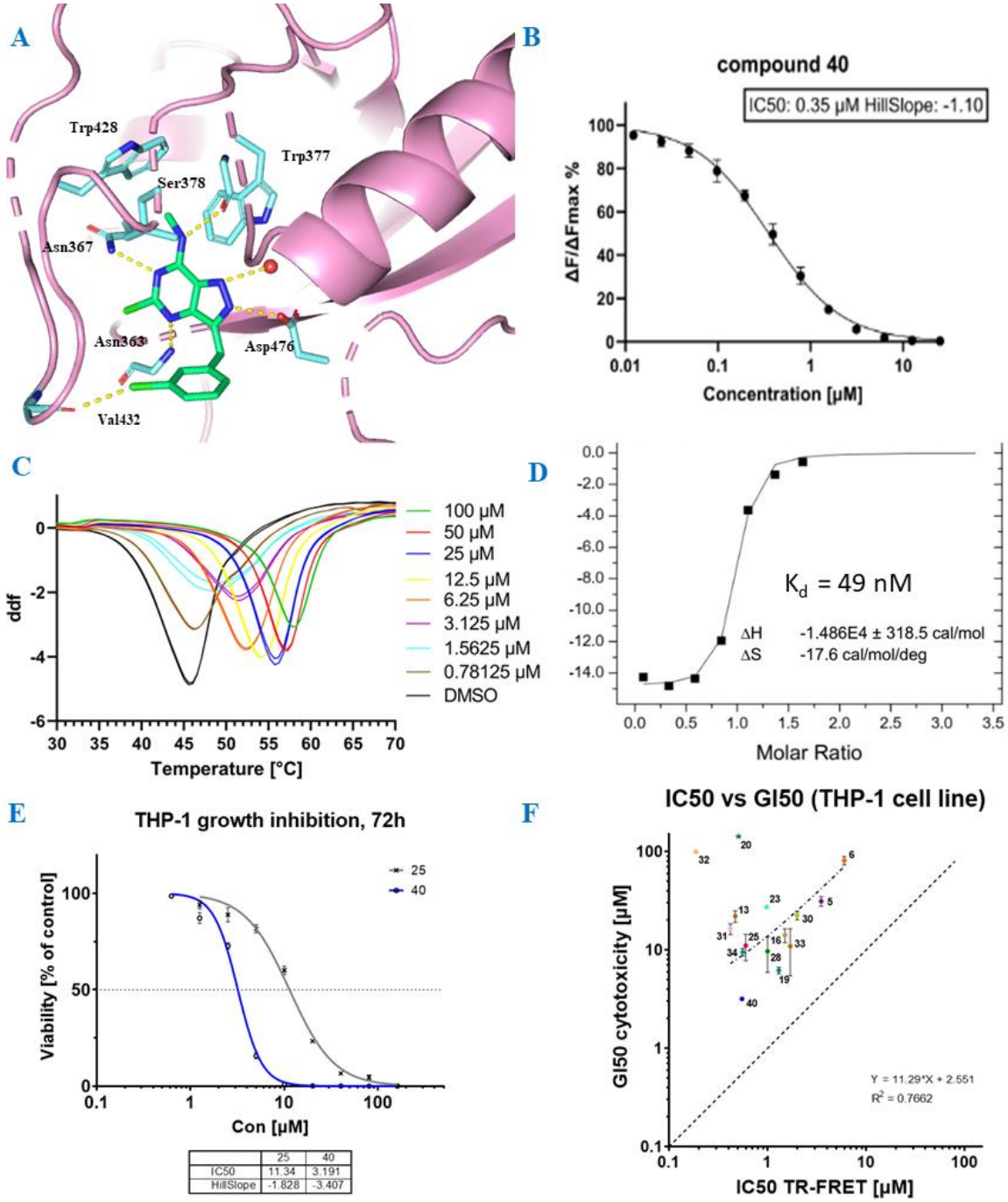
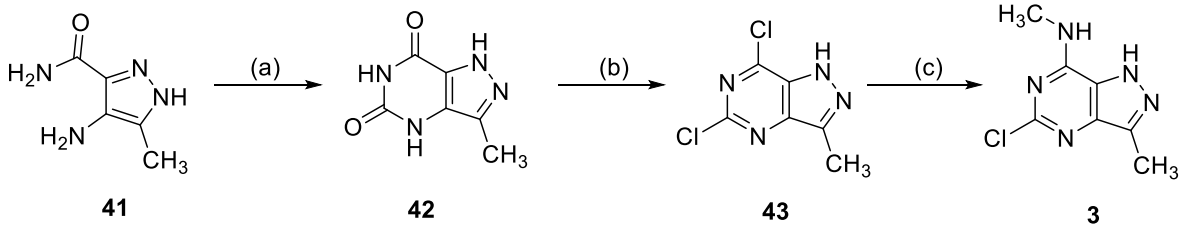


Figure 3. Biochemical and cellular evaluation of compound **40**. (A) Binding pose of compound **40** and interactions with YTHDC1 (PDB code 8Q4W). (B) HTRF dose-response curves of

YTHDC1 and compound **40**. (C) A dose-response thermal shift of YTHDC1 in the presence of compound **40**. (D) Isothermal titration calorimetry curve for YTHDC1 and compound **40**. (E) Dose-response curves for the antiproliferative effect of compounds **25** and **40** against THP-1 cell line. (F) Scatter plot of GI_{50} values for THP-1 vs. biochemical IC_{50} values which provides evidence of target engagement in the cell for this series of compounds. Note that compounds **20** and **32** were excluded from the fitting as their carboxylic acid hinders passage through the cell membrane.

Chemistry. During the initial stages of the campaign focused on fragment optimization, we successfully developed a synthetic route to prepare compound **3**. Compounds **1** and **2** had been purchased during the screening phase prior to this campaign.[21] To synthesize chloropyrazolopyrimidine heterocycle **3**, first, we performed a cyclization reaction of **41** by heating it in the presence of urea. The following deoxygenation and chlorination of the pyrimidine ring was achieved by $POCl_3$. Finally, we carried out regioselective S_NAr using an ethanolic solution of $MeNH_2$ (33%), ultimately yielding the desired fragment **3** with an overall yield of 9 % (Scheme 1).

Scheme 1. Preparation of Compound 3[a]

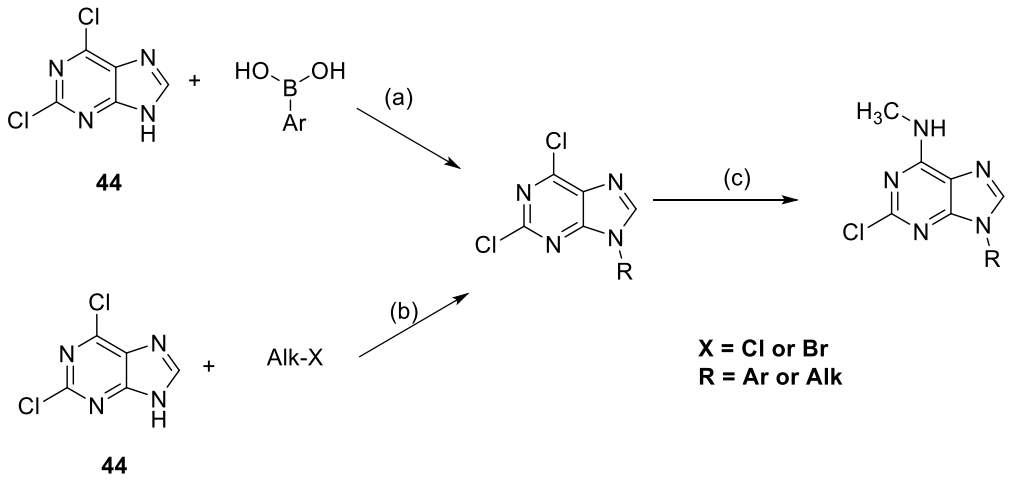


[a]Reagents and conditions: (a) Urea, 195 °C, neat (b) $POCl_3$, DIPEA, rt. (c) 33 % $MeNH_2$ in EtOH, EtOH, rt.

The main synthetic part comprises the preparation of N^9-substituted-4-methylaminopurines as promising YTHDC1 inhibitors. The strategy was based on the convenient purine N^9 derivatization

followed by a regioselective S_NAr. To achieve N^9 arylations, we used reported Chan-Lam coupling between commercially available 2,6-dichloropurine and boronic acid derivatives.[29] For the N^9 alkylation, and corresponding alkyl halides were used. During these reactions, we observed a formation of both N^9 and N^7 regioisomers. However, the desired N^9 isomer was formed as the main product and could be easily separated during the purification. The final synthetic step, S_NAr, was carried out using $MeNH_2$, except for compound **37**, where cyclopropylamine was used instead. If not stated otherwise, all the final molecules were synthesized according to the General synthetic scheme (Scheme 2). However, compounds **8**, **9** and **40** were acquired from a commercial supplier.[30] For the preparation of compound **7**, the General synthetic procedure could not be applied. Instead, the targeted compound was obtained by a two-step reaction process using pTSA catalyzed reaction of 2,6-dichloropurine **44** and 3,4-dihydropyran followed by nucleophilic aromatic substitution.

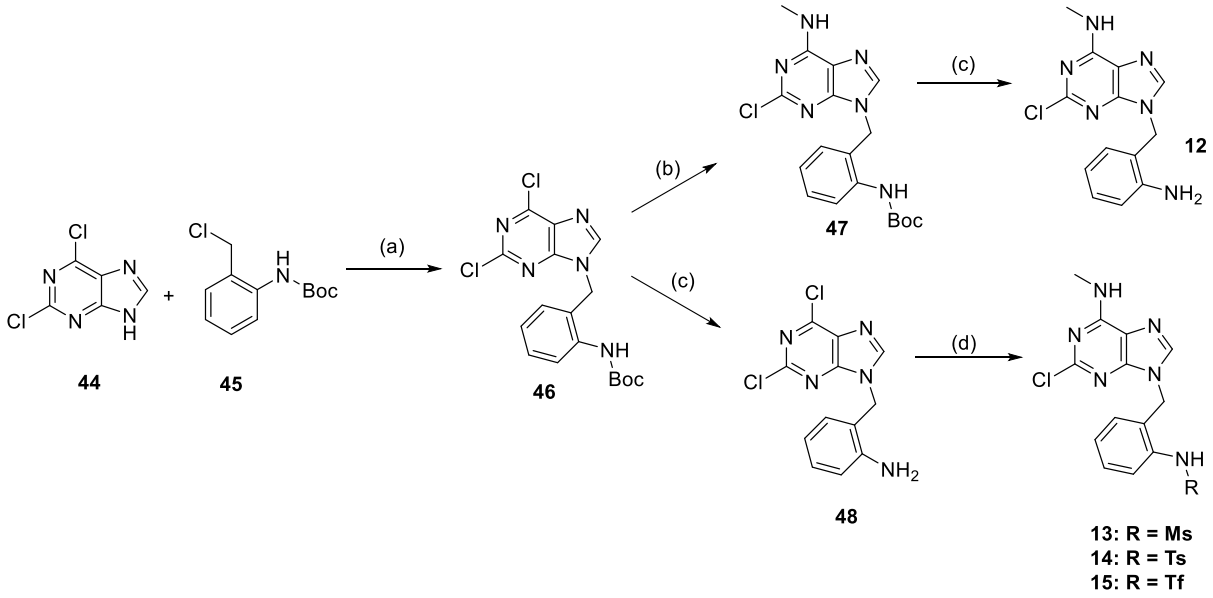
Scheme 2. General synthetic scheme for N^9-substituted purines[a]



[a]Reagents and conditions: (a) $Cu(OAc)_2$, 1,10-phenantroline, DCM, MS 4Å, rt; (b) K_2CO_3, DMF, rt; (c) 33% $MeNH_2$ in EtOH, EtOH, rt

Compounds **12-15** were prepared from a shared intermediate **46**. The synthesis of **12** followed the reaction sequence outline in the General synthetic scheme (Scheme 2), with subsequent removal of the Boc protecting group from **47**. On the other hand, the preparation of compounds **13-15** involved the initial Boc removal and formation of **48**, followed by free amino group modification using standard conditions for sulfonation and acylation, respectively. The S_NAr was carried out as the final step of the reaction sequence (Scheme 3). On the contrary, compound **31**, which also contains a sulfonamide moiety, was prepared according to the General synthetic scheme (Scheme 2). The 2,6-dichloropurine **44** alkylation was carried out with an intermediate already containing the sulfonamide group. Even though, this approach provided the final compound **31** in limited yield (6 %, after three steps), the reaction sequence was not further optimized. These findings suggest, that the modified reaction sequence, used for compounds **13-15**, shall be preferentially used in combination with other sulfonamide derivatives.

Scheme 3. Synthetic route for Compounds **12, 13, 14** and **15**[a]

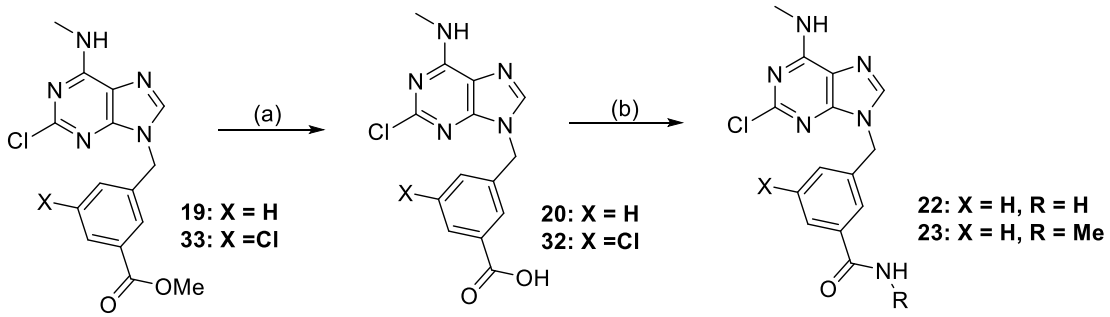


[a]Reagents and conditions: (a) K_2CO_3, DMF, rt; (b) 33% $MeNH_2$ in EtOH, EtOH, rt; (c)

TFA, DCM, rt (d) TsCl, MsCl or TFAA, Py, DCM, 0°C

Compounds **20**, **29** and **32** that contain carboxylic groups, as well as compounds **22**, and **23**, with amide moiety, were synthesized from corresponding alkyl esters **19**, **28** and **33**, respectively. The methyl ester hydrolysis was achieved by heating the reaction mixture in the presence of 37% HCl for compounds **19**, **28**, **33**. In the case of compound **10**, its *tert*-butyl ester analog was subjected to hydrolysis with TFA. For the synthesis of compounds **22**, and **23**, a combination of COMU® as the coupling agent along with the presence of NH_3 or $MeNH_2$ was used (Scheme 4).

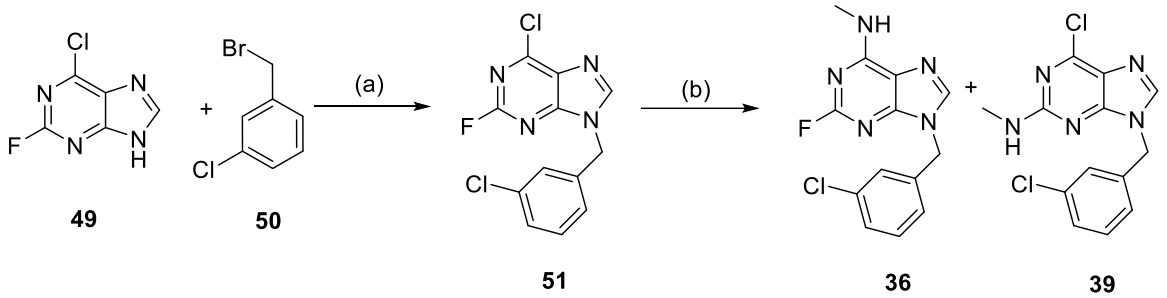
Scheme 4. Synthetic route for the preparation of carboxylic acid derivatives[a]



[a]Reagents and conditions: (a) 37 % HCl, dioxane, reflux; (b) RNH_2, COMU, DIPEA, DMF, 0°C to rt

Lastly, due to the nature of fluorine substituent, the nucleophilic aromatic substitution of **51** was not regioselective. As a result, this reaction led to the formation of two products **36** and **39** that were isolated and tested against YTHDC1 (Scheme 5).

Scheme 5. Synthetic route for Compounds **36** and **39**[a]



[a]Reagents and conditions: (a) K_2CO_3, DMF, rt; (b) 33% $MeNH_2$ in EtOH, EtOH, rt;

Conclusions. In summary, we used structure-based design to develop a small-molecule inhibitor targeting YTHDC1. Through step-by-step ligand optimization and leveraging the most potent substituents identified on a purine heterocycle, we successfully combined the structural features together with the 5-chloropyrazolopyrimidine scaffold. Supporting the campaign with high-resolution X-ray data, we evaluated the efficacy of the most potent compound **40** in the antiproliferative assay using TSA and ITC assays. Compound **40** exhibits a K_D value of 49 nM, very good LE and LLE values, and antiproliferative activity against acute myeloid leukemia cell lines (THP-1, MOLM-13, NOMO1). Compound **40** is selective against the cytoplasmic YTHDF1-3 readers according to two different assays (HTRF assay, TSA) and against YTHDC2 according to TSA. The correlations between the biochemical SAR (FRET IC_{50} values) and cellular SAR (THP-1 GI_{50} values) for the series that led to ligand **40** provide evidence of target engagement in the cell. Thus, we propose compound **40** as a chemical probe to study the role of the YTHDC1 m^6A-reader in AML.

Material and methods

HTRF assay (IC_{50}). GST-YTHDC1, GST-YTHDF1, GST-YTHDF2, and GST-YTHDF3 were purified as previously reported.[31] The HTRF assay was assembled and run as detailed in Ref.[21]

with the only difference being that the starting concentration of the dose-response experiments used for the IC_{50} determination was variated dependently from the tested compound. The same protocol applies to the four proteins. The competitive inhibition data of GST-YTHDC1 in the presence of the compounds were normalized using a blank assembled with all the components of the assay, including DMSO, except for GST-YTHDC1. The competitive inhibition data of GST-YTHDF1, GST-YTHDF2, and GST-YTHDF3 in the presence of compound 40 were normalized using a blank assembled with all the components of the assay, minus the protein, and 2-fold serial dilutions of compound 40. This measure was adopted to mitigate the interference arising from high concentrations of compound 40. In all cases, the signal was measured using a Spark plate reader (Tecan), with a 320 nm excitation filter and 620 nm (measurement 1) or 665 (measurement 2) emission filters, a dichroic 510 mirror, 75 flashes, and applying a lag time of 100 μs and an integration time of 400 μs.

Thermal shift assay. The YTH domain of YTHDF1 (residues 361-559), YTHDF2 (residues 383-579), YTHDF3 (residues 387-585), YTHDC1 (residues 345-509) and YTHDC2 (residues 1285-1424) were cloned into pET-based vector harbouring N-terminal hexa-histidine tag and a TEV cleavage site. All recombinant proteins were overexpressed at 20 °C in *E. coli* BL21 (DE3) upon induction with 0.4 mM IPTG and purified by HiTrap nickel column. The His tag was cleaved by the addition of TEV protease (1:100) to the purified recombinant protein while dialysis to remove imidazole at 4 °C overnight. The samples were then passed through nickel column and further purified by size exclusion chromatography. YTHDC1, YTHDF1, YTHDF2, YTHDF3, and YTHDC2 were buffered in 50 mM HEPES pH 7.5, 150 mM NaCl, and tested in a white 96 well plate at a final concentration of 2 μM. SYPRO Orange dye was added to the mix with a volume ratio of 1:1000. Compound **40** was also added to the mix and tested as a set of 2-fold dilutions.

The fluorescence monitoring was performed using a LightCycler® 480 System. The temperature was set up to increase with a ramp rate of 0.06 °C/s from 20°C to 85°C and 10 acquisitions per °C were taken in dynamic integration time mode and using red 610 (498-610) filter combination. The melting curves were calculated using the Tm calling analysis of the LightCycler® 480 software release 1.5.1.62 SP3.

Isothermal titration calorimetry. The Isothermal titration calorimetry (ITC) experiment was carried out at 18 °C using MicroCal ITC200 (GE Healthcare). Protein and compound were dissolved in 20 mM Tris pH 7.4, 150 mM NaCl along with 0.2% DMSO. Protein at the concentration of 100 µM was titrated into the sample cell containing 10 µM compound. After an initial injection of 1 µL, 13 injections of 3.0 µL each were performed. The raw data were integrated and analyzed using a single-binding site model, provided in the MicroCal Origin software package.[32]

Crystallography. The crystals of YTHDC1 YTH domain were obtained by mixing 1 µL protein solution at 10 mg/mL with mother liquor containing 0.1 M Bis-Tris at pH 6.5, 0.2 M ammonium sulfate and 25% PEG 3350 at 22°C in a hanging drop vapor diffusion setup. To obtain crystals of protein complexed with fragments, the crystals were transferred to a 1 µL drop containing 50-200 mM (depending on the solubility) fragment directly dissolved in 0.1 M Bis-Tris at pH 6.5, 0.2 M ammonium sulfate and 30% PEG 3350, soaked overnight at 22 °C, harvested and frozen in liquid nitrogen without additional cryoprotection. Diffraction data were collected at the Swiss Light Source (Villigen, Switzerland) using the beamline X06DA (PXIII), and processed using XDS.[33] The structures were solved by molecular replacement using Phaser program[34] from the Phenix package.[35] The unliganded structure of YTHDC1 (PDB ID: 4R3H) was used as a search model. The model building and refinements were performed using COOT[36] and phenix.refine.[37]

Cell culture. THP-1, MOLM-13 and NOMO1 cell lines were obtained from DSMZ-German Collection of Microorganisms and Cell Cultures GmbH. Cells were cultured in RPMI 1640 medium (11875093, Thermo Fisher Scientific) containing 10 % FBS (16140071, Thermo Fisher Scientific) and 1 % penicillin-streptomycin (15140122, Thermo Fisher Scientific) in 5 % CO2 at 37 °C in a humidified incubator. Cell lines were tested negative formycoplasma contamination (PCR-based assay by Microsynth, Switzerland)

Cytotoxicty (GI_{50}). Cells were seeded in white clear-bottom 96-well plates at a density of 6 × 103 cells/well in 50 µL of the complete RPMI medium and treated with 50 µL of increasing concentrations of the indicated compounds dissolved in DMSO (final concentration of compounds 0.6 − 160 µM) or DMSO only (0.5 % (v/v)) as a negative control and incubated for 72 h at 37°C with 5 % CO2. Cell viability was determined using a CellTiter-Glo luminescent cell viability assay (Promega) based on the detection of ATP according to manufacturer's instructions. 100 µL of the reagent was added to each well and incubated for 10 min at room temperature on an orbital shaker. The luminescence was recorded using a Tecan Infinite 3046 M1000 microplate reader from the top. Background luminescence value was obtained from wells containing the CellTiter-Glo reagent and medium without cells. Cell viability curves were plotted in GraphPad Prism 9 and fitted with nonlinear regression, from which GI50 values were determined.

Chemistry. All reagents were purchased from commercial suppliers and used as received. Reactions run at elevated temperature were carried out in the oil bath. Our research group successfully synthesized all the compound as described, except for compounds **1, 2a, 2b, 8, 9** and **40** which were obtained from commercial supplier.[30] All compounds have >95% purity (HPLC). All reactions were monitored by thin-layer chromatography (Aluminium plates coated with silica gel 60 F_{254}). Flash column chromatography was carried out over silica gel (0.040-0.063 mm). ^{1}H

and ^{13}C {^{1}H} NMR spectra were recorded on AV2 400 MHz and AV600 Bruker spectrometers (400 MHz, 101 MHz and 600 MHz, 150 MHz, respectively) in DMSO or $CDCl_3$ Chemical shifts are given in ppm and their calibration was performed to the residual ^{1}H and ^{13}C signals of the deuterated solvents. Multiplicities are abbreviated as follows: singlet (s), doublet (d) multiplet (m), and broad signal (bs). The purity was acquired by Liquid chromatography high resolution electrospray ionization mass spectrometry (LC-HR- ESI-MS): Acquity UPLC (Waters, Milford, USA) connected to an Acquity eλ diode array detector and a Synapt G2 HR-ESI-QTOF-MS (Waters, Milford, USA); injection of 1 µL sample (c = ca. 10-100 µg mL−1 in the indicated solvent); Acquity BEH C18 HPLC column (1.7 µm particle size, 2 × 50 mm, Waters) kept at 30 °C elution at a flow rate of 400 µL/min with A: H2O + 0.02% TFA and B: CH3CN + 0.02% TFA, linear gradient from 10–95% B within 3 min, then isocratic 95% B for 2 min; UV spectra recorded from 190–300 nm at 1.2 nm resolution and 20 points s-1UV spectra recorded from 200–600 nm at 1.2 nm resolution and 20 points s−1; ESI: positive ionization mode, capillary voltage 3.0 kV, sampling cone 40V, extraction cone 4V, N2 cone gas 4 L h−1, N2 desolvation gas 800 L min−1, source temperature 120°C; mass analyzer in resolution mode: mass range 100–2'000 m/z with a scan rate of 1 Hz; mass calibration to <2 ppm within 50–2'500 m/z with a 5mM aq. soln. of HCO2Na, lockmasses: m/z 195.0882 (caffein, 0.7 ng mL−1) and 556.2771(Leucine-enkephalin, 2 ng mL−1).

5-chloro-N,3-dimethyl-1H-pyrazolo[4,3-d]pyrimidin-7-amine (3)

To a powder of 4-amino-3-methyl-1H-pyrazole-5-carboxamide **41** (0.13 g, 0.97 mmol), which was prepared following the reported procedure[38], was added urea (389 mg, 6.4 mmol). The neat reaction mixture was heated and stirred at 195 °C for 5 h. Upon the temperature increase, the solid reactants melted and after the product formation, the reaction mixture solidified. The reaction

vessel was cooled to rt and the crude product **42** was used in the next step without further purification.

The pyrazolo[4,3-d]pyrimidine-5,7(6H)-dione **42** was suspended in $POCl_3$ (4.6 mL) followed by the addition of DIPEA (0.403 mL, 2.3 mmol). The reaction mixture was heated at 70 °C for 14 hours. The volatiles were removed *in vacuo* and the residue was poured over ice. The mixture was extracted into EtOAc (3 x 6 mL) and the combined organic layers were dried over $MgSO_4$ and filtered. Activated charcoal was added to the filtrate and the mixture was stirred for 10 minutes. After the charcoal removal (filtration paper), the solvent was removed under reduced pressure. The crude product **43** was dissolved in EtOH and 33% $MeNH_2$ in EtOH (0.2 mL) was added into the reaction vessel. The reaction mixture was stirred at rt for 1 hour and after the reaction completion (TLC), the volatiles were removed *in vacuo*. The crude product **3** was purified using flash column chromatography (SiO_2; EtOAc/MeOH = 10 : 1) and the desired compound was obtained as a white solid (0.018 g, 9 % after three steps). ^{1}H NMR (400 MHz, MeOH - d^4) δ 3.12 (s, 3H), 2.49 (s, 3H). ^{13}C NMR (126 MHz, MeOH - d^4) δ 155.8, 154.2, 138.6, 136.7, 128.3, 27.8, 9.3. LRMS (ESI) m/z: [M + H]+ calcd for $C_7H_9ClN_5$; 198.054 found, 198.019.

General procedure 1 (N^9-alkylation of purines)

2,6-dichloropurine **44** (1 equiv) was dissolved in DMF (0.5 M) and K_2CO_3 (2 equiv) was added. Corresponding alkyl halide was subsequently added to the reaction mixture (1 equiv). The resulting reaction mixture was stirred at rt until a reaction completion (Monitored by TLC). The reaction mixture was quenched by an addition of water and extracted into EtOAc. Combined organic layers were washed by 10 % aq. sol. of LiCl. Then dried over $MgSO_4$, filtrated, and evaporated.

General procedure 2 (Regioselective S_NAr with R-NH_2)

9-alkyl-2,6-dichloropurine (1 mmol) was suspended in EtOH (0.5 M) and 33 % $MeNH_2$ in EtOH (2 mL) and stirred at rt. After the reaction completion (TLC) the volatiles were removed *in vacuo.* The crude product was purified using flash column chromatography.

2-chloro-N-methyl-9-phenyl-9H-purin-6-amine (4)

The final compound was prepared following the General method 2 from corresponding 9-phenyl-2,6-dichloro-9H-purine (0.02 g, 0.075 mmol) that was prepared according to reported Chan-Lam coupling procedure.[29] The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1.2 : 1) and the desired compound was obtained as a white solid (0.017 g, 87 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 7.98 (s, 1H), 7.67 – 7.64 (m, 2H), 7.57 – 7.53 (m, 2H), 7.46 – 7.42 (m, 1H), 6.05 – 5.93 (bs, 1H), 3.22 (s, 3H); ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 156.4, 155.5, 149.7, 139.3, 134.6, 130.1, 128.4, 123.6, 119.6, 27.9. LRMS (ESI) m/z: [M + H]+ calcd for $C_{12}H_{11}ClN_5$; 260.070 found, 260.070.

9-benzyl-2-chloro-N-methyl-9H-purin-6-amine (5)

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.2 g, 1.06 mmol) and BnBr (0.126 mL, 1.06 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1.2 : 2) and the desired compound was obtained as a white solid (0.160 g, 54 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 8.05 (s, 1H), 7.43 – 7.36 (m, 3H), 7.31 (m, 2H), 5.41 (s, 2H); ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 153.4, 153.3, 152.1, 145.7, 134.1, 130.8, 129.6, 129.3, 128.3, 48.2.

The final compound was prepared following the General method 2 from corresponding 9-benzyl-2,6-dichloro-9H-purine (0.08 g, 0.286 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 2.5 : 1 -> 4 : 1) and the desired compound was

obtained as a white solid (0.06 g, 76 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 7.63 (bs, 1H), 7.38 – 7.32 (m, 3H), 7.31 – 7.28 (m, 2H), 5.90 – 5.88 (bs, 1H), 5.32 (s, 2H), 3.19 (bs, 3H); ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 156.1, 154.9, 150.1, 139.8, 135.3, 129.1, 128.5, 128.0, 118.7, 47.2, 29.7. LRMS (ESI) m/z: [M + H]+ calcd for $C_{13}H_{13}ClN_5$; 274.085 found, 274.085.

2-chloro-N-methyl-9-(pyridin-4-ylmethyl)-9H-purin-6-amine (6)

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.2 g, 1.06 mmol) and 4-bromoethylpyridine hydrobromide (0.267 g, 1.06 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/MeOH = 2 : 0.2) and the desired compound was obtained as a white solid (0.094 g, 32 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 8.64 – 8.62 (m, 2H), 8.11 (s, 1H), 7.08 (d, *J* = 6.0 Hz, 2H), 5.45 (s, 2H). ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 153.7, 153.3, 152.5, 151.0, 145.4, 143.0, 130.8, 122.2, 46.8.

The final compound **6** was prepared following the General method 2 from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.069 g, 0.246 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/MeOH = 2 : 0.2 -> 2 : 0.8) and the desired compound was obtained as a white solid (0.045 g, 66 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.53 – 8.52 (m, 2H), 8.29 – 8.27 (m, 1H), 8.26 (s, 1H), 7.16 – 7.15 (m, 2H), 5.42 (s, 2H), 2.93 (d, *J* = 4.6 Hz, 3H); ^{13}C NMR (101 MHz, DMSO – d^6) δ 155.6, 153.5, 150.0, 149.5, 145.5, 141.3, 121.8, 118.3, 45.2, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{12}H_{12}ClN_6$; 275.081 found, 275.081.

2-chloro-N-methyl-9-(tetrahydro-2H-pyran-2-yl)-9H-purin-6-amine (7)

The final compound **7** was prepared following the General method 2 from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.2 g, 0.073 mmol) that was prepared following reported procedure[39]. The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 2 : 1) and the desired compound was obtained as a white solid (0.165 g, 84 %). ^{1}H

NMR (400 MHz, DMSO – d^6) δ 8.35 (s, 1H), 8.25 – 8.22 (m, 1H), 5.56 (dd, J = 11.0, 2.2 Hz, 1H), 4.03 – 3.96 (m, 1H), 3.75 – 3.61 (m, 1H), 2.92 (d, J = 4.6 Hz, 3H), 2.37 – 2.18 (m, 1H), 2.08 – 1.92 (m, 2H), 1.86 – 1.72 (m, 1H), 1.68 – 1.56 (m, 2H). ^{13}C NMR (101 MHz, DMSO – d^6) δ 155.5, 153.4, 148.9, 139.2, 118.2, 80.9, 67.7, 30.0, 27.2, 24.5, 22.3. LRMS (ESI) m/z: [M + H]+ calcd for $C_{11}H_{15}ClN_5O$; 268.096 found, 268.097.

3-(2-chloro-6-(methylamino)-9H-purin-9-yl)benzoic acid (10)

The final compound **10** was prepared following the General method 2 from corresponding tert-butyl 3-(2,6-dichloro-9H-purin-9-yl)benzoate (0.02 g, 0.055 mmol) that was prepared following reported Chan-Lam coupling procedure.[29] The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 2 : 1). and the desired compound was obtained as a white solid (0.018 g, 91 %). ^{1}H NMR (400 MHz $CDCl_3$ - *d*) δ 8.21 (t, J = 1.9 Hz, 1H), 8.07 – 8.02 (m, 2H), 7.93 (dd, J = 8.1, 1.2 Hz, 1H), 7.61 (t, J = 7.9 Hz, 1H), 6.14 – 6.03 (bs, 1H), 3.29 – 3.15 (bs, 3H), 1.62 (s, 9H). ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 164.6, 156.4, 155.6, 139.0, 134.8, 134.0, 130.0, 129.2, 127.5, 124.1, 119.7, 82.1, 77.4, 28.3.

The tert-butyl ester (0.014, 0.039 mmol) was dissolved in DCM (0.5 mL) and TFA was added (0.02 mL). The reaction was stirred at rt overnight and neutralized with DIPEA. The volatiles were evaporated *in vacuo* and the crude product was recrystalized from H_2O. The final product was obtained as white solid (0.011 g, 93 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.64 (s, 1H), 8.41 – 8.37 (m, 1H), 8.35 – 8.34 (m, 1H), 8.07 – 8.01 (m, 2H), 7.76 – 7.72 (m, 1H), 2.96 (d, J = 4.5 Hz, 3H); ^{13}C NMR (101 MHz, DMSO – d^6) δ 166.5, 155.8, 153.9, 149.0, 140.1, 134.9, 132.3, 130.0, 128.5, 127.6, 123.9, 118.9, 27.3.LRMS (ESI) m/z: [M + H]+ calcd for $C_{13}H_{11}ClN_5O_2$; 304.057 found, 304.039.

2-(2-chloro-6-(methylamino)-9H-purin-9-yl)-N-phenylacetamide (11)

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.2 g, 1.06 mmol) and *N*-phenylchloroacetamide (1.0 g , 5.9 mmol), that was prepare following reported procedure[40]. The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 2:1 to 4:1) and the desired compound was obtained as a white solid (0.54 g, 28 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 10.53 (s, 1H), 8.88 (s, 1H), 7.57 – 7.54 (m, 2H), 7.35 – 7.31 (m, 2H), 7.11 – 7.09 (m, 1H), 5.42 (s, 1H). ^{13}C NMR (101 MHz, DMSO – d^6) δ 164.8, 163.1, 153.5, 151.0, 143.4, 138.3, 128.9, 123.9, 122.4, 119.3, 49.4.

The final compound **11** was prepared following the General method 2 from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.05 g, 0.155 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 4 : 1 to 10:1) and the desired compound was obtained as a white solid (0.038 g, 77 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 10.45 (bs, 1H), 8.23 – 8.22 (m, 1H), 8.13 (s, 1H), 7.59 (d, J = 7.3 Hz, 2H), 7.35 – 7.31 (m, 2H), 7.08 (t, J = 7.4 Hz, 1H), 5.05 (s, 2H), 2.94 (d, J = 4.6 Hz, 3H). ^{13}C NMR (101 MHz, DMSO – d^6) δ 164.9, 155.5, 153.3, 149.8, 142.2, 138.5, 128.9, 123.7, 119.1, 117.9, 45.7, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{14}ClN_6O$; 317.091 found, 317.092.

9-(2-aminobenzyl)-2-chloro-N-methyl-9H-purin-6-amine (12)

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.531 g, 2.81 mmol) and tert-butyl (2-(chloromethyl)phenyl)carbamate **45** that was prepared following reported procedure[41] (0.68 g, 2.81 mmol). The crude product **46** was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1 : 1) and the desired compound was obtained as a white solid (0.346 g, 31 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.93 (s, 1H), 8.62 (s, 1H), 7.33 – 7.28 (m, 2H), 7.15 – 7.10 (m, 1H), 7.08 (d, J = 7.0 Hz, 1H), 5.48 (s, 2H), 1.41 (s, 9H); ^{13}C NMR

(101 MHz, DMSO – d^6) δ 154.1, 154.0, 151.5, 150.2, 148.9, 136.5, 130.8, 130.2, 129.1, 129.0, 126.6, 125.9, 79.6, 44.4, 28.5.

The next step was performed using the General method 2 from corresponding 9-alkyl-2,6-dichloro-9H-purine **46** (0.15 g, 0.38 mmol). The crude product **47** was purified using flash column chromatography (SiO_2; EtOAc/Hept = 2 : 1) and the desired compound was obtained as a white solid (0.121 g, 81 %). ^{1}H NMR (400 MHz, DMSO – d^6) 9.06 (s, 1H), 8.28 – 8.22 (m, 1H), 8.11 (s, 1H), 7.38 (d, J = 8.7 Hz, 1H), 7.29 (td, J = 7.6, 1.6 Hz, 1H), 7.11 (td, J = 7.5, 1.4 Hz, 1H), 7.00 (d, J = 6.8 Hz, 1H), 5.32 (s, 2H), 2.92 (d, J = 4.6 Hz, 3H), 1.46 (s, 9H). ^{13}C NMR (101 MHz, DMSO – d^6) δ 155.6, 153.6, 153.3, 149.5, 141.1, 136.0, 130.1, 128.4, 125.6, 125.2, 118.1, 79.2, 43.0, 28.1, 27.3.

The final compound **12** was prepared by deprotection of **47** (0.05 g, 0.128 mmol) in DCM (0.6 mL) using TFA (0.097 mL, 1.28 mmol). The reaction mixture was stirred at room temperature for 6 h. The reaction was quenched by aq. sol. of Na_2CO_3 and extracted into EtOAc. Combined organic layers were dried over $MgSO_4$. The volatiles were removed *in vacuo* and the crude product was purified using flash column chromatography (SiO_2; EtOAc/MeOH = 2 : 1 → 10 : 1) and the desired compound was obtained as a brownish solid (0.024 g, 64 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.26 – 8.21 (m, 1H), 8.13 (s, 1H), 7.01 (td, J = 7.6, 1.6 Hz, 1H), 6.78 (dd, J = 7.6, 1.6 Hz, 1H), 6.69 (dd, J = 8.0, 1.2 Hz, 1H), 6.51 (td, J = 7.4, 1.2 Hz, 1H), 5.16 (s, 2H), 2.92 (d, J = 4.6 Hz, 3H). ^{13}C NMR (101 MHz, DMSO – d^6) δ 155.54, 153.23, 149.44, 145.93, 141.10, 128.80, 128.62, 119.68, 118.11, 116.53, 115.46, 43.09, 27.18. LRMS (ESI) m/z: [M + H]+ calcd for $C_{13}H_{14}ClN_6$; 289.096 found, 289.096.

N-(2-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)phenyl)methanesulfonamide (13)

The compound **46** (0.47 g, 1.19 mmol) was dissolved in DCM (5 mL) followed by addition of TFA (0.45 mL). The reaction mixture was stirred at rt for 16 hours. After the reaction completion, the reaction was quenched by aq. sol. Na_2CO_3 and extracted into DCM. Combined organic layers were dried over $MgSO_4$ and the volatiles were removed *in vacuo*.. The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1 : 1) and the intermediate **48** was obtained as a white solid (0.346 g, 31 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.93 (s, 1H), 8.62 (s, 1H), 7.33 – 7.28 (m, 2H), 7.15 – 7.10 (m, 1H), 7.08 (d, J = 7.0 Hz, 1H), 5.48 (s, 2H), 1.41 (s, 9H); ^{13}C NMR (101 MHz, DMSO – d^6) δ 154.1, 154.0, 151.5, 150.2, 148.9, 136.5, 130.8, 130.2, 129.1, 129.0, 126.6, 125.9, 79.6, 44.4, 28.5.

To a solution of Compound **48** (0.067 g, 0.227 mmol) in DCM (2.2 mL) was added pyridine (0.024 mL, 0.295 mmol). The solution was cooled down to 0 °C and MsCl (0.021 mL, 0.274 mmol) was added dropwise after 30 min. After the reaction completion (TLC), the volatiles were removed *in vacuo*. The crude product was subsequently used in the next step without further purification. The final product **13** was prepared following general method for S_NAr. The crude product was purified using flash column chromatography (SiO_2; EtOAc/MeOH = 4 : 1 → 4 : 0) and the desired compound was obtained as a white solid (0.025 g, 30 % after two steps). ^{1}H NMR (400 MHz, DMSO – d^6) δ 9.50 (s, 1H), 8.31- 8.25 (m, 1H), 8.18 (s, 1H), 7.40 (dd, J = 7.9, 1.5 Hz, 1H), 7.35 (td, J = 7.5, 1.5 Hz, 1H), 7.22 (td, J = 7.4, 1.5 Hz, 1H), 6.87 (dd, J = 7.8, 1.5 Hz, 1H), 5.45 (s, 2H), 3.09 (s, 3H), 2.92 (d, J = 4.6 Hz, 2H); ^{13}C NMR (101 MHz, DMSO – d^6) δ 155.6, 153.3, 149.6, 141.4, 134.8, 133.2, 128.7, 128.0, 127.0, 126.8, 118.2, 42.9, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{12}H_{12}N_5$; 367.073 found, 367.075.

N-(2-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)phenyl)-4-methylbenzene sulfonamide (14)

To a solution of Compound **48** (0.1 g, 0.339 mmol) in DCM (2 mL) was added pyridine (0.032 mL, 0.407 mmol). The solution was cooled down to 0 ℃ and TsCl (0.064 mL, 0.339 mmol) was added portion wise after 30 min. After the reaction completion (TLC), the volatiles were removed *in vacuo*. The crude product was subsequently used in the next step without further purification. The final product **14** was prepared following general method for S_NAr. The crude product was purified using flash column chromatography (SiO_2; EtOAc:Hept = 2 : 1) and the desired compound was obtained as a white solid (0.078 g, 52 % after two steps). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) 7.74 (d, *J* = 8.0 Hz, 2H), 7.72 – 7.68 (bs, 1H), 7.29 – 7.21 (m, 4H), 7.18 – 7.10 (m, 1H), 6.27 – 6.13 (bs, 1H), 4.93 (s, 2H), 3.20 – 3.03 (bs, 3H), 2.40 (s, 3H). ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 156.0, 154.7, 149.0, 143.5, 139.4, 138.2, 135.9, 130.7, 130.5, 130.2, 129.8, 127.5, 127.2, 1276.0, 118.7, 44.2, 27.8, 21.7. LRMS (ESI) m/z: [M + H]+ calcd for $C_{20}H_{20}ClN_6O_2S$; 443.105 found, 443.099.

N-(2-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)phenyl)-2,2,2-trifluoroacetamide (15)

To a solution Compound **48** (0.07 g, 0.237 mmol) in DCM (2 mL) was added pyridine (0.024 mL, 0.308 mmol). The solution was cooled down to 0 ℃ and trifluoroacetic anhydride (0.039 mL, 0.280 mmol) was added dropwise after 30 min. After the reaction completion (TLC), the volatiles were removed *in vacuo*. The crude product was subsequently used in the next step without further purification. The final compound **15** was prepared following the General method 2 for S_NAr. The crude product was purified using flash column chromatography (SiO_2; EtOAc/MeOH = 2 : 1 → 3 : 1) and the desired compound was obtained as a white solid (0.030 g, 30 % after two steps). ^{1}H NMR (400 MHz, DMSO – d^6) δ 11.26 (s, 1H), 8.28 – 8.26 (m, 1H), 8.10 (s, 1H), 7.42 – 7.31 (m, 3H), 7.18 (dd, J = 7.6, 1.5 Hz, 1H), 5.33 (s, 2H), 2.92 (d, J = 4.5 Hz, 1H); ^{13}C NMR (101 MHz,

DMSO – d^6) δ 155.8 (q, J = 36.7 Hz), 155.5, 153.4, 149.4, 141.0, 132.6, 132.3, 128.8, 128.7, 128.1, 127.4, 118.1, 117.4, 114.5, 42.7, 27.2. ^{19}F NMR (376 MHz, DMSO) δ -73.90. LRMS (ESI) m/z: [M + H]+ calcd for $C_{15}H_{12}ClF_3N_6O$; 385.079 found, 385.078.

2-chloro-9-(2-(difluoromethyl)benzyl)-N-methyl-9H-purin-6-amine (16)

2-(difluoromethyl)benzoic acid (0.500 g, 2.90 mmol) was dissolved in THF (5 mL) and BH_3 . DMS (0.687 mL, 7.25 mmol) was added dropwise. The reaction mixture was stirred under dinitrogen atmosphere at rt overnight. After the reaction completion (monitored by TLC), the reaction was quenched by aq. sol. of $NaHCO_3$. The mixture was extracted into EtOAc (3 x 7 mL). The combined organic layers were dried over $MgSO_4$, filtrated and evaporated. The resulting alcohol was dissolved in DCM (5.2 mL) together with one drop of DMF, followed by addition of $SOCl_2$ (0.315 mL, 4.35 mmol). The reaction mixture was stirred at rt overnight. The volatiles were removed *in vacuo* and the resulting alkyl chloride was used in the next step without further purification.

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.48 g, 2.54 mmol) and 1-chloromethyl-2-difluoromethylbenzen (0.449 g, 2.54 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1 : 1) and the desired compound was obtained as a white solid (0.210 g, 25 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 8.06 (s, 1H), 7.54 – 7.46 (m, 3H), 7.34 – 7.32 (m, 1H), 6.84 (t, *J* = 54.8 Hz, 1H), 5.63 (s, 2H); ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 153.4, 152.2, 145.9 (t, *J* = 2.6 Hz), 132.7, 132.1 (t, *J* = 2.0 Hz) , 131.9 (t, *J* = 21.6 Hz), 130.8, 130.7, 129.4, 128.3 (t, *J* = 8.2 Hz), 116.0 (t, *J* = 238.9 Hz), 44.4 (t, *J* = 2.7 Hz).

The final compound **16** was prepared following the General method 2 for S_NAr starting from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.080 g, 0.243 mmol). The crude product was

purified using flash column chromatography (SiO_2; EtOAc/Hept = 1.1 : 1) and the desired compound was obtained as a white solid (0.071g, 90 %). ^{1}H NMR (400 MHz, , $CDCl_3$ - *d*) δ 7.65 (s, 1H), 7.54 – 7.51 (m, 1H), 7.47 – 7.40 (m, 2H), 7.23 (d, *J* = 8.3 Hz, 1H), 6.87 (t, *J* = 54.9 Hz, 1H), 6.09 – 5.94 (bs, 1H), 5.52 (s, 2H), 3.28 – 3.08 (bs, 3H); ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 156.2, 140.1, 133.98 (t, *J* = 2.2 Hz), 131.86 (t, *J* = 21.4 Hz), 131.8 (t, *J* = 2.1 Hz), 130.1, 128.8, 127.63 (t, *J* = 8.1 Hz), 115.49 (t, *J* = 238.8 Hz), 43.5. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{13}ClF_2N_5$; 324.082 found, 324.082.

2-chloro-N-methyl-9-(2-(trifluoromethyl)benzyl)-9H-purin-6-amine (17)

2-(trifluoromethyl)benzoic acid (0.500 g, 2.63 mmol) was dissolved in THF (5.2 mL) and BH_3 . DMS (0.748 mL, 7.89 mmol) was added dropwise. The reaction mixture was stirred under dinitrogen atmosphere at rt overnight. After the reaction completion (monitored by TLC), the reaction was quenched by aq. sol. of $NaHCO_3$. The mixture was extracted into EtOAc (3 x 7mL). The combined organic layers were dried over $MgSO_4$, filtrated and evaporated. The resulting alcohol was dissolved in DCM (5.2 mL) together with one drop of DMF, followed by addition of $SOCl_2$ (0.504 mL, 6.95 mmol). The reaction mixture was stirred at rt overnight. The volatiles were removed *in vacuo* and the resulting alkylchloride was used in the next step without further purification.

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.525 g, 2.78 mmol) and 1-chloromethyl-2-trifluoromethylbenzen (0.540 g, 2.78 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1 : 1.2) and the desired compound was obtained as a white solid (0.234 g, 24 % after three steps). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 8.02 (s, 1H), 7.77 (d, *J* = 7.0 Hz, 1H), 7.58 – 7.47 (m, 2H), 7.29 (d, *J* = 7.6 Hz, 1H), 5.63 (s, 2H); ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 153.6, 153.5, 152.3, 133.1, 132.5,

130.7, 130.5, 129.4, 128.5 (q, J = 30.7 Hz), 128.3, 126.9 (q, J = 5.6 Hz), 125.6, 122.9, 44.5 (q, J = 2.9 Hz).

The final compound was prepared following the General method 2 for S_NAr from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.1 g, 0.288 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1 : 1) and the desired compound was obtained as a white solid (0.045 g, 46 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.33 – 8.27 (m, 1H), 8.17 (s, 1H), 7.82 (d, J = 7.7 Hz, 1H), 7.61 (t, J = 7.7 Hz, 1H), 7.54 (t, J = 7.8 Hz, 1H), 6.85 (d, J = 7.7 Hz, 1H), 5.55 (s, 2H), 2.94 (d, J = 4.6 Hz, 3H); ^{13}C NMR (101 MHz, DMSO – d^6) δ 156.1, 154.01, 150.1, 141.9, 135.0, 133.7, 128.8, 128.5, 126.77 (q, J = 5.7 Hz), 126.6, 126.3, 126.1, 126.0, 123.4, 118.8, 43.6, 27.7. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{12}ClF_3N_5$; 342.073 found, 342.074.

2-chloro-9-(2-methoxybenzyl)-N-methyl-9H-purin-6-amine (18)

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.2 g, 1.06 mmol) and 1-chloromethyl-2-methoxybenzen (0.165 g, 1.06 mmol) that was prepared following reported procedure[42]. The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1.2 : 1) and the desired compound was obtained as a white solid (0.103 g, 31 %). ^{1}H NMR (400 MHz, $CDCl_3$ - d) δ 8.14 (s, 1H), 7.39 (dd, J = 7.5, 1.7 Hz, 1H), 7.35 (td, J = 7.9, 1.7 Hz, 1H), 6.96 (td, J = 7.5, 1.1 Hz, 1H), 6.91 (d, J = 8.2 Hz, 1H), 5.38 (s, 2H), 3.86 (s, 3H); ^{13}C NMR (101 MHz, $CDCl_3$ - d) δ 157.5, 153.4, 152.9, 151.5, 146.6, 131.1, 131.0, 130.7, 122.3, 121.2, 110.9, 55.6, 43.8.

The final compound **18** was prepared following the General method 2 starting from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.048 g, 0.155 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 3 : 1) and the desired compound was obtained as a white solid (0.036 g, 76 %). ^{1}H NMR (400 MHz, $CDCl_3$ - d) δ 7.72

(s, 1H), 7.31 (td, J = 7.9, 1.7 Hz, 1H), 7.26 (dd, J = 7.5, 1.7 Hz, 1H), 6.94 – 6.8 (m, 2H), 6.22 – 6.02 (bs, 1H), 5.30 (s, 2H), 3.86 (s, 3H), 3.27 – 3.06 (bs, 3H); ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 157.4, 156.1, 140.6, 130.6, 130.5, 130.3, 127.7, 123.5, 121.0, 118.6, 110.7, 55.5, 42.7, 27.8. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{15}ClN_5O$; 304.096 found, 304.097.

methyl 3-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)benzoate (19)

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.825 g, 4.37 mmol) and methyl-3-bromomethylbenzoate (1.0 g, 4.37 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 2 : 1) and the desired compound was obtained as a white solid (0.465 g, 31 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 8.08 (s, 1H), 8.04 (dt, J = 7.3, 1.7 Hz, 1H), 8.01 – 8.00 (bs, 1H), 5.46 (s, 2H), 3.92 (s, 3H); ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 166.2, 153.4, 153.1, 152.1, 145.4, 134.5, 132.5, 131.4, 130.7, 130.3, 129.6, 129.1, 52.4, 47.6.

The final compound **19** was prepared following the General method 2 from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.465 g, 1.38 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 3 : 1) and the desired compound was obtained as a white solid (0.374g, 82 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.28 (s, 1H), 8.29 – 8.21 (m, 1H), 7.90 – 7.87 (m, 2H), 7.56 – 7.49 (m, 2H), 5.43 (s, 2H), 3.83 (s, 3H), 2.92 (d, *J* = 4.6 Hz, 1H); ^{13}C NMR (101 MHz, DMSO – d^6) δ 165.9, 155.6, 153.5, 149.5, 141.2, 137.5, 132.3, 130.1, 129.4, 128.6, 128.0, 118.3, 52.3, 45.9, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{15}H_{15}ClN_5O_2$; 332.091 found, 332.091.

3-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)benzoic acid (20)

The corresponding methyl ester **19** (0.1 g, 0.301 mmol) was dissolved in dioxane (3 mL) followed by the addition of 38 % HCl (2 mL). The reaction mixture was refluxed for 5 hours. The

reaction mixture was cooled to 0 °C and left for 4 hours at this temperature. The white precipitate was filtered off and dried on air. The carboxylic acid **20** was isolated as a white solid (60 mg, 63 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.32 (s, 1H), 8.32 – 8.25 (bs, 1H), 7.88 – 7.83 (m, 2H), 7.54 – 7.47 (m, 2H), 5.43 (s, 2H), 2.92 (d, J = 3.9 Hz, 3H). δ ^{13}C NMR (101 MHz, DMSO – d^6) δ 166.9, 155.4, 153.6, 149.4, 141.2, 137.2, 131.9, 131.2, 129.1, 128.8, 128.1, 117.9, 46.0, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{13}ClN_5O_2$; 318.075 found, 318.075.

9-(3-(2H-tetrazol-5-yl)benzyl)-2-chloro-N-methyl-9H-purin-6-amine (21)

To a solution of **26** (0.05 g, 0.167 mmol) in DMSO (1.6 mL) was added anhydrous $CuSO_4$ (0.066 g, 0.417 mmol) and NaN_3 (0.010 g, 0.167 mmol). The reaction mixture was heated at 100 °C for 16 h. The reaction mixture was quenched with 10 % HCl and extracted into EtOAc. Combined organic layers were washed with 10 % aq. sol. of LiCl and dried over $MgSO_4$ and concentrated under reduced pressure. The crude product was purified using flash column chromatography (SiO_2; DCM/MeOH = 5 : 1 → 1 : 1) and the desired compound was obtained as a white solid (0.021 g, 36 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.28 (s, 1H), 8.26 – 8.23 (m, 1H), 7.89 (d, J = 7.7 Hz, 1H), 7.84 (s, 1H), 7.35 (t, J = 7.7 Hz, 1H), 7.15 (d, J = 7.6 Hz, 1H), 5.39 (s, 2H), 2.92 (d, J = 4.7 Hz, 2H).^{13}C NMR (101 MHz, DMSO – d^6) δ 160.3, 155.6, 153.5, 149.5, 141.3, 136.8, 133.1, 128.8, 125.8, 125.2, 124.5, 118.3, 46.3, 27.2.

3-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)benzamide (22)

Corresponding carboxylic acid **20** (0.064 g, 0.201 mmol) was suspended in dry DMF (1.6 mL) followed by the addition of DIPEA (0.052 mL, 0.301 mmol). The reaction mixture was cooled to 0 °C and COMU (0.128 g, 0.301 mmol) was added after 30 min. To the reaction mixture was added 7N NH_3 in THF (0.2 mL) after additional 30 min. The reaction was slowly warmed to rt and stirred overnight. The reaction mixture was then extracted into EtOAc and combined organic layers were

dried over $MgSO_4$ and evaporated. The crude product was purified using flash column chromatography (SiO_2; EtOAc/MeOH = 2 : 0.1 → 2 : 0.2) and the desired compound was obtained as a white solid (0.015 g, 24 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.25 (s, 1H), 8.25 – 8.20 (bs, 1H), 7.98 (s, 1H), 7.79 (dt, J = 7.3, 1.7 Hz, 1H), 7.75 (s, 1H), 7.46 – 7.35 (m, 3H), 5.39 (s, 2H), 2.92 (d, J = 4.6 Hz, 2H); ^{13}C NMR (101 MHz, DMSO – d^6) δ 168.0, 156.0, 153.9, 149.9, 141.7, 137.4, 135.2, 130.6, 129.2, 127.2, 127.0, 118.7, 46.5, 27.7. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{14}ClN_6O$; 317.091 found, 317.091.

3-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)-N-methylbenzamide (23)

Corresponding carboxylic acid **20** (0.027 g, 0.084 mmol) was suspended in dry DMF (0.6 mL) followed by the addition of DIPEA (0.021 mL, 0.126 mmol). The reaction mixture was cooled to 0 °C and COMU (0.053 g, 0.126 mmol) was added after 30 min. To the reaction mixture was added 2M solution of $MeNH_2$ in THF (0.15 mL) after additional 30 min. The reaction was slowly warmed to rt and stirred overnight. The reaction mixture was then extracted into EtOAc and combined organic layers were dried over $MgSO_4$ and evaporated. The crude product was purified using flash column chromatography (SiO_2; EtOAc/MeOH = 2 : 0.1 → 2 : 0.2) and the desired compound was obtained as a white solid (0.017 g, 60 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.44 – 8.43 (m, 1H), 8.28 (s, 2H), 7.75 – 7.71 (m, 2H), 7.44 (t, J = 7.4 Hz, 1H), 7.39 (dt, J = 7.7, 1.6 Hz, 1H), 5.39 (s, 2H), 2.92 (d, J = 4.7 Hz, 3H), 2.76 (d, J = 4.6 Hz, 3H). ^{13}C NMR (101 MHz, DMSO – d^6) δ 166.8, 156.0, 153.9, 149.9, 141.7, 137.4, 135.5, 130.4, 129.2, 126.7, 126.6, 118.7, 46.5, 27.7, 26.7. LRMS (ESI) m/z: [M + H]+ calcd for $C_{15}H_{16}ClN_6$; 331.107 found, 331.106.

9-(3-bromobenzyl)-2-chloro-N-methyl-9H-purin-6-amine (24)

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.5 g, 2.65 mmol) and 1-bromo-3-bromomethylbenzen (0.661 g, 2.65 mmol). The crude

product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1.2 : 2) and the desired compound was obtained as a white solid (0.35 g, 37 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.84 (s, 1H), 7.63 – 7.61 (bs, 1H), 7.54 – 7.51 (m, 1H), 7.34 – 7.29 (m, 2H), 5.50 (s, 2H); ^{13}C NMR (101 MHz, DMSO – d^6) δ 153.4, 151.1, 149.8, 148.4, 138.2, 131.0, 131.0, 130.6, 130.5, 126.8, 121.9, 46.4.

The final compound **24** was prepared following the General method 2 starting from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.150 g, 0.419 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1 : 1) and the desired compound was obtained as a white solid (0.12 g, 81 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.26 (s, 1H), 7.54 – 7.49 (m, 2H), 7.32 (t, J = 7.7 Hz, 1H), 7.24 (d, J = 7.8 Hz, 1H), 5.35 (s, 2H), 2.92 (d, J = 4.7 Hz, 3H). ^{13}C NMR (101 MHz, DMSO – d^6) δ 155.6, 153.5, 149.4, 141.1, 139.4, 131.0, 130.7, 130.2, 126.5, 121.9, 118.3, 45.6, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{13}H_{12}BrClN_5$; 351.995 found, 351.996.

2-chloro-9-(3-chlorobenzyl)-N-methyl-9H-purin-6-amine (25)

The final compound was prepared following the General method 2 starting from corresponding compound **38** (0.272 g, 0.867 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 3 : 1) and the desired compound was obtained as a white solid (0.191 g, 71 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.26 (s, 1H), 8.28 – 8.21 (bs, 1H), 7.40 – 7.35 (m, 3H), 7.23 – 7.18 (m, 1H), 5.35 (s, 2H), 2.92 (d, J = 4.7 Hz, 3H); ^{13}C NMR (101 MHz, DMSO – d^6) δ 155.6, 153.5, 149.4, 141.2, 139.2, 133.3, 130.8, 127.9, 127.4, 126.1, 118.3, 45.7, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{13}H_{12}Cl_2N_5$; 308.045 found, 308.046.

3-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)benzonitrile (26)

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.337 g, 1.79 mmol) and 3-(bromomethyl)benzonitrile (0.35 g, 1.79 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 2.5 : 1) and the desired compound was obtained as a white solid (0.15 g, 28 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 8.11 (s, 1H), 7.67 (dt, *J* = 7.1, 1.7 Hz, 1H), 7.61 – 7.60 (m, 1H), 7.57 – 7.41 (m, 2H), 5.47 (s, 2H). ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 153.7, 153.2, 152.5, 145.2, 135.9, 132.8, 132.3, 131.4, 130.8, 130.5, 117.9, 113.8, 47.2.

The final compound **26** was prepared following the General method 2 from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.082 g, 0.262 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/MeOH = 10 : 1) and the desired compound was obtained as a white solid (0.057 g, 71 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.29 – 8.21 (m, 2H), 7.80 – 7.75 (m, 2H), 7.58 – 7.55 (m, 2H), 5.41 (s, 2H), 2.92 (d, *J* = 4.6 Hz, 1H). ^{13}C NMR (101 MHz, DMSO – d^6) δ 155.6, 153.4, 149.4, 141.1, 138.2, 132.3, 131.7, 131.1, 130.0, 118.5, 118.3, 111.6, 45.5, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{12}ClN_6$; 299.081 found, 299.080.

2-chloro-9-(3-methoxybenzyl)-N-methyl-9H-purin-6-amine (27)

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.2 g, 1.06 mmol) and 1-bromomethyl-3-methoxybenzen (0.212 g, 1.06 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 2 : 1.2) and the desired compound was obtained as a white solid (0.115 g, 35 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 8.02 (s, 1H), 7.25 (d, *J* = 13.5, 5.6 Hz, 1H), 6.87 – 6.80 (m, 3H), 5.33 (s, 2H), 3.75 (s, 3H); ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 160.4, 153.3, 153.3, 152.0, 145.7, 135.5, 130.8, 130.6, 120.3, 114.4, 114.1, 55.5, 48.1.

The final compound **27** was prepared following the General method 2 starting from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.050 g, 0.161 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 2 : 1) and the desired compound was obtained as a white solid (0.034 g, 69 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 7.64 (s, 1H), 7.28 – 7.24 (m, 1H), 6.87 – 6.81 (m, 3H), 6.20 – 6.01 (bs, 1H), 5.27 (s, 2H), 3.77 (s, 3H), 3.25 – 3.08 (bs, 3H). ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 160.2, 156.2, 155.0, 150.2, 139.9, 136.8, 130.3, 120.2, 118.7, 114.0, 113.8, 55.4, 47.3, 27.8. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{15}ClN_5O$; 304.096 found, 304.097.

methyl 4-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)benzoate (28)

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.2 g, 1.06 mmol) and methyl-4-(bromomethyl)benzoate (0.242 g, 1.06 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1 : 2) and the desired compound was obtained as a white solid (0.145 g, 40 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 8.08 (s, 1H), 8.05 – 8.03 (m, 2H), 7.38 – 7.34 (m, 2H), 5.47 (s, 2H), 3.91 (s, 3H). ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 166.3, 153.5, 153.3, 152.2, 145.5, 138.9, 131.0, 130.8, 130.7, 128.0, 52.5, 47.7.

The final compound **28** was prepared following the General method 2 from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.052 g, 0.154 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 2 : 1) and the desired compound was obtained as a white solid (0.038 g, 74 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.26 (s, 2H), 7.93 (d, 2H), 7.35 (d, 2H), 5.45 (s, 2H), 3.83 (s, 3H), 2.92 (bs, 3H); ^{13}C NMR (101 MHz, DMSO) δ 165.9, 155.6, 153.5, 149.5, 142.0, 141.3, 129.6, 129.0, 127.4, 118.3, 52.2, 45.9, 27.2. HRMS (ESI) m/z: [M + H]+ calcd for $C_{15}H_{15}ClN_5O_2$; 332.091 found, 332.091.

3-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)benzoic acid (29)

The corresponding methyl ester **28** (0.2 g, 0.602 mmol) was dissolved in dioxane (6 mL) followed by the addition of 38 % HCl (4 mL). The reaction mixture was refluxed for 5 hours. The reaction mixture was cooled to 0 ℃ and left for 4 hours at this temperature. The white precipitate was filtered off and dried on air. The carboxylic acid **29** was isolated as a white solid (164 mg, 85 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.52 (s, 1H), 8.48 – 8.38 (bs, 1H), 7.92 – 7.88 (m, 2H), 7.35 (d, J = 8.1 Hz, 2H), 5.47 (s, 2H), 2.93 (s, 3H); ^{13}C NMR (101 MHz, DMSO) δ 167.0, 155.1, 154.0, 149.3, 141.2, 141.0, 130.4, 129.8, 127.5, 116.8, 46.4, 27.3. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{13}ClN_5O_2$; 318.075 found, 318.075.

(4-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)phenyl)methanol (30)

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.2 g, 1.06 mmol) and (4-(bromomethyl)phenyl)methanol (0.212 g, 1.06 mmol) that was prepared following reported procedure [43]. The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 3 : 1 → 5 : 1) and the desired compound was obtained as a white solid (0.12 g, 37 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.83 (s, 1H), 8.74 (s, 1H), 7.32 – 7.27 (m, 4H), 5.47 (s, 2H), 5.24 – 5.07 (bs, 1H), 4.47 (s, 2H); ^{13}C NMR (101 MHz, DMSO) δ 153.4, 151.1, 149.8, 148.4, 142.6, 133.9, 130.5, 127.5, 126.8, 62.5, 47.0.

The final compound **30** was prepared following the General method 2 from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.05 g, 0.264 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/MeOH = 1 : 0.1) and the desired compound was obtained as a white solid (0.075 g, 92 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.22 (s, 1H), 8.23 – 8.18 (bs, 1H), 7.28 (d, J = 8.1 Hz, 2H), 7.22 (d, J = 8.2 Hz, 2H), 5.31 (s, 2H), 5.17 (t, J = 5.7 Hz, 1H), 4.45 (d, J = 5.7 Hz, 1H), 2.91 (d, J = 4.6 Hz, 3H); 13C NMR (101 MHz, DMSO) δ ^{13}C NMR (101 MHz,

DMSO) δ 155.6, 153.4, 149.5, 142.2, 141.2, 135.1, 127.2, 126.8, 118.3, 62.6, 46.2, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{14}ClN_5O$; 304.096 found, 304.096

N-(4-chloro-2-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)phenyl) methanesulfonamide (31)

To a solution of N-(4-chloro-2-(hydroxymethyl)phenyl)methanesulfonamide (0.747 g, 3.17 mmol) in DCM (6.3 mL), prepared following reported procedure[44], was added $SOCl_2$ (0.342 mL, 4.75 mmol) dropwise. The reaction mixture was stirred at rt for 90 min. The volatiles were removed *in vacuo* and the crude product was dissolved in DMF (6.2 mL), followed by addition of 2,6-dichloropurine **44** (0.599 g, 3.17 mmol) and K_2CO_3 (0.649 g, 4.71 mmol). The reaction was quenched by aq. sol. of NH_4Cl after 5 hours. The reaction mixture was extracted into EtOAc (3 x 12 mL) and combined organic layers were dried over $MgSO_4$, filtrated and evaporated. The residue was dissolved in EtOH (4 mL) and 33 % $MeNH_2$ in EtOH (1 mL) was added. The reaction mixture was stirred at rt for 1 hour. After the reaction completion, the volatiles were removed *in vacuo*. The crude compound was purified using flash column chromatography (SiO_2; EtOAc/Hept = 4 : 1 → 6 : 1) and the desired compound was obtained as a white solid (0.074 g, 6 % after three steps). ^{1}H NMR (400 MHz, DMSO – d^6) δ 9.61 (s, 1H), 8.32 - 8.27 (m, 1H), 8.21 (s, 1H), 7.45 – 7.40 (m, 2H), 6.98 – 6.96 (bs, 1H), 5.43 (s, 2H), 3.10 (s, 3H), 2.93 (d, J = 4.6 Hz, 3H); ^{13}C NMR (101 MHz, DMSO – d^6) δ 155.6, 153.3, 149.4, 141.3, 135.6, 133.8, 133.1, 128.8, 128.6, 118.2, 42.6, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{15}Cl_2N_6O_2S$; 401.035 found, 401.035.

3-chloro-5-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)benzoic acid (32)

The corresponding methyl ester **33** (0.06 g, 0.161 mmol) was dissolved in dioxane (2.5 mL) followed by the addition of 38 % HCl (1 mL). The reaction mixture was refluxed for 5 hours. The reaction mixture was cooled to 0 °C and left for 4 hours at this temperature. The white precipitate

was filtered off and dried on air. The carboxylic acid **32** was isolated as a white solid (0.045 g, 79 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.36 (s, 1H), 8.35 – 8.28 (bs, 1H), 7.83 (t, J = 1.8 Hz, 1H), 7.77 (t, J = 1.6 Hz, 1H), 7.68 (t, J = 1.9 Hz, 1H), 5.44 (s, 2H), 2.92 (d, J = 3.6 Hz, 3H). ^{13}C NMR (101 MHz, DMSO – d^6) δ 165.7, 155.4, 153.6, 149.3, 141.1, 139.5, 133.6, 133.3, 131.7, 128.3, 126.9, 117.8, 45.5, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{12}Cl_2N_5O_2$; 352.036 found, 352.036.

methyl 3-chloro-5-((2-chloro-6-(methylamino)-9H-purin-9-yl)methyl)benzoate (33)

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.459 g, 2.43 mmol) and methyl 3-(bromomethyl)-5-chlorobenzoate (0.641 g, 2.43 mmol) that was prepared following reported procedure [45]. The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1:1) and the desired compound was obtained as a white solid (0.51 g, 56 %). ^{1}H NMR (400 MHz, $CDCl_3$ - d) δ 8.10 (s, 1H), 8.02 – 8.00 (m, 1H), 7.88 – 7.86 (m, 1H), 7.48 (t, J = 1.9 Hz, 1H), 5.44 (s, 2H), 3.92 (s, 3H). ^{13}C NMR (101 MHz, $CDCl_3$ - d) δ 165.2, 153.7, 153.2, 152.4, 145.3, 136.5, 135.9, 133.1, 132.3, 130.8, 130.5, 127.2, 52.9, 47.2.

The final compound **33** was prepared following the General method 2 from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.337 g, 0.906 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 4 : 1) and the desired compound was obtained as a white solid (0.176 g, 53 %). %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.32 – 8.24 (m, 2H), 7.86 – 7.81 (m, 2H), 7.71 – 7.70 (m, 1H), 5.44 (s, 2H), 3.84 (s, 3H), 2.92 (d, J = 4.6 Hz, 3H); ^{13}C NMR (101 MHz, DMSO – d^6) δ 164.7, 155.6, 153.5, 149.4, 141.1, 139.8, 133.7, 132.2, 132.0, 128.2, 126.8, 118.3, 52.6, 45.4, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{15}H_{14}Cl_2N_5O_2$; 366.052 found, 366.052.

2-chloro-9-(3-chloro-4-methoxybenzyl)-N-methyl-9H-purin-6-amine (34)

The compound was prepared following the general procedure using 2,6-dichloropurine **44** (0.566 g, 3.00 mmol) and 2-chloro-4-(chloromethyl)-1-methoxybenzene (0.573 g, 3.00 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 3 : 2) and the desired compound was obtained as a white solid (0.71 g, 69 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 8.05 (s, 1H), 7.34 (d, J = 2.2 Hz, 1H), 7.21 (dd, J = 8.5, 2.3 Hz, 1H), 6.90 (d, J = 8.5 Hz, 1H), 5.32 (s, 2H), 3.88 (s, 3H); ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 155.7, 153.3, 153.1, 152.0, 145.5, 130.8, 130.2, 128.0, 127.0, 123.4, 112.6, 56.4, 47.2.

The final compound **34** was prepared following the General method 2 from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.134 g, 0.397 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 2 : 3 → 1 : 4) and the desired compound was obtained as a white solid (0.08 g, 59 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.24 (s, 1H), 8.26 – 8.18 (bs, 1H), 7.44 (d, *J* = 2.2 Hz, 1H), 7.25 (dd, *J* = 8.5, 2.2 Hz, 1H), 7.12 (d, *J* = 8.5 Hz, 1H), 5.26 (s, 2H), 3.82 (s, 3H), 2.91 (d, *J* = 4.6 Hz, 3H); ^{13}C NMR (101 MHz, DMSO – d^6) δ 155.5, 154.2, 153.4, 149.3, 141.0, 129.8, 129.3, 127.9, 121.0, 118.3, 113.0, 56.1, 45.3, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{14}Cl_2N_5O$; 338.057 found, 338.057.

2-chloro-9-(3,5-dichloro-4-methoxybenzyl)-N-methyl-9H-purin-6-amine (35)

3,5-dichloro-4-methoxybenzoic acid (0.500 g, 2.26 mmol) was dissolved in THF (5 mL) and BH_3 . DMS (0.529 mL, 5.65 mmol) was added dropwise. The reaction mixture was stirred under dinitrogen atmosphere at rt overnight. After the reaction completion (monitored by TLC), the reaction was quenched by aq. sol. of $NaHCO_3$. The mixture was extracted into EtOAc (3 x 5 mL). The combined organic layers were dried over $MgSO_4$, filtrated and evaporated. The resulting alcohol was dissolved in DCM (3 mL) together with one drop of DMF, followed by addition of $SOCl_2$ (0.261 mL, 3.63 mmol). The reaction mixture was stirred at rt overnight. The volatiles were

removed *in vacuo* and the resulting 1,3-dichloro-5-(chloromethyl)-2-methoxybenzene was used in the next step without further purification.

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.457 g, 2.42 mmol) and 1,3-dichloro-5-(chloromethyl)-2-methoxybenzene (0.449 g, 2.54 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1 : 1) and the desired compound was obtained as a white solid (0.325 g, 35 %). ^{1}H NMR (400 MHz, DMSO – d^6) δ 8.81 (s, 1H), 7.54 (s, 2H), 5.46 (s, 2H), 3.80 (t, 3H); ^{13}C NMR (101 MHz, DMSO – d^6) δ 153.5, 151.4, 151.1, 149.8, 148.3, 133.7, 130.7, 128.8, 128.5, 60.6, 45.6.

The final compound **35** was prepared following the General method 2 for S_NAr starting from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.080 g, 0.211 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1.1 : 1) and the desired compound was obtained as a white solid (0.064g, 81 %). ^{1}H NMR (400 MHz, , DMSO – d^6) δ 8.29 – 8.23 (bs, 1H), 8.25 (s, 1H), 7.45 (s, 2H), 5.31 (s, 2H), 3.80 (s, 3H), 2.92 (d, J = 4.7 Hz, 1H. ^{13}C NMR (101 MHz, DMSO – d^6) δ 155.6, 153.5, 151.2, 149.3, 141.1, 134.8, 128.5, 128.5, 118.3, 60.6, 44.9, 27.2. LRMS (ESI) m/z: [M + H]+ calcd for $C_{14}H_{13}Cl_3N_5O$; 372.018 found, 372.018.

9-(3-chlorobenzyl)-2-fluoro-N-methyl-9H-purin-6-amine (36) ZA540b and 6-chloro-9-(3-chlorobenzyl)-N-methyl-9H-purin-2-amine (39)

The N^9 alkylation was performed following the General procedure 1 using 6-chloro-2-fluoro-9H-purine **49** (0.578 g, 3.35 mmol) and 1-(bromomethyl)-3-chlorobenzene **50** (0.688 g, 3.35 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 1 : 1) and the desired compound was obtained as a white solid (0.178 g, 18 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 8.08 (s, 1H), 7.36 – 7.28 (m, 3H), 7.20 (dt, J = 6.7, 1.9 Hz, 1H), 5.36 (s, 2H). ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 157.66 (d, J = 220.8 Hz), 153.73 (d, J = 16.9 Hz), 153.24 (d, J

= 17.5 Hz), 145.46 (d, *J* = 3.3 Hz), 136.1, 136.4, 130.8, 130.34 (d, *J* = 4.9 Hz), 129.5, 128.2, 126.2, 47.5.

The final compounds **36** and **39** were prepared following the General method 2 for S_NAr starting from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.080 g, 0.211 mmol). The reaction provided both products that were separated using flash column chromatography (SiO_2; EtOAc/Hept = 1.2 : 1 to 2:1) and the desired compound was obtained as a white solids (**36:** 0.021 g, 20 %, **39**: 0.042 g, 40 %). **36:** ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 7.66 (s, 1H), 7.30 – 7.15 (m, 3H), 7.17 – 7.14 (m, 1H), 6.13 – 6.05 (bs, 1H), 5.26 (s, 2H), 3.24 – 3.09 (bs, 3H). ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 159.98 (d, J = 208.6 Hz), 157.31 (d, *J* = 20.5 Hz), 139.63 (d, *J* = 3.1 Hz), 137.5, 135.1, 130.5, 128.9, 127.9, 126.0, 118.2, 46.7, 27.8. ^{19}F NMR (376 MHz, $CDCl_3$ - *d*) δ -49.37. LRMS (ESI) m/z: [M + H]+ calcd for $C_{13}H_{12}ClFN_5$; 292.076 found, 292.077. **39**: ^{1}H NMR (400 MHz, CDCl3) δ 7.70 (s, 1H), 7.34 – 7.27 (m, 3H), 7.18 – 7.14 (m, 1H), 5.28 – 5.22 (bs, 1H), 5.23 (s, 2H), 3.04 (d, *J* = 5.0 Hz, 1H). ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 160.0, 154.0, 151.5, 141.2, 137.5, 135.1, 130.5, 128.9, 128.3, 126.1, 124.4, 46.7, 29.0. LRMS (ESI) m/z: [M + H]+ calcd for $C_{13}H_{12}Cl_2N_5$; 308.046 found, 308.048.

2-chloro-9-(3-chlorobenzyl)-N-cyclopropyl-9H-purin-6-amine (37)

The compound was prepared following general method for S_NAr starting from corresponding 9-alkyl-2,6-dichloro-9H-purine (0.07 g, 0.223 mmol) and cyclopropylamine (0.03 mL, 0.433 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 3 : 1) and the desired compound was obtained as a white solid (0.071 g, 95 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 7.66 (s, 1H), 7.32 – 7.28 (m, 2H), 7.24 (s, 1H), 7.17 – 7.12 (m, 1H), 6.18 – 5.95 (bs, 1H), 5.29 (s, 2H), 3.25 – 2.98 (bs, 1H), 1.87 (s, 1H), 0.96 – 0.91 (m, 2H), 0.67 – 0.63 (m, 2H); ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 156.6, 155.0, 140.00, 137.4, 135.1, 130.6, 128.9, 128.0, 126.1,

118.7, 46.7, 24.2, 7.7. LRMS (ESI) m/z: [M + H]+ calcd for $C_{15}H_{14}Cl_2N_5$; 334.062 found, 334.062.

2,6-dichloro-9-(3-chlorobenzyl)-9H-purine (38)

The N^9 alkylation was performed following the General procedure 1 using 2,6-dichloropurine **44** (0.5 g, 2.65 mmol) and 1-bromomethyl-3-chlorbenzen (0.543 g, 2.65 mmol). The crude product was purified using flash column chromatography (SiO_2; EtOAc/Hept = 2 : 1) and the desired compound was obtained as a white solid (0.495 g, 59 %). ^{1}H NMR (400 MHz, $CDCl_3$ - *d*) δ 8.07 (s, 1H), 7.37 – 7.29 (m, 3H), 7.19 (dt, J = 6.8, 1.8 Hz, 1H), 5.39 (s, 2H). ^{13}C NMR (101 MHz, $CDCl_3$ - *d*) δ 153.6, 153.2, 152.3, 145.4, 136.1, 135.5, 130.9, 130.8, 129.5, 128.2, 126.2, 47.5. LRMS (ESI) m/z: [M + H]+ calcd for $C_{12}H_8Cl_3N_4$; 312.981 found, 312.982.

ASSOCIATED CONTENT

Supplementary information – Figures with dose-response curves for antiproliferative, TR-FRET and thermal shift measurements. ^{1}H, ^{19}F and ^{13}C NMR spectra of final molecules and the HPLC purity trace.

Accession codes – The PDB codes of YTHDC1 in complex with the following compounds are listed herein: 8Q2S (4), 8Q2T (5), 8Q2U (6), 8Q2V (7), 8Q2W (8), 8Q2X (10), 8Q2Y (11), 8Q31 (12), 8Q32 (13), 8Q33 (15), 8Q35 (16), 8Q37 (18), 8Q38 (19), 8Q39 (21), 8Q3A (22), 8Q3G (23), 8Q4M (25), 8Q4N (26), 8Q4P (27), 8Q4Q (29), 8Q4R (30), 8Q4T (31), 8Q4U (36), 8Q4V (37), 8Q4W (40).

AUTHOR INFORMATION

Corresponding Author

Amedeo Caflisch − Department of Biochemistry, University of Zurich, Zurich CH-8057, Switzerland; orcid.org/0000-0002-2317-6792; Phone: +41 44 635 55 21; Email:caflisch@bioc.uzh.ch

Authors

František Zálešák − Department of Biochemistry, University of Zurich, Zurich CH-8057, Switzerland; orcid.org/0000-0003-1469-5248

Francesco Nai − Department of Biochemistry, University of Zurich, Zurich CH-8057, Switzerland; orcid.org/0000-0002-4258-3174

Rajiv K. Bedi − Department of Biochemistry, University of Zurich, Zurich CH-8057, Switzerland; orcid.org/0000-0002-8193-9006

Yaozong Li − Department of Biochemistry, University of Zurich, Zurich CH-8057, Switzerland; orcid.org/0000-0002-5796-2644

Marcin Herok − Department of Biochemistry, University of Zurich, Zurich CH-8057, Switzerland; orcid.org/0000-0003-0294-4040

Elena Bochenkova − Department of Biochemistry, University of Zurich, Zurich CH-8057, Switzerland; orcid.org/0000-0002-2314-6673

Francesco Errani − Department of Biochemistry, University of Zurich, Zurich CH-8057, Switzerland; orcid.org/0009-0004-4658-9022

The manuscript was written through the contributions of all authors. All authors have approved the final version of the manuscript.

Notes The authors declare no competing financial interest

ACKNOWLEDGEMENTS

We thank Beat Blattmann for help with crystallization and thePaul Scherrer Institute(PSI) staff and PXI and PXIII beamline scientists for acquisition of diffraction data. This work was supported financially by the Swiss National Science Foundation to A.C. (Grant Number 310030_212195) and the Swiss Cancer Research Foundation to A.C. (Grant number KFS 5748-02-2023). We also thank the Olga Mayenfisch Stiftung for financial support.

ABBREVIATIONS

HTRF assay, Homogeneous Time Resolved Fluorescence; DIPEA, *N,N*-Diisopropylethylamine; rt, room temperature; S_NAr, nucleophilic aromatic substitution; pTSA, *para*-toluenesulfonic acid; MS, molecular sieves; DMF, *N,N*-dimethylformamide; TFAA, trifluoroacetic acid anhydride; Py, pyrimidine; MsCl, mesyl chloride; TsCl, tosyl chloride; TFA, trifluoroacetic acid; COMU, (1-Cyano-2-ethoxy-2-oxoethylidenaminooxy)dimethylamino-morpholino-carbenium hexafluorophosphate.

REFERENCES

(1) Jiang, X.; Liu, B.; Nie, Z.; Duan, L.; Xiong, Q.; Jin, Z.; Yang, C.; Chen, Y. The Role of m^6A modification in the biological functions and diseases. *Signal Transduct. Target. Ther.* **2021**, *6* (1), 74. DOI: 10.1038/s41392-020-00450-x

(2) Barbieri, I.; Kouzarides, T. Role of RNA modifications in cancer. *Nat. Rev. Cancer* **2020**, *20* (6), 303. DOI: 10.1038/s41568-020-0253-2.

(3) Wiener, D.; Schwartz, S. The Epitranscriptome beyond m^6A. *Nat. Rev. Genet.* **2021**, *22*, 119. DOI: 10.1038/s41576-020-00295-8

(4) Boo, S. H.; Kim, Y. K. The Emerging Role of RNA Modifications in the Regulation of mRNA Stability. *Exp. Mol. Med.* **2020**, *52*, 400. DOI: 10.1038/s12276-020-0407-z.

(5) Song, D.; Shyh-Chang, N. An RNA Methylation Code to Regulate Protein Translation and Cell Fate. *Cell Prolif.* **2022**, *55* (5). DOI: 10.1111/cpr.13224.

(6) Ozato, K.; Dodagatta-Marri, E.; Zhou, Y.; Zhang, C.; Fu, J. A Review in Research Progress Concerning M6A Methylation and Immunoregulation. *Front. Immunol.* **2019**, *1*, 922. DOI: 10.3389/fimmu.2019.00922.

(7) Liu, J.; Yue, Y.; Han, D.; Wang, X.; Fu, Y.; Zhang, L.; Jia, G.; Yu, M.; Lu, Z.; Deng, X.; Dai, Q.; Chen, W.; He, C. A METTL3–METTL14 Complex Mediates Mammalian Nuclear RNA N^6-Adenosine Methylation. *Nat. Chem. Bio.* **2014**, *10*, 93 – 95. DOI: 10.1038/nchembio.1432.

(8) Xu, Y.; Zhang, Y.; Luo, Y.; Qiu, G.; Lu, J.; He, M.; Wang, Y. Novel Insights into the METTL3-METTL14 Complex in Musculoskeletal Diseases. *Cell Death Discov.* **2023**, *9*, 170. DOI: 10.1038/s41420-023-01435-9.

(9) Zaccara, S.; Ries, R. J.; Jaffrey, S. R. Reading, Writing and Erasing mRNA Methylation. *Nat. Rev. Mol. Cell. Biol*. **2019**, *20*, 608. DOI: 10.1038/s41580-019-0168-5.

(10) Zhao, X.; Yang, Y.; Sun, B.-F.; Shi, Y.; Yang, X.; Xiao, W.; Hao, Y.-J.; Ping, X.-L.; Chen, Y.-S.; Wang, W.-J.; Jin, K.-X.; Wang, X.; Huang, C.-M.; Fu, Y.; Ge, X.-M.; Song, S.-H.; Jeong, H. S.; Yanagisawa, H.; Niu, Y.; Jia, G.-F.; Wu, W.; Tong, W.-M.; Okamoto, A.; He, C.; Rendtlew Danielsen, J. M.; Wang, X.-J.; Yang, Y.-G. FTO-Dependent Demethylation of N^6-Methyladenosine Regulates mRNA Splicing and Is Required for Adipogenesis. *Cell. Res.* **2014**, *24*, 1403. DOI: 10.1038/cr.2014.151.

(11) Dolbois, A.; K. Bedi, R.; Bochenkova, E.; Müller, A.; V. Moroz-Omori, E.; Huang, D.; Caflisch, A. 1,4,9-Triazaspiro[5.5]Undecan-2-One Derivatives as Potent and Selective METTL3 Inhibitors. *J. Med. Chem.* **2021**, *64* (17), 12738. DOI: 10.1021/acs.jmedchem.1c00773.

(12) Moroz-Omori, E. V; Huang, D.; Kumar Bedi, R.; Cheriyamkunnel, S. J.; Bochenkova, E.; Dolbois, A.; Rzeczkowski, M. D.; Li, Y.; Wiedmer, L.; Caflisch, A. METTL3 Inhibitors for Epitranscriptomic Modulation of Cellular Processes. *ChemMedChem* **2021**, *16*, 3035. DOI: 10.1002/cmdc.202100291.

(13) Xu, Y.; Zhang, W.; Shen, F.; Yang, X.; Liu, H.; Dai, S.; Sun, X.; Huang, J.; Guo, Q. YTH Domain Proteins: A Family of m6A Readers in Cancer Progression. *Front. Oncol.* **2021**, *11*. DOI:10.3389/fonc.2021.629560.

(14) Shi, R.; Ying, S.; Li, Y.; Zhu, L.; Wang, X.; Jin, H. Linking the YTH Domain to Cancer: The Importance of YTH Family Proteins in Epigenetics. *Cell Death Dis.* **2021**, *12*, 346. DOI: 10.1038/s41419-021-03625-8.

(15) Sikorski, V.; Selberg, S.; Lalowski, M.; Karelson, M.; Kankuri, E. The Structure and Function of YTHDF Epitranscriptomic m6A Readers. **2023**. *Trends Pharmacol. Sci.* 2023, *44* (6), 335. DOI: 10.1016/j.tips.2023.03.004.

(16) Fu, Y.; Zhuang, X. M6A-Binding YTHDF Proteins Promote Stress Granule Formation. *Nat. Chem. Biol*. **2020**, *16*, 955. DOI: 10.1038/s41589-020-0524-y.

(17) Qiao, Y.; Sun, Q.; Chen, X.; He, L.; Wang, D.; Su, R.; Xue, Y.; Sun, H.; Wang, H. Nuclear m^6A Reader YTHDC1 Promotes Muscle Stem Cell Activation/Proliferation by Regulating MRNA Splicing and Nuclear Export. *eLife* **2023**, *12*, 82703. DOI: 10.7554/eLife.82703.

(18) Yan, H.; Zhang, L.; Cui, X.; Zheng, S.; Li, R. Roles and Mechanisms of the m^6A Reader YTHDC1 in Biological Processes and Diseases. *Cell Death Discov.*. **2022**, 8, 237. DOI: 10.1038/s41420-022-01040-2.

(19) Tan, B.; Zhou, K.; Liu, W.; Prince, E.; Qing, Y.; Li, Y.; Han, L.; Qin, X.; Su, R.; Pokharel, S. P.; Yang, L.; Zhao, Z.; Shen, C.; Li, W.; Chen, Z.; Zhang, Z.; Deng, X.; Small, A.; Wang, K.; Leung, K.; Chen, C.-W.; Shen, B.; Chen, J. RNA N^6-Methyladenosine Reader YTHDC1 Is Essential for TGF-Beta-Mediated Metastasis of Triple Negative Breast Cancer. *Theranostics* **2022**, *12* (13), 5727. DOI: 10.7150/thno.71872.

(20) Cheng, Y.; Xie, W.; Pickering, B. F.; Chu, K. L.; Savino, A. M.; Yang, X.; Luo, H.; Nguyen, D. T.; Mo, S.; Barin, E.; Velleca, A.; Rohwetter, T. M.; Patel, D. J.; Jaffrey, S. R.; Kharas, M. G. N6-Methyladenosine on mRNA Facilitates a Phase-Separated Nuclear Body That Suppresses Myeloid Leukemic Differentiation. *Cancer Cell* **2021**, *39* (7), 958. DOI: 10.1016/j.ccell.2021.04.017.

(21) Li, Y.; Bedi, R. K.; Nai, F.; Von Roten, V.; Dolbois, A.; Zálešák, F.; Nachawati, R.; Huang, D.; Caflisch, A. Structure-Based Design of Ligands of the M6A-RNA Reader YTHDC1. *Eur. J. Med. Chem. Rep.* **2022**, *5*, 100057. DOI: 10.1016/J.EJMCR.2022.100057.

(22) Nai, F.; Nachawati, R.; Zálešák, F.; Wang, X.; Li, Y.; Caflisch, A. Fragment Ligands of the m^6A-RNA Reader YTHDF2. *ACS Med. Chem. Lett.* **2022**, *13*, 41. DOI: 10.1021/acsmedchemlett.2c00303.

(23) Hong, Y.-G.; Yang, Z.; Chen, Y.; Liu, T.; Zheng, Y.; Zhou, C.; Wu, G.-C.; Chen, Y.; Xia, J.; Wen, R.; Liu, W.; Zhao, Y.; Chen, J.; Gao, X.; Chen, Z. The RNA m^{6}A Reader YTHDF1 Is Required for Acute Myeloid Leukemia Progression. *Cancer Res.* **2023**, *83* (6), 845. DOI: 10.1158/0008-5472.can-21-4249.

(24) Zou, Z.; Wei, J.; Chen, Y.; Kang, Y.; Shi, H.; Chen, S.; Zhou, Y.; Sepich-Poore, C.; Zhuang, X.; Zhou, X.; Jiang, H.; Wen, Z.; Jin, P.; Luo, C.; He, C. FMRP Phosphorylation Modulates Neuronal Translation through YTHDF1 1. *bioRxiv* **2022**. DOI: 10.1101/2022.11.29.518448.

(25) Micaelli, M.; Dalle Vedove, A.; Cerofolini, L.; Vigna, J.; Sighel, D.; Zaccara, S.; Bonomo, I.; Poulentzas, G.; Rosatti, F.; Cazzanelli, G.; Alunno, L.; Belli, R.; Peroni, D.; Dassi, E.; Murakami, S.; Jaffrey, S. R.; Fragai, M.; Mancini, I.; Lolli, G.; Quattrone, A.; Provenzani, A. Small-Molecule Ebselen Binds to YTHDF Proteins Interfering with the Recognition of N^6-Methyladenosine-Modified RNAs. *ACS Pharmacol. Transl. Sci.* **2022**, *5*(10), 872. DOI: 10.1021/acsptsci.2c00008.

(26) Yang, S.; Zhang, H.; Li, Y.; Wang, F.; Lin, G.; Niu, T.; Li, H.; Yi, Y.; Zhou, H.; Yang, R.; Yao, R.; Zhou, P.; Li, Y.; Wu, M.; Chen, M.-X.; Xu, H. Discovery of a Selective YTHDC1 Inhibitor That Targets Acute Myeloid Leukemia. *Res. Sq.* **2023.** DOI: 10.21203/rs.3.rs-2644364/v1.

(27) Li, Y.; Kumar Bedi, R.; Wiedmer, L.; Sun, X.; Huang, D.; Caflisch, A. Atomistic and Thermodynamic Analysis of N^6-Methyladenosine (M6A) Recognition by the Reader Domain of YTHDC1. *J. Chem. Theory Comput.* **2021**, *17* (2), 1240. DOI: 10.1021/acs.jctc.0c01136.

(28) Sander, T.; Freyss, J.; Korff, V.; Rufener, C. DataWarrior: An Open-Source Program For Chemistry Aware Data Visualization And Analysis. *J. Chem. Inf. Model.* **2015**, *5*(2), 460. DOI: 10.1021/ci500588j.

(29) Bakkestuen, A. K.; Gundersen, L. L. Regioselective N-9 Arylation of Purines Employing Arylboronic Acids in the Presence of Cu(II). *Tetrahedron Lett.* **2003**, *44* (16), 3359. DOI: 10.1016/S0040-4039(03)00576-8.

(30) Compounds purchased from https://chem-space.com/

(31) Wiedmer, L.; Eberle, S. A.; Bedi, R. K.; Śledź, P.; Caflisch, A. A Reader-Based Assay for m^{6}A Writers and Erasers. *Anal. Chem.* **2019**, *91* (4), 3078. DOI: 10.1021/acs.analchem.8b05500.

(32) Bedi, R. K.; Huang, D.; Wiedmer, L.; Li, Y.; Dolbois, A.; Aleksandra Wojdyla, J.; Sharpe, M. E.; Caflisch, A.; Sledz, P. Selectively Disrupting m^{6}A-Dependent Protein−RNA Interactions with Fragments. *ACS Chem. Biol.* **2020**, *15*, 625. DOI: 10.1021/acschembio.9b00894.

(33) Kabsch, W. *Acta Cryst.* **2010**, *66* (2), 125. DOI: 10.1107/S0907444909047337.

(34) McCoy, A. J.; Grosse-Kunstleve, R. W.; Adams, P. D.; Winn, M. D.; Storoni, L. C.; Read, R. J. Phaser Crystallographic Software. *J. Appl. Cryst.* **2007**, *40* (4), 658. DOI: 10.1107/S0021889807021206.

(35) Liebschner, D.; Afonine, P. V; Baker, M. L.; Bunkó, G.; Chen, V. B.; Croll, T. I.; Hintze, B.; Hung, L.-W.; Jain, S.; Airlie, }; Mccoy, J.; Moriarty, N. W.; Oeffner, R. D.; Poon, B. K.; Prisant, M. G.; Read, R. J.; Richardson, J. S.; Richardson, D. C.; Sammito, M. D.; Sobolev, O. V; Stockwell, D. H.; Terwilliger, T. C.; Urzhumtsev, A. G.; Videau, L. L.; Williams, C. J.; Adams, P. D. Macromolecular Structure Determination Using X-Rays, Neutrons and Electrons: Recent Developments in Phenix. *Acta Cryst.* **2019**, *75*, 861. DOI: 10.1107/S2059798319011471.

(36) Emsley, P.; Lohkamp, B.; Scott, W. G.; Cowtan, K. Biological Crystallography Features and Development of Coot. *Acta Cryst.* **2010**, *66*, 486. DOI: 10.1107/S0907444910007493.

(37) Afonine, P. V; Grosse-Kunstleve, R. W.; Echols, N.; Headd, J. J.; Moriarty, N. W.; Mustyakimov, M.; Terwilliger, T. C.; Urzhumtsev, A.; Zwart, P. H.; Adams, P. D. Biological Crystallography Towards Automated Crystallographic Structure Refinement with Phenix.Refine. *Acta Cryst.* **2012**, *68*, 352. DOI: 10.1107/S0907444912001308.

(38) Robins, R. K.; Holum, L. B.; Furcht, F. W. Synthesis of Some 3-Methyl-5,7-Substituted Pyrazolo[4,3,*d*]-pyrimidines *J. Org. Chem.* **1956**, 21 (8), 833. DOI: 10.1021/jo01114a001

(39) Zhang, Q.; Hu, X.; Wan, G.; Wang, J.; Li, L.; Wu, X.; Liu, Z.; Yu, L. Discovery of 3-(((9H-Purin-6-yl)amino)methyl)-4,6-Dimethylpyridin-2(1H)-One Derivatives as novel tubulin polymerization inhibitors for treatment of cancer. *Eur. J. Med. Chem.* **2019**, 184, 111728. DOI: 10.1016/j.ejmech.2019.111728.

(40) Li, X.; Han, J.; Bujaranipalli, S.; He, J.; Kim, E. Y.; Kim, H.; Im, J. H.; Cho, W.-J. Structure-Based Discovery and Development of Novel O-GlcNAcase Inhibitors for the Treatment of Alzheimer's Disease. *Eur. J. Med. Chem.* **2022**, *238*, 114444. DOI: 10.1016/j.ejmech.2022.114444.

(41) Hovey, M. T.; Check, C. T.; Sipher, A. F.; Scheidt, K. A. N-Heterocyclic-Carbene-Catalyzed Synthesis of 2-Aryl Indoles. *Angew. Chem. Int. Ed.* **2014**, *53*, 9603. DOI: 10.1002/anie.201405035.

(42) Amin, S.; S. Hecht, S.; Hoffman, D. Synthesis of Angular Ring Methoxy-5-Methylchrysenes and 5-Methylchrysenols. *J. Org. Chem.* **2002**, *46* (11), 2394–2398. DOI: 10.1021/jo00324a039.

(43) Allard, E.; Delaunay, J.; Cousseau, J. C_{60}^{2-} Chemistry: C_{60} Adducts Bearing Two Ester, Carbonyl, or Alcohol Groups. *Org. Lett.* **2003**, *5* (13), 2239. DOI: 10.1021/ol034545k.

(44) Wu, S.; Liu, C.; Luo, G.; Jin, Z.; Zheng, P.; Chi, Y. R. NHC-Catalyzed Chemoselective Reactions of Enals and Aminobenzaldehydes for Access to Chiral Dihydroquinolines. *Angew. Chem. Int. Ed.* **2019**, *58* (51), 18410. DOI: 10.1002/ANIE.201909479.

(45) Bobileva, O.; Bobrovs, R.; Sirma, E. E.; Kanepe, I.; Bula, A. L.; Patetko, L.; Ramata-Stunda, A.; Grinberga, S.; Jirgensons, A.; Jaudzems, K. 3-(Adenosylthio)Benzoic Acid Derivatives as SARS-CoV-2 Nsp14 Methyltransferase Inhibitors. *Molecules* **2023**, *28* (2), 768. DOI: 10.3390/MOLECULES28020768/S1.