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METTL3 Inhibitors for Epitranscriptomic Modulation of Cellular Processes

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Abstract: The methylase METTL3 is the writer enzyme of the N6 -methyladenosine (m6 A) modification of RNA. Using a structure-based drug discovery approach, we identified a METTL3 inhibitor with potency in a biochemical assay of 280 nM, while its enantiomer is 100 times less active. We observed a dose-dependent reduction in the m6 A methylation level of mRNA in several cell lines treated with the inhibitor already after 16 h of treatment, which lasted for at least 6 days. Importantly, the prolonged incubation (up to 6 days) with the METTL3 inhibitor did not alter levels of other RNA modifications (i. e., m1 A, m6 Am , m7 G), suggesting selectivity of the developed compound towards other RNA methyltransferases. **Keywords:** METTL3; RNA methyltransferase inhibitor; epitranscriptomics; leukemia; m6A.

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Special
Issue

METTL3 Inhibitors for Epitranscriptomic Modulation of Cellular Processes

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The methylase METTL3 is the writer enzyme of the N⁶-methyladenosine (m⁶A) modification of RNA. Using a structure-based drug discovery approach, we identified a METTL3 inhibitor with potency in a biochemical assay of 280 nM, while its enantiomer is 100 times less active. We observed a dose-dependent reduction in the m⁶A methylation

level of mRNA in several cell lines treated with the inhibitor already after 16 h of treatment, which lasted for at least 6 days. Importantly, the prolonged incubation (up to 6 days) with the METTL3 inhibitor did not alter levels of other RNA modifications (i. e., m¹A, m⁶A_m, m⁷G), suggesting selectivity of the developed compound towards other RNA methyltransferases.

Introduction

The finely organized network of gene expression comprising RNA transcription, splicing, transport, translation, and degradation is often perturbed in cancer.^[1,2] In addition to previously known regulators of gene expression, such as epigenetic modifications or miRNAs, the recently discovered layer of regulation based on co- and post-transcriptional RNA modifications gave rise to a new field named epitranscriptomics.^[2,3] While over 160 different RNA modifications have been discovered to date, one of the most abundant modifications, N⁶-methyladenosine or m⁶A (comprising 0.1–0.4% of all adenosine in mRNA), is involved in most of the aspects of RNA regulation, i. e., alternative polyadenylation, splicing (controls about 3% of alternatively spliced exons^[4]), nuclear export, stability, and translation initiation.^[2,3] It is also found in other RNA species, including lncRNAs,^[5] rRNAs,^[6] and snRNAs.^[7] Dysregulated m⁶A deposition is directly involved in the development of acute myeloid leukemia (AML) and lymphomas, difficult-to-treat blood cancers,^[1,8–12] as well being associated with other types of cancer (e. g., bladder, lung, ovarian, colorectal, bone, liver, gastric).^[13–21]

In mRNAs and lncRNAs, most of the m⁶A modifications are installed in [G/A/U][G > A]m⁶AC[U > A > C] consensus sites

(GGm⁶ACU is the most prevalent)^[22] by the so-called m⁶A-METTL complex (MAC), consisting of methyltransferase-like protein 3 (METTL3) and METTL14.^[3] MAC is assisted by a regulatory complex (named MACOM, m⁶A-METTL-associated complex) composed of WTAP, RBM15/B, VIRMA, ZC3H13 and HAKAI.^[23] The crystal structure of the METTL3/14 complex was resolved in 2016,^[24–26] revealing the function of each component. While METTL14 plays a scaffolding role in substrate RNA recognition, forming an RNA-binding groove at the interface of the two subunits, METTL3 carries out the catalytic transfer of the methyl group from a cofactor S-(5'-adenosyl)-L-methionine (SAM) onto the N⁶ atom of the adenine.^[24–26]

To date, there are no cell-permeable inhibitors of METTL3/14 with a disclosed chemical structure except for the universal nucleoside analogue sinefungin, which inhibits most methyltransferases and is not selective for METTL3.^[27,28] Therefore, all published studies of the role of m⁶A in cancer have relied on the knockdown/overexpression of the writers, erasers, and readers. In contrast to this approach, using small-molecule inhibitors preserves the function of the target enzyme to act as a scaffold for protein-protein interactions that would otherwise be disrupted by RNAi, thus enabling discrimination between the enzymatic and structural roles. Indeed, it has been demonstrated that both a catalytically active and a non-functional METTL3 lead to a reduction in p-AKT levels highlighting the m⁶A-independent function of METTL3.^[10] In addition, small-molecule inhibitors enable titration experiments across a range of concentrations to be performed, ranging from mild perturbation to near-complete inhibition of the METTL3 enzymatic activity, which can reveal a spectrum of phenotypes. Generally, inhibitors can inactivate their targets rapidly, allowing precise temporal control over METTL3 function. This is particularly important for studying downstream effects of m⁶A depletion on gene and protein expression and will help to elucidate the m⁶A regulatory network in health and disease.

Here we report a nanomolar inhibitor of METTL3 (UZH1a) which is selective and cell-permeable, while its enantiomer UZH1b is essentially inactive. The crystal structure of the

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complex shows several favourable interactions of the **UZH1a** chemical probe with METTL3, some of which are not observed in the METTL3/SAM complex and thus provide selectivity against other SAM-dependent methyltransferases. Furthermore, the characterization of **UZH1a** in biochemical and cellular assays provides evidence that **UZH1a** has a good potential for the development of a lead compound against METTL3-dependent cancers.

Results and Discussion

Our initial efforts to develop a potent and selective METTL3 inhibitor were focused on screening an adenine-based library, which yielded several hits with micromolar potency in a homogenous time-resolved fluorescence (HTRF) enzyme inhibition assay.^[29] Protein structure-based optimization coupled with compound potency evaluation in our HTRF assay^[30] resulted in the high-nanomolar inhibitor **UZH1a** (Figure 1a). The potency of **UZH1a** in the HTRF assay was 280 nM, while its enantiomer **UZH1b** was 100-fold less active (Figure 1b). We confirmed the binding of **UZH1a** but not its enantiomer **UZH1b** to METTL3 by X-ray crystallography (Figure 1c). The **UZH1a** inhibitor fills the pocket of the adenosine moiety of SAM but not the pocket of the SAM methionine. Besides the favourable van der Waals contacts, there are several hydrogen bonds between polar groups of the **UZH1a** inhibitor and METTL3 (yellow dashed lines in Figure 1c). The pyrimidine moiety is engaged in two hydrogen bonds with the backbone NH groups of Ile378 and Asn549. It is also involved in an edge-to-face π -stacking arrangement with the phenol of Phe534 and π interactions with the amide group of the Asn549 side chain. The hydroxyl group is involved as the donor in a hydrogen bond with the side chain oxygen atom of Asn549 (distance of 2.5 Å) and in water-bridged polar interactions (distance of 2.8 Å) with the backbone NH and CO groups of residues 550 and 535, respectively. There is also an intra-inhibitor hydrogen bond between the amide NH and phenolic oxygen of **UZH1a** (green dashed line in Figure 1c). Notably, the binding of **UZH1a** results in a conformational rearrangement (6 Å displacement of the amino group) of the Lys513 side chain with respect to the Lys513 orientation in the complex of METTL3 with SAM or sinefungin (Figure 1d, PDB code 6Y4G). In detail, the tertiary amino group of **UZH1a** replaces the primary amine of Lys513 in the salt bridge with Asp395. As a consequence, the amine of Lys513 replaces the amino group of SAM in the salt bridge with Glu532. These castling-like changes of ligand/METTL3 and intra-METTL3 ionic interactions are likely to provide selectivity against other SAM-dependent methyltransferases. Indeed, **UZH1a** demonstrated high selectivity towards other protein methyltransferases (as well as several promiscuous protein kinases), with remaining enzymatic activity of over 75% at 10 μ M concentration of **UZH1a** (Table 1).

UZH1a possesses favourable physicochemical properties and is therefore compatible with cell-based experiments. Indeed, its molecular weight is relatively low (558 g/mol), and the octanol-water partition coefficient $\log D_{7.4}$ of 2.6 is optimal for

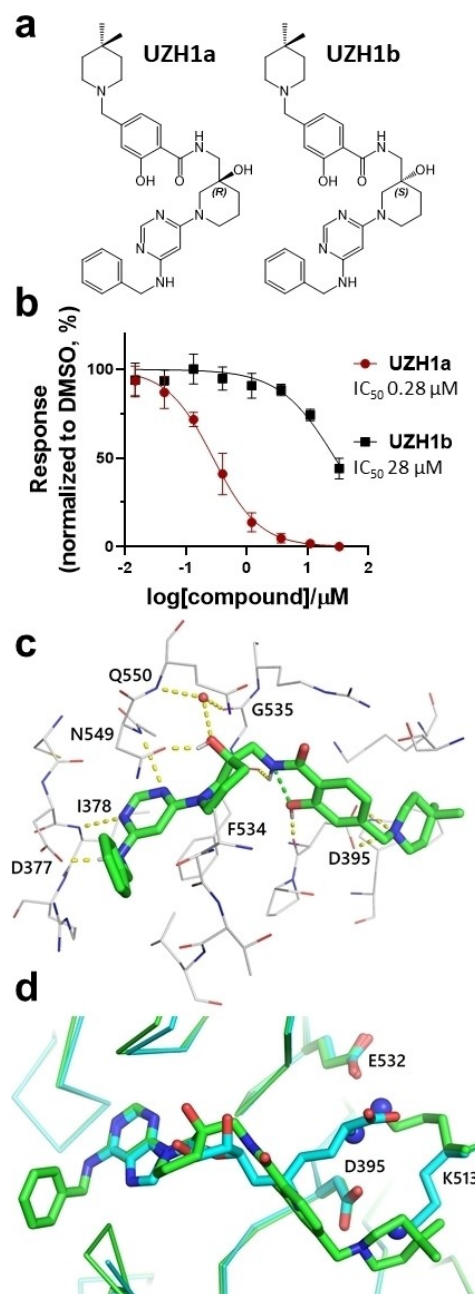


Figure 1. *In vitro* characterization of METTL3 inhibitors. (a) Chemical structures of **UZH1a** and **UZH1b**; absolute configuration was determined by X-ray crystallography. (b) Enzymatic activity assay based on HTRF, mean \pm SD, N=4. (c) Crystal structure of METTL3/14 in complex with **UZH1a** (carbon atoms in green, PDB code 7ACD). Hydrogen bonds (yellow and green dashed lines) and a water molecule in contact with the inhibitor (red sphere) are shown. (d) Overlay of crystal structures of METTL3/14 in complex with **UZH1a** (carbon atoms in green) and METTL3/14 in complex with sinefungin (carbon atoms in cyan, PDB code 6Y4G). The amino groups involved in the castling-like conformational change are highlighted (blue spheres on N atoms).

cell uptake. **UZH1a** was highly permeable ($P_{app} > 1 \cdot 10^{-5}$ cm/s) in a Caco-2 permeability assay used to evaluate intestinal uptake (Table 2). However, the efflux ratio of 2.3 suggests that **UZH1a** may be subject to active efflux. Importantly, the large difference in biochemical potency of **UZH1a** and **UZH1b** (> 100-fold,

Table 1. Kinases and protein methyltransferases selectivity profile of **UZH1a** at 10 μM .

Type	Protein	Remaining activity [%] ^[a]		Control IC ₅₀ [μM]	Control compound ^[b]
		Replicate 1	Replicate 2		
methyltransferase	DOT1L	100	98	0.278	SAH
	G9a	96	99	1.95	SAH
	MLL4 complex	88	89	2.11	SAH
	PRDM9	101	101	3.98	chaetocin
	PRMT1	91	88	0.140	SAH
	SETD2	99	103	7.39	SAH
	SMYD3	88	92	18.7	SAH
kinase	Abl	80	74	–	–
	CDK2/cyclinA	83	72	–	–
	cKit	95	101	–	–
	DDR1	96	85	–	–
	Flt3(D835Y)	80	68	–	–
	Fms	92	102	–	–
	JAK1	101	102	–	–
	PDGFR α	88	81	–	–
	Pim-1	100	95	–	–
	ROCK-I	86	80	–	–

[a] The remaining activity is the percentage of enzymatic activity in the presence of 10 μM **UZH1a** with respect to the buffer solution containing DMSO. The closer to 100% are these values, the weaker the inhibitory potency of **UZH1a**. [b] SAH = S-(5'-adenosyl)-L-homocysteine.

Table 2. Permeability of reference compounds and **UZH1a** in Caco-2 cell assay.

Compound	Direction	P _{app} [10 ⁻⁶ cm/s]		P _{app} (B–A)/P _{app} (A–B)
		Replicate 1	Replicate 2	
Metoprolol	A–B	33.7	31.4	1.1
	B–A	34.7	33.3	
Atenolol	A–B	0.5	0.4	2.0
	B–A	0.9	1.0	
Erythromycin	A–B	0.3	0.2	62.9
	B–A	18.1	17.5	
UZH1a	A–B	12.4	11.8	2.3
	B–A	25.6	28.8	

P_{app}—apparent permeability coefficient, A—apical, B—basal.

Figure 1b) allows for discrimination of non-specific effects due to the chemical properties of the compounds and makes them highly suitable for mechanistic cellular studies.

Higher potency of **UZH1a** in comparison to **UZH1b** in the biochemical assay (Figure 1b) is consistent with higher thermal stabilization of the METTL3/14 construct by **UZH1a** than **UZH1b** in a protein thermal shift assay (Figure 2a, $\Delta T_m = 2.5^\circ\text{C}$ and 0.5°C at 10 μM , respectively). This increased stabilization towards thermal denaturation allowed us to study **UZH1a** binding to its protein target directly in cells. HEK293T cells were transiently transfected with a plasmid encoding METTL3 catalytic domain (residues 356–580) tagged on its N-terminus with ProLabel® (ePL) peptide (42 aa) to investigate the **UZH1a** and **UZH1b** binding in a cellular thermal shift assay based on the InCell Pulse platform (Figure 2b). We observed increased fusion protein abundance at 46 $^\circ\text{C}$ upon incubation with **UZH1a**, but not with **UZH1b**, with EC₅₀ of 4 μM .

In line with the biochemical and cellular target engagement assays, we observed a dose-dependent reduction in m⁶A methylation level in mRNA from the human leukemia cell line MOLM-13 treated with **UZH1a** (IC₅₀ of 7 μM), while **UZH1b** was less active at concentrations up to 100 μM in these cells, as determined by triple-quadrupole LC mass spectrometry (Fig-

ure 3a). However, the effect of the m⁶A level depletion upon **UZH1a** treatment was attenuated by at least one order of magnitude in cells in comparison to the biochemical assay. This could be the consequence of cellular efflux (Table 2) or competition with high intracellular SAM/SAH levels. Interestingly, m⁶A level reduction in mRNAs followed a first-order kinetics, with a maximum m⁶A inhibition level of 70% and a half-decay time τ of 1.8 h (Figure 3b).

The latter value is lower than the average mRNA half-life in cells (≈ 10 h)^[31] and is in line with the previously observed rapid degradation of m⁶A-modified mRNAs.^[4,32,33] We have verified that m⁶A level reduction was not caused by a reduction in METTL3 protein level (Figure 3c,d). Using **UZH1a**, we could, for the first time, directly estimate the average half-life of m⁶A-modified RNA in the cells. This result illustrates an important advantage of chemical inhibition of METTL3 over genetic manipulation, as it allows precise temporal control over METTL3 function due to the rapid cellular penetration of small-molecule probes. One has to bear in mind that this method does not allow the decay of m⁶A-modified mRNA to be distinguished from the potential demethylation of the transcripts *via* cellular ALKBH5. The latter, however, has been shown to affect only a small percentage of mRNAs after transcription.^[4]

We confirmed that **UZH1a** was able to reduce m⁶A/A levels in mRNA fraction not only in the leukemia cell line MOLM-13 but also in at least two other cell lines (*i. e.*, osteosarcoma U2OS cells and immortalized human embryonic kidney cell line HEK293T, Figure 4a). However, the m⁶A level reduction IC₅₀ values were slightly higher in these cell lines in comparison to MOLM-13 cells (9 and 15 μM vs. 7 μM , respectively). Importantly, the ability to inhibit cellular m⁶A levels in mRNA fraction was preserved for at least 6 days (Figure 4b).

Interestingly, siRNA-mediated METTL3 knockdown in both HEK293T and U2OS failed to adequately reduce m⁶A levels in mRNA despite the strong reduction of METTL3 protein level (over 80%, Figure 4c,d). It has previously been observed that

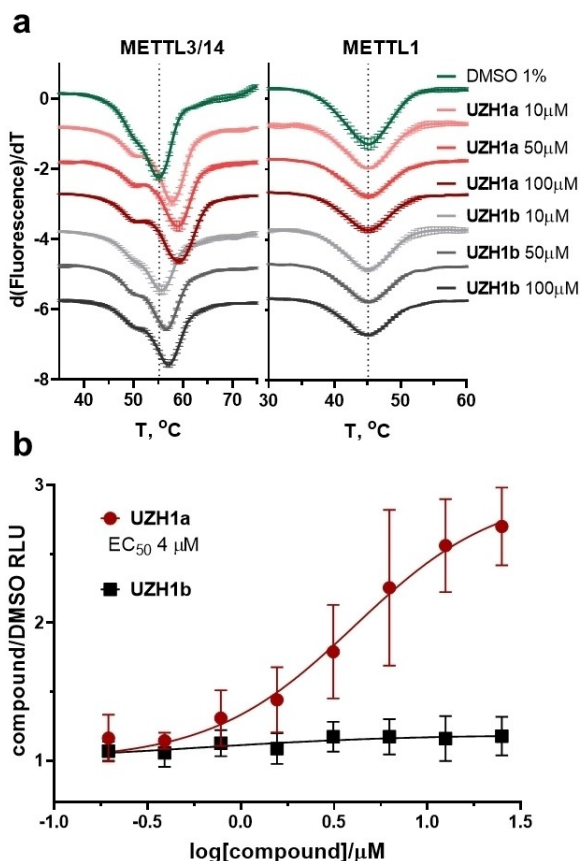


Figure 2. Protein thermal shift assay for METTL3/14 and METTL1 and cellular target engagement assay of UZH1a and UZH1b in HEK293T cells. (a) Results of the differential scanning fluorimetry (protein thermal shift assay) for METTL3/14 and METTL1 constructs. Derivative plot of the thermal denaturation curves of METTL3/14 and METTL1 in the presence of UZH1a (red), UZH1b (gray), or DMSO (green). Protein concentrations were 2 μ M, mean \pm SD, N = 3 (technical replicates). (b) Results of InCELL Pulse assay in HEK293T cells transfected with METTL3 construct and incubated with increasing concentrations of UZH1a and UZH1b for 1 h at 37 °C, followed by heating at 46 °C for 3 min. The soluble METTL3 fraction was quantified in luminescence-based assay, DMSO-treated cells served as normalization control, mean \pm SD, N = 3–4 (biological replicates).

even a low residual level of METTL3 expression can maintain m⁶A/A ratio in cells.^[34] These findings illustrate yet another advantage of our inhibitor that is able to rapidly reduce m⁶A levels up to 75% in a dose-dependent manner in a range of cell lines, and, therefore, comprises a valuable tool to probe various cellular functions of m⁶A-modified mRNAs.

In order to evaluate the selectivity of our METTL3 inhibitor towards other RNA methyltransferases in cells, we analysed the levels of m⁶A as well as other RNA modifications (*i.e.*, m⁶A_m, m¹A, m⁷G) in total RNA. It is known that m⁶A is present in other RNA species^[6,7,35–37] and is installed in rRNAs by enzymes other than METTL3 (*i.e.*, METTL5/TRMT112 for 18S and ZCCHC4 for 28S ribosomal subunits, respectively).^[35,36] The average half-life of rRNAs and tRNAs, presenting the major RNA component in cells, is much longer than that of mRNA.^[31,38] Therefore, prolonged incubation times are necessary to evaluate potential non-specific inhibition of the methyltransferases modifying

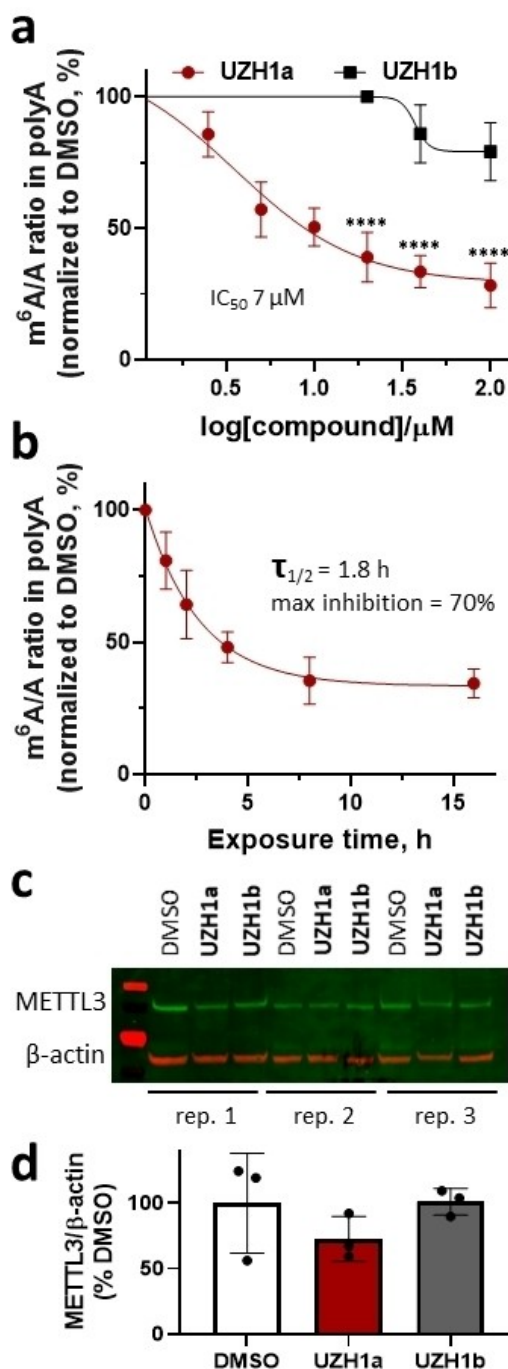


Figure 3. Cellular activity of the METTL3 inhibitor UZH1a in MOLM-13 cells. (a) Results of UPLC-MS/MS assay of m⁶A level in mRNA from MOLM-13 cells upon UZH1a and UZH1b treatment at a dose range of 2.5 to 100 μ M for 16 h, mean \pm SD, N = 3–7 (biological replicates), *****p* < 0.0001 vs. UZH1b. (b) Kinetics of m⁶A level reduction in mRNA from MOLM-13 cells upon UZH1a treatment at a dose of 40 μ M for 1 to 16 h determined in UPLC-MS/MS assay, mean \pm SD, N = 3 (biological replicates), normalized to DMSO-treated samples for each time point. (c) Western blotting of METTL3 expression level in MOLM-13 cells after treatment with 40 μ M of UZH1a and UZH1b for 16 h, N = 3 (biological replicates). (d) Quantification of the METTL3 signal intensity relative to β -actin in western blot images of compound treated versus DMSO only treated samples, mean \pm SD, N = 3 (biological replicates).

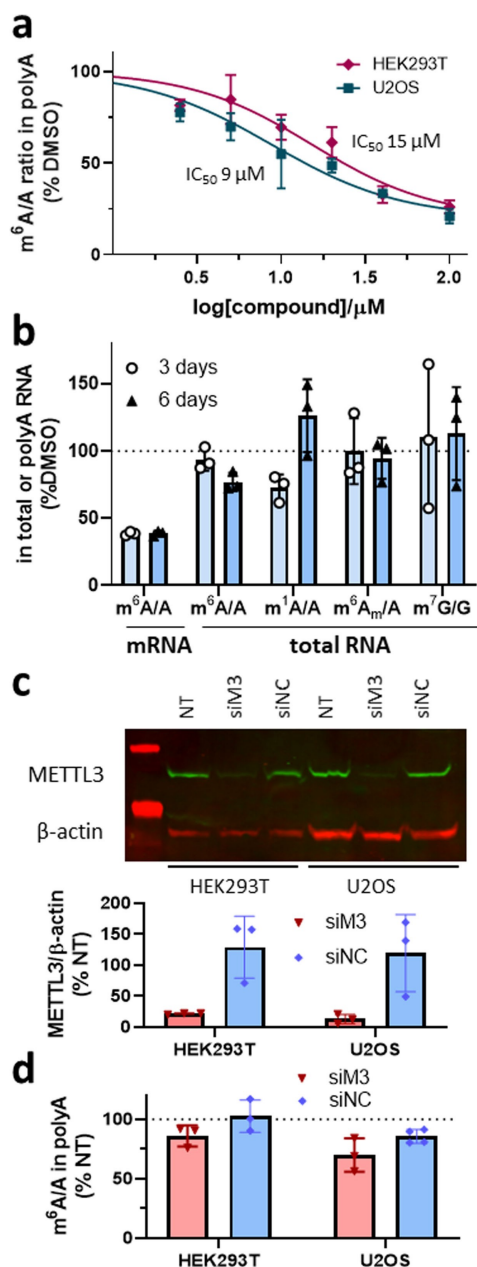


Figure 4. Cellular activity of the METTL3 inhibitor **UZH1a** and siRNA-mediated METTL3 knockdown in HEK293T and U2OS cells. (a) Results of UPLC-MS/MS assay of m⁶A level in mRNA from U2OS and HEK293T cells upon **UZH1a** treatment at a dose range of 2.5 to 100 μM for 16 h, mean ± SD, N = 3 (biological replicates), normalized to DMSO-treated samples for each cell line. (b) Results of UPLC-MS/MS assay of modified nucleosides level in mRNA fraction and in total RNA from U2OS cells upon 40 μM **UZH1a** treatment for 3 and 6 days, mean ± SD, N = 3 (biological replicates), normalized to DMSO-treated samples. (c) Representative results of western blotting analysis of METTL3 protein level in HEK293T and U2OS cells after 72-h transfection with 50 nM of siRNA targeting METTL3 (siM3), negative control siRNA (siNC), or in non-transfected cells (NT). Quantification of the METTL3 signal intensity relative to β-actin in western blot images of siRNA-treated versus non-transfected (NT) samples, mean ± SD, N = 3 (biological replicates). (d) Results of UPLC-MS/MS assay of m⁶A level in mRNA from HEK293T and U2OS cells after 72-h transfection with 50 nM of siRNA targeting METTL3 (siM3) or negative control siRNA (siNC), normalized to non-transfected cells (NT), mean ± SD, N = 3–4 (biological replicates).

these RNA types. We studied the levels of m⁶A, m⁶A_m, m¹A, m⁷G in total RNA of U2OS cells treated with 40 μM of **UZH1a** for 3 and 6 days and observed virtually no changes in these modifications (Figure 4b). In addition to the LC-MS/MS analysis of m⁷G methylation levels in total RNA after **UZH1a** treatment (Figure 4b), we used mechanistic protein thermal shift assay to confirm the absence of binding of our compound **UZH1a** to METTL1 (Figure 2a), an enzyme responsible for m⁷G installation on tRNA,^[39,40] mRNA,^[41] and miRNA.^[42] Overall, these results point out the exceptional selectivity of **UZH1a** towards other RNA methyltransferases.

Conclusion

Chemical probes that target proteins involved in the m⁶A modification of RNA are essential for understanding the complexity of this epitranscriptomic regulatory network. Furthermore, they are also valuable starting points for designing novel drugs against several diseases ranging from cancer to viral infections. Here, we have characterized a small-molecule inhibitor of METTL3 by protein crystallography, a biochemical binding assay, and a battery of cellular assays. Our METTL3 inhibitor **UZH1a** shows high-nanomolar potency in the biochemical assay, good selectivity against a panel of protein methyltransferases and kinases, and is active in cells. Importantly, METTL3 chemical inhibition reduced the m⁶A/A ratio in mRNAs of three different cell lines (AML MOLM-13 cells, osteosarcoma U2OS cells, and the embryonic kidney cell line HEK293T). In addition, we confirmed the selectivity of our compound towards other RNA methyltransferases in a protein thermal shift assay as well as in living cells. The crystal structure of METTL3/14 in complex with **UZH1a** has revealed its binding mode and is used as the basis for the development of more potent inhibitors. Future work will focus on the implications of METTL3 inhibition in various disease models.

Experimental Section

Compound characterization

UZH1a

Purity >95%, confirmed by supercritical fluid chromatography (SFC).

¹H NMR (500 MHz, CD₃OD) δ 7.95 (s, 1H), 7.82 (d, *J* = 8.0 Hz, 1H), 7.31–7.24 (m, 4H), 7.23–7.18 (m, 1H), 6.90 (s, 1H), 6.86 (d, *J* = 8.0 Hz, 1H), 5.58 (s, 1H), 4.38 (s, 2H), 3.64–3.37 (m, 8H), 2.52 (s, 4H), 1.84–1.74 (m, 2H), 1.72–1.59 (m, 2H), 1.43 (t, *J* = 5.6 Hz, 4H), 0.94 (s, 6H).

¹³C NMR (126 MHz, CD₃OD) δ 170.7, 164.4, 163.8, 161.8, 158.2, 144.0, 140.4, 129.9, 129.6, 128.3, 128.1, 120.8, 120.0, 117.2, 82.6, 71.8, 63.5, 52.8, 50.9, 47.1, 45.9, 45.5, 39.0, 35.3, 29.1, 28.4, 22.6. Five carbon signals are missing due to overlapping of equivalent signals.

HRMS (ESI): *m/z*: calculated for C₃₂H₄₃N₆O₃⁺ [*M* + H⁺]: 559.33912 found: 559.33851.

UZH1b

Purity > 95%, confirmed by SFC.

$^1\text{H NMR}$ (500 MHz, CD_3OD) δ 7.95 (s, 1H), 7.83 (d, $J=8.0$ Hz, 1H), 7.31–7.24 (m, 4H), 7.23–7.18 (m, 1H), 6.91 (s, 1H), 6.88 (d, $J=8.0$ Hz, 1H), 5.58 (s, 1H), 4.38 (s, 2H), 3.64–3.37 (m, 8H), 2.54 (s, 3H), 1.84–1.74 (m, 2H), 1.72–1.59 (m, 2H), 1.44 (t, $J=5.6$ Hz, 4H), 0.94 (s, 6H).

$^{13}\text{C NMR}$ (126 MHz, CD_3OD) δ 170.7, 164.4, 163.7, 161.7, 158.2, 143.7, 140.4, 129.9, 129.6, 128.3, 128.1, 121.0, 120.0, 117.2, 82.5, 71.8, 63.4, 52.8, 50.9, 47.1, 45.9, 45.5, 38.9, 35.3, 29.1, 28.5, 22.6. Five carbon signals are missing due to overlapping of equivalent signals.

HRMS (ESI): m/z : calculated for $\text{C}_{32}\text{H}_{43}\text{N}_6\text{O}_3^+$ [$\text{M} + \text{H}^+$]: 559.33912 found: 559.33848.

IR (cm^{-1} , neat): 418, 426, 438, 452, 471, 488, 498, 513, 547, 555, 592, 608, 668, 697, 733, 748, 778, 798, 826, 873, 977, 1027, 1114, 1150, 1171, 1226, 1260, 1306, 1329, 1362, 1431, 1453, 1496, 1505, 1548, 1595, 2922, 2948, 3323.

Protein expression and purification

Recombinant METTL3₃₅₄₋₅₈₀-METTL14₁₀₆₋₃₉₆ complex constructs for crystallization and for the use in enzymatic activity and protein thermal shift assays were expressed using the baculovirus/Sf9 insect cell expression system as previously described.^[25]

The plasmid expressing the N-terminally hexahistidine-tagged METTL1 protein was a gift from Cheryl Arrowsmith (Addgene ID: 25264). The protein was overexpressed for 12 hours at 20 °C in *Escherichia coli* BL21 (DE3) cells upon induction with 0.2 mM IPTG. The cells were harvested and resuspended in the lysis buffer containing 100 mM Tris-HCl at pH 8.0 and 500 mM NaCl. The cells were lysed by sonication and the cell lysate was clarified by centrifugation at 50'000 g for two hours and loaded onto Ni-NTA affinity column (5 mL HisTrap FF from GE Healthcare). After extensive washing with the wash buffer containing 100 mM Tris-HCl at pH 8.0, 500 mM NaCl and 50 mM imidazole the target protein was eluted with elution buffer containing 100 mM Tris-HCl at pH 8.0, 500 mM NaCl and 250 mM imidazole. The N-terminal hexahistidine-tag was removed by cleavage with tobacco etch virus (TEV) protease at 1:50 ratio. The excess imidazole was removed by overnight dialysis and the sample was subjected to secondary subtractive Ni-NTA affinity chromatography step to remove the protease and uncleaved protein. Finally, the protein was subjected to a gel filtration step using Superdex 75 16/60 column in a buffer containing 10 mM Tris-HCl at pH 8, 150 mM NaCl.

Differential scanning fluorimetry (thermal shift assay)

Thermal shift assay was performed as previously described with slight modifications.^[43] Briefly, protein (METTL1 or METTL3/14) at a final concentration of 2 μM was mixed with **UZH1a** or **UZH1b** at final concentrations of 10, 50, 100 μM in a final volume of 20 μL in a buffer system consisting of 50 mM HEPES at pH 7.5 and 150 mM NaCl. DMSO concentration was kept at 1% (v/v). SYPRO Orange was added at a final dilution of 1:1000 (v/v) as a fluorescence probe (ex/em 465/590 nm). Differential scanning fluorimetry was performed on a LightCycler[®] 480 Instrument II (Roche Diagnostics, Indianapolis, IN). The temperature was raised with a step of 3.6 °C per minute from 20 °C to 85 °C and fluorescence readings were taken at each interval. The reported values (ΔT_m) are calculated as the difference between the transition midpoints of an individual sample and the average of the reference wells (containing the protein and the DMSO only) in the same plate.

Reader-based HTRF assay of METTL3 inhibition in vitro

Compound potencies were evaluated by using a previously reported METTL3 inhibition assay.^[30] Briefly, the level of $m^6\text{A}$ in the oligoribonucleotide substrate after the reaction catalyzed by METTL3-METTL14 was quantified by measuring specific binding of modified oligoribonucleotide to the $m^6\text{A}$ reader YTHDC1₃₄₅₋₅₀₉ by homogeneous time-resolved fluorescence (HTRF). Tested compounds that inhibit METTL3 decrease the $m^6\text{A}$ level and, thus, reduce the HTRF signal. Response curves were plotted in GraphPad Prism 8.4 and fitted with nonlinear regression "log(inhibitor) vs. normalized response-variable slope", from which IC_{50} values were determined. The IC_{50} values are given as an average of at least three independent measurements for each compound.

Protein crystallization

The protein crystals of METTL3₃₅₄₋₅₈₀-METTL14₁₀₆₋₃₉₆ were obtained as previously described.^[25] Briefly, purified METTL3/14 complex was diluted to 5 mg/mL in 10 mM Tris-Cl pH 8.0, 200 mM NaCl. Crystals were obtained using the hanging drop vapor diffusion by mixing 1 mL complex solution with 1 mL reservoir solution containing 20% PEG3350 and 400 mM of magnesium acetate. The soaking experiment was carried out by transferring crystals to a 1 μL drop containing 100 mM compound directly dissolved in the buffer containing 30% PEG3350, and 400 mM magnesium acetate. After 16-h incubation at 22 °C, the crystals were harvested and flash-frozen in liquid nitrogen.

Data collection and structure solution

Diffraction data were collected at the PXIII beamline at the Swiss Light Source (SLS) of the Paul Scherrer Institute (PSI, Villigen, Switzerland) and processed using XDS as previously described.^[29] The crystal structures were solved by molecular replacement by employing the 5L6D structure as the search model in the Phaser program^[44] (Phenix package). In the crystals not subjected to soaking, clear electron density for product cofactor *S*-adenosyl-L-homocysteine (SAH) is visible. Therefore, in this soaking experiment setup test compounds competed with SAH for the *S*-adenosyl-L-methionine (SAM) binding site. In the crystal structure of **UZH1a**, the electron density due to the homocysteine part of SAH was no longer visible. All of the crystallographic models were constructed through iterative cycles of manual model building with COOT^[45] and refinement with phenix.refine.^[46] Default XYZ (reciprocal-space), XYZ (real-space), individual B-factors and occupancies refinement parameters appropriate for the resolution range were utilized. During the first run of the refinement update water was used in phenix.refine followed by addition of the missing water molecules manually.

Selectivity profiling

For kinase inhibition testing, Merck-Millipore KinaseProfiler[™] assay (radiometric, activity testing) was performed by Eurofins (Luxembourg). **UZH1a** was tested in single dose mode, in duplicate, at 10 μM , concentration of ATP was 10 μM . The required volume of the 50 \times stock of **UZH1a** in DMSO was added to the assay well, before a reaction mix containing an enzyme and substrate was added. The reaction was initiated by the addition of ATP. There was no pre-incubation of the compound with the enzyme/substrate mix prior to ATP addition. For further details of each individual assay, please refer to the company's website.

For protein methyltransferase inhibition testing, HotSpot Methyltransferase Profiling assay was performed by Reaction Biology

Corporation (Malvern, PA). **UZH1a** was tested in single dose mode, in duplicate, at 10 μM . Control compounds, SAH (S-(5'-adenosyl)-L-homocysteine) or chaetocin, were tested in 10-dose IC_{50} mode with 3-fold serial dilution starting at 100 or 200 μM . Reactions were carried out at 1 μM SAM.

Caco-2 cell permeability assay

The **UZH1a** intestinal permeability has been evaluated in Caco-2 cell permeability assay by ChemPartner (Shanghai, China) following standard procedures. Briefly, Caco-2 monolayer cells were cultured in Millicell-24[®] cell culture plates (Merk Millipore; surface area of the membrane = 0.7 cm^2 , $V_A = 0.8$ mL (A-to-B) or 0.4 mL (B-to-A)). The cell permeability of **UZH1a** along with the reference compounds (*i.e.*, erythromycin, metoprolol, and atenolol) was tested at 10 μM in HBSS buffer containing 0.4% DMSO (*v/v*) final concentration. The compounds were placed either in apical or basolateral chamber, and their concentrations were evaluated in both compartments after the 90-min incubation at 37 $^{\circ}\text{C}$ using the API 4000[™] LC/MS/MS system (Applied Biosystems, Waltham, MA). The Caco-2 monolayer's leakiness after the treatment was evaluated by measuring concentrations of a fluorescent dye Lucifer Yellow with low cell permeability (5 μM initial concentration) using a fluorometer (at Ex/Em of 485/535 nm). Lucifer Yellow P_{app} values were lower than $1 \cdot 10^{-6}$ cm/s , and transepithelial electrical resistance (TEER) values were higher than 400 $\Omega \cdot \text{cm}^2$, verifying that the Caco-2 monolayers were intact. Mass recovery of **UZH1a** was 82.4% (A-to-B) and 92.4% (B-to-A), indicating low non-specific adsorption to the assay chambers. Compound permeability was evaluated in duplicates.

Compound permeability was calculated according to the following equation, where V_A is the volume in the acceptor well, area is the surface area of the membrane and time is the total transport time in seconds:

$$P_{\text{app}} = \frac{V_A \times [\text{Compound}]_{\text{acceptor}} \times \text{Dilution Factor}}{\text{Area} \times \text{Time} \times [\text{Compound}]_{\text{donor, initial}}}$$

Lucifer Yellow permeability was calculated according to the following equation:

$$P_{\text{app}} = \frac{V_A \times ([\text{RFU}]_{\text{acceptor}} - [\text{RFU}]_{\text{blank}}) \times \text{Dilution Factor}}{\text{Area} \times \text{Time} \times ([\text{RFU}]_{\text{donor, initial}} - [\text{RFU}]_{\text{blank}})}$$

Compound recovery was calculated according to the following equation:

$$\text{Recovery (\%)} = \frac{\text{Total compound}_{\text{donor, 90 min}} + \text{Total compound}_{\text{acceptor, 90 min}}}{\text{Total compound}_{\text{donor, initial}}} \times 100\%$$

Transepithelial electrical resistance (TEER) was calculated according to the following equation:

$$\text{TEER } (\Omega \cdot \text{cm}^2) = (\text{Resistance}_{\text{sample}} - \text{Resistance}_{\text{blank}}) \times \text{Area}$$

Cell culture

U2OS and HEK293T cells were maintained in DMEM medium supplemented with 4.5 g/L glucose, GlutaMAX[™], 10% Gibco[™] FBS, and 1% penicillin/streptomycin (complete medium) in 5% CO_2 at

37 $^{\circ}\text{C}$ in a humidified incubator, with regular passaging twice a week using 1:5 split ratio. MOLM-13 cells were cultured in RPMI 1640 medium containing 10% Gibco[™] FBS and 1% penicillin/streptomycin (complete medium) in 5% CO_2 at 37 $^{\circ}\text{C}$ in a humidified incubator, with maintained cell densities at $0.6\text{--}2 \cdot 10^6$ cells/mL. All cell lines have been authenticated by cell line typing (external service provided by Microsynth, Switzerland). All cell lines were tested negative for mycoplasma contamination (PCR-based assay by Microsynth, Switzerland).

Cellular target engagement assay InCELL Pulse

Gene encoding METTL3 catalytic domain (residue 356–580) was amplified using primers listed below:

pICP-N-M3-F: ATATGAATTCACACAGAGTGTCCGAGGTGATTCC

pICP-N-M3-R: ATGCTCTAGATTATAAATCTTAGGTTTAGAGATGATAC-CATCTGGG

The amplified gene was ligated into the pICP-ePL-N vector between EcoRI and XbaI restriction sites

Cellular target engagement of the compounds was evaluated using InCELL Pulse assay according to the manufacturer's instructions. Briefly, HEK293T cells were seeded at $1 \cdot 10^6$ cells/well in 2 mL complete medium in a 6-well plate and transfected with 1.5 μg of the plasmid encoding enhanced ProLabel[®] (ePL) enzyme fragment (42 aa) fused to the N-terminus of the METTL3 catalytic domain (residues 354–580) using Lipofectamine RNAiMAX for 24 h. After the transfection, cells were washed once with Opti-MEM medium, detached from the plate by gentle washing and counted. Forty microliters of Opti-MEM medium containing 5000 transfected cells were mixed with 10 μL of 5 \times compound (**UZH1a** or **UZH1b**) solution in Opti-MEM or 2.5% (*v/v*) of DMSO (5 \times) as a negative control and incubated for 1 h at 37 $^{\circ}\text{C}$. After the incubation, cells were heated at 46 $^{\circ}\text{C}$ for 3 min followed by incubation at 22 $^{\circ}\text{C}$ for 3 min, and the non-aggregated METTL3-ePL protein was quantified using luminescence-based assay according to the InCELL Pulse kit's instructions.

Quantification of m⁶A/A ratio in polyadenylated RNA and total RNA by UPLC-MS/MS analysis

MOLM-13 cells were seeded into 6 well plates at a density of $1 \cdot 10^6$ cells/mL in 2 mL of complete RPMI medium. U2OS cells were seeded at $5 \cdot 10^5$ cells/well in a 6-well plate, whereas HEK293T cells were seeded at $1 \cdot 10^6$ cells/well. After 24 h, cells were treated with increasing concentrations of **UZH1a** or **UZH1b** in DMSO or DMSO alone as a negative control (final concentration of DMSO 0.5% (*v/v*)) for indicated time points. For the long-term incubation studies, U2OS cells were seeded at $1.5 \cdot 10^5$ cells/well in a 6-well plate. After 24 h, cells were treated with 40 μM of **UZH1a** in DMSO or DMSO alone as a negative control (final concentration of DMSO 0.5% (*v/v*)) for 3 days or, after reseeding at $1.5 \cdot 10^5$ cells/well and adding fresh compound, 6 days in total. Following incubation in cell culture incubator, cells were washed once with PBS, and total RNA was extracted using 0.5 mL of GENEzol[™] reagent according to manufacturer's instructions. The final volume of 50 μL of total RNA eluate was subjected to two rounds of purification using 25 μL Sera-Mag magnetic oligo(dT) particles per sample. The polyadenylated RNA was eluted with nuclease-free water in a final volume of 25 μL , and its concentration was determined using NanoDrop. One hundred nanograms of mRNA was digested to nucleosides and dephosphorylated in a one-pot reaction using 0.5 μL of nucleoside digestion mix in 25 μL of total reaction volume for 4 hours at 37 $^{\circ}\text{C}$.

The samples were used for UPLC-MS/MS analysis without further purification steps.

Nucleoside UPLC-MS/MS was performed at the Functional Genomics Centre Zurich following previously described procedures with slight modifications.^{[47][48]} Briefly, the samples were diluted 10 or 100-fold in solvent A (0.1% (v/v) formic acid in water), separated using a Waters Aquity M class (Waters) UPLC and detected with a TSQ Quantiva (Thermo Scientific) mass spectrometer by using a nano electrospray ionization (ESI) source in SRM mode. Analytes were separated on an HSS T3 column (particle size 1.8 μm, dimensions 0.15 × 60 mm). The mobile phases used for elution were A 0.1% (v/v) formic acid in water and B 0.1% (v/v) formic acid in acetonitrile at a flow rate of 2 μL/min, column temperature was ambient, sample temperature was 8 °C, injection volume was set to 1 μL, and sample loop volume was 1 μL. The analytes were separated following a gradient of 0–4 min 12.5% B, 4–4.1 min 99% B, 4.1–4.6 min 12.5% B, 6 min 12.5% B. Adenosine, m⁶A, m¹A, and m⁶A_m eluted at 3.30 min, 3.44 min, 2.89 min, and 3.56 min, respectively. Guanosine and m⁷G eluted at 3.34 min and 3.27 min, respectively. The mass spectrometer was operated using the following parameters: ESI positive ion mode, SRM acquisition mode, capillary voltage 3 kV, source temperature 300 °C, collision energy of 15 V for adenosine and guanosine, 20 V for m⁷G, and 25 V for m⁶A, m¹A, and m⁶A_m. Adenosine, m⁶A, m¹A, and m⁶A_m were detected by monitoring precursor to product transitions of m/z = 268.1 → 136.1, m/z = 282.1 → 150.1, m/z = 282.1 → 150.1, and m/z = 296.1 → 152.1, respectively. Guanosine and m⁷G were detected by monitoring precursor to product transitions of m/z = 284.1 → 136.1 and m/z = 298.1 → 166.1, respectively. Absolute concentrations of adenosine, guanosine, m⁶A, m¹A, m⁶A_m, and m⁷G in the samples were determined based on an external calibration curve generated using adenosine (Sigma-Aldrich, 01890-5G), guanosine (Sigma-Aldrich, G6264-1G), m⁶A (Chemie Brunschwig, CBLQB-1055-1 g), m¹A (MedChem Express, MCE-HY-113081), m⁶A_m (Toronto Research Chemicals, D447415), and m⁷G (Sigma-Aldrich, M0627-100MG) nucleoside standards. The modified-to-nonmodified nucleoside ratio of compound-treated samples were normalized to the corresponding value of DMSO-treated negative control. All measurements were performed in technical duplicates and repeated in at least three biological replicates on different days. Inhibition curves were plotted in GraphPad Prism 8.4 and fitted with nonlinear regression “log(inhibitor) vs. normalized response-variable slope” with the maximal inhibition fixed at 80%, from which IC₅₀ values were determined.

RNAi-mediated knockdown of METTL3

For western blotting experiments, HEK293T or U2OS cells were seeded in 6-well plates at a density of 1.5 · 10⁵ cells/well in 2 mL of complete DMEM medium, transfected with total 50 nM of METTL3-targeting siRNA mix (at equimolar ratio) or negative control siRNA using Lipofectamine[®] RNAiMAX according to manufacturer's instructions, and incubated at 37 °C with 5% CO₂. The sequences of siRNAs are provided in the materials section. After 72 h, cells were analyzed for METTL3 protein expression using western blotting as described in the western blotting section.

To evaluate the effect of METTL3 knockdown on m⁶A/A level in polyadenylated RNA, HEK293T or U2OS cells were seeded in 6-well plates and transfected as described above. After 72 h, m⁶A/A ratio reduction in polyadenylated RNA was evaluated as described in UPLC-MS/MS analysis section. The assay was repeated three times on different days.

Western blotting

For immunodetection of METTL3 expression level after the compound exposure, MOLM-13 cells were seeded at 1 · 10⁶ cells/mL in a 6-well plate in complete RPMI media. The next day, cells were treated with the indicated doses of compounds. After 16 hours of treatment, cells were collected by centrifugation, washed once with PBS, and resuspended in 200 μL of RIPA buffer (150 mM sodium chloride, 1% (v/v) IGAPAL CA630, 0.5% (w/v) sodium deoxycholate, 0.1% (v/v) sodium dodecyl sulphate, 50 mM Tris-HCl) with added protease inhibitor cocktail (1 × final concentration). After incubating for 30 minutes on ice, cell lysates were centrifuged for 15–20 minutes at 16000 × g at 4 °C to remove cell debris, and the supernatant containing proteins was collected. The protein concentration was quantified with Pierce[™] Coomassie (Bradford) protein assay kit, and 30 μg of protein was loaded per well on a SurePAGE 12% Bis-Tris polyacrylamide gel. Following electrophoresis, the proteins were transferred to methanol-activated low fluorescence PVDF membrane using Trans-Blot[®] Turbo[™] Transfer System (Bio-Rad, Hercules, CA) according to manufacturer's instructions. The membrane was then blocked with 5% non-fat milk and incubated with anti-METTL3 rabbit (1:1000) and anti-β-actin mouse (1:5000) antibodies overnight at 4 °C. The next day, the membrane was washed 3 times with PBST (PBS with 0.1% (v/v) Tween-20) and incubated with IRDye[®] 800CW goat anti-rabbit IgG and IRDye[®] 680RD donkey anti-mouse IgG secondary antibodies (1:10000) for 2 hours at room temperature. After incubation, the membrane was again washed three times in PBST, and fluorescence signal was detected on Odyssey[®] CLx Imaging System (LI-COR). The band intensity in each lane was quantified using Image Studio Lite Version 5.2.5 (LI-COR). For the detection of METTL3 protein in U2OS and HEK293T cells following the siRNA-mediated knockdown, the cells were washed with PBS and lysed in 60 μL of lysis buffer. Cell lysates were probed for METTL3 expression as described above.

Statistical analysis

The statistical analysis was performed using GraphPad Prism software. Multiple experimental groups were compared pairwise using the one-way analysis of variance (ANOVA) followed by Tukey's post-hoc test unless stated otherwise, normality was assessed using Shapiro-Wilk test. The differences between treatment groups were considered statistically significant at *p*-values lower than 0.05.

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Conflict of Interest

The authors declare no conflict of interest.

Keywords: METTL3 · RNA methyltransferase inhibitor · epitranscriptomics · m6A · leukemia

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