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**S-Adenosyl-L-methionine (SAM) is crucial for methylation and tightly controlled in cells. We examined SAM-III riboswitch response to 17 SAM analogues and used a Spinach/SAM aptasensor to monitor their enzymatic formation *in situ*. Most SAM analogues were recognized, unless they featured an *ortho*-substituted benzyl ring, indicating potential regulatory effects via SAM riboswitches.**

Riboswitches are regulatory elements found primarily in the 5'-untranslated regions of RNA from biosynthetic genes.<sup>1</sup> Upon binding to their ligand, they undergo structural changes that affect gene expression levels. One of the most prominent natural riboswitches senses *S*-adenosyl-L-methionine (SAM)<sup>1,2</sup> and regulates gene expression in the sulfur and SAM metabolism, including methionine adenosyltransferase (MAT).<sup>3</sup> Currently eight distinct riboswitch classes have been reported for SAM, which can be divided into the SAM-I superfamily (SAM-I,<sup>4,5</sup> SAM-IV,<sup>6</sup> SAM-I/IV<sup>7</sup>), the SAM-II superfamily (SAM-II,<sup>8,9</sup> SAM-V<sup>10,11</sup>), the SAM-III<sup>12</sup> riboswitches, the SAM-VI<sup>13</sup> riboswitches and the SAM/SAH<sup>14</sup> riboswitches. All SAM riboswitches recognize their ligand by hydrogen bonding to the nucleobase moiety and strong electrostatic interactions with the sulfonium center (Table S1, ESI†). However, the shape of the binding cleft surrounding the methyl group differs considerably among the classes. Few SAM-like metabolites, like *S*-adenosyl-L-homocysteine (SAH), have been tested regarding binding to SAM riboswitches, however, no data on SAM analogues (also called AdoMet analogues) with larger alkyl or benzyl groups instead of methyl were published.<sup>15,16</sup>

Aptasensors consist of a metabolite-binding aptamer domain and a signaling domain. Fluorescent light-up aptamers (FLAPs) are particularly useful and popular signaling domains.<sup>17–20</sup> SAM-dependent aptasensors with various FLAPs are available

## SAM-III aptamer enables detection of enzymatic SAM analogue generation†

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and have been used to monitor SAM dynamics and SAM levels in cells.<sup>15,21</sup>

SAM is the primary methylation agent and second most abundant cofactor after ATP. SAM analogues, in which the methyl group has been replaced by other residues, allow site-specific alkylation or benzylation by promiscuous methyltransferases. In particular, the enzymatic formation from methionine (analogues) and ATP catalyzed by methionine adenosyltransferase (MAT) has been demonstrated also in cells.<sup>22</sup> Recent work on engineered MATs greatly expanded the scope of enzymatically available SAM analogues to now include a plethora of benzylic groups.<sup>23,24</sup> However, monitoring MAT activity for SAM analogue synthesis requires HPLC analysis. Alternative approaches, like detection by phosphomolybdate, suffer from inaccuracy.<sup>25</sup> Methods for fast and easily applicable analysis of SAM analogue detection, preferably for monitoring their *in situ* formation, are therefore highly required. Aptasensors could provide a convenient approach, but no data is available for the acceptance of SAM analogues.

In this study, we investigated the promiscuity of the SAM-III riboswitch for 17 SAM analogues and implemented the Spinach/SAM aptasensor for *in situ* detection of chemo-enzymatic synthesis of SAM analogues by an engineered MAT enzyme.

We first asked which SAM riboswitch domains may exhibit promiscuity for SAM analogues by examining their crystal structures (Fig. S1, ESI†). Among all SAM riboswitches, the SAM-III riboswitch, also known as the S<sub>MK</sub> box, has the largest binding cleft surrounding the methyl group of SAM (Fig. 1A).<sup>26</sup> The methyl group is situated adjacent to the P2 helix, while helices P1 and P4 form the binding pocket. These helices are positioned further from the methyl group, creating a binding cleft that is open toward the riboswitch surface. This unoccupied space suggests the potential to accommodate larger substituents at the sulfonium center.

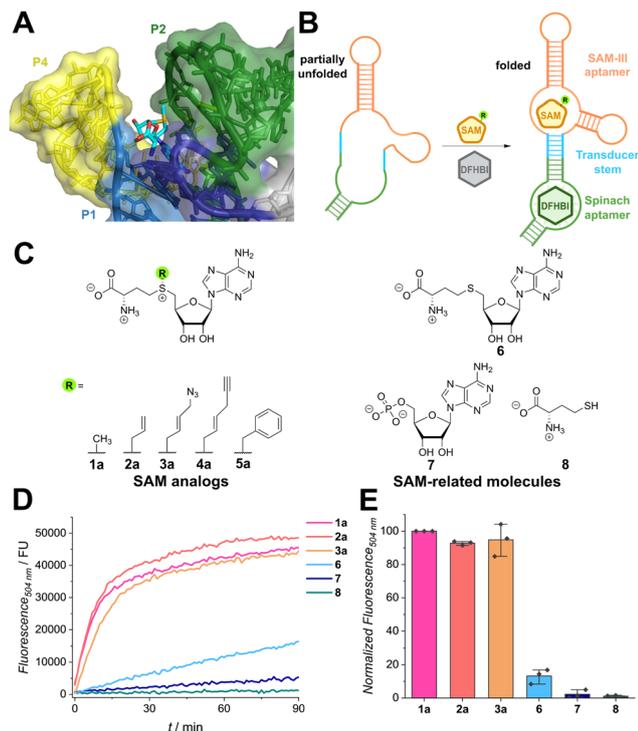
To investigate the promiscuity, we tested the Spinach/SAM aptasensor designed by Jaffrey *et al.* that is based on the S<sub>MK</sub> box from *Enterococcus faecalis*.<sup>15</sup> When SAM is bound, the transducer hybridizes, allowing the Spinach aptamer to adopt its active conformation and bind the fluorophore DFHBI

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**Fig. 1** Recognition of SAM analogues by the Spinach/SAM aptasensor. (A) Crystal structure of the SAM-III aptamer of *Enterococcus faecalis* with SAM bound (shown in sticks) (PDB: 3E5C). The methyl group at the sulfonium center (yellow) is in proximity to the P2 helix (green) and framed by the P1 (blue) and P4 helices (yellow). (B) Scheme illustrating the function of the aptasensor. Binding of SAM (analogue) induces hybridization of the transducer stem and folding of the Spinach aptamer domain. DFHBI ((Z)-5-(3,5-difluoro-4-hydroxybenzylidene)-2,3-dimethyl-3,5-dihydro-4H-imidazol-4-one) can then bind and the fluorescence signal increases. (C) Chemical structures of SAM analogues (**1–5**) and structurally related molecules used as controls (**6–8**). (D) Time course showing activation of fluorescence of the Spinach/SAM aptasensor by indicated SAM analogues (500 μM) and controls (500 μM) during incubation at 25 °C. Fluorescence was measured in 1 min intervals. The net fluorescence without ligand was subtracted from the raw data to exclude analogue-independent signal increase by fluorophore binding. Assay conditions: 120 μM DFHBI, 3 μM Spinach sensor, 100 flashes, manual z-position 21 709 μm, manual gain 200 and with  $\lambda_{\text{Ex}} = 460 \text{ nm}/\lambda_{\text{Em}} = 504 \text{ nm}$ . (E) Bar plot depicting intensity measurements recorded after 30 min. Fluorescence intensities were normalized to SAM and repeated in  $n = 3$  independent experiments.

(i.e., ((Z)-5-(3,5-difluoro-4-hydroxybenzylidene)-2,3-dimethyl-3,5-dihydro-4H-imidazol-4-one)). Binding leads to suppression of intramolecular rotation and thus an increase in fluorescence (Fig. 1B). We first measured binding affinities for SAM and a panel of chemically synthesized SAM analogues (**1a–5a**, Fig. S3, ESI†) by incubating the analogues for 30 min with the aptasensor. The obtained  $\text{EC}_{50}$ -value of  $2.26 \mu\text{M} \pm 0.10 \mu\text{M}$  is slightly higher than the literature value for the natural riboswitch ( $K_{\text{D}} = 0.57 \mu\text{M}$ ).<sup>16</sup> This difference may be attributed to alterations of the transducer and Spinach domains. All tested SAM analogues were accepted by the aptasensor and with growing size of the moiety, the binding affinity decreased ( $\text{EC}_{50} = \mathbf{1a}: 2.26 \mu\text{M} < \mathbf{2a}: 4.1 \mu\text{M} < \mathbf{4a}: 11.0 \mu\text{M} < \mathbf{3a}: 11.6 \mu\text{M} < \mathbf{5a}: 39.5 \mu\text{M}$ ). Interestingly, SAM analogue concentrations of approximately

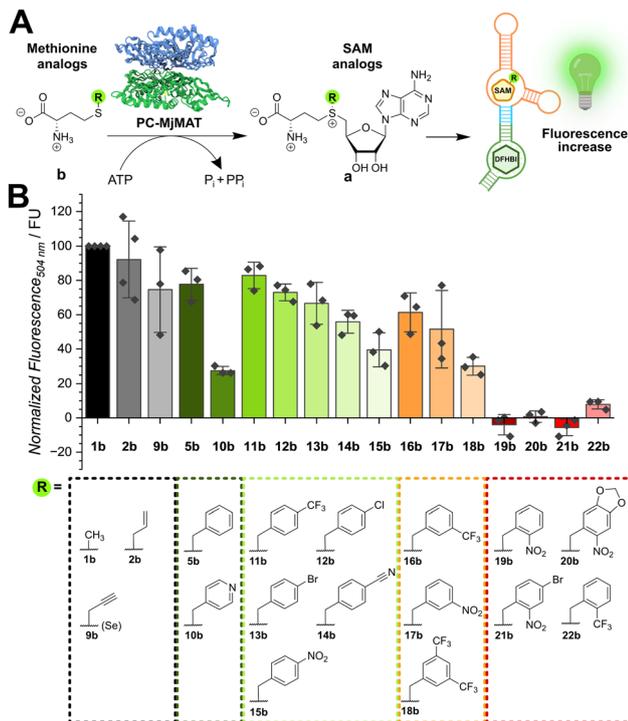
1 mM led to a signal decrease. Similar observations were reported by Chen *et al.* for the Spinach aptasensor, but not for the Pepper-based construct, which might hint on SAM-interference with the Spinach aptamer.<sup>27</sup>

To prove that SAM-related metabolites do not interfere with aptasensor signaling, several metabolites were tested (Fig. 1D and E). To this end, the metabolites and the aptasensor were directly mixed and time-dependent measurements were performed. While SAM analogues show a fast increase and reach a plateau after 15 min, SAH (**6**) and AMP (**7**) only show a slow increase in fluorescence, indicating weak binding to the RNA (Fig. 1D). The amino acid L-methionine (**8**) does not induce fluorescence. Strong preference for SAM in contrast to SAM-related molecules after 30 min of incubation (Fig. 1E) is in line with experiments reported by Paige *et al.*<sup>15</sup>

We then asked if the aptasensor can be used to monitor direct *in situ* generation of SAM analogues. PC-MjMAT is a promiscuous MAT variant from *Methanocaldococcus jannaschii* that cannot only react methionine and ATP, but also various methionine analogues (Fig. 2A).<sup>23</sup> PC-MjMAT has been shown to synthesize SAM analogues bearing small alkyl and a range of benzylic groups, including the photocleavable *ortho*-nitrobenzyl group.<sup>23,24</sup> PC-MjMAT and the aptasensor were combined in one-pot reactions and a panel of 17 methionine analogues was tested (Fig. 2B). Methionine and two small analogues containing the allyl or propargyl group (**2b**, **9b**) and the selenium (introduced for higher stability)<sup>28</sup> were well tolerated. When we tested the benzylic methionine analogues (**5b**, **10b–22b**), we observed that out of the 14 compounds tested, 10 were accepted by the aptasensor, leading to a fluorescent signal (Fig. 2B). We observed that the position of the substituent provides the major contribution to their acceptance by the riboswitch, following the order *para* > *meta* > *ortho*. One exception is the pyridinylmethyl **10b** with a lower aptasensor activation that might form an additional hydrogen bond with the aptasensor. While all *para*- and *meta*-modified analogues bound to the aptamer, *ortho*-modified analogues did not show significant binding to the aptasensor. This included *ortho*-nitrobenzyl groups of the photo-cleavable compounds **19b–21b**, which could be interesting for reversible blocking of methyltransferase target sites.<sup>29,30</sup> To exclude chemically specific effects or quenching by the nitro group, the *ortho*-trifluoromethyl-modified analogues **11** (*para*), **16** (*meta*) and **22** (*ortho*) were tested. Again, modification with an *ortho*-modified substituent inhibited binding to the RNA whereas *para*- and *meta*-modified analogues were accepted. The *ortho*-substituent is closer to the nucleic acid than *meta*- and *para*-substituents, suggesting steric hindrance as main reason preventing binding.

This study investigated for the first time how SAM riboswitches respond to analogues of the cofactor SAM. The results show that the  $\text{S}_{\text{MK}}$  box can accommodate a diverse panel of SAM analogues harboring larger alkyl or various benzyl groups. This is in line with the crystal structure showing a large solvent-exposed binding-cleft, which had led us to hypothesize that diverse SAM analogues might be accepted. Out of the 19 tested compounds, 15 were binding and led to a good signal in the





**Fig. 2** *In situ* detection of SAM analogue formation in enzymatic reactions. (A) Scheme for the chemo-enzymatic synthesis of SAM analogues by PC-MjMAT starting from methionine analogues. (B) Screening of SAM analogue acceptance by the SAM aptasensor by PC-MjMAT chemoenzymatic synthesis starting from methionine analogues. The bar plot shows normalized fluorescence intensities after 90 min of measurement for all SAM analogues tested. Graphs for the time-dependent fluorescence increase for all analogues are shown in Fig. S4 (ESI†). Conditions: 1 mM L-methionine analogue, 1 mM ATP, 100  $\mu$ M PC-MjMAT, 120  $\mu$ M DFHBI, 3  $\mu$ M aptasensor, MAT/ aptasensor buffer (50 mM HEPES, 100 mM KCl, 10 mM MgCl<sub>2</sub>, pH = 7.4), 25 °C. The control signal without analogue was subtracted to exclude SAM analogue independent signal increase by fluorophore binding. As **15b** is a fluorophore, the control signal here included the methionine analogue but no PC-MjMAT to correct for background fluorescence. The assay was performed  $n = 3$  times for each analog.

SAM/Spinach aptasensor. We conclude that steric rather than electronic effects provide the key contribution to recognition of SAM analogues with benzylic groups by the SAM riboswitch. This is supported by the observation that *ortho*-modified benzylic analogues are not accepted.

We implemented the aptasensor for *in situ* detection of SAM analogues generated in enzymatic reactions. Formation of SAM analogues led directly to a fluorescent signal that was detected in 384 well format. This assay can be a valuable tool to assess the promiscuity of SAM-synthesizing enzymes and facilitate their engineering, which currently requires HPLC analyses. As the fluorescence signal by the riboswitch can be reverted by depletion of SAM (analogue), the assay might also be used to monitor the activity of SAM (analogue) consuming enzymes. In future work, the assay may also be implemented in flow cytometry to allow for high-throughput screening. As SAM-aptasensors have been utilized to measure SAM and in cells, both in bacteria and eukaryotes,<sup>15,27</sup> our results suggest that imaging of SAM

analogues may also be possible, although careful controls accounting for natural SAM would have to be performed.

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## Data availability

The data supporting this article have been included as part of the ESI.†

## Conflicts of interest

There are no conflicts to declare.

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