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Designing fluorescent biosensors using circular permutations of riboswitches

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ABSTRACT

RNA-based fluorescent (RBF) biosensors have been applied to detect a variety of metabolites *in vitro* and in live cells. They are designed by combining the ligand sensing domain of natural riboswitches with *in vitro* selected fluorogenic aptamers. Different biosensor topologies have been developed to accommodate the diversity of riboswitch structures. Here we show that circular permutation of the riboswitch ligand sensing domain also gives functional biosensors, using the SAM-I riboswitch as our model. We reveal that this design can enhance fluorescence turn-on and ligand binding affinity compared to the non-permuted topology.

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1. Introduction

Riboswitches have been applied for classic synthetic biology applications including inducible gene regulation and metabolic engineering [1,2]. More recently, detailed structural information of riboswitch secondary and tertiary structure has enabled their combination with *in vitro* selected fluorogenic aptamers to generate RNA-based fluorescent (RBF) biosensors for metabolite imaging and detection [3]. RBF biosensors are designed so that binding of a target ligand induces the riboswitch conformational change that allows a dye to bind and turn on fluorescence. In this way, the biosensor generates a fluorescent signal only in the presence of a target metabolite. To date, RBF biosensors have been applied for high-throughput screening of enzyme activity *in vitro* and *in vivo*, imaging metabolites and signals in Gram positive and Gram negative bacteria, and anaerobic imaging of bacterial signals [3].

Different biosensor topologies have been explored to broaden the rational design strategy to accommodate the diversity of riboswitch and other aptamer folds. The first riboswitch-based biosensors utilized “transducer modules” consisting of randomized stem sequences to connect to the dye-binding aptamer called Spinach, which binds the profluorescent dye 3,5-difluoro-4-hydroxy benzylidene imidazolinone (DFHBI) [4]. One improvement our group made to biosensor design was to use the natural P1 stem from native riboswitches, which simplified the transducer module

design and permitted libraries of riboswitch aptamers to be screened for finding the optimal biosensor [5–7]. The conventional topology (RS-Sp, Fig. 1A) thus uses a P1-P2' stem architecture for the transducer module, where P1 refers to the first pairing stem of the ligand-sensing riboswitch and P2' stem refers to the second pairing stem of the dye-binding aptamer, which could be either Spinach or an improved fluorogenic aptamer variant, Spinach2 [8].

The conventional riboswitch-Spinach fusion design is effective for many riboswitch aptamers, but is not suitable when the terminal ends of the ligand-sensing domain do not form a P1 stem. For example, the *S*-adenosyl-L-homocysteine (SAH) riboswitch contains a terminal pseudoknot that places the 5' and 3' ends far apart. To address this challenge, our group constructed a biosensor by making a circular permutation of the Spinach2 aptamer and fusing it to the P2 stem of the SAH riboswitch (RS-cpSp, Fig. 1B) [9]. An alternate strategy is to incorporate Spinach to mimic the expression platforms found in native riboswitches. The ligand-free riboswitch aptamer interferes with the Spinach aptamer folding to block DFBHI binding, but target ligand binding restores the dye-binding aptamer (Fig. 1C) [10].

Here we showcase another strategy to access new biosensor designs that uses circular permutations of a riboswitch aptamer (cpRS-Sp, Fig. 1D). To our knowledge, there are no natural examples of circularly permuted riboswitches. However, circularly-permuted hammerhead ribozyme variants are found in diverse genomes [11] and some pre-tRNAs form circularized intermediates before maturation [12]. To investigate these new biosensor topologies, the SAM-I riboswitch was chosen as a model 4-way junction RNA fold to explore novel permuted architectures. This riboswitch

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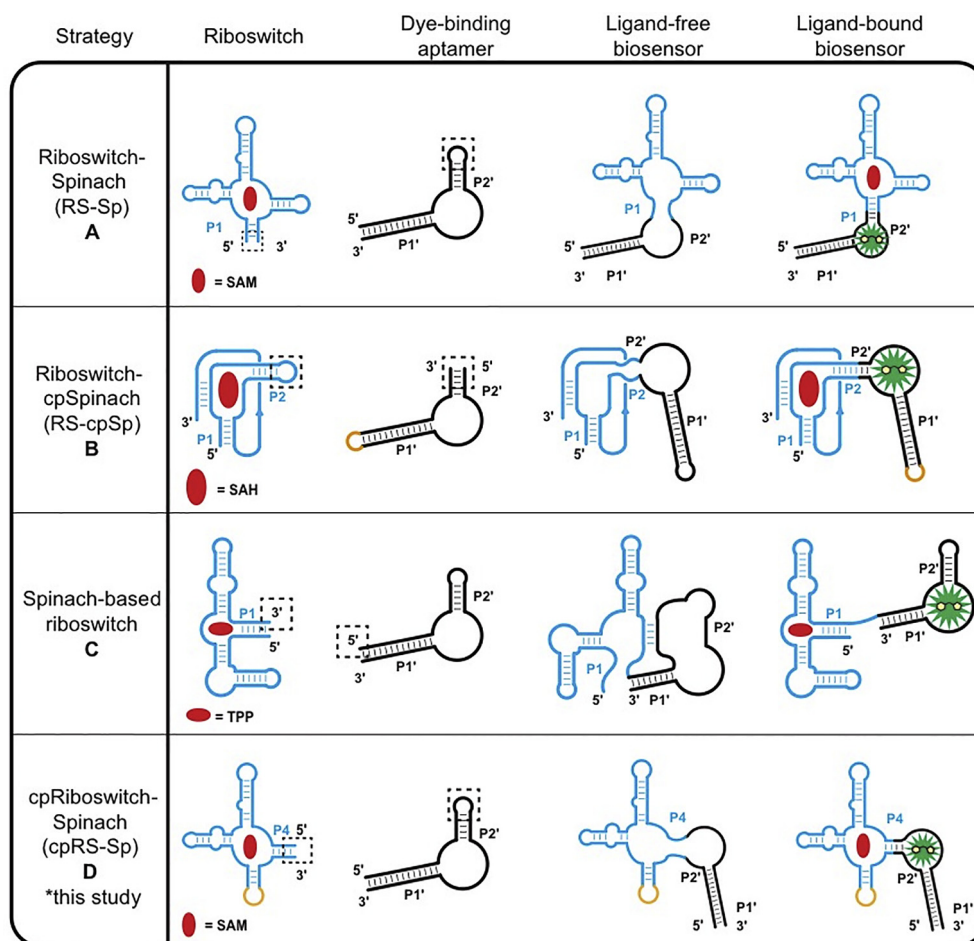


Fig. 1. Design strategies for riboswitch-based fluorescent biosensors. (A) The riboswitch–Spinach (RS-Sp) fusion strategy for biosensor development involves grafting the P1 stem of a riboswitch sensor domain to the P2' stem of Spinach [4–7]. (B) For riboswitches with non-pairing 5' and 3' ends, an internal stem of the riboswitch can be fused to a circular permutation of Spinach2 to generate a riboswitch–circularly permuted Spinach2 design (RS-cpSp) [9]. (C) Spinach riboswitches can be created by replacing the gene regulatory expression platform of a riboswitch with the Spinach aptamer [10]. (D) A new biosensor strategy for any riboswitch with multiple pairing stems is to circularly permute the riboswitch and fuse it to the P2' stem of Spinach2 to generate a circularly-permuted riboswitch–Spinach2 (cpRS-Sp) design. Riboswitch domains (blue) and their corresponding metabolites (red) are depicted with either the Spinach/Spinach2 aptamer (black) and 3,5-difluoro-4-hydroxybenzylidene imidazolinone (DFHBI, green). “Sp” represents either Spinach or Spinach 2, which are used in different studies. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

class has been extensively characterized with structural, biochemical, and biophysical studies [13–15], which suggested that a circularly-permuted SAM-I riboswitch could be fused with a fluorogenic aptamer via different pairing stems. These constructs were generated, screened, and characterized as new biosensors utilizing circularly-permuted riboswitch topologies, then were compared to traditional biosensor architectures.

2. Materials & methods

2.1. Reagents and oligonucleotides

DNA oligonucleotides for biosensor constructs (Table S1) were purchased as Ultramers from Integrated DNA Technologies (Coralville, IA) and other DNA oligonucleotides were purchased from Elim Biopharmaceuticals (Hayward, CA). S-Adenosyl-L-methionine (SAM), S-adenosyl-L-homocysteine (SAH) were purchased from Sigma-Aldrich (St Louis, MO). DFHBI was synthesized following previously described protocols [16] and was stored as a 10 mM stock in DMSO at –20 °C. Chemically competent BL21

(DE3) Star cells were purchased from Life Technologies (Carlsbad, CA).

2.2. In vitro transcription procedure

DNA templates for *in vitro* transcription were prepared by PCR amplification using Phusion DNA polymerase (NEB) from sequence-confirmed plasmids or Ultramer oligonucleotides (for screening experiment only) using primers that added the T7 polymerase promoter sequence at the 5' end. PCR products were purified either by a 96-well format ZR-96 DNA Clean-up kit (Zymo Research) for screening experiments or by QIAquick PCR purification kit (Qiagen) for characterization and application experiments. RNA was transcribed from DNA templates using T7 RNA polymerase in 40 mM Tris-HCl, pH 8.0, 6 mM MgCl₂, 2 mM spermidine, and 10 mM DTT. RNAs were either purified by a 96-well format ZR-96 Clean & Concentrator (Zymo Research) or by denaturing (7.5 M urea) 6% PAGE. RNAs purified by PAGE were visualized by UV shadowing and extracted from gel pieces using Crush Soak buffer (10 mM Tris-HCl, pH 7.5, 200 mM NaCl and 1 mM EDTA, pH 8.0). Purified RNAs were precipitated with ethanol, dried, and then resuspended in TE buffer (10 mM Tris-HCl, pH 8.0, 1 mM EDTA).

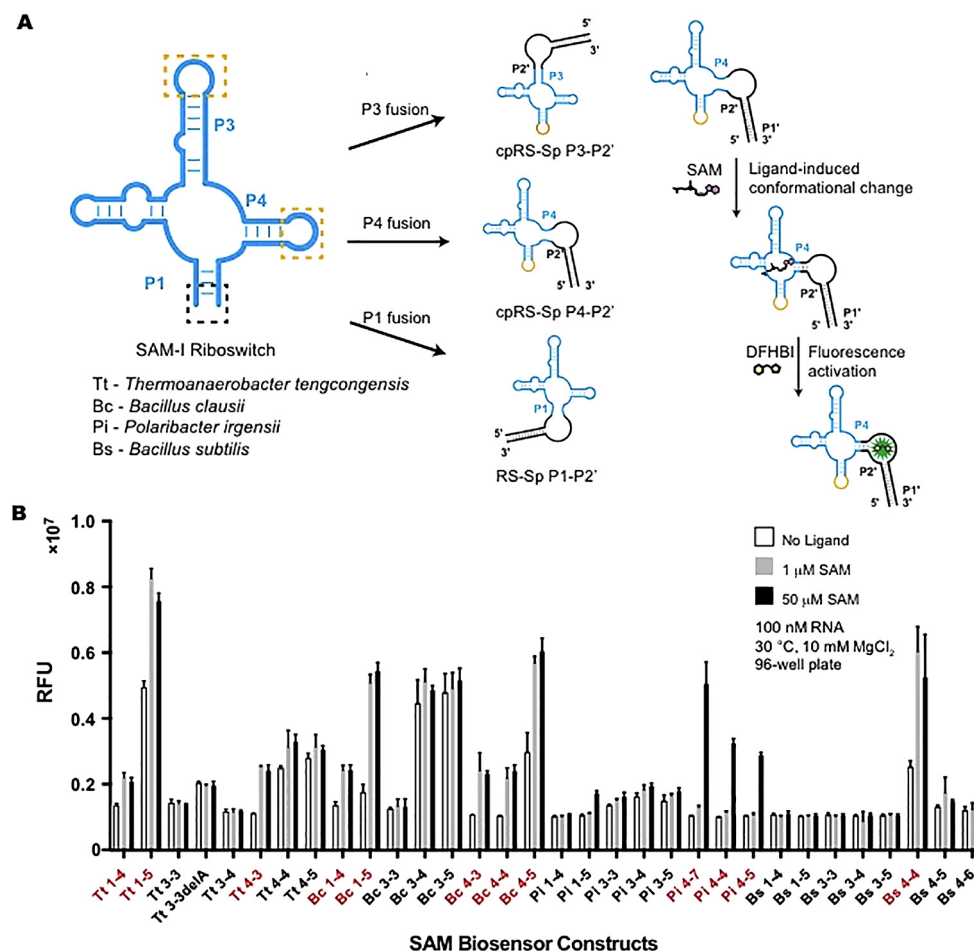


Fig. 2. *In vitro* screen of cPRS-Sp biosensors derived from the SAM-I riboswitch class. (A) Design of different SAM biosensor architectures and mechanism for fluorescence turn-on is depicted for the P4-P2' biosensor. (B) *In vitro* fluorescence response of SAM-I biosensors to SAM ligand. The nomenclature is based on the species origin of the sequence, the pairing stem of the SAM-I riboswitch that is fused to Spinach, and number of base pairs retained from the riboswitch stem, e.g. Bs 4-4 stands for SAM-I riboswitch from *Bacillus subtilis*, with Spinach fused to the P4 stem of the riboswitch that retains 4 base pairs. SAM biosensors chosen for further characterization (with greater than 1.5 $\times$ fluorescence increase at 50 μ M SAM) are indicated in red. Data shown are mean values $\pm$ standard deviation taken from two independent replicates. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

Accurate RNA concentrations were determined by measuring the absorbance at 260 nm after performing a hydrolysis assay to eliminate the hypochromic effect due to RNA secondary structure [17].

2.3. General procedure for *in vitro* fluorescence assays

In vitro fluorescence assays were carried out in binding buffer containing 100 nM RNA, 10 μ M DFHBI, 40 mM HEPES, pH 7.5, 125 mM KCl, and 3 or 10 mM $MgCl_2$, as indicated in the figures. Other conditions, including temperature and concentration of ligand, were varied in different experiments as indicated. The RNA was renatured by heating to 72 °C for 3 min in the binding buffer then cooled to ambient temperature for 5 min prior to addition to the reaction solution. DFHBI was added to the solution containing buffer and RNA, and then ligand (or water for no ligand control) was added before fluorescence measurement. Binding reactions were performed in 100 μ L volumes and were incubated at the indicated temperature in a Corning Costar 3915 96-well black plate or a Greiner Bio-One 384-well black plate in a Molecular Devices SpectraMax Paradigm Multi-Mode detection platform plate reader (Sunnyvale, CA). The fluorescence emission was measured during 30–60 min total with the following instrument parameters: 448 nm excitation, 506 nm emission.

2.4. Fluorescence polarization (FP) binding assays

Fluorescence polarization readings were carried out using a QuantaMaster spectrofluorometer (Photon Technology International) at excitation 646 nm, emission 662 nm. Samples were prepared in 50 mL of TBM buffer (90 mM Tris base, 89 mM boric acid, and 10 mM $MgCl_2$, pH 7.0) containing 1 mM of a Cy5-labeled SAM analog (C₈-Cy5) [18] and saturation binding experiments were performed with RNA concentrations ranging from 0 to 40 μ M. RNA was added successively to the sample cuvette, and concentration values were corrected for added volume. Samples were equilibrated at 30 °C for 2 min prior to each FP measurement using tubing connecting the cuvette holder to a water bath.

2.5. Binding affinity analysis of SAM biosensors

To measure the binding affinities of SAM biosensors, fluorescence assays were performed with the following conditions: binding buffer with 10 mM $MgCl_2$, 37 °C, 100 nM RNA, 10 μ M DFHBI, and ligand SAM concentrations from 10 nM to 100 μ M. The fluorescence of the sample with DFHBI but no RNA was subtracted as background to determine relative fluorescence units.

2.6. In vivo fluorescence assays by flow cytometry

Preparation of cell samples for flow cytometry was carried out by inoculating 3 mL of LB/carb media with 150 μ L of an overnight culture of BL21 (DE3) Star cells containing either the pET31b-RS-Sp or pET31b-cpRS-Sp constructs (Table S2). Cells were grown aerobically to an $OD_{600} \sim 0.5$ – 0.6 , then induced with 1 mM IPTG at 37 °C for 2 h. Cell density was measured by OD_{600} , and assuming that there are 1×10^9 cells/mL for an OD_{600} of 1, 4×10^8 cells were sampled and pelleted at rt for 4 min at 3700 rcf, washed once with PBS media at pH 7.0, then resuspended in PBS media containing 100 μ M DFHBI. Cellular fluorescence was measured for 30,000 cells using an Attune NxT Acoustic Focusing Cytometer (Life Technologies).

3. Results

3.1 Screen of different biosensor topologies

The SAM-I riboswitch has a 4-way junction architecture. Biochemical and X-ray crystallographic information have shown that

the pairing stem element P2 is involved in formation of a pseudoknot that is important for SAM binding [14]. Thus, modifications at the P2 stem were excluded from our biosensor design, but the other pairing stems P1, P3, and P4 became candidates for the transducer stem.

Biosensors that fuse Spinach2 to the riboswitch P1 stem are the conventional RS-Sp design. By contrast, biosensors that fuse Spinach2 to the riboswitch P3 or P4 stems would represent two new cpRS-Sp designs, which require the P1 stem to be closed by a loop sequence (GCAA, Fig. 2A). In all cases, the riboswitch or circularly permuted riboswitch was fused to the P2' stem of Spinach2, thus the nomenclature for the different topologies is P1-P2', P3-P2' or P4-P2'. A small library of 32 biosensor candidates were designed based on these criteria: 1) Four SAM-I riboswitches that have been previously characterized were chosen [13,14,19], as it has been shown that sampling riboswitches from diverse phylogeny can generate highly fluorescent and well-folded RNA biosensors [7]; 2) For each riboswitch sequence, the three biosensor topologies described above were designed; 3) Based on empirically derived rules for length of the transducer stem [4], each P1-P2' construct had two possible stem lengths and each P3-P2' and P4-P2' had three possible stem lengths.

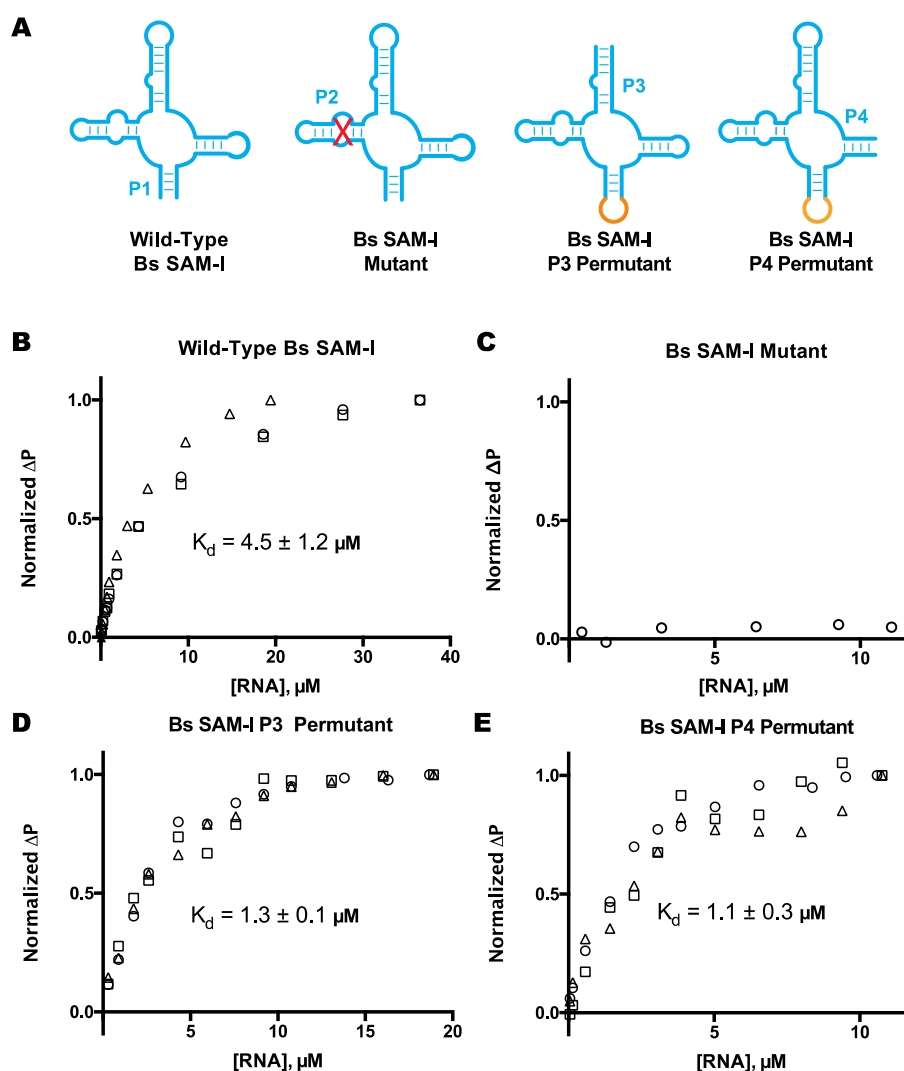


Fig. 3. Ligand binding analysis of SAM-I riboswitch aptamers. (A) Schematic of SAM-I riboswitch aptamer, mutant, and permutants. Fluorescence polarization data is shown for Cy5-labeled SAM analog with (A) wild-type Bs SAM-I (P1-7 stem), (B) Bs SAM-I mutant (P2 disruptive mutation), (C) Bs SAM-I P3 permutant (P3-5 stem), and (D) Bs SAM-I P4 permutant (P4-6 stem). Different symbols indicate independent replicates. Change in polarization (ΔP) was normalized to the maximal difference observed at the highest RNA concentration. The reported K_d value is the average of those calculated from best-fit curves with margin of error reported as standard deviation. Data from part (B) and (C) were reproduced with permission from [18].

The 32 biosensors were ordered as commercial DNA oligonucleotides, amplified by PCR, synthesized by *in vitro* transcription, and tested for fluorescence response to ligand, all in a 96-well format. In this initial high-throughput screen, 12 candidate biosensors showed a response to SAM with greater than $1.5\times$ fluorescence activation (Fig. 2B). Interestingly, 8 hits were cpRS-Sp P4-P2' designs and the remaining hits were the conventional RS-Sp P1-P2' design. None of the P3-P2' constructs were functional as biosensors, even though we separately verified that both P3 and P4 permutants of the *Bs* riboswitch aptamer retain good binding affinity to SAM (Fig. 3).

Further *in vitro* analysis of the 12 hits was performed at physiologically relevant conditions (37 °C and 3 mM Mg^{2+}) to predict their performance in bacteria (Fig. 4A). Three initial hits of the cpRS-Sp P4-P2' design with varying stem lengths derived from *Polaribacter irgensii* SAM-I did not give fluorescence turn-on in response to SAM in this secondary screen until higher ligand concentrations (200 μ M, Fig. 4B), which is consistent with their observed poorer affinity in the initial screen (Fig. 2B). The remaining biosensors still retained activity with similar fold activations.

3.2. Selectivity and sensitivity of cpRS-Sp biosensors for SAM

To compare cpRS-Sp and RS-Sp biosensor designs, constructs Bc 4-5 and Bc 1-5 were chosen as representative models. Both biosensors were derived from the same parent SAM-I riboswitch from *Bacillus clausii* and use the same base pair length in the transducer (Fig. 5A). We tested their selectivity for SAM versus *S*-adenosyl-*L*-homocysteine (SAH), which differs in one methyl group from the natural ligand. Previous studies have shown that SAM-I riboswitches are 550-fold selective for SAM over SAH [20]. The two riboswitch-based biosensors also appear fully selective for SAM over SAH at equimolar concentrations. Bc 4-5 and Bc 1-5 were measured to have dissociation constants (K_D) for SAM of ~ 0.7 and ~ 1.7 μ M, respectively. Furthermore, Bc 4-5 had ~ 2 -fold higher

maximal fluorescence than Bc 1-5. Thus, Bc 4-5, a cpRS-Sp design, has better performance than Bc 1-5, a conventional RS-Sp design.

3.3. *In vivo* detection of SAM with a cpRS-Sp biosensor using flow cytometry

While the secondary screen at physiologically relevant conditions suggests these biosensors would function in a cellular context, we wanted to verify their activity in bacterial cells. Efforts to image metabolites such as SAM are important as it is the required cofactor for enzymatic methylation and is a key metabolite in cellular sulfur metabolism [21]. Additionally, *E. coli* have been shown to contain sub-millimolar levels of SAM [22] and this high abundance would allow us to measure maximal fluorescence turn-on response in cells expressing biosensor constructs. To validate that cellular fluorescence was due to ligand-dependent turn on of the biosensors, mutant biosensors were generated that carry mutations that ablate ligand binding in the SAM-I riboswitch [13]. Additionally, Spinach2 and a Spinach2 mutant [8] that cannot bind DFHBI were included to compare the maximal fluorescence of our biosensors relative to the parent dye-binding aptamer Spinach2 and the background fluorescence of biosensor mutants to the inactive Spinach2. Each of the constructs were inserted into a tRNA scaffold on a pET31b plasmid [23]. *E. coli* cells transformed with pET31b plasmids containing each construct were analyzed by flow cytometry. In all cases, robust fluorescence was observed for functional biosensors while their corresponding mutants showed close to background fluorescence (Fig. 6). Similar to the *in vitro* results, Bc 4-5 was brighter *in vivo* than Bc 1-5. Furthermore, two cpRS-Sp designs, Bs 4-4 and Bc 4-5, possess mean fluorescence intensity values higher than the parent Spinach2 aptamer. While their signal is saturated in this *E. coli* strain, these biosensors may be useful in auxotrophic strains to study biosynthetic enzyme mutants or SAM transporters. As we have seen previously [7], by surveying diverse riboswitch sequences and a limited set of transducer stem lengths, we obtained biosensors with a range of affinities.

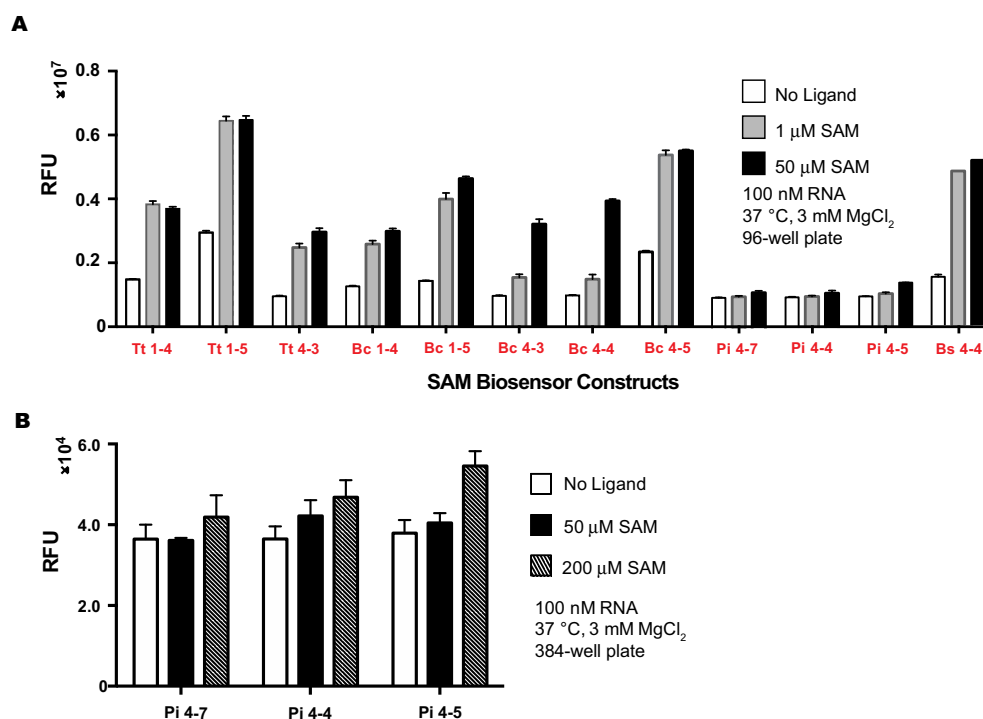


Fig. 4. Re-screen of cpRS-Sp biosensors at physiologically relevant conditions. (A) *In vitro* fluorescence response of SAM-I biosensors to SAM ligand at 37 °C and 3 mM $MgCl_2$. (B) Pi biosensors were re-screened at higher SAM concentrations. Data shown are mean values $\pm$ standard deviation taken from two independent replicates.

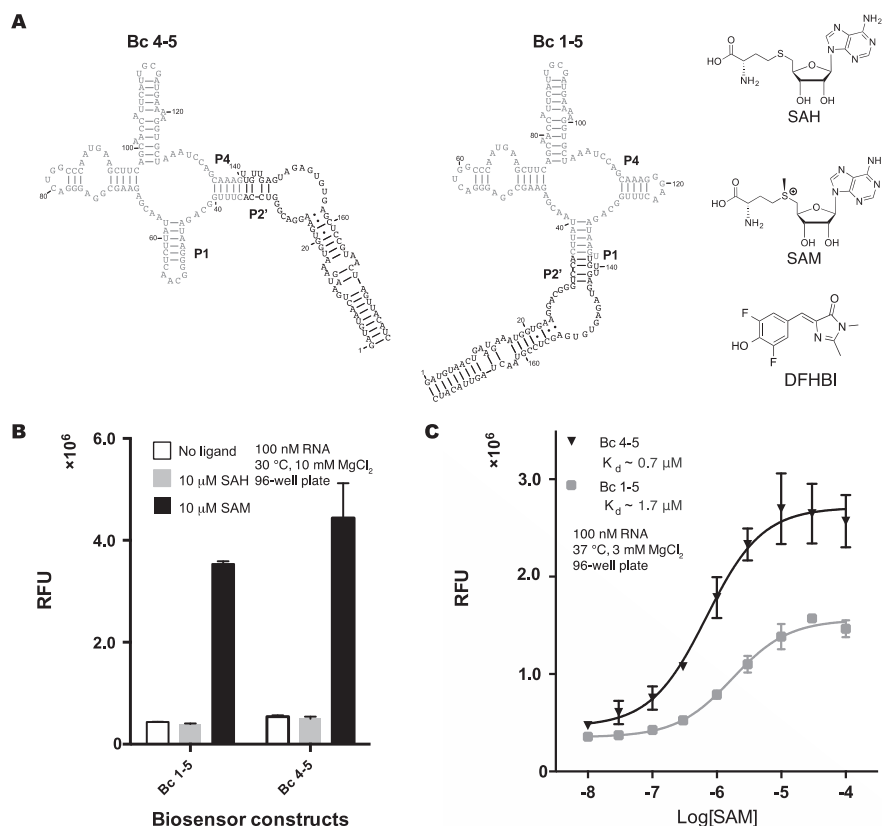


Fig. 5. Selectivity and sensitivity of cpRS-Sp biosensors. (A) Secondary structure models for Bc 1-5 and Bc 4-5 biosensors and chemical structures of the biosensor ligand, S-adenosyl-L-methionine (SAM), the structural analog S-adenosyl-L-homocysteine (SAH), and dye DFHBI. (B) Selectivity of biosensors for SAM over SAH. (c) *In vitro* analysis of biosensor binding affinities for SAM. Data shown are mean values $\pm$ standard deviation taken from at least two independent replicates.

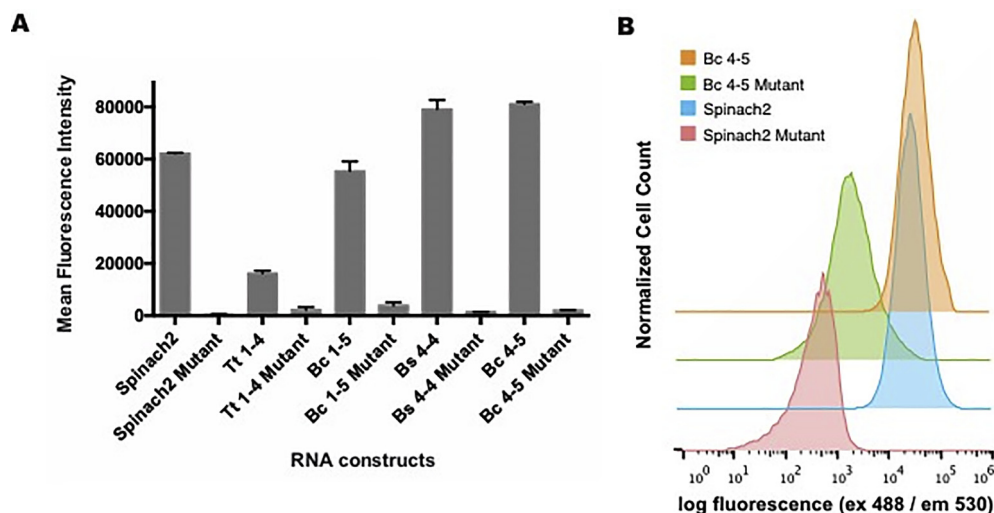


Fig. 6. *In vivo* performance of SAM cpRS-Sp biosensors. (A) Live cell fluorescence measured by flow cytometry for *E. coli* BL21* cells expressing plasmid encoding biosensors and incubated in media containing DFHBI. Mean fluorescence intensity was determined by analyzing 30,000 cells per replicate. All error bars represent standard deviation between technical replicates. (B) Representative flow histograms for cells expressing biosensors or controls.

3.4. An unexpected effect of SAM on biosensor and Spinach2 fluorescence

While re-investigating biosensor function at higher SAM concentrations, we observed that fluorescence turn-on was decreased at 500 μ M and higher ligand concentrations (Fig. 7). These results were reminiscent of a prior observation we made, that 3 mM and higher ATP reduced Spinach2 fluorescence [9], likely due to com-

petitive displacement of DFHBI from the dye-binding pocket. Similarly, we found that Spinach2 fluorescence was significantly decreased at 500 μ M and higher SAM. These surprising results lead us to suggest that the Spinach2 aptamer may have general affinity for adenosine-containing metabolites. Although the K_D for SAM or ATP is much poorer (500-fold or higher) than for DFHBI, the intracellular concentrations of these metabolites are in the range that they can compete for binding. This explains at least in part why

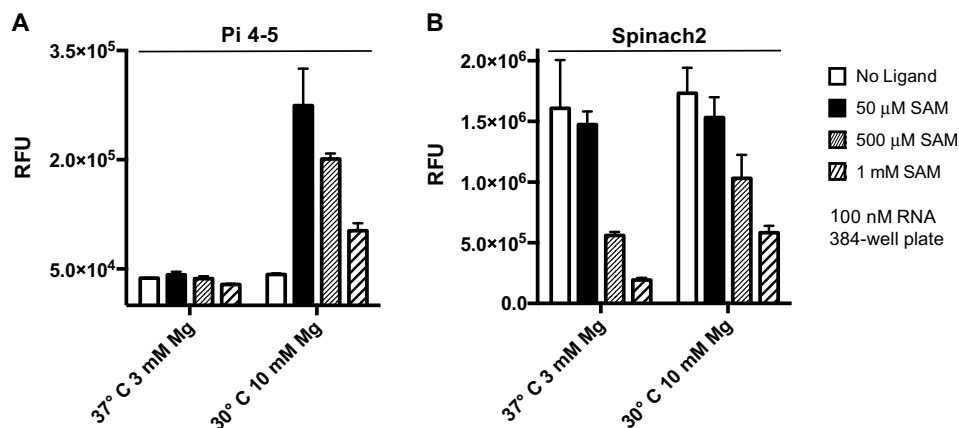


Fig. 7. Effect of SAM on biosensor and Spinach2 fluorescence. *In vitro* fluorescence activation of (A) biosensor Pi-4-5 and (B) Spinach2 at 30 °C, 10 mM MgCl₂ and 37 °C, 3 mM MgCl₂ in the presence of 0, 50, 500, and 1000 μ M SAM. Data shown are mean values $\pm$ standard deviation taken from two independent replicates.

observed fluorescence turn-on is generally lower *in vivo* than *in vitro*. In practice, we have found that using 50 or 100 μ M DFHBI gives sufficient signal for experiments in *E. coli*. Also, we always perform *in vivo* experiments with Spinach2 as a control to compare to our Spinach2-based biosensors. Biosensor function is validated for sensing changes in metabolite levels only if the same conditions result in no fluorescence change in the Spinach2 control.

4. Discussion and conclusions

This study shows that circularly-permuted riboswitches can provide an additional scaffold for designing functional RNA-based fluorescent biosensors. In fact, our results suggest it may be possible to generate more sensitive and higher fluorescent biosensors from a parent riboswitch by surveying different transducer stem topologies. There are a number of natural riboswitches or synthetic aptamers besides the SAM-I class that possess multi-stem junctions, such as the PreQ1 or FMN riboswitches [24,25] for which this strategy could be adapted to generate biosensors for their respective ligands. Also, the ability to exploit different connections to riboswitch folds may be beneficial to make biosensors with other dye-binding aptamers besides the Spinach scaffold, such as the Mango and SRB-2 aptamers for example [26,27] or to make other RNA devices [2]. Although naturally circularly-permuted riboswitches have not yet been found, our results show that permuted aptamer domains can retain or even improve on ligand binding and conformation switching, which suggests that this method may be promising for making synthetic riboswitches for gene regulation as well.

While P4-P2' cPRS-Sp constructs were functional, biosensors utilizing a P3-P2' topology were not. We verified at least in one example that the circularly permuted riboswitch aptamers still bind SAM with good affinities. There are two other potential explanations for this result. One is that the SAM-I riboswitch aptamer does not undergo a significant conformational change in the P3 stem upon ligand binding. A detailed study of the *B. subtilis* yitJ SAM-I riboswitch discovered key differences in conformational changes with and without the expression platform [28]. The riboswitch aptamer with the expression platform displayed modest structural changes in the P3 stem upon ligand addition, but the aptamer alone showed little to no modulation in the presence of SAM. Our biosensor design incorporates only the SAM-I riboswitch aptamer, so likely does not modulate at the P3 stem efficiently to alter fluorescence output. Alternatively, even though the aptamer alone is capable of binding SAM, it is possible that the positioning

of the Spinach2 aptamer at the P3 stem sterically hinders the riboswitch folding, or vice versa.

In vivo analysis by flow cytometry shows that cPRS-Sp sensors with P4-P2' topologies, Bs 4-4 and Bc 4-5, have higher mean fluorescence intensities than the Spinach2 aptamer alone (Fig. 5A). This directly contrasts with RS-cpSp sensors for SAH, which exhibit lower fluorescence than the fluorogenic aptamer alone [9]. This suggests that the placement of circular permutation on the fluorogenic aptamer hinders maximal fluorescence more than creating a circular permutation on the riboswitch.

Additionally, there have been two previously developed biosensors for SAM that use the RS-Sp and Spinach-based riboswitch topologies (Fig. 1A, C). The first SAM biosensor fused the original Spinach aptamer to the *Enterococcus faecalis* SAM-III riboswitch, which is from a different riboswitch class that utilizes a 3-way junction [4,29]. The second biosensor was a Spinach-based riboswitch generated by fusing Spinach to the *B. subtilis* yitJ SAM-I riboswitch [5], which is one of the native riboswitches used in this study. Interestingly, the only functional biosensor construct derived from this riboswitch from our designs was Bs 4-4. The Bs 4-4 biosensor representing the cPRS-Sp topology has a dissociation constant of 1.0 μ M, which is comparable to the reported dissociation constant of 1.2 μ M for the Spinach-based riboswitch [10].

Overall, this work validates our concept that circular permutations of riboswitch aptamers can add to existing design strategies for creating RNA-based fluorescent biosensors. To our knowledge, there has not been any previous biosensor designs which utilize circular permutations on the riboswitch aptamer. Furthermore, our success with modulating biosensor fluorescence via ligand-induced structural changes in the P4 stem suggests that SAM riboswitches with this alternate topology have the potential to regulate gene expression as well.

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Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at <https://doi.org/10.1016/j.ymeth.2018.02.014>.

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