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Highly selective DNA aptamer sensor for intracellular detection of coenzyme A†

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Detecting Coenzyme A (CoA) in cells is vital for understanding its role in metabolism. DNA aptamers, though widely used for monitoring many other molecules, have not been effective for CoA detection, as previous attempts at obtaining DNA aptamers for CoA using SELEX resulted in aptamers that only recognize the adenine moiety of CoA. This “tyranny” of adenine dominating in SELEX has, therefore, hampered the SELEX of aptamers specific for CoA. To meet this challenge, we employed a capture SELEX method by incorporating rigorous counter selections against adenine, adenosine, ATP, pantetheine, and pantothenic acid, resulting in a highly specific DNA aptamer for CoA over adenosine, ATP and other related metabolites such as NADH, with a dissociation constant of 48.9 μM . This aptamer was then converted to a fluorescent sensor for CoA across pH 6.4–8.0. Confocal microscopy showed its ability to visualize CoA in living cells, with fluorescence changes observed upon manipulating CoA levels. This method broadens SELEX’s application and presents a promising approach for studying and understanding CoA dynamics.

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Introduction

Aptamers, single-stranded DNA (ssDNA) or RNA molecules can fold into specific three-dimensional structures to selectively bind their target molecules.^{1,2} Aptamers are smaller and more cost-effective than protein-based sensors, and their target binding can readily be translated into a DNA/RNA hybridization melting temperature change that is quite predictable and often results in a large signal change when they are labeled with a Förster resonance energy transfer (FRET) pair. Because of these advantages, aptamers have been increasingly applied in sensing and imaging in biological systems.^{3–22}

Another major advantage of the aptamers is that they can be selected to be highly specific to a wide range of targets using

Systematic Evolution of Ligands by Exponential Enrichment (SELEX).^{23–32} In fact, the SELEX is so powerful, it is considered to be generally applicable to almost any target. However, despite the perception, applying SELEX to obtain aptamers for some targets remains a significant challenge. One example of challenging metabolites is coenzyme A (CoA), a molecule with a 3'-phosphoadenosine diphosphate structure that closely resembles adenine-containing metabolites like ATP (Fig. 1). This “tyranny” of adenine dominating in SELEX has, therefore, hampered the SELEX of aptamers specific for CoA, as previous attempts yielded only aptamers that bind the adenine moiety of CoA, resulting in poor selectivity over adenosine, ATP, and other adenosine-containing metabolites such as NADH.^{33–35} As a result, despite decades of research and advances in aptamer

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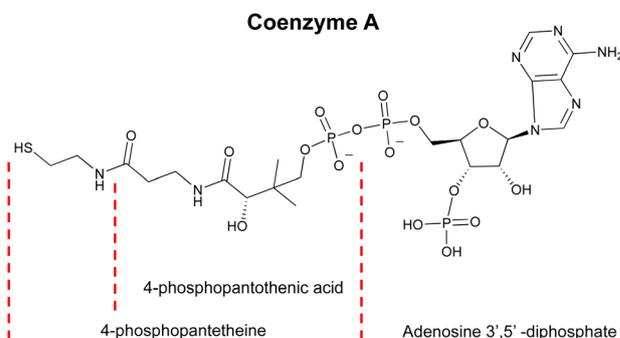


Fig. 1 Chemical structure of Coenzyme A (CoA), showing key structural motifs including adenosine 3',5'-diphosphate, 4-phosphopantetheine, and 4-phosphopantothenic acid.



technology, there is no aptamer ideal for detecting CoA due to selectivity challenges.^{33–35}

CoA is a vital metabolite central to acyl group transfer and enzymatic regulation.^{36,37} It functions both as a substrate and allosteric modulator in critical pathways, including post-translational modifications like acetylation of histones and other proteins, which influence gene expression and cellular function.^{38,39} Maintaining CoA homeostasis is essential for cellular health, as disruptions are linked to conditions including Pantothenate Kinase-Associated Neurodegeneration (PKAN),⁴⁰ CoA Synthase Protein-Associated Neurodegeneration (CoPAN),⁴¹ cataracts, cardiomyopathy, and other CoA synthesis pathway diseases.³⁷ Real-time monitoring of CoA in living cells is therefore essential to understanding its metabolic roles and disease implications.

Real-time detection of CoA in living cells remains challenging due to its structural complexity and similarity to other biologically relevant small molecules.^{37,42,43} Traditional methods, such as high-performance liquid chromatography (HPLC) and mass spectrometry, while sensitive, are unsuitable for real-time and spatially resolved measurements in living cells.^{44–46} Semisynthetic biosensors have been developed to couple a green fluorescent protein-HaloTag fusion protein with a CoA-dependent fluorescent ligand to produce Förster resonance energy transfer (FRET) signals.⁴⁷ Despite progress, it is challenging to convert CoA binding into the FRET signals, as this conversion requires careful system design to translate the binding of the small CoA molecule into conformation changes in a relatively larger protein.⁴⁸ As a result, the FRET signals are often weak, making CoA detection difficult in the complex cellular environment.⁴⁹ Thus, a rapid, simple, sensitive, specific, and cost-effective method for monitoring CoA levels in cells is still needed.

In this study, we address the issue of selectivity in CoA detection *via* aptamers by utilizing a capture SELEX method specifically designed to isolate DNA aptamers that are highly selective for CoA. We incorporate rigorous counter-selections against a variety of adenine-containing molecules, allowing us to overcome previous limitations. The resulting aptamer was then converted into a fluorescent sensor for the real-time detection and visualization of intracellular CoA levels. This approach provides not only a sensitive, specific, and cost-effective tool for studying CoA metabolism, but also represents a significant advancement in the selection and development of aptamer-based detection technologies.

Results and discussion

SELEX of DNA aptamers for CoA against adenine or other molecules containing the adenine moiety

Previous SELEX efforts to isolate CoA-binding aptamers resulted in those that primarily bind the adenine portion of CoA. As a result, the isolated aptamers can also bind ATP.^{33,34} In these SELEX processes, CoA was attached to a solid support at a different attachment point, and a DNA library was then added to the solid support containing the CoA to isolate sequences that bind CoA. We hypothesize that the attachment of CoA to

the solid support can significantly impact the selection outcome, often imposing steric constraints that eliminate desirable binding structures. To address this issue, we employed a capture SELEX method that leverages the aptamer's inherent ability to transition between two distinct conformations: a duplex formation with an antisense DNA strand and a complex structure binding its specific target, CoA.^{50–52} This SELEX method was achieved by immobilizing the structural-switching DNA library onto a solid support, followed by introducing CoA that remains free in its native state in solution. This library-immobilized SELEX strategy ensured that all the epitopes of CoA were exposed to the DNA library and could participate in aptamer binding. The DNA molecules that bound to CoA were subsequently released from the support. This approach aimed to remove steric constraints and facilitate the isolation of a more diverse and selective population of CoA-binding aptamers. More importantly, we enriched the DNA pools with an affinity for the (*R*)-pantetheine, a subunit of CoA, and then employed in SELEX for CoA. Moreover, starting at the 6th round of selection, we incorporated rigorous counter-selection steps against other molecules, including adenine and molecules with adenine as a component of their structures, such as ATP. Specifically, the CoA aptamer was selected using a DNA library containing 82 nucleotides (Fig. 2a). This library includes 40-nucleotide randomized sequences (red) flanked by two constant sequences (blue) on each end that include 5'-GCAGTCGGCGTCCGACAG-3' and 5'-CTGTCCGACGATGTAACGCTTCAC-3', which serve as primers for PCR amplification. To ensure successful selection without altering the chemical properties of CoA, we chose not to covalently link the CoA to a solid support as in previous CoA SELEX studies.^{33–35} Instead, we immobilized the DNA library on microbeads through a capture strand DNA that can hybridize to the underlined sequences in one of the two flanking regions, 5'-GCAGTCGGCGTCCGACAG-3'. Only the DNA molecules with sequences that bound to CoA and subsequently altered their secondary structure were released from the microbeads into the solution (Fig. 2a).

To ensure high selectivity for CoA against adenine and other molecules containing the adenine moiety, the first four SELEX rounds were initially carried out using (*R*)-pantetheine, a subunit of CoA, to enrich the DNA pools with an affinity for the (*R*)-pantetheine portion of CoA (Table S1†). At round 5, the selection target was switched to CoA to isolate DNA aptamers that bind CoA. The concentrations of CoA used in the selection were gradually decreased from 5 mM to 0.1 mM to enrich aptamers with a strong binding affinity for CoA. To achieve high selectivity, counter-selections against adenine, adenosine triphosphate (ATP), pantetheine, and pantothenic acid were introduced from the 6th round of selection, with gradually increasing concentrations. Specifically, from the 6th round of selection, 1 mM adenosine and 9 mM ATP were introduced as counter-selection targets and any DNA molecules eluted with adenosine or ATP were discarded from the selection. Additionally, 150 μM pantetheine was introduced as a counter-selection target starting from round 8 through round 13 and 150 μM of pantothenic acid was introduced as a counter-selection target starting from round



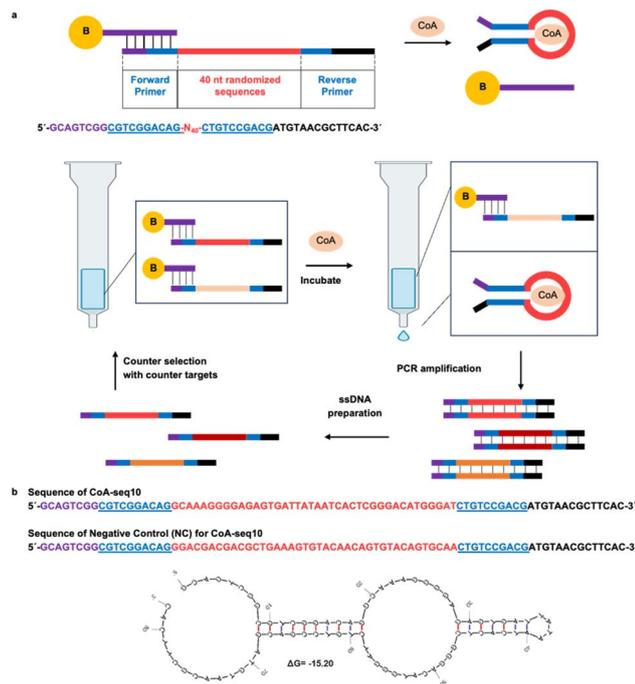


Fig. 2 Selection of CoA aptamer. (a) *In vitro* selection process for obtaining the CoA aptamer. Each DNA library strand contains a 40-nucleotide randomized sequence (red) flanked by two constant sequences (purple, blue, and black) at both ends, serving as PCR primers. The binding region between the capture strand and the DNA library is highlighted in purple. The forward primer consists of purple and blue regions, and the reverse primer is composed of blue and black regions. After immobilizing the DNA library onto beads (yellow), CoA is introduced into the solution. DNA strands that bind to CoA undergo a structural switch, detaching from the capture strand and entering the solution, forming a stem-loop structure with sequences represented by blue color. These eluted strands are then PCR amplified, and ssDNA libraries are prepared. After re-immobilizing the DNA library onto the beads, counter-selection molecules are added to the solution, and the non-specific DNA strands are removed. The DNA library is then incubated with CoA, and the eluted CoA-aptamer DNA is collected. This selection process is repeated multiple times. (b) CoA-seq10 aptamer sequence and its predicted secondary structure, generated using UNAFold software.⁵⁶ The sequence is illustrated with different colors, whose function is explained in legends of (a). The UNAFold prediction illustrates the secondary structure of the CoA aptamer upon binding to CoA.⁵⁶

12. These counter-selections were introduced to remove any aptamers that interacted with molecules that bear similarities to only a component of the CoA, rather than CoA itself. To monitor the selection progress, we employed qPCR to determine the elution yield from each selection round.⁵³ The elution yield is defined as the amount of ssDNA bound to CoA and eluted from the microbeads into solution, divided by the total amount of ssDNA added to the microbeads (Fig. S1†). When the elution yield stopped increasing, high-throughput sequencing (HTS) was utilized to analyze the sequences from round 9 to round 13, enabling us to track the enrichment of individual sequences. After sorting the sequences by sequencing quality and identifying those with intact forward and reverse primer sequences, we used the FASTAptamer analysis toolkit to find sequences or

clusters that were conserved across all rounds.^{54,55} Specifically, we used the FASTAptamer-count and FASTAptamer-cluster functions to analyze the abundance and similarity of individual sequences, tracking changes across different SELEX rounds.⁵⁴ We then applied the FASTAptamer-enrich function to monitor changes in sequence distribution across selection rounds and identified representative enriched aptamer sequences (Table S2†).⁵⁴ We then tested their binding affinity to CoA by labeling the aptamer candidates with a 5'-FAM fluorophore and the capture strand that attached to the microbeads, as shown in Fig. 2a, with a 3'-Black Hole Quencher®-1 (BHQ1), measuring the increase in fluorescence after the candidates bind CoA, which causes its strand displacement from the capture strand (Fig. 3d). CoA-seq10, a most active sequence and continuously enriched during SELEX, was chosen for sensor development (Fig. 2b and S2†). As illustrated in Fig. 2b, CoA-seq10 is predicted to form a stem-loop secondary structure, as determined using UNAFold software.⁵⁶

Characterization of the CoA-seq10 aptamer and its conversion into a fluorescent sensor for CoA

To characterize the binding of CoA-seq10 to CoA and its affinity, isothermal titration calorimetry (ITC) was employed, revealing

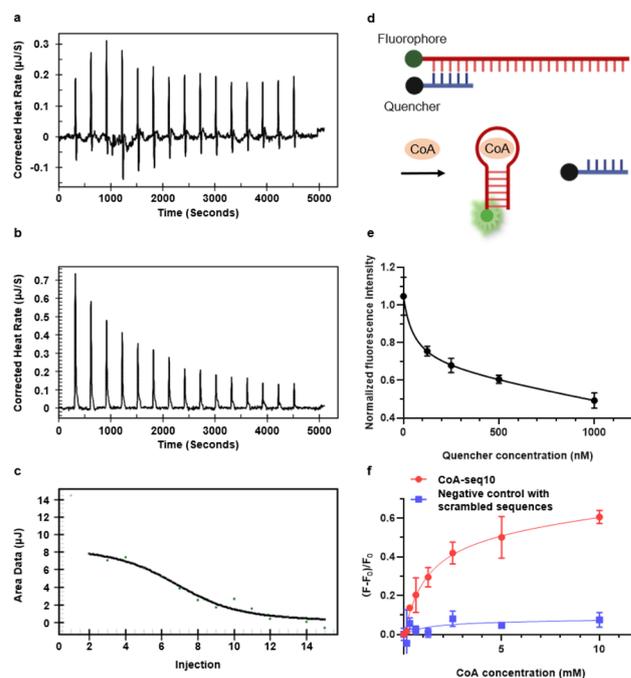


Fig. 3 Characterization of CoA-seq10 aptamer sensor. (a) Thermogram for the ITC titration of 300 μM CoA-seq10 titrated by 3.2 mM CoA in aptamer binding buffer; (b) thermogram for the ITC titration of aptamer binding buffer by 3.2 mM CoA in the same buffer; (c) integrated heat of the ITC titration for CoA-seq10 and CoA, the black line represents the binding curve fitted with the 'one set of binding sites' model; (d) general setup of the CoA aptamer sensor's structure switching mechanism. (e) Normalized fluorescence intensity of different aptamer:quencher ratios, when aptamer concentration is 50 nM. (f) Normalized fluorescence of CoA-seq10 aptamer sensor versus a negative control with scrambled sequences.



a dissociation constant (K_d) of 39.8 μM (Fig. 3a–c), which is comparable to other small molecule aptamers, like the ATP aptamer with a K_d of 6 μM .³ To convert the CoA aptamer into a fluorescent sensor, we utilized an intramolecular strand displacement fluorescence sensor design (Fig. 3d). Specifically, a FAM fluorophore was conjugated to the 5' end of the aptamer CoA-seq10, and a Black Hole Quencher®-1 (BHQ1) was attached to the 3' end of the quencher strand. When the aptamer strand hybridizes with the quencher strand ($T_m = 62.3$ °C), it brings the quencher next to the fluorophore, resulting in the quenching of the FAM fluorophore by BHQ1. The binding to CoA by the CoA-seq10 aptamer weakens this hybridization (T_m decreases to 0 °C), allowing the quencher strand to dehybridize from the aptamer strand, causing a significant increase in fluorescence.

To minimize background fluorescence, we optimized the ratio between aptamer and quencher and found an aptamer:quencher ratio of 1 : 5 to be optimal (Fig. 3e). With this aptamer:quencher ratio, the fluorescence of the aptamer sensor increased with increasing concentrations of CoA, with no significant further increase observed at 10 mM. The limit of detection (LOD) was determined to be 0.126 mM (Fig. 3f), based on $3\sigma_b/\text{slope}$, where σ_b is the standard deviation of three blank samples. Additionally, replacing the CoA aptamer with a DNA of the same length but with a scrambled sequence did not increase fluorescence signals in the presence of CoA. This result underscores the specificity of the aptamer in sensor development. The fluorescent sensor can also serve as an alternative method to determine K_d values for the aptamer-CoA interaction. In this method, a K_d for the quencher with the aptamer in the absence of CoA was determined to be 55.38 nM (Fig. 3e), while a K_d for the CoA-induced aptamer–quencher dissociation was found to be 1.133 mM (Fig. 3f). Therefore, using a formula reported previously,⁵⁷ the K_d for the aptamer-CoA interaction was calculated from these two K_d 's to be 48.9 μM . This fluorescent method provides an independent method to evaluate the K_d , which is similar to the K_d (39.8 μM) obtained by ITC. Additionally, additional mutations were introduced into the CoA-seq10 aptamer to identify the critical regions involved in its binding and fluorescence response to CoA. The data in Fig. S3† show that these mutations in the aptamer sequence resulted in a decrease in the fluorescence response to CoA, indicating that these specific regions of the aptamer are crucial for both CoA recognition and the conformational change needed to trigger the fluorescence response.

To further validate the specificity of the CoA-aptamer against other intracellular metabolites, we tested its fluorescence response to various concentrations of CoA and compared it to the responses elicited by other structurally similar molecules, such as acetyl coenzyme A (acetyl-CoA), ATP, and D-pantothenic acid. As shown in Fig. 4a, other metabolites, including acetyl-CoA, ATP, and D-pantothenic acid, did not cause any significant increase in fluorescence intensity. In contrast, the sensor exhibited a marked increase in fluorescence intensity as the concentration of CoA increased. Further testing involved incubating the CoA aptamer sensor to a variety of intracellular metabolites at physiologically relevant concentrations (Fig. 4b). The metabolites include: (1) acetyl-CoA, (2) D-pantothenic acid,

(3) nucleotides (ATP, CTP, GTP, UTP), (4) other adenine-based metabolites ADP, nicotinamide adenine dinucleotide (NAD), NADH, NADP, NADPH, (5) CoA derivatives, such as propionyl-CoA, butyryl-CoA, hexanoyl-CoA, malonyl-CoA, succinyl-CoA, 3-hydroxy-3-methylglutaryl (HMG-CoA), lauroyl-CoA, myristoyl-CoA, and oleoyl-CoA, and (6) divalent cations like Ca^{2+} and Mg^{2+} . Additionally, to rule out the possibility that the CoA aptamer can still bind the competing targets of CoA without structural changes and thus no detachment of the quencher-labeled oligo, we used ITC to measure the Can10 aptamer's binding with ATP and found no evidence of binding (Fig. S4†). At physiologically relevant concentrations, none of these metabolites induced an increase in the fluorescence signal, confirming the high specificity for CoA aptamer sensor against all other metabolites, including those that share a similar component (e.g., adenine) as CoA.

Cellular environments vary significantly in pH, ranging from acidic compartments such as lysosomes to more alkaline areas like the mitochondrial matrix. To demonstrate that the CoA aptamer sensor can work across different pH levels in diverse physiological conditions, we evaluated its pH sensitivity across a range of physiologically relevant pH values. As shown in Fig. S5,† the CoA aptamer sensor maintained a consistent fluorescence response to CoA, with fluorescence intensity increasing when CoA concentration increases, across the tested pH range (pH 6.4–8.0). These results indicate that the aptamer can function and generate fluorescence signals across these physiological pH conditions.

Application of the aptamer sensor to monitor CoA in living cells

To demonstrate the applicability of the CoA aptamer sensor in visualizing CoA distribution and abundance in living cells, we delivered it into HeLa cells as a representative cell line and visualized it using confocal laser-scanning microscopy (CLSM). Although FAM is a commonly used fluorophore for studies in test tubes, we did not use it to label the CoA aptamer sensor for cellular studies since FAM is sensitive to different pH environments inside the cells. Instead, we used Cy5.5 to label the CoA aptamer strand and Iowa Black® RQ to label the quencher strand and then delivered them into HeLa cells using lipofectamine 3000. As shown in Fig. 5, S6 and S7,† bright signals from the CoA aptamer sensor were observed in all HeLa cells. In contrast, using a negative control (NC) of a DNA oligo with the same length as the CoA aptamer but the aptamer sequence replaced with a scrambled sequence (see Fig. 2b) resulted in a reduced fluorescence signal. This negative control (5Cy5/ATGCAGTCGGCGTCGGACAGGGACGACGACGCTGAAAGTGTA CAACAGTGTACAGTGCAACTGTCCGACGATGTAACGCTTCAC) highlights the essential role of the aptamer in recognizing CoA using a DNA sequence of the same length as the CoA aptamer but with a scrambled sequence—where the region that binds to the quencher strand is preserved while the CoA-binding portion is disrupted—resulted in a reduced fluorescence signal. This negative control highlights the essential role of the aptamer in recognizing CoA. To assess the duration over which the



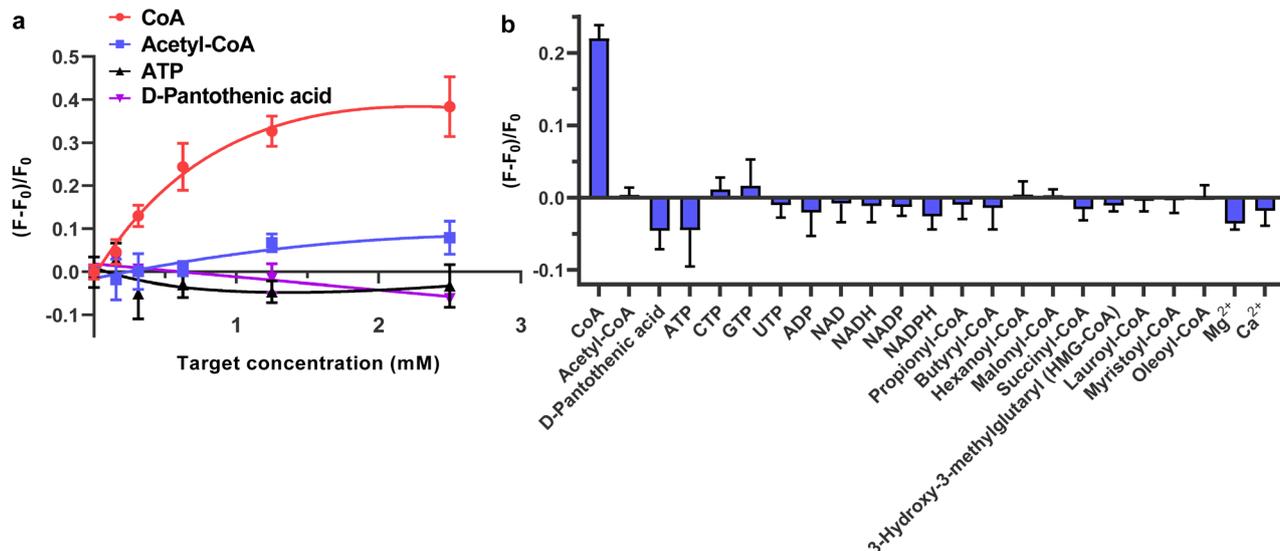


Fig. 4 CoA aptamer sensor's selectivity against similar molecules to CoA. (a) Normalized fluorescence of the CoA-seq10 aptamer sensor's selectivity with concentration dependence of analytes. (b) Normalized fluorescence of the CoA-seq10 aptamer sensor's selectivity with physiologically relevant concentrations of analytes.

fluorescence signal from the CoA-seq10 aptamer sensor can be distinguished from the negative control (NC) sensor after transfection, a time-course study was performed. As shown in Fig. S6 and S7,[†] 3.5 hours after the transfection, the fluorescence signal from the CoA aptamer is significantly higher than that of the NC sensor for up to 8 hours. Although the fluorescence signal from the CoA-seq10 aptamer decreases over time, it remains distinguishable from the NC sensor. To ensure our sensor can monitor CoA selectively, we added either Pantazine 2891 (PZ-2891),⁵⁸ which is an allosteric PANK activator that increases free CoA concentration inside cells, presumably by disrupting pantothenate kinase (PanK) feedback regulation^{59,60} or hopantenate (HoPan), an analog of pantothenate, which is used to chemically inhibit CoA biosynthesis.^{61,62} After treating the HeLa cells with either 3 μ M PZ-2891 or 1 mM HoPan to

regulate the cellular CoA levels, we added Cy5.5-labeled aptamer sensors and compared them with untreated cells. As shown in Fig. 5, compared with untreated cells, PZ-2891 treated cells exhibited a significant fluorescence increase in the red channel, while HoPan-treated samples showed a lower red fluorescence signal. Furthermore, the fluorescence intensity of the negative control with scrambled sequence produced relatively weak fluorescence signals compared to untreated cells (Fig. 5). In addition, we conducted a time-course drug treatment with 3 μ M PZ-2891 in HeLa cells to assess the dynamic response of the CoA-seq10 aptamer sensor under drug exposure, measuring the fluorescence signal at multiple time points post-treatment. As shown in Fig. S7 and S8,[†] the fluorescence intensity of the CoA-seq10 aptamer sensor is significantly higher than the no-treatment control after 6 hours of incubation with 3 μ M PZ-

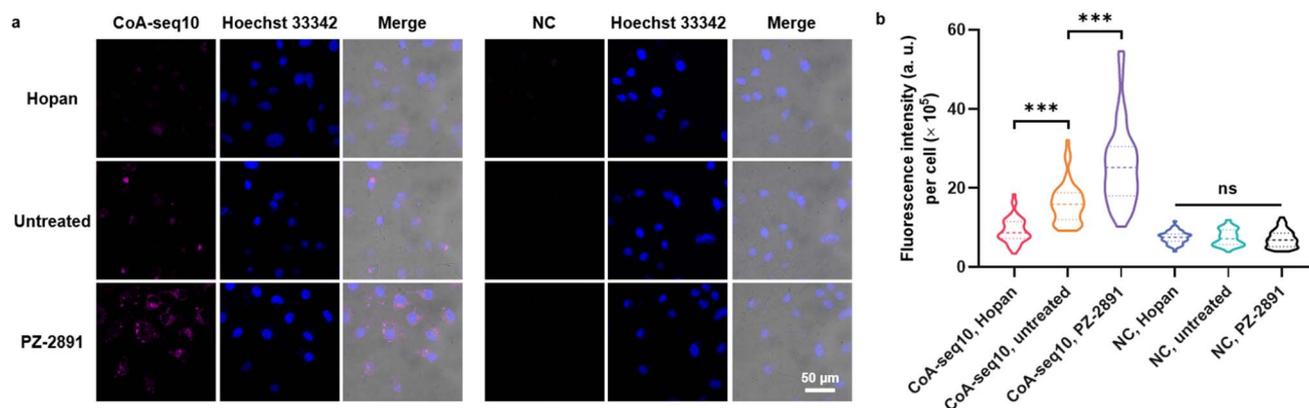


Fig. 5 CoA aptamer sensor detects CoA regulation in HeLa cells. (a) CLSM images of CoA using the CoA-seq10 sensor and NC sensor in live HeLa cells during CoA regulation. (b) Quantification of average fluorescence intensity (arbitrary units, a.u.) per cell shown in (a); *** $P < 0.0001$. Data in (a) and (b) represent three independent experiments; $n = 5$ frames. Scale bar is 50 μ m. Data are shown as mean \pm s.d. Statistical significance was determined by unpaired two-tailed Student's t -test; NS, not significant ($P > 0.05$); *** ($P < 0.001$).



2891. A statistically significant difference is observed between the CoA-seq10 aptamer sensor and the NC sensor at each drug treatment time point. These results collectively demonstrate that the CoA aptamer sensor can be applied to monitor intracellular CoA.

Conclusions

CoA is an essential metabolite involved in key biological processes, including the synthesis and oxidation of fatty acids and the citric acid cycle for energy production. Despite its critical role, sensors for CoA are limited. While SELEX has been a widely used method for generating DNA and RNA aptamers for many targets, efforts to isolate aptamers for CoA have largely failed, with most aptamers binding only the adenine moiety of CoA, highlighting a major limitation of the SELEX methods to obtain aptamers with high selectivity. To claim that SELEX can be generally applied to almost any target, we need to develop methods to overcome this major limitation.

To meet this challenge, we have improved the SELEX process by incorporating rigorous counter-selections against adenine, ATP, pantetheine, and pantothenic acid. This approach led to the successful identification of a highly selective DNA aptamer, CoA-seq10, making this aptamer capable of monitoring CoA at the physiological concentration of CoA in the cytoplasm (0.02 to 0.14 mM) and mitochondria (2.2 to over 5 mM).³⁶ These results were confirmed using ITC and fluorescence assays, with a limit of detection (126 μ M). The aptamer's specificity allowed it to distinguish CoA from other intracellular metabolites and function effectively across different physiological pH levels. Confocal microscopy further demonstrated the aptamer sensor's ability to visualize CoA in living cells, with significant fluorescence changes upon regulating cellular CoA levels. In summary, this work contributes to SELEX methodology, overcoming some of its limitations and enabling the development of a CoA-selective aptamer that opens new opportunities for studying CoA metabolism in living cells.

Data availability

All data are available in the main text or the ESI†.

Author contributions

Conceptualization: Yuan Ma, Whitney Lewis, Yi Lu. Methodology: Yuan Ma, Whitney Lewis, Peng Yan, Linggen Kong, Quanbing Mou, Xiangli Shao, Weijie Guo. Investigation: Yuan Ma, Whitney Lewis, Peng Yan, Linggen Kong (*in vitro* selection of CoA aptamers); Yuan Ma, Whitney Lewis, Quanbing Mou (aptamer characterization experiments and data analysis); Yuan Ma, Whitney Lewis, Xiangli Shao, Weijie Guo (cell experiments and data analysis). Writing – original Draft: Yuan Ma, Whitney Lewis, Yi Lu. Writing – review & editing: all authors. Supervision: Yi Lu. Equal Contribution: Yuan Ma and Whitney Lewis contributed equally to this work.

Conflicts of interest

There are no conflicts to declare.

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