A cAMP Sensor Based on Ligand-Dependent Protein Stabilization

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ABSTRACT: cAMP is a ubiquitous second messenger with many functions in diverse organisms. Current cAMP sensors, including FRET-based and single-wavelength-based sensors, allow for real-time visualization of this small molecule in cultured cells and in some cases in vivo. Nonetheless, the observation of cAMP in living animals is still difficult, typically requiring specialized microscopes and ex vivo tissue processing. Here we used ligand-dependent protein stabilization to create a new cAMP sensor. This sensor allows specific and sensitive detection of cAMP in living zebrafish embryos, which may enable new understanding of the functions of cAMP in living vertebrates.

Cyclic adenosine 3′–5′-monophosphate (cAMP) is an essential second messenger that amplifies environmental signals received by G-protein-coupled receptors (GPCRs). The importance of cAMP is underscored by the multitude of physiological processes that it regulates including heartbeat, learning, and memory, and it can initiate a wide range of cellular responses including proliferation, differentiation, and death. The kinetics and function of cAMP depend on many factors including the cell type and subcellular compartment where it accumulates and interacts with other signaling molecules, which highlights the importance of sensors to visualize cAMP in cultured cells and intact organisms.

Fluorescent sensors for cAMP based on Förster resonance energy transfer (FRET) respond rapidly to local changes in cAMP concentration. Although useful in cultured cells, FRET-based sensors are characterized by low signal-to-noise ratio, are prone to photobleaching, and are hampered by light scattering in intact tissues, limiting their use for in vivo imaging. Single wavelength cAMP sensors using circularly permuted fluorescent proteins, such as Pink Flamindo, R-FlincA, and cAMPr, can mitigate some of the disadvantages of FRET-based sensors. Single wavelength sensors have been used to track fast cAMP dynamics in dissected Drosophila brains, in C. elegans neurons, and in mouse astrocytes. However, there is often need for dissection and an ex vivo imaging, and it is not clear whether the current tools are useful for long-term imaging.

Ligand-dependent protein stabilization is a strategy that can be used to generate single-wavelength biosensors for small molecules. This approach is based on a protein that is stable only when bound to its cognate ligand. This sensor is metabolically unstable and degraded by the proteasome in its unbound state, but engagement by its cognate ligand prevents degradation and leads to a dose-dependent fluorescent signal when the sensor is fused to an appropriate partner such as GFP. Here we report the application of this approach to develop a genetically encoded single-wavelength cAMP sensor for in vivo imaging using the zebrafish model system (Figure 1; Figure S1).

We sought to identify conditionally stable mutants of the cyclic nucleotide binding domain (CNBD) from the MlotiK1 bacterial channel, which binds cAMP with high specificity and sensitivity in mammalian cells. We started with a codon-optimized CNBD sequence and generated a library of sequence variants using error-prone PCR. This library of CNBD domains was then fused to GFP to enable a cell-based screen for mutants stabilized by cAMP (Figure S1A,B). NIH3T3 cells were stably transduced with the library and subjected to serial rounds of FACS sorting after application or withdrawal of the adenylyl cyclase agonist forskolin (FSK). We selected clones with high GFP signal in the presence of FSK and very low GFP signal in the absence of FSK (Figure S1B). DNA sequencing of selected clones identified many variants encoding unique missense mutations in CNBD, and two variants in particular displayed 3-fold (N41) to 4-fold (N49) increases in GFP intensity upon forskolin treatment (Figure 1B; Figure S1C,D). The N49 variant contains a single mutation in the CNBD domain (Figure S1C,D; blue), whereas N41 contains four mutations.

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To further characterize this cAMP sensor, we generated stable transgenic fish lines expressing DDcAMP under control of the regulatory sequences from the ubiquitin gene (abbreviated as Tg(ubi:DDcAMP), Figure 2A–C), which drives widespread expression in the embryo.\(^2\) As controls, we generated transgenic fish lines expressing ubiquitin-GFP with no CNBD (Figure 2D–F) or the cAMP-insensitive variant DDcAMP\(^{R307Q}\) (Figure 2G–I). In situ hybridization demonstrated that the transgenes expressed GFP mRNA at similar levels at 24 h postfertilization (hpf) (Figure 2A,D,G), whereas confocal imaging revealed differences in GFP signal. In contrast to the control transgenes (Figure 2E,F,H,I), embryos expressing the sensor displayed discrete GFP signal localized to the horizontal myoseptum of the somites (Figure 2B,C, white arrows). Quantification confirmed that a region containing muscle pioneers displayed much more GFP signal than a reference area not containing muscle pioneers in Tg(ubi:DDcAMP) transgenic embryos, but not in embryos expressing the cAMP-insensitive variant cAMP-DDcAMP\(^{R307Q}\) (Figure S2C). In Tg(ubi:DDcAMP) embryos from different transgenic founders and in transgenic embryos expressing the N49 sensor variant Tg(ubi:N49-GFP), GFP expression was enriched in the muscle pioneers and in the slow superficial fibers (Figure S2Ba–c; yellow arrowheads), cells known to require cAMP signaling for proper specification and differentiation.\(^2\) Thus, the comparison of sensor and control transgenic animals provides evidence that DDcAMP is specifically stabilized in a subpopulation of muscle cells.

To test the specificity and sensitivity of DDcAMP to cAMP, we treated Tg(ubi:DDcAMP) transgenic embryos with FSK for 20 h starting at 4–5 hpf and imaged fluorescent signal in anterior somites using confocal microscopy (Figure 3A,B). FSK significantly increased GFP expression in Tg(ubi:DDcAMP) but not in Tg(ubi:DDcAMP\(^{R307Q}\)) embryos, and the overall GFP signal in Tg(ubi:DDcAMP\(^{R307Q}\)) embryos was low (Figure 3B). Transgenic sensor animals Tg(ubi:DDcAMP) showed increased GFP intensity in the enveloping layer (EVL) of the tail (Figure 3C) when treated with increasing concentrations of the cell-permeant cAMP analog 8-cpt-cAMP for 24 h (Figure 3D). The sensor was not activated by treatment with the cGMP analog 8-cpt-cGMP, and treatment with 8-cpt-cAMP did not increase signal in Tg(ubi:DDcAMP\(^{R307Q}\)) control animals (Figure 3D, Figure S3B). Taken together, these experiments provide evidence that DDcAMP is a specific and sensitive sensor for cAMP in the developing embryo.

Time course and time lapse analyses of Tg(ubi:DDcAMP) indicate that GFP signal appeared in adaxial cells of developing somites at 15 hpf (Movie 1; Figure 3E, red arrow; Figure S3C, white arrows). The signal increased in the somites over time (Figure S3C), pointing to a potential time window of cAMP production during somitogenesis. Further kinetic analyses on Tg(ubi:DDcAMP) animals showed that the sensor provides a 3-fold increase in fluorescence intensity when treated with FSK for 4 h starting from 14 to 18 hpf (Figure 3F). To compare DDcAMP with the existing cAMP sensor cAMPR,\(^1\) we

spread across the CNBD protein structure (Figure S1C,D; pink).

To test the forskolin responsivity of these CNBD variants in vivo, we injected synthetic mRNA encoding the N41-GFP and N49-GFP sensors into zebrafish embryos, allowing ubiquitous and transient expression of the sensor proteins (Figure 1C,D). Four or five hours after injection of mRNA, we treated embryos with 20 μM FSK for 20 h and subsequently imaged the GFP signal (Figure 1E). Both variants displayed a visible fluorescent signal when expressed in zebrafish embryos, but only N41 increased in response to FSK treatment (Figure 1F).

Figure 1. (A) Ribbon diagram of the cAMP sensor composed of CNBD-GFP protein that is unstable without cAMP but stabilized upon cAMP binding: solution structure of a bacterial cyclic nucleotide-activated K\(^+\) channel binding domain in cAMP-free form on the left (PBD 2K0G) and bound to cAMP on the right (PBD 2K0G). (B) NIH 3T3 cells stably expressing CNBD-GFP derived from error-prone PCR were treated with 1% serum or 20 μM forskolin for 17 h. (C) Synthetic mRNA encoding the sensor variants N41 and N49 CNBD and the corresponding cAMP-insensitive controls (R307Q). (D) mRNA was injected into zebrafish embryos at the one-cell stage. Embryos were treated at 4–5 hpf with FSK or DMSO for 20 h, then imaged at 24–25 hpf. (E) Representative images showing 24 hpf embryos injected with N41-GFP and N41\(^{R307Q}\)-GFP mRNA at one-cell stage and treated with DMSO or 20 μM FSK starting at 4 hpf. (F) Each dot represents mean GFP intensity of one embryo. Error bars indicate SD: \(*p < 0.5\) by one-way ANOVA (with Sidak’s multiple comparisons), \(n = 13–22\) embryos each condition; ns = not significant. AU = arbitrary unit.

protein, because DMSO-treated N41\(^{R307Q}\)-GFP embryos displayed greater GFP signal than DMSO-treated embryos expressing N41-GFP (Figure 1E,F). These analyses provided evidence that the N41-GFP sensor protein is selectively stabilized upon binding to cAMP in vivo, and we hereafter refer to this engineered sensor as DDcAMP (destabilized detector of cAMP).

To test whether cAMP binding is required for the response of the N41 variant to forskolin, we compared FSK treatment of developing embryo.

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generated a stable transgenic fish line expressing cAMP driven by the *ubi* promoter *Tg(ubi:cAMPr)*. Transgenic cAMPr originating from one founder expressed GFP in slow muscle cells, similar to DDCAMP (Figure 3G), whereas cAMPr transgenic fish from another line expressed detectable GFP in skin but not muscle cells (Figure S3D). Signal from cAMPr increases after few minutes of FSK treatment in cell culture, skin but not muscle cells (Figure S3D). Signal from cAMPr originating from one founder expressed GFP in slow muscle cells, similar to DDcAMP (Figure 3G), whereas cAMPr originating from another founder expressed GFP in fast muscle cells, similar to DDcAMP (Figure 3G), whereas cAMPr originating from one founder expressed GFP in slow muscle cells (Figure S4A). In contrast to *Tg(ubi::DDcAMP)*, GFP intensity did not significantly increase in response to FSK in *Tg(ubi::cAMPr)* animals at any of these time points (Figure 3H).

Previous studies show that cAMP regulates muscle cell specification. In zebrafish, secretion of Sonic Hedgehog (Shh) from the notochord instructs adjacent adaxial cells to differentiate into muscle pioneers, slow muscle fibers, or medial fast fibers. The patterning of somitic cells depends in part on the level and timing of cAMP activity, which inhibits the response to Shh.

Analysis of doubly transgenic embryos expressing both *Tg(ubi::DDcAMP)* and the Shh reporter *Tg(glim:mCherry-NLS)* revealed that N41-GFP signal appeared in Shh-responsive muscle pioneers (mp) and the Shh reporter *Tg(glim:mCherry-NLS)* was expressed in developing muscle (Figure S4B). Imaging indicated that DDCAMP accurately reports on the reduced levels of endogenous cAMP produced by Gpr161 in developing muscle (Figure 4B,C). Imaging indicated that DDCAMP is detectable in the nucleus and cytoplasm, whereas Gpr161 is most active in cilia. In an effort to quantify cAMP levels in cilia, a cilium-localizing peptide was appended to the N-terminus of DDCAMP (Arl13b-DDDCAMP). The cilia-localized Arl13b-DDDCAMP displayed similar levels of GFP signal to the unliganded DDcAMP protein. ciliary DDcAMP signal was similar in wild-type and gpr161 mutants (Figure 4D,E). These results indicate that the sensor does not specifically detect Gpr161-dependent cAMP in the cillum, perhaps because ciliary localization impedes efficient degradation of the unliganded DDCAMP protein.

There is evidence that the ciliary GPCR Gpr161 negatively regulates Shh signaling by increasing cAMP concentration in the cilium, and *gpr161b* is expressed in developing adaxial cells of zebrafish embryos. To determine if the DDCAMP sensor detects cAMP generated by Gpr161 in developing muscle cells, we used CRISPR-Cas9 to generate presumed null mutations in the duplicated genes encoding the Gpr161a and Gpr161b proteins (*gpr161a* and *gpr161b*, Figure S4C). In accordance with previous work, the number of Shh responsive cells in the somites was increased in the double mutant for *gpr161ad1129129* and maternal zygotic MZgpr161b*128/128*, hereafter referred to as “gpr161 mutants” (Figure 4A, magenta). In gpr161 mutants, the *Tg(ubi::DDcAMP)* GFP signal in somites was reduced, providing evidence that DDCAMP accurately reports on the reduced levels of endogenous cAMP produced by Gpr161 in developing muscle (Figure 4B,C). Imaging indicated that DDCAMP is detectable in the nucleus and cytoplasm, whereas Gpr161 is most active in cilia. In zebrafish, we used ligand-dependent protein stabilization to develop protein-based sensors for secondary metabolites such as progesterone in yeast and digoxin in plants. Given that existing cAMP sensors are not suitable for imaging in living zebrafish, we used ligand-dependent protein stabilization to generate a genetically encoded single wavelength sensor that allows long-term imaging of endogenous cAMP in zebrafish embryos. We identified DDCAMP as a protein sensor that is...
Figure 3. (A) Diagram of DMSO and 20 μM FSK treatment on Tg(ubi:DDcAMP) and Tg(ubi:DDcAMP<sup>307Q</sup>) embryos beginning at 4−5 hpf for 20 h. Anterior somites from treated embryos were imaged at 24 hpf, and GFP intensity was measured in somites 7−11. (B) Images of somites from DMSO and FSK-treated Tg(ubi:DDcAMP) and Tg(ubi:DDcAMP<sup>307Q</sup>) embryos at 24 hpf. The graph represents the mean fluorescence intensity of five somites per embryo, and each point corresponds to one embryo. Error bars indicate SD; ***p < 0.001 by two-way ANOVA (Sidak’s multiple comparisons), n = 6−11 animals for each condition. ns = not significant. AU = arbitrary unit. (C) Diagram of DMSO and 8-cpt-cAMP treatment of Tg(ubi:DDcAMP) starting at 12−13 somite stage (12−13ss) and imaged after 24 h of incubation. The tip of the tail was imaged for GFP intensity and the EVLs were quantified. (D) Tg(ubi:DDcAMP) embryos from the same clutch were treated with DMSO, 100 μM 8-cpt-cAMP, or 100 μM 8-cpt-cGMP; signal from 20 EVL cells was averaged per animal. Each dot in the graph represents one animal. Error bars indicate SD; ****p < 0.0001 one-way ANOVA (with Bonferroni’s multiple comparisons), n = 6−9 animal per condition. AU = arbitrary unit. (E) Frames from a confocal time lapse image with Airyscan 2 processing of Tg(ubi:DDcAMP) embryo starting at 15 hpf (t = 0 min). One frame every 20 min is shown as representation of the time lapse. GFP-expressing muscle cells are indicated with red arrows. (F) Graph represents the fold change of mean fluorescence intensity measured in five somites per embryo at time 0 and after 2 and 4 h of treatment. Error bars indicate SEM; **p < 0.01 two-way ANOVA (with Sidak’s multiple comparisons), n = 6−10 animals for each condition. ns = not significant, FC = fold change. (G) Confocal images of somites from Tg(ubi:DDcAMP) and Tg(ubi:cubePr) embryos after 4 h of FSK and DMSO treatment show GFP expression in muscle cells but different response to FSK treatment. (H) Graph represents the mean fluorescence intensity measured in five somites per embryo at the time points indicated. Error bars indicate SEM; there is no significance among the groups; n = 4−5 animals for each condition.
Specifically stabilized by cAMP, and we created transgenic fish that ubiquitously express DDcAMP. DDcAMP specifically responds to the production of cAMP in developing muscle cells. Our analyses suggest that DDcAMP turnover is regulated by cAMP concentration and that this sensor produces a strong GFP signal specifically in response to cAMP. These features enable in vivo time lapse imaging without phototoxicity and bleaching; however, DDcAMP is relatively slow to respond to changes in cAMP levels. There is a 3-fold increase in DDcAMP GFP intensity after 4 h of exposure to 20 μM FSK. This time lag presumably reflects the time required for newly translated DDcAMP to be stabilized by cAMP. Thus, DDcAMP is suitable for imaging endogenous cAMP levels in living zebrafish embryos, but it may not be ideal for monitoring changes in cAMP levels that occur over rapid time scales or in subcellular compartments with restricted access to the ubiquitin-proteasome system. DDcAMP has complementary features to other recently developed cAMP sensors, and it will enable new approaches to analyze cAMP in different experimental settings, cell types, and model organisms.

ASSOCIATED CONTENT

Supporting Information
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Cloning and genotyping primers and sgRNA sequences (XLSX)
Model organisms and strains, recombinant DNA resources, and software and algorithms (XLSX)
Confocal time lapse image of Tg(ubi:DDcAMP) embryo starting at 15 hpf (t = 0). The embryo is oriented dorsally, with the anterior developing head on top. Developing adaxial cells gradually increase GFP intensity in the somites. The images were taken every 8 minutes for over 7 hours of time lapse (MP4)
Experimental information on CNBD structure, transgene, and pharmacological and genetic validation, details on the experimental procedures for the CNBD selection, transgene generation, and image acquisition (PDF)

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Figure 4. (A) Confocal images of cAMP (green) and Shh reporter (magenta) signals in control fish and double mutants for gpr161a (st129 allele) and MZgpr161b (st128 allele). MZ = maternal zygotic. Control fish include gpr161a/+; gpr161b/+, and gpr161a+/+; gpr161b+/. (B) Graphs show GFP fluorescence intensity measured in the somite area corresponding to the muscle pioneers in Tg(gli:mCherry-NLS); Tg(ubi:DDcAMP) (control) and in Tg(gli:mCherry-NLS); Tg(ubi:DDcAMP); gpr161a+/+; MZgpr161b+/+ (gpr161 mut) embryos. Error bars indicate SD, ****p < 0.0001 by t-test. (C) Mean of the pixel values in each image acquired in panel A plotted as a function of their frequency. gpr161 double mutant fish display darker values compared to the controls. n = 4–5 animals per genotype. (D) Magnification of somites from 24 hpf Tg(gli:mCherry-NLS) control and Tg(gli:mCherry-NLS); gpr161a+/+; MZgpr161b+/+ (gpr161 mut) embryos. (E) Mean of the pixel values in each image acquired in panel A plotted as a function of their frequency. gpr161 double mutant fish display darker values compared to the controls. n = 4–5 animals per genotype. (D) Magnification of somites from 24 hpf Tg(gli:mCherry-NLS) control and Tg(gli:mCherry-NLS); gpr161a+/+; MZgpr161b+/+ (gpr161 mut) embryos.
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Notes

The authors declare no competing financial interest.

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