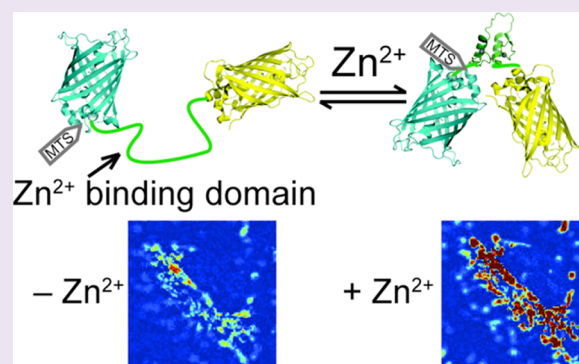


New Sensors for Quantitative Measurement of Mitochondrial Zn²⁺J. Genevieve Park,[†] Yan Qin,[†] Domenico F. Galati,[‡] and Amy E. Palmer^{*,†}[†]Department of Chemistry and Biochemistry and BioFrontiers Institute and [‡]Department of Molecular, Cellular, and Developmental Biology, University of Colorado Boulder, Boulder, Colorado 80309, United States

S Supporting Information

ABSTRACT: Zinc (Zn²⁺) homeostasis plays a vital role in cell function, and the dysregulation of intracellular Zn²⁺ is associated with mitochondrial dysfunction. Few tools exist to quantitatively monitor the buffered, free Zn²⁺ concentration in mitochondria of living cells ([Zn²⁺]_{mito}). We have validated three high dynamic range, ratiometric, genetically encoded, fluorescent Zn²⁺ sensors that we have successfully used to precisely measure and monitor [Zn²⁺]_{mito} in several cell types. Using one of these sensors, called mito-ZapCY1, we report observations that free Zn²⁺ is buffered at concentrations about 3 orders of magnitude lower in mitochondria than in the cytosol and that HeLa cells expressing mito-ZapCY1 have an average [Zn²⁺]_{mito} of 0.14 pM, which differs significantly from other cell types. These optimized mitochondrial Zn²⁺ sensors could improve our understanding of the relationship between Zn²⁺ homeostasis and mitochondrial function.



Zn²⁺ is a micronutrient that is required for human life, and deficiency can lead to impaired cognition, immune dysfunction, diarrhea, and death, particularly in children under the age of 5 years.¹ Although Zn²⁺ is essential for cell function, accumulation of Zn²⁺ to toxic levels leads to cell death. The human genome encodes two dozen Zn²⁺-specific transporters and many metal-buffering proteins, which are expressed in a tissue-specific manner.² Our current understanding of Zn²⁺ homeostasis is that intracellular Zn²⁺ is distributed into a large pool of structural or catalytic Zn²⁺ that is tightly bound, and two smaller pools of Zn²⁺: free Zn²⁺ and exchangeable Zn²⁺ bound loosely to small molecule or protein partners.^{3,4} Zn²⁺ homeostasis can be altered in diseases, such as neurodegeneration.⁵ In order to effectively study Zn²⁺ biology, we must be able to observe and manipulate Zn²⁺ specifically with subcellular resolution.

Cellular Zn²⁺ homeostasis affects mitochondrial function through poorly understood mechanisms. Zn²⁺ toxicity can lead to the release of cytochrome *c* from the intermembrane space, caspase activation, and apoptosis.^{6–8} Changes in Zn²⁺ availability can affect metabolism, including oxidative phosphorylation.^{9,10} Intracellular Zn²⁺ can depolarize mitochondria and decrease mitochondrial movement.^{11–13} Lastly, it is likely that mitochondria are a source and sink of Zn²⁺ in neurons and other cells.^{3,14–16} To understand how Zn²⁺ homeostasis affects mitochondrial function, we must be able to measure and monitor mitochondrial Zn²⁺.

Few tools exist to observe mitochondrial Zn²⁺ homeostasis in living cells with high specificity. Small molecule fluorescent probes are arguably the most popular tools. FluoZin-3 increases in intensity upon binding Zn²⁺ and has been used to observe free Zn²⁺ in isolated mitochondria.^{17,18} Positively charged

probes, such as RhodZin-3, concentrate within mitochondria of intact cells due to the negative mitochondrial inner membrane potential ($\Delta\psi_m$).^{17,19} Consequently, it is problematic to monitor Zn²⁺ in depolarized mitochondria using such probes because a decrease in fluorescence intensity could be caused by either a decrease in Zn²⁺ or in $\Delta\psi_m$. Other mitochondrial sensors consist of both a small molecule fluorophore and a protein component, which can be genetically targeted to mitochondria. This approach was used to target the fluorescent Zn²⁺ probe Zinpyr1 to mitochondria²⁰ and to exclusively express an excitation ratiometric Zn²⁺ sensor derived from carbonic anhydrase in mitochondria.²¹ Other genetically encoded Zn²⁺ sensors, such as the eCALWY family, have not yet been targeted to mitochondria.²² Our group previously constructed mito-ZifCY1 (renamed from mito-Cys₂His₂), a genetically encoded, ratiometric, Zn²⁺-specific sensor targeted to mitochondria, but its measurements are limited by its small dynamic range.¹⁴

In this study, we show that increasing the dynamic range of genetically encoded Zn²⁺ sensors improves their precision. We make novel comparisons of the [Zn²⁺]_{mito} of different cell types using improved sensors. We find [Zn²⁺]_{mito} is about 3 orders of magnitude lower than the cytosolic free Zn²⁺ concentration and varies considerably among different cell types.

Sensor Design and Validation. The Zn²⁺ sensors constructed in this study are variants of previously published cytosolic ZapCY1 and ZifCY1 sensors (Figure 1, panel a; Supplementary Figure 1, Tables 1 and 2), which respond

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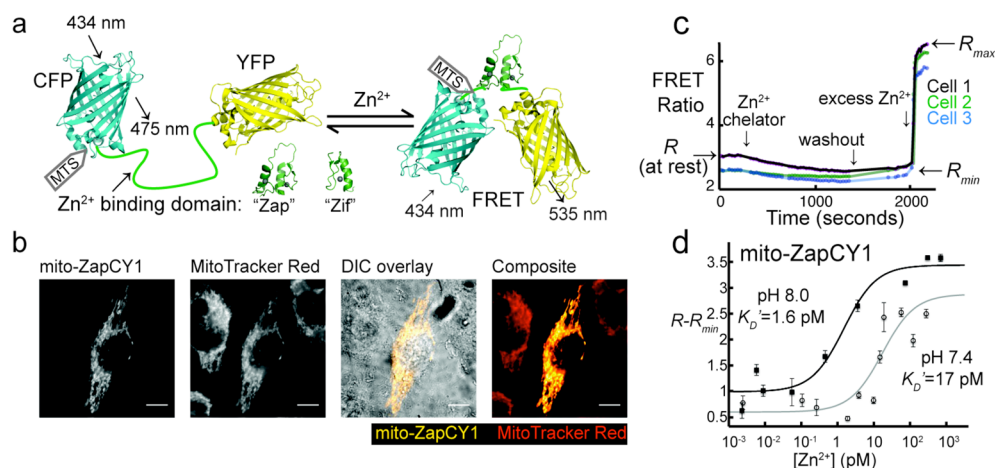


Figure 1. Design of genetically encoded mitochondrial Zn^{2+} sensors mito-ZapCY1 and mito-ZifCY1. (a) These sensors undergo a conformational change upon binding Zn^{2+} , which leads to a change in FRET. The Zn^{2+} -binding domain used in the “Zap” and “Zif” sensors consists of the first two Zn^{2+} fingers of the *Saccharomyces cerevisiae* protein Zap1 and the Zn^{2+} finger from the mammalian protein Zif268, respectively. The mitochondria targeting sequence (MTS) is appended to the N-terminus of the sensor. (b) mito-ZapCY1 co-localizes with MitoTracker Red in living HeLa cells; Pearson’s coefficient 0.938; scale bars represent 10 μm . (c) The fractional saturation of the sensor mito-ZapCY1 was determined in HeLa cells by measuring R (at rest), R_{\min} , and R_{\max} . (d) The *in situ* K_D' of mito-ZapCY1 was determined in HeLa cells at pH 7.4 and 8.0. Each point represents the average $(R - R_{\min})$ of at least 3 cells in a single experiment at a specific free $[\text{Zn}^{2+}]$.

specifically to Zn^{2+} over other biologically relevant divalent cations, including calcium, magnesium, iron, and copper.^{14,23} The conformational change upon Zn^{2+} binding changes the Förster resonance energy transfer (FRET) efficiency, and thus the sensor’s fluorescence emission reports the proportion of bound sensor. We report the magnitude of FRET as the FRET ratio (R), which is the fluorescence intensity of the acceptor fluorescence protein (FP) divided by that of the donor FP when only the donor FP is excited. Estimation of free Zn^{2+} is possible when the sensor’s affinity for Zn^{2+} is known and an *in vivo* sensor calibration is performed. Importantly, these sensors cannot estimate total Zn^{2+} .

Sensors were targeted to the mitochondrial matrix by appending an N-terminal mitochondrial targeting sequence and display excellent co-localization with MitoTracker Red in HeLa cells (Figure 1, panel b) and other cells tested (Supplementary Figure 2).

In order to identify an improved mitochondrial Zn^{2+} sensor, several new sensors were constructed with different Zn^{2+} binding domains. Details of sensor design are included in Supplementary Methods. Specifically, we sought to identify sensors that have an appropriate affinity (K_D') for Zn^{2+} allowing detection of both decreases and increases in Zn^{2+} concentration and with an improved dynamic range (defined as the maximum R (R_{\max}) divided by the minimum R (R_{\min})). In most cases, the Zn^{2+} -saturated sensor reports R_{\max} and Zn^{2+} -free sensor reports R_{\min} , but some display inverted responses.²² Measurement of R_{\max} and R_{\min} was achieved by performing *in situ* calibrations of single HeLa cells expressing each sensor (Figure 1, panel c). In a typical *in situ* calibration, R is measured in living cells treated with 150 μM of the Zn^{2+} chelator *N,N,N',N'*-tetrakis-(2-pyridylmethyl)-ethylenediamine (TPEN), followed by 0.75 μM pyrithione (a Zn^{2+} ionophore) and 10 μM ZnCl_2 . These calibrations are performed on single cells because the fractional saturation ($(R - R_{\min})/(R_{\max} - R_{\min})$) varies less than R_{\min} and R_{\max} from cell to cell (Supplementary Table 4). We also hoped to identify partially saturated sensors, which are best for quantitative measurements. Supplementary Table 3 reports the dynamic range and

fractional saturation of all the sensors tested and reveals mito-ZapCY1 as a robust sensor with a resting fractional saturation of $8.7 \pm 5.8\%$ and the current largest dynamic range of 3.2, significantly better than the previously reported mito-ZifCY1 sensor,¹⁴ which has a dynamic range of 1.2 (Supplementary Figure 3).

Estimates of mitochondrial free Zn^{2+} in HeLa cells based on mito-ZapCY1 dramatically differ from those based on mito-ZifCY1. Using mito-ZifCY1 (*in vitro* $\text{ZifCY1 } K_D' = 1.7 \pm 0.2 \mu\text{M}$), we previously observed a fractional saturation of $\sim 41\%$, identical to what we observe in the present study (Supplementary Table 3) leading to an estimate of $[\text{Zn}^{2+}]_{\text{mito}}$ of $680 \pm 140 \text{ nM}$ in HeLa cells. In contrast, measurements using mito-ZapCY1 (*in vitro* $\text{ZapCY1 } K_D' = 2.53 \text{ pM}$, pH 7.4) estimate a $[\text{Zn}^{2+}]_{\text{mito}}$ of 0.22 pM in HeLa cells. We hypothesized that the poor dynamic range of mito-ZifCY1 (Supplementary Figure 3) results in unreliable estimates of $[\text{Zn}^{2+}]_{\text{mito}}$ and set out to improve its dynamic range. Circular permutation of FPs, which involves relocating the N- and C-termini to different loops, has been shown to impact the dynamic range of FRET sensors by changing the orientation of the two FPs (Figure 2, panel a).^{24–26} We screened 5 variants of circularly permuted Venus (cpV), each permuted at a different location, within the framework of mito-ZifCY1. Figure 2 presents the dynamic range and fractional saturation of these sensors, and Supplementary Figure 4 presents representative calibration traces. The 5 variants yielded significantly different fractional saturation and dynamic ranges, resulting in two sensors with increased dynamic ranges, which were named mito-ZifCV1.49 and mito-ZifCV1.173, where the “49” denotes that Venus was circularly permuted at amino acid position 49. The high dynamic ranges of mito-ZifCV1.49 and mito-ZifCV1.173, which display inverted responses to Zn^{2+} , result from increased FRET in the unbound state and little change in FRET in the Zn^{2+} -bound state. We found that the fractional saturation decreases as the dynamic range increases, even though the Zn^{2+} binding domains are identical. It has been demonstrated that incorporation of cpV can decrease the K_D' 2- to 10-fold,²⁵ but even variation of the K_D' cannot explain the strong correlation

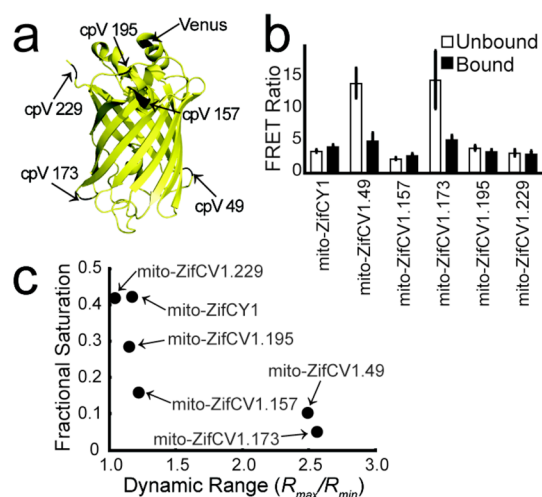


Figure 2. Circular permutation of mito-ZifCY1 dramatically increases its dynamic range. (a) The N- and C-termini of cpV FPs are relocated to 5 different loops of the original Venus FP at the amino acid positions 49, 157, 173, 195, and 229. (b) The average R of the unbound and bound sensors expressed in at least 10 cells from 2 or more independent experiments, acquired using identical exposure times, are summarized. (c) High dynamic range sensors report lower fractional saturation.

between the fractional saturation and dynamic range, suggesting that as hypothesized, measurements made using low dynamic range sensors can be inconsistent.

Converting the fractional saturation to $[Zn^{2+}]_{mito}$ requires estimation of the K_D' . The molecular environment of mitochondrial matrix differs from cytosol, and the pH and redox balance can change significantly under different conditions. The pH in the mitochondrial matrix is typically ~ 8.0 without perturbation but can vary from about 6.5 to 8.5 under different conditions.²⁷ Using Mito-pHRed,²⁷ we estimated the mitochondrial pH in HeLa cells to be ~ 8.0 . Therefore, we calibrated mito-ZapCY1 in mitochondria of living cells by adding Zn^{2+} buffered at different free concentrations to Ca^{2+} , Mg^{2+} , and phosphate-free imaging media in the presence of $50 \mu g mL^{-1}$ alamethicin. Alamethicin is an antimicrobial peptide that can permeabilize the mitochondrial inner membrane to small molecules only.²⁸ Using this method, we found mito-ZapCY1 has an *in situ* K_D' of 1.6 pM and 17 pM at pH 8.0 and 7.4, respectively (Figure 1, panel d). These *in situ* affinities are comparable to the affinities measured *in vitro* at different pH (Supplementary Figure 5). We estimate $[Zn^{2+}]_{mito}$ to be 0.14 pM in HeLa cells based on this *in situ* titration of mito-ZapCY1 at pH 8.0.

Comparison of Mitochondrial to Cytosolic Zn^{2+} . Next, we confirmed that two high-dynamic-range sensors with different Zn^{2+} binding domains yield consistent estimates of $[Zn^{2+}]_{mito}$. As shown in Figure 3, the fractional saturations of mito-ZapCY1 and mito-ZifCV1.173 in HeLa cells were $16 \pm 10\%$ and $5.8 \pm 3.1\%$, respectively, while the corresponding sensors in the cytosol were almost completely saturated ($90 \pm 7.3\%$ and $95 \pm 1.1\%$, respectively). These data indicate that under resting conditions in HeLa cells, the free Zn^{2+} is buffered at concentrations about 3 orders of magnitude lower in mitochondria than in the cytosol.

If, in fact, $[Zn^{2+}]_{mito}$ is buffered at a lower concentration than free cytosolic Zn^{2+} , the addition of a Zn^{2+} ionophore, in the absence of extracellular Zn^{2+} , should dissipate the Zn^{2+} gradient

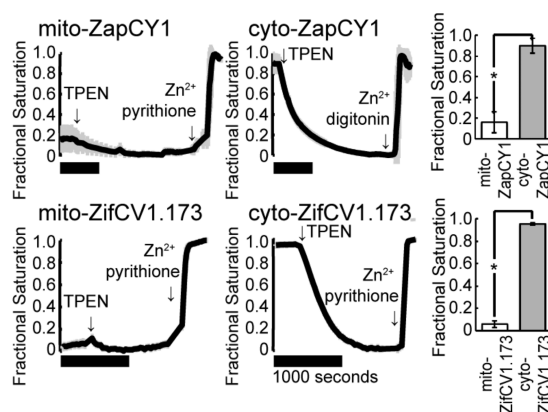


Figure 3. The fractional saturation of Zn^{2+} sensors is lower in the mitochondria than in the cytosol. Representative calibrations of ZapCY1 and ZifCV1.173 expressed in the mitochondrial matrix or the cytosol of HeLa cells are shown. Comparison of the fractional saturation of each sensor illustrates the difference in mitochondrial and cytosolic Zn^{2+} ($*p < 0.0001$, Student's t test). Horizontal black bars represent 1000 s. At least 3 cells were measured in each experiment.

between these two subcellular compartments. Indeed, treatment of several cell types expressing mito-ZapCY1 with $5 \mu M$ pyrithione caused a rapid increase in mitochondrial Zn^{2+} , which was reversed by the addition of $150 \mu M$ TPEN (Supplementary Figure 6). Our interpretation of these results is that pyrithione moves Zn^{2+} into mitochondria from other subcellular compartments, such as the cytosol, which buffer free Zn^{2+} at higher concentrations.

Comparison of $[Zn^{2+}]_{mito}$ in Different Cell Types. We used mito-ZapCY1 to quantitatively compare $[Zn^{2+}]_{mito}$ in different cell types and different environmental conditions. Mito-ZapCY1 was expressed in HeLa cells, MIN6 cells (a mouse insulinoma cell line), primary cortical neurons, and HC11 cells (a mouse mammary epithelial cell line). Resting R , R_{min} , and R_{max} were measured in individual cells to determine the fractional saturation of mito-ZapCY1 in each of the different cell types. Figure 4 presents a summary of mito-ZapCY1's fractional saturation in different cell types and pseudocolor images of cells at rest, upon TPEN treatment, and upon addition of Zn^{2+} /pyrithione. Significant differences among cell types were observed, including lower fractional saturation of mito-ZapCY1 in HeLa cells ($8.7 \pm 5.8\%$) than in MIN6 cells ($41 \pm 18\%$) or neurons ($59 \pm 20\%$). We also compared the fractional saturation of mito-ZapCY1 in HC11 cells grown in basal media ($-prolactin$) to those in lactogenic media ($+prolactin$) and found that $[Zn^{2+}]_{mito}$ is significantly higher in non-lactogenic HC11 cells than in lactogenic HC11 cells ($p = 0.0022$, Student's t test). This is particularly intriguing given that HC11 cells have been shown to undergo massive redistribution of intracellular Zn^{2+} pools and alterations in Zn^{2+} transporter expression upon lactogenic stimulation.^{29,30} These results suggest that $[Zn^{2+}]_{mito}$ is regulated differently in several cell types and under different environmental conditions. While at this point we do not know how cells maintain different levels of mitochondrial Zn^{2+} , we speculate that they will likely exhibit different levels of Zn^{2+} transporters, buffers, and other regulatory proteins that are necessary to ensure appropriate Zn^{2+} management for each specialized cell.

In summary, we constructed and validated three genetically encoded, high dynamic range mitochondrial Zn^{2+} sensors. Although sensors with low dynamic range are capable of

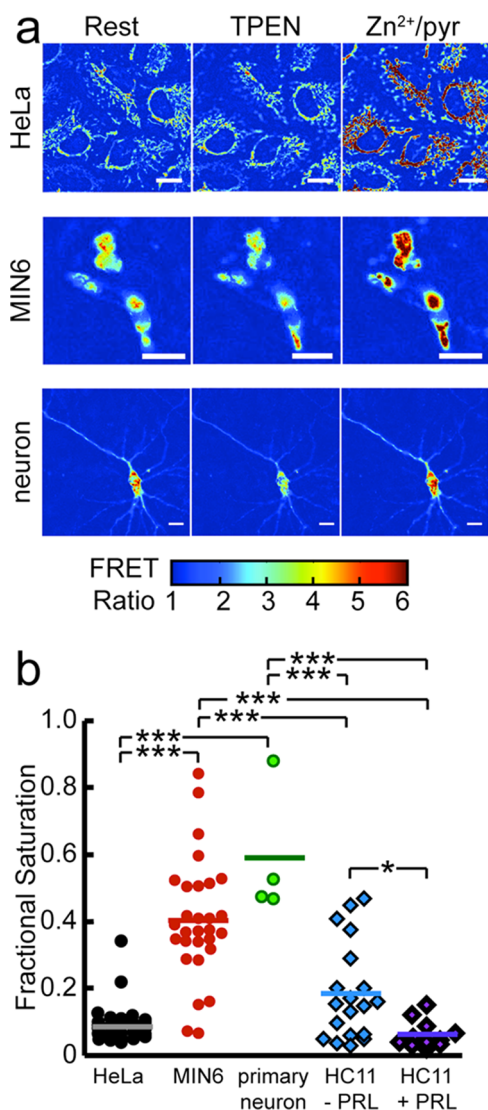


Figure 4. Quantitative comparison of $[\text{Zn}^{2+}]_{\text{mito}}$ in different cell types. (a) Pseudocolor images of the FRET ratio of mito-ZapCY1 in HeLa cells, MIN6 cells, and a primary cortical neuron illustrate changes in the FRET ratio in response to treatment with $150 \mu\text{M}$ TPEN or with $0.75 \mu\text{M}$ pyrithione and $10 \mu\text{M}$ ZnCl_2 . Scale bars represent $10 \mu\text{m}$. (b) $[\text{Zn}^{2+}]_{\text{mito}}$ differs significantly among cell types. Each marker shows the fractional saturation of mito-ZapCY1 in a single cell ($*p < 0.0022$, Student's *t* test; $***p < 0.0001$, ANOVA, Tukey's HSD posthoc test).

detecting relative changes in Zn^{2+} , high dynamic range sensors are necessary for making consistent and quantitative comparisons of Zn^{2+} between different cell types. Using mito-ZapCY1, we estimate $[\text{Zn}^{2+}]_{\text{mito}}$ to be 0.14 pM in HeLa cells. A recent study reports a similar $[\text{Zn}^{2+}]_{\text{mito}}$ of 0.2 pM measured with a different ratiometric Zn^{2+} biosensor in the PC12 rat pheochromocytoma line.²¹ We believe that these mitochondrial Zn^{2+} sensors can be used to address the complex interplay between Zn^{2+} homeostasis and mitochondrial function.

Although FRET-based genetically encoded sensors are capable of making quantitative measurements in intact cells, it is poorly understood if and how Zn^{2+} sensor expression changes the total Zn^{2+} concentration and the Zn^{2+} -buffering capacity of the cell. The fractional saturation of mito-ZapCY1 does not change as the sensor expression increases (Supplementary Figure 7), suggesting that in contrast to small molecule

sensors,^{4,31} these sensors do not deplete the mitochondrial Zn^{2+} pool. The lack of perturbation of resting Zn^{2+} also suggests that, similar to the cytosol, mitochondrial Zn^{2+} is buffered and the sensor concentration is much lower than the concentration of the buffer.

A potential limitation of the Zn^{2+} -finger binding domains of the mito-ZapCY1 and mito-ZifCV1 sensors is their sensitivity to oxidation and changes in pH. However, we observe little perturbation due to acidification by acetic acid or treatment with H_2O_2 . We performed parallel experiments with the sensors mito-pHRed and mito-RoGFP2 and observed little change in pH or oxidation in the process of sensor calibration (Supplementary Figure 8).^{27,32}

In conclusion, we created improved mitochondrial Zn^{2+} sensors to measure the buffered set point of free Zn^{2+} with superior accuracy and precision. These new sensors complement and expand our fluorescent toolbox for studying Zn^{2+} and other ions in complex, biological systems.

METHODS

Sensor Construction. Four repeats of the coding sequence for the first 29 amino acids of the human cytochrome *c* oxidase subunit 8a (mitochondrial precursor; accession number NP_004065) precede the coding sequence of each Zn^{2+} sensor in the mammalian expression vector pcDNA3.1, as previously described.¹⁴ Citrine FP in "CY" sensors was replaced with cpV FPs to make "CV" sensors using 5' *SacI* and 3' *EcoRI* restriction sites. ZapCY1 was expressed in *E. coli* and purified for use in *in vitro* titrations as previously described.²³ Please refer to Supplemental Methods for further details.

Cell Culture. HeLa cells were maintained in high glucose Dulbecco's Modified Eagle Medium (DMEM) with 10% fetal bovine serum (FBS) and penicillin/streptomycin (pen/strep). MIN6 cells were cultured in D-MEM supplemented with 10% (v/v) FBS, L-glutamine, sodium pyruvate, β -mercaptoethanol, and penicillin/streptomycin. HC11 cells were maintained, as previously published, in non-lactogenic medium, which was replaced with lactogenic medium 24–48 h prior to imaging.³⁰ All cells were transfected 48–72 h before imaging with Mirus TransIT-LT or electroporated the Neon system (Life Technologies). Primary cortical neurons were obtained, cultured, and transfected as described in Supplemental Methods. Transfected cells were stained with MitoTracker Red (Life Technologies) for co-localization studies. During all experiments except *in situ* titrations, cells were imaged in phosphate-free HEPES-buffered Hanks Balanced Salt Solution (HHBSS), pH 7.4.

mito-ZapCY1 Titrations. Zn^{2+} solutions used for the titration of mito-ZapCY1 were buffered using Zn^{2+} chelators (EGTA, EDTA, and HEDTA) using a previously published method²³ with the following modifications. Solutions for both *in vitro* and *in situ* titrations were adjusted to the specified pH, and the free $[\text{Zn}^{2+}]$ in each Zn^{2+} /chelator buffered solution was calculated for different pH. One HeLa cell imaging experiment was performed at each Zn^{2+} concentration and the average *R* of ≥ 3 cells was used to calculate K_D' . Cells were imaged in phosphate-free HHBSS, pH 7.4 throughout Zn^{2+} chelation with $150 \mu\text{M}$ TPEN, which was then replaced with phosphate-, Ca^{2+} -, and Mg^{2+} -free HHBSS, pH 7.4 or pH 8.0 with $125 \mu\text{M}$ dithiothreitol (to prevent sensor oxidation) and Zn^{2+} /chelator to buffer the free $[\text{Zn}^{2+}]$. Cells and mitochondria were permeabilized using $50 \mu\text{g mL}^{-1}$ alamethicin. All chemicals were purchased from Sigma.

Imaging. Image acquisition and analysis were performed as previously published.²³ Microscope filter combinations for FRET and CFP: 430/24 nm excitation filter, 455 nm dichroic mirror, 535/25 nm and 470/24 nm emission filters, respectively. Images were analyzed using ImageJ and Matlab software.

Statistical Analysis. Statistical analysis was performed using the Student's *t* test or ANOVA with Tukey's HSD posthoc test in the KaleidaGraph program.

■ ASSOCIATED CONTENT**■ Supporting Information**

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Author Contributions

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Notes

The authors declare no competing financial interest.

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