

# Loss of Pde6 reduces cell body $\text{Ca}^{2+}$ transients within photoreceptors

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Modulation of  $\text{Ca}^{2+}$  within cells is tightly regulated through complex and dynamic interactions between the plasma membrane and internal compartments. In this study, we exploit *in vivo* imaging strategies based on genetically encoded  $\text{Ca}^{2+}$  indicators to define changes in perikaryal  $\text{Ca}^{2+}$  concentration of intact photoreceptors. We developed double-transgenic zebrafish larvae expressing GCaMP3 in all cones and tdTomato in long-wavelength cones to test the hypothesis that photoreceptor degeneration induced by mutations in the phosphodiesterase-6 (Pde6) gene is driven by excessive  $[\text{Ca}^{2+}]_i$  levels within the cell body. Arguing against  $\text{Ca}^{2+}$  overload in Pde6 mutant photoreceptors, simultaneous analysis of cone photoreceptor morphology and  $\text{Ca}^{2+}$  fluxes revealed that degeneration of *pde6c*<sup>w59</sup> mutant cones, which lack the cone-specific cGMP phosphodiesterase, is not associated with sustained increases in perikaryal  $[\text{Ca}^{2+}]_i$ . Analysis of  $[\text{Ca}^{2+}]_i$  in dissociated *Pde6β*<sup>rd1</sup> mouse rods shows conservation of this finding across vertebrates. *In vivo*, transient and Pde6-independent  $\text{Ca}^{2+}$  elevations ('flashes') were detected throughout the inner segment and the synapse. As the mutant cells proceeded to degenerate, these  $\text{Ca}^{2+}$  fluxes diminished. This study thus provides insight into  $\text{Ca}^{2+}$  dynamics in a common form of inherited blindness and uncovers a dramatic, light-independent modulation of  $[\text{Ca}^{2+}]_i$  that occurs in normal cones.

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Photoreceptor degeneration is a devastating disease that affects millions of people worldwide. Mutations that alter cyclic guanosine-mono-phosphate (cGMP) levels are a common cause of photoreceptor degeneration; mutations in guanylate cyclase, guanylate cyclase-activating proteins, phosphodiesterase-6 (Pde6) and the Pde6 chaperone, aryl hydrocarbon receptor interacting protein-like 1, all cause loss of rods and/or cones (Retinal Information Network, <https://sph.uth.edu/retnet/home.htm>). The precise mechanism of cGMP-mediated cell death is poorly defined, and thus animal models are particularly relevant to study as a tool to help cure and treat this disease.

The most studied animal model for photoreceptor degeneration due to elevated cGMP is the *rd1* mouse, identified more than 90 years ago.<sup>1,2</sup> *rd1* mice have a mutation in the beta subunit of rod cGMP Pde6, a key mediator of the light response.<sup>3,4</sup> Normally, light activates a molecular cascade that causes Pde6 to hydrolyze cGMP, resulting in closure of cation channels within the photoreceptor outer segment (OS). In the absence of Pde6 beta in *rd1* mice, cGMP levels remain elevated and rod photoreceptors start to degenerate after their differentiation and before eye opening.<sup>3</sup> Cone photoreceptors lacking Pde6c also have elevated cGMP in their OSs.<sup>5</sup> Elevated cGMP is proposed to result in sustained increases in OS  $\text{Ca}^{2+}$ , triggering death. Consistently, elevated OS  $\text{Ca}^{2+}$  was measured in hypomorphic Pde6

mutant rods<sup>6</sup> and rod viability was recently shown to improve in *Cngb1*<sup>(-/-)</sup> × *Pde6β*<sup>rd1</sup> double-mutant mice.<sup>7</sup> However, the degeneration processes take place within the perikaryon, not the OS. Thus, a key challenge has been to determine the signaling pathways through which elevated cGMP and  $\text{Ca}^{2+}$  within OSs of Pde6 mutant cells affect photoreceptor cell biology downstream from the OS.

Although events associated with non-apoptotic cell death pathways such as poly-ADP-ribose-polymerase and calpain activation occur in rods dying due to elevated cGMP,<sup>8,9</sup> the molecular cascade activated in degenerating Pde6 mutant photoreceptors is unknown. Specifically, involvement of  $\text{Ca}^{2+}$  in the *rd1* pathology is a major unsolved question.<sup>10</sup> It is often assumed that elevated  $\text{Ca}^{2+}$  within OSs triggers cell death by activating  $\text{Ca}^{2+}$ -dependent mechanisms within the cell body.<sup>11</sup> However, cGMP and  $\text{Ca}^{2+}$  diffusion from the OS to the inner segment are likely to be limited by anatomical constraints (the ciliary bottleneck), buffering and the ellipsoid mitochondrial barrier.

To ascertain the role of  $\text{Ca}^{2+}$  in photoreceptor degeneration caused by loss of Pde6, it is necessary to measure this ion within the inner segment. We therefore generated here a double-transgenic zebrafish, transgenic (*Tg; Trβ2:tdTomato; TαCP:GCaMP3*), in which the  $\text{Ca}^{2+}$  indicator GCaMP3 is selectively expressed in the photoreceptor cell body, not the OS. We recorded  $\text{Ca}^{2+}$  dynamics within intact cones from

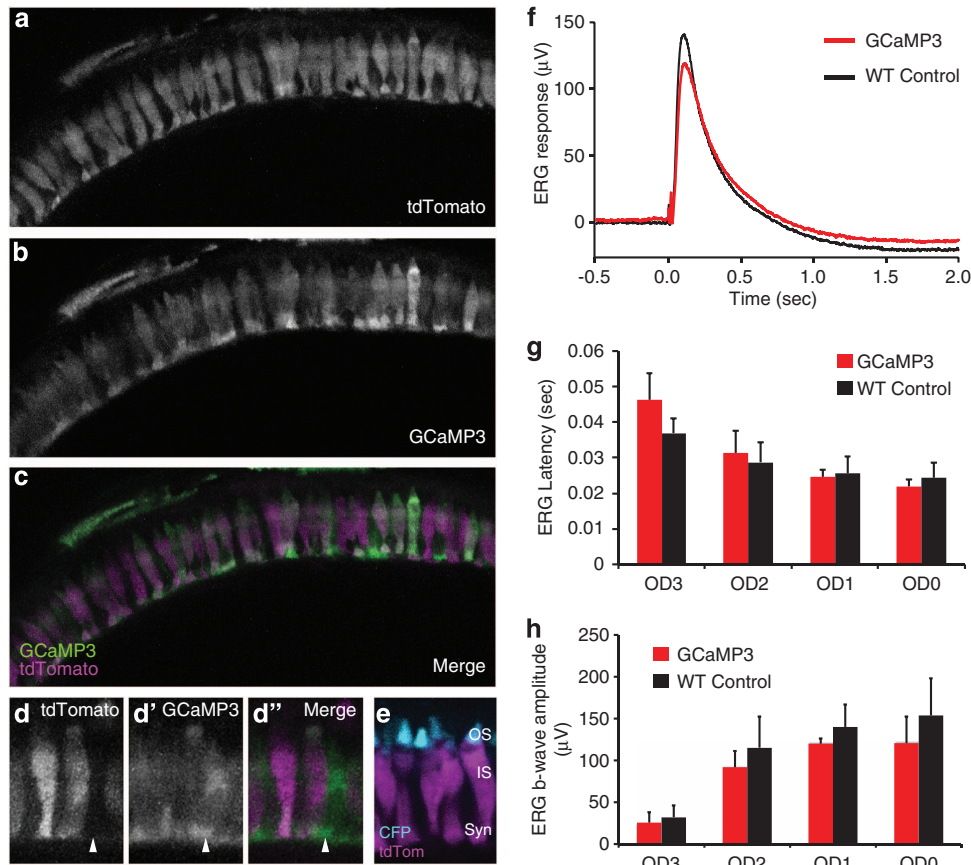
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**Abbreviations:** cGMP, cyclic guanosine-mono-phosphate; Pde6, phosphodiesterase-6; OS, outer segment; d.p.f., days post fertilization; WT, wild type; RP, retinitis pigmentosa; PTU, 1-phenyl 2-thiourea; ERG, electroretinogram; *TαCP*, cone  $\alpha$ -transducin promoter;  $[\text{Ca}^{2+}]_i$ , intracellular calcium; ROI, region of interest; Tg, transgenic; *Trβ2*, thyroid hormone receptor  $\beta$ 2; GFP, green fluorescent protein; mCFP, membrane cyan fluorescent protein

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**Figure 1** GCaMP3 is expressed in all cone photoreceptors and does not interfere with light responses. (a–c) Retinal slice projections from a representative 5 d.p.f. *Tg(Trl32:tdTomato;T $\alpha$ CP:GCaMP3)* double-transgenic *pde6c<sup>w59</sup>* mutant zebrafish larvae displaying intact photoreceptors prior to their degeneration. The tdTomato (a) is expressed only in long-wavelength cones, whereas the GCaMP3 (b) is expressed in all cone photoreceptors. Panel c shows both tdTomato (magenta) and GCaMP3 (green) channels merged. (d) Magnified view of cone photoreceptors in the *Tg(Trl32:tdTomato;T $\alpha$ CP:GCaMP3)* double transgenic. Arrow indicates a cell expressing GCaMP3 (green) but not tdTomato (magenta). Less expression of both GCaMP3 and tdTomato is observed in the cone outer segments (OS), whose location within the photoreceptor layer is clearly defined by the *T $\alpha$ CP:membrane CFP* (membrane cyan fluorescent protein (mCFP); cyan) transgene (e) Retinal section from a *Tg(Trl32:tdTomato; T $\alpha$ CP:mCFP)* double-transgenic fish expressing tdTomato in long-wavelength cones (magenta) and membrane CFP (cyan) in all cones. The outer segments (OS), inner segments (IS), and synapses (syn) are as indicated. (f–h) ERG waveforms (f), b-wave latencies (g), and b-wave amplitudes (h) for 5 d.p.f. WT non-transgenic versus *Tg(T $\alpha$ CP:GCaMP3)* larvae show no significant differences in their light responses over three orders of magnitude.  $n = 3$  fish per condition; bars = S.D.

wild type (WT) and mutant (*pde6c<sup>w59</sup>*) retinas using multi-photon time-lapse imaging. *pde6c<sup>w59</sup>* cones degenerate starting immediately after their differentiation at 4 days post fertilization (d.p.f.), and are mostly lost by 7 d.p.f.<sup>12,13</sup> This mutation is thus analogous to the well-characterized rod *Pde6 $\beta$ <sup>rd1</sup>* mutation.<sup>3,4</sup> To confirm our results in the mammalian model, we extended our approach to record intracellular  $Ca^{2+}$  ( $[Ca^{2+}]_i$ ) levels in *rd1* mouse rods.

By combining the power of zebrafish genetics with live, noninvasive imaging, we were able to follow  $Ca^{2+}$  signaling within degenerating vertebrate cones in real-time. Surprisingly, we did not detect sustained  $[Ca^{2+}]_i$  increases in *pde6c<sup>w59</sup>* mutant cones nor were abnormalities in baseline  $[Ca^{2+}]_i$  observed in degenerating *rd1* mouse rods.  $Ca^{2+}$  signals in non-mutant and *pde6c<sup>w59</sup>* mutant zebrafish cones include recurring transient  $[Ca^{2+}]_i$  increases ('flashes'), consistent with complex regulation of the  $Ca^{2+}$  homeostatic apparatus within the photoreceptor regions downstream from the OS. Our results thus challenge the prevailing view that *rd1* degeneration is driven by global  $Ca^{2+}$  elevations that

overload the endogenous  $Ca^{2+}$  buffering and clearance mechanisms within the cell body, and thus have implications for the development of neuroprotection strategies in retinitis pigmentosa (RP) models of retinal degeneration.

## Results

We constructed a transgenic line expressing GCaMP3 specifically in zebrafish cones using the cone transducin promoter (*T $\alpha$ CP*) that we previously isolated.<sup>14</sup> GCaMP3 is a fluorescent  $Ca^{2+}$  indicator consisting of circularly permuted green fluorescent protein (GFP), calmodulin, and the  $Ca^{2+}$ -calmodulin target peptide M13.<sup>15</sup> GCaMP3 has increased fold fluorescence change (increased  $F_{max}/F_{min}$ ) and higher  $Ca^{2+}$  affinity ( $660 \pm 19$  nm) than many other genetically encoded indicators.<sup>15</sup> It is widely used as an indicator of *in vivo* changes in intracellular  $Ca^{2+}$ .<sup>16–18</sup> In zebrafish cones, GCaMP3 localizes throughout the cone cytoplasm, but is less abundant in the OS (Figures 1b–e). We then generated double transgenics by crossing the *Tg(T $\alpha$ CP:GCaMP3)* strain

with a transgenic fish line expressing the fluorescent protein tdTomato selectively in long-wavelength cone photoreceptors using the thyroid hormone receptor  $\beta 2$  promoter, *Tg(Tr $\beta$ 2:tdTomato)* (Williams et al.<sup>19</sup> and Figures 1a and c–e). The tdTomato fluorescence in long-wavelength cones has several uses. Uniform cytoplasmic distribution of tdTomato facilitates visualization of the overall cell shape so that morphological changes during cell death can be analyzed. Further, its expression in a subset of discretely spaced cones defines these cells for visual analysis, and serves as a photo-bleaching control. Double-transgenic fish (*Tg(Tr $\beta$ 2:tdTomato; T $\alpha$ CP:GCaMP3)*) were used for live imaging experiments, where we directed our observation of Ca<sup>2+</sup> dynamics to the cone photoreceptor cell body and synaptic terminal.

**ERG analysis of transgenic GCaMP3 cones.** A potential problem when expressing fluorescent indicators of intracellular Ca<sup>2+</sup> is the disruption of Ca<sup>2+</sup> homeostasis due to abnormal buffering. To ensure that GCaMP3 expressed in cones does not alter normal fluxes in Ca<sup>2+</sup>, we measured electroretinogram (ERG) responses from our transgenic line and compared these with WT ERG responses. ERGs were recorded from excised eyes as previously described.<sup>20</sup> If introduction of excess buffer affected the photoreceptor [Ca<sup>2+</sup>]<sub>i</sub>, the delay and/or reduction in synaptic signaling would be detected as a diminished and/or delayed ERG b-wave. We found that the ERG waveforms appeared similar, and no significant differences in b-wave amplitude or latency between transgenic and non-transgenic WT cones were observed (Figures 1f–h). These results indicate that the GCaMP3 transgene does not significantly alter the visual responses of photoreceptors.

**Ca<sup>2+</sup> dynamics in dying zebrafish cones.** Our previous characterization of the *pde6c*<sup>w59</sup> mutant used multiphoton imaging to analyze morphological changes associated with cone photoreceptor cell death that occur *in vivo*.<sup>12</sup> Mutant cells differentiate normally and appear morphologically normal, with small OSs and extended synaptic processes, until ~4 d.p.f. The mutant cells then proceed through a series of morphological changes prior to their clearance from the retina. Typically, the synapse is first retracted, followed by the contraction of the OS and the rounding of the cell body. These morphological changes occur asynchronously across the retina over 2–3 days; by 7 d.p.f. the vast majority of cones are gone in the mutant retina. In an individual cell, morphological changes occur over ~8 h (Lewis et al.<sup>12</sup> and Figure 2b). Based on recent studies in the *rd1* mouse, morphological changes associated with cell death in *pde6c*<sup>w59</sup> cones are likely late in the cell death cascade.<sup>21</sup>

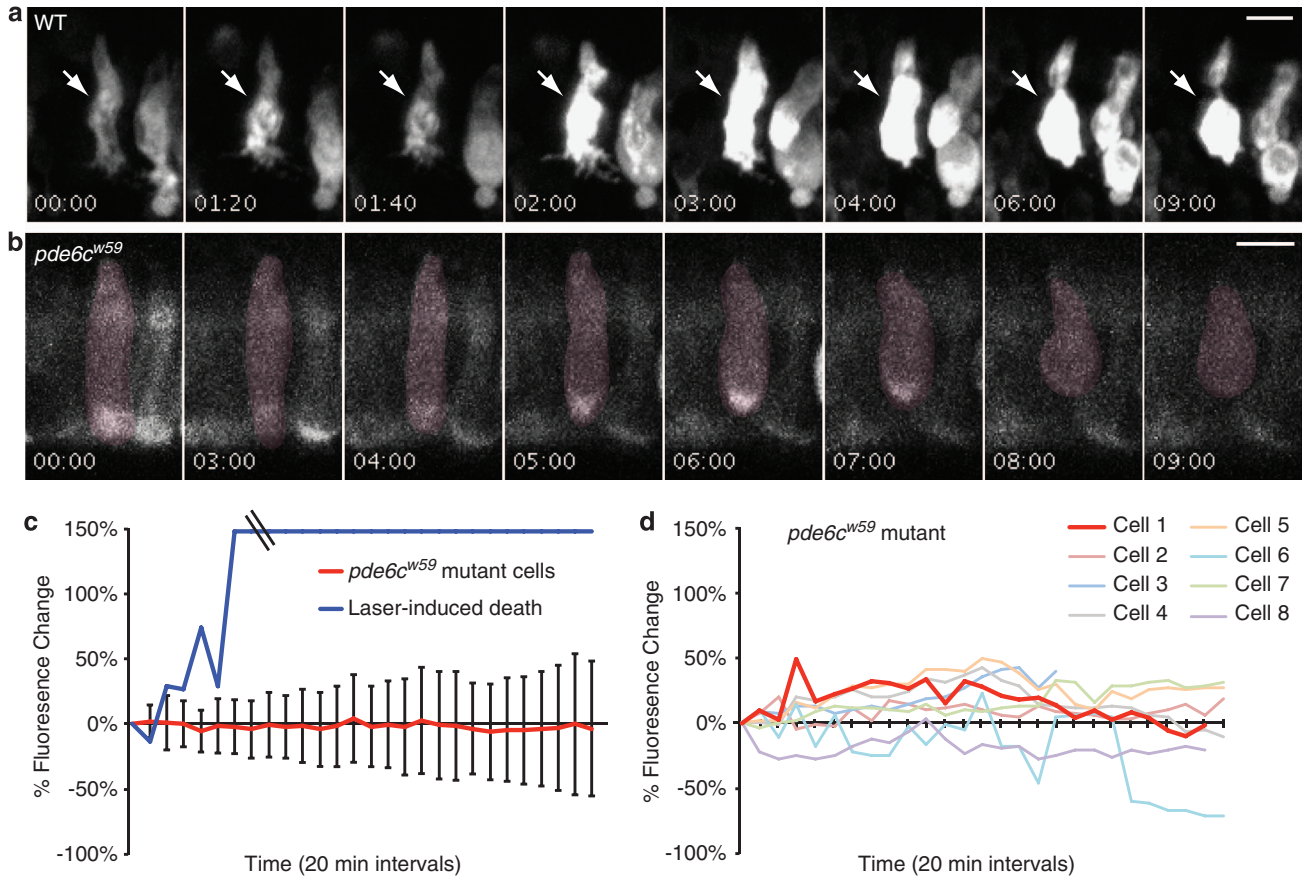
To examine Ca<sup>2+</sup> dynamics during cell death, we monitored both changes in GCaMP3 fluorescence and in cell shape at 20 min intervals in time-lapse experiments that extended up to 9 h. We then quantified changes in GCaMP3 fluorescence in dying cells that initially appeared morphologically normal at the beginning of the time-lapse, but had completely rounded up by the end of the experiment. As a proof of principle for this study, we initially caused death of non-mutant cells by intense infrared laser illumination via multiphoton time-lapse imaging. This method led to profound

increases in GCaMP3 fluorescence that saturated our detectors (Figures 2a and c). During the dying process of these laser-damaged cells, dramatic localized increases in GCaMP3 fluorescence typically around the nucleus were observed. These initial GCaMP3 changes were either transient or sustained, but were consistently followed by dramatic persistent increases that extended throughout the entire cell body as the photoreceptor rounded up (Figure 2a).

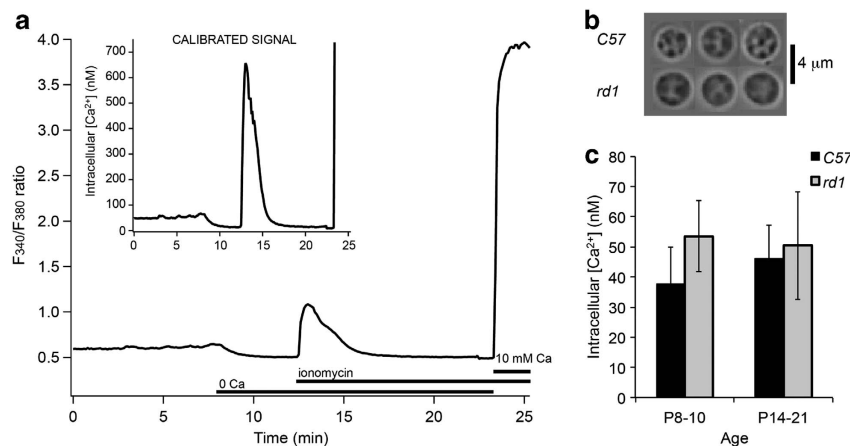
We used optimized multiphoton imaging conditions that do not damage homozygous *pde6c*<sup>w59</sup> cones<sup>12</sup> to monitor GCaMP3 fluorescence changes in mutant cells that degenerate during the 9 h time-lapse. In contrast to the dramatic changes observed with laser-induced death, we did not detect pronounced intracellular increases in GCaMP3 fluorescence in the *pde6c*<sup>w59</sup> mutant dying photoreceptors (Figure 2b). We quantified intracellular fluorescent changes in 19 individual cells from four different fish, and none of these cells showed sustained increases in GCaMP3 fluorescence over time (Figure 2c). Even though the average change in cell body fluorescence centered at zero for these 19 cells, the specific changes in individual cells were unique. Some cells showed very little change, some decreased in overall fluorescence, whereas others showed recurring small transient increases in GCaMP3 fluorescence (Figure 2d and Figure 4).

**Ca<sup>2+</sup> dynamics in dying mouse rods.** To determine whether our finding represented a general feature of RP-like photoreceptor degeneration across vertebrates, we extended our measurements to intracellular Ca<sup>2+</sup> in rod photoreceptors from *rd1* mice, arguably the most widely used model of Pde6 dysfunction.<sup>7–11</sup> Intracellular Ca<sup>2+</sup> was assessed in acutely isolated *rd1* rods and age-matched WT cells at the start of degeneration (postnatal day 8–10), and at the peak of degeneration after eye opening (postnatal day 14–21). The cells were identified by shape, size and/or expression of GFP (Figure 3b).<sup>22</sup> Figure 3a illustrates averaged data from 19 simultaneously recorded *rd1* rods loaded with the Ca<sup>2+</sup> indicator Fura-2. The calibration protocol shows baseline ratio values for dissociated light-adapted rods, followed by decreased ratio in Ca<sup>2+</sup> free saline (possibly mediated by the closure of store-operated channels)<sup>22</sup> and then ionomycin-induced Ca<sup>2+</sup> release from internal stores. The inset displays calibrated [Ca<sup>2+</sup>]<sub>i</sub> with stable baseline levels maintained at ~50 nM. [Ca<sup>2+</sup>]<sub>i</sub> levels in developing rods were comparable to concentrations measured in adult WT cells.<sup>22</sup> Consistent with *in vivo* measurements in intact teleost cones, baseline [Ca<sup>2+</sup>]<sub>i</sub> in dissociated *rd1* rod somata was not significantly elevated before or during cGMP-driven rod degeneration even at ages P14–21 during massive rod loss (Figure 3c).

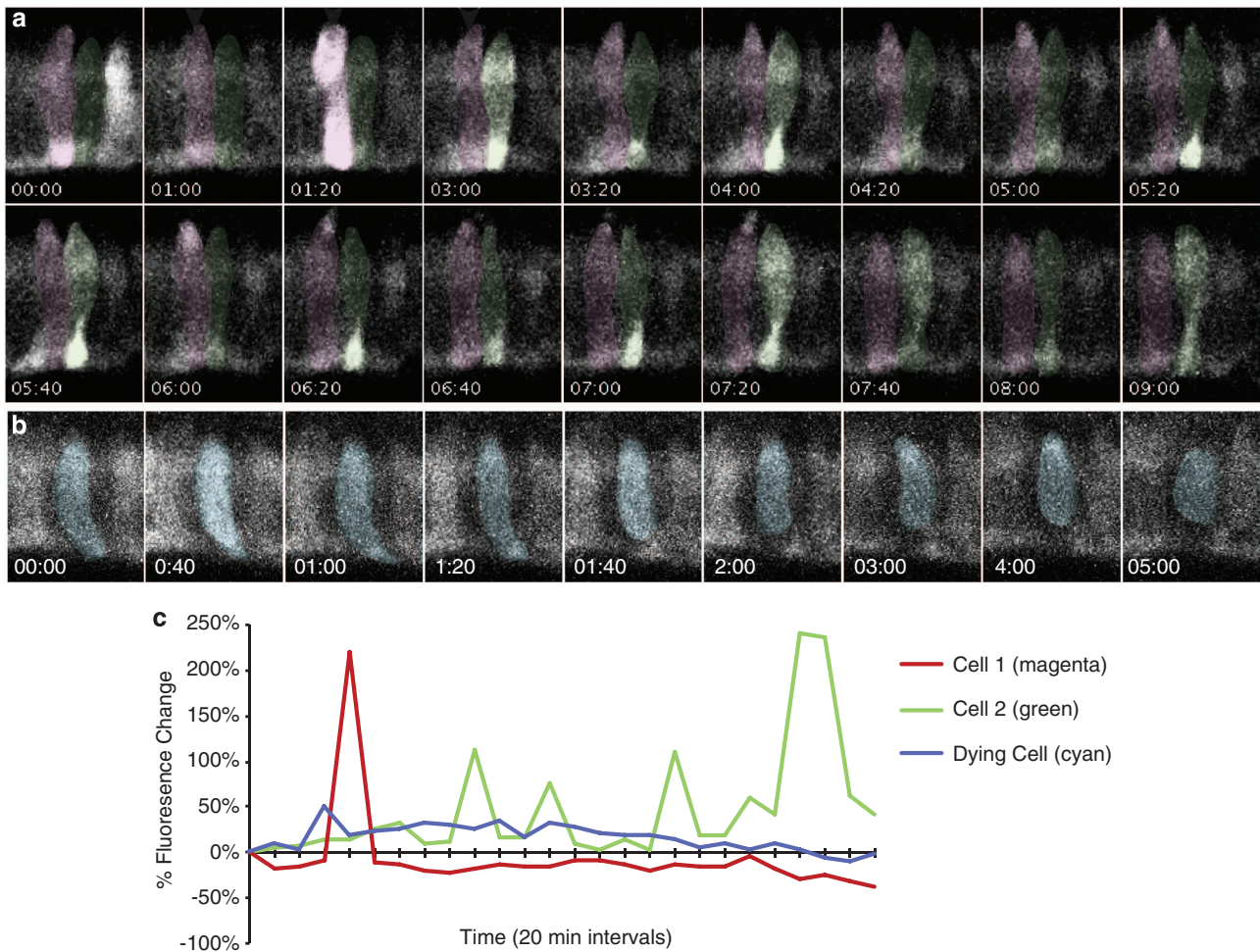
**Ca<sup>2+</sup> oscillations in *pde6c*<sup>w59</sup> zebrafish cone photoreceptors.** We next analyzed the Ca<sup>2+</sup> oscillations that we detected in zebrafish cones to determine how these related to the cell death process. We found that 27% of *pde6c*<sup>w59</sup> mutant cones ( $n=84$ ) exhibited oscillations in GCaMP3 fluorescence during our 9-hr time-lapse experiments. The intensity of oscillations varied from cell to cell and even within cells, thus we used a change in intensity of ~18% as a lower limit for our definition of an increase.



**Figure 2** Modest and variable changes in  $Ca^{2+}$  occur during the death of *pde6c<sup>w59</sup>* cones. Photoreceptors from 5–6 d.p.f. larvae expressing  $T\alpha CP:GCaMP3$  were imaged every 20 min for 9 h. (a, b) Montages of representative cells at select time points show GCaMP3 fluorescence and morphology changes in dying photoreceptors for both *pde6c<sup>w59</sup>* heterozygous (a) and *pde6c<sup>w59</sup>* homozygous (b) photoreceptors. Time = (h:min). (c) Graph showing mean percentage of fluorescence change of GCaMP3 from 19 *pde6c<sup>w59</sup>* mutant cells undergoing cGMP-mediated death (bars = S.D.) and the single representative laser-induced dying cell shown in a. Laser power and time of exposure was increased to stimulate laser-induced death of non-mutant cones. In addition, these cells were more sensitive than the homozygous *pde6c<sup>w59</sup>* cones to prolonged laser exposure. For homozygous *pde6c<sup>w59</sup>* mutants, laser conditions were optimized to not cause morphological defects during the extended time-lapses. (d) Graph showing variations in percentage of GCaMP3 fluorescence changes in eight individual cones from a single *pde6c<sup>w59</sup>* mutant fish. The red line (cell 1) indicates the percentage of change in GCaMP3 fluorescence of the representative dying *pde6c<sup>w59</sup>* mutant cone shown in b



**Figure 3**  $Ca^{2+}$  measurements in dying mouse rods. (a) Average ( $n = 19$ ) fluorescence ratio and calibrated signal during baseline calcium measurements from acutely isolated *Pde6 $\beta$ <sup>rd1</sup>* (*rd1*) mouse rod perikarya at P10. Ionomycin was used to clamp intracellular calcium levels to external zero calcium (containing 1 mM EGTA to obtain Rmin) or 10 mM calcium level (to obtain Rmax), allowing for a cell-by-cell calibration of the fluorescent signal. (b) Typical morphology of *C57Bl6J* (*C57*) or *rd1* mouse rod photoreceptors. (c) No significant differences in absolute baseline  $Ca^{2+}$  concentration of acutely isolated *C57Bl6J* (*C57*) or *Pde6 $\beta$ <sup>rd1</sup>* (*rd1*) mouse rod perikarya at ages between 8 and 10 days and 14 and 21 days were observed. Shown are group averages  $\pm$  S.E.M. p8–10:  $37.7 \pm 12.2$  (*C57*),  $53.5 \pm 11.8$  (*rd1*); p14–21:  $46.2 \pm 11.0$  (*C57*),  $50.4 \pm 17.8$  (*rd1*)



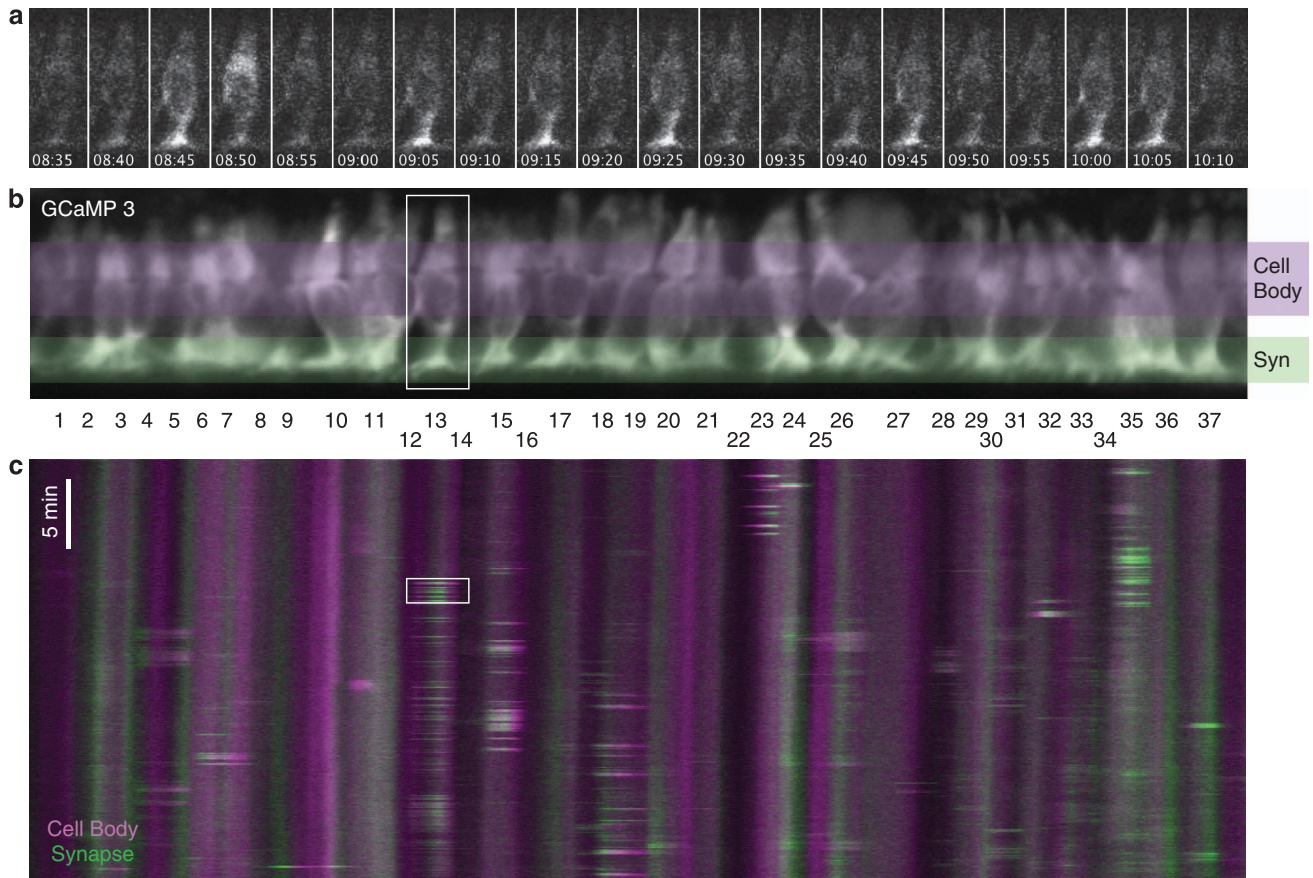
**Figure 4** Ca<sup>2+</sup> oscillations occur in morphologically normal and dying *pde6c<sup>w59</sup>* cones. Photoreceptors from homozygous *pde6c<sup>w59</sup>* double-transgenic animals *Tg(Tri32:tdTomato;TαCP:GCaMP3)* were imaged every 20 min. Only the GCaMP3 channel is shown. (a, b) Montages of representative morphologically normal (a) and degenerating (b) *pde6c<sup>w59</sup>* cones show transient increases in GCaMP3 fluorescence. Time = (h:min). (c) Graph of percentage change in GCaMP3 fluorescence intensity for outlined cells shown in a and b. Oscillations in GCaMP3 fluorescence occur at random intervals throughout the time-lapse in morphologically normal cones, whereas transient increases in dying cells usually occur near the beginning of the time-lapses (c and Figure 2d)

GCaMP3 increases occurred both simultaneously throughout the cell body and synapse, or were often localized only to the synaptic region (Figure 4a). Both morphologically normal cells (Figures 4a and c) and those undergoing morphological changes (Figures 4b and c) exhibited Ca<sup>2+</sup> oscillations. For the dying cells that eventually rounded up, the most significant increase(s) in [Ca<sup>2+</sup>]<sub>i</sub> occurred at the beginning of the time-lapse before any major cell shape changes were observed (Figures 4b and c).

To analyze the temporal and spatial characteristics of these recurring transient GCaMP3 increases in greater detail, we conducted shorter time-lapse experiments (ranging from 15–30 min) and acquired images every 5 s. Image acquisition every 5 s allowed sampling of a single retinal plane containing ~30–50 cells. As the overall experiment was short, we focused our analysis on the *pde6c<sup>w59</sup>* mutant cone cells that appeared morphologically normal throughout the experiment. A representative experiment and subsequent data analysis is shown in Figure 5. Using this rapid imaging paradigm, rapid recurring Ca<sup>2+</sup> increases (‘flashes’) were observed in many

cells within this imaged population (see Supplemental Movie). For example, in the representative cell shown in Figure 5a, a bright increase is observed in the synapse at 0905 hours, but is not seen in the time point before (0900 hours) nor after (0910 hours). Another synaptic increase is seen in the same cell at 0915 hours, is gone at 0920 hours, and reoccurs at 0925 hours. This analysis indicates that these particular flashes lasted for 5 s or less. To optimize our analysis of multiple *pde6c<sup>w59</sup>* photoreceptor cells in a single retinal population, we straightened our retinal images (Figure 5b) and generated kymographs (Figure 5c). Two kymographs were generated for each time-lapse; one for the synapse and one for the cell body. These kymographs are overlaid in two different colors in Figure 5c. Green shows GCaMP3 fluorescence increases localized to the synapse regions and magenta shows Ca<sup>2+</sup> changes throughout the cell body (Figure 5c).

Using our kymographs, we graphed the GCaMP3 signal intensity over time for each photoreceptor and established general categories describing the types of flashes we



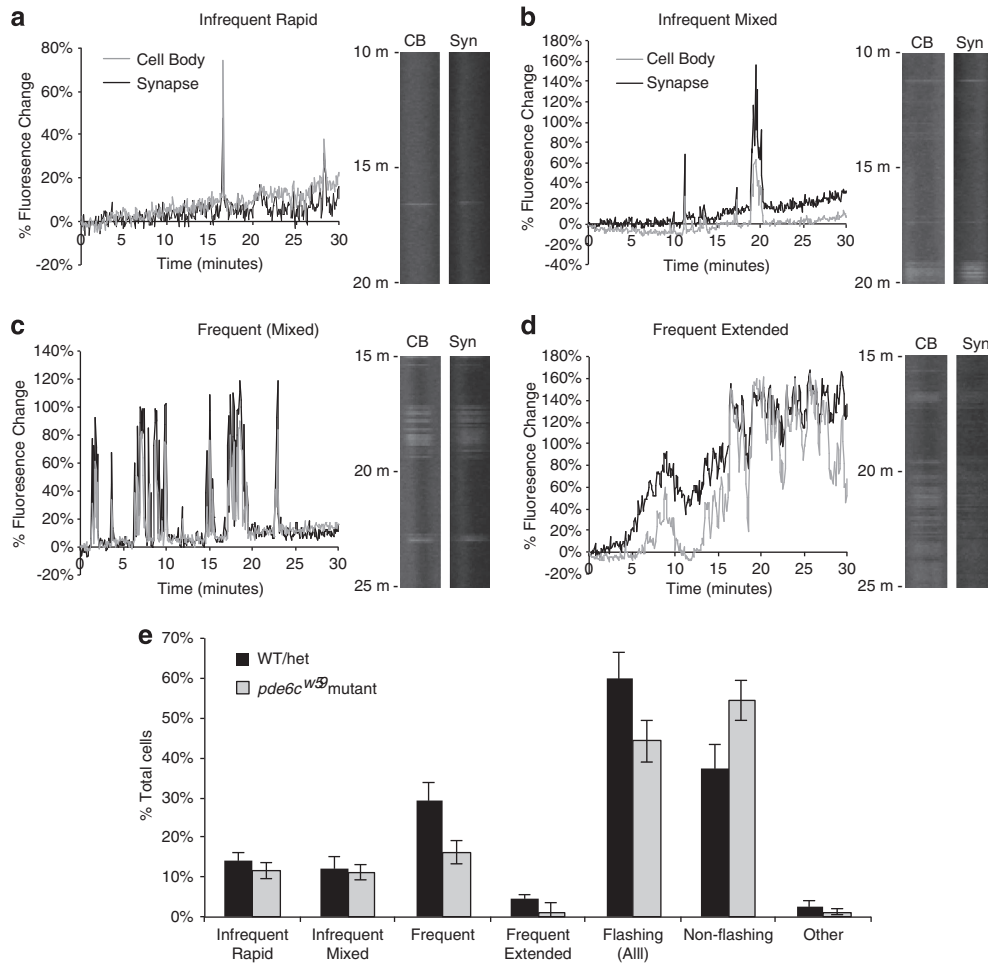
**Figure 5** Recurrent  $\text{Ca}^{2+}$  flashes are variable, rapid, asynchronous, and prevalent. Photoreceptors from homozygous double-transgenic *Tg(Trb2:tdTomato);T $\alpha$ :CP:GCaMP3* *pde6c*<sup>w59</sup> fish were rapidly imaged every 5 s for 15–30 min. Only the GCaMP3 channel is shown. (a) Montage of a representative cell (cell 13; see b and c) at select time points show rapid transient increases in GCaMP3 fluorescence occurring in both the cell body and synapse. Time = (min:s). (b) Representative image (time) projection from a straightened *Tg(Trb2:tdTomato);T $\alpha$ :CP:GCaMP3* *pde6c*<sup>w59</sup> retina shows GCaMP3 fluorescence in all cones. The shaded areas represent the areas used to generate the kymograph (c) for the cell body (magenta) and for the synapse (syn; green). All identifiable cells are numbered below and correspond to a column on the kymograph in c. (c) Representative kymograph of GCaMP3 signal across all the time points for both the cell body (magenta) and synapse (green). The boxed areas on both the kymograph (c) and the straightened retina (b) represent the cell (cell 13) and time period shown in the montage in a

detected. Cells that flashed fewer than 10 times over 30 min were labeled as ‘infrequent,’ whereas those flashing more than 10 times were categorized as ‘frequent’. Cells that flashed infrequently were also subdivided based upon the duration of the flashes. Cells whose infrequent flashes lasted for only 1–3 time points were deemed as ‘infrequent rapid’, whereas ‘infrequent mixed’ cells exhibit flashes that were sustained over several time points (Figures 6a and b). Cells with frequent flashes exhibited transient GCaMP3 increases of mixed duration (Figure 6c), but a few cells showed recurring transient increases that lasted over an extended period of time (categorized as ‘frequent extended’), often with varying intensities within the elevated GCaMP3 signal (Figure 6d). A few cells also showed a change in fluorescence, either increasing or decreasing, that persisted throughout the time-lapse. This small number of cells was categorized in the ‘other’ category. From our analysis of 447 cells from 11 *pde6c*<sup>w59</sup> fish, we determined that 46% of cells flash during the 15–30 min time-lapse experiments, and that there were roughly equal number of cells that flash infrequently (both rapid and mixed) and those that flash frequently. Consistent with our longer

time-lapse experiments, extended increases in fluorescence were detected in only a few cells (Figure 6e).

**$\text{Ca}^{2+}$  dynamics in non-mutant cone photoreceptors.** As  $\text{Ca}^{2+}$  flashes within photoreceptors have not been reported previously, we speculated that elevated cGMP and consequent metabolic changes in *pde6c*<sup>w59</sup> cones disrupt  $\text{Ca}^{2+}$  homeostasis, causing transient increases in  $[\text{Ca}^{2+}]_i$ . To examine this possibility, we analyzed flashes in photoreceptors from both WT and *pde6c*<sup>w59</sup> heterozygous fish, siblings of the *pde6c*<sup>w59</sup> mutants previously analyzed, and imaged these photoreceptors *in vivo* at 5 s intervals for 15–30 min. We noted that the light-sensitive cones from WT and *pde6c*<sup>w59</sup> heterozygous fish were more sensitive to laser exposure than the light-insensitive homozygous *pde6c*<sup>w59</sup> photoreceptors. Slight laser-induced morphological abnormalities were detected in some photoreceptors during imaging, and data were not analyzed from these cells.

To our surprise, kymograph analysis of cone photoreceptors from WT and *pde6c*<sup>w59</sup> heterozygous fish ( $n = 263$  cells; 6 fish) revealed the same categories of flashes as observed in



**Figure 6** Characterization of heterogeneous Ca<sup>2+</sup> flashes show fewer *pde6c<sup>w59</sup>* mutant cones with frequent flash patterns. Individual cones from kymographs of the rapid 5 s time-lapse imaging experiments were analyzed and categorized according to their pattern of GCaMP3 flashes. (a–d) Representative graphs and kymograph segments of GCaMP3 flash patterns in both the cell body (CB) and synapse (syn) of cones. Flashing cells were categorized into four main categories: ‘infrequent rapid’ (a), ‘infrequent mixed’ (b), ‘frequent’ (c), and ‘frequent extended’ (d). See text for more details regarding flash categorization. (e) Graph showing the percentage of various categories of flashing cells in both *pde6c<sup>w59</sup>* mutant and non-mutant (WT/het) cones. The ‘other’ category consists of a small number of non-flashing cells that showed persistent increasing or decreasing GCaMP3 fluorescence throughout the duration of the time-lapse (e). Overall, fewer cells exhibiting Ca<sup>2+</sup> flashes were observed in *pde6c<sup>w59</sup>* mutants than in WT/het larval retinas (‘Flashing (All)’), largely due to a reduction in the ‘frequent’ category. No significant difference in mutant and non-mutant cones was observed in the other categories. For *pde6c<sup>w59</sup>* mutants: *n* = 447 cells; 11 fish. For WT/het: *n* = 263 cells; six fish. Bars = S.E.M.

the *pde6c<sup>w59</sup>* mutant photoreceptors. Further, our results indicated that altered metabolism due to elevated cGMP did not increase the number of cells flashing or the flash frequency of individual cells, but rather decreased these; 44% of mutant cells flashed *versus* 60% of non-mutant cones (*P* = 0.09), and this reduction in flashing *pde6c<sup>w59</sup>* mutant cones was primarily cells categorized as ‘frequent’ (16% mutant *versus* 29% non-mutant, *P* = 0.03, Figure 6e).

Taken together, these data show that dynamic modulation of [Ca<sup>2+</sup>]<sub>i</sub> occurs in WT cone photoreceptors and that this modulation is mostly retained in light-insensitive *pde6c<sup>w59</sup>* mutant cones prior to their degeneration, although fewer frequently flashing cells were observed. However, at late stages of death when mutant cells degenerate, Ca<sup>2+</sup> transients are substantially reduced or cease and catastrophic Ca<sup>2+</sup> release into the cytoplasm does not occur.

## Discussion

In this study, we report an *in vivo* analysis of Ca<sup>2+</sup> dynamics in cone photoreceptors within a host animal. We exploit a double-transgenic zebrafish that enables simultaneous analysis of cytoplasmic changes in Ca<sup>2+</sup> and of overall changes in cell morphology, and extend our analysis of Ca<sup>2+</sup> regulation in *pde6c<sup>w59</sup>* cones to the commonly studied *rd1* mouse model. We discovered that (1) *pde6c<sup>w59</sup>* mutant cone photoreceptors and *rd1* rods do not exhibit sustained intracellular Ca<sup>2+</sup> increases during degeneration; (2) transient Ca<sup>2+</sup> increases are prevalent in *pde6c<sup>w59</sup>* mutant and non-mutant cone photoreceptors; (3) these Ca<sup>2+</sup> flashes occur throughout the cell body and synapse, or are only localized to the synapse; (4) Ca<sup>2+</sup> transients vary in frequency and duration; and (5) fewer cells in the *pde6c<sup>w59</sup>* mutant retinas show a pattern of frequent recurring Ca<sup>2+</sup> flashes compared with non-mutant retinas. Our findings shed light on the mechanism of cell death occurring in *pde6c<sup>w59</sup>*

mutant cones and *rd1* rods, and highlight a potentially functionally important measure of normal cone photoreceptor physiology.

Pde6 is a key mediator of the well-established vertebrate phototransduction cascade. Activated by GTP-bound  $\alpha$ -transducin, it hydrolyzes cGMP, resulting in closure of cation channels in the OS plasma membrane. Loss of Pde6 activity increases OS  $Ca^{2+}$  levels<sup>6</sup> possibly causing cell death, a conjecture supported by the observation that reducing  $Ca^{2+}$  influx by knocking out cGMP-gated cation channels (*Cngb1*<sup>(-/-)</sup> × *Pde6 $\beta$ <sup>rd1</sup>*) improves viability of *rd1* rods.<sup>7</sup> However, to understand the physiological coupling between  $[Ca^{2+}]_i$  and the degeneration process, it is necessary to ascertain  $Ca^{2+}$  dynamics within the inner segment. We therefore developed a novel zebrafish transgenic line, *Tg(Tr $\beta$ 2:tdTomato;T $\alpha$ CP:GCaMP3)*, and compared  $Ca^{2+}$  signals obtained in degenerating *pde6c<sup>w59</sup>* cones with  $[Ca^{2+}]_i$  levels within the *Pde6 $\beta$ <sup>rd1</sup>* rods. To avoid interference from signals emanating from over-activation of  $Ca^{2+}$  fluxes through cGMP channels,<sup>7</sup> *pde6c<sup>w59</sup>* mutants were designed to express the genetically encoded  $Ca^{2+}$  indicator GCaMP3 within cell compartments downstream from the OS.

One possible mechanism for *pde6c<sup>w59</sup>* and *rd1* photoreceptor degeneration was mitochondrial  $Ca^{2+}$  overload and a consequent increase in cytosolic  $[Ca^{2+}]_i$  within the perikaryon.<sup>23</sup> Using live imaging, we detected modest changes in intracellular  $Ca^{2+}$  that varied between individual dying cells; some cells displayed decreases in overall GCaMP3 fluorescence and several displayed small  $Ca^{2+}$  oscillations. Nonetheless, sustained increases in  $Ca^{2+}$  in cell bodies of *pde6c<sup>w59</sup>* cones undergoing morphological changes of cell death were not detected during our *in vivo* time-lapse experiments nor were elevated perikaryal  $[Ca^{2+}]_i$  levels observed in *Pde6 $\beta$ <sup>rd1</sup>* mouse rods. This finding is in stark contrast to the pronounced  $[Ca^{2+}]_i$  increases elicited in non-mutant cones by excessive laser exposure (Figure 2). Hence, our observations argue against a major role of depolarization, voltage-activated  $Ca^{2+}$  influx, mitochondrial overload and  $Ca^{2+}$  release from internal stores in cone and rod death driven by Pde6 mutations. Consistent with this, elimination of  $Ca^{2+}$  influx through L-type channels slightly delayed, but did not prevent, *rd1* degeneration.<sup>24</sup>

We observed dramatic transient increases in  $[Ca^{2+}]_i$  in WT and in pre-degeneration *pde6c<sup>w59</sup>* cones. This finding indicates that  $Ca^{2+}$  homeostasis in photoreceptors is dynamically regulated and similar in light-adapted and unstimulated photoreceptors. Thus, the  $Ca^{2+}$  changes that we report appear independent of phototransduction. Although future studies will explore possible sources for such  $Ca^{2+}$  flashes, potential sources include voltage-gated  $Ca^{2+}$  channels, receptor-mediated and/or store-operated  $Ca^{2+}$  influx from the extracellular space channels, as well as  $Ca^{2+}$  release from internal stores gated by ryanodine/inositol triphosphate release  $Ca^{2+}$  channels.<sup>22</sup> Numerous studies focusing on non-excitabile and excitabile cells<sup>25</sup> suggest that internal compartments represent major contributors to spontaneous elevations in  $[Ca^{2+}]_i$ . ‘Sparks’ or ‘puffs’ have been proposed to regulate cardiac and skeletal muscle excitation–contraction coupling, vascular tone regulation, membrane excitability and neuronal secretion, but were also implicated in

regulation of developmental changes, plasticity or cell to cell communication.<sup>25</sup> Ryanodine-sensitive  $Ca^{2+}$  stores are present in photoreceptors and have been proposed to help maintain  $Ca^{2+}$  levels within a physiological range optimal for normal function under dim lights,<sup>26</sup> whereas store-operated calcium entry and transient receptor potential channels might have a more significant function under bright and prolonged illumination.<sup>22,27</sup>

In conclusion, we here document new features of  $Ca^{2+}$  signaling in cone photoreceptors and address the long-standing question whether Pde6 mutations trigger photoreceptor degeneration through  $Ca^{2+}$ -dependent activation of proapoptotic pathways driven by cGMP-induced  $Ca^{2+}$  overloads within the cell body. We found that, surprisingly, Pde6 loss does not result in sustained increases in  $[Ca^{2+}]_i$  in either mouse rods or zebrafish cones. Our data show that *in vivo* analysis of photoreceptor physiology represents a powerful new tool to study function in normal and diseased retinal tissue.

## Materials and Methods

**Zebrafish maintenance and mutant isolation.** Adult fish and larvae were maintained at 28.5 °C in reverse-osmosis distilled water reconstituted for fish compatibility by addition of salts and vitamins<sup>28</sup> on a 10/14 h dark/light cycle. Unless otherwise noted, all zebrafish strains are homozygous for the *roy orbison* (*roy*) mutation.<sup>29</sup> Fish used in live imaging experiments were maintained in embryo media<sup>28</sup> containing 0.003% 1-phenyl 2-thiourea (PTU) starting at 8–24 h post fertilization (h.p.f.). The *pde6c<sup>w59</sup>* mutant was originally isolated in a screen of mutagenized-zebrafish using the optokinetic response behavioral assay as described previously.<sup>13</sup> The *pde6c<sup>w59</sup>* mutation is maintained as heterozygotes and intercrossed to generate homozygous mutants.

**Construction of transgenic zebrafish.** To express GCaMP3 specifically in cones, we used the Gateway-based Tol2kit<sup>30</sup> and inserted GCaMP3 downstream of the 3.2Kb of the *T $\alpha$ CP*.<sup>14</sup> The resulting construct was injected into WT zebrafish embryos at the 1–2-cell stage together with Tol2 transposase mRNA.<sup>31</sup> Mosaics were identified at 4 d.p.f. and raised to adulthood. Germline carriers were identified in the F1 generation. To visualize cone morphology, we generated a transgenic line *Tg(Tr $\beta$ 2:tdTomato)* expressing tdTomato in the long-wavelength cones (this paper and Suzuki *et al.*, in preparation). The *Tr $\beta$ 2* promoter isolation, clone construction, and the general expression were described previously.<sup>19</sup> Germline carriers were identified from mosaic fish injected with this construct. Both *Tg(T $\alpha$ CP:GCaMP3)* and *Tg(Tr $\beta$ 2:tdTomato)* were crossed into the *pde6c<sup>w59</sup>* mutant lines. The transgenic line *Tg(T $\alpha$ CP:membrane CFP)* was described previously.<sup>12</sup>

**Pde6c<sup>w59</sup> genotyping.** After imaging, zebrafish larval genotypes were verified by PCR analysis followed by restriction enzyme digest, as previously described.<sup>28</sup> Primers were used to create a restriction site for BsaXI in only the mutant locus (forward: 5'-TTGGCCTCTGGAATACTGGCT-3'; reverse: 5'-GTTTGACCAGAACCCGGAAG-3'). PCR products were digested with BsaXI and genotyped according to its restriction profile.

**ERG recordings.** ERGs were recorded as described previously.<sup>32</sup> Briefly, 5 and 6 d.p.f. larvae were anesthetized in 0.02% tricaine, and eyes were removed using a fine tungsten wire loop. Excised eyes were then placed in an oxygenated Ringer's solution (in mM; 130 NaCl, 2.5 KCl, 20 NaHCO<sub>3</sub>, 0.7 CaCl<sub>2</sub>, 1.0 MgCl<sub>2</sub>, and 20 glucose), and a glass electrode was positioned directly onto the cornea. After 3 min (min) of dark adaptation, eyes were exposed to white light flashes, and their electrical responses were recorded. Data were acquired and processed as described previously.<sup>33</sup> Peak values are listed as the mean ± S.D. from three animals.

**Live time-lapse imaging in zebrafish.** For time-lapse imaging, transparent PTU-treated 5–6 d.p.f. larvae were embedded in a drop of 1% low melting point agarose in embryo media. After solidifying, the agarose was covered



with embryo media containing 0.003% PTU and 0.02% (w/v) tricaine. Live time-lapse imaging experiments were performed on an Olympus FV1000MPE multiphoton laser scanning microscope equipped with a  $\times 25$  (NA 1.0) long working distance water objective. Images of the *Tg(Trf2:tdTomato); T $\alpha$ :CP:GCaMP3* double transgenics were acquired using a red/green filter cube at 890 nm wavelength. For the extended time-lapse experiments, images stacks of 30–40  $\mu\text{m}$  (1  $\mu\text{m}$  per slice) were taken every 20 min for 9 h. Single image slices were taken for the shorter time-lapse experiments, which require rapid sampling every 5 s for 15–30 min.

**Image processing and analysis.** Image processing and analysis was carried out using the ImageJ (Fiji; <http://fiji.sc/Fiji>) software.<sup>34</sup> Dying *pde6c<sup>W59</sup>* cells were identified by examining images collected from the extended 9-hour time-lapse experiments. Images were Z-projected using 7–10 stacks at 1  $\mu\text{m}$  depth, for a total of 7–10  $\mu\text{m}$  depth depending on the shape of the cell. The Z-projected images were adjusted to correct for xy drift using the 'stack reg' function, and further corrected for bleaching using the EMBL tools 'bleach correction' function with the tdTomato image selected as a reference (using the 'simple ratio' setting with background set to 0.0). These adjustments were done using the entire image prior to cropping to focus on the dying cell(s). Regions of interest (ROIs) were placed on the cell body of the dying cell and the mean intensity (gray value) measurements were taken. Due to the shape changes these cells undergo, we used the center of the cell body to ensure that the cell continued to be inside the ROI throughout the dying process. Percent change in fluorescence is calculated as  $((F_x - F_0)/F_0 \times 100\%)$  where  $F_x$  is the mean fluorescence intensity at time point  $x$  and  $F_0$  is the starting fluorescence at time = 0.

For generating kymographs, retina time stacks were first straightened out using the 'Straighten' function in Fiji (Edit > Selection > Straighten) on a segmented line that followed the retina curvature. The synapse and the cell body layers were then separately selected, resliced (Image > Stack > Reslice), and Z-projected (Sum slices) to generate two separate kymographs for each time-lapse. These two kymographs were then combined in two different channels (represented in magenta and green in Figure 5c) and analyzed for flashes. Individual flashing cells, represented as striped columns in the kymographs, were analyzed using the 'Plot Profile' function in Fiji (Analyze > Plot Profile) with options set to vertical profile. Adobe Photoshop CS5 (San Jose, CA, USA) was used to process the images further. Final figures were assembled using Adobe Illustrator CS5.

**Ca<sup>2+</sup> measurements in mouse rods.** [Ca<sup>2+</sup>]<sub>i</sub> in acutely isolated mouse rods was measured as described previously.<sup>22</sup> Briefly, postnatal day P8–P10 and P14–P21, *C57Bl6J* (C57) and *Pde6 $\beta$ <sup>rd1</sup>* (*rd1*) mice (Jackson Laboratories, Bar Harbor, ME, USA) were killed, enucleated and the retinas were treated with papain (30–50 U/ml; Worthington, Freehold, NJ, USA; 1 h at room temperature). Acutely isolated photoreceptor cells with intact somas were identified based on their size and morphology (Figure 3b;<sup>22</sup>). The number of mice ( $N$ ) and cells ( $n$ ) in each group was: C57 (P8–10):  $N = 3$ ,  $n = 122$ ; *rd1* (P8–10):  $N = 7$ ,  $n = 186$ ; C57 (P14–21):  $N = 3$ ,  $n = 126$ ; *rd1* (P14–21):  $N = 3$ ,  $n = 27$ . Following trituration, cells were plated on concanavalin A-coated (0.2 mg/ml; Sigma, St. Louis, MO, USA) coverslips, loaded with fura-2 AM (1–5  $\mu\text{M}$ ; Invitrogen, Life Technologies, Grand Island, NY, USA) for 30 min, and washed for 30 min in dye-free L-15 medium at room temperature. Fluorescence was measured using 340 and 380 nm excitation filters (Chroma, Brattleboro, VT, USA), whereas emission was high-pass filtered at 510 nm and captured with cooled digital CCD cameras (HQ2, Photometrics, Tucson, AZ, USA). Data acquisition (10 pairs of images/min) and fluorescence ratio calculations with background subtraction in ROIs encompassing the rod perikaryon were performed with NIS Elements (Melville, NY, USA). Calibration of cytosolic free [Ca<sup>2+</sup>]<sub>i</sub> was carried out by using 10  $\mu\text{M}$  ionomycin in combination with 0 Ca<sup>2+</sup>/1 mM EGTA followed by 10 mM Ca<sup>2+</sup> (Figure 3a;<sup>35</sup>). The apparent free [Ca<sup>2+</sup>]<sub>i</sub> was determined from the equation  $[\text{Ca}^{2+}]_i = ((R - R_{\min}) / (R_{\max} - R)) \times \beta \times K_d$ , where  $R$  is the ratio of emission intensity at 510 nm evoked by 340 nm excitation versus emission intensity at 510 nm evoked by 380 nm excitation;  $R_{\min}$  is the ratio at zero free Ca<sup>2+</sup>;  $R_{\max}$  is the ratio at saturating Ca<sup>2+</sup>;  $K_d$  the dissociation constant for Ca<sup>2+</sup> binding to fura-2 in the presence of millimolar Mg<sup>2+</sup> was taken from the literature (224 nM;<sup>35,36</sup>);  $\beta = (F_{380_{\max}}/F_{380_{\min}})$ .  $R$ ,  $R_{\min}$ ,  $R_{\max}$ , and  $\beta$  were measured and calculated for each ROI (cell) separately. Baseline values were established for each cell by averaging baseline [Ca<sup>2+</sup>]<sub>i</sub> over a 5 min period. Group baseline values are averages of individual cell data  $\pm$  S.E.M.

## Conflict of Interest

The authors declare no conflict of interest.

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## Author Contributions

EM, AL, and GS conducted zebrafish experiments. PB made calcium measurements from mouse rods. SS generated the *Trf2:tdTomato* transgenic zebrafish. SEB conceived and directed research. EM, AL, PB, DK and SEB analyzed data. Paper was written by SEB with major contributions from EM, DK, AL, and PB.

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