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<td>Grimley, Joshua S.; Li, Li; Wang, Weina; Wen, Lei; Beese, Lorena S.; Hellinga, Homme W.; Augustine, George James</td>
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Visualization of Synaptic Inhibition with an Optogenetic Sensor Developed by Cell-Free Protein Engineering Automation

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We describe an engineered fluorescent optogenetic sensor, SuperClomeleon, that robustly detects inhibitory synaptic activity in single, cultured mouse neurons by reporting intracellular chloride changes produced by exogenous GABA or inhibitory synaptic activity. Using a cell-free protein engineering automation methodology that bypasses gene cloning, we iteratively constructed, produced, and assayed hundreds of mutations in binding-site residues to identify improvements in Clomeleon, a first-generation, suboptimal sensor. Structural analysis revealed that these improvements involve halide contacts and distant side chain rearrangements. The development of optogenetic sensors that respond to neural activity enables cellular tracking of neural activity using optical, rather than electrophysiological, signals. Construction of such sensors using in vitro protein engineering establishes a powerful approach for developing new probes for brain imaging.

Introduction

Optogenetic sensors (OSs) based on the green fluorescent protein (GFP), combined with advances in optical imaging methods, have enabled quantitative monitoring of cellular processes with high spatiotemporal resolution (Okumoto, 2010). Many such OSs have been developed to track subcellular localization (Flach et al., 1994) or monitor protein–protein interactions (Xia et al., 2001), and have great potential for measurement of neuronal activity by enabling precise, cellular-level reporting of neural circuit activity (Zhang et al., 2010; Mancuso et al., 2011) based on optical rather than electrophysiological signals (Siegel and Isacoff, 1997; Kuner and Augustine, 2000; Hires et al., 2008). However, engineering OSs that monitor the small and rapid changes in the concentrations of signaling molecules typically associated with neuronal activity has been challenging (Miyawaki et al., 1997; Kuner and Augustine, 2000; Okumoto et al., 2005; Hires et al., 2008; Arosio et al., 2010).

Presynaptic release of GABA opens postsynaptic GABA receptors, resulting in Cl− fluxes that underlie synaptic inhibition. Fluorescent OSs that monitor intracellular Cl− concentration, [Cl−], therefore could report on this important class of synaptic action. A first-generation Cl− sensor, Clomeleon, has been used to monitor such dynamic changes in [Cl−], in cultured neurons (Kuner and Augustine, 2000) and in brain tissue (Berglund et al., 2006). Clomeleon is a fusion of the yellow fluorescent protein (YFP), which contains a serendipitous Cl−/H11002 sensitive cyan fluorescent protein (CFP). Halide binding to YFP quenches fluorescence emission (Wachtler and Remington, 1999), altering fluorescence resonance energy transfer (FRET) between the CFP donor and the YFP acceptor (Kuner and Augustine, 2000). The use of FRET enables ratiometric determination of [Cl−], which is unaffected by variations in fluorescence emission intensity associated with differences in indicator concentration, optical path length, or excitation intensity (Bright et al., 1989; Miyawaki, 2005). However, it remains extremely challenging to detect GABA-induced [Cl−] changes with Clomeleon because the Cl− affinity of Clomeleon (~100 mM) is well beyond the physiological range of [Cl−] (~5–6 mM in most adult neurons; Berglund et al., 2006). This limits the signal-to-noise ratio (s/n), requiring averaging of multiple trials before a response can be detected via fluorescence imaging.
Here we report the use of recently developed protein engineering automation techniques (Cox et al., 2007; Allert et al., 2010) to improve the fluorescence response of Clomeleon by manipulating its halide affinity and fluorophore characteristics. The resulting sensor, SuperClomeleon, has a significantly improved s/n in the relevant range of postsynaptic [Cl\(^{-}\)] compared to the parent protein.

Materials and Methods

Protein engineering. Oligonucleotides (79–80 bases) were synthesized by solid phase oligonucleotide synthesis (MerMade 192; BioAutomation) and assembled into full-length open reading frames (ORFs) by automation (Cox et al., 2007), which were reamplified by PCR using 5'-biotinylated primers. Protein was produced by in vitro-coupled transcription and translation (TnT) reactions (Allert et al., 2010); 2.4 μg of linear, biotinylated dsDNA was added to 30 μl of a BL21 Star (DE3; Invitrogen) cell lysate and a solution of amino acids, nucleotide triphosphates, and cofactors and water (final volume 120 μl) in 96-well PCR plates, which were covered with a breathable seal for optimum aerobic expression and chromophore maturation (Heim et al., 1994), and incubated with shaking for 8 h at 30°C, followed by 6 h at 4°C.

All proteins had combined C-terminal Flag and His\(_6\) peptides for affinity purification. Proteins were purified at 4°C using TnT reactions with anti-Flag M2 agarose (Sigma) and Cl\(^{-}\)-free buffer: four (15 min) washes of agarose-bound FPs with 20 mM HEPES, pH 7.1, removed Cl\(^{-}\) and water (final volume 120 μl) in 96-well PCR plates, which were covered with a breathable seal for optimum aerobic expression and chromophore maturation (Heim et al., 1994), and incubated with shaking for 8 h at 30°C, followed by 6 h at 4°C.

The resulting sensor, SuperClomeleon, has a significantly improved s/n in the relevant range of postsynaptic [Cl\(^{-}\)] compared to the parent protein.
ments): 25°C, 4 ml quartz cuvettes, exciting at 450 ± 1 nm, collecting emission spectra (460–550 nm), and 4 nm slit widths.

Thermal stabilities were measured following fluorescence emission intensities in a real-time PCR instrument (LightCycler 480 II; Roche), 384-well microtiter plates, heating at 1.1°C/min (20–95°C), 10 acquisitions/s, 1 s integration time, and a 498 ± 40 nm excitation filter and 580 ± 20 nm emission filter. The denaturation transition midpoint temperature (Tm) values were obtained from first-derivative transforms of the data. Conventionally cloned and expressed single-domain Q69T/V163A variant (79.2 ± μs in 20 mM HEPES, pH 7.1) was diluted to 3 μs in 2 ml of 20 mM MES, pH 6.0, in a quartz cuvette with a Teflon-coated stir bar and stopper. Coincidence of thermal melts monitored by fluorescence and by circular dichroism (CD) was determined using an Aviv model 202 CD spectrophotometer, the sample was heated to 98–20°C while simultaneously monitoring the CD (222 nm) and fluorescence (513 nm excitation monochromator and 530 nm Schott high-pass filter) signals.

Changes in error, δS/δS, (Marvin et al., 1997), of variant i relative to a reference species, ref, at the midpoint of analytic fluctuations, S0, can be defined as follows:

\[
\frac{\delta S}{\delta S} = \frac{S_{i0} + K_{d, i} \cdot F_{\text{max}, i}}{S_{i0} + K_{d, ref} \cdot F_{\text{max}, ref}} \cdot \frac{S_{ref} \cdot \delta K_{d, i}}{S_{ref} \cdot \delta K_{d, ref}}
\]

where \( \delta P_{\text{M}} > 1 \) for improved variants (for [Cl]−, sensing, the midpoint, M, is 5.5 mM).

X-ray crystallography. The Q69T/V163A variant was purified by IMAC (see above), followed by gel filtration (Superdex200; GE Healthcare), eluting with 50 mM Tris and 150 mM NaCl, pH 7.5. The protein was concentrated, dialyzed, spin concentrated (10 kDa cutoff; Sartorius) to 384-well microtiter plates, heating at 1.1°C/min (20°C–95°C), 10% PEG 3K, 150 mM NH4OAc at pH 5.4) for 3 days at 17°C. The crystals were transferred to cryoprotectant (22% (w/v) PEG 2K, 50 mM NaOAc at pH 4.6, 90 mM MgCl2; or 10% PEG 3K, 150 mM NH4OAc at pH 5.4) for 3 days at 17°C. The crystals were transferred to cryoprotectant (22% (w/v) PEG 2K, 50 mM NaOAc at pH 4.6) with 20% ethylene glycol and either water (apo) or 100 mM KI (Wachter et al., 2000), equilibrated at room temperature (2 h), and flash frozen in liquid nitrogen. X-ray diffraction data were collected at the Advanced Light Source SIBYLS beam line at 1.1109 Å (apo, 3ST0) and flash frozen in liquid nitrogen. X-ray diffraction data were collected at the Advanced Light Source SIBYLS beam line at 1.1109 Å (apo, 3ST0) and 1.1169 Å (iodide, 3SV5). The diffraction images were processed using HKL2000 (Otwinowski and Minor, 1997). The crystal structure was solved by molecular replacement using Phaser (McCoy et al., 2007) and the YFP structure 2O24 (Arosio et al., 2007) as the search model. Initial model building and refinement was done in REFMAC (Murshudov et al., 1997), further refinement including ligand occupancy was achieved using HKL2000 (Otwinowski and Minor, 1997). The crystal structure was solved by molecular replacement using Phaser (McCoy et al., 2007) and the YFP structure 2O24 (Arosio et al., 2007) as the search model. Initial model building and refinement was done in REFMAC (Murshudov et al., 1997), further refinement including ligand occupancy was achieved using HKL2000 (Otwinowski and Minor, 1997). The crystal structure was solved by molecular replacement using Phaser (McCoy et al., 2007) and the YFP structure 2O24 (Arosio et al., 2007) as the search model. Initial model building and refinement was done in REFMAC (Murshudov et al., 1997), further refinement including ligand occupancy was achieved using HKL2000 (Otwinowski and Minor, 1997). The crystal structure was solved by molecular replacement using Phaser (McCoy et al., 2007) and the YFP structure 2O24 (Arosio et al., 2007) as the search model. Initial model building and refinement was done in REFMAC (Murshudov et al., 1997), further refinement including ligand occupancy was achieved using HKL2000 (Otwinowski and Minor, 1997).

Cellular imaging. Neurons were obtained from cortex or hippocampus of postnatal day 0–2 mice (Mus musculus) of either sex and cultured according to standard procedures (Kuner and Augustine, 2000). Neurons were transfected 3–4 d after culturing, via Lipofectamine 2000 (Invitrogen; Giller et al., 2004).

OS FRET was imaged in the cell body of transfected neurons by using an upright two-photon microscope (Prairie Technology) equipped with a 60× water-immersion objective (Olympus), collecting fluorescence emission of CFP (485 ± 20 nm) and YFP (535 ± 15 nm) in parallel. The CFP donor was excited by 840 nm light from a Ti: sapphire laser (Chameleon; Coherent) with a Pockels cell to attenuate laser power. Images were acquired and analyzed with Prairie View software. Illumination and detection conditions were fixed for all measurements to facilitate comparisons between measurements made on different OIs. The experiments shown in Figure 9, D–F, were done similarly, but used an Olympus FV-1000 two-photon microscope with a 25× objective. It is important to note that the absolute values of FRET ratios depend on the acquisition conditions, such as photomultiplier gain and filter properties, and will therefore differ between microscopes.

To vary [Cl−], neurons (3–4 days in vitro) were transfected with OS variants, placed in a perfusion chamber, and treated with ionophores (10 μM nigericin and 5 μM tributyltin acetate) to clamp [Cl−], and intracellular pH to extracellular levels. Physiological salines containing various [Cl−] were perfused through the chamber at a rate of 1 ml/min. High-[Cl−] saline contained 161 mM Cl−, 10 mM HEPES, 20 mM d(-)-glucose, 2 mM Na-EGTA, and 4 mM MgCl2. Cl−-free saline consisted of 10 mM HEPES, 20 mM d(+)-glucose, 48 mM Na-glucosate, 105 mM K-glucosate, 2 mM Na-EGTA, and 4 mM Mg (glucosate)2. Intermediate [Cl−] solutions were prepared by mixing the two salines. The KF saline used to fully quench YFP fluorescence contained 10 mM HEPES, 20 mM d(-)-glucose, 48 mM NaF, 105 mM KF, 2 mM Na-EGTA, and 4 mM Mg (glucosate)2. All salines were adjusted to pH 7.1.

Quantifying FRET measurements. FRET-based Cl− indicators such as Clomeleon and SuperClomeleon have emission spectra that change shape as a function of [Cl−]. These changes in the shape of the emission spectrum are quantified by measuring fluorescence emission at two wavelengths, λ1 and λ2, corresponding to the maximum for the CFP donor and YFP acceptor, respectively. An emission ratio, R, is then calculated from the fluorescence emission (F) measured at these two wavelengths, F1 and F2, as follows:

\[
R = \frac{F_2}{F_1}
\]

Cl− decreases the fluorescence of the YFP acceptor, F2. As a result, this ratio declines from a maximum, [Cl−]max in the absence of Cl− to a minimum, [Cl−]min in saturating [Cl−]. To convert the fluorescence emission ratio into [Cl−], the calibration scheme presented in Grynkiewicz et al. (1985) must be modified to take into account the fact that the titration curves decrease as a function of [Cl−]. Here we recapitulate the scheme of Grynkiewicz et al. (1985), in terms appropriate for measurement of [Cl−], by indicators such as Clomeleon and SuperClomeleon.

The starting point in this scheme is to state the fluorescence intensity at the two emission wavelengths in terms of four proportionality constants, s, representing the relative fluorescence of the Cl− bound (subscript b)
and Cl\textsuperscript{−} free (subscript f) forms of indicator at emission wavelengths \( \lambda_1 \) and \( \lambda_2 \). The total fluorescence of the indicator at each wavelength is then:

\[
F_1 = S_f C_f + S_b C_b
\]

(3)

\[
F_2 = S_f C_f + S_b C_b
\]

(4)

where \( c \) indicates the concentrations of the Cl\textsuperscript{−} bound (subscript b) and Cl\textsuperscript{−} -free (subscript f) species. \( s_f \) and \( s_b \) represent the fluorescence of the CFP donor in Cl\textsuperscript{−} -free and saturating conditions, respectively. The relative fractions of the Cl\textsuperscript{−} bound and Cl\textsuperscript{−} -free forms of the indicator vary with [Cl\textsuperscript{−}] according to the Cl\textsuperscript{−} dissociation constant, \( K_d \), of the indicator:

\[
K_d = \frac{c_f}{c_b}([Cl^-]).
\]

(5)

Combining Equations 3a and b allows \( R \) to be expressed in terms of \( s \) and \( c \):

\[
R = \frac{s_f C_f + s_b C_b}{s_f C_f + s_b C_b}.
\]

(6)

Combining Equations 4 and 5 then yields:

\[
R = \frac{s_f + \frac{s_f C_f}{K_d}}{s_b + \frac{s_f C_f}{K_d}}.
\]

(7)

This can be rearranged into the following expressions:

\[
[Cl^-] (R_{Sb} - s_b) = K_d (s_f - R s_f)
\]

(8)

\[
[Cl^-] = K_d \frac{\left(\frac{s_f}{s_b} - R \right) \left(s_f\right)}{R - \frac{s_b}{s_b} \left(s_b\right)}.
\]

(9)

The limiting values for the titration curve, \( R_{max} \) and \( R_{min} \), can be restated in terms of their component \( s \) constants:

\[
R_{max} = \frac{s_f}{s_f}
\]

(10)

\[
R_{min} = \frac{s_b}{s_b}
\]

(11)

Substituting these into Equation 8 yields:

\[
[Cl^-] = K_d \left(\frac{R_{max} - R}{R_{min} - R}\right) \left(\frac{s_f}{s_b}\right).
\]

(12)

Note that although the form of this equation closely resembles Equation 5 of Grynkiewicz et al. (1985), many of the terms are different.

It is worth emphasizing that while \( K_d \) is independent of the specific instrument used to measure \( R \), all other variables on the right side of Equation 11 are instrument-specific. Thus, \( R_{max}, R_{min}, s_f, \) and \( s_b \) must be determined empirically for each instrument used to measure \( R \) for Clomeleon, SuperClomeleon, or other FRET-based Cl\textsuperscript{−} indicators. This can be done by varying [Cl\textsuperscript{−}] using the procedures described above.

Electrophysiology. Neurons were patch clamped via glass pipettes (5–6 MΩ resistance) filled with an internal solution containing 5 mM Cl\textsuperscript{−}: 140 mM K-glucuronate, 10 mM HEPES, 4 mM Na-ATP, 0.4 mM Na-GTP, 5 mM EGTA, 0.5 mM CaCl\textsubscript{2}, and 2 mM MgCl\textsubscript{2}. Extracellular saline contained 150 mM NaCl, 3 mM KCl, 2 mM CaCl\textsubscript{2}, 2 mM MgCl\textsubscript{2}, 20 mM D(+)-glucose, and 10 mM HEPES, pH 7.4. Electrophysiological signals were recorded using a Multiclamp 700B amplifier (Molecular Devices) and digitized with a Digidata 1440 (Molecular Devices) A-D converter. Data were acquired and analyzed with pClamp10.1 software (Molecular Devices). GABA was locally applied onto neurons via pressure ejection. For this purpose, a glass pipette containing GABA (1 mM) was connected to a Picospritzer II (General Valve) to provide 100 ms pulses. IPSCs were evoked by stimulating presynaptic neurons using a concentric bipolar electrode (FHC). A train of extracellular current pulses (75 μA, 10 Hz, 2 s) were applied near the neuron being recorded from, while blocking EPSCs by adding 20 μM CNQX and 50 μM APV to the extracellular solution.

Throughout this paper, error determinations represent the SEM, unless indicated otherwise. Statistical comparisons were made with the Student’s t test (Press et al., 2007, p. 729).

Results

Matching sensor affinity with [Cl\textsuperscript{−}]

Structural studies of YFP (Wachter et al., 2000; Griesbeck et al., 2001; Fig. 1A) have shown that its halide-binding site comprises the chromophore and nine nearby residues (Fig. 1B). The principles of molecular recognition determining halide binding are not well defined. Consequently, it is impossible to predict a priori how mutations in the halide-binding site contribute to binding affinity. We therefore used recently developed cell-free protein engineering methods (Cox et al., 2007; Allert et al., 2010) that bypass cloning steps (Fig. 2) to rapidly explore the Cl\textsuperscript{−} binding and fluorescence properties of single, double, and triple mutants in the crystallographically defined halide-binding site. We mutated seven of these residues, avoiding Arg96, which catalyzes chromophore maturation (Barondeau et al., 2003; Sniegowski et al., 2005) and Tyr203, which tunes the spectral characteristics of YFP (Dickson et al., 1997). Starting with all 133 single mutants at these positions, double and triple mutations were iteratively constructed in CT in the absence of the CFP FRET donor (Kuner and Augustine, 2000). Another iterative set of mutants beginning with the 133 single mutations were also tested in CT H148Q, a mutation previously used to improve Cl\textsuperscript{−} affinity (Jayaraman et al., 2000; Galietta et al., 2001; Markova et al., 2008) but with diminished fluorescence (Elslinger et al., 1999).

In noncooperative binding to a single site, changes in fluorescence emission intensity are directly proportional to the fraction of binding sites occupied by Cl\textsuperscript{−} (\( j \)). Sensor responses are maximal when the dissociation constant, \( K_d \), matches the midpoint of the relevant [Cl\textsuperscript{−}] range (Marvin et al., 1997); in a mature neuron, [Cl\textsuperscript{−}], rises from ~5 to ~6 mM during synaptic inhibition (Berglund et al., 2006) with a midpoint of 5.5 mM. For a single-site hyperbolic binding isotherm, over such a range the theoretical maximal signal is 46.6% (\( K_d = 5.5 \text{ mM}, \Delta y_{max} = y_1 - y_0 = 0.046 \)) of the difference in fluorescence intensities observed in the absence and saturating presence of Cl\textsuperscript{−} (\( \Delta F_{max} \)). For Clomeleon (\( K_d = 119 \text{ mM} \)), the maximal signal is 0.8% of \( \Delta F_{max} \); matching
Cl⁻ affinity therefore could achieve a 5.9-fold improvement in signal, if \( \Delta F_{\text{max}} \) remained constant.

Clomeleon variants were rapidly screened using a cell-free protein engineering methodology that obviates cloning steps (Fig. 2). Linear DNA fragments encoded ORFs flanked by T7 RNA polymerase promoter and terminator sites were assembled robotically from synthetic oligonucleotides (Cox et al., 2007). The ORF mRNA sequence of CT was optimized computationally for \( E. \text{coli} \) expression (Allert et al., 2010). The resulting DNA fragments were then used to produce proteins by combined \textit{in vitro} transcription and translation in 120 aliquots of \( E. \text{coli} \) extracts, purified by small-scale affinity purification using beads, and assayed for Cl⁻ binding in microtiter plates.

A total of 424 variants, encoded in genes constructed from 344 unique oligonucleotides, were tested in this manner. The genes encoding the initial 266 single-point mutants were assembled from 269 oligonucleotides. Following an initial triage of these single-point variants based on fluorescence and expression levels, the Cl⁻ affinities of 131 variants were determined by Cl⁻ titration using a fluorescence plate reader. Their response to Cl⁻ was evaluated using a metric that takes into account both fluorescence and affinity matching, \( \rho_{\text{s,5}} \) (see Materials and Methods) Twelve single mutants showed performance improvements (\( \rho_{\text{s,5}} > 1 \)) with Q69T the only one having \( \rho_{\text{s,5}} > 2 \) (Fig. 3A,B).

In subsequent assembly rounds, we constructed 150 double mutants from a subset of 29 single mutants and included single-point variant with \( \rho_{\text{s,5}} < 1 \) to test whether improvements can arise by combining suboptimal intermediates. Out of these 150 double mutants, the five best (\( \rho_{\text{s,5}} \geq 2.32 \)) were combinations of only six single mutations (Q69T, V150A/I, and V163A/G/T), from which the set of all six possible triple mutants was constructed. With this automated gene assembly method, only 75 additional oligonucleotides were needed to construct the genes encoding the 156 higher-order mutants.

The average \( K_d \) values for Cl⁻ binding of the 285 characterized CT variants improved from 101 mM (single) to 77.5 mM (double) to 33.3 mM (triple mutants), with a concomitant increase in average \( \rho_{\text{s,5}} \) values from 0.43 to 0.83 to 2.21. Based on their \( \rho_{\text{s,5}} \) values, 63 variants (22%) were improvements over CT (Fig. 3B) and had mutations in at least one of six binding site residues. The five highest affinity variants (\( K_d < 10 \text{ mM} \)) were single mutations at Gln183; these exhibited diminished fluorescence brightness and a reduced \( \Delta F_{\text{max}} \) response, precluding their use in cellular measurements (Fig. 3A,B). The 13 best variants had \( \rho_{\text{s,5}} > 2 \) and included mutations at three positions (Gln69, Val150, and Val163). The best CT mutant, Q69T/V163A,
was composed of individual single mutants each with $p_{S,5} > 1$ and was almost three times more sensitive than CT in vitro (Table 1).

Using conventionally cloned and expressed protein, we assayed the binding and stability of 3 of the top 13 variants, as well as the original CT, the highest affinity variant Q183A, chloride-insensitive mutants Q69H and Q69M (Griesbeck et al., 2001), and four H148Q variants. Cl$^-$ affinities were determined by titration in a fluorescence plate reader. For most variants, the results were similar to the in vitro screening observations (Fig. 3C; Table 1). The Cl$^-$ dependence of thermal stabilities, which is thermodynamically linked to binding (Isom et al., 2010; Layton and Hellinga, 2010), was used to confirm the binding properties determined by titration. To determine thermal stabilities, the temperature dependence of the chromophore fluorescence was followed, which was coincident with the thermal stability determined by circular dichroism (Fig. 4A). Cl$^-$ increased the stability of chloride-responsive proteins, whereas the nonresponsive variants Q69H and Q69M showed minimal changes (Fig. 4B), consistent with the presence or absence of Cl$^-$-binding sites inferred from fluorescence observations. No consistent trend in relative affinities for F$^-$, Br$^-$, and I$^-$ was observed (Table 1), indicating that molecular recognition of halides is quite complex (Wachter and Remington, 1999; Jayaraman et al., 2000; Kuner and Augustine, 2000). The pH dependence of fluorescence response and halide affinity (Fig. 5A,B) reveals linkage between chromophore ionization and binding affinity (see Discussion).

**Structural analysis**

High-resolution x-ray structures of the halide-free (apo) and I$^-$ complexes were determined for the Q69T/V163A variant that was used in cellular measurements (Fig. 6, Table 2). With the exception of some distant loops, the backbones of these two structures and YFP (Wachter et al., 2000) are essentially the same. Two I$^-$ sites were observed by anomalous scattering in the vicinity of the chromophore (Fig. 6A–E). One site (I) was observed previously (Wachter et al., 2000). The second site (II) is located 9 Å away, where I$^-$ replaces an interior water observed in the apo-protein. Small rearrangements of Thr63 and Thr108 side chains in the complex and apo-protein (Fig. 6F), relative to YFP, enable I$^-$ binding at Site II. The two side chain hydroxyls interact with the bound I$^-$. Ethylene glycol (EG, used as cryoprotectant at 36% in structure determination) also bound to Site I (Fig. 6C,E). Both I$^-$ and EG were observed at less than full occupancy in the complex, suggesting that these two ligands compete. EG binding neither was detectable by fluorescence quenching nor affected Cl$^-$ affinity and therefore EG is a low-affinity ligand. Formate was also observed to bind in the vicinity of the chromophore (Fig. 6G); it is likely to originate from ultrafiltration membranes used in protein purification.

**FRET performance**

We fused the affinity-tuned CT variants with a CFP FRET donor to reconstruct improved versions of Clomeleon. The performance of these OSs was evaluated in vitro (Table 3) and in a series of cellular FRET measurements. The Cl$^-$ affinities of these sensors were determined by Cl$^-$ titration experiments (Fig. 7), using two-photon imaging to measure sensor FRET in the cell bodies of individual cultured hippocampal neurons transfected with the OS of interest (Fig. 7A). [Cl$^-$], was varied by permeabilizing the cell membrane to external Cl$^-$ via ionophores (Berglund et al., 2006). The resulting changes in FRET emission (Fig. 7B) yielded titration curves such as those shown in Figure 7C. These curves were fit using Equation 11 to determine the $K_d$ for Cl$^-$ binding. When imaging synaptic inhibition in neurons, the most important parameter is s/n. We therefore also estimated the s/n for the novel OSs, using the approach described above: “signal” corresponded to the change in FRET as [Cl$^-$], was raised from 5 to 6 mM, and “noise” was the SD of the FRET ratio determined at 5 mM [Cl$^-$], for each cell.
Table 1. Chloride binding to single domain YFP variants

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<th>YFP mutation</th>
<th>Cell-free Cl(^-) screen</th>
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<td>(\Delta F_{\text{max}})</td>
<td>(K_s)</td>
<td>(F^-)</td>
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<td>((\text{mW}))</td>
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<td>(K_a) ((\text{mM}))</td>
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<td>CT</td>
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<td>Q69M</td>
<td>0.36</td>
<td>0.17</td>
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</table>

Chloride binding to single domain YFP variants. All halide titration and thermal stability data determined at pH 7.1. \(\Delta F_{\text{max}}\) values are scaled relative to Clomeleon Topaz. \(F^-\) endpoint was not included in the halide titration fits of conventionally prepared protein.

FRET depends upon both the distance between, and the orientation of, the fluorescent donor and acceptor. As a result, the linker between the donor and acceptor is an important determinant of FRET efficiency in GFP-based OSs (Evers et al., 2006). Therefore, to improve FRET efficiency, we used the strategy of Shimozono et al. (2006) to truncate the linker between the CFP and YFP of Clomeleon-based OSs: 40 residues (11 residues from the C terminus of CFP, the 24-residue linker, and 5 residues from the N terminus of YFP) were replaced with two residues. This truncation (short linker) did increase FRET efficiency, evident as an increase in the difference in FRET ratios (\(\Delta F_{\text{max}}\)) observed in the absence of Cl\(^-\) and in the presence of saturating concentrations of Cl\(^-\) (or \(F^-\); Fig. 7C). This version of Clomeleon exhibited enhanced Cl\(^-\) binding affinity (Fig. 7C, inset) and improved s/n (Fig. 7D; \(n = 4\)). The mean s/n measured for this version was significantly better than that of Clomeleon (Student’s \(t = 52.1, p = 0.000\)).

Because donor and acceptor fluorescence also play important roles in determining s/n, we next introduced into both the donor and acceptor fluorophores a mutation (S30R) that increases brightness and stability (Pédelaq et al., 2006). This pair of mutations produced the largest improvement in \(\Delta F_{\text{max}}\) (Fig. 7C) and further improved s/n (Fig. 7D; \(n = 18\)), but had no effect on Cl\(^-\) affinity (Fig. 7C, inset).

After making these improvements to the FRET properties of Clomeleon, we next replaced the YFP FRET acceptor with several affinity-tuned CT variants. All of these variants increased the Cl\(^-\) affinity of Clomeleon: V163S (27.0 ± 0.1 mM; \(n = 8\)), I152L/V163A (4.8 ± 0.2 mM; \(n = 4\)), V150A/V163A (4.6 ± 0.2 mM; \(n = 5\)), and V163A/L201I (2.5 ± 0.6 mM; \(n = 5\)). All of these also improved s/n above that produced by the Clomeleon variant with the short linker and S30R mutations (Fig. 7D). We also tested an OS with YFP H148Q/I152L/V163S, a variant previously reported to have improved properties (Markova et al., 2008). However, for FRET-based measurements, this OS had s/n properties no better than those of Clomeleon (Fig. 7D). This is due to its relatively poor Cl\(^-\) affinity (27.0 ± 5.0 mM; \(n = 4\)) and the diminished fluorescence of the FRET acceptor (Table 1).

The largest improvement in s/n was observed when the affinity-tuning mutations in the best CT variant from the cell-free screen, Q69T/V163A, were included in the OS FRET acceptor.
This sensor had a Cl\(^{-}\) affinity of 8.1 ± 0.5 mM. \(\Delta R_{\text{max}}\) for this version was smaller than for the short linker/S30R version of Clomeleon, but was still approximately twice as large as the value measured for Clomeleon (Fig. 7C). Most importantly, s/n for this variant was substantially larger than that measured for any of the previous versions of Clomeleon (Fig. 7D; \(n = 21\)).

In the last step, Clomeleon was further improved by replacing CFP with a brighter donor, Cerulean (Rizzo et al., 2004). This substitution decreased \(\Delta R_{\text{max}}\) somewhat (Fig. 7C) but did not affect Cl\(^{-}\) affinity. s/n for this construct was the best of any of the Clomeleon variants that we produced (Fig. 7D) and is signifi-

![Figure 6.](image)

**Table 2. Crystallographic data collection and refinement statistics**

<table>
<thead>
<tr>
<th></th>
<th>Q69T/V163A apo</th>
<th>Q69T/V163A iodide-bound</th>
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<tbody>
<tr>
<td><strong>Space group</strong></td>
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<td>P2(_1)2(_1)2(_1)</td>
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<tr>
<td><strong>Cell parameters</strong></td>
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<td>(I/\sigma(I))</td>
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Crystallographic data collection and refinement statistics. Values in parentheses refer to the highest resolution shell. All atoms are included in the \(B\)-factor calculations.

**Table 3. In vitro chloride binding to fusion proteins**

<table>
<thead>
<tr>
<th>Fusion protein components</th>
<th>In vitro</th>
<th>(K_d) (mM)</th>
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<tbody>
<tr>
<td>Donor(^a)</td>
<td>Acceptor</td>
<td>Linker(^b)</td>
</tr>
<tr>
<td>CFP(^d)</td>
<td>CT</td>
<td>Long</td>
</tr>
<tr>
<td>CFP</td>
<td>Q69T/V163A</td>
<td>Long</td>
</tr>
<tr>
<td>Cerulean</td>
<td>Q69T/V163A</td>
<td>Long</td>
</tr>
<tr>
<td>CFP</td>
<td>CT</td>
<td>Long</td>
</tr>
<tr>
<td>Cerulean</td>
<td>Q69T/V163A</td>
<td>Long</td>
</tr>
<tr>
<td>CFP</td>
<td>CT</td>
<td>Short</td>
</tr>
<tr>
<td>Cerulean</td>
<td>Q69T/V163A</td>
<td>Short</td>
</tr>
<tr>
<td>Cerulean</td>
<td>Q69T/V163A</td>
<td>Short</td>
</tr>
</tbody>
</table>

In vitro chloride binding to fusion proteins. \(^a\)CFP corresponds to the FRET donor in Clomeleon (Kuner and Augustine, 2000). \(^b\)Constructs with a long linker tested in vitro contained a scrambled protease cleavage site. \(^c\)S30R mutation was in both FPs. \(^d\)This is the original Clomeleon (Kuner and Augustine, 2000). \(^e\)SuperClomeleon.

Detecting GABA-mediated changes in [Cl\(^{-}\)]\(\text{in}\)

We determined the ability of SuperClomeleon to detect transient [Cl\(^{-}\)]\(\text{in}\), changes produced in neurons by brief applications of the inhibitory neurotransmitter GABA. Cultured hippocampal neurons were transfected with SuperClomeleon and GABA was locally applied from a pipette positioned near the neuron (Fig. 8A). The neurons were voltage clamped to monitor Cl\(^{-}\) flux and as a change in SuperClomeleon FRET (Fig. 8B, C). The magnitude and direction of both current and FRET responses varied with membrane potential according to the direction of Cl\(^{-}\) flux. At a holding potential of −40 mV, Cl\(^{-}\) influx...
was observed as a large, outward current and a decrease in SuperClomeleon FRET (Fig. 8B, left). At more negative potentials, where the electrochemical driving force on Cl\textsuperscript{−}/H\textsubscript{11002}\textsuperscript{+} influx was reduced, the currents and FRET responses became smaller and reversed their polarities at \textit{H}\textsubscript{11002}\textsuperscript{+} \textit{90 mV (Fig. 8B, right). The reversal potential of approximately \textit{H}\textsubscript{11002}\textsuperscript{+} 80 mV was consistent with the Cl\textsuperscript{−}/H\textsubscript{11002}\textsuperscript{+} equilibrium potential of \textit{H}\textsubscript{11002}\textsuperscript{+} \textit{89 mV predicted by the Nernst equation.}

To define the time course and magnitude of the GABA-induced Cl\textsuperscript{−}/H\textsubscript{11002}\textsuperscript{+} flux, we integrated the GABA-induced current over time (transmembrane charge). Comparison of the rising phases of the transmembrane charge and the change in SuperClomeleon FRET ratio indicated that the FRET response lags behind the Cl\textsuperscript{−} flux by 1–2 s (Fig. 8C). This is consistent with previous observations made with Clomeleon (Kuner and Augustine, 2000). For observations made on the single neuron shown in Figure 8B, there was a linear relationship between the Cl\textsuperscript{−} flux, as measured by the charge, and the integrated FRET signal (Fig. 8D). This correlation was also evident in 49 measurements from 10 neurons (Fig. 8E, red) and indicates that SuperClomeleon reliably reports the Cl\textsuperscript{−} fluxes and resulting changes in [Cl\textsuperscript{−}], produced by activation of GABA receptors. Much smaller FRET responses were observed in neurons expressing Clomeleon (Fig. 8E, black). The slope of the relationship between transmembrane charge and integrated FRET signal indicated that SuperClomeleon responses were 4.8-fold greater than those of Clomeleon for a given amount of Cl\textsuperscript{−} flux (Fig. 8E). This indicates that SuperClomeleon improves FRET signal amplitude 5-fold compared with Clomeleon. FRET signals declined with time constants of 6.9 ± 0.6 s (\textit{n} = 38) for SuperClomeleon and 4.9 ± 0.7 s (\textit{n} = 18) for Clomeleon, reflecting slow Cl\textsuperscript{−} efflux from neurons (Kuner and Augustine, 2000; Berglund et al., 2006). By analogy with fluorescent calcium indicators (Regehr and Atluri, 1995), the slower time course of decay observed by SuperClomeleon is probably due to the improved ability of this indicator to detect small and persistent changes in [Cl\textsuperscript{−}].

**Imaging GABA-mediated synaptic inhibition with SuperClomeleon**

One of the potentially most important applications of OSs is imaging the spatiotemporal dynamics of synaptic inhibitory circuits (Mancuso et al., 2011). While Clomeleon is capable of reporting the small changes in [Cl\textsuperscript{−}], associated with synaptic inhibition (Berglund et al., 2006, 2008), its low Cl\textsuperscript{−} sensitivity and very low s/n in the desired concentration range has prevented routine imaging of synaptic inhibition. We therefore evaluated the ability of SuperClomeleon to perform this de-
manding task in cultured hippocampal neurons by activating inhibitory synapses via electrical stimulation of nearby interneurons (Fig. 9A).

Postsynaptic neurons expressing Clomeleon or SuperClomeleon were voltage clamped (−50 mV) to measure the postsynaptic Cl− currents associated with IPSCs. Under these conditions, brief trains of electrical stimuli produced IPSCs, which decreased during the train (Fig. 9B, bottom) due to synaptic depression (Gitler et al., 2004b). In neurons expressing Clomeleon, inhibitory synaptic activity rarely produced detectable FRET changes (Fig. 9B, left). However, similar stimuli reliably produced transient reductions in SuperClomeleon FRET (Fig. 9B, right), revealing that this indicator is capable of detecting the relatively small changes in [Cl−], associated with synaptic inhibition. To compare the performance of the two OSs, we measured synaptic responses in eight cells expressing SuperClomeleon and in five cells expressing Clomeleon. While 100% (16/16) of the trials yielded detectable SuperClomeleon responses, Clomeleon reported responses only 50% (8/16) of the time.

Responses to inhibitory synaptic activity were quantified to provide a direct comparison of s/n for inhibitory synaptic responses reported by SuperClomeleon and Clomeleon. For the examples shown in Figure 9B, the s/n ratio was −5.8 for the neuron expressing SuperClomeleon and was too small to be determined reliably in the neuron expressing Clomeleon. This was further quantified by first calculating the s/n by integrating the stimulus-induced change in FRET signal and dividing this value by the baseline FRET noise (as done in Fig. 8D). Because of substantial cell-to-cell variations in IPSC amplitude, these responses were then normalized for the amount of synaptic Cl− flux by dividing their values by the synaptic charge produced by the stimulus. The cumulative probability plots shown in Figure 9C show the results of this analysis, with the dashed vertical line indicating values where there was no measurable response (i.e., a s/n of 0). This analysis revealed that, on average, the s/n of SuperClomeleon responses (median = 0.15 ± 0.01 s·pC−1) was 5.4-fold larger than those of Clomeleon (median = 0.028 ± 0.008 s·pC−1).

For many potential applications of SuperClomeleon, it is likely that the postsynaptic neuron would not be voltage clamped. Indeed, in many applications it is possible that there would be no electrophysiological measurements of postsynaptic activity at all. FRET responses should be smaller when the neurons are not voltage clamped, because the electrochemical driving force on Cl− will change as the membrane potential changes during the IPSP. Thus, we completed our analysis by comparing the ability of SuperClomeleon and Clomeleon to detect inhibitory activity in postsynaptic neurons that were not voltage clamped.

Under current-clamp conditions, the same brief trains of inhibitory synaptic activity produced a series of IPSPs (Fig. 9D, bottom). In neurons expressing Clomeleon, detectable FRET signals were observed in only three of nine experiments (Fig. 9D, left). In contrast, a similar amount of inhibitory synaptic activity produced detectable changes in FRET in every case (nine of nine experiments) for neurons expressing SuperClomeleon (Fig. 9D, right). The mean integrated change in FRET signal measured with SuperClomeleon was 5.4-fold larger than for the integrated change in FRET signal measured in neurons expressing Clomeleon (Fig. 9E). The difference in the performance of Clomeleon and SuperClomeleon for detecting changes in [Cl−], associated with IPSPs was even greater when considering s/n, rather than integrated change in FRET signal: the s/n for SuperClomeleon was >6 times larger than that measured for Clomeleon (Fig. 9F). In summary, our measurements under both voltage-clamp and current-clamp conditions indicate that the s/n was ~6-fold better for SuperClomeleon than for Clomeleon. We therefore conclude that SuperClomeleon is superior for measurements of changes in [Cl−], associated with synaptic inhibition.
Discussion
We used a multistage strategy to improve Cl\textsuperscript{−} sensing by Clomeleon. First, using a cell-free automation methodology to rapidly prototype variants in vitro, we improved YFP by more closely matching its Cl\textsuperscript{−} affinity to postsynaptic [Cl\textsuperscript{−}]. Next, we incorporated this affinity-tuned single domain into Clomeleon for FRET-based sensing, enhancing s/n in cellular imaging experiments. Improvements in affinity matching, while retaining fluorescence, resulted in the largest enhancement of cellular imaging performance. Shortening the interdomain linker and introducing brightening mutations significantly improved the FRET response. This process culminated in identification of SuperClomeleon, a variant that greatly enhanced observation of synaptic inhibition in neurons.

Choosing single-point mutants with clearly improved $p_{S,5}$ values was a critical step for success; incremental improvements also arose from combinations of initially suboptimal variants. Nevertheless, given that the number of variants that need to be examined in an enumerative strategy can increase approximately exponentially with each iteration that includes more positions, carrying forward only clearly improved variants balances experimental load and likelihood of success. We observed good agreement between the affinities determined in the initial screen and the proteins produced conventionally. Cell-free protein engineering therefore is a powerful tool for generating and prototyping mutant proteins.

Our structural analysis revealed that the effects of the mutations are complex. The halide-binding site of SuperClomeleon resides within the internal cavity that accommodates the conformational change required for chromophore cyclization (Barondeau et al., 2003). This internal cavity accommodates two halide-binding sites, as well as a molecule of EG and a molecule of formate. The appearance of a second halide-binding site accompanied by motions of the hydrogen-bonding side chains in the otherwise rigid structure shows that the lining of this cavity is mobile and can accommodate binding sites through subtle rearrangements. The presence of small organics, such as ethylene glycol and formate, suggests that this cavity can be further diversified to bind and detect small molecules.

The YFP chromophore exists in four possible charge states (anionic, cationic, zwitterionic, and neutral), of which only the anionic form is fluorescent (Elsäger et al., 1999). Bound Cl\textsuperscript{−} and the anionic chromophore interact unfavorably by electrostatic repulsion, so that Cl\textsuperscript{−} binding is favored in the protonated, neutral, nonfluorescent state. These interactions link [Cl\textsuperscript{−}] and affinity with pH and chromophore $p_K_c$, setting up a trade-off between Cl\textsuperscript{−} affinity and sensor brightness. Improvements in Cl\textsuperscript{−} affinity correlate with raised chromophore $p_K_c$ values relative to CT, which decreases the concentration of anionic chromophore in the halide-free protein at a given pH, thereby diminishing sensor brightness. This decreased fluorescence reduces $\Delta F_{\text{max}}$ because Cl\textsuperscript{−} binding quenches emission. The highest affinity variant Q183A has a raised $p_K_c$ value and diminished fluorescence brightness. Similarly, other mutations, including H148Q, increase affinity at the expense of brightness (Jayaraman et al., 2000; Wachter et al., 2000; Galiotta et al., 2001; Markova et al., 2008; Fig. 3 A, B). In constructing SuperClomeleon these trade-offs could be balanced sufficiently to improve sensing substantially; that is, SuperClomeleon offers improved sensitivity to physiological Cl\textsuperscript{−} fluxes despite a higher interference by protons.

Affinity matching together with other enhancements improved the s/n characteristics of SuperClomeleon in cellular imaging. As a result, SuperClomeleon represents a significant improvement over Clomeleon and consistently outperformed Clomeleon in all of our neuronal imaging experiments: Cl\textsuperscript{−} titrations, activation of GABA receptors via exogenous GABA, and electrically evoked synaptic inhibition. Quantitatively, our GABA application experiments indicate a 4.8-fold improvement in signal for SuperClomeleon relative to Clomeleon, while the synaptic inhibition data indicate a s/n that is 5- to 6-fold larger than that
measured for Clomeleon. The key qualitative improvement is that SuperClomeleon robustly detects inhibitory synaptic activity in single neurons even under the relatively noisy conditions of two-photon imaging; this was not the case for Clomeleon (Berglund et al., 2006, 2008). Of particular note is that SuperClomeleon can detect responses to brief IFSP trains, an advantage that will prove useful for many future applications and will enable optogenetic imaging of the spatiotemporal dynamics of synaptic inhibitory circuits at the cellular level of resolution (Mancuso et al., 2011).

In conclusion, we have partnered automated in vitro protein engineering technology with cellular imaging to develop an OS with improved ability to image inhibitory synaptic activity. This development not only enables novel brain imaging experiments but also establishes a general approach for developing and improving many other types of OSs.

References

Grimley, Li et al. “Engineering SuperClomeleon”


