Directly light-regulated binding of RGS-LOV photoreceptors to anionic membrane phospholipids

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We report natural light–oxygen–voltage (LOV) photoreceptors with a blue light-switched, high-affinity (Kd ∼ 10−7 M), and direct electrostatic interaction with anionic phospholipids. Membrane localization of one such photoreceptor, BcLOV4 from Botrytis cinerea, is directly coupled to its flavin photocycle, and is mediated by a polybasic amphipathic helix in the linker region between the LOV sensor and its C-terminal domain of unknown function (DUF), as revealed through a combination of bioinformatics, computational protein modeling, structure–function studies, and optogenetic assays in yeast and mammalian cell line expression systems. In model systems, BcLOV4 rapidly translocates from the cytosol to plasma membrane (∼1 second). The reversible electrostatic interaction is nonselective among anionic phospholipids, exhibiting binding strengths dependent on the total anionic content of the membrane without preference for a specific headgroup. The in vitro and cellular responses were also observed with a BcLOV4 homolog and thus are likely to be general across the dikarya LOV class, whose members are associated with regulator of G-protein signaling (RGS) domains. Natural photoreceptors are not previously known to directly associate with membrane phospholipids in a light-dependent manner, and thus this work establishes both a photosensory signal transmission mode and a single-component optogenetic tool with rapid membrane localization kinetics that approaches the diffusion limit.

Significance

Light–oxygen–voltage (LOV) domain photoreceptors are found ubiquitously in nature and possess highly diverse signaling roles and mechanisms. Here, we show that a class of fungal LOV proteins dynamically associates with anionic plasma membrane phospholipids by a blue light-switched electrostatic interaction. This reversible association is rapidly triggered by blue light and ceases within seconds when illumination ceases. Within the native host, we predict that these proteins regulate G-protein signaling by the controlled recruitment of fused regulator of G-protein signaling (RGS) domains; in applied contexts, we anticipate that engineered chimeric versions of such proteins will be useful for rapid optogenetic membrane localization of fused proteins through direct interaction with the membrane itself, without requiring additional components to direct subcellular localization.


The authors declare no conflict of interest.

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Data deposition: The plasmid for mammalian codon-optimized BcLOV4 fused to mCherry and the plasmid for native sequence BcLOV4 fused to mCherry were deposited with AddGene (https://addgene.org) (accession nos. 114595 and 114596).

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Results

Domain Topology and in Vitro Photophysical Characterization. Conserved domain analyses by us and others (3, 17–20) report a consensus RGS-LOV architecture with a low-complexity region and RGS domain located N-terminal to a single LOV domain. Secondary-structure predictions and structural modeling here indicate that there is an additional C-terminal domain of unidentified function (DUF) with mixed α-helix/β-sheet content as well (Fig. 1 A and B). The LOV and DUF domains are connected via a predicted LOV ζα-helix linker, which extends into a polybasic amphipathic helix (AH) and is known to mediate LOV signaling elsewhere (4–6). RGS-LOV candidates from five organisms were chosen for either their previously hypothesized functional roles by others (17, 18) or their short length, and were assessed for solubility as full-length proteins in bacterial expression systems (SI Appendix, Fig. S1). One of these, BcLOV4 (also named BcRGS1) from the noble rot fungus Botrytis cinerea (18, 21) (GenBank accession number CCD53251.1), could be produced in good yield as a dark-adapted oligomer and was used for further analyses (SI Appendix, Fig. S2).

BcLOV4 had an optical absorbance peak at λ_{max} = 450 nm with triplet-peak fine structure (Fig. 1C) indicative of a LOV-bound flavin mononucleotide (FMN) cofactor (see SI Appendix, Fig. S2, for cofactor isolation). BcLOV4 photocycled with rapid thermal reversion kinetics (τ_{off} = 18.5 ± 2 s) (Fig. 1D). However, stable photocycle measurements required in vitro stabilization by high salinity alone (0.5–1 M NaCl) or in combination with glycerol (10%), or immobilization on solid-phase supports (see SI Appendix, Fig. S3, for photocycling summary). In the absence of such stabilization, illuminated BcLOV4 quickly aggregated into turbid solutions of micrometer-scale colloids as measured by dynamic light scattering (DLS) (Fig. 1E). This in vitro photoaggregation was preventable with high stabilization (e.g., 1 M NaCl), reversible with intermediate stabilization (e.g., 0.5 M NaCl), and irreversible in normal-salinity PBS, eventually precipitating from solution (SI Appendix, Fig. S4). This phenomenon was dependent on flavin photocycling, since it was abolished for a photochemically inactive C292A BcLOV4 mutant. This mutant is still a holoprotein, but the C292A mutation prevents the formation of the critical cysteinyl-flavin photoadduct (23) that initiates canonical LOV signaling, such that it mimics a permanently dark-adapted protein even in the presence of blue light.

Rapid Membrane Localization in Cells in Response to Blue Light. Light-activated aggregation has not previously been reported for LOV proteins, although oligomerization into photobodies is known among natural phytochromes, cryptochromes, and their engineered optogenetic variants (24, 25). Thus, to functionally probe whether BcLOV4 forms photobodies in cells, BcLOV4 was visualized by fluorescence microscopy when heterologously expressed in mammalian cells (Fig. 2), which we used because B. cinerea is pathogenic and less genetically tractable than HEK cells. Cells expressing 3×-FLAG-tagged BcLOV4 were fixed in blue light or the dark, and then stained with fluorescent dye-labeled anti-FLAG monoclonal antibody. To our surprise, BcLOV4 did not primarily form photobodies in cells but instead localized to the plasma membrane in a blue light-dependent manner (Fig. 2B). Both the dynamic membrane localization in cells and in vitro photoaggregation were also observed with the homolog from the black yeast Cypellophora europaea (hereon called CeRGS) (GenBank accession number ETN36999.1) (SI Appendix, Fig. S5). Thus, importantly, the in vitro and cellular phenomena are likely general to the class of RGS-LOV proteins. Due to low purified recombinant protein yield and heterologous expression levels of CeRGS in cellular assays, data reported hereafter focus on BcLOV4.

To determine the dynamics of this translocation process, mCherry-tagged BcLOV4 variants were directly visualized, using cotransfected isoprenylated GFP as a plasma membrane marker in HEK cells.

Fig. 1. Bioinformatics annotation and photochemical competence of BcLOV4. (A) Secondary-structure conservation across 66 candidate RGS-LOV-DUF proteins, where height represents information content at a given position, in bits. Gray scale, bit score in fifths. (B) Consensus secondary-structure prediction and domain architecture of BcLOV4, from JPred, phyre2, PSI-PRED, and i-TASSER (secondary structures), IUPRED (disorder), Helixplot (amphipathic helices), and Pfam hidden Markov models database (domains, HMM = match in database). (C) Representative flavin photocycling of BcLOV4 stabilized by 1 M NaCl and 10% glycerol to prevent photoinduced aggregation, measured by absorbance spectroscopy, illumination, 15 mW/cm^2; λ = 455 nm. Time indicates postillumination recovery period. (D) Recovery kinetics monitored at λ = 450-nm absorbance (A450). Black, exponential fit. Gray, mean ± SD (n = 3). (E) In vitro aggregation of BcLOV4 in direct response to blue light. The C292A mutant is unable to form a covalent cysteinyl-flavin photoaduct and is thus photochemically inactive. Illuminated samples become turbid but can be stabilized by high-salinity and/or molecular crowding agent. Illumination, 15 mW/cm^2; λ = 455 nm. Particle size by DLS (mean ± SD).

Here, we report that these cytosolic LOV proteins dynamically and reversibly associate with the plasma membrane by directly light-regulated and high-affinity binding to anionic phospholipids, as revealed through a combination of bioinformatics, computational protein modeling in Rosetta, in vitro structure-function studies with purified recombinantly expressed protein, and optogenetic assays in multiple eukaryotic heterologous expression systems. The photosensory phenomenon was found to be directly coupled to flavin photocycling and is likely general across RGS-LOVs. This study establishes a significant signaling mechanism relevant to natural photoceptors, and broadly applicable to single-component optogenetic tools for dynamic membrane localization.
The functional kinetics of membrane association and undocking measured by live-cell imaging were fast (HEK $\tau_{on} = 1.11$ s, $\tau_{off} = 89.1$ s) (Fig. 2 B–F). The membrane association was on the timescale of diffusion to the inner leaflet (Fig. 2 C and E) (~0.7–1.6 s; see Materials and Methods for timescale estimate determination), and such kinetics is indicative of a high-affinity interaction between BcLOV4 and its membrane target. We hypothesized that such a light-switched interaction could occur directly between BcLOV4 and membrane lipids, because the photosensory signal-transmitting J-helix linker is fused to a polybasic amphipathic helix (AH1 in Fig. 1A, from residues 403–416) similar to those involved in membrane association in other systems (26–28).

**Directly Light-Regulated and High-Affinity Interaction with Anionic Phospholipids.** Initial protein–lipid overlay screening assays suggested that BcLOV4 bound anionic lipids but not zwitterionic ones, but this assay tests for headgroup interactions without recapitulating a membrane interface. Thus, to further test for a direct protein–lipid interaction with a more realistic membrane target (Fig. 3), we created droplets of water-in-oil (w/o) emulsions (29) containing purified recombinant BcLOV4-mCherry in the dispersed/aqueous phase, and phospholipid monolayers at the droplet interface to emulate the plasma membrane inner leaflet (Fig. 3A). Artificial membranes were composed of the zwitterionic phosphatidylcholine (PC) mixed with anionic phospholipids of varying concentration and headgroup charge density.

These droplet assays allowed for complete control over illumination conditions and membrane compositions without complications introduced by the presence of other proteins. The facile customization and ability to multiplex on an automated fluorescence microscope made the system highly useful for screening and cross-validating other methods for establishing binding interactions, like surface plasmon resonance (SPR). As seen in 20% phosphatidylserine (PS)-containing emulsions of similar PS composition to mammalian membranes (Fig. 3 B and C), illuminated BcLOV4-mCherry primarily localized to the phospholipid interface, instead of aggregating as observed in lipid-free bulk solution experiments. Conversely, BcLOV4 formed colloids in lieu of binding pure zwitterionic PC interfaces with only positively charged headgroups. Both light-activated localization and aggregation within the aqueous compartment diminished as salinity increased (SI Appendix, Fig. S6), suggesting an electrostatic basis for these phenomena.

Localization was diminished in blue light for the photochemically inactive C292A mutant (32), confirming direct coupling of the phenomenon to signaling initiation by flavin photocycling as opposed to an unknown blue light interaction (Fig. 3D). Conversely, localization was persistent in the absence of illumination with a constitutively active Q355N mutant that structurally mimics the signal-transducing conformation of the LOV J-helix linker region as if it were in a permanently lit or active signaling state, even in the absence of illumination (30–32) (Fig. 3D). The constitutively active mutant also retained the BcLOV4 binding preference of light-activated wild-type BcLOV4 for net anionic lipids over purely zwitterionic PC interfaces. Thus, optical activation of the interaction with membrane phospholipid is consistent with known structure–function determinants of LOV signaling with respect to flavin photocycling and signal transmission via the J-helix.

Dissociation constants for BcLOV4 binding to immobilized liposomal bilayers were next measured by SPR. The measurements were made with the photochemically inactive C292A and constitutively active Q355N mutants, since controlled illumination within the instrument was not possible. BcLOV4-mCherry variants were used for SPR assays both to maintain consistency with droplet assays, and for improved solubility and protein yield. The BcLOV4 constitutively active mutant affinity for 20% PS liposomal bilayers was $K_{D_{Q355N}} = 130$ nM, or >20-fold enhanced vs. the photochemically inactive mutant $K_{D_{C292A}} = 3.2$ μM (Fig. 3E). Thus, consistent with biophysical inferences from cellular kinetics, BcLOV4 indeed
Fig. 3. In vitro binding to anionic membrane lipids. (A) Schematic of BcLOV4 in lipid-stabilized w/o emulsions. (B) Fluorescence micrographs of wild-type BcLOV4 fused to mCherry. Translocation to the inner leaflet-like interface is observed with increasing anionic PS content, but not with purely zwitterionic PC interfaces. (C) Phospholipid interface binding curves, calculated as the membrane interface/dispersed phase ratio (normalized) of BcLOV4 in the light and dark. n = 20–75 droplets; error, SEM. (D) Constitutively active BcLOV4 Q35SN structurally mimics the photoactivated signaling state, is localized to the interface in the dark, and retains its preference for net anionic phospholipids over zwitterionic ones. The photochemically inactive C292A mutant cannot form a covalent cysteinyl-flavin photoadduct and remains in the aqueous dispersed phase even upon illumination. (E–G) Fluorescence micrographs of wild-type and mutants in lipid-stabilized w/o emulsions. A constitutively active protein (with 1x PBS, 1x PBS, and 1x PBS) possesses a high-affinity light-switched interaction with anionic phospholipids. Binding increased with total anionic content (with PS, Fig. 3F), but there were minimal differences between phospholipids of different headgroup charge density under conditions of matching total charge (Fig. 3G). Thus, BcLOV4 membrane binding is charge dependent but nonspecific to headgroup identity, unlike the well-established preference of pleckstrin homology (PH) domains for certain phosphatidylinositol phosphates (PIPs) (33), or similar lactadherin-C2 domain-specificity for PS (34).

Key Structure–Function of the Dynamic Protein–Lipid Interaction. Having identified the light-switched interaction partner as an anionic phospholipid, we next sought to determine the protein binding site (Fig. 4) and focused on the polybasic amphipathic helix in the linker region that is largely conserved among the fungal homologs (AH1) (Fig. 4A). In BcLOV4, this helix possesses a conserved “FFK” motif (residues 412–414) found in membrane-interacting amphipathic helices of Bcl-2-associated death promoter (BAD) (35), kinase suppressor of RAS (KSR) (36), and cepocin anti-microbial peptides (37), and a “FFK” sequence (residues 408–410) found at the membrane interface of the M2 proton channel of influenza A [Protein Data Bank ID code 2rlf]. In such motifs, aromatic side chains putatively insert into the phospholipid bilayer, while the proximal lysine side chains electrostatically bind anionic lipids enriched in the inner leaflet without great headgroup specificity (27) (Fig. 4B).

Thus, candidate phenylalanine and tyrosine residues within this region were mutated to alanines (Fig. 4C–E). The BcLOV4-AH1 mutant (amphipathic helix mutant) photocycled similarly to wild-type protein (SI Appendix, Fig. S3). In SPR assays, the BcLOV4-AH1 constitutively active mutant showed a 10-fold reduction in affinity, $K_{d_{\text{AH1-Q35SN}}} = 1.4 \mu M$, for 20% PS liposomal bilayers (Fig. 4C), providing evidence that light-induced exposure of the specific lipid-binding motifs drives membrane association. BcLOV4-AH1 also showed reduced binding to anionic phospholipids in droplets of w/o emulsions, and largely remained in the aqueous compartment/dispersed phase when illuminated with blue light (Fig. 4D and E). The downward shift in the droplet-based phospholipid interface binding curves of the AH1 mutant from wild-type levels confirmed the direct lipid-binding roles of the aromatic side chains in the FFK and FFK motifs.

In vitro truncation analyses were performed to establish the relative contributions of the N-terminal and C-terminal domains in signal transmission from the LOV blue light sensor to the lipid-binding regions (Fig. 4F–H). mCherry-fused RGS-truncated BcLOV4Δ1–240 or “LOV-DUF,” aggregated in the absence of lipids in the dark (32 ± 43 nm by DLS) and exhibited an upward shift in the phospholipid interface binding curve vs. full-length (both as photochemically inactive mutants; Fig. 4G). Deletion of the unstructured N terminus alone, BcLOV4Δ1–96, had no such effects. These data suggest that, first, the RGS domain serves an inhibitory role when dark-adapted and, second, that the LOV-DUF alone is sufficient for membrane association. While C-terminal truncations of BcLOV4 (i.e., ADUF) were insoluble, the isolated DUF region, which included the putative lipid-interacting AH1, was soluble as an in vitro refolded product (SI Appendix, Fig. S7). The DUF bound anionic phospholipids in protein–lipid overlay assays (Fig. 4H), further implicating the region C-terminal to the LOV sensor in lipid binding. In totality, these findings suggest that BcLOV4 signal transmission is mediated...
by light-induced structural rearrangements that expose a critical polybasic amphipathic helix at the LOV-DUF linker that is inhibited by the RGS domain in the dark (Fig. 4f).

**Blue Light-Dependent Membrane Localization in Fungus.** To determine whether the fungal-derived BcLOV4 associates with membranes in a blue-light-inducible manner in fungus, dynamic localization assays were performed in *Saccharomyces cerevisiae* yeast (Fig. 5). Such confirmation of the photosensory signal in fungal cells is important because they possess high cytoplasmic salinity and anionic membrane lipid content (38–40) that may influence the electrostatic interaction based on the binding studies here (Fig. 3f and SI Appendix, Fig. S6).

BcLOV4-mCherry in yeast indeed translocated from cytoplasm to plasma membrane (Fig. 5A) in response to blue light and in a dark-reversible manner in confocal microscopy analysis of agar-immobilized transformed cells. The measured kinetics in yeast (y0 = 1.20 s, t0ff = 84.9 s) (Fig. 5B and C) were similar to those measured in mammalian cells, although one should note that the similarity in dissociation timescales between the two eukaryotic expression systems here may be purely coincidental, given the electrostatic differences in cellular milieu. Membrane association kinetics on the timescale of intracellular diffusion to membrane (~0.5–1.0 s; see Materials and Methods for determination of estimates) further confirmed the high-affinity membrane–lipid interaction. Thus, RGS-LOV photosensory signal transmission was consistent across all contexts studied, from pure in vitro systems to fungal and mammalian expression systems.

**Discussion**

Our cumulative findings suggest a photosensory signal transduction mode by RGS-LOV of rapidly blue light-inducible, and reversible, membrane association mediated by electrostatic interactions with anionic phospholipids. While other membrane-binding proteins contain PAS domain sensors (41, 42) related to LOV domains, such as PhoQ, Aer, and LuxQ (43, 44), these are ligand-regulated transmembrane proteins unlike the cytoplasmic photoreceptors here reported. We underscore that this signaling mode was not anticipated from hidden Markov model-based bioinformatics searches for conserved domains, which found nothing C-terminal to the LOV sensor. De novo secondary-structure and Rosetta (45) structural predictions suggest that the DUF in this region may adopt a PAS-like fold with antiparallel β-sheets (SI Appendix, Fig. S7D), and thus it is possible that the LOV–DUF interaction is an evolutionarily conserved PAS/PAS interaction as observed in other systems. It should be noted that other lipid interaction sites may exist beyond the critical amphipathic helix between the LOV and DUF domains. A future high-resolution structure of the lipid-bound state will greatly inform the proposed biophysical model, as well as conclusively determine whether the DUF is indeed a PAS domain.

Membrane localization is a known prerequisite for the canonical GAP activity of RGS proteins (46, 47). Therefore, it is plausible that the signal transmission mode proposed in Fig. 4f—and the photosensory response that was consistently observed across in vitro and cellular systems (including in yeast) and across different RGS-LOV proteins (BcLOV4 and CeRGS)—serves to regulate...
interactions with cognate Gα proteins at the membrane in-host. Little is known about the photobiological role of BcLOV4 (18), its interactions with the three Gα proteins of Botrytis (48), and the physiological roles of the latter. However, the isolated (or truncated) RGS domain of MoRGS5 (GenBank accession number EHA46884.1), the RGS-LOV of *Magnaporthe oryzae* (rice blast fungus), does interact in yeast two-hybrid assays with its cognate Gα protein, MagB (17), which is involved with hydrophobic sensing and plant infectivity (49, 50). Genetic knockout of MoRGS5 (ΔMoRGS5) results in increased intracellular cAMP levels, further implicating a role for RGS-LOV proteins in cell signaling (17, 18). ΔMoRGS5 strains, however, exhibit no pronounced organism-level phenotypic difference from wild-type strains. Thus, while a conclusive photobiological role has yet to be established for RGS-LOV proteins (51), which of note have not been shown to be photochemically active to date, the findings here provide a potential biophysical mechanism by which they may affect fungal physiology: through light-regulated and reversible membrane association of a Gα-interacting photoreceptor to fine-tune Gα-dependent cAMP signaling (18).

In an applied context, BcLOV4 also contributes a useful single-component optogenetic system for photoinducible membrane localization that is compatible in yeast and mammalian expression systems. Its translocation kinetics was apparently limited by diffusion in a cellular context and thus approaches a practical limit for rapid optogenetic membrane localization. Unlike the indirect membrane binding of optogenetic tools that rely on heterodimerization between cytosolic and membrane-bound partners (52, 53), BcLOV4 as a single-component system is insensitive to heterogeneity in relative expression level tuning of two components, and is more facile in transgene delivery.

A common goal in optogenetics, and a common motivation for establishing fundamental structure–function that may beget new molecular engineering principles for creating better protein tools, is the identification of photoreceptor mutations that confer beneficial kinetic properties. For example, lengthening the photocycle of the sensor may extend the active signaling duration across the whole protein in some cases, thereby decreasing the stimulation fluence/duration required for sustained activity (54, 55). LOV sensor engineering to tune the photocycle, however, does not guarantee concomitant tuning of functional signaling outputs (56), such as membrane localization for BcLOV4. Screening mutations known to alter the photocycle in other LOV proteins (30, 55–57, 61) revealed that a BcLOV4 C258I mutation in the α-helix to β-strand transition (21) impairs membrane residence in HEK cellular assays (HEK $\tau_{mem} = 622.7$ s) (SI Appendix, Fig. S8), as well as the in vitro photocycle of the salt-stabilized mutant ($\tau_{mem} = 586.5$ s). Similar to wild-type protein, membrane localization in this mutant persists longer than its photocycle duration. Given that the membrane undocking time of BcLOV4 is longer than its in vitro photocycle, thermal reversion of the photoexcited LOV may not disrupt all lipid interactions of the membrane-bound state across the whole multidomain protein, resulting in the overall longer time constant for undocking than thermal revision alone. This C258I residue is an interesting candidate for further structure–function studies on how flavin photochemistry couples to the signaling state in diverse LOV. More broadly, the direct readout of membrane localization assays makes RGS-LOV an interesting LOV class for such structure–function studies at large.

In summary, the myriad results presented here establish a photosensory signaling mode by RGS-LOV through a directly light-regulated, reversible, and high-affinity electrostatic interaction between anionic plasma membrane phospholipids and a polybasic amphipathic helix at the LOV interface with its C-terminal DUF. This work highlights the utility of convergent approaches that link bioinformatics, in vitro structure–function, and functional assays in live cells to define a mechanism by which photoreceptors dynamically regulate cellular physiology in response to sensory cues.

**Materials and Methods**

**Genetic Constructs and Protein Expression.**

**Bacterial genetic constructs.** For protein expression, genes fragments encoding for BcLOV4 (GenBank accession number CDD5325.1), *Calyptophora cyphellophora* europa LOV (EHN36999.1), *Marsannina brunnea* LOV (EKD10672.1), *Magnaporthe oryzae* LOV (EHA46884.1), and *Exophila dermatitis* LOV (EHY60593.1) were ordered from Integrated DNA Technologies as gBlocks and assembled by Gibson cloning or PCR assembly. Transgenes were cloned into a pET21/28-derived bacterial expression vector. C-terminal mCherry fusions with a GGGS linker were generated by Gibson cloning. Genetic constructs were transformed into competent *Escherichia coli* (C2984H; NEB Turbo). Mutants were generated by QuikChange site-directed mutagenesis. All sequences were verified by Sanger sequencing.

**Mammalian genetic constructs.** DNA sequence of BcLOV4 was human codon-optimized (Genscript). The C-terminal mCherry fusion was created as described above. The mCherry-free variant with a C-terminal "3xFLAG" tag (Sigma Aldrich) had a GGGS linker. Transgenes were cloned into the pCNA3.1 mammalian expression vector (Invitrogen).

**Yeast genetic constructs.** BcLOV4-mCherry was cloned into a pRS316 yeast expression vector with uracil auxotrophic marker (plasmid #3546; Addgene), and transformed into *S. cerevisiae* (ATCC 201388 strain BY4741) competent cells prepared using a Zymo Research Frozen-EZ Yeast Transformation II Kit. Cells were cultured in uracil dropout medium (Sigma-Aldrich). Recombinant protein expression, isolation, and purification. Recombinant proteins were expressed in *E. coli* BL21(DE3). Cells were shaken (250 rpm) post-induction for 18–22 h at 18 °C in complete darkness, harvested by centrifugation, and dissolved in 50 mL of lysis buffer (50 mM sodium phosphate, 500 mM NaCl, 0.5% Triton X-100, pH 6.5) per liter of harvested culture. Samples were homogenized through a 21-gauge needle, sonicated, and clarified by centrifugation, all at ≤4 °C. His6-tagged protein was affinity-purified by fast protein liquid chromatography (FPLC) (AKTA Basic) on Ni-NTA (GE HiTrap FF) columns in darkness, using a stringent column wash (20–200 mM imidazole linear gradient). Protein was eluted with 500 mM imidazole and buffer exchanged into PBS using PD-10 desalting columns and centrifuged to pellet insoluble debris.

**In vitro refolding.** His6-Gb1-tagged BcLOV4 DUF (∆1–356) was expressed in BL21(DE3) *E. coli*. After lysing cells with a French Pressure Cell (Avestin)
Emulsiflex-C5) and centrifuging, the pellet was resuspended in protein solubilization buffer (50 mM Tris-HCl (pH 8), 500 mM NaCl, 0.5% Triton X-100, 0.5 mM DTT, 6 M guanidine HCl, 2 mM EDTA). After denaturation at 4°C for 5–10 min, supernatant was added drop-by-drop to 500 mL of dilution buffer (50 mM Tris-HCl (pH 8), 500 mM NaCl) over a 2-h period to refold the protein. Protein was concentrated via Amicon stirred cell and FPLC-purified on a Superdex 75 or 200 size exclusion column.

**Eukaryotic Cellular Assays.**

**Mammalian cell culture and transfection.** HEK293T cells were cultured in D10 media and maintained at 37 °C in a 5% CO₂ incubator. Cells were seeded onto collagen-treated or poly-l-lysine-treated glass bottom dishes or into 24-well glass-bottom plates, and transfected at ~20–30% confluence using the TransIT-T293 transfection reagent. Cells were fixed in 4% paraformaldehyde in PBS at 15 min after the addition of the reagent. Yeast strains were immobilized on agarose pads before imaging, as reported by others (63).

**Optical microscopy.** Fluorescence microscopy was performed on an automated Leica DMi6000B fluorescence microscope as described previously (64, 65). When needed, isoprenylated GFP (66) was cotransfected as a plasma membrane marker. After a 5-s-long blue light pulse (15 μW/cm²), BLoV4-mCherry images were collected every 200 ms (membrane association) or 5 s (membrane dissociation). Localization kinetics was measured for single cells by line section analysis in ImageJ and MATLAB. Spinning-disk confocal microscopy was performed as described by others (65). HEK cell fixation and immunocytochemistry. Cells were fixed in 4% paraformaldehyde in PBS for 15 min at room temperature, under dim red light (dark-adapted) or strobed illumination (Mightex; λexc = 455 nm, ≥15 μW/cm², 5 s on/25 s off). Immunocytochemistry analysis of 3×-FLAG-tagged protein was performed by standard methods with Alexa 488-conjugated anti-2×FLAG (5407; Cell Signaling Technology) or anti-3×-FLAG antibody (#8146; Cell Signaling Technology) followed by an Alexa 488-conjugated secondary antibody (#4408; Cell Signaling Technology).

**Diffusion estimates.** The intracellular diffusion constant (Dintra) of BLoV4-mCherry was calculated as 8.88±2 cm²/s, assuming an in vitro dark-adapted hydrodynamic radius of 10 nm measured by DLS and an intracellular viscosity of 2.5 centipoise in mammalian cells (67). The timescale for diffusion to the plasma membrane was considered the time to travel a length of 1.6× the hydrodynamic radius (rdiff) in two dimensions (t = rdiff²/(4Dtra)) (68), assuming a 5–7.5-μm radius for HEK cells. The diffusion timescale in HEK was estimated as ∼0.7–1.6 s. Diffusion timescales in yeast (estimated as t = 0.5–1.0 μs) were calculated similarly assuming a 2- to 3-μm radius and cytoplasmic viscosity of 10 centipoise (69).

**In Vitro Protein Analysis: Nonlipid Interactions.**

**Absorbance spectroscopy and photocycling measurements.** Absorbance scans were measured on a CytoScan UV spectrophotometer. Photocycle kinetics was measured by monitoring the absorbance at 450 nm (A450); after 15 s of baseline measurements, samples were stimulated with a collimated LED (Mightex; 10 μW/cm², 550 nm). Recovery was monitored in the dark. For solid-state photocycling measurements, 40 ng of His6-tagged protein was nutated with 0.5 mg of magnetic Ni-NTA beads (resin 88221; Thermo Fisher) in 400 μL of PBS for 1 h, washed, and resuspended in 200 μL of PBS; flavin fluorescence scans were made on a Tecan Infinite M200 plate reader (λexc = 450; λem = 505), similar to absorbance scans. Protein fluorescence and flavin incorporation determination. Flavin and holoprotein concentration were determined by A450 measurements (fequiv=20 = 12,500 M⁻¹ cm⁻²). To estimate protein concentration from A280 measurements, the optical density contributions of flavin, mCherry, and photoaggregates were subtracted. A450-derived flavin concentration was converted to A280con 

**Bioinformatics.**

Maximum-likelihood phylogenetic tree construction. The tree was constructed by aligning all candidate sequences with MUSCLE, building a phylogenetic tree with PhyML, and rendering a tree with TreeDyn through the phylogeny.fr webserver (www.phylogeny.fr) (74). Taxonomic class assignments were made with the Interactive Tree of Life (iTOL) server (itol.embl.de) (75).

**Secondary-structure modeling and consensus annotation.** Candidate amino acid sequences were submitted individually to I-TASSER (76), I-TASSER (76), Phyre (78), and P3SPRED (79). A consensus secondary-structure prediction was generated by equally weighting α-helix and β-sheet predictions from the four algorithms at every amino acid residue and requiring two of four programs to agree on any given structural element. Amphiarchic helices were predicted with the HelixQuest web server (80).

**In vitro energy minimization modeling in Rosetta.** De novo structural predictions were made with Rosetta (81, 82) on 100 Intel ES-2665 2.4-GHz Xeon processors using the Abinitio Relax protocol. The consensus secondary-structure prediction was used throughout the process to filter out trajectories that were unlikely to converge to the supplied secondary structure. Near-native topologies were identified by determining the most frequently sampled conformations using clustering with rmsd as the distance metric. The lowest energy trajectory of the largest cluster was hypothesized to be the closest approximation of the native structure.

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