

Spectroscopic Characterization of Radical Pair Photochemistry in Nonmigratory Avian Cryptochromes: Magnetic Field Effects in GgCry4a

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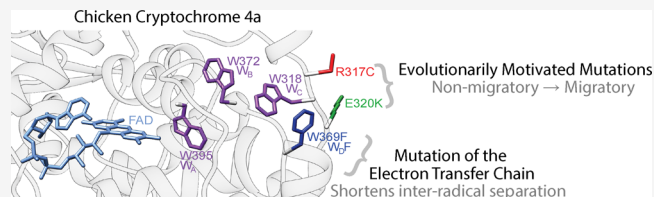
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ABSTRACT: The magnetic compass sensor in night-migratory songbirds is thought to be a flavin-tryptophan radical pair formed by blue-light excitation of the protein cryptochrome-4a (Cry4a) localized in photoreceptor cells in the birds' retinas. The effects of applied magnetic fields on the photochemistry of purified Cry4a from the migratory European robin are well characterized, but it is less clear what, if anything, distinguishes the magnetic responses of the Cry4a proteins from migratory and nonmigratory species. We present here a detailed study of the magnetic sensitivity of Cry4a from the nonmigratory chicken. The wild-type protein is compared with two mutants in which either Arg317 or Glu320, both close to the tryptophan radical, were replaced by the amino acids Cys and Lys, respectively, found in Cry4a from robins and other night-migratory passerines. These sites had previously been identified as probably facilitating the evolution of an optimized magnetic sensor for nocturnal orientation in songbirds. Neither of these mutations was found to affect the reaction kinetics or magnetic sensitivity of the radical pairs, suggesting that any differences in Cry4a between robin and chicken must stem from their ability to transmit magnetic information, for example via protein–protein interactions. In contrast, a Trp → Phe mutation at the end of the tryptophan-tetrad electron transfer chain in both cryptochromes led to a large increase in magnetic sensitivity, suggesting different sensing and signaling roles for the third and fourth tryptophans.



INTRODUCTION

Discovered by Kaptein and Oosterhoff, the radical pair mechanism (RPM) governs the magnetic field-sensitive reaction dynamics of certain chemical reactions, namely those proceeding via short-lived radical pair intermediates.¹ Crucially, it was proposed by Schulten et al. in 1978 that avian magnetoreception, the ability of certain birds to use the Earth's magnetic field for their annual migration, might have its origin in RPM-driven, light-induced reactions, now believed to occur within proteins located in the birds' eyes.^{2–7} More than a decade after Schulten's original proposal, cryptochromes (Cry) were discovered, first in the model plant *Arabidopsis thaliana* (*At*) and later in most kingdoms of life, including animals.^{8–11} As blue-light-sensitive flavoproteins, cryptochromes are closely related both structurally and genetically to their ancestors, the DNA photolyases (PL).^{8–11} Their discovery sparked great interest from zoologists and physical scientists alike, resulting in an array of behavioral animal studies^{4,12–17} and *in vitro* spectroscopic studies on the purified proteins, testing the so-called cryptochrome hypothesis of magnetoreception within the RPM framework.^{18–25}

Observations of magnetic field-sensitive, RPM-driven, light-induced reactions in the cryptochrome/photolyase family of proteins, such as *AtCry1*, *E. coli* photolyase (*EcPL*) and *Drosophila melanogaster* Cry (*DmCry*) have added significant weight to the cryptochrome hypothesis.^{19–21} However, it was not until 2021 that the magnetic field sensitivity of a cryptochrome from a migratory animal, the European robin (*Erithacus rubecula*, *Er*), was demonstrated.¹⁸ A detailed spectroscopic study of *ErCry4a* and two *Cry4a* proteins from nonmigratory species, pigeon (*Columba livia*, *Cl*) and chicken (*Gallus gallus*, *Gg*), drew several important conclusions.¹⁸

- (i) All three wild-type (WT) avian *Cry4a*'s form photo-generated spin-correlated radical pairs (SCRPs) via

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sequential electron transfer between the flavin adenine dinucleotide (FAD) cofactor and tryptophan (W) residues (see Figure 1).

- (ii) In *ErCry4a* WT, electron transfer proceeds along a tetrad of tryptophan residues (which we shall label W_A , W_B , W_C , and W_D in order of increasing distance from the FAD cofactor).
- (iii) The kinetics and yields of the radical pair reaction in all three proteins are sensitive to applied magnetic fields *in vitro*.
- (iv) In *ErCry4a*, replacement of the fourth tryptophan (W_{369} , W_D) by a redox-inactive phenylalanine (F) leads to significantly enhanced magnetosensitivity.
- (v) The magnetic field sensitivity of *ErCry4a* WT seems to differ from that observed for *ClCry4a* WT and *GgCry4a* WT.

Having studied in detail the spectroscopic and magnetic characteristics of *ErCry4a* and several of its mutants,¹⁸ the purpose of the work presented here is to provide an equally detailed, and hence, comparative study of a cryptochrome from a nonmigratory bird, namely the chicken. We expressed and studied mutant proteins which provide crucial links between

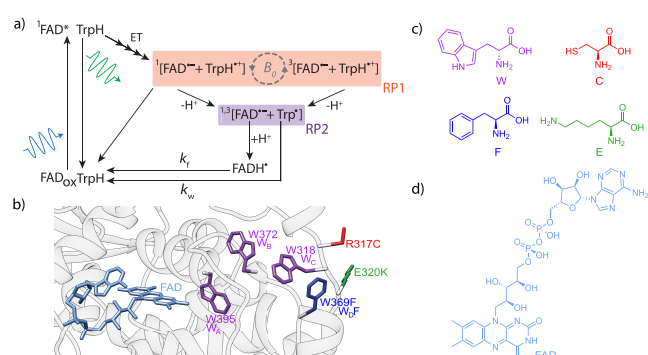


Figure 1. a) Photoreaction scheme proposed for avian cryptochromes. RP1 (orange) formed in an initial singlet state by electron transfer (ET) between FAD (blue in panel b)) and neighboring tryptophan residues (purple in panel b)) can undergo hyperfine driven spin mixing into the triplet state. This mixing is sensitive to an external magnetic field (B_0). Both spin states of RP1 can undergo a deprotonation reaction of the tryptophan radical cation ($\text{TrpH}^{\bullet+}$) to yield a radical pair (RP2, purple) which lacks spin correlation. Recombination of RP1 to the ground state by back-electron transfer is only possible from the singlet state. Magnetic field effects on the efficiency of singlet–triplet mixing (gray dashed arrows) thus impact the quantum yield of RP2 and hence FADH^{\bullet} as well as FAD_{ox} . The magnetic field effect on the latter is detected by field-sensitive fluorescence (green curly arrow) from the $^1\text{FAD}^*$ excited state. Long-lived FADH^{\bullet} and Trp^{\bullet} radicals return to their ground states by independent oxidation and reduction reactions, respectively. The rate constants for FADH^{\bullet} oxidation and reduction are k_F and k_W , respectively. b) Homology model structure (Swiss-model)^{29–33} of *GgCry4a*, with residues 317 and 320 exchanged for their counterparts in *ErCry4a*, R317C and E320K, and residue 369 exchange for phenylalanine, W369F. The model is based on the crystal structure obtained for *ClCry4a*, with the C-terminus removed (PDB entry 6PU0).³⁴ The moieties involved in the electron transfer chain (FAD and tryptophans) are shown in light blue and purple, respectively. Note that only single point mutants, where the mutation is introduced at either position 317, 320 or 369, were studied in this work. c) Structures of the amino acids tryptophan, phenylalanine, cysteine and lysine. d) Structure of FAD.

migratory and nonmigratory birds in an attempt to identify any mutations resulting in different spectroscopic, kinetic or magnetic properties. We have also further developed our experimental procedures to allow for efficient recycling of long-lived radicals using the more biologically relevant oxidant, molecular oxygen (see Section SM.3–7 in the Supporting Information) instead of potassium ferricyanide used in previous work.^{18,21}

The starting point of any such study must be a close inspection of the different species' *Cry4a* sequences, noting any differences involving or neighboring the electron transfer chain.¹¹ The tetrad of tryptophans, which provides the electron transfer chain in *ErCry4a* and *ClCry4a*,^{18,26} is conserved widely across avian species, but not in all. Both the robin and pigeon are known from behavioral experiments to use the Earth's magnetic field.^{4,12,13,27,28} Chickens, by contrast are neither migratory nor homing birds. Herein, we verify the identity of the electron transfer chain that leads to the formation of SCRP in *GgCry4a*.

A detailed phylogenetic analysis of 363 bird genomes, by Langebrake et al., indicated positive selection for cysteine (C) and lysine (K) residues at positions 317 and 320, respectively, in the *Cry4a* sequence of migratory passerines.¹¹ From homology models¹⁸ of the *ClCry4a* crystal structure,³⁴ these residues are thought to be in close proximity to the two distal tryptophans, W318 and W369 (W_C and W_D) of the electron transfer chain.^{18,26} Given the apparent importance of both tryptophan residues to the spin chemistry of these proteins (see points (i) to (v) above), a study of the roles of these cysteine and lysine residues in the photochemistry of *Cry4a* is clearly of interest. To further illustrate the evolutionary conservation of these residues, the amino acid sequences of the *Cry4a* WT proteins from two night-migratory passerines (*ErCry4a* and *Sylvia atricapilla*, *SaCry4a*), and two non-migratory, nonpasserine birds (*GgCry4a* and *ClCry4a*) are provided in the Supporting Information (Figure S2), with the latter two both lacking the conserved residues (C, K) found in the migratory passerines.

Here, we study in detail the effects of single-point mutations of *GgCry4a*, with the arginine (R) and glutamic acid (E) residues in positions 317 and 320 exchanged for their respective counterparts from *ErCry4a* (C and K, respectively). Additionally, we also explore the effect of exchanging the fourth tryptophan residue (W_{369} , W_D) for a redox inactive phenylalanine (F). Figure 1b–d provides a summary of the mutations, nomenclature, and amino acids discussed in this paper. For this study, we chose to make *GgCry4a* “more robin-like” (by means of the R317C and E320K mutations) instead of starting with *ErCry4a* and making it “more chicken-like” (C317R and K320E) because the higher expression yields of the chicken protein allowed a greater range of high-quality measurements than would have been possible with robin *Cry4a*.

RESULTS

All the proteins used in this study were expressed and purified using the procedure for *GgCry4a* WT in Xu et al.,¹⁸ except for some improvements made later and described in the Supporting Information (SM.1 and Table S1). The protocol for expressing and purifying *GgCry4a* W369F (W_D F) has been described in Golesworthy et al.³⁵ Full details for expression of *GgCry4a* R317C and E320K, produced for the first time for this study, are included in the Supporting Information (SM.1

and Table S1). Before any spectroscopic measurements commenced, the diluted protein samples were investigated using native mass spectrometry (SM.2 for experimental details) which confirmed that the proteins were present almost exclusively in their monomeric states and contained the FAD cofactor (Figure S3). Vibrational fine-structure in the UV-visible spectrum in the 300–500 nm region (Figure S4) confirmed that the FAD cofactor was bound within the protein. The absence of any features above 500 nm along with the spectral positions of the vibrational maxima furthermore demonstrated that the proteins contained only the fully oxidized form (FAD_{ox}) of the FAD cofactor.

Electron Paramagnetic Resonance. The first blue-light photosensitive proteins to demonstrate any magnetic field sensitivity were *AtCry1* and *EcPL*.^{19,20} These proteins differ from avian Crys in that the electron transfer cascade, which leads to the formation of a magnetically sensitive SCRPs (termed RP1), proceeds along a triad, rather than a tetrad, of tryptophan residues.^{18,20,24,26,36} However, the identity of RP1 is the same in all proteins, comprising a flavin semiquinone radical (FAD^{•-}) and a tryptophan radical cation (TrpH^{•+}) as shown in Figure 1.^{18,20,24,37} Photoinduced electron transfer to FAD in *DmCry*, *ErCry4a*, and *ClCry4a* proceeds along a tetrad of tryptophans, extending the electron transfer cascade to W_D.^{18,24,26} Electron Paramagnetic Resonance (EPR) played the pivotal role in identifying the terminal radical pair in all five proteins. It has been exploited here again to characterize the SCRPs in *GgCry4a* and its mutants. Experimental and technical details can be found in the Supporting Information (SM.3 and Figure S5).

Briefly, we have applied two EPR techniques common in the study of SCRPs: transient EPR (TrEPR) and out-of-phase electron spin echo envelope modulation (out-of-phase ESEEM).^{38–41} Together, they provide information on radical–radical distances and the identities, spin polarization, and spin correlation of the radicals.

The TrEPR spectra of SCRPs (see SM.3) show distinctive antiphase (e.g., emission–absorption, *e-a*) line-shapes and have been recorded for a number of cryptochromes as well as for *EcPL*.^{18,24,26,42–44} Additional features contain information on the interaction of the electron spins with the magnetic field, with each other and with nuclear spins in the two radicals.^{18,26,42,43,45} Upon blue-light photoexcitation of *GgCry4a* WT, a TrEPR spectrum is obtained whose prevailing antiphase line-shape confirms the formation of a singlet-born SCRPs with the fine structure arising from electron–nuclear spin hyperfine couplings (Figure 2a, purple). The spectra obtained for the two mutants, R317C and E320K, are also shown in Figure 2a and, within signal-to-noise, are indistinguishable from that of the WT protein. In contrast, the TrEPR spectrum of the W369F (W_DF) mutant, lacking the W_D residue, is distinctly different (Figure 2b). The antiphase signature, characteristic of a SCRPs, is still observed but the splittings are now less well resolved. The different spectrum provides a first indication that mutation of W369 (W_D) to F interrupts the redox chain significantly, as previously observed for *DmCry* W394F, *ErCry4a* W369F, and *ClCry4a* W369F.^{18,24,26}

We have simulated the TrEPR data (Figure S8), following the treatment outlined in ref 46 and the full set of simulation parameters is provided in the Supporting Information (Table S2). The *g*-factors and hyperfine couplings used for these simulations correspond well with those published elsewhere for

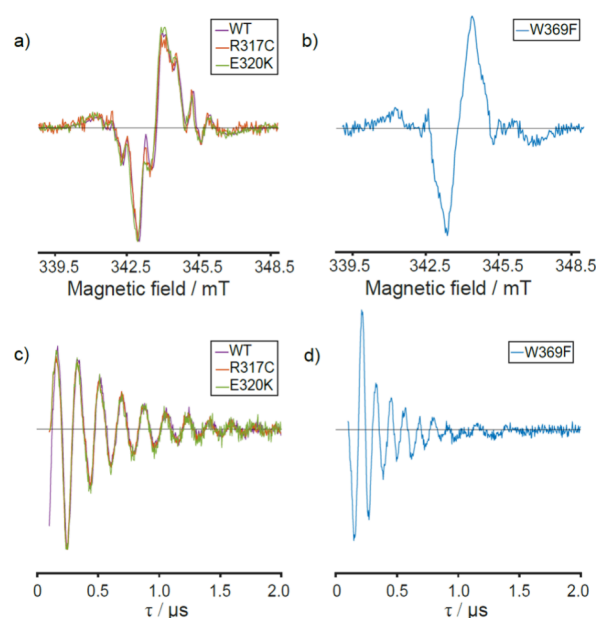


Figure 2. a) X-band (9.75 GHz) TrEPR spectra obtained for *GgCry4a* WT (purple), *GgCry4a* R317C (red) and *GgCry4a* E320K (green) averaged over 0.5–1 μ s after a 450 nm laser flash. b) Same as a) for *GgCry4a* W369F. The positive and negative signals in all four spectra arise from absorptive (a) and emissive (e) transitions, respectively. c) Out-of-phase ESEEM traces obtained for *GgCry4a* WT (purple), *GgCry4a* R317C (red) and *GgCry4a* E320K (green) at Q-band frequency (34 GHz). d) Same as c) for *GgCry4a* W369F. All EPR data shown here were acquired in the presence of 5 mM $K_3Fe(CN)_6$ to promote flavin reoxidation. This was found to have a negligible impact on the TrEPR and out-of-phase ESEEM traces obtained (detailed comparisons in the presence and absence of $K_3Fe(CN)_6$ for both the TrEPR and out-of-phase ESEEM data are shown in Figure S6 and Figure S7, respectively).

FAD and tryptophan radicals.^{24,26} Note that the value of the dipolar coupling, *D*, was fixed during the simulation to the value obtained from the out-of-phase ESEEM experiments.

The simulations of the TrEPR spectra of *GgCry4a* WT and all three mutants are consistent with the formation of a FAD^{•-}–TrpH^{•+} SCRPs upon blue-light photoexcitation (Figure S8). While the simulation parameters for the three proteins with an unperturbed tryptophan tetrad are identical (Table S2), the W369F (W_DF) mutant returns a significantly larger dipolar (through-space) coupling, *D*, between the FAD^{•-} and TrpH^{•+} radicals. The r^{-3} dependence of *D* on the inter-radical separation, *r*, (in addition to the simulation parameters in Table S2) suggests a shortening of the electron transfer chain in W369F ($r \approx 17$ Å) compared to the WT ($r \approx 21$ Å) and R317C and E320K mutants. The difference in the fine structure of the TrEPR spectra of *GgCry4a* WT and W369F is, therefore, mainly attributed to a broadening of the spectrum due to this increase in the inter-radical dipolar coupling (*D*), in addition to a change in the relative orientation of the radicals, based on our simulations (see Figure S8 and Table S2). A truncated electron transfer cascade in *GgCry4a* W369F mirrors the TrEPR results in *DmCry* W394F, *ErCry4a* W369F, and *ClCry4a* W369F.^{18,24,26}

Full clarification of the identity of the terminal electron donor is provided by the gold standard technique for the determination of inter-radical distances in SCRPs, out-of-phase ESEEM. Figure 2c shows the out-of-phase ESEEM data

obtained for GgCry4a WT, R317C, and E320K (purple, red, and green traces, respectively). The traces are, within signal-to-noise, identical, exhibiting clear modulations whose frequency is determined by the interactions between the two electron spins (SM.3). Full details of the out-of-phase ESEEM simulations in Figure 2c and 2d is provided in the Supporting Information (SM.3 and Figure S9), but Table 1 summarises the crucial parameters extracted here: the strengths of the dipolar coupling, D , and the resulting radical separations related by D (MHz) = $-7.8 \times 10^4/[r(\text{\AA})]^3$. For all three proteins containing the full tryptophan tetrad, we obtain radical–radical separations of $21 \pm 0.1 \text{\AA}$.

For comparison, center-to-center separations between FAD and each W residue in the tetrad, as well as a neighboring tyrosine (Y), were extracted from a homology model of GgCry4a WT, based on the crystal structure of ClCry4a (Table S3).³⁴ It is clear that electron transfer in GgCry4a WT, R317C, and E320K, occurs along the entire tryptophan tetrad, as in DmCry WT, ClCry4a WT and ErCry4a WT, and in contrast to the triad of tryptophan residues in AtCry1 and EcPL.^{18,24–26,36}

Finally, Figure 2d shows the out-of-phase ESEEM trace for GgCry4a W369F. The higher modulation frequency is the result of a larger dipolar coupling from the shorter radical separation (18 Å) consistent with a radical pair containing FAD and W318 (W_C , see Table S3).

Transient Absorption Spectroscopy. Having established the identity of the RP1 radicals (Figure 1) in all four proteins, transient absorption (TA) spectroscopy was employed to trace the kinetics of various species produced by monitoring ΔA ($= A_{h\nu} - A_{GS}$) as a function of the pump–probe delay time (see SM.4). $A_{h\nu}$ and A_{GS} are the absorbance of the sample in the presence and absence (ground state absorbance) of a 450 nm photoexcitation beam, respectively.

The four panels in Figure 3 depict the time evolution of the wavelength-resolved ΔA spectra of the four proteins following laser flash illumination at time $t = 0$. Three dominating features are common to the spectral profiles of all four proteins (see also Figure S10a): (1) The negative ground state bleach (GSB) between 400 and 500 nm, matching the absorption profile of FAD_{OX} (with vibronic fine-structure indicating protein-bound FAD); (2) Positive ΔA signals indicative of the formation (and subsequent decay) of $TrpH^{*+}$ and FAD^{*-} radicals (RP1), identified by their characteristic absorbance bands below 400 nm (FAD^{*-}) and above 500 nm (initially mostly $TrpH^{*+}$, with a minor contribution from FAD^{*-}), respectively (cf. Figure S1). These radicals form on a femtosecond time scale beyond the time resolution of the spectrometer.⁴⁷ (3) The subsequent decay of absorption

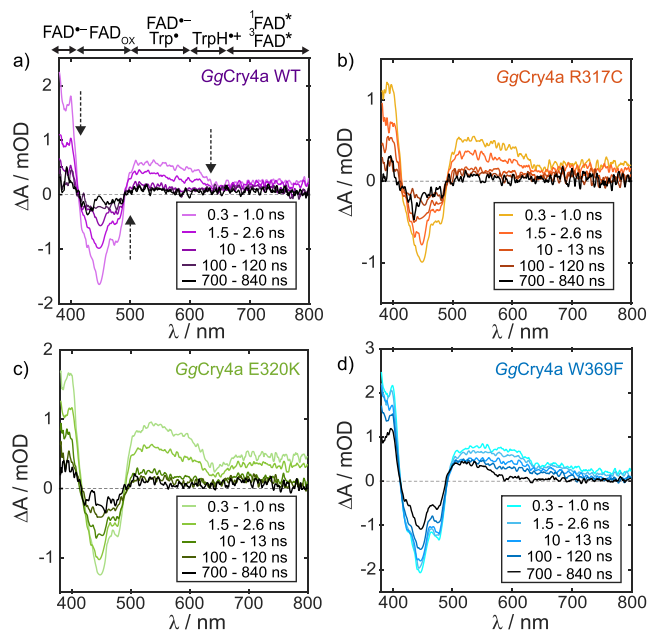


Figure 3. Wavelength-resolved transient absorption spectra, $\Delta A(\lambda)$, at different times following the 450 nm laser pulse at $t = 0$, for the air-equilibrated GgCry4a WT (a), GgCry4a R317C (b), GgCry4a E320K (c), and GgCry4a W369F (d) proteins. All spectra were averaged over the time-windows given in the corresponding legend. Above panel (a), a rough guide is provided indicating which species in the photoreaction scheme (Figure 1) contribute most to the spectra at a given wavelength. The feature between 500 and 650 nm is ascribed to absorption by FAD^{*-} and $TrpH^{*+}$ (see Figure S1). The signals at wavelengths beyond 650 nm are assigned to singlet ($^1FAD^*$, 750–800 nm) and triplet ($^3FAD^*$, 650–720 nm) excited states of FAD with the former decaying more rapidly than the latter. We believe these signals derive from a small fraction of misfolded protein in which electron transfer along the Trp-tetrad is significantly slower than in the native state, allowing $^1FAD^*$ and, after intersystem crossing, $^3FAD^*$ to be detected. Similar signals were observed for European robin Cry4a by Xu et al. (ref 18, Figure S5). The assignment of the 650–720 nm signal to $^3FAD^*$, rather than a radical intermediate, is supported by the absence of a magnetic field effect between 700 and 800 nm in GgCry4a W369F (see Figure S11).

signals on a submicrosecond time scale along with recovery in the GSB region are consistent with the recombination of radical pairs.

The time evolution of the GgCry4a W369F spectrum (Figure 3d) is markedly different from the others and shows a longer-lived ($t > 100$ ns) component in the 500–600 nm region, which is a clear indication of Trp^* formation by deprotonation of the $TrpH^{*+}$ radical. This results in the spin-uncorrelated radical pair, RP2 (see Figure 1).^{18,20,37} This feature is not observed in the TA spectra of the three proteins that have an intact electron transfer chain. The comparatively slow ground state recovery and radical absorbance decay above 500 nm in GgCry4a W369F, is thus a direct consequence of its efficient RP2 formation which, due to $TrpH^{*+}$ deprotonation, cannot recombine directly to the ground state.

The temporal evolution of the ΔA signals for all four proteins, averaged over 500–550 nm, is shown in Figure 4a (and, with a logarithmic time-axis, in Figure S10b). The kinetic profiles in this wavelength region predominantly reflect the decay pathways of FAD^{*-} and $Trp^*/TrpH^{*+}$, and can be reliably fitted with a biexponential model for all four proteins

Table 1. Dipolar (D) Coupling Constants Obtained from Least-Squares Fitting to the Out-of-Phase ESEEM Traces Shown in Figure 2c and 2d^a

Protein	D /MHz	r /Å
GgCry4a WT	-8.37 ± 0.06	21.04 ± 0.05
GgCry4a R317C	-8.46 ± 0.07	20.97 ± 0.06
GgCry4a E320K	-8.51 ± 0.07	20.93 ± 0.06
GgCry4a W369F	-13.95 ± 0.14	17.75 ± 0.06

^aThe inter-radical separations were calculated assuming the validity of the point-dipole approximation (see SM.3). The exponential damping, quantified by relaxation time, T_v , and the exchange couplings, J , are provided in Table S4.

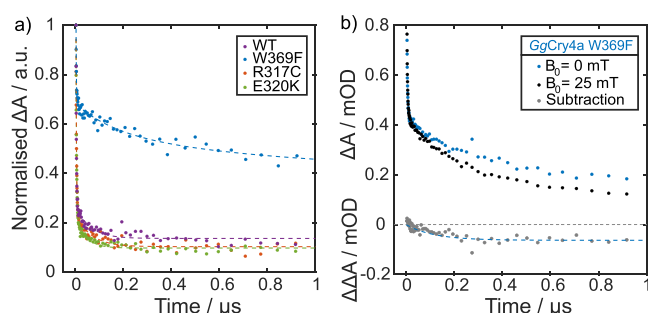


Figure 4. a) Normalized $\Delta A(t, B_0 = 0)$ following the 450 nm laser pulse at $t = 0$, for air-equilibrated GgCry4a WT (purple), GgCry4a W369F (blue), GgCry4a R317C (red), and GgCry4a E320K (green) proteins, wavelength-averaged over 500–550 nm. The ΔA in this region is dominated by the absorbance of $\text{FAD}^{\bullet-}$, $\text{TrpH}^{\bullet+}$, and Trp^{\bullet} . The dashed lines are kinetic fits to the data set of the same color. Fitting parameters are provided in Table 2. b) $\Delta A(t)$ traces of air-equilibrated GgCry4a W369F, averaged over 500–650 nm, obtained in the absence (blue) and presence (black) of a 25 mT external magnetic field. The $\Delta\Delta A(t, B_0)$ obtained from these data using eq 1 is shown in gray, below the $\Delta A(t)$ data. A monoexponential fit to the data including a constant offset (blue dashed line) yields a rise-time of 116 ± 16 ns for the development of the MFE. Further experimental details can be found in SM.4.

plus a constant offset to account for the formation of long-lived photoproducts (Table 2).

The faster (~ 2 ns) components (τ_1) should be interpreted with some caution as they approach the time resolution of the spectrometer. Nonetheless, they are similar in all four proteins and are assigned to the recombination of $\text{FAD}^{\bullet-}$ and the $\text{TrpH}^{\bullet+}$ at site 372 (W_B). This is plausible as the efficiency of the electron transfer steps forming the terminal radical pair in *ErCry4a* WT has recently been shown to decrease as charge separation proceeds along the tryptophan tetrad.⁴⁷ The equivalent decay in the *ErCry4a* W318F (W_C F) mutant, also exhibited a dominant lifetime of 2.5 ns.¹⁸

The longer kinetic decay component (τ_2) can be assigned to $\text{FAD}^{\bullet-}$ and $\text{TrpH}^{\bullet+}$ recombination involving the terminal tryptophan in each protein. As expected from the data in Figure 3d, τ_2 is significantly larger in GgCry4a W369F compared to the other three proteins. The higher yield of species living beyond 1 μs reflects the efficient formation of RP2 in the GgCry4a W369F mutant.

The time constants for the decay of $\text{FAD}^{\bullet-}$ and $\text{TrpH}^{\bullet+}$ as well as the FAD_{ox} recovery in GgCry4 W369F may be estimated from the 370–400 nm, 600–650 nm, and 450–480 nm spectral regions, respectively (Figure S12a–c). This allows

Table 2. Lifetimes Obtained for All Four Proteins from Kinetic Fitting of the ΔA Decay Curves, Averaged over 500–550 nm, Shown in Figure 4a^a

Protein	τ_1/ns	τ_2/ns
GgCry4a WT	1.5 ± 0.5	60 ± 4.9
GgCry4a R317C	2.0 ± 0.5	64 ± 5.2
GgCry4a E320K	1.9 ± 0.5	40 ± 2.4
GgCry4a W369F	1.7 ± 0.5	299 ± 15

^aAll kinetic decays were fitted to bi-exponentials, including a constant offset to account for long-lived photoproducts. The uncertainty in τ_1 is dominated by the pump pulse duration (1 ns) and that of τ_2 is calculated from the error in the fit (see SM.3 for further details).

an estimate of the $\text{FAD}^{\bullet-}$ lifetime ($\tau_2, \text{FAD}^{\bullet-}$), the recovery time of the GSB ($\tau_2, \text{FAD}_{\text{ox}}$), and lifetime of W318 (W_C) deprotonation ($\tau_2, \text{TrpH}^{\bullet+}$). All kinetic traces can be fitted to a biexponential decay plus a constant offset (Table S5). The short (~ 2 ns) and long (275 ns, 193 ns, 270 ns, respectively) time constants agree well with those in Table 2. As both recombination and deprotonation time scales agree reasonably with the rate of GSB recovery, it appears that deprotonation of $\text{TrpH}^{\bullet+}$ and recombination of RP1 occur on similar time scales, and we assign the 299 ns component in Table 2 to a combination of these processes. These assignments are corroborated by the individual species' kinetics of GgCry4 W369F (Figure S13) which clearly show the formation of Trp^{\bullet} from $\text{TrpH}^{\bullet+}$ in conjunction with a partial decay in $\text{FAD}^{\bullet-}$ and GSB recovery, all on a time scale of ca. 200 ns (τ_2 in Table S6).

Transient absorption spectroscopy has also been applied to study the field sensitivity of the radical pair dynamics. TA spectra were recorded in both the absence ($\Delta A(t, B_0 = 0)$) and presence ($\Delta A(t, B_0)$) of a static magnetic field, with the field effect quantified as,

$$\Delta\Delta A(t) = \Delta A(t, B_0) - \Delta A(t, B_0 = 0) \quad (1)$$

As the proteins used in this work form singlet-born radical pairs (Figure 1) the application of fields greater than ~ 1 mT (the effective hyperfine coupling) impedes singlet–triplet mixing, leading to an enhanced recombination yield and yield and lower radical concentrations.^{48–50} $\Delta\Delta A(t)$ is, therefore, negative for a singlet-born radical pair.^{19,20,35}

Figure 4b shows the time-resolved $\Delta A(t, B_0 = 25$ mT) (blue) and $\Delta A(t, 0$ mT) (black) traces for GgCry4a W369F, averaged over 500–650 nm as well as their difference ($\Delta\Delta A(t)$, gray). Three features of these temporal profiles are noteworthy. First, the application of a 25 mT field indeed leads to a significant decrease in the radical absorption over time. Second, this suppression in radical concentration develops after the sharp initial drop (τ_1) in radical absorbance, and on a time scale comparable to τ_2 ($\tau_{\Delta\Delta A} = 116 \pm 16$ ns). This is not surprising as the fields applied here (25 mT, corresponding to 700 MHz electron Larmor frequency) have insufficient time to affect spin-mixing processes on the shorter ($\tau_1 \sim 2$ ns) time scale. Third, the field effect plateaus after ca. 400 ns indicating formation of a photoproduct downstream of RP1 with a lifetime exceeding 1 μs (RP2).

No field effects were observed by TA in the proteins with intact tryptophan tetrads. This is explained by the fact that the majority (>80%) of the radicals produced in these proteins decay within 5 ns, and the recombination of the terminal tryptophan (W_D) radical is also fast (40 ns). To characterize the field effects, we have utilized other optical techniques with superior sensitivity.

Broadband Cavity-Enhanced Absorption Spectroscopy. We have developed broadband cavity-enhanced absorption spectroscopy (BBCEAS) to measure spectrally resolved magnetic field effects as low as 1% ($\Delta\Delta A \approx 1 \mu\text{OD}$).^{21,51} A sample contained within an optical cavity is continually excited at 455 nm and changes in the absorbance monitored using a pseudocontinuous broadband supercontinuum probe beam. Field effects are measured with on/off magnetic fields. BBCEAS boasts a much-improved sensitivity over TA for accumulation of data on longer-lived photoproducts. As in TA spectroscopy, the ΔA and $\Delta\Delta A$ signals quantify the effects of

photoexcitation and applied magnetic fields on the cryptochrome samples, respectively.^{18,21}

Figure 5a shows the evolution of the ΔA spectrum for GgCry4a WT, under constant illumination conditions. The GSB and radical absorbance both increase over the first 10 s of illumination as the photostationary equilibrium of the system is driven toward the photoproduct(s). Global spectral fitting of the kinetics (see Figure 5a inset and SM.8) confirm that the dominant absorbers on this time scale in the 500–750 nm region are the $FADH^{\bullet}$ and Trp^{\bullet} radicals that comprise RP2. The species-resolved accumulation kinetics in Figure 5b shed light on the subsequent fate of the radicals: $FAD^{\bullet-}$ protonation on a ~ 1 s time scale leads to steady accumulation of $FADH^{\bullet}$ over the illumination period, while the Trp^{\bullet} concentration rises quickly before reaching a steady-state for $t > 1$ s. Similar behavior is observed in all four proteins studied (see Figure S14).

Figure 6a shows the wavelength-resolved, time-averaged $\Delta\Delta A$ spectra ($B_0 = 30$ mT) of all four proteins as recorded by BBCEAS. All exhibit negative MFEs across the 500–750 nm wavelength region confirming the formation of singlet-born SCRP in each case. Under the O_2 -reoxidising conditions employed here, all $\Delta\Delta A$ spectra were dominated by $FADH^{\bullet}$ with a smaller Trp^{\bullet} contribution (see Figure S15).^{51–53} On this longer time scale, the 30 mT magnetic field has the largest

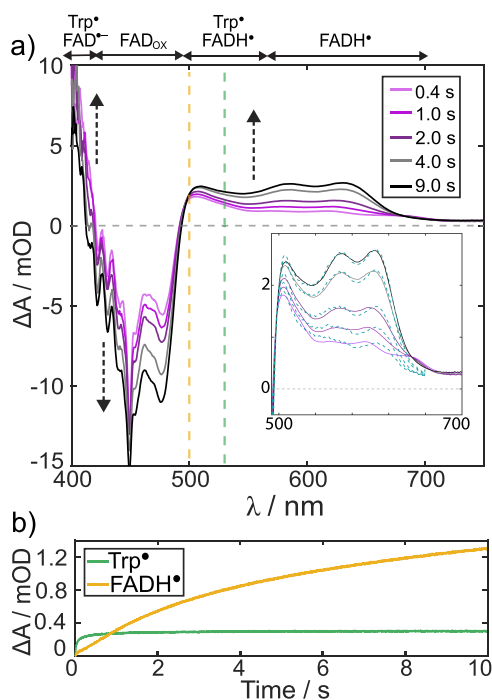


Figure 5. a) ΔA spectra obtained by BBCEAS for GgCry4a WT at the different time windows indicated in the legend. The yellow and light green dashed lines give the wavelengths at which the kinetics profiles are extracted in b). Dashed black arrows indicate the time evolution of the spectra. As in Figure 3a, a rough guide is provided indicating which species in the photoreaction scheme (Figure 1) contribute most to the spectra at a given wavelength. Inset: Spectral fit (teal dashed line) obtained by global analysis (see SM.8) of the BBCEAS ΔA spectra for GgCry4a WT (same as Figure 5a), using the literature spectra of FAD_{ox} , $FAD^{\bullet-}$, $FADH^{\bullet}$, and Trp^{\bullet} as basis spectra. b) ΔA accumulation kinetics of $FADH^{\bullet}$ at 500 nm (yellow) and Trp^{\bullet} at 530 nm (light green) extracted by global fitting of the ΔA spectra (see inset and dashed lines) over the duration of the illumination period.

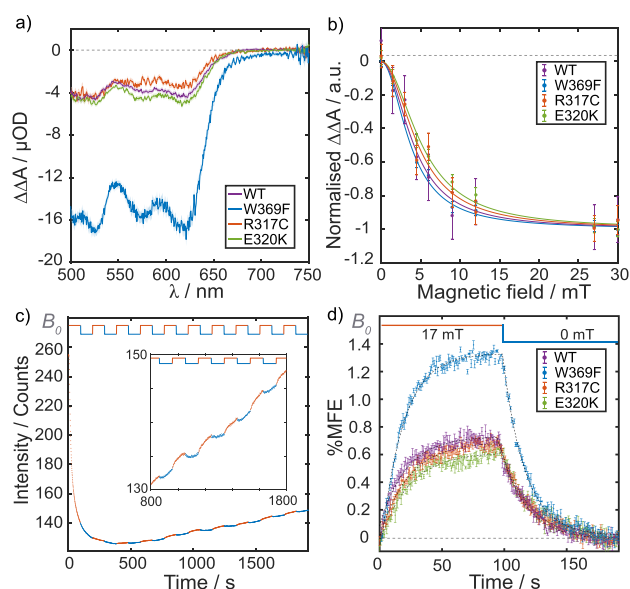


Figure 6. a) $\Delta\Delta A$ spectra obtained by BBCEAS on all four proteins, averaged over 10 s, using an applied magnetic field strength of 30 mT and a field switching period of 200 ms. b) Magnetically altered reaction yield curves (solid circles), obtained by BBCEAS as in a), with varied applied magnetic field strength. The data were fit to a single Lorentzian (solid lines). c) Time evolution of the fluorescence intensity of GgCry4a WT, obtained by CM, averaged over the entire field of view ($100 \times 100 \mu m^2$), following onset of illumination (448 nm) at $t = 0$. A magnetic field with a period of 191.8 s was switched between 0 and 17 mT during the acquisition leading to modulations (see inset) in the fluorescence intensity during the field-on (red fluorescence segments) and field-off (blue fluorescence segments) periods. The applied magnetic field (B_0) is shown schematically above the data, again with field-on and field-off periods denoted by red and blue. d) Time-averaged %MFE for all four proteins, obtained by CM, calculated using eq 2, from background subtracted data (Figure S16). The black dashed lines are single-exponential fits to the mean %MFE over both the field-on and field-off periods. The time constants (τ_{eMFE}) reported in Table 3 were obtained from these fits.

effect on the concentration of the neutral flavin radical, $FADH^{\bullet}$, formed by protonation of $FAD^{\bullet-}$ following electron transfer.

The GgCry4a WT, R317C, and E320K proteins all exhibit very similar MFEs ($\Delta\Delta A_{500-630 \text{ nm}} \approx -4 \mu OD$). The field effect on the W369F mutant, is substantially larger ($\Delta\Delta A_{500-630 \text{ nm}} \approx -16 \mu OD$), consistent with the differences between ErCry4a WT and ErCry4a W369F reported previously.^{18,35}

Figure 6b shows the wavelength-averaged (500–630 nm) $\Delta\Delta A$ signal for all four proteins as a function of the strength of the applied static magnetic field. All four species exhibit a Lorentzian-like dependence on B_0 , characteristic of hyperfine coupling-driven singlet–triplet mixing in a SCRP. These magnetically altered reaction yield (MARY) curves are commonly characterized by their half width at half-maximum, a parameter known as $B_{1/2}$. Broadly, this parameter reflects the extent of hyperfine coupling within the SCRP via the Weller formula, which, for an FAD-Trp SCRP, predicts a $B_{1/2}$ of approximately 3 mT.⁵⁴ Wong et al. have estimated an improved lower bound for $B_{1/2}$ in cryptochromes under our *in vitro* conditions of 2.46 mT.⁵⁰ All four species exhibit similar values of $B_{1/2}$ (≈ 4 mT, Table S6). This is consistent with an $FAD^{\bullet-}/TrpH^{\bullet+}$ SCRP with some spin relaxation impacting the

dynamics during the radical pair lifetime, leading to a larger $B_{1/2}$ than the idealized values mentioned above.^{54,55} In summary, the sensitivity of BBCEAS helps confirm the formation of magnetically sensitive radical pairs in all four proteins with very similar $B_{1/2}$ values. Furthermore, mutations near the tryptophan chain have little detectable effect on the size of the MFE and removal of W_D leads to a significant increase in the size of the field effect, mirroring earlier results for *ErCry4a*.¹⁸

Confocal Microscopy. While EPR and the direct absorption techniques above probe the photoproducts (right-hand side of the photoreaction scheme in Figure 1a), Confocal Microscopy (CM) yields complementary information on the fluorescence of the flavin excited state. Photoexcitation of FAD is followed by rapid (< ns) intramolecular electron transfer along the tryptophan chain resulting in a very small fluorescence quantum yield.^{56,57} However, pseudocontinuous photoexcitation of the sample accumulates ground state FAD_{OX} that has undergone repeated RP singlet–triplet mixing cycles. Any magnetic field effect on RP1 directly impacts the FAD fluorescence intensity.^{58–60} Combined with sensitive fluorescence detection, this makes CM a powerful method for studying MFEs on cryptochromes.^{60,61}

Following the approach described in ref 60, protein samples were subject to continuous 448 nm (raster-scanned) laser excitation (see SM.6). An external magnetic field was switched on and off repeatedly during the acquisition to probe the MFE on the sample fluorescence. Importantly, increased RP1 recombination arising from an applied magnetic field increases the concentration of ground state FAD_{OX} , and, hence, the observed fluorescence. This approach to field-effect detection of a flavoprotein was first demonstrated with *AtCry1*.^{58–60} Figure 6c depicts the CM-detected fluorescence intensity of *GgCry4a* WT, averaged over the entire field of view, as a function of time following the onset of 448 nm photo-illumination at $t = 0$. The fluorescence intensity exhibits an initial sharp drop ascribed to photobleaching of the FAD_{OX} cofactor. It then slowly increases again as a result of two processes: photoinduced degradation and diffusion of fresh protein into the field-of-view, with the former process dominating at long times.^{58,60} The former process results in the release of unbound FAD, which, lacking a nearby electron donor, exhibits a much increased fluorescence quantum yield (ca. ten times higher relative to protein-bound FAD).^{56,62} Superimposed on the time evolution described above is a shallow modulation of the fluorescence intensity, in synchrony with the switching of an applied static field (Figure 6c, top) between 17 mT (red) and 0 mT (blue).

The background fluorescence was subtracted (Figure S16a–b) to obtain a %MFE, defined as

$$\%MFE = \frac{I_F(B_0) - I_F(B_0 = 0)}{I_F(B_0 = 0)} \times 100 \quad (2)$$

where $I_F(B_0)$ and $I_F(B_0 = 0)$ are the fluorescence intensities in the presence and absence of a magnetic field, respectively.

Close inspection of Figure 6c reveals a consistent increase of the fluorescence intensity with an applied magnetic field. Averaging of several “field on–off” cycles (Figure S16c) gives insight into the dynamics of the field effect evolution. Figure 6d shows the time-averaged %MFE data for all four proteins. As expected for singlet-born radical pairs, the applied magnetic field hinders singlet–triplet mixing leading to more singlet

recombination to the ground state and an increase in fluorescence. The %MFE of all four *Cry4a* proteins displays a delayed response to the application of a magnetic field. This is characteristic of enhanced magnetic field effects (eMFEs), leading to biphasic kinetics of the MFE; a prompt fluorescence (increase) upon magnetic field application followed by a slow rise to a constant value. eMFEs were first studied in the solution phase in ref 58 but have since been reported in both fluorophore-doped protein crystals⁶⁰ and human cells⁶³ (note that the failure of ref 64 to reproduce the latter has been rebutted by the authors of the original study in ref 65). This is, however, the first example of eMFEs in an avian cryptochrome. These eMFEs are similar to those previously reported for *AtCry1* and are consistent with $k_F > k_W$ (see Figure 1).⁵⁸

The eMFEs of all four proteins can be fit well by a single exponential function with a time constant, τ_{eMFE} (Table 3), which is the same for the field-on and field-off steps. All four eMFEs saturate by ca. 50 s after magnetic field switching, with a reproducibly faster τ_{eMFE} for *GgCry4a* W369F relative to the other three proteins, which are very similar to one another.

The three proteins containing a complete tryptophan tetrad exhibit the same %MFEs within signal-to-noise. *GgCry4a* W369F (Figure 6d, blue), by contrast exhibits a significantly more pronounced field sensitivity, paralleling the results from BBCEAS and TA. The trends in the relative magnitudes of the MFEs of all four samples studied in this work are, therefore, reproducible between all spectroscopic techniques employed here, probing complementary parts of the photocycle.

Continuous Blue-Light Illumination. All the methods described thus far rely on the recovery of the system back to the ground state before re-excitation of the flavin leading to formation of the magnetosensitive radical pair, RP1. To test the samples for their behavior under long-term illumination, we exposed the proteins to 1 min of light of 450 nm (ca. 25 W m⁻²) and followed their recovery using UV–visible spectroscopy. Figure 7a (purple line) shows the UV–visible spectrum of *GgCry4a* WT after 1 min illumination. The pronounced 500–700 nm absorption reflects the generation of the stable protonated flavin semiquinone ($FADH^\bullet$).^{18,26}

This assignment is confirmed by a global analysis fit (black dashed line, see SM.8) obtained assuming a single first-order recovery from $FADH^\bullet$ (dashed blue spectrum) to FAD_{OX} (solid teal spectrum). The blue and teal spectra are decay associated spectra (DAS) obtained by the global analysis procedure. As with other avian cryptochromes, these systems are efficiently photoreduced under moderate light intensity.^{18,26}

Figure 7b shows the recovery kinetics of $FADH^\bullet$, obtained at 298 K (25 °C) by averaging over the 605–615 nm region. The relative $FADH^\bullet$ radical yield is larger in *GgCry4a* WT (purple) and R317C (red) than in W369F (blue) or E320K (green). This suggests that the replacement of W_D by

Table 3. Time Constants Obtained by Fitting an Exponential to the Delayed Component of the %MFE Determined by CM for All Four Proteins

Protein	τ_{eMFE}/s
<i>GgCry4a</i> WT	21.0 ± 0.4
<i>GgCry4a</i> R317C	23.3 ± 0.3
<i>GgCry4a</i> E320K	23.3 ± 0.5
<i>GgCry4a</i> W369F	17.7 ± 0.2

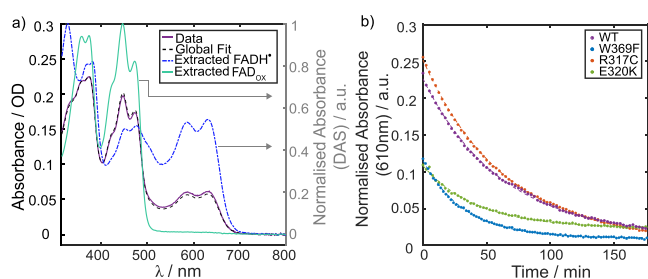


Figure 7. a) UV–visible spectrum of GgCry4a WT (purple) after 1 min illumination at 450 nm. The spectrum was successfully modeled by global analysis (black dashed line) assuming a single first-order recovery (see SM. 8). The normalized decay associated spectra (DAS) obtained from kinetic fitting are consistent with recovery from FADH[•] (blue dashed spectrum) to FAD_{ox} (teal spectrum). b) Recovery kinetics of FADH[•], averaged over 605–615 nm, for GgCry4a WT (purple), GgCry4a W369F (blue), GgCry4a R317C (red), and GgCry4a E320K (green), with photoexcitation ceasing at $t = 0$. The kinetics are normalized to the absorbance maximum of FAD_{ox} at 450 nm, prior to sample illumination. Single exponential fits to each kinetic trace are given by dashed lines of the same color as the data.

phenylalanine and the presence of K320 affects the efficiency of formation and/or lifetime (see Table 4) of the semiquinone radical. This observation is consistent with other measurements on single-point mutants of *DmCry*, producing long-lived FAD^{•−}, due to differences in the redox states stabilized by these proteins.^{37,66} In the *DmCry* measurements, mutations to the tryptophan chain as well as residues neighboring the terminal tryptophan decreased both the rate of photoreduction under steady-state conditions and the overall photoreduction yield.^{25,37,66}

The lifetimes obtained by kinetic fitting of the FADH[•] recovery data to a single exponential including constant offsets (Table 4, and Figure 7b) agree well with those obtained by global analysis (Table S7). The fastest recoveries were observed for GgCry4a W369F and GgCry4a E320K ($\tau_{\text{rec}} = 33$ min and $\tau_{\text{rec}} = 43$ min, respectively). The time scale on which these proteins recover is in general agreement with those obtained in ref 17, with differences due to the choice of buffer and illumination conditions (see SM.7).

DISCUSSION

The focus of this study was a comparative, spectroscopic investigation of four Cry4a avian cryptochromes, namely wild-type GgCry4a and three of its mutants, each with a single-point mutation. These were W369F, in which W_D is replaced by a phenylalanine, and R317C and E320K, both chosen after sequence comparison had highlighted amino acid positions 317 and 320 as distinct between migratory (C317 and K320) and nonmigratory birds, such as the chicken (R317 and

Table 4. Time Constants (τ_{rec}) Obtained by Monoexponential Fitting of the Decay of FADH[•] Generated by Continuous Blue-Light Illumination, Shown in Figure 7b

Protein	$\tau_{\text{rec}}/\text{min}$
GgCry4a WT	61.4 ± 0.21
GgCry4a R317C	63.9 ± 0.15
GgCry4a E320K	42.8 ± 0.32
GgCry4a W369F	32.6 ± 0.15

E320).¹¹ The most notable observations of our comprehensive study are:

1. All Four Proteins Form Spin-Correlated Radical Pairs. The TrEPR data of all four proteins (Figure 2) show the typical (fingerprint) antiphase structure characteristic for SCRP. Furthermore, these spectra are consistent with an FAD^{•−} TrpH^{•+} radical pair (RP1), with differences between the spectra of GgCry4a WT/R317C/E320K relative to GgCry4a W369F attributed to different relative orientations of the FAD^{•−} and the terminal TrpH^{•+} radicals. The inter-radical separation between FAD^{•−} and the terminal TrpH^{•+} is markedly shorter in GgCry4a W369F than the other proteins, as confirmed by out-of-phase ESEEM measurements.^{18,24} This demonstrates that electron transfer in wild-type GgCry4a proceeds along the whole tetrad of tryptophans as in *DmCry*, *CiCry4a*, and *ErCry4a*.^{18,20,24–26} EPR also confirms that the inter-radical separation and coupling parameters ($|D| = 8–13$ MHz, $J \approx 0$ MHz) are sufficiently small in all four proteins to allow efficient singlet–triplet interconversion at the field strengths employed here leading to magnetic field effects on the radical recombination yields and kinetics.

2. Shortening of the Electron Transfer Chain Increases the Lifetime of the Radical Pair, RP1. TA spectroscopy (Figure 3 and Table 2, τ_2) indicates that the radical pair lifetime of RP1 in the (W369F) mutant protein with its triad of tryptophan residues exceeds that of the proteins with the full tetrad, *in vitro*. This result seems counterintuitive, given the shorter inter-radical distance of the terminal FAD^{•−} TrpH^{•+} radical pair which leads to faster radical recombination. One explanation for this unexpected result is that the back electron transfer, which returns RP1 to the ground state, is in the Marcus inverted region.⁶⁷ Thus in GgCry4a W369F the radical recombination is slowed by an increase in the driving force for back-electron transfer or a significant decrease in the reorganization energy (due e.g. to a change in the solvation of the tryptophan radical). Few studies of electron transfer in cryptochromes have been reported, but detailed ultrafast investigations of related flavoproteins (including *EcPL* and *AtCry3*) have found that all back electron transfer processes involving the tryptophan triad are in the Marcus inverted region.^{68,69} This work supports an analogous process in avian Cry4a.

3. Shortening of the Electron Transfer Chain Increases the Effect of Applied Magnetic Fields on the Radical Kinetics and Yields. In agreement with previous results on *ErCry4a* W369F, shortening of the tryptophan chain to three residues in GgCry4a W369F increases the protein's magnetic field sensitivity *in vitro*, as measured by all the spectroscopy methods employed here.¹⁸ As observed previously for *EcPL*, this results from similar time scales of protonation and terminal tryptophan recombination (Figure 4a, Figure S12) in GgCry4a W369F, resulting in larger MFEs than the other three proteins (Figure 4b, Figure 6a,d).²⁰ The similar deprotonation time scales in GgCry4a W369F and *DmCry* W394F mutants (both lacking their W_D, with W318 (W_C) acting as the terminal donor) supports the kinetic assignments of the TA spectra (Figure S12, Table S5).²⁴ The rate of deprotonation is, however, an order of magnitude shorter than the reported RP2 formation rate in *EcPL* (with its tryptophan triad).^{20,25,44} However, the study on *EcPL* used higher glycerol concentration than the work conducted here, which could, in part, account for the prolonged lifetime of TrpH^{•+}. It may also indicate that TrpH^{•+} at residue 318 (W_C)

in avian Cry4a's is more liable to deprotonation than the corresponding 306 (W_C) position in EcPL.

In contrast to AtCry1 and EcPL²⁰ no clear evidence was found here for Low Field Effects (in which the sign of the MFE is inverted when the magnetic field is weaker than the effective hyperfine interaction).⁷⁰ The same is true for ErCry4a.¹⁸ Compared to ref 20, the measurements reported here were performed under conditions closer to those expected *in vivo*, specifically higher temperature (278 K vs 250 K), which may have prevented the emergence of a sign change. It is possible that a strong Low Field Effect only occurs when the protein is aligned by complexation to membrane proteins in photoreceptor cells in the retina.⁵⁰

4. Mutations of Amino Acid Positions 317 and 320 Do Not Impact Significantly on Radical Pair Kinetics or Magnetic Field Effects; Minor Changes Are Observed for FADH[•] Reoxidation. All spectroscopies employed here confirm that mutation of neither R317 nor E320 to cysteine and lysine, respectively, has significant impact on either the radical pair kinetics or the magnetic field effects *in vitro*.

In continuous illumination experiments, minor differences in the FADH[•] lifetime are observed for the E320K mutant compared to the WT and R317C (Figure 7b). This could be due to the proximity of the lysine to the W_D altering the reduction potential of Trp[•], such that it is less easily reduced, thus promoting bimolecular recombination of FADH[•] with Trp[•]. Support for such a mechanism comes from the fact that lysine residues modify the redox chemistry of iron–sulfur clusters in *E. coli* fumarate reductase.⁷¹ Further work on this hypothesis is presently ongoing but is outside the scope of this paper.

In W369F at 298 K (25 °C), FADH[•] recovers more rapidly than in the WT following 1 min photoillumination (Figure 7b, Table 4). Additionally, a smaller τ_{eMFE} relative to the other three proteins is observed by CM. This is perhaps expected, given that the environment of the Trp[•] is significantly perturbed in this mutant and that τ_{eMFE} is determined by the lifetimes of the flavin and tryptophan radicals. The τ_{eMFE} values obtained here are significantly longer than those reported for AtCry1 using a wide-field microscopy.⁵⁸ We attribute this to raster scanning of the excitation laser employed in CM, slowing the MFE kinetics. It is noteworthy that a prolonged τ_{eMFE} is also observed for AtCry1 in CM (Figure S17).

The faster recovery of W369F in both CM and continuous illumination experiments can be explained by concomitant recovery of FADH[•] and Trp[•], with any solvent mediated recovery of Trp[•] suppressed, as W318 (W_C) is less solvent exposed than W369 (W_D). The shorter τ_{eMFE} provides an additional explanation for the much larger MFE observed in GgCry4a W369F by BBCEAS. While the field switching time employed in BBCEAS is too fast to resolve eMFEs, the time evolution of the MFE over the 100 ms field-on time still contributes to the overall BBCEAS $\Delta\Delta A$ signature. A faster τ_{eMFE} in this system leads to an amplification of the MFE, relative to the other three proteins.

5. Oxygen Efficiently Recycles Cry4a Proteins under Continuous or Repeated Light Exposure. Our results provide the first demonstration that photoreduced Cry4a's can be effectively recycled by molecular oxygen *in vitro* (see SM.3–7 for details) and that MFEs can be detected without adding oxidizing agents, e.g. ferricyanide. The exact reoxidation mechanism is currently unknown. Studies of AtCry1 have suggested superoxide ($O_2^{\bullet-}$) as a possible reaction partner for

FADH[•], generating hydrogen peroxide as a byproduct of flavin oxidation,⁷² consistent with similar models proposed for flavoenzymes, including monooxygenases.^{73,74} A test of this hypothesis could involve addition of peroxide scavengers to samples of Cry4a exposed to different O_2 concentrations and/or varied duration and intensity of light exposure. Such measurements have been conducted on DmCry using *in vitro* fluorescence assays confirming the generation of reactive oxygen species upon blue-light illumination.⁷⁵ However, it is unlikely that the superoxide radicals would contribute to the spin dynamics of the systems due to rapid spin-relaxation of any resulting radical pair, driven by spin–orbit coupling in $O_2^{\bullet-}$.⁷⁶ Kattinig and colleagues^{77,78} have proposed a mechanism that could circumvent this issue via a paramagnetic scavenger that could react spin-selectively with one of the components of a flavin-superoxide radical pair. Probably the best way to test this idea would be to attach a stable radical to the protein with appropriate redox properties at a position predicted by spin dynamics simulations.^{77,78}

CONCLUSION

Despite their evolutionary conservation, the cysteine and lysine residues in positions 317 and 320 in ErCry4a have little impact on either the spin dynamics or radical pair kinetics when introduced into GgCry4a. Any differences in the magnetic sensitivity of robin and chicken Cry4a's (and, more generally, between Cry4a proteins from migratory and nonmigratory birds) must therefore lie in their magnetic *signaling*, rather than *sensing*, properties. Positions 317 and 320 are both solvent-exposed and so could facilitate the binding of Cry4a to intracellular signaling partners, some of which have been provisionally identified.^{79,80} More information regarding the entire signal transduction process will be required to reach definite conclusions on the role of these sequence differences. The lysine residue does affect recovery times of FADH[•], and flavin redox states such as FADH[•] and FADH⁻ are known to act as biologically relevant states in AtCry1/AtCry2 and EcPL, respectively. This residue could be used to tune the flavin lifetime for this purpose.⁸¹ Other residues may be involved in light-induced structural rearrangements, e.g. in the C-terminal domain.⁸²

Mutation of the fourth tryptophan to a phenylalanine (GgCry4a W369F) has a dramatic effect in all spectroscopic studies employed here. EPR demonstrates a shortening of the electron transport chain in W369F compared to the WT and confirms that RP1 is FAD^{•-} paired with the fourth (W_D) tryptophan in the WT protein. Furthermore, all the techniques used here find an increase in the protein's magnetic-field sensitivity for the W369F mutant. The same result was observed for ErCry4a,¹⁸ and suggests that, while nature has seemingly optimized the avian magnetic compass to use four tryptophan residues, this is at the expense of the magnetic sensitivity, at least *in vitro*. This may point to a role for the solvent-exposed tryptophan residue, W_D , in signaling, with suggestions that Cry4a may stabilize a composite radical pair, in which the magnetic sensitivity is optimized through the spin-dependent reactions of W318 (W_C) and signaling involving W369 (W_D). These possibilities have been discussed in detail in refs. 18 and 83. More detailed experimental investigations, including temperature-dependent out-of-phase ESEEM studies and single-point mutations (e.g., tryptophan to tyrosine) which could perturb the energetics of the equilibrium would be required to confirm such a hypothesis.

Finally, we believe our work represents an important step forward, providing a new gold standard for spectroscopic investigations of Cry4a proteins under more biologically relevant conditions, avoiding the harsh oxidizing conditions employed (such as the use of potassium ferricyanide).^{18,21,66}

■ ASSOCIATED CONTENT

SI Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/jacs.4c14037>.

Experimental and Data Analysis Methods, protein characterization by native mass spectrometry and UV–vis spectroscopy, additional EPR, TA and BBCEAS spectra and CM data, global analysis fitting procedure and results for BBCEAS and TA spectroscopy, comparison of Cry4a sequences of migratory and nonmigratory birds, comparison of EPR and BBCEAS data acquired with and without potassium ferricyanide (PDF)

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Notes

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