

6-21-2011

Human cryptochrome exhibits light-dependent magnetosensitivity

Lauren E. Foley

University of Massachusetts Medical School

Robert J. Gegear

University of Massachusetts Medical School

Steven M. Reppert

University of Massachusetts Medical School

Follow this and additional works at: http://escholarship.umassmed.edu/gsbs_sp



Part of the [Neuroscience and Neurobiology Commons](#)

Repository Citation

Foley, Lauren E.; Gegear, Robert J.; and Reppert, Steven M., "Human cryptochrome exhibits light-dependent magnetosensitivity" (2011). *GSBS Student Publications*. 1770.

http://escholarship.umassmed.edu/gsbs_sp/1770

This material is brought to you by eScholarship@UMMS. It has been accepted for inclusion in GSBS Student Publications by an authorized administrator of eScholarship@UMMS. For more information, please contact Lisa.Palmer@umassmed.edu.

ARTICLE

Received 6 Apr 2011 | Accepted 24 May 2011 | Published 21 Jun 2011

DOI: 10.1038/ncomms1364

Human cryptochrome exhibits light-dependent magnetosensitivity

Lauren E. Foley¹, Robert J. Gegear^{1,†} & Steven M. Reppert¹

Humans are not believed to have a magnetic sense, even though many animals use the Earth's magnetic field for orientation and navigation. One model of magnetosensing in animals proposes that geomagnetic fields are perceived by light-sensitive chemical reactions involving the flavoprotein cryptochrome (CRY). Here we show using a transgenic approach that human CRY2, which is heavily expressed in the retina, can function as a magnetosensor in the magnetoreception system of *Drosophila* and that it does so in a light-dependent manner. The results show that human CRY2 has the molecular capability to function as a light-sensitive magnetosensor and reopen an area of sensory biology that is ready for further exploration in humans.

¹Department of Neurobiology, University of Massachusetts Medical School, Worcester, Massachusetts 01605, USA. †Present address: Department of Biology and Biotechnology, Worcester Polytechnic Institute, Worcester, Massachusetts 01609, USA. Correspondence and requests for materials should be addressed to S.M.R. (email: Steven.Reppert@umassmed.edu).

In vertebrates, such as migratory birds and sea turtles, the ability to sense the Earth's magnetic field is clearly important for positional and directional information during their long-distance migrations^{1–3}. In many animals, magnetoreception is thought to depend on light-sensitive chemical reactions involving the flavo-protein cryptochrome (CRY)^{3–5}.

There are two major groups of CRYs found in animals, based on both their phylogenetic position and function in the regulation of circadian clocks⁶. *Drosophila*-like type 1 CRYs (which have been found only in invertebrates) are sensitive to blue light and function mainly as circadian photoreceptors. On the other hand, vertebrate-like type 2 CRYs (which are found in both vertebrates and invertebrates) are thought to function mainly as negative regulators of the clockwork's transcriptional feedback loop and do so in a light-independent manner⁷.

Genetic studies have shown that a light-dependent magnetic sense in *Drosophila* is indeed mediated by its type 1 CRY^{8,9}. Furthermore, using the *Drosophila* GAL4-UAS transgenic approach for targeted gene expression, it has been shown recently that type 1 CRY proteins from either the fly itself or the monarch butterfly, and an insect version of type 2 CRY from the monarch can each restore magnetosensitivity and its light dependency in CRY-deficient *Drosophila*¹⁰.

Humans are widely assumed not to have a magnetic sense³. For example, the extensive behavioural studies by Robin Baker^{11–15}, suggesting a link between non-visual navigation and magnetoreception in humans, are controversial. However, there is consistent evidence of an influence of geomagnetic fields on the light sensitivity of the human visual system^{16,17}. Moreover, it has been proposed recently that light-sensitive magnetic responses are not only used for directional information, but may also aid visual spatial perception in mammals, by providing a spherical coordinate system for integrating spatial position³. We therefore evaluated the light-dependent magnetosensing potential of human CRY. Here we show using a transgenic approach, that human CRY can function as a magnetosensor in the magnetoreception system of *Drosophila* and that it does so in a light-dependent manner. Thus, human CRY has the molecular capability to function as a light-sensitive magnetosensor, and this finding may lead to a renewed interest in human magnetoreception.

Results

Human CRY2 rescues light-dependent magnetosensitivity. Although there are two versions of type 2 CRYs in humans (hCRY1 and hCRY2), we focused our transgenic study on hCRY2, because previous anatomical studies have shown that it is expressed at ca. 11-fold higher levels than hCRY1 in the retina^{18,19}. Thus hCRY2 is nicely poised to receive the light information necessary to initiate a light-sensitive magnetic response in humans.

In *Drosophila*, magnetosensitivity can be reliably examined using a behavioural assay, in which groups of individuals experience an electric-coil-generated magnetic-field gradient ranging from ambient to eight times the Earth's strength on one arm of a two arm T-maze^{8,10}. The directional response of flies to the field is tested by comparing the number of flies in the arm that faces the coil that generates a magnetic field with the number in the opposite arm that faces the coil that is not magnetized. Flies are tested for their direction response in either the naive state or after training when the field is paired with a sucrose reward. Using this behavioural assay, both wild-type Canton-S flies and CRY-deficient flies expressing the *Drosophila cry* transgene under the control of the *tim-GAL4* driver, which expresses the transgene where *cry* is normally expressed, manifest significant naive directional avoidance of the field (negative preference index value) and a significant directional trained preference for the field (positive preference index value) (Fig. 1a, white and black bars, respectively)^{8,10}.

Remarkably, CRY-deficient flies expressing the hCRY2 transgene under the control of the *tim-GAL4* driver also showed comparably

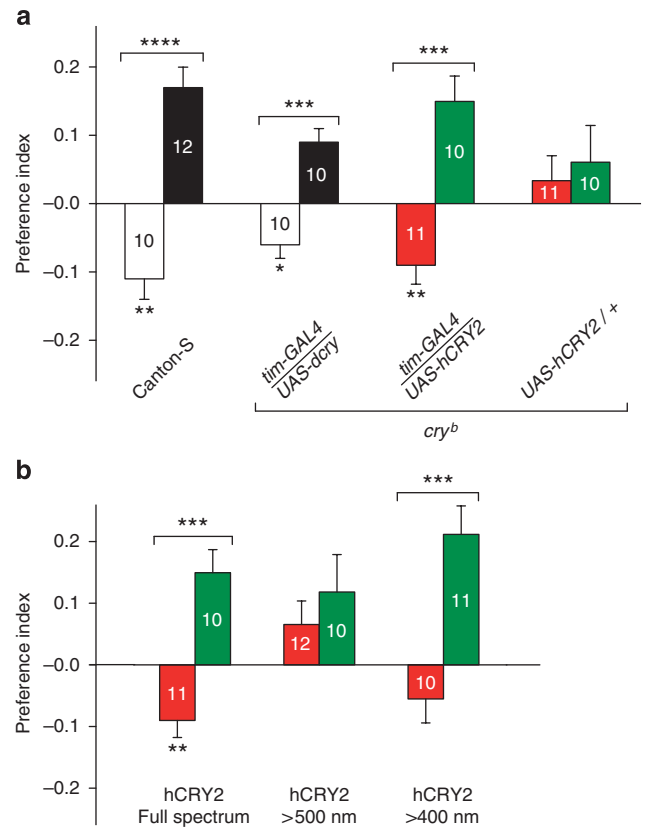


Figure 1 | Human CRY2 rescues light-dependent magnetoreception in CRY-deficient flies. (a) A *tim-GAL4*-driven human CRY2 transgene (*tim-GAL4/UAS-hCRY2*) rescues magnetic responses in the CRY loss-of-function *cry^b* mutant background. For comparison, naive and trained responses to the magnetic field are shown for wild-type Canton-S flies (left bar set; reproduced from ref. 8) and for a *tim-GAL4*-driven *Drosophila cry* transgene in *cry^b* flies (*tim-GAL4/UAS-dcry*) (second from left bar set; reproduced from ref. 10). The *UAS-hCRY2/+* transgene alone (without the *tim-GAL4* driver) did not result in significant magnetosensitive responses ($P > 0.05$; right bar set). Bars show PI values for naive (white, *dcry*; or red, hCRY2) and trained (black, *dcry*; or green, hCRY2) groups. To test whether flies responded to the experimental magnetic field, we either used a one-sample *t*-test to compare naive PI values with zero (that is, PI value expected with no response to the magnetic field) or a Student's *t*-test to compare PI values between trained and naive groups. Numbers represent groups tested. Values are mean \pm s.e.m. * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$. Genotypes in parentheses: *tim-GAL4/UAS-dcry* (*y w; tim-GAL4/UAS-mycdcry; cry^b*); *tim-GAL4/UAS-hCRY* (*y w; tim-GAL4/UAS-mychCRY2; cry^b*); and *UAS-dcry/+* (*y w; UAS-mychCRY2/+; cry^b*). (b) Light dependence of magnetic responses rescued by human (h)CRY2 (*y w; tim-GAL4/UAS-mychCRY2; cry^b*). The full-spectrum data are the same as those depicted in a. The irradiance curves for the three light conditions used are the same as those used previously⁸. Bars show PI values for naive (red) and trained (green) groups. To test whether flies responded to the experimental magnetic field, we either used a one-sample *t*-test to compare naive PI values to zero (that is, PI value expected with no response to the magnetic field) or a Student's *t*-test to compare PI values between trained and naive groups. Numbers represent groups tested. Values are mean \pm s.e.m. *** $P < 0.01$, **** $P < 0.001$.

robust naive and trained responses to a magnetic field under full-spectrum light (Fig. 1a, red and green bars, respectively), while CRY-deficient fly lines expressing either the GAL4 driver alone¹⁰ or the *UAS-hCRY2/+* transgene alone (Fig. 1a) did not respond. Moreover, the hCRY2 rescue of magnetosensitivity was light sensitive (Fig. 1b);

the hCRY2-induced magnetic responses were blocked when only long-wavelength light (>500 nm) was available, and were partially restored when light >400 nm was available. Thus, light in the blue range is necessary for the hCRY2 magnetic response.

Discussion

Our results show that hCRY2, the prototype type 2 CRY^{18,19}, has the molecular capability to function in a light-dependent magnetoreception system, either as a light-sensitive magnetosensor or as part of a magnetosensing pathway. However, we do not yet know whether this capability is translated into a downstream biological response in the human retina. Nonetheless, the transgenic findings with hCRY2, together with its anatomical location in the human retina and previous work showing field effects on the visual system, suggest that a reassessment of human magnetosensitivity may be in order. The results also define a clear light-sensitive response for a mammalian CRY, as predicted in 1998 (ref. 18). Additional research on magnetosensitivity in humans at the behavioural level, with particular emphasis on the influence of magnetic fields on visual function, rather than non-visual navigation, would be informative.

Methods

Fly lines. Fly stocks were raised on standard cornmeal/agar medium at 25 °C and 60% relative humidity under a 12-h light:12-h dark lighting cycle. The *cry^b* and *UAS-dcry* (*UAS-mycdcry*) were a gift from Patrick Emery. The *tim-GAL4* (*tim-GAL4/CyO*)²⁰ driver line was used for transgene expression in *cry^b* flies.

For generating *UAS-mychCRY2* transgenic lines, the *hCRY2* open reading frame was amplified by PCR from *pfmh2* and subcloned into the *BglII* and *NofI* sites of the *pUASTattBmyc* vector. The *myc-pUASTattB* vector was generated by cloning a *BamHI-myc-BglII* fragment, created using two oligos followed by primer extension, into the *BglII* site of *pUASTattB*. The construct contained a full Kozak sequence. The construct was sequenced before injection into *y w, nos-phiC31/+*; *attP40* or *y w, nos-phiC31/Y; attP40/+* embryos by Genetic Services. During balancing, the *y w, nos-phiC31*-containing X chromosome was replaced with the *y w*-containing X chromosome.

Behavioural apparatus. The apparatus consisted of a choice chamber and an illuminated black box containing the two-coil system connected to an adjustable DC power supply⁸. The choice chamber was comprised of a one-tube training section and a two-tube choice section (T-maze) for testing the relative response of flies to a magnetic field. An elevator section was used to transfer flies between training and T-maze sections (see below). The housing box was constructed such that the choice chamber could be placed between the two coils in either an upright position (used during training) or when rotated to the horizontal (used during testing). In this way, flies were exposed to the same lighting conditions and magnetic field intensity during training and test trials.

The two-coil system allowed for the production of a magnetic field on one side, while producing no field on the opposite side and alternation of the field between sides; heat was generated equally in either coil, with or without generating a magnetic field. Coils were positioned directly under the tube ends at 45° to the horizontal. We adjusted the power supply so that the magnetic field intensity ranged from 0.1 G at the tube entrance to 5 G at the end of the tube. The box containing the coil system was open on the top so that the chamber, regardless of position, could be illuminated by one ZooMed Reptisun 10.0 UVB fluorescent tube (F17T8). Irradiance measurements were taken from inside one arm of the T-maze portion of the choice chamber apparatus to ensure that the lighting conditions measured represented those experienced by the flies while they were being either trained or tested. Wavelength dependence of magnetosensitivity was examined by either covering the top of the box with a long-wavelength filter that transmitted wavelengths of light >500 nm (Edmund Optic) or >400 nm (E400 from Gentex).

Behavioural procedure. Flies were starved for 18 h before testing their magnetic response. All experiments were performed between 0800 and 1500 hours EST. Flies in the trained group were moved into the training section facing one of two adjustable coils with no field for 2 min, and then were reloaded into the training tube containing sucrose reinforcement and a magnetic field for an additional 2 min. After being held for 1 min in the elevator section, flies were tested for their magnetic preference by transferring them to the T-maze section and allowing them to choose between the sides with or without a magnetic field for 2 min. Flies in the naive group were loaded into the elevator section and immediately transferred to the T-maze and allowed to choose between the sides with or without a magnetic field for 2 min. Trained and naive groups were tested consecutively with the magnetic field on the same side. As an additional control for side preferences independent of magnetic stimuli, we tested flies in the absence of a magnetic field.

For each population of flies tested (100–150 individuals), we calculated a preference index (PI) value based on the following equation: $(P_M - 0.5)/0.5$, where P_M is the proportion of flies on the magnetic field side of the T-maze. To test whether flies responded to the experimental magnetic field, we either used a one-sample *t*-test to compare naive PI values to zero (that is, PI value expected with no response to the magnetic field) or a Student's *t*-test to compare PI values between trained and naive groups.

References

- Wiltschko, W. & Wiltschko, R. Magnetic orientation and magnetoreception in birds and other animals. *J. Comp. Physiol. A* **191**, 675–693 (2005).
- Lohmann, K. J., Lohmann, C. M. F. & Putman, N. F. Magnetic maps in animals: nature's GPS. *J. Exp. Biol.* **210**, 3697–3705 (2007).
- Phillips, J. B., Muheim, R. & Jorge, P. E. A behavioral perspective on the biophysics of the light-dependent magnetic compass: a link between directional and special perception? *J. Exp. Biol.* **213**, 3247–3255 (2010).
- Ritz, T., Adem, S. & Schulten, K. A model for photoreceptor-based magnetoreception in birds. *Biophys. J.* **78**, 707–718 (2000).
- Lohmann, K. J. Q&A: animal behaviour: magnetic-field perception. *Nature* **464**, 1140–1142 (2010).
- Yuan, Q., Metterville, D., Briscoe, A. D. & Reppert, S. M. Insect cryptochromes: gene duplication and loss define diverse ways to construct insect circadian clocks. *Mol. Biol. Evol.* **24**, 948–955 (2007).
- Griffin, E. A. Jr., Staknis, D. & Weitz, C. J. Light-independent role of CRY1 and CRY2 in the mammalian circadian clock. *Science* **286**, 768–771 (1999).
- Gegeer, R. J., Casselman, A., Waddell, S. & Reppert, S. M. Cryptochrome mediates light-dependent magnetosensitivity in *Drosophila*. *Nature* **454**, 1014–1018 (2008).
- Yoshii, U., Ahmad, M. & Helfrich-Forster, C. Cryptochrome mediates light-dependent magnetosensitivity of *Drosophila*'s circadian clock. *PLoS Biol.* **7**, 813–819 (2009).
- Gegeer, R. J., Foley, L. E., Casselman, A. & Reppert, S. M. Animal cryptochromes mediate magnetoreception by an unconventional photochemical mechanism. *Nature* **463**, 804–807 (2010).
- Baker, R. R. *Human Navigation* (Manchester University Press, 1989).
- Fildes, B. N., O'Laughlin, B. J., Bradshaw, J. L. & Evans, W. J. Human orientation with restricted sensory information: no evidence for magnetic sensitivity. *Perception* **13**, 229–326 (1984).
- Westby, G. W. M. & Partridge, K. J. Human homing: still no evidence despite geomagnetic controls. *J. Exp. Biol.* **120**, 325–331 (1986).
- Baker, R. R. Human navigation and magnetoreception: the Manchester experiments do replicate. *Anim. Behav.* **35**, 691–704 (1987).
- Finney, B. A role for magnetoreception in human navigation. *Curr. Anthropol.* **36**, 500–506 (1995).
- Thoss, F., Bartsch, B., Fritzsche, B., Tellschaft, D. & Thoss, M. J. The magnetic field sensitivity of the human visual system shows resonance and compass characteristic. *Comp. Physiol. A* **186**, 1007–1010 (2000).
- Thoss, F., Bartsch, B., Tellschaft, D. & Thoss, D. The light sensitivity of human visual system depends on the direction of view. *J. Comp. Physiol. A* **188**, 235–237 (2002).
- Miyamoto, Y. & Sancar, A. Vitamin B2-based blue-light photoreceptors in the retinohypothalamic tract as the photoactive pigments for setting the circadian clock in mammals. *Proc. Natl. Acad. Sci. USA* **95**, 6097–6102 (1998).
- Thompson, C. L. *et al.* Expression of the blue-light receptor cryptochrome in the human retina. *Invest. Ophthalmol. Vis. Sci.* **44**, 4515–4521 (2003).
- Emery, P., So, W. V., Kaneko, M., Hall, J. C. & Rosbash, M. CRY, a *Drosophila* clock and light-regulated cryptochrome, is a major contributor to circadian rhythm resetting and photosensitivity. *Cell* **95**, 669–679 (1998).

Acknowledgments

We thank Aziz Sancar for providing the *hCRY2* *pfmh2* construct, Andrea Pereira for help with constructing the *UAS-hCRY2* clone and Shruti Basil and Andrew Dowd for technical assistance. Supported by NIH grant R01 GM086794.

Author contributions

All authors contributed to experimental design, execution, data analysis and writing the paper.

Additional information

Competing financial interests: The authors declare no competing financial interests.

Reprints and permission information is available online at <http://npg.nature.com/reprintsandpermissions/>

How to cite this article: Foley, L. E. *et al.* Human cryptochrome exhibits light-dependent magnetosensitivity. *Nat. Commun.* 2:356 doi: 10.1038/ncomms1364 (2011).

License: This work is licensed under a Creative Commons Attribution-NonCommercial-Share Alike 3.0 Unported License. To view a copy of this license, visit <http://creativecommons.org/licenses/by-nc-sa/3.0/>