

UC Berkeley

UC Berkeley Previously Published Works

Title

Quantum entanglement in photosynthetic light harvesting complexes

Permalink

<https://escholarship.org/uc/item/7mj263d7>

Authors

Sarovar, Mohan
Ishizaki, Akihito
Fleming, Graham R
et al.

Publication Date

2009-05-22

Peer reviewed

Quantum entanglement in photosynthetic light-harvesting complexes

Mohan Sarovar^{1,2*}, Akihito Ishizaki^{2,3}, Graham R. Fleming^{2,3} and K. Birgitta Whaley^{1,2}

Light-harvesting components of photosynthetic organisms are complex, coupled, many-body quantum systems, in which electronic coherence has recently been shown to survive for relatively long timescales, despite the decohering effects of their environments. Here, we analyse entanglement in multichromophoric light-harvesting complexes, and establish methods for quantification of entanglement by describing necessary and sufficient conditions for entanglement and by deriving a measure of global entanglement. These methods are then applied to the Fenna–Matthews–Olson protein to extract the initial state and temperature dependencies of entanglement. We show that, although the Fenna–Matthews–Olson protein in natural conditions largely contains bipartite entanglement between dimerized chromophores, a small amount of long-range and multipartite entanglement should exist even at physiological temperatures. This constitutes the first rigorous quantification of entanglement in a biological system. Finally, we discuss the practical use of entanglement in densely packed molecular aggregates such as light-harvesting complexes.

Unlike the case in classical physics, within quantum mechanics we can have maximal knowledge of a composite physical system and still not be able to assign a definite state to its constituent elements without reference to their relation to one another^{1,2}. Such systems are called entangled, and entanglement is a characteristic quantum mechanical effect that has been widely investigated in recent years^{3,4}. Entanglement is often viewed as a fragile and exotic property, and in the quantum information context, where it is used as a resource for information processing tasks, precisely engineered entangled states of interest can indeed be both fragile and difficult to manufacture. However, it has also been recognized that entanglement is a natural feature of coherent evolution, and recently there has been an effort to expand the realms in which entanglement can be shown to exist rigorously, particularly in ‘natural’ systems—that is, not ones manufactured in laboratory conditions. Signatures of entanglement, a characteristically quantum feature, have been demonstrated in thermal states of bulk systems at low temperatures and between parties at macroscopic length scales⁵. Additionally, several recent studies have focused on the dynamics of entanglement in damped, driven or generally non-equilibrium quantum systems^{6–9}. The dynamics of entanglement in open systems can be extremely non-trivial—especially in many-body systems—and the precise influence of non-Hamiltonian dynamics on entanglement is poorly understood. In a result particularly relevant to this work, it is shown in ref. 8 that entanglement can be continuously generated and destroyed by non-equilibrium effects in an environment where no static entanglement exists. The possibility of entanglement in noisy non-equilibrium systems at high temperatures intimates the question of whether we can observe entanglement in the complex non-equilibrium chemical and biological processes necessary for life. Here we present strong evidence for answering this question in the affirmative by determining the timescales and temperatures for which entanglement is

observable in a protein structure that is central to photosynthesis by green anoxygenic bacteria.

Light-harvesting complexes and entanglement

Recent ultrafast spectroscopic studies have revealed the presence of quantum coherence at picosecond timescales in biological structures, specifically in light-harvesting complexes (LHCs; refs 10–13). These studies demonstrate that, in moderately strongly coupled, non-equilibrium systems, quantum features can be observed even in the presence of a poorly controlled, decohering environment. During the initial stage of photosynthesis, light is captured by pigment–protein antennas, known as light-harvesting complexes, and the excitation energy is then transferred through these antennas to reaction centres, where photosynthetic chemical reactions are initiated. Different LHCs vary in their detailed structure but all consist of densely packed units of pigment molecules, and all are spectacularly efficient at transporting excitation energy in disordered environments¹⁴. Average interchromophore separations on the scale of ~ 15 Å are fairly common in LHCs. At these distances, the dipole coupling of these molecules is considerable and leads to coherent interactions at observable timescales^{15–17}. It is this ‘site’ coherence (coherence between spatially separated pigment molecules) that prompts us to examine entanglement in these systems and to consider the timescales and temperatures at which entanglement can exist.

We begin by noting that in natural conditions many LHCs contain at most one excitation at any given time¹⁴. This is especially true of LHCs in photosynthetic bacteria, which are among the ones most heavily studied, because these bacteria receive very little sunlight in their natural habitat. Given this, we can treat each chromophore as a two-level system and the natural Hilbert space of the LHC quantum states will further be restricted to the zero- and one-excitation subspaces of the full tensor-product

¹Berkeley Center for Quantum Information and Computation, Berkeley, California 94720, USA, ²Department of Chemistry, University of California, Berkeley, California 94720, USA, ³Physical Bioscience Division, Lawrence Berkeley National Laboratory, Berkeley, California 94720, USA.

*e-mail: msarovar@berkeley.edu.

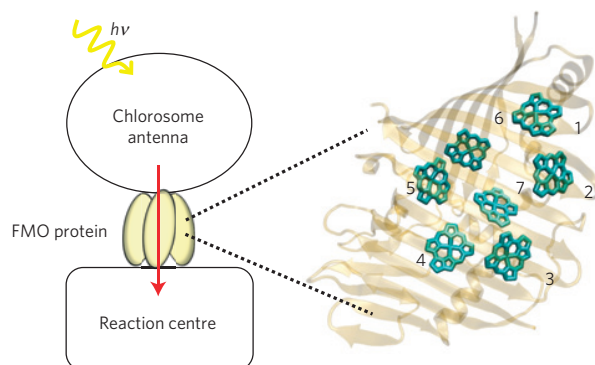


Figure 1 | The light-harvesting apparatus of green sulphur bacteria and the FMO protein. The schematic on the left illustrates the absorption of light by the chlorosome antenna and transport of the resulting excitation to the reaction centre through the FMO protein. On the right is an image of a monomer of the FMO protein, showing also its orientation relative to the antenna and the reaction centre^{22,24}. The multiring units are BChla molecules and the surrounding β sheets and α helices form the protein environment in which the BChla molecules are embedded. The numbers label individual BChla molecules, also referred to as ‘sites’ in the main text.

space derived from N chromophores. Furthermore, there is no coherence between states in these different subspaces, because under natural conditions all processes connecting the two subspaces are incoherent. A state in the single-excitation manifold is written in the site basis as

$$\rho(t) = \sum_{i=1}^N \rho_{ii}(t) |i\rangle \langle i| + \sum_{i=1}^N \sum_{j>i}^N \rho_{ij}(t) |i\rangle \langle j| + \rho_{ij}^*(t) |j\rangle \langle i|$$

where $|i\rangle$ represents the state where only the i th chromophore (site) is excited and all other chromophores are in their electronic ground states. The density matrix ρ is unnormalized—that is, $\sum_i \rho_{ii}(t) \leq 1$ —because the single-excitation state has a finite lifetime owing to trapping by the reaction centre complex and radiative decay of the excitation.

Given an unnormalized density matrix ρ representing a single-excitation state, we wish to calculate the amount of entanglement in the state. Entanglement in the site basis refers to non-local correlations between the electronic states of spatially separated chromophores. This is analogous to the entanglement of electromagnetic modes carrying a single photon in the dual-rail representation¹⁸. Mixed-state, multipartite entanglement is notoriously difficult to quantify⁴, but the single-excitation restriction enables us to formulate two useful and simple measures. First, for bipartite entanglement between two chromophores, i and j , we use a standard measure of entanglement, concurrence¹⁹, which, if ρ is the single-excitation density matrix for the whole complex, takes the form $C_{ij} = 2|\rho_{ij}|$. Second, to quantify global entanglement in the LHC, we derive (see Methods section) a readily computable expression for mixed-state entanglement in the physically relevant zero- and single-excitation subspaces, namely

$$E[\rho] = - \sum_{i=1}^N \rho_{ii} \ln \rho_{ii} - S(\rho) \quad (1)$$

where $S(\rho) = -\text{tr} \rho \ln \rho$ is the von Neumann entropy of the state ρ . In Supplementary Information we show that this measure, which is based on the relative entropy of entanglement²⁰, is a true entanglement monotone over the physically relevant zero- and single-excitation states and hence provides a natural quantitative measure of entanglement in LHCs.

Entanglement dynamics in the Fenna–Matthews–Olson (FMO) complex

A commonly studied LHC is the FMO protein from green sulphur bacteria, such as *Chlorobium tepidum*²¹. The FMO complex is a trimer formed by three identical monomers that each bind seven bacteriochlorophyll-*a* (BChla) molecules. We shall restrict our study to a single monomer, which is shown in Fig. 1, because the monomers function independently. The site energies of the BChla molecules and coupling between molecules are well characterized; we use the *Chlorobium tepidum* site energies and coupling strengths from ref. 22 to form a Hamiltonian that describes the closed-system dynamics of an FMO monomer excitation (see the Supplementary Information for details). The structure of the Hamiltonian indicates that some pairs of chromophores are moderately strongly coupled (owing to their close proximity and favourable dipole orientations) and hence effectively form dimers. The wavefunctions of the system’s energy eigenstates, usually called Frenkel excitons, are primarily delocalized across these dimers. The dominant dimers are formed by chromophore pairs: 1–2, 5–6 and 3–4.

In addition to the reversible (Hamiltonian) dynamics, there are interactions between each chromophore and the protein environment in which it is embedded. These interactions couple the protein dynamics to the FMO energy levels, resulting in static and dynamic disorder, and thereby lead to dephasing of the FMO electronic excitation state. Coherence properties of the FMO protein are then dictated by the interplay between coherent dynamics of the complex and decoherence effects due to environmental interactions. Distinctly quantum properties such as entanglement rely critically on coherence, and therefore it is essential that a model that accurately accounts for environmental effects on coherence be used to make predictions about entanglement in the FMO protein. In this work we use a recently developed non-perturbative, non-Markovian quantum master equation³⁴ to simulate excitation dynamics in the FMO complex. This dynamical model is particularly suited to describing the complex exciton–phonon interactions in LHCs and incorporates the dynamics of phonon reorganization in a realistic manner. See the Methods section for a brief review and discussion of this model, and the physical parameters we use in the simulations presented below.

Analysis of the FMO energy levels and its physical structure indicates that sites 1 and 6 interface with the chlorosome, transmitting energy to the complex²² (see Fig. 1), and recent experiments have confirmed that these two sites are the first to become excited in the FMO protein (ref. 24). In Fig. 2 we show the time evolution of the global measure of entanglement given by equation (1) when the initial state is an excitation on site 1 or 6, for two temperatures: 77 K and 300 K. In addition to simulations at the physiologically relevant temperature of 300 K, we also carry out simulations at 77 K because the ultrafast spectroscopy experiments that probe LHCs are commonly carried out at this temperature^{10,11} (although very recently room-temperature experiments have also been performed^{12,13}). The inset to this graph also includes the trace of the single-excitation density matrix to show the total population in the single-excitation subspace. A general feature of the global entanglement measure in all the scenarios depicted in Fig. 2 is that its value rises rapidly for short times and then after ~ 30 – 50 fs decays with varying amounts of oscillation. An explanation for this behaviour is that entanglement increases rapidly for short times owing to quick delocalization of the excitation caused by large 1–2 and 5–6 site coupling terms in the FMO complex Hamiltonian. Then, as the excitation begins exploring other sites, the global entanglement decreases owing to incoherent transport and rapid dephasing. For both initial conditions and temperatures, there is finite entanglement in the system up to 5 ps. At 300 K the short-time behaviour is qualitatively similar to the low-temperature case. At long times, however, the

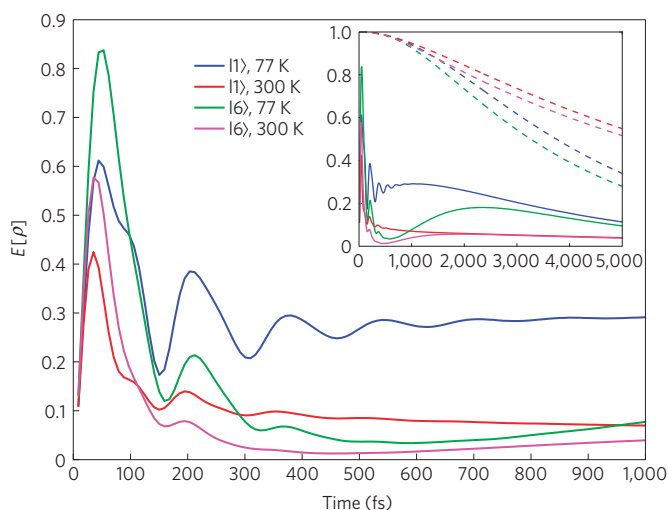


Figure 2 | Global entanglement in the FMO protein. Time evolution of the global entanglement measure given in equation (1) for the two initial states $|1\rangle$ and $|6\rangle$, at low ($T = 77$ K) and high ($T = 300$ K) temperatures. The inset shows the long-time evolution of the same quantities, together with the trace of the single-excitation density matrix as dashed curves (identical colour coding, and the same units on the axes as in the main figure).

initial state dependence of the global entanglement is suppressed, in contrast to the low-temperature case.

To further elucidate the dynamics and structure of entanglement in the FMO monomer under realistic conditions, we examine pairwise entanglement in the system, as measured by the bipartite concurrence between two sites: $C_{ij} = 2|\rho_{ij}|$ for any two sites i, j . Figures 3 and 4 show the time evolution of pairwise concurrence when the initial state is $|1\rangle$ and $|6\rangle$, respectively, at both 77 K and 300 K. For clarity, we have only shown the most significant and illustrative concurrence curves, but it should be noted that there is finite bipartite entanglement between almost all chromophores, especially for short timescales. Supplementary Information contains plots that provide a complete depiction of bipartite entanglement in this complex.

When the initial excitation is on site 1, as shown in Fig. 3, the bipartite entanglement in the complex is primarily between sites 1, 2, 3 and 4. Most of this bipartite entanglement is between pairs of moderately strongly coupled chromophores that form dimers and hence have large amounts of coherence. However, a surprising aspect of the short-time behaviour is that there is considerable concurrence between non-nearest-neighbour sites, as indicated by the 1–5 and 1–3 curves. Perhaps most strikingly, there is a large amount of entanglement within 1 ps between chromophores 1 and 3, which are almost the furthest apart in the FMO complex (separated by ~ 28 Å), and are very weakly coupled. This long-range entanglement is mediated by chromophores connecting these sites, and is a sign of multipartite entanglement in the system. At 77 K bipartite entanglement persists for more than 5 ps, and seems limited only by the energy trapping rate—essentially there is non-negligible entanglement as long as the excitation has not been trapped by the reaction centre. Bipartite entanglement persists for long periods of time at 300 K also, but it diminishes at a faster rate compared with low temperatures. However, note that at short times (< 600 fs) temperature has little effect on the amount of bipartite entanglement in the system—qualitatively, the concurrence is scaled by $\sim 3/4$ when the temperature is increased from 77 K to 300 K.

When the initial excitation is on site 6, as shown in Fig. 4, bipartite entanglement exists between several chromophores at short times; there is non-negligible entanglement between any two

of the sites 4, 5, 6 and 7. Note that not all of these are dimerized chromophores with large couplings. Furthermore, the presence of finite entanglement across any bipartite partition of the sites 4, 5, 6 and 7 indicates that the FMO complex contains genuine multipartite distributed entanglement within ~ 600 fs.

To summarize, realistic simulations of FMO dynamics indicate that considerable multipartite entanglement is present in the FMO complex at timescales of ~ 5 ps at 77 K and ~ 2 ps at 300 K. For both realistic initial states there is multipartite entanglement between all sites involved in excitation transport.

Significance and implications of LHC entanglement

The previous section presented numerical evidence for the existence of entanglement in the FMO complex for picosecond timescales—essentially until the excitation is trapped by the reaction centre. This is remarkable in a biological or disordered system at physiological temperatures. It illustrates that non-equilibrium multipartite entanglement can exist for relatively long times, even in highly decoherent environments. Although the length scales over which entanglement was shown to persist were restricted to $\lesssim 30$ Å because of the relatively small size of the FMO complex, we expect that such long-lived, non-equilibrium entanglement will also be present in larger light-harvesting antenna complexes, such as LH1 and LH2 in purple bacteria. This is because they contain the key necessary ingredient: moderately strongly coupled chromophores that can lead to significant coherent delocalization of electronic excitations^{16,25}. In larger light-harvesting antennae it may also be possible to take advantage of the ability to create and support multiple excitations to access a richer variety of entangled states.

Our numerical results are based on a non-perturbative, non-Markovian dynamical model particularly formulated to model the actual conditions in LHCs. The precise dynamics of entanglement at short times is dependent on the non-Markovian dynamics of the system, and a correct picture of which chromophores are entangled at long times relies on an accurate approach to thermal equilibrium. The dynamical model we use realistically models both these limits. We note, however, that the long-time survival of entanglement in the FMO complex is also predicted by less sophisticated dynamical models (see Supplementary Information) and therefore is a robust feature of these systems.

We emphasize that our prediction of entanglement in the FMO complex is experimentally verifiable because the timescales on which the entanglement exists can readily be probed using nonlinear femtosecond spectroscopy techniques. Quantum-state tomography on LHCs is not possible at present, but techniques are under development that will enable individual density-matrix elements to be extracted. The primary obstacle to this task is the determination of excitonic transition dipoles in the complex because the spectroscopy signals representing elements of the density matrix are scaled by the magnitude of these dipoles. It has recently been demonstrated that polarization-dependent two-dimensional spectroscopy is capable of determining these dipole moments²⁶, and hence the dynamical estimation of density-matrix elements is within reach. An entanglement witness, \mathcal{W} , identified in the Methods section, simplifies the experimental verification of entanglement because it implies that monitoring a small subset of site basis coherences (for example, the ones expected to be the largest in magnitude) is sufficient for detecting entanglement.

It is interesting to consider the biological significance of this entanglement. The FMO complex is an unusual light-harvesting component because its primary role is as a wire, to transport excitations between the main LHC of green sulphur bacteria and their reaction centres. Most other light-harvesting structures have dual roles of light capture and excitation transport and are as a consequence larger and more complex. This study demonstrating the essential features of entanglement in LHCs focused on the FMO

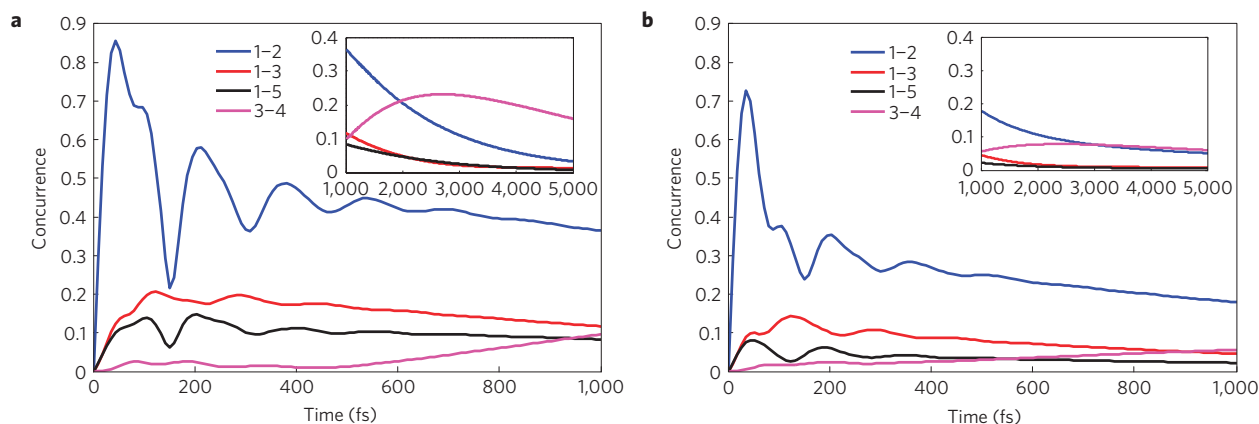


Figure 3 | Bipartite entanglement in the FMO protein when the initial state is an excitation localized on site 1. **a,b**, Time evolution of the concurrence measure of bipartite site entanglement, $C_{ij} = 2|\rho_{ij}|$, at 77 K (**a**) and 300 K (**b**). Only curves for the most entangled chromophores are shown. The insets show the long-time behaviour (identical colour coding, and the same units on the axes as in the main figures).

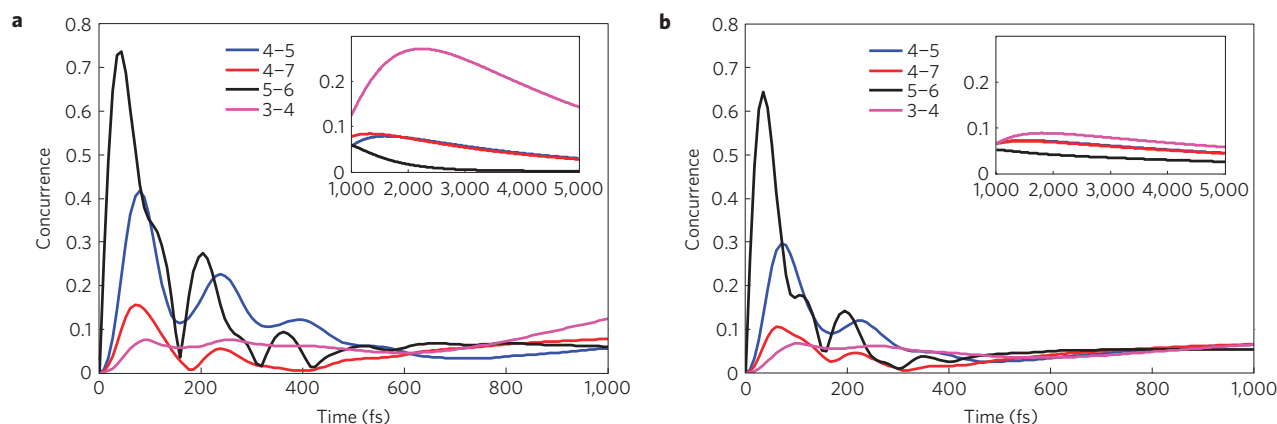


Figure 4 | Bipartite entanglement in the FMO protein when the initial state is an excitation localized on site 6. **a,b**, Time evolution of the concurrence measure of bipartite site entanglement, $C_{ij} = 2|\rho_{ij}|$, at 77 K (**a**) and 300 K (**b**). Only curves for the most entangled chromophores are shown. The insets show the long-time behaviour (identical colour coding, and the same units on the axes as in the main figures).

complex because it is very well characterized. Although FMO may be too small to take advantage of the non-classical correlations implied by entanglement, we expect that such entanglement has functional implications in more complex light-harvesting structures, particularly in photosynthetic units with several reaction centres. It is possible that strong correlations between distant sites may enhance properties of excitation transport (for example, robustness, efficiency, regularity) through such light-harvesting networks. Further study is required to determine the precise role the non-classical correlations that are realized by this entanglement play in light-harvesting dynamics of larger systems. We note that a functional role for entanglement in LHCs is consistent with studies of the effects of intersite electronic coherence on kinetic properties of LHCs, which have found that such coherence benefits efficiency^{27–32} and robustness^{33,34} of excitation transport.

Our framework for quantification of entanglement in LHCs is an essential step in the precise characterization of quantum resources in organic structures, and is of particular relevance to the endeavour of constructing quantum devices from such structures. Although coherence is a feature of quantum mechanical evolution whose quantification is not directly operationally material, the non-classical correlations embodied by entanglement constitute an operational resource. Entanglement resulting from delocalization of a single particle among experimentally accessible locations (for example, an excitation among the chromophores of an LHC)

represents non-local correlations with compelling technological applications^{18,35}. In particular, the presence of entanglement in LHCs sets the stage for investigating the applicability of entanglement-enhanced measurement³⁶ in biological systems. Quantum metrology enabled by the use of entangled states has been demonstrated using photons and ions, and these engineered systems have been used to measure frequencies and phases to unprecedented precision (for example, refs 37,38). The application of ideas from entanglement-enhanced metrology to the improved measurement of biological properties in these naturally robust quantum-coherent systems is likely to be fruitful. Within this context, note that the observed enhanced radiative decay of delocalized electronic excitation states of LH1 and LH2 (ref. 16), which by the arguments of this paper are also entangled states in the single-excitation subspace, is consistent with the fact that entangled states can be more sensitive probes of the environment than separable states.

In addition to quantum-enhanced metrology, densely packed molecular aggregates such as LHCs have potential for constructing naturally robust quantum devices. For example, ultrafast quantum-state transfer facilitated by excitation migration along engineered or self-assembled chromophoric arrays could be a possible realization of the ‘quantum wires’ that are much sought after in quantum technology. Entanglement between subunits in such an array is essential for high-fidelity quantum transport, and, as our study

shows, such entanglement is possible in molecular aggregate structures even at room temperature. This conclusion is reinforced by recent studies at room temperature that have presented evidence for delocalization of excitations along molecular wires formed by self-assembling J -aggregates³⁹. Integration of such molecular aggregates with solid-state devices (for example, cavities⁴⁰) opens up the possibility of engineering controllable quantum-coherent soft-matter structures that can be used to distribute quantum states or entanglement. It should be noted that the control overhead for quantum-state transfer in such wires is not overly prohibitive—it has been shown that the only requirement for high-fidelity state transfer in randomly coupled linear chains is complete control (for example, optical addressability) over the end points of the linear chain⁴¹.

We conclude by returning to the foundational significance of entanglement: it represents a uniquely quantum form of strong correlation between physical systems. The identification of entanglement—which was referred to by Schrödinger as ‘the characteristic trait of quantum mechanics, the one that enforces its entire departure from classical lines of thought’⁴²—between spatially distinct components of a biological system under natural functioning conditions further expands the field of physical systems for which non-trivial and uniquely quantum signatures become manifest.

Methods

Measures of entanglement in LHCs. The single-excitation assumption valid for many LHCs enables us to formulate simple measures of entanglement in these systems. First, note that if ρ represents a single-excitation state in the site basis, we have the following proposition:

$$\rho \text{ entangled} \iff \rho_{ij} \neq 0 \text{ for some } i \neq j \quad (2)$$

Here, the right-hand side simply means that ρ has some coherence. Hence, coherence (in the site basis) in the single-excitation subspace is necessary and sufficient for entanglement.

The forward implication of proposition (2) is clearly seen from its contrapositive form: $\rho_{ij} = 0 \forall i \neq j \implies \rho$ separable. If $\rho_{ij} = 0 \forall i \neq j$, then the state has the form $\rho = \sum_i \rho_{ii} |i\rangle\langle i|$, which is clearly a separable state because each $|i\rangle$ is separable. To prove the backward implication, again consider its contrapositive form: ρ separable $\implies \rho_{ij} = 0 \forall i \neq j$. If ρ is separable it contains no entanglement, and specifically no bipartite entanglement between any two chromophores. The well-known mixed-state bipartite-entanglement measure for two-level systems, concurrence¹⁹, can be computed between chromophores 1 and 2. In the single-excitation subspace, it evaluates to $C_{12} = 2|\rho_{12}|$. Hence, zero bipartite entanglement between these chromophores implies $\rho_{12} = 0$. Similarly, by setting the bipartite entanglement between all pairs of chromophores to zero, we find that $\rho_{ij} = 0 \forall i \neq j$.

This property of states in the single-excitation subspace implies that an effective entanglement witness^{33,44} for these states is simply the sum of all coherences:

$$\mathcal{W} = \sum_{i < j} |\rho_{ij}|$$

That is, $\mathcal{W} > 0 \iff \rho$ entangled. Note that unlike most entanglement witnesses, which only present sufficient conditions for entanglement, in this case $\mathcal{W} > 0$ is both necessary and sufficient for entanglement. Practically, this property can therefore be used in experiments to detect entanglement: any non-zero off-diagonal component of the LHC single-excitation density matrix in the site basis is a signature of entanglement. Simplifications of this witness, which become sufficient conditions for entanglement, can also be formulated by restricting the sum to be over a subset of the off-diagonal elements—this could be useful in situations where accessing all off-diagonal elements is experimentally prohibitive.

Last, we can also construct a measure of global entanglement in this system on the basis of the relative entropy of entanglement, which is defined as²⁰

$$E[\rho] = \min_{\sigma \in D} S(\rho || \sigma)$$

where D is the set of separable states and S is the relative entropy function: $S(\rho || \sigma) = \text{tr}(\rho \ln \rho - \rho \ln \sigma)$. As the physically relevant Hilbert space for LHCs is restricted to the zero- and single-excitation subspaces, we define the natural photosynthetic entanglement to be the restricted relative entropy of entanglement in which the minimization is performed over $\sigma \in D^*$, the set of separable states in the zero- and single-excitation manifold. We know from Proposition 1 above

that the set of separable states in this restricted Hilbert space has diagonal form. Consequently, the minimization problem becomes

$$E[\rho] = \min_{p_i} \text{tr}(\rho \ln \rho - \rho \ln \sigma)$$

$$\text{subject to the constraint } \sum_{i=1}^N p_i = \text{tr} \rho$$

where $\sigma = \text{diag}(p_1, p_2, \dots, p_N)$ and the optimization constraint derives from the fact that we are only considering separable states with the same normalization as ρ . As the cost function $E[\rho]$ is convex in p_i and the constraint is linear, this optimization is easily solved, for example, by using a Lagrange multiplier to combine the cost function and constraint into a Lagrangian and then finding its stationary point. This results in an explicit expression for the closest separable state, namely $\sigma^* = \text{diag}(p_1^*, p_2^*, \dots, p_N^*)$ with $p_i^* = \rho_{ii}$, as well as the following measure of global entanglement in the state ρ :

$$E[\rho] = - \sum_{i=1}^N \rho_{ii} \ln \rho_{ii} - S(\rho)$$

with $S(\rho) = -\text{tr} \rho \ln \rho$ the von Neumann entropy of ρ . We note that, consistent with our discussion above, $E[\rho]$ is also a measure of coherence in the system, because the first term on the right-hand side is the entropy of a state for which all coherences ρ_{ij} are artificially set to zero and the von Neumann entropy implicitly contains all coherences. We show in the Supplementary Information that this photosynthetic entanglement measure satisfies the properties of an entanglement monotone⁴ within the restricted zero- and single-excitation Hilbert space.

Realistic modelling of excitation dynamics in LHCs. Electronic excitation dynamics is governed by a Hamiltonian of the form $H = H_{\text{el}} + H_{\text{el-env}} + H_{\text{env}}$, where

$$H_{\text{el}} = \sum_{i=1}^N E_i |i\rangle\langle i| + \sum_{i=1}^N \sum_{j>i}^N J_{ij} (|i\rangle\langle j| + |j\rangle\langle i|)$$

describes the closed system dynamics of the N LHC chromophores, including on-site energies, E_j , and coupling terms, J_{ij} , that describe the coupling between the chromophores. The remaining terms in H describe the coupling between the electronic degrees of freedom of the LHC chromophore molecules and their environment, which typically consists of surrounding proteins, the electromagnetic field and reaction centre(s) to which the LHC is affixed. The dominant environmental perturbations are the phonons of the protein scaffolding around the LHC, and these are modelled as a diagonal coupling to a phonon bath:

$$H_{\text{el-env}} = \sum_{i=1}^N |i\rangle\langle i| \sum_{\xi} g_{\xi}^i Q_{\xi}^i \quad (3)$$

where Q_{ξ}^i is a phonon mode indexed by ξ , which is coupled to chromophore i with coupling strength g_{ξ}^i . Obtaining a faithful reduced description of the effective dynamics of just the electronic excitation state ρ requires a physically accurate averaging over the external degrees of freedom, and quantum master-equation approaches are commonly used for this averaging. In particular, the standard Redfield equation⁴⁵ is often used to explore the dynamics of photosynthetic excitation-energy transfer. This equation is valid when the exciton-phonon coupling—which can be specified by the magnitude of the reorganization energy—is much smaller than the interchromophoric coupling, because the equation is derived on the basis of a second-order perturbative truncation with respect to the exciton-phonon coupling. However, in many LHCs, the reorganization energies are not small in comparison with the interchromophoric coupling: for example, in the FMO protein the electronic coupling strengths span a wide range, 1–100 cm^{-1} , and the reorganization energies span a similar range^{22,26,46,47}. Hence, the standard Redfield-equation approach might lead to erroneous insights and incorrect conclusions regarding quantum-coherent effects in the FMO complex⁴⁸. Two of the present authors have recently presented a reliable theoretical framework to describe photosynthetic excitation-energy transfer that takes into account the phonon-relaxation dynamics associated with each chromophore in chromophore-protein complexes³⁴. This framework can describe quantum-coherent wave-like motion and incoherent hopping motion in a unified manner, and reduces to the standard Redfield theory and Förster theory⁴⁵ in their respective limits of validity. The ability of the framework to interpolate between these two limits is significant because photosynthetic excitation-energy transfer commonly occurs between these two perturbative regimes, as in the case of the FMO complex. The only assumptions used in this framework are that first, the exciton-phonon coupling is bilinear (for example, equation (3)), second, the environmental fluctuations are described as Gaussian processes, third, the total system is in a factorized (product) initial state and fourth, an overdamped Brownian oscillator model for the phonon environment, which results in exponentially (time-) correlated phonon fluctuations. The hierarchical expansion technique⁴⁹ is employed to obtain a practical expression for numerical calculations³⁴.

For the simulations in this work we take the reorganization energy of the molecular environment to be 35 cm^{-1} , a value that is consistent with experimentally extracted reorganization energies for this complex^{22,26,46,47}. We choose a phonon

relaxation time of 100 fs, and a reaction-centre trapping rate associated with site 3 of $(4 \text{ ps})^{-1}$, both of which are consistent with the literature on LHCs (refs 22,28,47). We do not assume any form of spatial correlations in the phonon fluctuations. This is primarily because there are no experimental data on spatially correlated phonon fluctuations in the FMO protein and the goal here is for the simulations to be as close as possible to what is currently known about the FMO environment. We note that, at short times, spatially correlated fluctuations are expected to increase coherence in the site basis, and hence will naturally increase the entanglement present at initial times. Therefore, the predictions in this work, which are based on simulations that assume no spatial correlations in the environment, could be viewed as a lower bound on the amount of entanglement in the FMO complex.

Finally, we have compared the entanglement predicted by the present framework with that predicted by standard Markovian Redfield models. This comparison is presented in the Supplementary Information.

Received 26 May 2009; accepted 13 March 2010; published online 25 April 2010

References

- Einstein, A., Podolsky, B. & Rosen, B. Can quantum-mechanical description of physical reality be considered complete?. *Phys. Rev.* **47**, 777–780 (1935).
- Schrödinger, E. Die gegenwärtige Situation in der Quantenmechanik. *Naturwissenschaften* **23**, 807–812 (1935).
- Amico, L., Fazio, R., Osterloh, A. & Vedral, V. Entanglement in many-body systems. *Rev. Mod. Phys.* **80**, 517–576 (2008).
- Horodecki, R., Horodecki, P., Horodecki, M. & Horodecki, K. Quantum entanglement. *Rev. Mod. Phys.* **81**, 865–942 (2009).
- Vedral, V. Quantifying entanglement in macroscopic systems. *Nature* **453**, 1004–1007 (2008).
- Vedral, V. Entanglement production in non-equilibrium thermodynamics. *J. Phys. Conf. Ser.* **143**, 012010–012018 (2007).
- Quiroga, L., Rodriguez, F. J., Ramirez, M. E. & Paris, R. Nonequilibrium thermal entanglement. *Phys. Rev. A* **75**, 032308 (2007).
- Cai, J., Popescu, S. & Briegel, H. J. Dynamic entanglement in oscillating molecules and potential biological implications. Preprint at <http://arxiv.org/abs/0809.4906> (2008).
- Thorwart, M., Eckel, J., Reina, J. H., Nalbach, P. & Weiss, S. Enhanced quantum entanglement in the non-Markovian dynamics of biomolecular excitons. *Chem. Phys. Lett.* **478**, 234–237 (2009).
- Engel, G. S. *et al.* Evidence for wavelike energy transfer through quantum coherence in photosynthetic systems. *Nature* **446**, 782–786 (2007).
- Lee, H., Cheng, Y.-C. & Fleming, G. R. Coherence dynamics in photosynthesis: Protein protection of excitonic coherence. *Science* **316**, 1462–1465 (2007).
- Collini, E. *et al.* Coherently wired light-harvesting in photosynthetic marine algae at ambient temperature. *Nature* **463**, 644–648 (2010).
- Panitchayangkoon, G. *et al.* Long-lived quantum coherence in photosynthetic complexes at physiological temperature. Preprint at <http://arxiv.org/abs/1001.5108> (2010).
- Blankenship, R. E. *Molecular Mechanisms of Photosynthesis* (Wiley-Blackwell, 2002).
- Pullerits, T., Chachisvilis, M. & Sundstrom, V. Exciton delocalization length in the B850 antenna of *Rhodobacter sphaeroides*. *J. Phys. Chem.* **100**, 10787–10792 (1996).
- Monshouwer, R., Abrahamsson, M., van Mourik, F. & van Grondelle, R. Superradiance and exciton delocalization in bacterial photosynthetic light-harvesting systems. *J. Phys. Chem. B* **101**, 7241–7248 (1997).
- van Amerongen, H., Valkunas, L. & van Grondelle, R. *Photosynthetic Excitons* (World Scientific, 2000).
- van Enk, S. J. Single-particle entanglement. *Phys. Rev. A* **72**, 064306 (2005).
- Hill, S. & Wootters, W. K. Entanglement of a pair of quantum bits. *Phys. Rev. Lett.* **78**, 5022–5025 (1997).
- Vedral, V., Plenio, M. B., Rippin, M. A. & Knight, P. L. Quantifying entanglement. *Phys. Rev. Lett.* **78**, 2275–2279 (1997).
- Camara-Artigas, A., Blankenship, R. E. & Allen, J. P. The structure of the FMO protein from *Chlorobium tepidum* at 2.2 Å resolution. *Photosynth. Res.* **75**, 49–55 (2003).
- Adolphs, J. & Renger, T. How proteins trigger excitation energy transfer in the FMO complex of green sulphur bacteria. *Biophysical J.* **91**, 2778–2797 (2006).
- Ishizaki, A. & Fleming, G. R. Unified treatment of quantum coherent and incoherent hopping dynamics in electronic energy transfer: Reduced hierarchy equations approach. *J. Chem. Phys.* **130**, 234111 (2009).
- Wen, J., Zhang, H., Gross, M. L. & Blankenship, R. E. Membrane orientation of the FMO antenna protein from *Chlorobaculum tepidum* as determined by mass spectrometry-based footprinting. *Proc. Natl Acad. Sci. USA* **106**, 6134–6139 (2009).
- Meier, T., Zhao, Y., Chernyak, V. & Mukamel, S. Polarons, localization, and excitonic coherence in superradiance of biological antenna complexes. *J. Chem. Phys.* **107**, 3876–3893 (1997).
- Read, E. L. *et al.* Visualization of excitonic structure in the Fenna–Matthews–Olson photosynthetic complex by polarization-dependent two-dimensional electronic spectroscopy. *Biophysical J.* **95**, 847–856 (2008).
- Mohseni, M., Rebentrost, P., Lloyd, S. & Aspuru-Guzik, A. Environment-assisted quantum walks in energy transfer of photosynthetic complexes. *J. Chem. Phys.* **129**, 174106 (2008).
- Plenio, M. B. & Huelga, S. F. Dephasing-assisted transport: Quantum networks and biomolecules. *New J. Phys.* **10**, 113019 (2008).
- Mukai, K., Abe, S. & Sumi, H. Theory of rapid excitation-energy transfer from B800 to optically-forbidden exciton states of B850 in the antenna system LH2 of photosynthetic purple bacteria. *J. Phys. Chem. B* **103**, 6096–6102 (1999).
- Scholes, G. D. & Fleming, G. R. On the mechanism of light harvesting in photosynthetic purple bacteria: B800 to B850 energy transfer. *J. Phys. Chem. B* **104**, 1854–1868 (2000).
- Jang, S., Newton, M. D. & Silbey, R. J. Multichromophoric Förster resonance energy transfer. *Phys. Rev. Lett.* **92**, 218301 (2004).
- Sumi, H. Bacterial photosynthesis begins with quantum-mechanical coherence. *Chem. Record* **1**, 480–493 (2001).
- Cheng, Y.-C. & Silbey, R. J. Coherence in the B800 ring of purple bacteria LH2. *Phys. Rev. Lett.* **96**, 028103 (2006).
- Ishizaki, A. & Fleming, G. R. Theoretical examination of quantum coherence in a photosynthetic system at physiological temperature. *Proc. Natl Acad. Sci. USA* **106**, 17255–17260 (2009).
- Lombardi, E., Sciarrino, F., Popescu, S. & De Martini, F. Teleportation of a vacuum-one-photon qubit. *Phys. Rev. Lett.* **88**, 070402 (2002).
- Giovannetti, V., Lloyd, S. & Maccone, L. Quantum-enhanced measurements: Beating the standard quantum limit. *Science* **306**, 1330–1336 (2004).
- Roos, C. F., Chwalla, M., Kim, K., Riebe, M. & Blatt, R. ‘Designer atoms’ for quantum metrology. *Nature* **443**, 316–319 (2006).
- Nagata, T., Okamoto, R., O’Brien, J. L., Sasaki, K. & Takeuchi, S. Beating the standard quantum limit with four-entangled photons. *Science* **316**, 726–729 (2007).
- Lagoudakis, P. G. *et al.* Experimental evidence for exciton scaling effects in self-assembled molecular wires. *Phys. Rev. Lett.* **93**, 257401 (2004).
- Tischler, J. R. *et al.* Solid state cavity QED: Strong coupling in organic thin films. *Org. Electron.* **8**, 94–113 (2007).
- Burgarth, D. & Bose, S. Perfect quantum state transfer with randomly coupled quantum chains. *New J. Phys.* **7**, 135–147 (2005).
- Schrödinger, E. Discussion of probability relations between separated systems. *Proc. Cambridge Phil. Soc.* **31**, 555–563 (1935).
- Horodecki, M., Horodecki, P. & Horodecki, R. Separability of mixed states: Necessary and sufficient conditions. *Phys. Lett. A* **223**, 1–8 (1996).
- Peres, A. Separability criterion for density matrices. *Phys. Rev. Lett.* **77**, 1413–1415 (1996).
- May, V. & Kühn, O. *Charge and Energy Transfer Dynamics in Molecular Systems* (Wiley-VCH, 2004).
- Brixner, T. *et al.* Two-dimensional spectroscopy of electronic couplings in photosynthesis. *Nature* **434**, 625–628 (2005).
- Cho, M., Vaswani, H. M., Brixner, T., Stegner, J. & Fleming, G. R. Exciton analysis in 2D electronic spectroscopy. *J. Phys. Chem. B* **109**, 10542–10556 (2005).
- Ishizaki, A. & Fleming, G. R. On the adequacy of the Redfield equation and related approaches to the study of quantum dynamics in electronic energy transfer. *J. Chem. Phys.* **130**, 234110 (2009).
- Tanimura, Y. Stochastic Liouville, Langevin, Fokker–Planck, and master equation approaches to quantum dissipative systems. *J. Phys. Soc. Jpn* **75**, 082001 (2006).

Acknowledgements

We are grateful to Y.-C. Cheng, J. Dawlaty, V. Vedral and M. Plenio for conversations and comments. This material is based on work supported by DARPA under award No N66001-09-1-2026. This work was supported by the Director, Office of Science, Office of Basic Energy Sciences of the US Department of Energy under contract No DE-AC02-05CH11231 and by the Chemical Sciences, Geosciences and Biosciences Division, Office of Basic Energy Sciences, US Department of Energy under contract DE-AC03-76SF00098. A.I. appreciates the support of a Japan Society for the Promotion of Science (JSPS) Postdoctoral Fellowship for Research Abroad.

Author contributions

Calculations were carried out by M.S. and A.I. All authors contributed extensively to the planning, discussion and writing up of this work.

Additional information

The authors declare no competing financial interests. Supplementary information accompanies this paper on www.nature.com/naturephysics. Reprints and permissions information is available online at <http://npq.nature.com/reprintsandpermissions>. Correspondence and requests for materials should be addressed to M.S.