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Dynamics and mechanism of light harvesting in UV photoreceptor UVR8†

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Photosynthetic pigments form light-harvesting networks to enable nearly perfect quantum efficiency in photosynthesis via excitation energy transfer. However, similar light-harvesting mechanisms have not been reported in light sensing processes in other classes of photoreceptors during light-mediated signaling. Here, based on our earlier report, we mapped out a striking energy-transfer network composed of 26 structural tryptophan residues in the plant UV-B photoreceptor UVR8. The spectra of the tryptophan chromophores are tuned by the protein environments, funneling all excitation energy to a cluster of four tryptophan residues, a pyramid center, where the excitation-induced monomerization is initiated for cell signaling. With extensive site-directed mutagenesis, various time-resolved fluorescence techniques, and combined QM/MM simulations, we determined the energy-transfer rates for all donor–acceptor pairs, revealing the time scales from tens of picoseconds to nanoseconds. The overall light harvesting quantum efficiency by the pyramid center is significantly increased to 73%, compared to a direct excitation probability of 35%. UVR8 is the only photoreceptor discovered so far using a natural amino-acid tryptophan without utilizing extrinsic chromophores to form a network to carry out both light harvesting and light perception for biological functions.

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Introduction

Sunlight is essential to living organisms both as the energy source for photosynthesis and as an environmental signal for biological functions. For the former, different photosynthetic organisms have evolved various light-harvesting complexes, including chlorosomes,¹ phycobilisomes,² FMO complexes,³ light-harvesting complex I (LHCI)⁴ and light-harvesting complex II (LHCII).⁵ These antenna complexes contain multiple spatially arranged photosynthetic pigments, which can absorb photon energy and then transmit the excitation energy to reaction centers for energy conversion, ensuring the efficiency of photosynthesis. For light-triggered signaling, living systems use

a variety of photoreceptors for the initial sensing of different regions of the solar spectrum, mediating critical biological processes of vision, circadian rhythm, photomorphogenesis and ultraviolet (UV) protection.^{6–10} In contrast with photosynthetic complexes, most photoreceptors contain only one chromophore, and thus, both the photon absorption and the subsequent photoreactions that lead to protein conformational changes occur at the same site without excitation energy transfer. Interestingly, a recently discovered plant UV-B (280–315 nm) photoreceptor, UVR8 (UV RESISTANCE LOCUS 8), is unique among all photoreceptor proteins, in which there is no external chromophore present, but multiple tryptophan residues in the protein sequence are believed to be its chromophores. UVR8 potentially provides the first example of photoreceptors that use highly organized light-harvesting networks.¹⁰

UVR8 forms a homodimer whose interface is decorated with multiple inter-subunit salt-bridges and hydrogen-bonding interactions in the ground or “dark” state.^{10–12} Upon absorption of UV-B irradiation, the UVR8 dimers spontaneously dissociate into monomers, which bind the key signaling protein Constitutively Photomorphogenic 1 (COP1) to trigger various protective mechanisms against UV damage.^{10–23} The high-resolution X-ray structure of *Arabidopsis thaliana* UVR8 has been solved in the dimeric form (Fig. 1a and b), revealing a 7-blade beta-propeller architecture of each monomer.^{11,12} Unlike other visible and infrared photoreceptors, UVR8 does not

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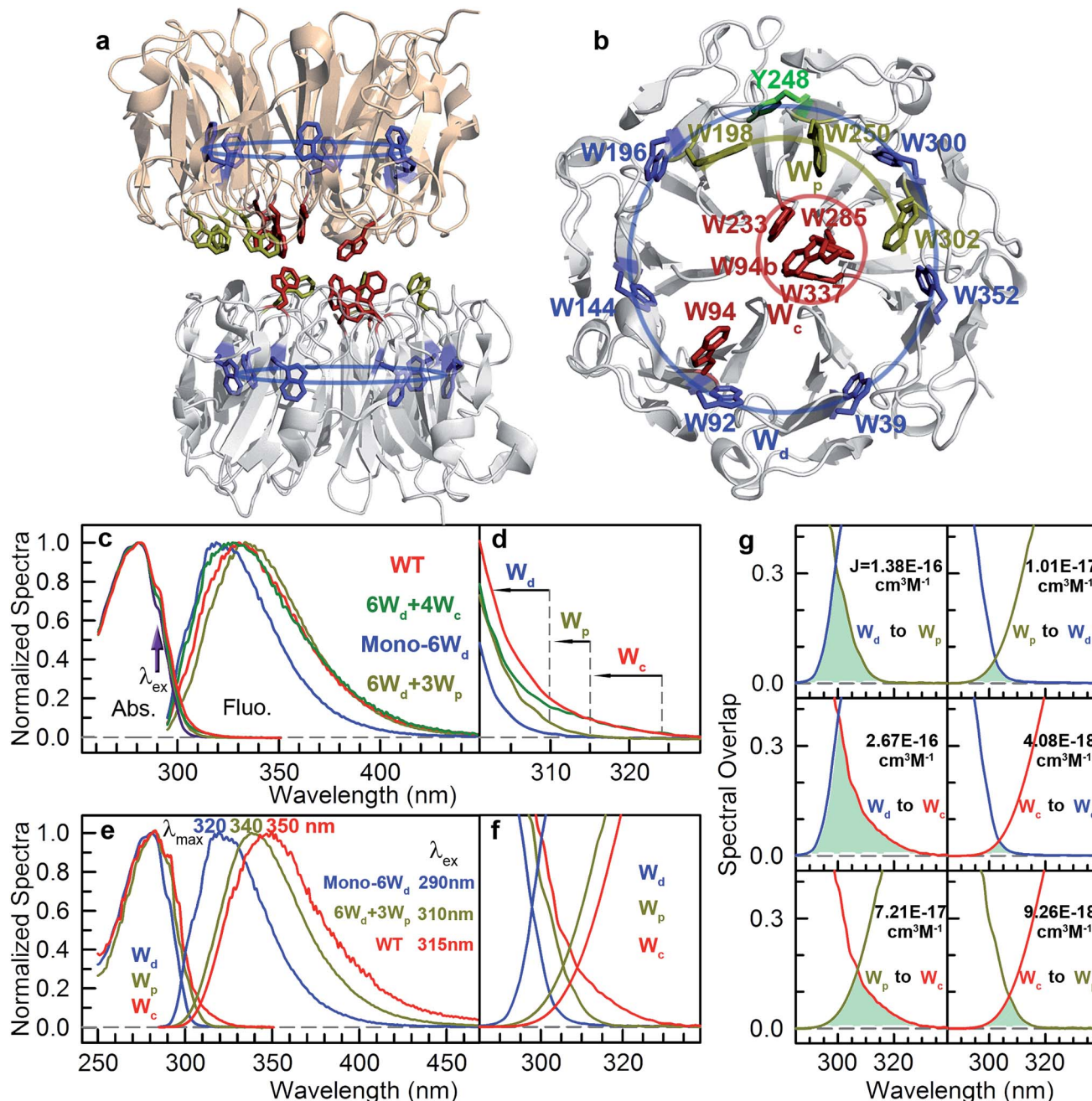


Fig. 1 UVR8 light-harvesting tryptophan and steady-state spectra. (a) A side view of UVR8 dimer with 3 groups of W residues highlighted in different colors: 6W_d, blue; 3W_p, dark yellow; and 4W_c, dark red. (b) A top view of one monomer shows symmetrical locations of 6W_d and relative positions of 3W_p/4W_c. W94b is from the other monomer. (c) Normalized absorption and emission spectra for UVR8 WT and 3 mutants. (d) The absorption spectra show differences beyond 300 nm wavelength. (e) Normalized absorption and emission spectra for W_d, W_p and W_c separately. Note the different excitation wavelengths for selected W_d (290 nm), W_p (310 nm) and W_c (315 nm) only. (f) A close view of the absorption-emission overlap region. (g) Spectral overlap integrals for energy transfer.

contain any external chromophores and utilizes the UV-absorbing natural amino acid tryptophan (Trp or W) to perceive UV-B. Each UVR8 monomer has 14 tryptophan residues, which, except one tryptophan in the flexible C-terminal region (not included in the X-ray structure), can be classified into 3 distinct groups (Fig. 1a and b): pyramid center Trp (W_c, 4 of the 13), peripheral Trp (W_p, 3 of the 13) and distal Trp (W_d, 6 of the 13). One UVR8 dimer has two symmetry-related pyramid

centers (4W_c); each is formed by van der Waals clustering of 3W_c (W285, W233 and W337) from one monomer and a fourth W_c (W94) from the opposing monomer. There are also three peripheral Trp residues, 3W_p (W198, W250 and W302), at the dimer interface surrounding the pyramid 4W_c. Furthermore, six Trp residues, 6W_d, are buried in the middle of the beta-propellers and form a highly symmetrical ring with another tyrosine residue (Fig. 1b). Among these tryptophan residues,

only two pyramid center Trp (W285 and W233) of the W_c pyramid are critical to the light-induced monomerization.^{11,12,24–30} Previously, we found a fast fluorescence quenching reaction in ~ 150 picoseconds among Trp residues of the pyramid cluster, likely using excitation energy on W_c to initiate dimer dissociation.²⁸ One key question is whether or not other tryptophan residues form a light-harvesting network to transmit excitation energy to the pyramid center ($4W_c$), enhancing UVR8 quantum yield for sensitive detection of the relatively low intensity of UV-B radiation. Previously, we reported a Trp-to-Trp energy transfer observation,²⁸ which was soon supported by other theoretical results.^{29,30} However, there is no direct experimental evidence of light harvesting; the time scales and scheme of the excitation energy transfer process are completely unknown.

In this work, we first designed tens of UVR8 mutants to remove certain tryptophan residues to isolate the system into three Trp spectral groups. We next used three different time-resolved fluorescence spectroscopy methods to completely measure the energy-transfer dynamics in various UVR8 mutants. Then, we applied combined quantum mechanical and molecular mechanical (QM/MM) simulations and Förster resonance energy transfer (FRET) theory to determine the energy transfer rates, providing further insights into the experimental data. These observations exclude a coherent energy-transfer behavior, as observed in photosynthesis, which is attributed to the large distances between the donors and acceptors in UVR8. Combining all experimental and theoretical results, we revealed a remarkable light-harvesting network in the soluble photoreceptor UVR8, which significantly increases the overall light-sensing efficiency.

Results and discussion

Unique spectra of the three tryptophan groups

Since excitation energy flows downhill, we first investigated the possible resonance energy transfer (RET) direction by obtaining the spectra of 3 Trp groups from mutagenesis studies (mutants summarized in Table S1†). We can knockout either the pyramid center (mutant W233/285/337/94F or $6W_d + 3W_p$) or the peripheral outlier (mutant W198/250/302F or $6W_d + 4W_c$) by replacing tryptophan with phenylalanine (Phe or F) residues to measure clean spectra of the three Trp groups. However, replacement of all seven interfacial tryptophan residues ($3W_p/4W_c$) to F leads to severe inclusion body formation with no functional protein yield, probably due to protein misfolding. We identified that the W302F mutant is the cause for this problem. We then used W94/233/285/337/198/250F or $6W_d + W302$ as a starting point and replaced the W302 position with all other 18 natural amino acids, but still failed to obtain any well-folded mutant. By further mutating an arginine (Arg or R) residue R286 involved in interfacial salt-bridges to a neutral alanine (Ala or A) residue and W302 to histidine (His or H), a well-folded monomeric protein (mono- $6W_d$) could be purified with all $3W_p/4W_c$ removed. Since previous FTIR experiments have suggested that the UVR8 dimer and monomer conformations have similar overall tertiary structure,³¹ the local

environments of buried distal tryptophan residues in the monomer mutant should be the same as those in dimers.

Fig. 1c shows the absorption and emission spectra of UVR8 WT and 3 mutants with certain tryptophan group(s) removed. Although absorption spectra of different protein samples have similar overall shapes with the peak at 282 nm and a shoulder at 290 nm, significant differences were observed in the red region beyond 300 nm (Fig. 1d). For WT and $6W_d + 4W_c$, in which the Trp pyramid center is intact, the absorption spectra extend to above 320 nm, which may result from the exciton couplings among the 4 closely packed W_c .^{11,32} For $6W_d + 3W_p$, although absorbance in the red tail region is much lower than WT due to mutation of the pyramid center, the spectrum still extends to

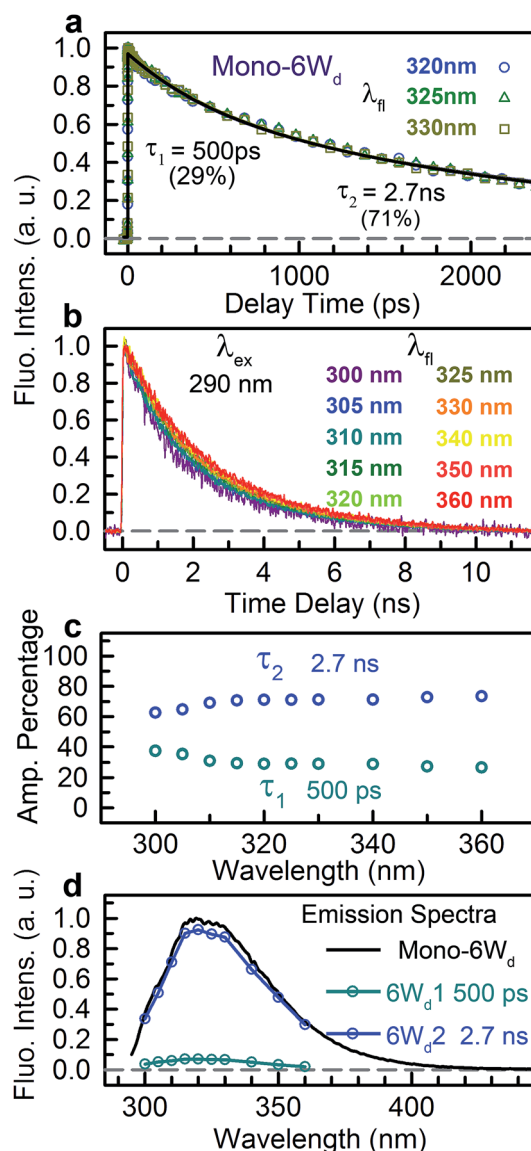


Fig. 2 Fluorescence lifetimes of $6W_d$. (a) Fluorescence transients at fs time resolution of mono- $6W_d$. (b) Fluorescence transients of mono- $6W_d$ measured by ps-resolved TCSPC. Note the fluorescence wavelengths labeled. (c) Amplitude percentages of 0.5 ns and 2.7 ns lifetimes at different fluorescence wavelengths. (d) Lifetime-associated spectra of 0.5 ns ($6W_d1$) and 2.7 ns ($6W_d2$) time constants.



315 nm. In contrast, the mutant mono-6W_d, by mutating all interfacial 3W_p/4W_c, has negligible absorbance beyond 310 nm, indicating that W_d exhibits the bluest absorption among the 3 groups.

From the absorption spectra of the protein samples, absorption spectra of 3 groups of tryptophan residues were readily obtained by spectral subtraction (Fig. 1e). The spectrum data indicate that light with wavelength beyond 310 nm can be used to selectively pump W_p/W_c without exciting W_d. We further measured the emission spectra at 3 different excitation wavelengths (290 nm, 300 nm and 310 nm) with mutants containing 6W_d and various W_p. As shown in Fig. S1,† the emission spectra gradually red shift as the excitation is moved to longer wavelengths, due to the change of excitation ratios of W_p to W_d at various wavelengths. Spectra at 310 nm excitation exhibit a similar shape for all 6W_d + W_p mutants (Fig. S1h†), suggesting that 3W_p have nearly the same emission spectra. Absorption of 4W_c is further red shifted, with a red tail extending to above 325 nm. Thus, 315 nm light can be used to selectively excite 4W_c in UVR8 WT or 6W_d + 4W_c to obtain clean W_c fluorescence; 310 nm excitation in 6W_d + 3W_p provides W_p emission; 290 nm light was applied to mono-6W_d to acquire the W_d spectrum.

Emission spectra of the 3 groups of tryptophan are dramatically different (Fig. 1e): W_d shows a very blue emission with

a peak at 320 nm; W_p has a fluorescence spectrum with a maximum at 340 nm (6W_d not excited at 310 nm); W_c exhibits the reddest emission (peak at 350 nm) among the 3 groups (only W_c excited at 315 nm). From our previous extensive studies of tryptophan fluorescence in proteins, W_d emission at 320 nm is unusually blue-shifted and typically from highly buried tryptophan, consistent with W_d locations in the middle of beta-propellers without water molecules within 5 Å. We also noticed that the W_d emission spectrum is structured and thus cannot be fitted with a lognormal function³³ that fits typically Trp fluorescence spectra. A structured emission spectrum was also observed for a highly buried Trp in azurin.³⁴ In contrast, W_p and W_c, located at the dimer interface, display redder emission compared to W_d. Water molecules and possibly nearby charged residues stabilize the tryptophan ¹L_a excited state, which has a large dipole moment.³⁵

As shown in Fig. 1f, there are overlapping regions between absorption and emission spectra of 3 Trp groups, leading to possible RET. The tryptophan spectra in Fig. 1e reveal an energetic ordering that favors excitation-energy flow from 6W_d to 3W_p/4W_c and from 3W_p to 4W_c. As shown in Fig. 1g, the spectral overlap integrals (*J*, eqn (6)) for W_d-to-W_p, W_d-to-W_c and W_p-to-W_c RET are $1.38 \times 10^{-16} \text{ cm}^3 \text{ M}^{-1}$, $2.67 \times 10^{-16} \text{ cm}^3 \text{ M}^{-1}$ and $7.21 \times 10^{-17} \text{ cm}^3 \text{ M}^{-1}$, respectively, whereas *J* values

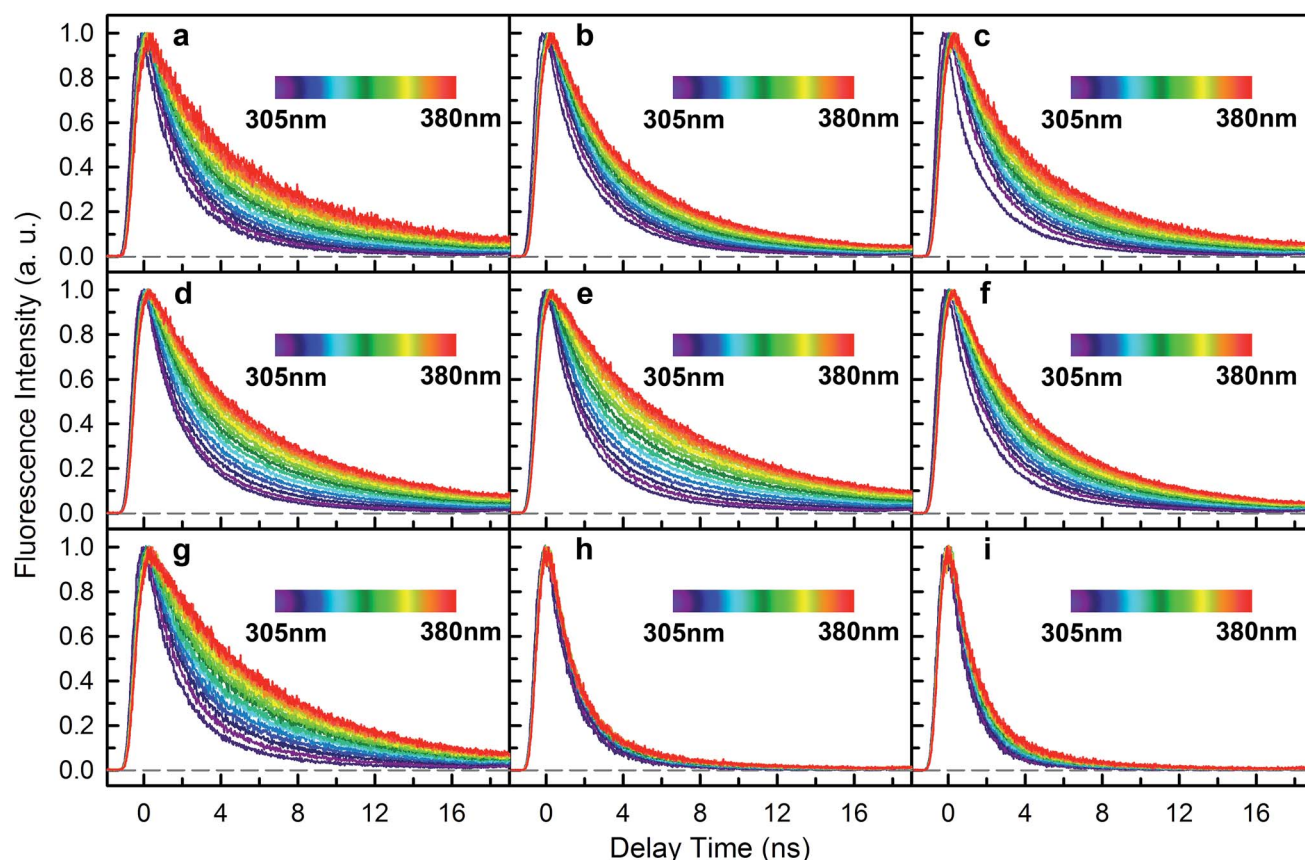


Fig. 3 Fluorescence decay transients ($\lambda_{\text{ex}} = 290 \text{ nm}$) from 305 nm to 380 nm (5 nm intervals) for UVR8 WT and 8 mutants measured with sub-nanosecond-resolved time correlated single photon counting (TCSPC). (a) 4W_c/198/250F (6W_d + W302). (b) 4W_c/198/302F (6W_d + W250). (c) 4W_c/250/302F (6W_d + W198). (d) 4W_c/198F (6W_d + W250/302). (e) 4W_c/250F (6W_d + W198/302). (f) 4W_c/302F (6W_d + W198/250). (g) 6W_d + 3W_p. (h) 6W_d + 4W_c. (i) WT.



for the corresponding backward transfer are $1.01 \times 10^{-17} \text{ cm}^3 \text{ M}^{-1}$, $4.08 \times 10^{-18} \text{ cm}^3 \text{ M}^{-1}$ and $9.26 \times 10^{-18} \text{ cm}^3 \text{ M}^{-1}$, respectively, which are much smaller than those for the forward transfer. Thus, the backward RET is negligible and the excitation energy is transferred in one direction from the $6W_d$ ring to the interface Trp, where the energy is then funneled to the critical $4W_c$ center. Also, the intra-group J values between $6W_d$ or $3W_p$ are much smaller than those between inter-groups. Photosynthesis light harvesting often uses a variety of pigments with different spectra to govern energy-transfer direction.³⁶ Here, UVR8 contains only one type of chromophore, the amino acid tryptophan, with spectral tuning by local environments to create a driving force for excitation energy flow.

Resonance energy transfer from distal W_d ring to interfacial W_p and W_c

We next experimentally studied RET dynamics from $6W_d$ to the dimer interface by determining lifetime changes of W_d with and without the W_p/W_c energy acceptors. First, we need to determine the lifetimes of mono- $6W_d$ and related emission spectra without RET. Fig. 2a shows femtosecond-resolved fluorescence transients of mono- $6W_d$, which have two lifetimes without RET and show no ultrafast component. Combining the fs- and ps-resolved (Fig. 2a and b) data, the two lifetimes of $6W_d$ were determined to be 500 ps and 2.7 ns, with amplitude percentages of 29% and 71%, respectively, at the emission peak of 320 nm.

As shown in Fig. 2c, the amplitude ratio of the 500 ps component gradually increases at shorter fluorescence wavelengths. With the fitted amplitudes from 300 nm to 360 nm, the associated spectra of the two W_d lifetimes were decomposed from the W_d total fluorescence spectrum (Fig. 2d, $6W_d1$ and $6W_d2$). The contribution from the 0.5 ns spectrum is minor; the 2.7 ns component is dominant due to a longer timescale and larger amplitude.

To investigate RET dynamics and determine the transfer timescales, we measured the fluorescence transients of a series of mutants containing $6W_d$ and various interfacial Trp residues (Table S1†): $6W_d + 1W_p$ (Fig. 3a–c), $6W_d + 2W_p$ (Fig. 3d–f), $6W_d + 3W_p$ (Fig. 3g), $6W_d + 4W_c$ (Fig. 3h) and WT (Fig. 3i). We expect to observe decreased W_d lifetimes due to additional RET channels. For each mutant, transients from 16 detection wavelengths (305 nm to 380 nm with 5 nm intervals) were measured using sub-nanosecond resolved TCSPC in order to follow the complete decay dynamics. For $6W_d + W_p$ mutants in Fig. 3a–g, transients at various wavelengths are dramatically different: fluorescence at the red side contains more of slow dynamics in 6–8 ns from W_p lifetimes; data at shorter wavelengths show more contribution from $6W_d$ dynamics because of their blue emission. In $6W_d + 4W_c$, however, all TCSPC transients look similar because the fluorescence in $4W_c$ is dominantly quenched in 80 ps (see below) and does not show significant contributions under this sub-ns time resolution, leaving the majority signal from $6W_d$ in all wavelengths. Significantly, we did not observe much 6–8 ns components (W_p lifetime without RET) in WT (Fig. 3i),

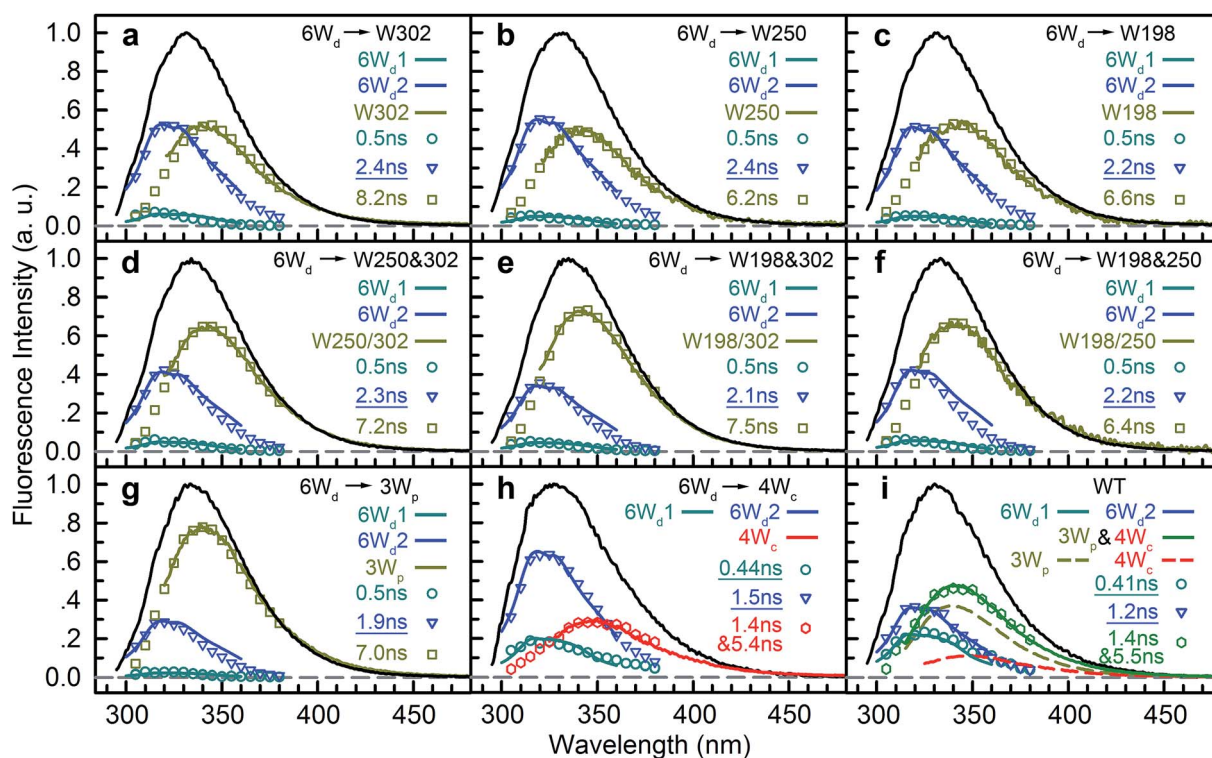


Fig. 4 Time scales from a global fitting and lifetime-associated spectra. (a) $4W_c/198/250F$ ($6W_d + W302$). (b) $4W_c/198/302F$ ($6W_d + W250$). (c) $4W_c/250/302F$ ($6W_d + W198$). (d) $4W_c/198F$ ($6W_d + W250/302$). (e) $4W_c/250F$ ($6W_d + W198/302$). (f) $4W_c/302F$ ($6W_d + W198/250$). (g) $6W_d + 3W_p$. (h) $6W_d + 4W_c$. (i) WT.



suggesting that W_p lifetimes become faster due to energy transfer to $4W_c$.

To determine W_d lifetimes with RET, the TCSPC data in Fig. 3 were analyzed using global fitting, in which transients at different detection wavelengths were fitted using the same set of time constants with varying amplitude percentages (fitting parameters in Tables S2–S10†). Lifetime associated spectra were constructed for the observed time scales (symbols in Fig. 4) and were then compared with directly measured spectra (solid lines) to determine their origin. The decay transients of the $6W_d + W_p$ mutants (Fig. 4a–g) were fitted with 3 exponential components: 0.50 ns, 1.9–2.4 ns and 6–8 ns. Since the 6–8 ns lifetimes were not observed in mono- $6W_d$ and their associated spectra are the same as W_p emission, they are from fluorescence lifetimes of W_p . Owing to different local environments, W198, W250 and W302 show time scales of 6.6 ns, 6.2 ns and 8.2 ns, respectively (Fig. 4a–c). The 0.50 ns lifetime ($6W_d1$) of W_d is much faster than RET time scales to W_p and does not change in the presence of the W_p acceptors. The 1.9–2.4 ns components come from the original 2.7 ns W_d lifetime ($6W_d2$) and are faster with additional RET channels to W_p . Moreover, this time scale decreases when more W_p are present, consistent with faster energy transfer with more RET acceptors. With the numbers from the three “ $6W_d + 1W_p$ ” mutants and ~3% fitting error, the RET to W198, W250 and W302 was calculated to be 12 ± 1.4 ns, 22 ± 4.3 ns and 24 ± 4.9 ns, respectively. By adding the 3 rates together, the total RET time is 5.9 ns, agreeing well with the 6.4 ± 0.6 ns time measured directly from the $6W_d + 3W_p$ mutant.

For $6W_d + 4W_c$ and WT (Fig. 4h and i), four decay components were needed to globally fit the transients. In $6W_d + 4W_c$, judging from the associated spectra, the 0.44 ns and 1.5 ns are the 2 lifetimes of W_d ; the 1.4 ns and 5.4 ns are from $4W_c$. The lifetime of $6W_d1$ is decreased to 0.44 ns from the previous 0.50 ns due to faster RET to $4W_c$ than to W_p . The $6W_d2$ lifetime changed to 1.5 ns from 2.7 ns, resulting in a 3.4 ± 0.2 ns RET time scale. In WT, the 0.41 ns associated spectrum is redder (peak at 330 nm) than the $6W_d1$ spectrum, suggesting possible W_p -to- W_c energy transfer on similar time scales, leading to mixing of the W_p emission in this component. The 2.7 ns lifetime changes to 1.2 ns by the 2.2 ± 0.2 ns RET to all interfacial Trp, the same as the total RET time calculated with separately measured 6.4 ± 0.6 ns transfer to $3W_p$ and 3.4 ± 0.2 ns RET to $4W_c$. WT shows the fastest RET in all samples, leading to a $44 \pm 0.6\%$ energy-transfer efficiency of $6W_d$.

To explain the two other time scales observed in $6W_d + 4W_c$ and WT, we studied $4W_c$ fluorescence decay dynamics. By 315 nm excitation with ps-resolved TCSPC in WT, we found that $4W_c$ has two lifetimes (Fig. 5a): 80 ps (75%) and 1.4 ns (25%). As shown in Fig. 5b, 315 nm and 320 nm excitation transients are similar since both only excite $4W_c$, whereas 290 nm and 310 nm light bring in W_d/W_p signals, leading to slower overall dynamics. In sub-ns resolved TCSPC data, where 80 ps cannot be resolved, 1.4 ns and 5.4 ns components were observed in $6W_d + 4W_c$ and their total spectrum agrees with $4W_c$ emission (Fig. 4h). The 1.4 ns component is consistent with the same time scale of $4W_c$ under ps time resolution, but the 5.4 ns component was not observed in WT 315 nm excitation data.

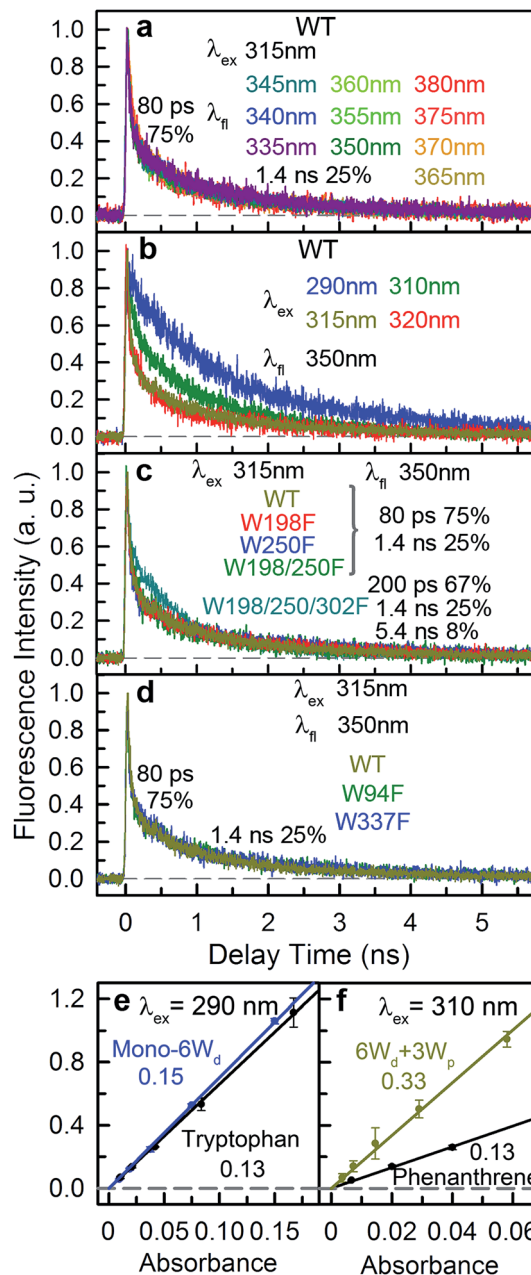


Fig. 5 Fluorescence lifetime of W_c and fluorescence quantum yields of W_d/W_p . (a) Fluorescence decay transients of $4W_c$ measured with WT ($\lambda_{ex} = 315$ nm). (b) Decay transients at 350 nm with various excitation wavelengths ($\lambda_{ex} = 290$ nm, 310 nm, 315 nm and 320 nm) for WT. (c) Fluorescence dynamics of the pyramid center measured for WT, W198F, W250F, W198/250F and W198/250/302F ($6W_d + 4W_c$). All transients are similar except that for $6W_d + 4W_c$ with an extra 5.4 ns slow decay constant. (d) Fluorescence dynamics of W_c measured by 315 nm excitation for WT, W94F and W337F. (e and f) Fluorescence quantum yields (Q_F) of (e) W_d and (f) W_p , measured using the tryptophan and phenanthrene standards, respectively.

From mutant studies in Fig. 5c, the 5.4 ns component in $6W_d + 4W_c$ was introduced by mutation of all $3W_p$, which probably causes local structural changes of $4W_c$, resulting in less efficient quenching. But the $4W_c$ absorption red tail (Fig. 1d) was not suppressed in $6W_d + 4W_c$ by this structural perturbation and



RET from $6W_d$ to $4W_c$ also seems unaffected. Interestingly, mutation of W198, W250 or both does not bring the 5.4 ns fluorescence dynamics to $4W_c$, suggesting important structural roles of W302, but not W198 or W250. In WT, although we observed similar time scales of 1.4 ns and 5.5 ns (Fig. 4i), their total spectrum is bluer than the $4W_c$ emission, clearly indicating that the latter is from W_p . The 5.5 ns is close to the 7.0 ns lifetime measured in $6W_d + 3W_p$ (Fig. 4g), suggesting W_p -to- W_c RET also has slow components. Energy-transfer rates depend on orientation factors (κ^2) and donor-acceptor distances (R), and both are sensitive to protein structural fluctuations.^{37,38} The flexible loop regions at the dimer interface can lead to broad rate distributions of W_p -to- W_c energy transfer. We further

investigated the reason for $4W_c$ double exponential decay by mutating two noncritical W_c : W94 and W337. Both mutants of W94F and W337F show similar decay dynamics to WT (Fig. 5d), indicating that the 1.4 ns component is not from any individual W_c and may result from structural fluctuations of $4W_c$, *i.e.*, the cluster can sample conformations with slower fluorescence quenching.

The time resolved experiments determined the overall RET rates of $6W_d$ but cannot provide energy-transfer time scales for individual donor-acceptor pairs or detailed RET rate distributions from structural fluctuations. To reveal this information, we next applied QM/MM computations and FRET theory to obtain the RET rates for all 84 possible donor acceptor pairs:

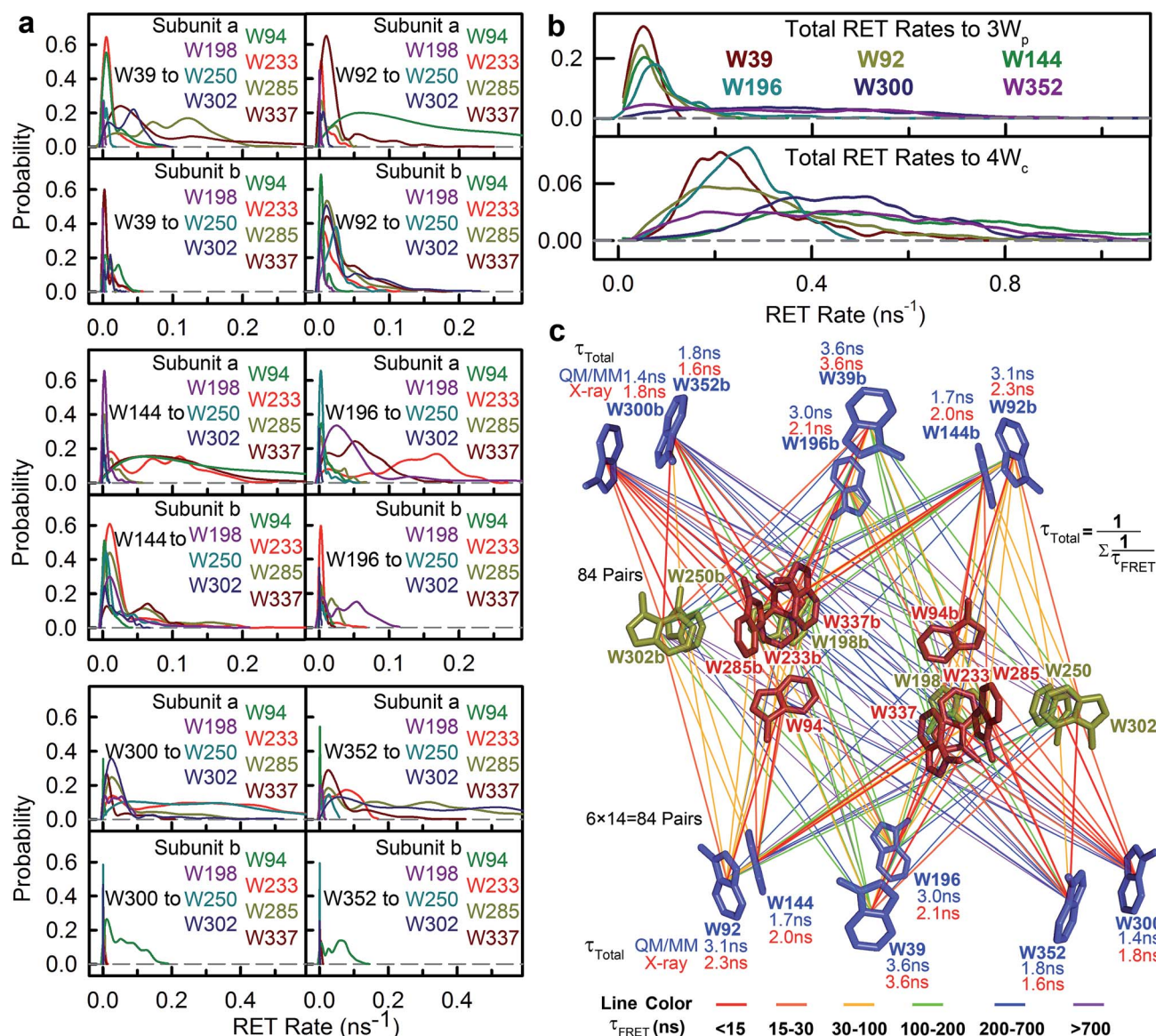


Fig. 6 RET from W_d to W_p/W_c . (a) RET rate distributions for all 84 donor-acceptor pairs from QM/MM calculations. (b) RET rate distributions for each W_d to total $3W_p$ (top) and to $4W_c$ (bottom) based on QM/MM methods. (c) The energy-transfer pathways from all W_d to 14 interfacial W residues ($3W_p + 4W_c$ on both subunits). The total RET time constants based on QM/MM (blue) and on X-ray structure (red) are shown near each W_d . Each line represents one energy-transfer pathway. Colors of the lines are based on effective RET time constants. The dominant paths are shown in red.



$6W_d$ to 14 interfacial W_p/W_c on both subunits. To model both fast and slow protein fluctuations, we conducted a 2 μ s MD simulation of UVR8 and evenly took 500 structures from the trajectory (4 ns interval) for RET rate calculations. In each structure, transition dipole moments were calculated with the TDDFT/MM level of theory for all tryptophan residues to obtain orientation factors (κ^2). With spectral overlap integrals (J in Fig. 1g), donor lifetimes (τ_D , Fig. 2), fluorescence quantum yields (Q_D , Fig. 5e and f), donor-acceptor distances (R) in MD structures and a refractive index of 1.33,³⁹ we obtained RET rate distributions for all 84 RET pathways (Fig. 6a). The energy-transfer rates span from 10^{-3} to 10^{-1} ns $^{-1}$ for different pairs, due to varying κ^2/R^6 values. For each donor-acceptor pair, we found that R values can fluctuate by 1–2 Å and κ^2 values vary significantly from 0 to 4 in different MD structures by protein fluctuations. Importantly, all W_d can transfer energy to interfacial Trp residues on the other subunit, leading to inter-subunit energy transfer. Among the $6W_d$, W39, W196, W300 and W352, prefer energy transfer to Trp on the same monomer, whereas W92 and W144 show significant RET to W_p/W_c on both monomers. It is also worth noting that $6W_d$ prefer direct energy transfer to W_c , especially to W285 and W233. As shown in Fig. 6b, the total RET rates to $4W_c$ are much faster than to $3W_p$, due to more favorable transition dipole moment coupling and greater spectral overlap between W_d and W_c . With the RET rate distributions from 500 MD structures and the 2.7 ns W_d lifetime, we simulated fluorescence decay transients for each W_d in the presence of each individual acceptor (84 transients shown in Fig. S2†) by averaging 500 single exponential decays curves. Since the decay time scales are close in various protein

conformations, all 84 simulated transients can be fitted with single exponential decays. Using the difference between these fitting values and the 2.7 ns original decay time, we calculated the effective RET time for each donor-acceptor energy-transfer pathway. We thus simplified broad rate distributions to single effective rates (Table S11†) that give the same W_d excited state decay dynamics and the same RET efficiency. The resultant effective RET times were schematically presented in Fig. 6c, revealing a complicated light-harvesting network. By adding up the rates of 14 parallel RET for each W_d (eqn (8)), total RET time constants (τ_{total}) were obtained as 3.6 ns, 3.1 ns, 1.7 ns, 3.0 ns, 1.4 ns and 1.8 ns for W39, W92, W144, W196, W300 and W352, respectively. We also calculated the 84 RET rates (Table S11† values in brackets) and τ_{total} with the static X-ray structure (Fig. 6c) for comparison with values from 500 MD snapshots. The two methods provide very different RET time scales for individual pathways as κ^2/R^6 is sensitive to tryptophan distances and orientations. But the resultant τ_{total} values are close because a sum of 14 rates reduces fluctuations, *i.e.*, one conformation that favors one RET pathway may impede another parallel channel, leaving the total rate nearly constant. Although the theoretical studies with fluctuations give very similar results as we obtained, such a picture may not explain an interface energy transfer from W_p to W_c as discussed in the following section of the paper.

To further compare the theoretical scheme with experimental data, we conducted numerical simulations of W_d fluorescence decay dynamics. The decay curves of the original 2.7 ns components were simulated for mutants and WT (Fig. 7 dashed lines) using the effective RET times from the theory (eqn (9) and

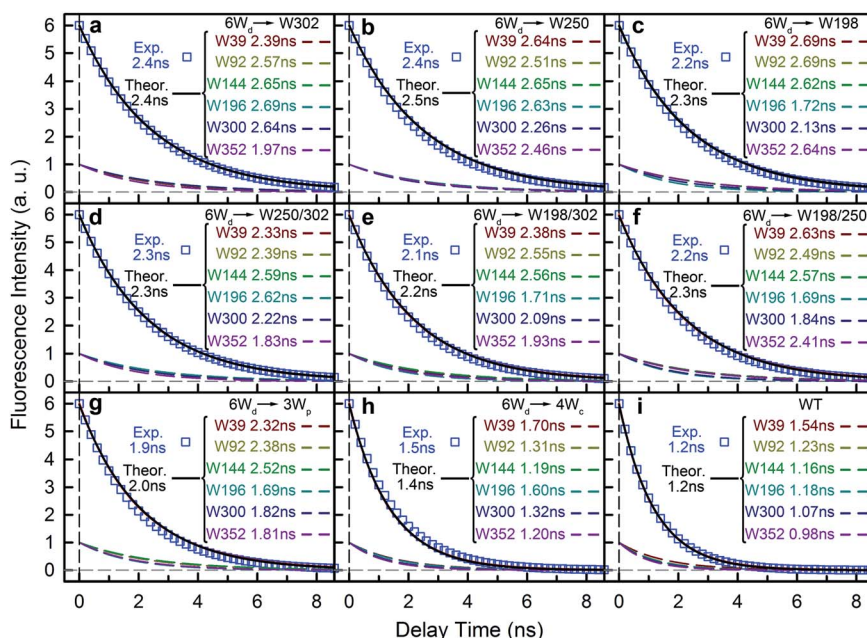


Fig. 7 Model simulations demonstrate that the total fluorescence decay dynamics of the $6W_d$ in UVR8 can be described with single exponential decays in various UVR8 protein mutants. (a) $4W_c/198/250F$ ($6W_d + W302$). (b) $4W_c/198/302F$ ($6W_d + W250$). (c) $4W_c/250/302F$ ($6W_d + W198$). (d) $4W_c/198F$ ($6W_d + W250/302$). (e) $4W_c/250F$ ($6W_d + W198/302$). (f) $4W_c/302F$ ($6W_d + W198/250$). (g) $6W_d + 3W_p$. (h) $6W_d + 4W_c$. (i) WT. The simulated total decay dynamics of the $6W_d$ (black solid lines, sum of the 6 dashed lines) can be fitted with a single decay time constant, which agrees well with experimental time scales.



(10)). In the presence of acceptors, the $6W_d$ donors have distinct RET rates due to asymmetrical locations of W_p/W_c , giving different decay curves (Fig. 7 dashed lines). However, by summing the 6 different single exponential decay curves, the total fluorescence decay of $6W_d$ can be perfectly described with a single time that agrees extremely well with experimental $6W_d$ lifetimes measured in Fig. 4, showing consistency between theory and experiments. The experimentally determined RET times mentioned above thus represent averaged RET rates of $6W_d$. Using the theoretical scheme in Fig. 6c, the overall energy-transfer efficiency for $6W_d$ was calculated to be 44% (Table S12†), in excellent agreement with the $44 \pm 0.6\%$ experimental value. These findings demonstrate that excitation energy flow from W_d to W_p/W_c follows the RET network shown in Fig. 6c, with tens of RET pathways from the buried W_d to the dimer interface.

Resonance energy transfer from the peripheral W_p outlier to the central W_c pyramid

To study W_p energy transfer, we applied the ps-TCSPEC method to measure the fluorescence lifetimes of the donors alone without RET using seven $6W_d + W_p$ mutants at 310 nm

excitation (shown in Fig. 8), and all transients exhibit typical double exponential decay of tryptophan fluorescence. With three $6W_d + 1W_p$ mutants (Fig. 8a–c), we determined fluorescence lifetimes of each individual W_p : W302 (1.1 ns, 13% and 8.2 ns, 87%); W250 (1.2 ns, 15% and 6.2 ns, 85%); W198 (1.2 ns, 18% and 6.6 ns, 82%). The second lifetimes in several nanoseconds are dominant and they completely agree with the time scales observed in sub-ns resolved TCSPC data (Fig. 4). The 1.1 ns and 1.2 ns fast components were not noticed in sub-ns data because of their minor percentages (<4%) in the overall signal with W_d , causing them to be mixed with other time scales. Fig. 8d–g show the fluorescence decay dynamics of W_p combinations measured with $6W_d + 2W_p$ and $6W_d + 3W_p$ mutants. Using the fitting parameters, we simulated the $2W_p$ and $3W_p$ fluorescence decay curves (Fig. 8h and k symbol), and they match the sum of individual W_p lifetime decay dynamics measured in $6W_d + 1W_p$ mutants (Fig. 8h and k), suggesting no significant RET among the $3W_p$.

With W_p lifetimes (Fig. 8a–c), the fluorescence quantum yield (Fig. 5f) and other values, we calculated RET rate distributions from W_p to W_c with 500 MD structures, following the procedure described in the previous section.

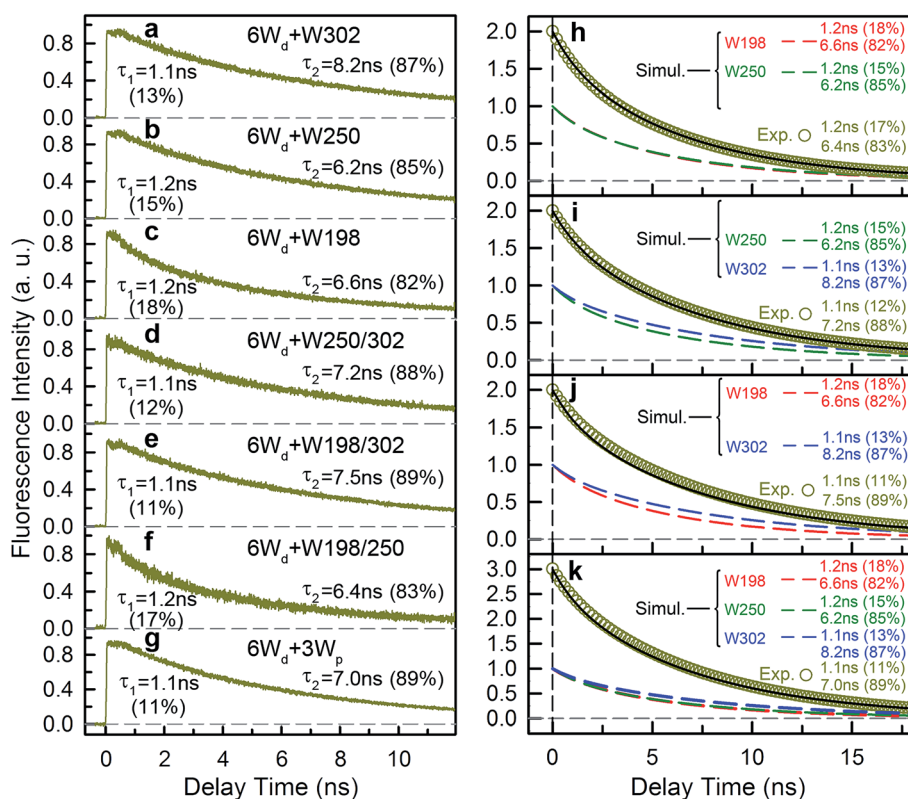


Fig. 8 W_p fluorescence lifetimes. (a–g) Fluorescence decay transients ($\lambda_{ex} = 310$ nm) at 340 nm fluorescence wavelength for 7 mutants: (a) $4W_c/198/250F$ ($6W_d + W302$). (b) $4W_c/198/302F$ ($6W_d + W250$). (c) $4W_c/250/302F$ ($6W_d + W198$). (d) $4W_c/198F$ ($6W_d + W250/302$). (e) $4W_c/250F$ ($6W_d + W198/302$). (f) $4W_c/302F$ ($6W_d + W198/250$). (g) $6W_d + 3W_p$. (h–k) Numerical simulations show that the W_p lifetimes observed in $6W_d + 2W_p$ mutants and $6W_d + 3W_p$ mutants match the sum of decay transients of $6W_d + 1W_p$. The mutants: (h) $6W_d + W198/250$; (i) $6W_d + W250/302$; (j) $6W_d + W198/302$; (k) $6W_d + 3W_p$. For each mutant, decay transients of individual W_p (colored dashed lines, red–W198; green–W250; blue–W302) were simulated using a double exponential decay model experimentally measured parameters from $6W_d + 1W_p$ mutants, which add up a total decay transient (black solid line). The total transients agree with the simulated curves (dark yellow circles) using time constants and ratios directly measured in $6W_d + 2W_p$ and $6W_d + 3W_p$ mutants.

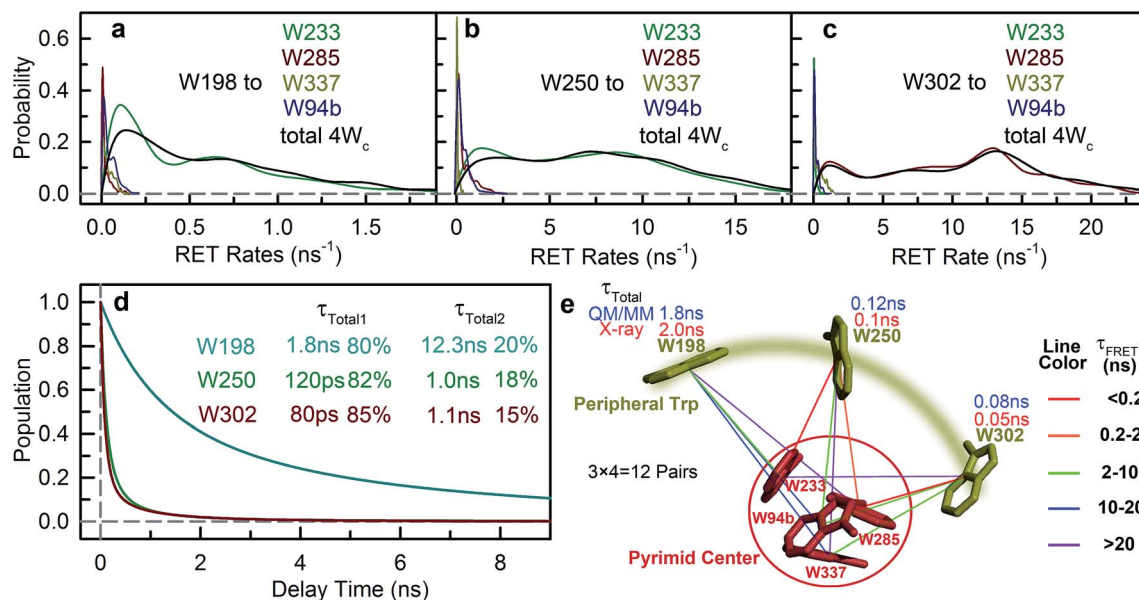


Fig. 9 RET rate distributions based on QM/MM calculations for each W_p–W_c pair. (a) W198 as the donor. (b) W250 as the donor. (c) W302 as the donor. In each figure, the distribution of the total RET rate to 4W_c is shown in black. (d) Simulation of energy transfer dynamics for 3W_p to 4W_c based on RET rate distributions. For all 3W_p, a fast energy transfer (τ_{Total1}) and a slow energy transfer (τ_{Total2}) are needed to fit the transients. (e) The 12 possible energy-transfer pathways from 3W_p to 4W_c as calculated with the FRET theory. Each line represents one energy-transfer pathway. Colors of the lines are based on calculated RET time constants. The total RET time constants are shown near each W_p.

Since W_p RET to the distant pyrimid center is negligible, we only show the 12 rate distributions for 3W_p to 4W_c in the nearby cluster (Fig. 9a–c). Due to shorter donor–acceptor distances, RET time scales here fall in the range of tens of picoseconds to nanoseconds, significantly faster than W_d RET. We found that W198, W250 and W302 mainly transfer energy to W233, W233 and W285, respectively, with the rate distributions close to their total RET rates to 4W_c. The RET pathways are consistent with their respective distances and orientation factors. Like W_d, RET dynamics of W_p were obtained by adding 500 different single exponential decay curves based on calculated RET times in 500 MD snapshots (Fig. 9d). For W_d–to–W_p/W_c channels, all 500 RET rates are small, and thus all 500 decay times are in a narrow range between 1.8 ns and 2.7 ns, making the averaged decay dynamics close to single exponential decays. However, for W_p–to–4W_c, the time scales range from tens of picoseconds to nanoseconds, unable to be described with a single decay time. For all 3W_p, the RET dynamics were effectively fitted with two timescales: a dominant fast component (τ_{Total1}) and a slow one (τ_{Total2}) with smaller amplitudes (Fig. 9d). Compared with W250 and W302, which have a τ_{Total1} of 120 ps and 80 ps, respectively, W198 has a dramatically slower τ_{Total1} of 1.8 ns, owing to unfavorable orientations and longer distances. τ_{Total2} are on nanosecond time scales, representing a small fraction of W_p with slow RET to the pyrimid center. Because both W_p to W_c are located in the flexible loop regions, UVR8 protein may sample conformations, resulting in distributions of RET. When using the X-ray structure for the calculations, single RET times of 2.0 ns, 100 ps and 50 ps

were obtained for W198, W250 and W302, respectively, which are in line with the τ_{Total1} fast RET times (Fig. 9e).

We then experimentally investigated the energy-transfer dynamics from W_p to W_c in various mutants, with ps-resolved TCSPC using 310 nm excitation wavelength to avoid the W_d signal, and then compared the fluorescence decays with theoretical RET model simulations. Since mutation of W302 brings an additional slow decay component to 4W_c that complicates data analysis, W302 was kept in the mutants. Transients measured at various fluorescence wavelengths from 330 nm to 380 nm are shown in Fig. S3–S6† with 340 nm data selected in Fig. 10a–d. Using calculated energy-transfer rate distributions, W_p lifetimes and absorbance values of W_p/W_c, we simulated these 310 nm pump transients, which agree well with experimental data for all different fluorescence wavelengths in all 4 mutants (Fig. 10 and S3–S6†). The total dynamics (green curve) consist of contributions from W_p (dark yellow) and 4W_c (red). Since W_p shows two lifetimes and two RET times, the W_p fluorescence signal is composed of four decay components corresponding to four subpopulations: short lifetime with fast RET (11–14%); long lifetime with fast RET (66–74%, most dominant); short lifetime with slow RET (2–3%, negligible); long lifetime with slow RET (13–16%). For W250/W302, the first two components are close to their τ_{Total1} in 120/80 ps since they are much faster than lifetimes; the third one is negligible; the last subpopulation decays in about 0.9 ns calculated from the total rate of 6.2/8.2 ns lifetime decay and the 1.0/1.1 ns τ_{Total2} . W198 shows decay components in 0.7 ns (14%), 1.4 ns (66%), 1.1 ns (3%) and 4.3 ns (17%). Signals of 4W_c contain the decays (80 ps and 1.4 ns) of directly excited 4W_c and



additional rise/decay components due to RET from W_p . The latter represent RET from the four W_p subpopulations to the two W_c components with complicated rise/decay dynamics, with the most dominant ones from the fast RET to the 80 ps and 1.4 ns $4W_c$ pools. Because of the slower formation (80–120 ps τ_{total1} in W250/W302) and faster decay (for the 80 ps $4W_c$ component), RET leads to a reverse kinetics of an 80 ps rise with an 80–120 ps decay. On the other hand, fast RET results in substantial accumulation of the 1.4 ns component in 80–120 ps, followed by 1.4 ns decay. The slow RET of W198 only brings negligible accumulation of the fast 80 ps component. By adding all these dynamics, RET causes a rise in 80–120 ps and additional decays in 1.4 ns, making the overall $4W_c$ fluorescence dynamics apparently slower than $4W_c$ alone. Fluorescence signals cannot be simulated well using single RET rates calculated for the X-ray structure, especially at long delay times, due to missing of the slow RET of τ_{total2} . These results suggest that energy-transfer rate distributions from W_p to W_c shown in Fig. 9 are accurate. We obtained RET efficiency of 63%, 94% and 96% for W198, W250 and W302 to $4W_c$, respectively, with an average value of 84%.

Using all the RET rates, including those from W_d , we simulated the fluorescence decay dynamics in WT at 290 nm excitation with all three Trp groups (Fig. 10e and S7†). $6W_d$ approximately follows a single-exponential decay in 1.2 ns as

discussed above. Besides the four-component decays of the initially excited W_p , due to W_d -to- W_p RET, W_p contains additional dynamics of formation and decay. The $4W_c$ signal is highly complicated, with directly excited $4W_c$, RET from W_p , direct RET from W_d and two-step transfer from W_d via W_p . Experimental data are consistent with simulation lines, validating our RET mechanism. We then integrated decay curves of different Trp groups and thus decomposed steady-state emission spectra to contributions of individual groups (Fig. 10f–j symbols). These spectra agree well with directly measured fluorescence spectra (Fig. 10f–j solid curves) in WT and all mutants, further confirming our model simulations.

Using all the RET time scales, we finally obtained the overall light sensing efficiency of UVR8. From spectra in Fig. 1e, the absorbance of W_d , W_p and W_c is 0.67, 0.85 and 0.89, respectively, at 290 nm. Using these values, for every 100 UV-B photons absorbed, 40 go to $6W_d$, 25 reach $3W_p$ and 35 excite $4W_c$ (Fig. 11). $6W_d$ transfers 31% and 13% of excitation energy to $4W_c$ and $3W_p$, respectively, donating 12.4 (40×0.31) photons to W_c and 5.2 (40×0.13) photons to W_p . $3W_p$ shows 84% RET efficiency, funneling additional 25.4 photons ($(25 + 5.2) \times 0.84$) to the cluster. With this delicate light-harvesting network, $4W_c$ finally perceives 73% ($0.35 + 0.124 + 0.254$) of the total number photons captured by the protein, more than double the 35% initial direct excitation.

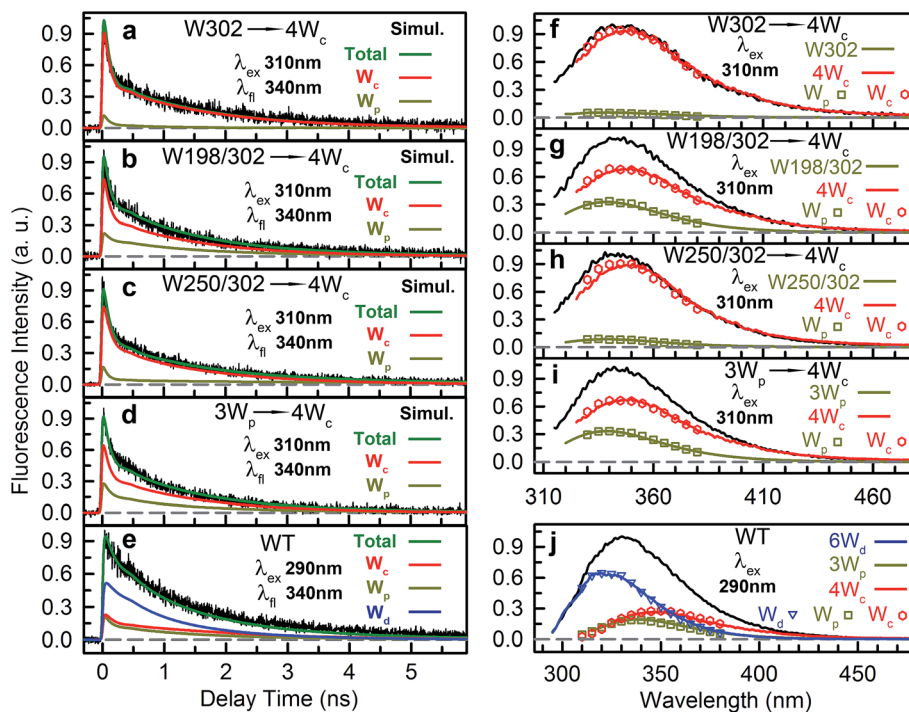


Fig. 10 Model simulations of fluorescence transients. (a–e) Simulations of typical fluorescence transients at 340 nm for mutants $6W_d$ + W302 + $4W_c$ (a), $6W_d$ + W198/302 + $4W_c$ (b), $6W_d$ + W250/302 + $4W_c$ (c), $6W_d$ + $3W_p$ + $4W_c$ (d) and WT (e). In each figure, the black solid line is the experimental data and the green solid line is the total simulation curve, which is the sum of contributions of W groups. (f–j) Comparison between directly measured and simulation-constructed spectra for $6W_d$ + W302 + $4W_c$ (f), $6W_d$ + W198/302 + $4W_c$ (g), $6W_d$ + W250/302 + $4W_c$ (h), $6W_d$ + $3W_p$ + $4W_c$ (i) and WT (j). In each panel, the black curve is the total emission. The colored symbols represent the spectra of three W groups decomposed from the total emission based on the time integrals of simulation curves. Solid color curves are the directly measured emission spectra of the three W groups.



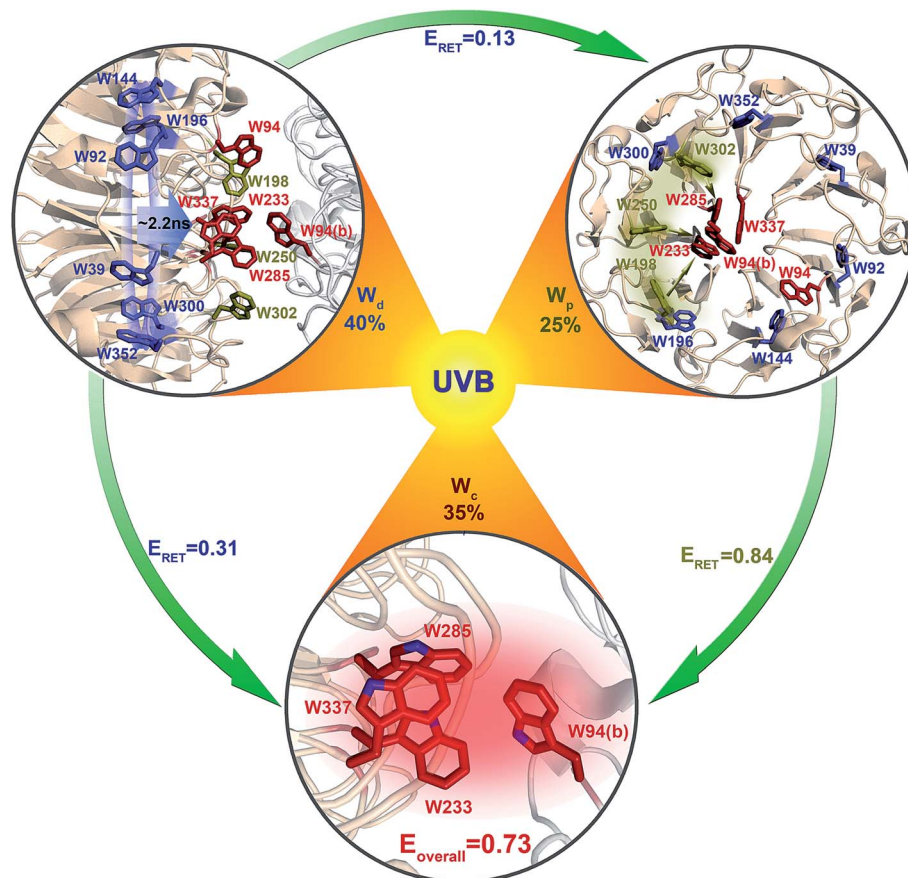


Fig. 11 A scheme of excitation energy-transfer networks in UVR8. Arrows show excitation-energy flow from the distal ring to the interfacial peripheral outliers and pyramid centers, and from the peripheral outlier to the pyramid center. The corresponding values are the branching fractions for their energy-transfer pathways.

Conclusion

Based on our previous brief report and summary,^{28,40} in this work, we described our detailed studies of the mechanism of light sensing by the UV-B photoreceptor *Arabidopsis thaliana* UVR8 and reported a striking light-harvesting network that involves all 26 structural tryptophan chromophores in the UVR8 dimer. With site-directed mutagenesis, we found that these Trp residues may be partitioned into three distinctive groups that exhibit significant difference in absorption spectra longer than 300 nm. The distal group of W_d of 6 Trp residues in the interior of UVR8 shows negligible absorbance beyond 310 nm; the absorption spectrum of the three peripheral residues W_p extends to 314 nm; and the absorption red-side tail of a pyramid cluster of four Trp residues, W_c , reaches beyond 320 nm. These differences allow selective excitation of W_d , W_p or W_c , revealing distinct emission spectra with peaks at 320 nm, 340 nm and 350 nm, respectively. This hierarchical arrangement, modulated by local environments, clearly separates the three groups into a ladder of energy landscapes, and allows RET from the highest energy, distal W_d to interfacial W_p/W_c and from W_p to the central, the lowest energy W_c cluster with negligible back energy transfer. The net result is to funnel all excitation energy to the pyramid center ($4W_c$).

For $6W_d$ RET to interfacial W_p/W_c , each W_d donor transfers energy to all 14 tryptophan residues on both subunits at the interface, resulting in a complicated network of 84 RET pathways. As investigated with 3 different time-resolved fluorescence methods and QM/MM calculations, these energy-transfer processes are on ns time scales and exhibit distributions of rates, instead of a single rate constant, due to protein structural fluctuations. $6W_d$ have an overall RET efficiency of 44%, with 31% to $4W_c$ and 13% to $3W_p$.

From time-resolved fluorescence and model simulations, the energy transfer from $3W_p$ to $4W_c$ also shows broad rate distributions, on tens of ps to ns time scales. Each W_p effectively contains a major population with dominant fast RET and a small fraction (10–15%) with slow RET, owing to structural flexibility in interfacial loop regions. W198, W250 and W302 have RET efficiency of 63%, 94% and 96%, respectively, with an overall efficiency of 85% for $3W_p$.

All the experimental energy-transfer rates perfectly agree with the QM/MM theoretical predictions, indicating that the light harvesting follows the classical excitation energy-transfer mechanism without any coherent energy-transfer character in such long transfer distances. Using the 26 structural tryptophans with a beautiful architecture, the delicate light harvesting network makes the tryptophan pyramid “reaction center”



achieve a quantum leap in efficiency of light sensing from 35% by direct exciton absorption to 73% through light harvesting. UVR8 is the only known photoreceptor that has a light-harvesting network, uniquely without the aid of external chromophores. Since UV-B only accounts for a minor portion of solar irradiation on the surface of the earth⁴¹ and the extinction coefficient of tryptophan is relatively low, the light harvesting in UVR8 is critical to the sensitive detection of lower intensity in UV-B light for initial signaling to biological functions.

Experiments and methods

Protein sample preparation

The purification of *Arabidopsis* UVR8 has been reported before.¹² Plasmids for mutant expression were constructed using MiniPrep kits (Qiagen) following the manufacturer's protocol. For all experiments, protein samples were kept in the following buffer condition: 150 mM NaCl, 25 mM Tris and pH = 8.0. The dimer/monomer state was determined with a 25 mL Superdex 200 size column (GE) and an AKTA FPLC system (GE). In all proteins, the distal unstructured W400 was mutated to phenylamine and we defined W400F as "the wild type (WT)" throughout this paper.

Steady-state absorption and emission spectra

All absorption spectra were measured using a Cary 50 UV-Vis spectrophotometer (Agilent). The OD at the absorption peak was adjusted to around 1.0 in 5 mm cuvettes for every sample. Absorption spectra were re-normalized to tryptophan numbers and then subtraction was conducted to obtain clean spectra of 3 tryptophan groups. The emission spectra were measured using a Fluoromax-3 spectrofluorometer (Horiba) at 2 μ M (290 nm excitation) or 10 μ M (310 nm and 315 nm excitation) protein concentrations. For fluorescence quantum yield (QY), fluorescence spectra of proteins at a series of concentrations were measured and integrated, and the integrals were then plotted *versus* absorbance at 290 nm or 310 nm excitation wavelengths. Tryptophan and phenanthrene standards were dissolved in water and absolute ethanol, respectively.^{42,43} For phenanthrene, the solution was purged with argon gas until no further increase in fluorescence was observed.⁴³ For each sample, 3 independent measurements were conducted, and the error bars were determined using the standard deviation of the 3 measurements.

Femtosecond-resolved fluorescence up-conversion method

The experimental layout has been detailed elsewhere.⁴⁴ Briefly, the 290 nm excitation beam was generated by OPA-800 (1 kHz, Spectra-Physics), and its pulse energy was attenuated to \sim 100 nJ. The instrument response time under the current noncollinear geometry is about 400 fs, and all data were taken at a magic angle (54.7°). The samples (150 μ M) were kept in spinning quartz cells during irradiation to avoid heating and photobleaching. Although this method provides femtosecond (fs) time resolution, the maximum window of measurement is limited by optical delay to 3 ns and a large amount (6–10 mg) of protein sample is required.

Picosecond-resolved time correlated single photon counting (TCSPC)

Fluorescence data were acquired using a commercially available FluoTime 200 system (Picoquant). For the excitation light, a tunable Tsunami laser (80 MHz, Spectra-Physics) was used to generate 100 fs laser pulses at 945 nm, 930 nm and 870 nm wavelengths, followed by third harmonic generation using a tripler (Miniopic) to produce 315 nm, 310 nm and 290 nm excitation pulses (100 fs, 80 MHz, 0.5 μ W). The full width at half maximum (FWHM) of the instrument response function (IRF) was 30 ps. All samples (10 μ M) were kept in 5 mm quartz cuvettes during measurements. No dissociation was observed after the experiments, as confirmed using the size column. This TCSPC method provides 15 ps time resolution with an exceptional signal-to-noise ratio and negligible sample damage by the weak excitation light. Time window is limited to 12 ns due to the 80 MHz repetition rate of the Tsunami laser. This method only needs a small amount (0.3 mg) of protein samples. The picosecond resolution is enough to investigate the RET dynamics among the three Trp groups. There might be ultrafast processes in UVR8, including the ultrafast energy delocalization in the pyramid center, and such dynamics is beyond the light harvesting reported here.

Sub-nanosecond-resolved time correlated single photon counting (TCSPC)

Fluorescence decay transients were acquired using a commercially available FluoTime 200 system (Picoquant) with PLS-290 pulsed LED (8 MHz, 290 nm, 1 μ W, Picoquant) as the excitation source. The FWHM of the IRF was 600–700 ps. All samples (10 μ M) were kept in 5 mm quartz cuvettes for measurements. For WT and 6W_d + 4W_c samples, little dissociation was observed after the experiments, as confirmed using a size column. This method provides a large time window up to hundreds of ns, covering the whole fluorescence decay process. But the time resolution is limited to 300 ps and excitation wavelength cannot be adjusted.

Data fitting and construction of lifetime associated spectra

All TCSPC data were fitted with convoluted multiple-exponential decay models using FluoFit software (Picoquant) and thus the fitting parameters (exponential decay times and their amplitudes) were obtained. Since the steady-state fluorescence spectrum is simply time integrals of exponential decays (eqn (1)), the spectrum of the m^{th} exponential component was given by eqn (2):

$$I(\lambda) = \int_0^{+\infty} \sum_i A_i e^{-\frac{t}{\tau_i}} dt = \sum_i A_i \tau_i \quad (1)$$

$$I_m(\lambda) = I(\lambda) \frac{A_m \tau_m}{\sum_i A_i \tau_i} \quad (2)$$

A_i and τ_i are the amplitude and time constant of the i^{th} exponential component from the transient probed at wavelength λ . $I(\lambda)$ is the intensity of the steady-state emission.



MD simulations

The X-ray structure of the UVR8 wild type (PDB entry: 4D9S) was solvated using TIP3P water in a cubic box of $104 \times 104 \times 104 \text{ \AA}^3$. Counter ions were added to neutralize charge and produce an ion concentration of approximately 0.15 M. The simulation was set up using CHARMM c38a2,⁴⁵ using the CHARMM27 force field⁴⁶ with CMAP correction.⁴⁷ After energy minimization, the system was gradually heated from 10 K to 300 K every 10 K using 100 ps NPT simulations at each temperature. The simulation was first carried out using CHARMM during initial 20 ns equilibration. Then a long-time simulation up to 2 μs was performed using GROMACS-4.6.5 molecular dynamics code⁴⁸ at 300 K and 1 atm pressure with a 2 fs time step. The non-pair list was updated every 10 steps. The grid neighbor searching method was applied in the simulation with a 10 \AA cutoff distance for the short-range neighbor list. Electrostatic interactions were treated by using the Partial-Mesh Ewald (PME) summation method⁴⁹ with a 14 \AA for long range and 10 \AA for short-range electrostatic cutoff, respectively. The short-range cutoff for van der Waals interactions during the simulation was 12 \AA . The isotropic pressure coupling was achieved by the Parrinello–Rahman method⁵⁰ with a compressibility of $4.5 \times 10^{-5} \text{ bar}^{-1}$.

QM/MM calculations

We extracted 500 snapshots (one snapshot in every 4 ns) from the 2 μs simulation for further QM/MM calculations using multistate density functional theory (MSDFT), which has been implemented in a locally modified version of GAMESS⁵¹ and CHARMM hybrid. In MSDF calculations, the electron singlet excited state can be written as⁵²

$$\Phi(S_1) = \underbrace{\Psi_1 \cdots \hat{A} \{ \Psi_a^{S_1} \} \cdots \Psi_N}_{N-1 \text{ fragments}} \quad (3)$$

where $\hat{A} \{ \Psi_a^{S_1} \}$ specifies an antisymmetric wave function for the locally excited residue *a*, which is electrostatically embedded in the rest of the system through a Hartree potential. In the present study, the excited state is represented by using time-dependent density functional theory (TDDFT). The remaining $N - 1$ fragments (amino acid residues, ions and water molecules) are approximated by using the CHARMM force field and TIP3P model for water. For every snapshot, the transition dipole moment was calculated for all 13 tryptophan chromophores using the time-dependent range-separated hybrid functional TD-CAM-B3LYP along with the 6-31+G(d) basis set. A total number of 6500 (13 Trp \times 500 snapshots) of QM/MM calculations were conducted.

Energy-transfer rate calculations with Förster resonance energy transfer (FRET) theory

According to Förster theory,^{37,38} the energy-transfer rate k_{RET} (in ns^{-1}) is given below:

$$k_{\text{RET}} = \frac{1}{\tau_{\text{D}}} \left(\frac{R_0}{R} \right)^6 = 8.79 \times 10^{23} \frac{\kappa^2 Q_{\text{D}} J}{R^6 n^4 \tau_{\text{D}}} \quad (4)$$

where R (in \AA) is the center-to-center distance between the donor and acceptor in the 500 MD snapshots. κ^2 is the orientation factor defined by

$$\kappa^2 = (\boldsymbol{\mu}_{\text{A}} \cdot \boldsymbol{\mu}_{\text{D}} - 3(\boldsymbol{\mu}_{\text{A}} \cdot \mathbf{R})(\boldsymbol{\mu}_{\text{D}} \cdot \mathbf{R}))^2 \quad (5)$$

where $\boldsymbol{\mu}_{\text{A}}$ and $\boldsymbol{\mu}_{\text{D}}$ are the normalized transition dipole moments of the acceptor and the donor, respectively, determined from QM/MM calculations. \mathbf{R} is the normalized vector connecting centers of donor and acceptor tryptophan residues in each structure. n is the refractive index within protein and was set to 1.33 based on the literature.³⁹ Q_{D} is the fluorescence quantum yield of the donor without acceptors as determined by experiments. The overlap integral J (in $\text{cm}^3 \text{ M}^{-1}$) was obtained using the following equation:

$$J = \frac{\int_0^\infty F_{\text{D}}(\lambda) \varepsilon_{\text{A}}(\lambda) \lambda^4 d\lambda}{\int_0^\infty F_{\text{D}}(\lambda) d\lambda} \quad (6)$$

where F_{D} is the emission spectrum of the donor (W_{d} or W_{p}). ε_{A} is the extinction coefficient of the acceptor (in $\text{M}^{-1} \text{ cm}^{-1}$), which is obtained by normalizing the peak extinction coefficient to $5600 \text{ M}^{-1} \text{ cm}^{-1}$.⁵³ The unit of wavelength λ was converted to centimeter to give J values in $\text{cm}^3 \text{ M}^{-1}$. τ_{D} in eqn (4) is the donor lifetime in nanoseconds. Since tryptophan has two lifetimes, the amplitude weighted average lifetime was used here:

$$\tau_{\text{D}} = \frac{A_1 \tau_1 + A_2 \tau_2}{A_1 + A_2} \quad (7)$$

In eqn (7), τ_1 and τ_2 are the two lifetimes. A_1 and A_2 are the corresponding amplitudes, respectively.

Finally, by plugging in all values to eqn (4), we obtained RET rates k_{RET} (in ns^{-1}) for all donor–acceptor pairs. Rate distributions were generated using calculated κ^2 from the 500 MD snapshots. The corresponding RET time scale τ_{RET} (in ns) is simply the reciprocal of k_{RET} .

Total RET rates and simulation of decay dynamics of the original 2.7 ns component

Every W_{d} transfers energy to all interfacial tryptophan residues on both UVR8 subunits. The total rate of parallel reactions is:

$$k_{\text{total},m} = \sum_{i=1}^N k_{\text{RET},mi} = 1/\tau_{\text{total},m} \quad (8)$$

$k_{\text{RET},mi}$ is the calculated effective RET rates from the m^{th} W_{d} to the i^{th} $W_{\text{p}}/W_{\text{c}}$. N is the number of interfacial tryptophan residues in the protein. The corresponding time constant is the reciprocal of k_{total} . Briefly, the RET rate was obtained for each W_{p} . The 2.7 ns lifetime of the m^{th} W_{d} in the presence of acceptors will decrease to $\tau_{\text{DA},m}$:

$$\tau_{\text{DA},m} = 1/(1/2.7 + k_{\text{total},m}) \quad (9)$$

By summing up the decay dynamics of all the 6 W_{d} , we obtained the total fluorescence dynamics:



$$I(t) = \sum_{m=1}^6 \exp(-t/\tau_{\text{DA},m}) \quad (10)$$

Numerical simulations of picosecond-resolved TCSPC transients and spectrum decomposition

The details are shown in the supplemental methods. Briefly, we applied a series of differential equations to simulate temporal evolution of excited state populations of various tryptophan residues and groups, using the energy-transfer model shown in this work. Signal contributions from three tryptophan groups were numerically simulated and added together to give the total simulation curve for comparison with ps-resolved TCSPC data. From time integrations of simulated time-resolved fluorescence curves of 3 tryptophan groups, the total steady-state emission spectra were decomposed to give emission spectra of 3 tryptophan groups.

RET efficiency calculations

RET efficiency can be defined with donor fluorescence lifetimes as follows.^{37,38}

$$E = 1 - \frac{\tau_{\text{DA}}}{\tau_{\text{D}}} = \frac{k_{\text{total}}}{\tau_{\text{D}}^{-1} + k_{\text{total}}} \quad (11)$$

τ_{DA} and τ_{D} are donor fluorescence lifetimes with and without the acceptors, respectively. τ_{D} was directly measured; τ_{DA} was calculated with τ_{D} and total energy transfer rates k_{total} (sum of k_{total} and the reciprocal of τ_{D}). For tryptophan chromophores, we treated two lifetimes as two different subpopulations and used the population weighted average efficiency:

$$E = R_1 E_1 + R_2 E_2 = (1 - R_2) E_1 + R_2 E_2 \quad (12)$$

R_1 and R_2 are the amplitude ratios of the two tryptophan lifetimes; E_1 and E_2 are the RET efficiencies of the two lifetime components as calculated using eqn (11). The overall light-harvesting efficiency of UVR8 was calculated based on RET efficiency of each W_{d} or W_{p} and absorbance values of the three Trp groups, as detailed in the ESI methods.

Author contributions

D. Z. designed the project; J. G. designed the computational studies; X. L., Z. L., M. K. and L. W. conducted the experiments; H. R. conducted MD simulations and QM/MM calculations. X. L., J. G. and D. Z. wrote the paper.

Conflicts of interest

The authors declare no conflict of interest.

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