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Regioselective photodimerization as a tool for light-regulated catalyst assembly

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While photoisomerization has dominated the design of photoswitchable catalysts, this work introduces an alternative approach: leveraging light-induced photodimerization to assemble catalytically active species. The adopted strategy is based on an acrylamidylpyrene derivative equipped with a TACN·Zn(II) catalytic unit. This system undergoes a visible-light-induced [2 + 2] cycloaddition, which is both regioselective and reversible, to form a catalytically active photodimer. While the *E*-to-*Z* photoisomerization of the monomer has no significant effect on catalysis, the photodimerization leads to a six-fold enhancement in catalytic activity. The photodimer's catalytic efficiency is attributed to the clustering of catalytic units, facilitating a more efficient transphosphorylation reaction. Notably, this system demonstrates the ability to temporally control catalytic reactivity, as the active dimer can be reverted to the monomers upon irradiation with UV light. This work highlights the potential of photodimerization as a robust alternative strategy for regulating catalytic activity and opens new avenues for light-responsive catalysis with temporal control.

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Introduction

Photoresponsive systems that couple catalytic function to light stimuli play a central role in fundamental biological processes, such as vision,^{1,2} light-directed growth and movement in plants.³ Additionally, plants leverage the conversion of light energy into chemical bonds to activate proteins *via* photodimerization,⁴ a process that is fundamental for regulating gene expression, circadian rhythms, and photomorphogenesis.^{5–7} Chemists have harnessed light to regulate catalytic activity, primarily exploiting photoisomerization processes to alter the geometry or electronic properties of catalyst architectures.⁸ Azobenzene scaffolds, for instance, have been widely explored to achieve photoswitchable catalysis by exploiting cooperative,^{9–17} electronic,^{18–21} or steric effects,^{22–25} or by inducing the self-assembly of molecules into supramolecular structures.^{26–28} Similarly, light-driven isomerization of molecular rotors^{29–31} and ring-closing/ring-opening reactions of photoactive dithienylethene^{32–35} and spiropyran³⁶ derivatives have been employed to design light-responsive catalysts.

In this work, we present an alternative strategy for photo-regulating catalytic reactivity. Instead of inducing conformational changes in a pre-existing catalyst, we harness light energy to construct an active catalyst *via* a photodimerization process. The [2 + 2] cycloaddition reaction—a widely studied photochemical process—forms cyclobutane adducts upon irradiation

across a broad wavelength range³⁷ and has been investigated in cinnamic acid,³⁸ thymine,³⁹ coumarin, stilbene,⁴⁰ and, more recently, acrydylpyrene derivatives. Acrydylpyrene offers unique advantages as one of the mildest and most orthogonal photochemical systems, with wavelengths for self-dimerization and cycloreversion at approximately 435 nm and 330 nm, respectively. These reversible photodimerization processes have been applied to polymer ligation,^{41–46} photodynamic hydrogels,^{47–49} and the photocontrol of duplex formation between artificial nucleic acids and RNA.⁵⁰

Here, we equipped an acrylamidylpyrene chromophore with a 1,4,7-triazacyclononane (TACN)·Zn(II) unit (Fig. 1), a moiety known to enhance catalytic activity in transphosphorylation reactions⁵¹ by promoting cooperative effects between neighbouring TACN·Zn(II) units.^{52–57} We demonstrated that irradiation at 420 nm induces both the *E*-to-*Z* isomerisation of the monomer, which did not affect the catalysis, and photodimerization into a catalytically active cyclobutane derivative. Prolonging the irradiation shifts the equilibrium towards the active dimer and the catalytic activity is reversibly regulated by alternating between 420 nm and 311 nm irradiation, which instead causes dissociation of the catalyst. Remarkably, the [2 + 2] cycloaddition is highly stereoselective, yielding a single cyclobutane product out of 11 theoretically possible regioisomers.

Results and discussion

After having synthesised the acrylamidylpyrene derivative *E*-1·Zn(II) (for details see SI, Section 2) we initially studied its

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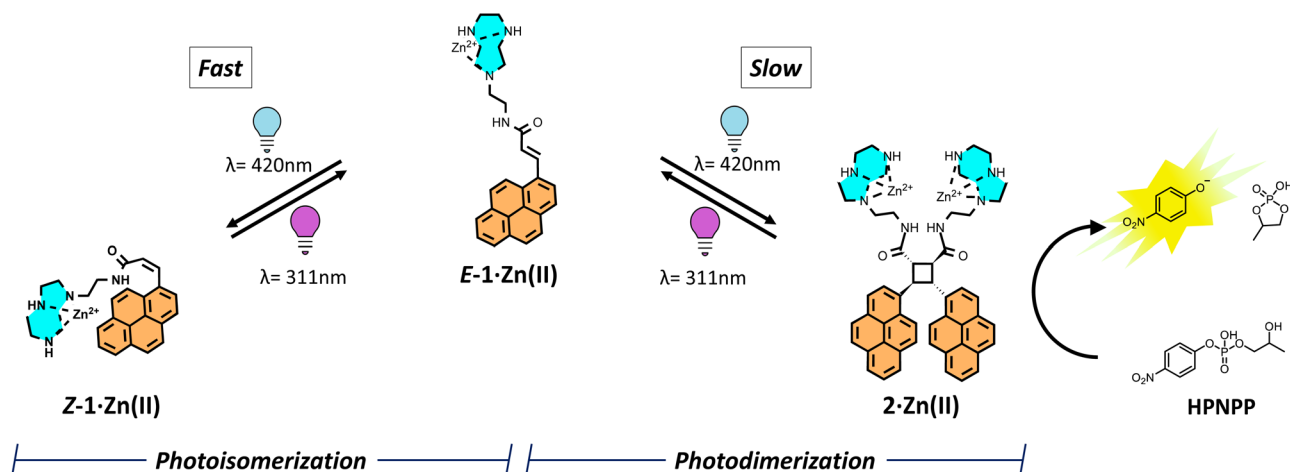


Fig. 1 Reaction scheme highlighting the studied processes: upon irradiation at 420 nm the $E-1 \cdot Zn(II)$ monomer can isomerize into $Z-1 \cdot Zn(II)$ (fast) or dimerize to give $2 \cdot Zn(II)$ (slow). The formed photodimer can catalyze the transphosphorylation of HPNPP into p -nitrophenol (PNP) and cyclic phosphate. The photoinduced processes can be reverted by irradiating with a wavelength of 311 nm.

photochemical behavior *via* $^1\text{H-NMR}$ spectroscopy, using $\text{D}_2\text{O} : \text{DMSO} 8 : 2$ as the solvent mixture, in order to prevent potential aggregation. The first process that we observed when irradiating a 0.5 mM solution of $E-1 \cdot Zn(II)$ at 420 nm for 3 min was the photoisomerization from the E to the Z isomer, reaching around 85% of the Z -isomer at the photostationary state (Fig. 2b, c and S28; for details on the choice of wavelengths see SI, Section 3.3). Upon irradiation at 311 nm for 2 hours reversible photoisomerization from Z into E isomer took place, and this process could be repeated for different cycles. However, it was observed that the total concentration of monomers was slowly decreasing

because of the formation of photodimer $2 \cdot Zn(II)$ (Fig. S29 and S30).

In fact, when a 0.5 mM solution of $E-1 \cdot Zn(II)$ in $\text{D}_2\text{O} : \text{DMSO} 8 : 2$ was irradiated for about 120 min at 420 nm, we observed that the $[2 + 2]$ cycloaddition prevailed over the photoisomerisation process (92% of dimer), which was confirmed by mass spectrometry and $^1\text{H-NMR}$ spectroscopy (see S23–S27 and S33–S34), that showed the disappearance of the olefinic doublets at 7.6 ppm and 6.5 ppm and the appearance of two multiplets at 5.1 ppm and 3.9 ppm, consistent with the cyclobutane ring formation (Fig. 3a) of just a single regioisomer (see below).⁵⁸ Additionally, $^1\text{H-NMR}$ analysis revealed significant upfield shifts of the TACN protons upon photodimerization (Fig. 3a, red squares), consistent with a closer proximity to the pyrene ring in the cyclobutane structure. COSY and HSQC experiments (see Fig. S26 and S27) further confirmed the splitting of TACN CH_2 protons into two populations, indicating magnetic anisotropy effects from the pyrene ring.⁵⁹ The UV-vis spectra (Fig. 3c) showed that the characteristic absorption band of the conjugated $E-1 \cdot Zn(II)$ ($\lambda_{\text{max}} = 372\text{ nm}$) undergoes a hypochromic shift in the isomer $Z-1 \cdot Zn(II)$ ($\lambda_{\text{max}} = 350\text{ nm}$), which is replaced by the typical structured absorption band of non-conjugated pyrene in the photodimer $2 \cdot Zn(II)$ ($\lambda_{\text{max}} = 269, 279, 334\text{ and }350\text{ nm}$). This behaviour is fully consistent with the successful $[2 + 2]$ cycloaddition.^{44,60,61} The mismatch between the monomer absorption maximum (375 nm) and the red-shifted photodimerization wavelength (420 nm) also aligns well with earlier findings.^{62–65} Reversibility of the dimerization can be achieved by irradiating the dimer solution at 311 nm (Fig. 3b and d). However, photodegradation hinders complete reversibility of the process (see S31 and S32), which regrettably occurs even increasing the irradiation wavelength to 365 nm.

We then decided to elucidate the structure of the obtained photodimer $2 \cdot Zn(II)$. The $[2 + 2]$ cycloaddition of acrylamidylpyrene can theoretically result in 11 different regioisomers (see Table S1 for details), depending on the nature of

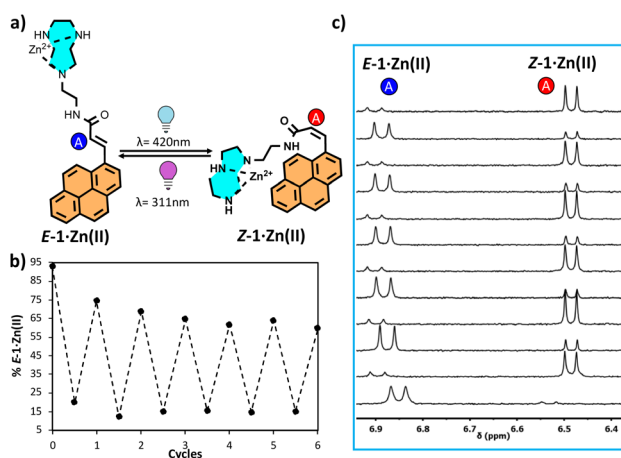


Fig. 2 (a) Reaction scheme of the reversible photoisomerization between $E-1 \cdot Zn(II)$ and $Z-1 \cdot Zn(II)$. (b) Graph showing the percentage of $E-1 \cdot Zn(II)$ isomer as a function of the isomerization cycles (420/311 nm) and (c) corresponding $^1\text{H-NMR}$ highlighting the doublet typical of the E and Z isomers. The coupling constants J^2 of the olefinic protons signals are consistent with the E and Z isomers (respectively 16 Hz (6.9 ppm) and 12 Hz (6.5 ppm), see S28). The concentration-dependent variation in the chemical shift of the doublets arising from proton A of $1 \cdot Zn(II)$ suggests the presence of weak intermolecular interactions involving mainly the E isomer—at least to some extent.

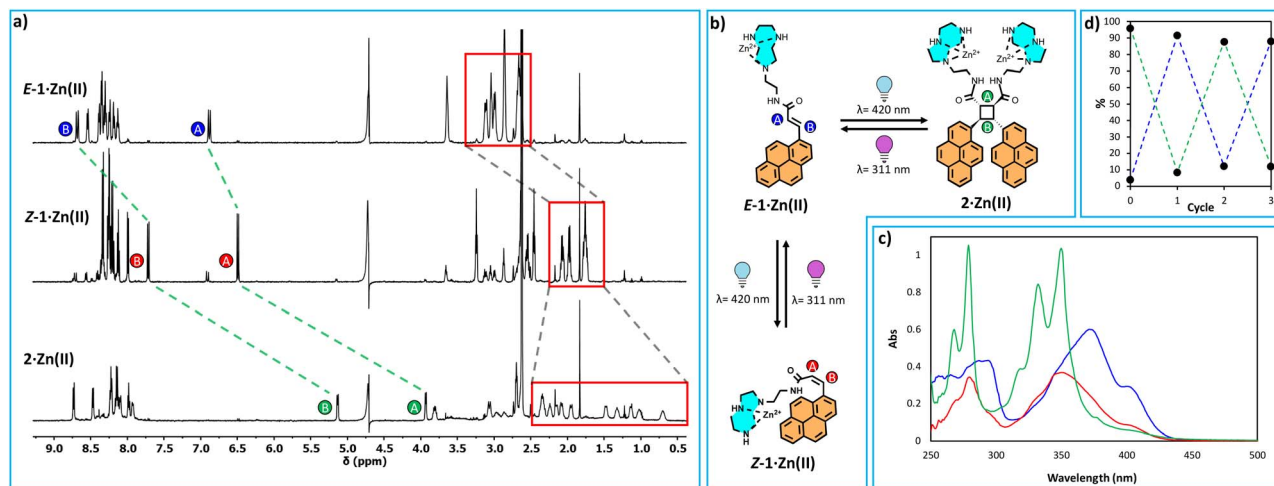


Fig. 3 (a) Comparison between ^1H -NMR of $E\text{-}1\cdot\text{Zn(II)}$, $Z\text{-}1\cdot\text{Zn(II)}$ and $2\cdot\text{Zn(II)}$; the styryl and the corresponding cyclobutane protons are highlighted with orange and green dots, while the CH_2 TACN $\cdot\text{Zn(II)}$ protons are highlighted with red squares. (b) Reaction scheme showing the reversible photoisomerization and photodimerization processes. (c) Comparison between UV-vis spectra of 20 μM water solutions of $E\text{-}1\cdot\text{Zn(II)}$ (blue), $Z\text{-}1\cdot\text{Zn(II)}$ (red) and $2\cdot\text{Zn(II)}$ (green). (d) Graph showing the percentage of $E\text{-}1\cdot\text{Zn(II)}$ (blue) and $Z\text{-}1\cdot\text{Zn(II)}$ (green) as a function of the irradiation cycles (420/311 nm) for a 0.5 mM solution of $E\text{-}1\cdot\text{Zn(II)}$ in $\text{D}_2\text{O}:\text{DMSO-}d_6$ 8 : 2.

the reactive monomers (E and/or Z) and on their orientation (head/tail, and syn/anti). Given that the ^1H -NMR pattern of the cyclobutane protons signals are a fingerprint of the isomer, the initial focus for establishing the structure of the dimer was placed on the cyclobutane protons at 5.1 ppm and 3.9 ppm (Fig. S47). The observed multiplet pattern, consistent with an $\text{AA}'\text{BB}'$ spin system, was in agreement with a cyclobutane ring exhibiting C_2 symmetry.^{66–69} This reduced the pool of possible candidates to only two isomers obtained by the head-to-head cycloaddition of either two $E\text{-}1\cdot\text{Zn(II)}$ or two $Z\text{-}1\cdot\text{Zn(II)}$ molecules (respectively EE-hh-anti and ZZ-hh-anti in Fig. 4a). Notably, NOE experiments cannot discriminate between these two isomers, as the distances between the $\text{A/A}'$ and $\text{B/B}'$ proton pairs are nearly identical in both isomers (2.4 Å for ZZ-hh-anti and 2.5 Å for EE-hh-anti). To resolve this ambiguity, we examined the coupling constants within the $\text{AA}'\text{BB}'$ spin system.⁶⁹ Since sterically hindered substituents can alter the geometry and the coupling networks of the cyclobutane ring, we employed DFT calculations to predict the J couplings for the cyclobutane protons of the EE-hh-anti and ZZ-hh-anti photodimers.

The optimized geometries of EE-hh-anti and ZZ-hh-anti dimers (Fig. 4b) were employed to calculate the coupling constants of the cyclobutane protons (NMR data are summarized in Tables S2–S4).

Comparison of the experimental ^1H -NMR signal of the protons resonating at $\delta_{\text{exp}} = 5.2$ ppm with simulated patterns based on DFT-calculated J values revealed a clear match for the EE-hh-anti isomer, while significant deviations were observed for the ZZ-hh-anti isomer (Fig. 4c and see Section S5 for details). This indicated that the photodimerization proceeded *via* the $[2 + 2]$ cycloaddition of two $E\text{-}1\cdot\text{Zn(II)}$ monomers, yielding the EE-hh-anti isomer as the single photoproduct (Fig. 4d).

Interestingly, the dimerizing monomer is the less concentrated species, as approximately 85% of the initial monomer has

already isomerized to the Z form. Although 20% DMSO prevents large-scale aggregation (see S39), ^1H -NMR experiments suggest that weak intermolecular interactions between $E\text{-}1\cdot\text{Zn(II)}$ monomers may still occur (see Fig. 2). These interactions could contribute to the high photodimerization efficiency observed even at low concentrations of the E -isomer (see S42). Additionally, it is worth noting that the photoisomerization rate is

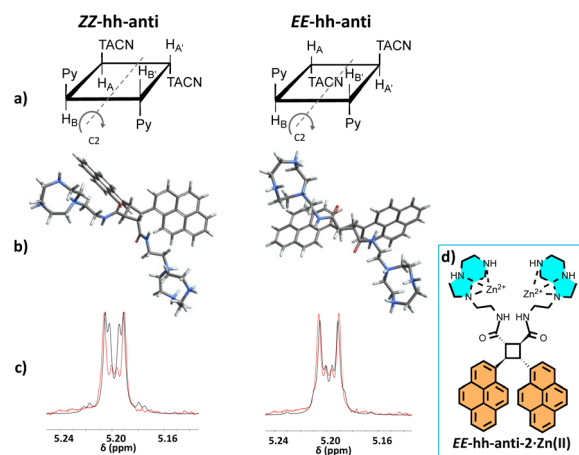


Fig. 4 (a) Schematic representations of the ZZ-hh-anti (left) and EE-hh-anti (right) isomers, with highlighted the C_2 symmetry axes. (b) Optimized structures of EE- and ZZ-hh-anti isomers. The cyclobutane ring protons of EE-hh-anti adopt quasi-*anti* conformations relative to each other, leading to relatively large 3J couplings. Conversely, in ZZ-hh-anti , one proton pair features a dihedral angle close to 90° , typically associated with small 3J couplings. (c) Overlap between experimental ^1H -NMR spectrum (red traces) and the simulated spectra from DFT-calculated NMR parameters for EE- and ZZ-hh-anti (black traces). The chemical shifts of calculated spectra were aligned to the experimental value of 5.20 ppm. (d) Molecular structures of the obtained $\text{EE-hh-anti-}2\cdot\text{Zn(II)}$ photodimer.

significantly faster than the photodimerization rate, allowing rapid *Z*-to-*E* isomerization to replenish the *E*-1·Zn(II) species as it is consumed during photodimerization.

After having established the molecular structure of the photodimer 2·Zn(II), we explored the catalytic properties of the system in the monomeric and dimeric states. The aim was to determine whether the photodimerization process enhances the system's catalytic activity toward the transesterification reaction of 2-hydroxypropyl-4-nitrophenylphosphate (HPNPP), which is a model substrate for RNA (Fig. 5a). Hence, the formation of *p*-nitrophenol (PNP) from a 1 mM solution of HPNPP was followed *via* UV/vis spectroscopy, in the presence of the monomers *E*-1·Zn(II) and *Z*-1·Zn(II) (100 μM), or 50 μM of the photodimer 2·Zn(II). These experiments (Fig. 5 and Section 4 of the SI) showed that the *E*-*Z* photoisomerization of 1·Zn(II) did not significantly affect the initial rate of the transesterification (0.75 mM h⁻¹ for *E*-1·Zn(II) and 0.93 mM h⁻¹ for *Z*-1·Zn(II)), but that the clustering of two TACN·Zn(II) units in 2·Zn(II) caused by photodimerization led to a 6-fold increase of the catalytic activity (4.22 mM h⁻¹ for 2·Zn(II)). The observed rate acceleration enhancement aligns with similar molecular systems containing two or more proximal TACN·Zn(II) units.^{70–72}

One of the most attractive features of the use of light for tuning the catalytic properties is the possibility to temporally control a chemical reaction.⁷³ Thus, because of the significant difference in catalysis between monomers and dimer, we decided to study how the catalytic activity varies depending on the irradiation time. We observed that the presence of HPNPP and the resulting nitrophenolate anion interfered with the photodimerization process. To circumvent this practical limitation of the model substrate and to maintain precise control over the photochemical step, we decoupled catalyst irradiation from substrate conversion.⁷⁴ This approach allowed us to conceptually investigate whether irradiation time can effectively regulate catalytic activity. Aliquots were taken after different irradiation times and studied by ¹H-NMR spectroscopy to determine the mixture composition and by UV-vis spectroscopy to determine the catalytic activity. In particular, a solution of *E*-1·Zn(II) was irradiated at 420 nm to form the [2 + 2] cycloadduct,

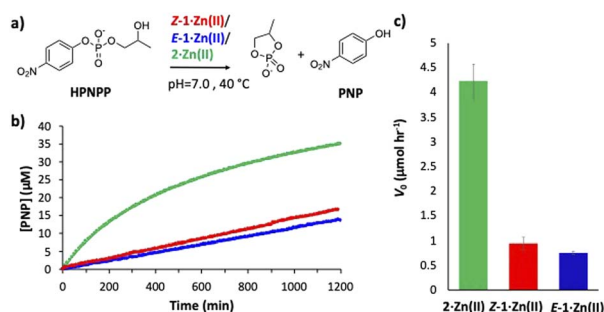


Fig. 5 (a) Reaction scheme of the studied HPNPP transesterification. (b) Concentration of the formed PNP as a function of the reaction time for a pH 7 solution of HPNPP 1 mM at 40 °C, in the presence of: *E*-1·Zn(II) (100 μM, blue line), or *Z*-1·Zn(II) (100 μM, red line), or 2·Zn(II) (50 μM, green line). (c) Graph highlighting the observed initial reaction rates.

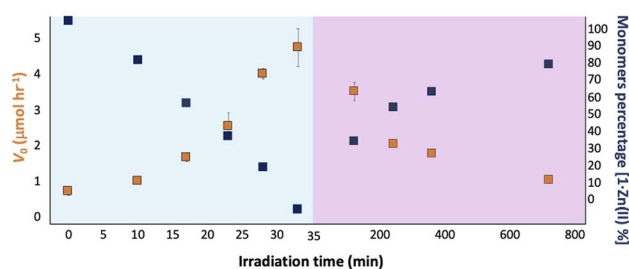


Fig. 6 Temporal control of HPNPP transesterification through photodimerization. The initial reaction rate of HPNPP transesterification (orange squares) and the percentage of monomers (1·Zn(II), blue squares) are shown as a function of the irradiation time. In particular, a 1 mM solution of *E*-1·Zn(II) was initially irradiated at 420 nm for 35 minutes (green background), and then at 311 nm for 700 minutes (pink background). Over the time, this solution was analysed *via* ¹H-NMR and aliquots (corresponding to 50 μM of 1·Zn(II)) were added to 1 mM HPNPP solutions, pH 7.0, 40 °C. The formation of PNP was followed *via* UV spectroscopy.

and the catalytic activity at different irradiation times was studied by taking 5 different aliquots of the irradiated mixture and adding these to a 1 mM HPNPP solution (Fig. 6, 0–35 minutes). In these experiments the irradiated *E*-1·Zn(II) solution was 1 mM instead of 0.5 mM (see Fig. 2 and 3), leading to a faster dimerization rate. Consequently, complete dimerization was achieved in just 33 minutes, and the irradiated wavelength was set to 311 nm to cause the dissociation of the photodimer 2·Zn(II). At 4 different times aliquots of the solution were taken and added to a 1 mM HPNPP solution. The initial rates of transesterification reaction were measured and plotted as a function of the irradiation time (Fig. 6, 0–800 minutes). The irradiation at 420 nm of a *E*-1·Zn(II) increased the transesterification rate of HPNPP about 6-fold, reaching the maximum after 33 minutes. This timing well correlates with the dimer formation rate, which indeed reaches the highest concentration after 33 minutes of irradiation (see also Fig. S43 and S44). On the other hand, subsequent irradiation at 311 nm caused a decrease in the reaction rate, which returned to initial values after around 700 minutes. Also, this trend well corresponds with the slower kinetic of the monomerization reaction, in fact 700 min of irradiation are needed to decrease the dimer concentration to 21%.

Conclusions

While photoisomerization processes are extensively explored for controlling system reactivity, the use of photodimerization reactions for this purpose remains largely unexplored. Here, we have described a styrylpyrene-based molecular system equipped with a TACN·Zn(II) catalytic unit that undergoes regioselective [2 + 2] cycloaddition upon visible light irradiation (420 nm). In addition to photodimerization, we observed *E*-to-*Z* photoisomerization, which does not hinder the process as prolonged irradiation drives the equilibrium toward the dimeric state. Even if photodegradation affects the efficiency, both the photoinduced processes are reversible, with irradiation at 311 nm restoring the original *E* isomer.

Remarkably, the photodimerization is highly regioselective, yielding only one photoisomer. Detailed $^1\text{H-NMR}$ spectral analysis, supported by DFT-calculations, confirmed the photodimer's *EE*-*hh*-anti geometry. Investigating the catalytic properties of this dimer toward HPNPP transesterification revealed a significant finding: while photoisomerization has no impact on catalysis, the $[2 + 2]$ cycloaddition creates a dimer whose clustered catalytic units result in a 6-fold enhancement of the reaction rate compared to the monomeric system.

These results illustrate the potential of visible light to drive the covalent assembly of catalytically active molecules, enabling temporal control over system reactivity. Remarkably, while many photoswitches suffer from the thermodynamic instability of their active form, the catalytically active photodimer is a robust product that does not spontaneously revert to its monomeric inactive state. By establishing photodimerization as a viable alternative to photoisomerization for regulating catalysis, this work opens new avenues for designing light-responsive networks and advancing temporal reaction control strategies.

Author contributions

The manuscript was written through contributions of all authors. T. M., A. N. and M. L. performed the synthesis, the NMR and UV-vis spectroscopic experiments. F. R. performed DFT calculation and NMR simulation. S. B. UV-vis experiments and review, LP writing – review & editing. L. G. designed the project, analysed the data, supervised the research and wrote the manuscript.

Conflicts of interest

There are no conflicts to declare.

Data availability

The data supporting this article have been included as part of the SI, including the synthetic procedures and the characterizations of all products reported in this study. Details on photoreactivity studies and DFT calculations are included as well. See DOI: <https://doi.org/10.1039/d5sc03858h>.

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