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## Enzyme-inspired flavin–polydopamine as a biocompatible nanoparticle photocatalyst†

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**A new approach aimed at designing an enzyme-inspired photocatalyst is presented that exploits the inherent photocatalytic activity of flavin and the facile polymerization of dopamine to afford hybrid cofactor-containing nanoparticles. The flavin–polydopamine system benefits from ease of synthesis, tunability in terms of size and activity, and excellent temporal control over the catalyzed reactions. This novel, versatile photocatalyst exhibits both photooxidation and photoreduction of chromogenic enzymatic substrates. In addition, the prepared hybrid nanoparticles are shown to be non-toxic, paving the way to their use in a wider range of applications beyond green catalysis, such as antioxidant adjuvants to various therapeutic approaches.**

Enzymes are not only protein catalysts that drive biochemical processes responsible for the sustenance of life, but due to their specificity, biodegradability, and efficiency both in terms of substrate conversion and energy usage, they are highly desirable for green chemistry applications. However, the key disadvantages of wider enzyme use out of the lab-setting are difficulties associated with their isolation in large quantities, post-reaction separation, low stability under non-physiological conditions such as high temperature or pH, and sensitivity to organic solvents.<sup>1,2</sup> Several approaches have been developed to address these issues, each with their own set of advantages and drawbacks. For example, more robust enzymes and enzyme-inspired systems have been designed either by immobilization or encasing natural enzymes,<sup>3</sup> by use of directed evolution to make efficient non-natural enzymes,<sup>4</sup> or by utilization of various materials to prepare enzyme mimics.<sup>5,6</sup> The latter has particularly benefited from developments in nanotechnology and the emergence of inorganic materials with enzyme-like activities known as nanozymes.<sup>7,8</sup> The most prominent

### New concepts

The use of nanomaterials as enzyme-like catalysts has risen in popularity over the last decade, with particular focus on inorganic nanomaterials (nanozymes). However, the versatility and efficiency of enzymes are difficult to match and there are continuous efforts to propose new design strategies. By taking inspiration from natural flavoenzymes, we designed a novel biocompatible nanoparticle system that is light-responsive and shows tuneable activity. Our approach employs polydopamine (PDA) nanoparticles as a flavin carrier, which mimics the tryptophan environment of natural flavin enzymes. Nanoparticles of tuneable size and flavin content are prepared *via* a facile co-polymerisation of dopamine and flavin–dopamine conjugates. Temporal control over both the oxidation and reduction is demonstrated using visible light irradiation and two different enzyme substrates. In addition, a preliminary *in vitro* assay confirmed a high level of biocompatibility, as well as potential antioxidant effect. Our findings open a new route to the design of biocompatible and tuneable nano-sized, enzyme-inspired photocatalysts, which exhibit a versatile reactivity profile, are tuneable both in terms of structure and reactivity, and are characterized by ease of fabrication and low toxicity. This profile makes them suitable for use in green catalysis, but also in nanomedicine and design of organic energy harvesting systems.

examples of nanozymes to date are iron oxide (Fe<sub>3</sub>O<sub>4</sub>) nanoparticles, with high catalytic activity similar to those of peroxidases and catalase, ceria nanoparticles (CeO<sub>2</sub>), and metallic nanoparticles such as gold (Au NP) and silver (Ag NPs). Over the past decade, these materials have been explored as low-cost and robust alternatives to natural enzymes, with the focus on industrial applications for chemical synthesis,<sup>9,10</sup> water remediation,<sup>11</sup> or as fuel additives to reduce pollution.<sup>12</sup> Lately, the attention of researchers working on nanozymes has shifted more towards nanomedicine and therapeutic applications in biofilm disruption, antioxidation, tumor removal and tissue engineering.<sup>8,13</sup> Despite their attractiveness due to ease of synthesis and storage, one of the drawbacks of nanozyme use is their poorly understood toxicological profile including their long-term toxicity and environmental impact.<sup>14,15</sup> This calls for new strategies and rational design approaches to afford a new generation of biocompatible nanozymes tailored to a particular application.<sup>16</sup> To get closer to this goal, we have set out to rationally design a new generation of hybrid

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nanozymes combining an organic enzyme component with a nanostructured material, which is not a passive carrier of an enzymatic centre, but actively engages in the catalytic process.

We took inspiration from flavoenzymes, an important group of oxidoreductases that catalyse a large variety of different types of redox reactions, thanks to their isoalloxazine-based flavin cofactors, flavin adenine mononucleotide (FAD) and flavin dinucleotide (FAD). As a result of their wide-ranging activity profile, flavoenzymes are implemented within a range of biological processes such as metabolic pathways, respiratory chain reactions and photochemical processes.<sup>17–20</sup> The latter has contributed to the development of flavin photocatalysts as an efficient, green and mild strategy to afford light-driven transformations such as *E/Z* isomerisation, [2+2] cycloaddition, electrophilic chlorination and sulfide oxidation.<sup>21</sup> However, although efficient homogenous systems have been explored, there are issues related to such systems such as limited separation of the products from catalysts hindering both the product purity and the reuse of the catalysts. There are only a few examples of heterogeneous flavin photocatalysts and they are yet to show efficiency and selectivity similar to their natural counterparts.<sup>22–24</sup> There are examples, however, of very efficient ‘dark’ catalytic flavin systems utilising artificial and biomimetic polymeric structures.<sup>25–27</sup> With this in mind, and to further advance the promising field of flavin photocatalysis, we aimed to develop an enzyme-inspired photocatalytic system by embedding flavin within an active nanostructured polymer matrix employing an easy-to-use copolymerization strategy.

We have chosen polydopamine (PDA) as the carrier polymer due to its reported biocompatibility,<sup>28,29</sup> and structural and

electronic properties.<sup>30</sup> Although PDA has extensively been used as a molecular adhesive because of its ability to form coatings on virtually any surface,<sup>31</sup> a more interesting property from a synthetic chemistry point of view is its intrinsic catalytic activity.<sup>32–34</sup> Composed of a sequence of extended  $\pi$ -systems, PDA can be considered an amorphous organic semiconductor and has already been employed as an electron gate for artificial photosynthesis,<sup>35</sup> and as a mediator in various catalytic systems based on metallic nanoparticles,<sup>36</sup> and even whole cells.<sup>37</sup> Although the exact mechanism of PDA’s action in these applications has not yet been deciphered, it has been shown that PDA accelerates the rate of electron transfer by acting as an electron acceptor.<sup>35</sup>

Based on these studies, we hypothesized that the combination of flavin and PDA would result in an interesting system that displays a wider activity profile significantly enhancing the current repertoire of catalytic flavin systems. Within natural flavoenzymes, function and catalytic activity are defined by specific interaction between a flavin cofactor and surrounding amino acids. For example, a key reaction enabling switching from a dark to light-adapted state of BLUF (sensors of blue light using FAD) photo-receptors, which guides biological signaling pathways, is the electron transfer between tyrosine and a flavin cofactor.<sup>38</sup> The structure of PDA (Fig. 1A) mimics the tyrosine moieties suggesting that in a flavin–PDA conjugate, the polymer can actively engage in the catalytic activity and electron transfer. We therefore set out to prepare a flavin–dopamine conjugate, and formulate our catalysts in the form of flavin–PDA nanoparticles (FLPDA NPs). Such nanoparticles are characterized by an increased



**Fig. 1** (A) Illustration of flavin–polydopamine (FLPDA) NPs highlighting the polydopamine structure. (B) Comparison of naturally occurring flavin cofactor FAD and FLDA synthesized in this paper. The STEM images of the NPs prepared using different ratios of FLDA and dopamine: (C) FLPDA-5, scale bar = 2  $\mu$ m, (D) FLPDA-10, scale bar = 200 nm.

surface area aiding the charge transfer (CT) and substrate interaction with the photo-excited flavin species whilst offering more flexibility in terms of future applications. For example, the surface of the nanoparticles could be additionally modified to contain particular cell targeting moieties or stabilizing shells to tailor the stability and physiochemical properties of the system to a desired application.

To prepare the **FLPDA** NPs, a flavin-dopamine monomer (**FLDA**, Fig. 1B, right) was first prepared that mimics the structure of the naturally occurring FAD (Fig. 1B, left). The ribityl chain was replaced with a triethylene glycol (TEG) moiety, and adenosine diphosphate with the dopamine analogue, 3,4-dihydroxyphenylacetic acid (DOPAC) (see ESI†, Section 1.3 for synthetic details). Following the synthesis of a monomer, a copolymerization of **FLDA** with dopamine (DA) was performed at room temperature using a water/ethanol solvent system in the presence of ammonia as previously described for bare polydopamine (PDA) NP formation.<sup>39</sup> As shown in Fig. 1C and D, spherical **FLPDA** NPs ranging from 100 to 200 nm were obtained depending on the ratio of used monomers.

In order to explore the effect of the amount of **FLDA** on the co-polymerisation and formation of NPs, 5 : 1, 10 : 1 and 20 : 1 molar ratios of DA to **FLDA** were used, resulting in NPs referred to as **FLPDA-5**, **FLPDA-10**, and **FLPDA-20** respectively. The prepared samples were washed extensively and lyophilized to avoid any thermal decomposition of the polymer that could occur through vacuum heating. They were noticeably browner in color than black unmodified PDA and their zeta potential values ranged from  $-35.7$  to  $-41.5$  mV for **FLPDA-5** and **FLPDA-20** respectively indicating high stability and dispersibility (see Table S1, ESI†). The size and shape of the **FLPDA** particles were further investigated by STEM (Fig. 1C and D and ESI†) showing that all the samples contained spherical NPs of similar size (approx. 200 nm) with relatively large size distribution ( $\pm \sim 50$  nm). Some conjoined particles and the largest size distribution was observed in **FLPDA-5** (see Table S1, ESI†). In contrast, PDA synthesized under the same conditions displays a narrow size distribution with an average size of  $110 \pm 18$  nm (see Fig. S4, ESI†), which indicates that the presence of flavin moieties affects the polymerization mechanism and oligomer aggregation to form particles with less defined shape and size. This is most likely due to H-bonding and electrostatic interactions between the flavin group and oligomeric units, and our ongoing work is focused on optimizing the polymerization procedure further.

To validate the presence of flavin moieties, particles were analyzed by UV-Vis absorption and fluorescence spectroscopy. The UV-Vis spectrum of our **FLDA** monomer shows absorption bands at  $\lambda_{\text{max}} = 445$  nm and 373 nm corresponding to the transitions from the ground state ( $S_0$ ) to the  $S_1$  ( $\lambda_{\text{max}} \sim 442$ – $450$  nm) and  $S_2$  ( $\lambda_{\text{max}} \sim 360$ – $375$  nm) excited states (see Fig. S6A, ESI†).<sup>18</sup> These bands are redshifted to  $\sim 456$  nm for the  $S_0 \rightarrow S_1$  transition and  $\sim 376$  nm for the  $S_0 \rightarrow S_2$  transition in **FLPDA** (Fig. S6B, ESI†). This can be explained by an increase in proton donation from PDA,<sup>40</sup> and by the electron-withdrawing inductive effects on the flavin moieties due to incorporation into the highly conjugated PDA system.<sup>41</sup>

The fluorescence emission spectrum of **FLDA** is characterized by emission maxima at  $\sim 527$  nm ( $\lambda_{\text{ex}} = 450$  nm) which correlates well to other known flavin compounds (see Fig. S7A, ESI†).<sup>42</sup> The **FLPDA** samples also exhibit this signal, confirming the presence of flavin moieties in the particles (see Fig. S7B, ESI†). An earlier study in which flavin compounds were incubated with melanin (structurally analogous to PDA) showed that the fluorescent properties of flavins did not change upon binding and there was no significant fluorescence quenching by the polymer.<sup>43</sup> That study, however, focused on the non-covalent binding of the flavin derivatives to melanin, and not on the covalent attachment used in our work. In order to validate the assumption that flavin content can be indeed quantified *via* fluorescence measurements of the particles, a DOPAC-free flavin analogue, **FLOH** was prepared and compared to the DOPAC conjugated monomer **FLDA** (see Fig. S8A, ESI†). We observed that the fluorescence intensity was much lower for **FLDA** when compared to the same concentration of **FLOH**, despite the absorbance of the compounds being very similar. This observation indicates some intramolecular isoalloxazine-catechol fluorescence quenching similar to that found for FAD and adenosine due to a possible formation of a 'closed' structure (see Fig. S8B and C, ESI†).<sup>44</sup> Kozik *et al.* suggested that the key phenolic and indolic groups in the melanin polymer do not incur any fluorescence quenching due to their inability to form CT complexes with the isoalloxazine ring.<sup>43</sup> Although we initially found out that there was a fluorescence decrease when **FLOH** was incubated with the PDA NPs, this was no longer valid after we corrected for the absorbance of PDA at the excitation and emission wavelengths. The fluorescence spectrum shows a slight redshift to 527 nm similar to that of **FLDA** and **FLPDA**, but at the same intensity as **FLOH** alone (Fig. S9, ESI†). This demonstrates that the flavin moiety bound covalently to the polymer experiences the same quenching as the monomer flavin bound to the dopamine since the formation of the CT complex between isoalloxazine and other units appears to be negligible.

Taking this into account, a fluorescence calibration curve was obtained using known concentrations of **FLDA** to approximate the flavin concentration within the nanoparticles. As expected, **FLPDA-5** contains a higher proportion of flavin than **FLPDA-10** and **FLPDA-20**; approximately 1.11, 0.84, and 0.71  $\mu\text{mol mg}^{-1}$  **FLPDA**, respectively (see Fig. S10 and Table S2, ESI†). Clearly, the relative amounts of flavin do not exactly match the initial monomer ratios used for the synthesis. This can be explained by base-catalyzed cleavage and/or hydrolysis of some flavin moieties during the polymerization, which was previously reported for other flavins.<sup>45,46</sup> The presence of flavin within the nanoparticles was additionally confirmed by FTIR measurements showing characteristic C–H stretching vibration bands at 2924 and 2855  $\text{cm}^{-1}$ , the sharp bands at 1545  $\text{cm}^{-1}$  and 1580  $\text{cm}^{-1}$  relating to  $\nu(\text{C}=\text{N})$  modes in the isoalloxazine ring and flavin carbonyl  $\nu(\text{C}=\text{O})$  and  $\nu(\text{C}=\text{C})$  at 1711 and 1680  $\text{cm}^{-1}$ , respectively (see Fig. S11, ESI† for more details).

Following the preparation and characterization of the **FLPDA** NPs, we set out to explore their photocatalytic activity. This was done by exploiting the fluorescent compound resorufin (**RF**, absorption  $\lambda_{\text{max}} = 572$  nm) and using the precursor



**Scheme 1** The photo-redox catalytic cycles of resorufin (RF) production starting from Amplex Red (AR photooxidation) and resazurin (RZ photoreduction) in the presence of FLPDA.

compound Amplex Red (AR) to monitor oxidation, and resazurin (RZ) to monitor the reduction reactions (Scheme 1). In a typical experiment, 1–10  $\mu\text{g mL}^{-1}$  FLPDA NPs were incubated with AR (100  $\mu\text{M}$ ) in 10 mM KPi buffer at pH 7.4 (2 mL) and then irradiated with a blue laser diode (458 nm), monitoring the absorption spectrum in real time (see Fig. 2A and ESI<sup>†</sup> for set up details). Our hypothesis that FLPDA would be able to carry out photooxidation was confirmed by conversion of AR to RF upon irradiation with FLPDA-5 showing the highest conversion rate of 3.69  $\mu\text{M h}^{-1}$  using an excitation intensity of 1.19  $\text{mW cm}^{-2}$ , due to the higher concentration of flavin moieties (see Fig. 2B).

Control experiments were also performed at the same excitation intensity using the FLDA monomer, unmodified PDA and a blank sample containing AR without any catalyst. The latter was carried out as it has been previously reported that trace amounts of excited state RF can catalyse the oxidation of AR.<sup>47</sup> In our case, however, the blank measurement showed a much slower rate of conversion (0.24  $\mu\text{M h}^{-1}$ ) and no reaction was observed for PDA (10  $\mu\text{g mL}^{-1}$ ) (Fig. 2B). The latter was expected due to high photon absorption capability of PDA, which has already been successfully utilized for the design of light-harvesting devices.<sup>48–50</sup>



**Fig. 2** (A) Plot of the absorbance changes observed for the photooxidation of AR to RF by FLPDA-5 (10  $\mu\text{g mL}^{-1}$ ) in KPi buffer (10 mM, pH 7.4). (B) Comparative plot of AR photooxidation in the presence of FLPDA photocatalysts and controls (10  $\mu\text{g mL}^{-1}$ , [FLDA] = 10 mol%). (C) Plot of the absorbance changes observed for the photoreduction of RZ to RF by FLPDA-5 (10  $\mu\text{g mL}^{-1}$ ) in the presence of EDTA (100  $\mu\text{M}$ ) in KPi buffer (10 mM, pH 7.4). (D) Plot showing the relative consumption of RZ and production of RF by FLPDA-5 (10  $\mu\text{g mL}^{-1}$ ) in the presence of EDTA (100  $\mu\text{M}$ ) in KPi buffer (10 mM, pH 7.4). (E) The comparative plot of RZ photoreduction in the presence of the FLPDA photocatalysts and controls (10  $\mu\text{g mL}^{-1}$ , [FLDA] = 10 mol%). All experiments were done using a 458 nm light source with 6.40  $\text{mW cm}^{-2}$  excitation intensity, apart from (B) which was obtained using 2.95  $\text{mW cm}^{-2}$  excitation intensity.

When compared to **FLPDA**, a much slower reaction, with only  $0.43 \mu\text{M}$  **RF** being produced per hour, was observed with the **FLDA** monomer (10 mol%). This large difference indicates a potential synergistic relationship between PDA and flavin within the hybrid material, which results in stabilization of reactive intermediates and/or enhancement of CT. To gain a better understanding of this effect, we further investigated the kinetic profile of **AR** photooxidation by **FLPDA-5** using the Michaelis–Menten model. This showed that our system does display enzyme-like kinetic behaviour indicating substrate binding, leading to reaction and subsequent product dissociation. The  $K_m$  value obtained from this fit was  $166.7 \mu\text{M}$  with  $V_{\text{max}}$  of  $1.66 \mu\text{M min}^{-1}$  (see Fig. S12A and B, ESI†). Compared to enzymes with high specificity to Amplex Red, such as horseradish peroxidase (HRP), these values indicate less specificity and efficiency ( $6.4 \mu\text{M}$  and  $2.2 \mu\text{M min}^{-1}$  respectively) but still in a range to be considered as an effective nanozyme.<sup>51</sup> Overall, we envisage that aromatic stacking between **AR**, PDA and flavin allows favourable orientation for catalysis, therefore, enhancing the activity of **FLPDA** over homogenous **FLDA** – similar to previously reported flavin–PPH<sub>3</sub>–functionalised gold nanoclusters.<sup>52</sup> It can also be assumed that non-specific binding of **AR** to the PDA polymer alone is responsible for a decrease in the activity for **FLPDA-10** (less flavin, more PDA) when compared with **FLPDA-5** (less PDA, more flavin).

We also observed that higher laser power accelerates the conversion process, which clearly demonstrates the photocatalytic character of **FLPDA** as one would expect more absorbed photons to lead to higher activity (see Fig. S13, ESI†). Also, no reaction occurred between the catalyst and substrate in the dark over a period of >18 hours (see Fig. S14, ESI†). It should also be noted that the STEM images of the recovered NPs do not show significant changes in the shape or morphology after irradiation (see Fig. S5C and D, ESI†).

Photoreductive capabilities of our catalytic system were also investigated due to the importance of catalytic reduction reactions in organic chemistry. Although these reactions are often used to yield a range of synthons and feedstocks, they traditionally employ costly and toxic rare earth metals or hydride complexes as catalysts.<sup>53</sup> Only recently, photo-biocatalytic approaches employing enzymes emerged as safer and milder alternatives. In one such example, reduction of the C=C bond was achieved using old yellow enzyme homologue from *B. subtilis* and ethylenediaminetetraacetic acid (EDTA) as an electron donor.<sup>54</sup> To explore the potential of our hybrid nanoparticles for photoreductions, we employed a blue dye resazurin (**RZ**, absorption  $\lambda_{\text{max}} = 602 \text{ nm}$ ) as a substrate and EDTA as the electron donor to afford **RF** (Scheme 1).

Similarly to previous experiments,  $1\text{--}10 \mu\text{g mL}^{-1}$  **FLPDA** NPs were irradiated with a blue laser diode (458 nm) at different excitation intensities but now in the presence of **RZ** ( $25 \mu\text{M}$ ) and EDTA ( $100 \mu\text{M}$ ) in degassed 10 mM KPi buffer at pH 7.4 (2 mL) under N<sub>2</sub> atmosphere. The spectral changes were then monitored in real time, following the decrease of **RZ** absorption at 602 nm and the increase of **RF** absorption at 572 nm (see Fig. 2C). Fast reduction of the **RZ** substrate and production of **RF** was observed

showing complete conversion after 4 min of irradiation (Fig. 2D). The decrease in **RF** concentration after 4 min can be attributed to bleaching. Similar to the **AR** photooxidation reactions, the STEM images of the recovered NPs do not show significant changes in the shape or morphology (see Fig. S5E and F, ESI†).

Control runs were also performed (Fig. 2E) showing that PDA ( $10 \mu\text{g mL}^{-1}$ ) displayed almost no activity whereas, some conversion of **RZ** to **RF** ( $0.58 \mu\text{M min}^{-1}$  compared to  $5.7 \mu\text{M min}^{-1}$  for **FLPDA-5**) was observed in the blank measurement containing EDTA, but no catalyst. As previously reported, this is due to the interaction of triplet state **RZ** and electron dense tertiary aliphatic amines.<sup>55</sup> Irradiation of **RZ** without EDTA led to **RZ** bleaching and no conversion to **RF** (data not shown). As with our photooxidation assay, the conversion rate was doubled for **FLPDA-5** compared to **FLPDA-10** due to a larger number of flavin active sites. The same amount of **FLDA** monomer showed 6 times slower conversion rate ( $0.95 \mu\text{M min}^{-1}$ ) providing further evidence of the enhanced activity of **FLPDA** through the substrate–PDA–flavin interactions.

Furthermore, we investigated the switchability of our catalytic system using **FLPDA** in the presence of the **AR** substrate. A clear increase in absorption of the product **RF** at 572 nm is observed when the system is irradiated (ON state) and no further reaction takes place when the laser is switched off for 30 min (OFF state) as evidenced by the absorbance plateau (see Fig. 3). This demonstrates that the catalytic activity of **FLPDA** towards oxidation of the substrate is light-dependent and has an excellent capacity for temporal control, a feature that is widely sought after in green chemistry to minimize the cost and maximize the efficiency of a reaction.<sup>56</sup> Another factor that is valuable in green catalysis is the recyclability of a catalyst. Accordingly, we tested the recyclability of **FLPDA-5** in the **RZ** photoreduction reaction by removing the catalyst post-reaction *via* centrifugation. The catalyst was then washed and reused in subsequent reactions, maintaining good activity for up to 4 runs (see Fig. S15, ESI†). Due to the inherent losses experienced during catalyst recovery by washing/centrifugation our future work is focussed on developing a continuous flow system using immobilised **FLPDA**.

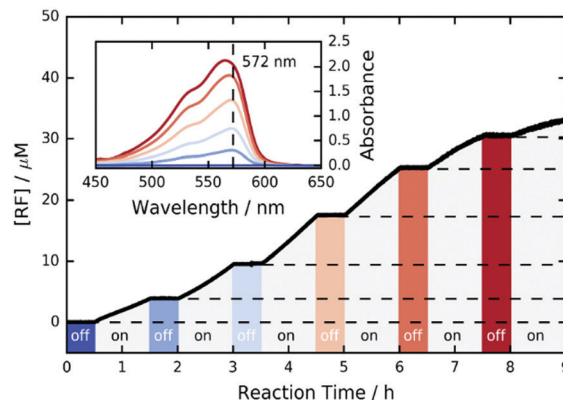


Fig. 3 ON/OFF cycle data for the photooxidation of **AR** by **FLPDA-5** ( $10 \mu\text{g mL}^{-1}$ ) in KPi Buffer (10 mM, pH 7.4) using  $2.95 \text{ mW cm}^{-2}$  excitation intensity (458 nm).

The promising results regarding **FLPDA**'s activity prompted us to explore the photodecomposition of the system. The photostability of flavin compounds is often an issue when these compounds are utilised in photocatalysis. Namely, upon irradiation the absorbance of flavin at  $\sim 445$  nm decreases due to dealkylation to either lumichrome or lumiflavin as well as scission of the alloxazine ring.<sup>57</sup> To see if the same happens with dopamine-conjugates, we irradiated the monomer **FLDA** and the analogue **FLOH** alone in a solution using 458 nm light (50.3 mW) under  $N_2$  atmosphere and in the air over 30 min at pH 7.4. Remarkably, in both air and under  $N_2$  atmosphere, **FLDA** shows almost no photodecomposition when compared to **FLOH** (see Fig. S16, ESI<sup>†</sup>). This could be explained by possible CT between alloxazine and catechol in a 'closed' conformation akin to FAD as rationalised based on our fluorescence data. The decrease in absorbance was far greater for **FLOH** in the air when compared to  $N_2$ , most likely due to the photoinduced ROS formation increasing the rate of decomposition.

The same study was also performed for **FLPDA-5** by recording the UV-Vis absorbance and fluorescence spectra of the particles before and after 30 minutes irradiation, as well as the supernatant after centrifugation of the particles (see Fig. S17, ESI<sup>†</sup>). The absorbance spectra of the particles in both air and nitrogen still showed the presence of characteristic flavin bands (Fig. S17A, ESI<sup>†</sup>). However, the fluorescence of the particles after the reaction had decreased (Fig. S17B, ESI<sup>†</sup>). Analysis of the supernatants showed emission at  $\sim 524$  nm most likely indicating the presence of the lumiflavin indicating that some flavins were cleaved (Fig. S12B, ESI<sup>†</sup>).<sup>57</sup> This could be remedied by the inclusion of a sacrificial electron donor such as EDTA which effectively limited the extent of this degradation (see Fig. S17A and B, ESI<sup>†</sup>).<sup>58</sup> Nevertheless, it is clear that the incorporation of PDA does enhance the photostability of the flavin moieties through the formation of CT complexes increasing the recyclability. Our current work is looking at different dopamine-flavin conjugation sites and the possible effects this could have on both the catalytic activity and the photodecomposition.

Finally, considering the potential uses of such a system in nanomedicine and water remediation, we performed a preliminary investigation of the inherent toxicity of the **FLPDA** nanoparticles using three *in vitro* assays. First, **FLPDA** was PEGylated to increase the stability in cell culture media (DMEM) and reduce immunogenicity when applied *in vivo* applications.<sup>59</sup> This was achieved through the covalent attachment of amino-terminated mPEG<sub>5000</sub> to **FLPDA** by incubation in 10 mM TRIS buffer at pH 8.5 for 18 h. Cell viability was investigated *via* an MTS assay after 24 h of incubation with the particles. As shown in Fig. 4, (black diamonds) **PEG-FLPDA** has a negligible effect on the cell proliferation over a wide dosage range until the concentration of particles is significantly increased (to  $1.0 \text{ mg mL}^{-1}$ ) at which point the viability is reduced to 54.6%. We also tested the cytotoxicity of **PEG-FLPDA** using a lactate dehydrogenase (LDH) assay, which measures the release of LDH and signals significant cell damage. This was only observed for very high concentration dosages of  $1.0 \text{ mg mL}^{-1}$  after incubation for 24 h (Fig. 4, blue dots), correlating well to the MTS assay.

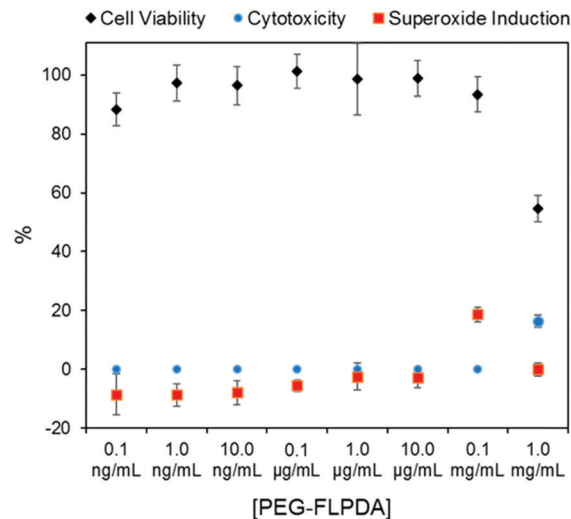


Fig. 4 Summary of *in vitro* assay data for A549 cells showing cell viability (*via* MTS assay), cytotoxicity (*via* LDH assay) and superoxide generation in the presence of **PEG-FLPDA** after 24 h.

To complement these assays, we additionally explored the effect of **PEG-FLPDA** on the cell oxidative stress using a superoxide detection assay to monitor the induction or scavenging of superoxide within the cells due to the presence of the particles. The data in Fig. 4 (red squares) show that the amount of superoxide decreases (relative to untreated controls) upon incubation with up to  $10.0 \mu\text{g mL}^{-1}$  **PEG-FLPDA**. This indicates that **FLPDA** has antioxidant properties similar to PDA and melanin.<sup>60–62</sup> However, at a concentration of  $0.1 \text{ mg mL}^{-1}$ , incubated cells showed an increased accumulation of superoxide, although this seems not to impact the viability of the cells. Decrease of the superoxide production again at  $1.0 \text{ mg mL}^{-1}$  is correlated to the widespread cell death at this high concentration. However, these three assays clearly indicate that the **FLPDA** NPs are non-toxic below concentrations of  $1.0 \text{ mg mL}^{-1}$ , and even show some antioxidative effect below  $0.1 \text{ mg mL}^{-1}$ . These first studies indicate the high potential of these flavin-containing hybrid systems not only for photocatalysis but also for potential use in water remediation and biomedical applications.

## Conclusions

In conclusion, we have designed a novel, effective and versatile organic photocatalyst based on the combination of two natural species, flavin and polydopamine. Such an enzyme-inspired catalyst can be prepared using a straightforward, green chemistry protocol, is easy to purify and handle, can be used both for photooxidation and photoreduction, and can be activated on demand. To our knowledge, this is the first example of an organic polymer nanoparticle with embedded flavin being used as a photocatalyst for oxidation and reduction of the organic species. *In vitro* assays performed to demonstrate the non-toxicity of the system have also indicated some antioxidative properties, prompting further studies exploring the use of this material for biomedical applications. We believe that this enzyme-inspired

material could find application beyond organic catalysis, in particular within biomedical research as an effective antioxidant adjuvant or drug nanocarrier, and in combination with suitable substrates in the design of organic photovoltaic systems.

## Conflicts of interest

There are no conflicts to declare.

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## References

- M. T. Reetz, *Chem. Rec.*, 2016, **16**, 2449–2459.
- S. Li, X. Yang, S. Yang, M. Zhu and X. Wang, *Comput. Struct. Biotechnol. J.*, 2012, **3**, e201209017.
- M. P. Thompson, I. Peñafiel, S. C. Cosgrove and N. J. Turner, *Org. Process Res. Dev.*, 2019, **23**, 9–18.
- F. H. Arnold, *Angew. Chem., Int. Ed.*, 2017, **57**, 4143–4148.
- R. Breslow, *Artificial Enzymes*, Wiley-VCH, Weinheim, 2006.
- E. Kuah, S. Toh, J. Yee, Q. Ma and Z. Gao, *Chem. – Eur. J.*, 2016, **22**, 8404–8430.
- H. Wei and E. Wang, *Chem. Soc. Rev.*, 2013, **42**, 6060–6093.
- X. Wang, Y. Hu and H. Wei, *Inorg. Chem. Front.*, 2016, **3**, 41–60.
- L. L. Chng, N. Erathodiyil and J. Y. Ying, *Acc. Chem. Res.*, 2013, **46**, 1825–1837.
- L. Pasquato, P. Pengo and P. Scrimin, *Supramol. Chem.*, 2005, **17**, 163–171.
- W. Zhang, *J. Nanopart. Res.*, 2003, **5**, 323–332.
- V. Sajith, C. B. Sobhan and G. P. Peterson, *Adv. Mech. Eng.*, 2010, **2**, 581407.
- J. Golchin, K. Golchin, N. Alidadian, S. Ghaderi, S. Eslamkhah, M. Eslamkhah and A. Akbarzadeh, *Artif. Cells, Nanomed., Biotechnol.*, 2017, **45**, 1069–1076.
- M. A. Maurer-Jones, I. L. Gunsolus, C. J. Murphy and C. L. Haynes, *Anal. Chem.*, 2013, **85**, 3036–3049.
- A. Sukhanova, S. Bozrova, P. Sokolov, M. Berestovoy, A. Karaulov and I. Nabiev, *Nanoscale Res. Lett.*, 2018, **13**, 1–21.
- B. Liu and J. Liu, *Nano Res.*, 2017, **10**, 1125–1148.
- C. Walsh, *Acc. Chem. Res.*, 1980, **13**, 148–155.
- P. F. Heelis, *Chem. Soc. Rev.*, 1982, **11**, 15–39.
- V. Massey, *Biochem. Soc. Trans.*, 2000, **28**, 283–296.
- E. Silva and A. M. Edwards, *Flavins Photochemistry and Photobiology*, Royal Society of Chemistry, Cambridge, 2006.
- B. König, S. Kümmel, E. Svobodová and R. Cibulka, *Phys. Sci. Rev.*, 2018, **3**, 20170168.
- H. Schmaderer, P. Hilgers, R. Lechner and B. König, *Adv. Synth. Catal.*, 2009, **351**, 163–174.
- J. Špačková, E. Svobodová, T. Hartman, I. Stibor, J. Kopecká, J. Cibulková, J. Chudoba and R. Cibulka, *ChemCatChem*, 2017, **9**, 1177–1181.
- M. Kurfiřt, J. Špačková, E. Svobodová and R. Cibulka, *Monatsh. Chem.*, 2018, **149**, 863–869.
- B. J. Jordan, G. Cooke, J. F. Garety, M. A. Pollier, N. Kryvokhyzha, A. Bayir, G. Rabani and V. M. Rotello, *Chem. Commun.*, 2007, 1248–1250.
- Y. Arakawa, K. Minagawa and Y. Imada, *Polym. J.*, 2018, **50**, 941–949.
- Y. Chevalier, Y. L. T. Ki, D. L. Nouen, J. P. Mahy, J. P. Goddard and F. Avenier, *Angew. Chem., Int. Ed.*, 2018, **57**, 16412–16415.
- S. Hong, K. Y. Kim, H. J. Wook, S. Y. Park, K. D. Lee, D. Y. Lee and H. Lee, *Nanomedicine*, 2011, **6**, 793–801.
- X. Liu, J. Cao, H. Li, J. Li, Q. Jin, K. Ren and J. Ji, *ACS Nano*, 2013, **7**, 9384–9395.
- Y. Liu, K. Ai and L. Lu, *Chem. Rev.*, 2014, **114**, 5057–5115.
- H. Lee, S. M. Dellatore, W. M. Miller and P. B. Messersmith, *Science*, 2007, **318**, 426–430.
- Z. Yang, J. Sun, X. Liu, Q. Su, Y. Liu, Q. Li and S. Zhang, *Tetrahedron Lett.*, 2014, **55**, 3239–3243.
- R. Mrówczyński, A. Bunge and J. Liebscher, *Chem. – Eur. J.*, 2014, **20**, 8647–8653.
- Y. Du, H. C. Yang, X. L. Xu, J. Wu and Z. K. Xu, *ChemCatChem*, 2015, **7**, 3822–3825.
- J. H. Kim, M. Lee and C. B. Park, *Angew. Chem., Int. Ed.*, 2014, **53**, 6364–6368.
- A. Kunfi, V. Szabó, Á. Mastalir, I. Bucsi, M. Mohai, P. Németh, I. Bertóti and G. London, *ChemCatChem*, 2017, **9**, 3236–3244.
- L. Wang, Z.-Y. Hu, X.-Y. Yang, B.-B. Zhang, W. Geng, G. Van Tendeloo and B.-L. Su, *Chem. Commun.*, 2017, **53**, 6617–6620.
- T. Mathes, I. H. M. Van Stokkum, M. Stierl and J. T. M. Kennis, *J. Biol. Chem.*, 2012, **287**, 31725–31738.
- K. Ai, Y. Liu, C. Ruan, L. Lu and G. Lu, *Adv. Mater.*, 2013, **25**, 998–1003.
- A. Kotaki, M. Naoi and K. Yagi, *J. Biochem.*, 1970, **68**, 287–292.
- L. N. Mataranga-Popa, I. Torje, T. Ghosh, M. J. Leitl, A. Späth, M. L. Novianti, R. D. Webster and B. König, *Org. Biomol. Chem.*, 2015, **13**, 10198–10204.
- A. Kotaki and K. Yagi, *J. Biochem.*, 1970, **68**, 509–516.
- A. Kozik, W. Korytowski, T. Sarna and A. S. Bloom, *Biophys. Chem.*, 1990, **38**, 39–48.
- G. Weber, *Biochem. J.*, 1950, **47**, 114–121.
- S. B. Smith and T. C. Bruice, *J. Am. Chem. Soc.*, 1975, **14**, 2875–2881.
- T. Harayama, Y. Tezuka, T. Taga and F. Yoneda, *J. Chem. Soc., Perkin Trans. 1*, 1987, 75–83.
- F. A. Summers, B. Zhao, D. Ganini and R. P. Mason, *Methods Enzymol.*, 2013, **526**, 1–17.
- M. Lee, J. U. Kim, J. S. Lee, B. Il Lee, J. Shin and C. B. Park, *Adv. Mater.*, 2014, **26**, 4463–4468.
- A. Xie, K. Zhang, F. Wu, N. Wang, Y. Wang and M. Wang, *Catal. Sci. Technol.*, 2016, **6**, 1764–1771.

- 50 M. Lee, J. U. Kim, K. J. Lee, S. Ahn, Y. B. Shin, J. Shin and C. B. Park, *ACS Nano*, 2015, **9**, 6206–6213.
- 51 H. S. Rho, A. T. Hanke, M. Ottens and H. Gardeniers, *PLoS One*, 2016, **11**, e0153437.
- 52 Y. Imada, M. Osaki, M. Noguchi, T. Maeda, M. Fujiki, S. Kawamorita, N. Komiya and T. Naota, *ChemCatChem*, 2015, **7**, 99–106.
- 53 R. A. Sheldon, I. W. C. E. Arends and U. Hanefeld, *Green Chemistry and Catalysis*, Wiley-VCH, Weinheim, 2007.
- 54 M. M. Grau, J. C. Van Der Toorn, L. G. Otten, P. Macheroux, A. Taglieber, F. E. Zilly, I. W. C. E. Arends and F. Hollmann, *Adv. Synth. Catal.*, 2009, **351**, 3279–3286.
- 55 C. Bueno, M. L. Villegas, S. G. Bertolotti, C. M. Previtali, M. G. Neumann and M. V. Encinas, *Photochem. Photobiol.*, 2002, **76**, 385–390.
- 56 R. Göstl, A. Senf and S. Hecht, *Chem. Soc. Rev.*, 2014, **43**, 1982–1996.
- 57 M. A. Sheraz, S. H. Kazi, S. Ahmed, Z. Anwar and I. Ahmad, *Beilstein J. Org. Chem.*, 2014, **10**, 1999–2012.
- 58 S. Alonso-De Castro, E. Ruggiero, A. Ruiz-De-Angulo, E. Rezabal, J. C. Mareque-Rivas, X. Lopez, F. López-Gallego and L. Salassa, *Chem. Sci.*, 2017, **8**, 4619–4625.
- 59 J. S. Suk, Q. Xu, N. Kim, J. Hanes and L. M. Ensign, *Adv. Drug Delivery Rev.*, 2016, **99**, 28–51.
- 60 Y. Liu, K. Ai, X. Ji, D. Askhatova, R. Du, L. Lu and J. Shi, *J. Am. Chem. Soc.*, 2017, **139**, 856–862.
- 61 M. D'Ischia, A. Napolitano, V. Ball, C. T. Chen and M. J. Buehler, *Acc. Chem. Res.*, 2014, **47**, 3541–3550.
- 62 K. Y. Ju, Y. Lee, S. Lee, S. B. Park and J. K. Lee, *Biomacromolecules*, 2011, **12**, 625–632.