

Visible-Light-Activated Carbon Monoxide Release from Porphyrin–Flavonol Hybrids

Andrea Ramundo, Jiří Janoš, Lucie Muchová, Mária Šranková, Jakub Dostál, Miroslav Kloz, Libor Vitek, Petr Slaviček,* and Petr Klán*



Cite This: *J. Am. Chem. Soc.* 2024, 146, 920–929



Read Online

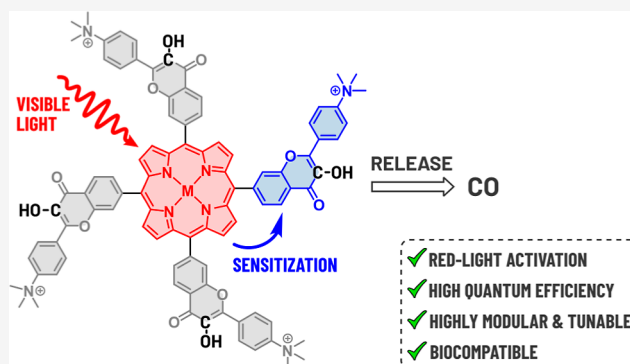
ACCESS |

Metrics & More

Article Recommendations

Supporting Information

ABSTRACT: We report on porphyrin–flavonol hybrids consisting of a porphyrin antenna and four covalently bound 3-hydroxyflavone (flavonol) groups, which act as highly efficient photoactivatable carbon monoxide (CO)-releasing molecules (photoCORMs). These bichromophoric systems enable activation of the UV-absorbing flavonol chromophore by visible light up to 650 nm and offer precise spatial and temporal control of CO administration. The physicochemical properties of the porphyrin antenna system can also be tuned by inserting a metal cation. Our computational study revealed that the process occurs via endergonic triplet–triplet energy transfer from porphyrin to flavonol and may become feasible thanks to flavonol energy stabilization upon intramolecular proton transfer. This mechanism was also indirectly supported by steady-state and transient absorption spectroscopy techniques. Additionally, the porphyrin–flavonol hybrids were found to be biologically benign. With four flavonol CO donors attached to a single porphyrin chromophore, high CO release yields, excellent uncaging cross sections, low toxicity, and CO therapeutic properties, these photoCORMs offer exceptional potential for their further development and future biological and medical applications.



INTRODUCTION

Carbon monoxide (CO), a small gaseous signaling molecule with inherent toxicity,¹ controls various critical cellular processes, with mitochondria being its primary target, and manifests anti-inflammatory, antiproliferative, and cytoprotective properties.^{2,3} It soon gained attention as a promising therapeutic agent in several settings, such as organ transplantation or cancer treatments.^{4–6} Remarkably, CO can stimulate cancer cells to switch to a pure oxidative metabolism in an anti-Warburg fashion,⁷ leading to growth inhibition and cell death.^{8,9} Many studies suggest that CO is a useful gaseous drug, but its acute toxicity poses severe limitations in its administration under a clinical scenario. A controlled, targeted release is therefore essential.

Photoactivatable CO-releasing molecules (photoCORMs) can serve as prodrugs that release CO after the absorption of a photon.⁴ Light as a control stimulus provides the required spatiotemporal control to circumvent CO toxicity.¹⁰ Activation wavelengths in the so-called “phototherapeutic window” (650–900 nm) allow for a deep light penetration without impairing cellular damage.¹¹ Therefore, the development of red- or near-infrared (NIR)-absorbing photoCORMs is desirable. The first examples of photoCORMs included transition metal–carbonyl complexes,¹² cyclopropenones,^{13,14} or 1,2-dioxolane-3,4-diones,¹⁵ all absorbing in the UV region.

α -Diketones¹⁶ ($\lambda_{\max} = 470$ nm) and a xanthene-based chromophore¹⁷ ($\lambda_{\max} = 488$ nm) have been developed to deliver CO under visible-light activation. A *meso*-carboxy BODIPY¹⁸ ($\lambda_{\max} = 652$ nm) was the first red-light activatable organic photoCORM, yet suffering from a poor uncaging cross-section ($\Phi_{\text{CO}}\epsilon_{\max}$).

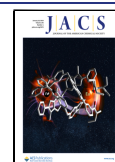
3-Hydroxyflavone **1** (flavonol; 3-hydroxy-2-phenylchromen-4-one, *Scheme 1A*) belongs to the family of flavonoids, a class of naturally occurring molecules.¹⁹ It undergoes photooxygenation to give CO and a salicylic acid derivative as the main photoproducts upon irradiation with UV light ($\lambda_{\max} = 355$ nm).^{20,21} Several attempts have been made to improve flavonol absorption properties. π -Extended flavonols with λ_{\max} in the 400–540 nm range have been developed,^{22–24} although still retaining the same mechanism of photodecarbonylation.²⁵ Our laboratory used an alternative approach of combining an established cyanine 7 scaffold with the flavonol moiety,

Received: October 14, 2023

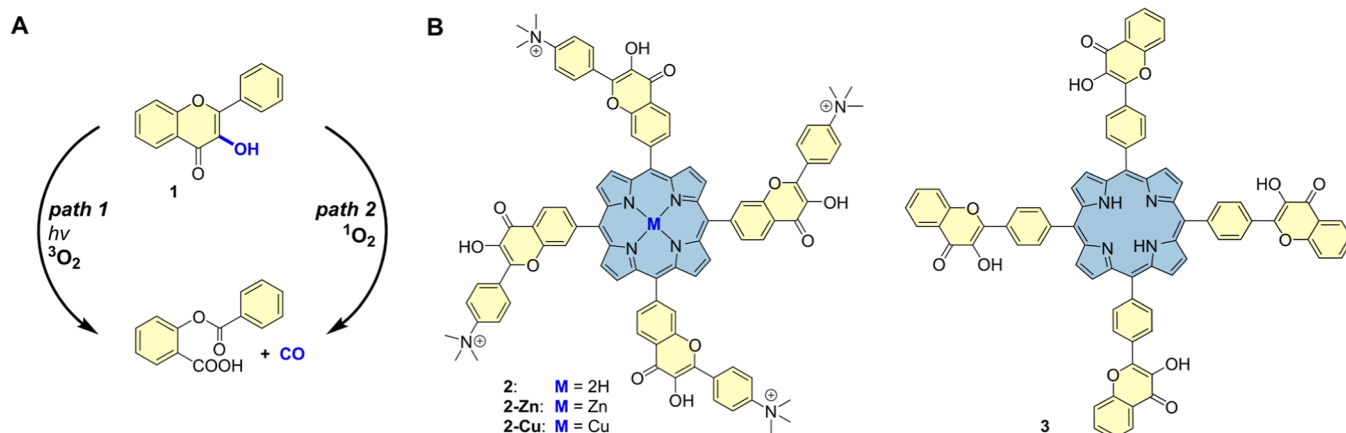
Revised: November 26, 2023

Accepted: December 11, 2023

Published: December 29, 2023



Scheme 1. Flavonol Chemistry and Studied Compounds; (A) (Photo)decarbonylation of Flavonol 1; (B) Structures of Porphyrin–Flavonol Hybrids 2 and 3 Composed of a Central Light-Harvesting Porphyrin Sensitizer (Blue) and Four CO-Releasing Flavonol Moieties (Yellow) Studied in This Work



pushing the absorption of the resulting cyanine-flavonol hybrids to the NIR region ($\lambda_{\max} = 750\text{--}790\text{ nm}$).^{26,27}

More recently, 3,4'-dihydroxyflavone derivatives have been embedded in a polymer matrix with 5-(4-hydroxyphenyl)-10,15,20-triphenylporphyrin used as a $^1\text{O}_2$ generator to trigger CO release.²⁸ Still, the combined requirements for a successful biological application of a photoCORM, such as water solubility, absorption near or within the phototherapeutic window, high quantum efficiency of CO release, and very low or no toxicity, remain a challenge.

In a recent study, some of us showed that porphyrins could be repurposed as a new class of photoactivatable molecules releasing biologically relevant substances, such as indibulin or methotrexate, attached to the porphyrin *meso*-methyl position upon visible-light activation.²⁹

In this work, we used a porphyrin moiety operating as an antenna and a triplet energy donor that transfers triplet energy to four CO-releasing flavonol moieties attached to the *meso* positions (2 and 3, Scheme 1B). The introduction of metal ions into the porphyrin core of 2 was aimed to affect the photochemical efficiencies, singlet oxygen production, and other physicochemical properties of the hybrids. Steady-state optical and transient absorption (TA) spectroscopies and quantum-chemical calculations were used to evaluate the reaction mechanism that involves the sensitization of flavonol by triplet-excited porphyrin. We also studied the systems' applicability in cell culture experiments through fluorescence microscopy, cytotoxicity determination, and high-resolution respirometry to assess mitochondrial metabolism.

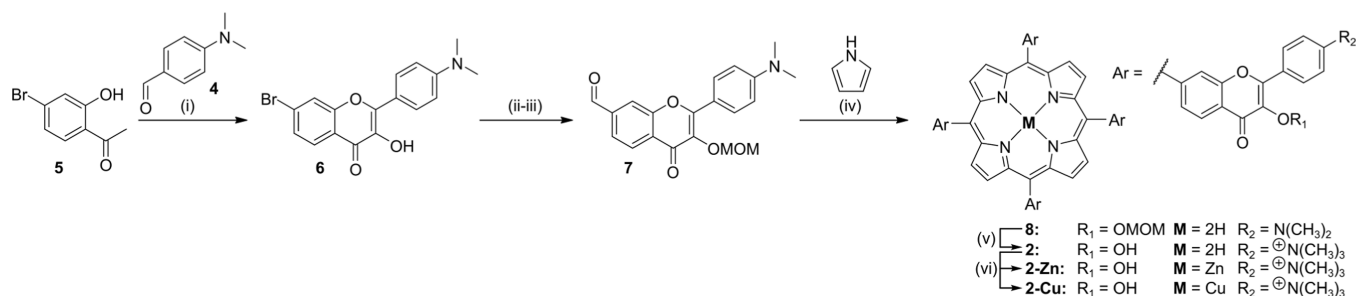
RESULTS AND DISCUSSION

In our search for photoCORMs operating at the edge of or within the phototherapeutic window,³⁰ we designed structures 2 and 3, consisting of the central porphyrin moiety as an antenna bearing four UV-absorbing flavonol units ($\lambda_{\max} = 355\text{ nm}$) in the four *meso* positions, to improve atom economy of the CO production (up to 4 equiv of CO released per one starting compound). The nature of this aromatic macrocycle, absorbing across the entire visible spectrum (400–670 nm),^{31,32} would allow for fine-tuning of its spectroscopic and photochemical properties by incorporating metal ions into the porphyrin core and other structural modifications.³³

To experimentally determine whether porphyrins can sensitize CO release from flavonols, methanol solutions containing either *meso*-(tetra-4-carboxyphenyl)porphyrin (TcPP) or its zinc complex (ZnTcPP) and a flavonol derivative (1, or its more water-soluble 4'-trimethylammonium analog—see the Supporting Information) were irradiated with red light (600 or 650 nm) where flavonol does not absorb. CO was liberated with excellent yields of up to 0.96 equiv, whereas only 0.08–0.14 equiv of CO were generated upon direct irradiation of flavonols at 365 nm in the absence of a porphyrin sensitizer (Table S1).

Two possible mechanisms of CO release from flavonol derivatives have been postulated.^{20,25} The first one involves the reaction between triplet-excited flavonol and ground-state oxygen ($^3\text{O}_2$, path 1, Scheme 1A) or ground-state flavonol and singlet oxygen ($^1\text{O}_2$) formed in situ by photosensitization (path 2). These two orthogonal mechanisms can operate simultaneously, but their relative efficiencies are driven by the acid/base equilibria related to the OH group ($\text{pK}_a = 8.7$).³⁴ For instance, path 2 becomes predominant only when the conjugate base of 1 is considered.²⁴ However, only the acid form of 1 was detected by UV–vis spectroscopy in methanol. Indeed, we found the reactivity of ground-state flavonol 1 toward $^1\text{O}_2$ to be very low ($k_{\Sigma} = 2.5 \times 10^5\text{ M}^{-1}\text{ s}^{-1}$). The calculated photodecarbonylation quantum yield due to the reaction with singlet oxygen was found to be 10^{-4} , which is lower by more than an order of magnitude than that of the CO production via direct irradiation (Supporting Information). In addition, ZnTcPP was irradiated as a singlet oxygen generator in the presence of 1 at 600 nm, where ZnTcPP absorbs exclusively, in a methanol solution with and without a large excess of NaN_3 as a singlet oxygen scavenger (Table S2). The CO production was the same, providing experimental evidence that $^1\text{O}_2$ is not involved in the release of CO from flavonol 1 when porphyrin is used as a sensitizer.

These encouraging results led to the design, synthesis, and studies of porphyrin–flavonol hybrids 2 and 3 (Scheme 1B). 4-(Dimethylamino)benzaldehyde (4) and 4'-bromo-2'-hydroxyacetophenone 5 were converted to substituted flavonol 6 via the Algar–Flynn–Oyamada reaction,³⁵ and the bromo substituent in 6 was used for installing an aldehyde group in 7 through a Pd-catalyzed formylation with CO/H_2 (Scheme 2). The methoxymethyl (MOM) protection of the 3-hydroxyl

Scheme 2. Synthesis of the Porphyrin–Flavonol Hybrids **2**, **2-Zn**, and **2-Cu**^a

^aReaction conditions: (i) NaOH, H₂O₂, CH₃OH, 55%; (ii) MOMBr, CH₂Cl₂, 0 °C, 89%; (iii) Pd(OAc)₂, cataCXium, TMEDA, CO/H₂, 8 atm, toluene, 100 °C, 54%; (iv) pyrrole, TFA, DDQ, CHCl₃, 23 °C, 18%; (v) CH₃OTf, HOTf, CH₃CN, 23 °C, 54%; (vi) **2-Zn**: Zn(OTf)₂, DMF, 130 °C, 77%. **2-Cu**: Cu(OTf)₂, CH₃OH, 23 °C, 73%.

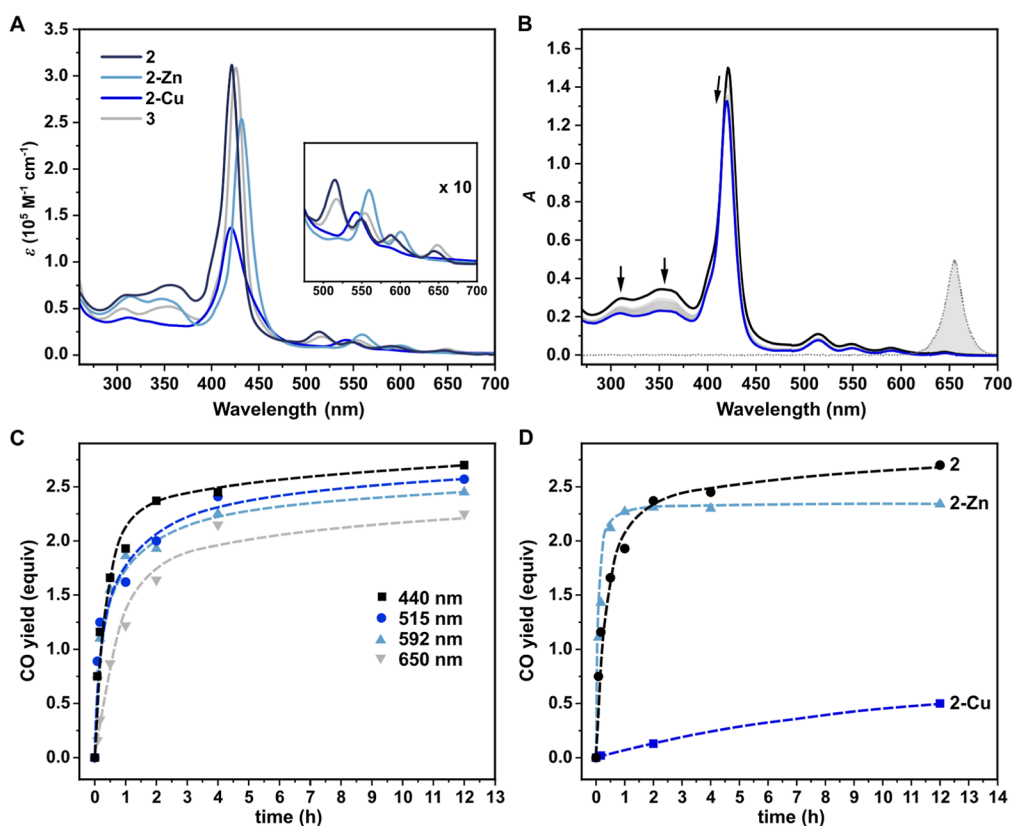


Figure 1. Photochemical properties of studied hybrids. (A) Absorption spectra of **2**, **2-Zn**, and **2-Cu** (methanol), and **3** (methanol/DMSO, 4:1, v/v). Inset: a 10-fold magnification of the Q-band region. (B) Irradiation of **2** in aerated methanol ($5 \mu\text{M}$) at 650 nm; the spectra were recorded every 60 min, and the spectra prior to (black line) and after (blue line) 12 h irradiation are highlighted. The emission spectrum of the irradiation source is shown as a gray dotted line. (C) Time-dependent CO evolution from **2** in aerated methanol ($5 \mu\text{M}$) upon irradiation at different wavelengths in the 0–12 h interval. (D) Time-dependent CO evolution in aerated methanol from **2**, **2-Zn**, and **2-Cu** upon irradiation at the Soret band in the 0–12 h interval. The yield of CO released was determined by headspace GC–MS and expressed as equivalents of CO released by 1 equiv of the photoCORM.

group was necessary as this functional group strongly interfered with macrocyclization. In the next step, the porphyrin–flavonol hybrid **8** was obtained by condensing pyrrole and the aldehyde **7** using the well-established Lindsay conditions.³⁶ Finally, one-pot methylation of the dimethylamino groups in **8** and concomitant MOM deprotection afforded the target molecule **2** on a gram scale. The use of column chromatography was necessary only after the formation of porphyrin **8** step (iv), while conventional purification methods (extraction and precipitation) were used during the other synthetic steps. Hybrid **3** was obtained

on a multigram scale using the same strategy, and no chromatographic methods were required throughout this synthetic sequence. Detailed synthetic procedures are provided in the [Supporting Information](#).

The absorption spectra of **2** and **3** are analogous to those of metal-free porphyrins³⁷ and exhibit strong and sharp bands in the blue region (Soret bands), along with four less intense bands in the 500–670 nm range (Q bands) (Figure 1A, Table 1). Two additional bands at 310 and 360 nm correspond to the electronic transitions of the flavonol groups in their conjugate acid forms. Hybrids **2** and **3** emit in the red-to-NIR region (λ_{em}

Table 1. Photophysical and Photochemical Properties of the Porphyrin–Flavonol Hybrids^{a,c}

compd	$\lambda_{\text{abs}}/\text{nm}$ ($\epsilon/10^4 \text{ M}^{-1} \text{ cm}^{-1}$) ^b	$\lambda_{\text{em}}/\text{nm}$	Φ_{f}^c	CO yield ^d /equiv	Φ_{CO}^e	$\Phi_{\text{CO}}\epsilon_{\text{max}}^f/\text{M}^{-1} \text{ cm}^{-1}$ at ($\lambda_{\text{abs}}/\text{nm}$)	Φ_{Δ}^g
2	421 (31.3), 514 (2.4), 549 (1.3), 590 (0.9), 645 (0.4)	649, 715	0.035	2.9 ± 0.1 2.7 ± 0.1 ^h	0.018 ± 0.001	(5.6 ± 0.3) × 10 ³ (421)	0.17
2-Zn	431 (25.6), 560 (2.6), 601 (0.9)	611, 660	0.017	2.3 ± 0.1 2.1 ± 0.1 ^h	0.040 ± 0.002	(1.0 ± 0.1) × 10 ⁴ (431)	0.1
2-Cu	421 (15.4), 543 (1.8), 582 (0.7)	ND	ND	0.8 ± 0.1	0.002 ± 0.001	(2.9 ± 0.8) × 10 ² (421)	<10 ⁻³
3	426 (33.9), 517 (2.0), 554 (1.5), 591 (0.8), 650 (0.7)	652, 717	0.036	3.3 ± 0.1 2.1 ± 0.1 ^h	0.003 ± 0.001	(1.0 ± 0.2) × 10 ³ (426)	0.32 ⁱ

^aAll measurements were performed in methanol for **2**, **2-Zn**, **2-Cu**, and methanol/DMSO (4:1, v/v) for **3**, unless stated otherwise. ^bMolar absorption coefficients ϵ . ^cFluorescence quantum yields Φ_{f} upon excitation at the Soret band. ^dTotal chemical yields of released CO monitored by GC headspace, obtained upon exhaustive irradiation at the Soret band. ^eAbsolute quantum yields of CO release determined using a Si-photodiode. ^fCO uncaging cross sections $\Phi_{\text{CO}}\epsilon_{\text{max}}$ at (λ_{max}). ^gSinglet oxygen production quantum yields obtained from the ¹O₂ luminescence. ^hIn PBS. ⁱIn DMF. ND = not detected.

= 650, 715 nm; Figure S4B) with a moderate fluorescence quantum yield (Φ_{f}) of 0.035. The absorption spectra of the hybrids resemble the sum of the absorption spectra of individual chromophores (Figure S4A).

Both compounds were found to be stable in solutions for days at room temperature in the dark (Figure S5). Irradiation of compound **2** or **3** ($c \sim 5 \mu\text{M}$) at 650 nm in methanol led to a decrease in the intensity of the bands at 310 and 360 nm, accounting for the decomposition of the flavonols units, and a decrease in the porphyrin Soret band, most likely due to self-sensitized oxidation of the macrocycle.³⁸ Analogous photochemical behavior was observed upon irradiation at 440, 515, and 600 nm (Figure S6). This consistent wavelength-independent behavior suggests that the entire visible range can be used for photoactivation.

Hybrid **2** irradiated in the range of 440–650 nm, liberated CO with maximum chemical yields of 2.9 ± 0.1 equiv in methanol and 2.7 ± 0.1 equiv in a phosphate buffer saline (PBS) solution. The yields of CO from **3** in methanol/DMSO (4:1, v/v) were higher (3.3 ± 0.1 equiv) but dropped to 2.1 ± 0.1 equiv in PBS. The time-dependent CO evolution in Figure 1C shows that **2** released most of CO in the first 30–60 min. The quantum yield of CO release (Φ_{CO}) for **2** in methanol, 0.018 ± 0.001, was found to be wavelength-independent across the entire visible range (Table S6) and was orders of magnitude higher than the photodecomposition of the porphyrin core observed at the Soret band. Accounting for the high molar absorption coefficient of the porphyrin core, an uncaging cross-section ($\Phi_{\text{CO}}\epsilon_{\text{max}}$) of 5600 M⁻¹ cm⁻¹ at $\lambda_{\text{max}} = 421 \text{ nm}$ is extraordinarily high, one of the highest values reported to date for a visible-light-activatable photoCORM.¹⁰ An uncaging cross-section of 70 M⁻¹ cm⁻¹ was found at 645 nm (see all Φ_{CO} and $\Phi_{\text{CO}}\epsilon_{\text{max}}$ values in Table S6).

The ability of porphyrins to incorporate metal cations gave us a tool to control and tailor the photochemical properties of the hybrids for a given application. We optimized a general and facile procedure for incorporating metal cations by reacting **2** with the corresponding metal triflates in methanol or DMF. Analytically pure metal complexes **2-Zn** and **2-Cu** were obtained by precipitation from a crude reaction mixture by slow addition of CH₂Cl₂. Upon metal insertion, the absorption spectra were only slightly affected, possessing molar absorption coefficients lower by 20–50% compared to that of **2**. The emission properties are more influenced as Φ_{f} decreased for **2-Zn** by a factor of 2, while no emission was detected for **2-Cu**. We also examined whether these hybrids do not form flavonolato complexes among individual flavonol moieties

and zinc or copper cations. The interpretation of the absorption, ¹H NMR, and MS spectra confirmed that the flavonol group is not complexed (e.g., Figures S39 and S68).

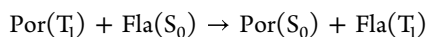
The incorporation of metals significantly impacted the system's ability to release CO (Figure 1D). Irradiation of **2-Zn** in methanol at 440 or 600 nm led to the efficient disappearance of absorption bands of both the flavonol and porphyrin moieties. A higher Φ_{CO} of 0.040 ± 0.002 (~2.2 times higher than that of **2**) resulted in an extraordinarily high uncaging cross-section of 10 230 M⁻¹ cm⁻¹ at λ_{max} and 380 M⁻¹ cm⁻¹ at 601 nm. The yield of released CO was 2.3 ± 0.1 equiv in methanol and 2.1 ± 0.1 equiv in PBS. In contrast, **2-Cu** showed reduced photoreactivity with a much less efficient CO release ($\Phi_{\text{CO}} = 0.002 \pm 0.001$) and a lower maximum chemical yield of 0.8 ± 0.1 equiv upon exhaustive irradiation at 440 nm. In our recent study, the porphyrin-Cu(II) derivatives bearing a leaving group in the *meso*-methyl position were found to be photochemically inactive due to an ultrafast internal conversion observed as the main deactivation pathway.²⁹

The products of the flavonol moiety degradation formed upon irradiation of **3** at 440 nm were identified as the anticipated esters of salicylic acid (MALDI-TOF; Figures S12 and S13). There was an evident sequential photodecarbonylation in all four flavonol units, but we were not able to assign the exact structures of products formed from porphyrin ring opening by MALDI-TOF and ESI-TOF. This degradation pathway, which is not relevant in the case of bimolecular flavonol sensitization mentioned above, is probably responsible for CO yields lower than 4 equiv in the studied hybrids (Table 1).

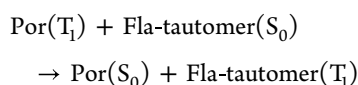
The essential role of oxygen²⁰ in the solution was confirmed by irradiating **3** in a degassed methanol/DMSO solution (4:1, v/v). The photodecomposition of the flavonol groups (detected at 355 nm) was 15 times slower, and only traces of CO were detected, even after prolonged irradiation (Figures S9 and S10).

We examined the role of ¹O₂ produced by porphyrin sensitization, but we did not observe any correlation between the Φ_{CO} and Φ_{Δ} values (Table 1). In addition, the CO production from **3** was not affected in the presence of a singlet oxygen trap (NaN₃ or α -terpinene, 5 mM; Table S7); thus, ¹O₂ is not directly involved in the reaction. On the other hand, the presence of a triplet quencher (cyclooctatetraene or 4-nitrobenzyl alcohol, 10 mM) reduced the CO release efficiency by ~20% (Table S8), indicating a productive triplet-excited state.

To examine the CO release mechanism from the hybrids, we first ruled out possible charge transfer from the triplet porphyrin to the flavonol as the initial state of the process (Supporting Information). We then hypothesized that triplet-excited porphyrin (Por) sensitizes flavonol (Fla) according to

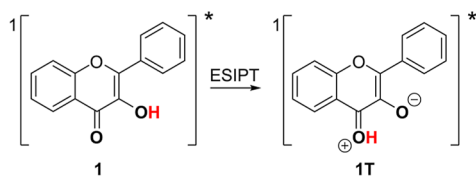


However, according to our calculations, this process is endergonic with a Gibbs free energy of 0.48 eV (calculated at the B3LYP/6-31+g* level; Scheme S2). This conclusion is also supported by the experimentally measured energetics: the triplet energies, E_T , of porphyrin and Zn-porphyrin, 1.44 and 1.60 eV, respectively,³⁹ are lower than that of flavonol, 1.79 eV.⁴⁰ The process is thus energetically forbidden. However, it would be exergonic with a Gibbs free energy of -0.40 eV (see Scheme S2) if the acceptor were flavonol in its tautomeric form (experimental $E_T = 0.99$ eV,⁴⁰) formed upon intramolecular proton transfer from the 3-OH group to the carbonyl group (tautomerization)

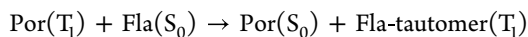


Such a tautomer is known to be generated rapidly upon excitation to the singlet excited state (<125 fs) via excited-state intramolecular proton transfer (ESIPT) to give a Stokes-shifted emitting phototautomer 1T (Scheme 3).^{41,42}

Scheme 3. Phototautomerization of Flavonol



Considering the ground-state flavonol **1** as the initial state and tautomer **1T** in the triplet state T_1 as the final state according to



the Gibbs free energy of the TEnT process decreased to 0.06 eV (see Scheme S2) as a consequence of the flavonol energy stabilization upon tautomerization in the triplet state (see Figure S1). This intermolecular energy transfer should thus be governed by the flavonol ESIPT either stepwise or simultaneously.

Electronic transitions in many critical processes are coupled to a nuclear motion, with proton-coupled electron transfer (PCET) being the most prominent example.⁴³ PCET plays an important role in many biological and chemical processes, such as proton-coupled redox processes occurring during tyrosine oxidation and reduction in photosystem II,⁴⁴ redox reactions of tyrosine in biological energy transduction, charge transport or enzymatic catalysis,⁴⁵ or in organic synthesis.⁴⁶ At higher energies, the nonlocal Auger-Meitner processes are mediated by proton transfer in the intermediate state.⁴⁷

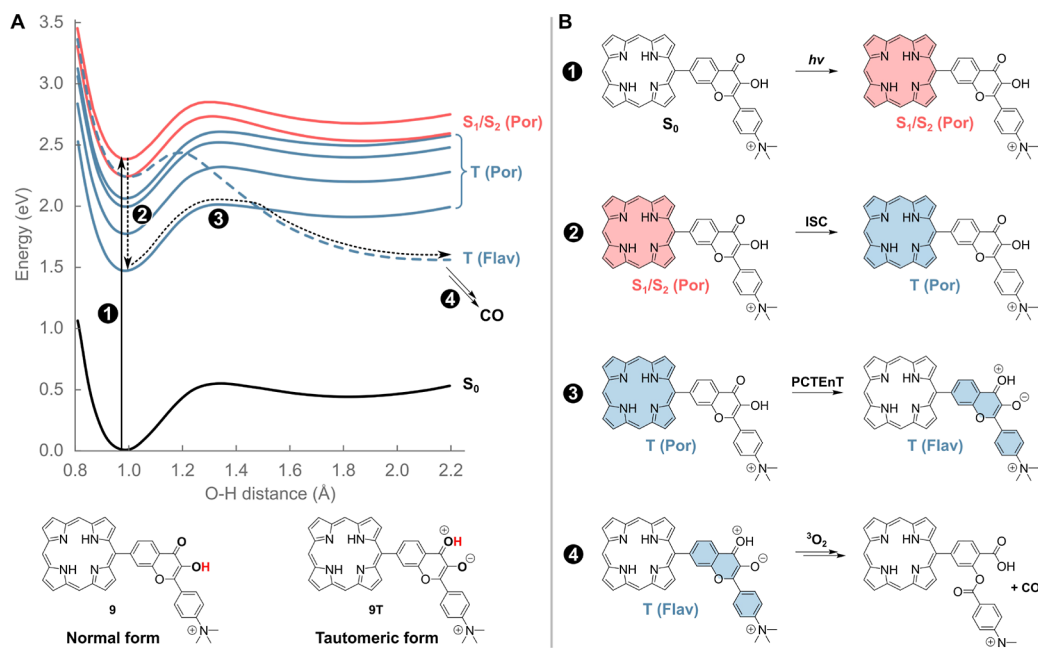
In 2019, Hammarström, Mayer, Hammes-Schiffer, and co-workers provided experimental evidence of the existence of inverted region behavior for concerted proton–electron transfer.⁴⁸ The same laboratories very recently reported a unique mechanism in anthracene–phenol–pyridine triads, in

which energy transfer is coupled to a nuclear motion, termed proton-coupled energy transfer (PCEnT).⁴⁹ In this process, the formation of a locally excited state on anthracene is followed by singlet–singlet energy transfer to the phenol–pyridine chromophore. Despite the lack of spectral overlap of the donor emission and acceptor absorption spectra required for the Förster resonance (dipole–dipole) mechanism,^{50,51} the process was made possible by coupling proton transfer in the phenol–pyridine unit with energy transfer that lowered its excited-state energy.

The question is whether such a mechanism would also be operational in the triplet electronic manifold as most antenna systems absorbing visible/NIR light exhibit a relatively efficient intersystem crossing (ISC) to produce longer-lived triplet states. Triplet energy transfer (TEnT) by electron exchange is a spin-allowed process in which the triplet energy of the donor should exceed that of the acceptor and requires appreciable overlap of their molecular orbitals.⁵²

In this work, we investigated the possibility that the mechanism of flavonol sensitization occurs through a simultaneous proton-coupled TEnT (PCTEnT). We conducted quantum-chemical calculations on model compound **9** (Scheme 4). Upon excitation into one of the absorption bands, a locally excited singlet state of porphyrin, corresponding to the S_1 and S_2 electronic states, is formed, while the flavonol unit remains in its nonproductive ground state (step 1, Scheme 4B). The singlet excitation localized on the flavonol chromophore and a mixed excitation between the porphyrin and flavonol units were found to be above the S_1 and S_2 energy levels (Table S3); thus, they do not interfere with the overall process. Analogous to the isolated porphyrin, efficient ISC to the triplet manifold and subsequent IC to the lowest triplet state²⁹ follow (step 2, Scheme 4). The triplet state with the excitation on the flavonol units is located ~0.8 eV above the lowest triplet state of porphyrin, and this electronic state is still not populated (note: electronic interactions between the two units are limited; the electronic states can be safely associated with excitations localized on the two different units; see Supporting Information and Figure S1C). Similar to the calculations on isolated flavonol, **1** (Figure S1A), tautomerization of flavonol via intramolecular ESIPT (step 3, Scheme 4) is the critical step in the transformation to produce tautomer **9T** with the triplet excitation localized on the flavonol unit, thus via the PCTEnT mechanism. Note that in the ground and first four triplet states of **9**, the potential energy curves follow that of the flavonol singlet state, and the energy increases upon proton transfer, whereas a higher flavonol triplet state (T_5) possesses a minimum for the excited flavonol-tautomer. Therefore, simultaneous proton and energy transfers should occur in the lowest triplet state—the proton transfer mediates the energy transfer taking place at the crossing of the two diabatic states (with the triplet excitation localized either on the flavonol or porphyrin moieties). The tautomerization barrier in the triplet state (the calculated value of 0.53 eV in protic methanol; Figure S3 and Table S4) allows the process to occur within the triplet-state lifetime. The relative energies of both tautomers in the excited (triplet) state are within the thermal energy, with the Gibbs free energy of the tautomeric form being higher by only 0.16 eV (Table S5). The triplet-excited flavonol unit can then undergo the reaction with triplet oxygen to release CO^{24,25} (step 4, Scheme 4); the triplet state of flavonol is more acidic than that in the ground state, yet it should not deprotonate in methanol (see the Supporting

Scheme 4. Proton-Coupled Triplet–Triplet Energy Transfer. (A) Potential Energy Curves and (B) the Proposed Mechanism of CO Photorelease from a Porphyrin (Por)-Flavonol (Flav) Model Hybrid 9 via Four Reaction Steps^a



^aThe potential energies are calculated at the TDDFT/B3LYP/6-31g* level. The red and blue colors correspond to the singlet (S) and triplet (T) excited states, respectively

Information). The amount of 9T is possibly very small because it is formed slowly across the barrier, while it is consumed in the rapid reaction with triplet oxygen. The whole process can thus be entropically driven despite the slightly endothermic nature of the PCTEnT process. The whole system is similar to other endothermic photochemical processes.^{53–55}

Directly confirming the PCTEnT process requires spectroscopic characterization of the flavonol tautomer in the triplet-excited porphyrin–flavonol complex. This is, however, very difficult, as the triplet tautomer is a reactive intermediate that undergoes a comparatively rapid decay, and its concentration must be very low. Furthermore, the TA signal of flavonol 1 appears in the 390–430 nm region (Figure S17 and the literature⁴³); thus, it is located within the region of porphyrin ground-state bleaching, making its direct detection difficult due to the strong signal overlap.

Instead, we aimed to study triplet–triplet energy transfer by comparing the triplet lifetimes of hybrid 3 and model porphyrins under different conditions. Femtosecond (fs) TA spectra of 3 in DMSO (Figure S14) showed four main species upon excitation at 387 nm (at this wavelength, both flavonol moieties and the porphyrin core absorb). The first three components are assigned^{39,40} to an ultrafast internal conversion from S₂ to S₁ (0.2 ps) and vibrational relaxation within the Q bands (0.8 and 9.9 ps). The final component is associated^{41,42} with singlet–triplet ISC ($k_{\text{ISC}} = 4.1 \times 10^8 \text{ s}^{-1}$). We performed TA experiments with 3 and tetra-4-ethynylphenylporphyrin (TEPP) in THF to selectively excite the porphyrin core at 520 nm. TA spectra in the 300–1050 nm region highlight a fast and a slow component in both compounds (Figure S15). The faster component, similar for both 3 and TEPP (8.6 and 11.8 ns, respectively, and not affected by the presence of oxygen), is assigned to singlet–triplet ISC. The slower component was sensitive to the oxygen content, with lifetimes 3 orders of magnitude longer in

degassed conditions. We assigned it to the triplet state. The triplet lifetime of hybrid 3 in degassed solutions (168 μs) was found to be shorter than that of TEPP by a factor of 2 (381 μs). This may suggest that a quenching process is operative. However, these data cannot be directly compared because the model porphyrin is not an exact model of the porphyrin core in 3.

Nanosecond (ns) TA spectra in degassed THF were compared for 3 and tetraphenylporphyrin (TPP) upon excitation at 532 nm (Figure S16). The spectra of 3 revealed an intense band in the 300–600 nm region, split into two bands by the ground state bleach at 425 nm. Both bands decayed with the same lifetime of 90 μs and are assigned to the triplet state⁴³ ($\tau = 292 \text{ ns}$ in an aerated solution). nsTA spectra of TPP under the same conditions lacked the prominent TA band in the UV region (flavonol) and showed a longer triplet lifetime (120 μs) again.

Finally, another indirect evidence of the proposed TEnT mechanism was provided by Stern Volmer analysis of the ZnTcPP triplet decay at different triplet quencher 1 concentrations. A bimolecular quenching rate k_q of $\sim 2 \times 10^5 \text{ M}^{-1} \text{ s}^{-1}$ (Figure S18) was found, and it correlates well with the predicted, slightly endergonic Gibbs energy of the TEnT process (0.06 eV), calculated for the flavonol energy stabilization upon tautomerization in the triplet state.

Porphyrins are well-known for their ability to produce singlet oxygen (¹O₂) by ground-state oxygen (³O₂) sensitization. A simultaneous presence of CO and ¹O₂ can be responsible for the synergistic enhancement of cytotoxicity of some PDT agents against cancer cells.⁵⁶ We measured the efficiency of ¹O₂ production (Φ_{Δ}) for all of the derivatives using the luminescence signal of ¹O₂. Φ_{Δ} of the metal-free hybrid 2 was found to be 0.17 in methanol. Metal insertion drastically reduced Φ_{Δ} to 0.10 for 2-Zn and $<10^{-3}$ for 2-Cu.

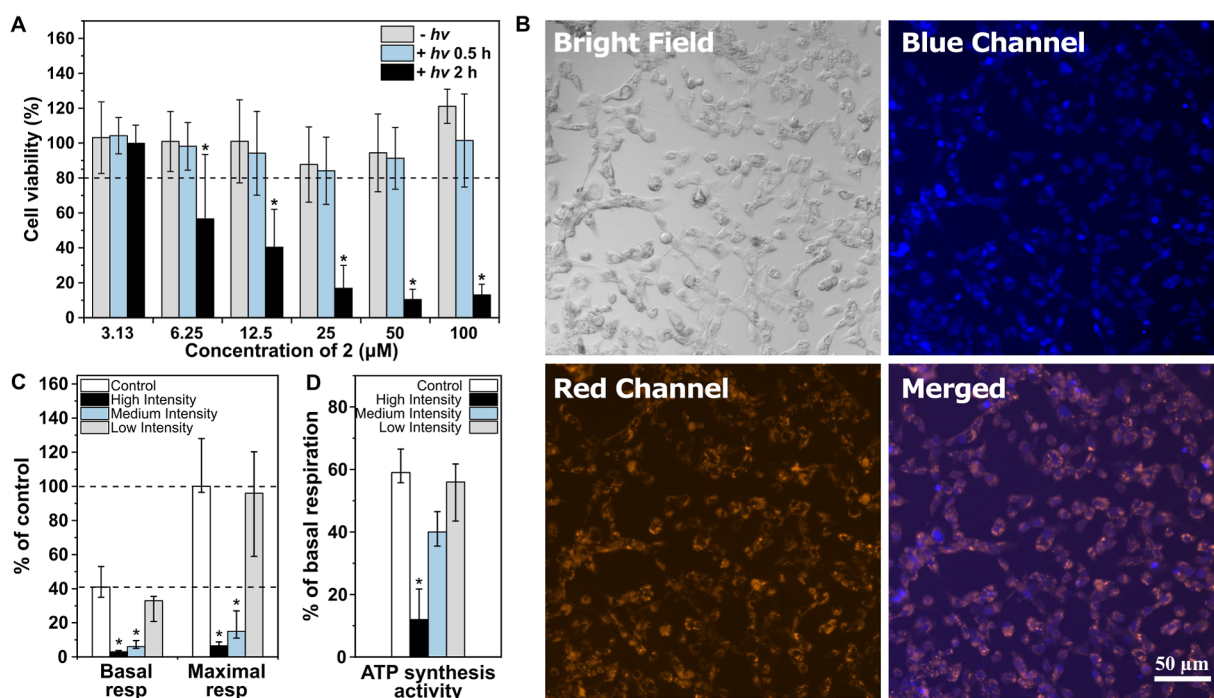


Figure 2. Biological experiments. (A) Effect of **2** on the viability of HepG2 cells after 24 h. Cytotoxicity was assessed by an MTT test after treating HepG2 cells with solutions of **2** in colorless MEM (1% DMSO) for 24 h without ($-hv$) or with ($+hv$) for 0.5 or 2 h of irradiation using white light (120 mW cm^{-2}). The values are expressed as % of untreated controls. * P -value ≤ 0.05 vs untreated control; $n \geq 8$. (B) Fluorescence microscopy of HepaG cells incubated with **2** ($100 \mu\text{M}$, 24 h). After incubation, the cells were washed with PBS and visualized by using white light (bright field) and a WIG filter (red channel). Nuclei were counterstained with DAPI and observed under UV light (blue channel). (C,D) Effect of irradiation intensity on the respiration of HepG2 cells. HepG2 cells were treated with **2** ($25 \mu\text{M}$) and irradiated with high-, medium-, or low-intensity white light (990 , 120 , or 8 mW cm^{-2} , respectively) for 30 min; basal and maximum respiration (the values are expressed as % of the maximum respiration level of untreated controls) and ATP synthesis activities were analyzed immediately after incubation; ATP synthesis activity was calculated as the ratio of basal respiration to respiration after oligomycin inhibition and expressed as % of the basal respiration level of the untreated controls. * P -value ≤ 0.05 ; $n \geq 4$.

We also evaluated the effect of compound **2** on cell viability. The human hepatoblastoma HepG2 cells were incubated with **2** within the concentration range of 1.5 – $100 \mu\text{M}$. No significant toxicity was observed after **2** (Figure S20) and 24 h (Figure 2A) incubation in the dark. Stable photoproducts obtained upon prolonged irradiation of a solution of **2** did not affect cell viability up to the concentration of $200 \mu\text{M}$ following 24 h incubation (Figure S21). Similarly, no toxicity of **2** was observed upon irradiation of cells for 30 min, but after 2 h, it resulted in a significant decrease in cell viability at the concentration of $6.25 \mu\text{M}$ and higher (Figure 2A). We attribute the observed delayed cytotoxicity to $^1\text{O}_2$ produced, thanks to oxygen photosensitization by primary photoproducts formed upon prolonged irradiation, as discussed above.

Mitochondria are an important target for CO as this gaseous signaling molecule can alter mitochondrial respiration and cytochrome *c* oxygenase activity or ROS production.⁵⁷ Whether these effects are cytotoxic or cytoprotective depends mainly on the CO concentration.⁵⁸ In our experimental system, HepG2 cells were incubated with **2** at a concentration of $25 \mu\text{M}$ and irradiated at different intensities of white light. It resulted in the CO release into the medium (Figures S22 and S23), which was manifested by differential effects on mitochondrial functions. The highest CO concentrations formed at high irradiation intensities strongly inhibited mitochondrial basal and maximum respiratory activities and ATP synthesis (Figure 2C,D). On the contrary, almost no effect on mitochondrial function was observed using low-

intensity irradiation related to low CO concentrations. This demonstrates the high tuneability and adjustability of our light-triggered CO release systems, allowing for better CO release control with a predictable biological outcome. No effect on cellular respiration was observed in cells exposed to **2** in the dark or upon irradiation of solutions without the active substances (Figure S24). Even though CO can freely pass cellular membranes, some of its biological effects may depend on whether it is released intra- or extracellularly.⁵⁹ Nevertheless, our fluorescence microscopy studies clearly showed the cytoplasmic localization of **2** (Figures 2B and S25) in human hepatic progenitor HeparG cells 24 h after incubation.

CONCLUSIONS

In this work, we developed efficient photoactivatable carbon monoxide donors (photoCORMs) consisting of a porphyrin antenna and a covalently bound flavonol moiety as an energy acceptor. They absorb throughout the whole visible part of the spectrum, including the tissue-transparent window. Incorporating a metal cation into the porphyrin moiety can modulate the photochemical properties. With four CO flavonol donors attached to a single porphyrin chromophore, excellent CO release yields and uncaging cross sections, and in vitro biocompatibility, these photoCORMs have high potential for their further development and future biological and medical applications. Their advantages include high CO yields (atom economy), intracellular CO delivery, tuneability of CO release, and combining biological effects of CO with those of the

porphyrin moiety. In this regard, it should be noted that photodynamic therapy using porphyrins⁶⁰ has been established as an effective anticancer treatment for specific human tumors, and the same applies to the anticancer effects of CO.⁶¹

We hypothesized that the mechanism of CO release is based on proton-coupled triplet energy transfer (PCEnT) between the excited porphyrin and flavonol tautomer, analogous to the recently discovered process of proton-coupled singlet energy transfer.⁴⁹ The viability of this mechanism was supported by our computational studies, but despite our efforts, none of our spectroscopic investigations provided direct evidence. Because the key intermediate (triplet-excited flavonol tautomer) formed during PCEnT is predicted to be energetically above the locally excited state, it may appear only at very low concentrations that time-resolved methods cannot detect. We will try to solve this problem in our future projects for which we will design new systems with longer-lived transients.

■ ASSOCIATED CONTENT

SI Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/jacs.3c11426>.

Materials and methods; synthesis; NMR, absorption, emission, and time-resolved spectroscopy; quantum-chemical calculations; and biological experiments (PDF)

■ AUTHOR INFORMATION

Corresponding Authors

Petr Slavíček – Department of Physical Chemistry, University of Chemistry and Technology, 16628 Prague 6, Czech Republic; orcid.org/0000-0002-5358-5538; Email: Petr.Slavicek@vscht.cz

Petr Klán – Department of Chemistry, Faculty of Science, Masaryk University, 62500 Brno, Czech Republic; RECETOX, Faculty of Science, Masaryk University, 62500 Brno, Czech Republic; orcid.org/0000-0001-6287-2742; Email: klan@sci.muni.cz

Authors

Andrea Ramundo – Department of Chemistry, Faculty of Science, Masaryk University, 62500 Brno, Czech Republic; RECETOX, Faculty of Science, Masaryk University, 62500 Brno, Czech Republic; orcid.org/0000-0002-1249-4840

Jiří Janoš – Department of Physical Chemistry, University of Chemistry and Technology, 16628 Prague 6, Czech Republic; orcid.org/0000-0001-5903-8538

Lucie Muchová – Institute of Medical Biochemistry and Laboratory Diagnostics, and 4th Department of Internal Medicine, General University Hospital in Prague and First Faculty of Medicine, Charles University, 12108 Prague 2, Czech Republic

Mária Šranková – Institute of Medical Biochemistry and Laboratory Diagnostics, and 4th Department of Internal Medicine, General University Hospital in Prague and First Faculty of Medicine, Charles University, 12108 Prague 2, Czech Republic

Jakub Dostál – ELI Beamlines Facility, The Extreme Light Infrastructure ERIC, 25241 Dolní Břežany, Czech Republic

Miroslav Kloz – ELI Beamlines Facility, The Extreme Light Infrastructure ERIC, 25241 Dolní Břežany, Czech Republic

Libor Vitek – Institute of Medical Biochemistry and Laboratory Diagnostics, and 4th Department of Internal

Medicine, General University Hospital in Prague and First Faculty of Medicine, Charles University, 12108 Prague 2, Czech Republic; orcid.org/0000-0002-5318-0151

Complete contact information is available at: <https://pubs.acs.org/doi/10.1021/jacs.3c11426>

Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

P. K. thanks the Czech Science Foundation (GA21-01799S) and the RECETOX Research Infrastructure (no. LM2018121) financed by the Ministry of Education, Youth and Sports, and the Operational Programme Research, Development, and Education (the CETOCOEN EXCELLENCE project no. CZ.02.1.01/0.0/0.0/17_043/0009632) for supportive background. This project was also supported by the European Union's Horizon 2020 Research and Innovation Programme under grant agreement no. 857560 (P. K.) and by Charles University (GAUK 314621; M.Š.). J.J. and P.S. thank the Czech Science Foundation for the financial support (Grant no. 23-07066S). J. J. was additionally supported by the grant of Specific university research—grant No A1\FCHI_2023_001. We acknowledge Proteomics of CIISB, Instruct-CZ Centre, supported by MEYS CR (LM2018127) for MALDI-TOF/TOF measurements and National Infrastructure for Chemical Biology (CZ-OPENSREEN, LM2023052) for NMR measurements. This research was also supported by the Czech Science Foundation (21-09692M and 21-05180S, M. K.). We thank Dominik Madea, Lubos Jilek, and Marek Martinek (Masaryk University) for their help with time-resolved experiments and Josef Filgas (University of Chemistry and Technology) for discussing the theoretical results.

■ REFERENCES

- (1) Romão, C. C.; Blättler, W. A.; Seixas, J. D.; Bernardes, G. J. Developing drug molecules for therapy with carbon monoxide. *Chem. Soc. Rev.* **2012**, *41*, 3571–3583.
- (2) Lu, W.; Yang, X.; Wang, B. Carbon monoxide signaling and soluble guanylyl cyclase: Facts, myths, and intriguing possibilities. *Biochem. Pharmacol.* **2022**, *200*, 115041.
- (3) Wang, B.; Otterbein, L. E. *Carbon Monoxide in Drug Discovery: Basics, Pharmacology, and Therapeutic Potential*; John Wiley & Sons: Hoboken, NJ, USA, 2022.
- (4) Ling, K.; Men, F.; Wang, W.-C.; Zhou, Y.-Q.; Zhang, H.-W.; Ye, D.-W. Carbon monoxide and its controlled release: therapeutic application, detection, and development of carbon monoxide releasing molecules (CORMs) miniperspective. *J. Med. Chem.* **2018**, *61*, 2611–2635.
- (5) Motterlini, R.; Otterbein, L. E. The therapeutic potential of carbon monoxide. *Nat. Rev. Drug Discovery* **2010**, *9*, 728–743.
- (6) Vitek, L.; Gbelcová, H.; Muchová, L.; Váňová, K.; Zelenka, J.; Koničková, R.; Šuk, J.; Zadinova, M.; Knejzlík, Z.; Ahmad, S.; et al. Antiproliferative effects of carbon monoxide on pancreatic cancer. *Dig. Liver Dis.* **2014**, *46*, 369–375.
- (7) Vander Heiden, M. G.; Cantley, L. C.; Thompson, C. B. Understanding the Warburg effect: the metabolic requirements of cell proliferation. *Science* **2009**, *324*, 1029–1033.
- (8) Wegiel, B.; Gallo, D.; Csizmadia, E.; Harris, C.; Belcher, J.; Vercellotti, G. M.; Penacho, N.; Seth, P.; Sukhatme, V.; Ahmed, A.; et al. Carbon Monoxide Expedites Metabolic Exhaustion to Inhibit Tumor Growth. *Cancer Res.* **2013**, *73*, 7009–7021.
- (9) Pinto, M. N.; Mascharak, P. K. Light-assisted and remote delivery of carbon monoxide to malignant cells and tissues:

Photochemotherapy in the spotlight. *J. Photochem. Photobiol., C* **2020**, *42*, 100341.

(10) Weinstain, R.; Slanina, T.; Kand, D.; Klan, P. Visible-to-NIR-light activated release: from small molecules to nanomaterials. *Chem. Rev.* **2020**, *120*, 13135–13272.

(11) Allison, R. R.; Sibata, C. H. Oncologic photodynamic therapy photosensitizers: a clinical review. *Photodiagn. Photodyn. Ther.* **2010**, *7*, 61–75.

(12) Carrington, S. J.; Chakraborty, I.; Bernard, J. M.; Mascharak, P. K. A theranostic two-tone luminescent PhotoCORM derived from Re (I) and (2-pyridyl)-benzothiazole: trackable CO delivery to malignant cells. *Inorg. Chem.* **2016**, *55*, 7852–7858.

(13) Martinek, M.; Filipova, L.; Galeta, J.; Ludvíková, L.; Klan, P. Photochemical formation of dibenzosilacyclohept-4-yne for Cu-free click chemistry with azides and 1, 2, 4, 5-Tetrazines. *Org. Lett.* **2016**, *18*, 4892–4895.

(14) Poloukhtine, A. A.; Mbua, N. E.; Wolfert, M. A.; Boons, G.-J.; Popik, V. V. Selective labeling of living cells by a photo-triggered click reaction. *J. Am. Chem. Soc.* **2009**, *131*, 15769–15776.

(15) Chapman, O.; Wojtkowski, P.; Adam, W.; Rodriguez, O.; Rucktaeschel, R. Photochemical transformations. XLIV. Cyclic peroxides. Synthesis and chemistry of α -lactones. *J. Am. Chem. Soc.* **1972**, *94*, 1365–1367.

(16) Peng, P.; Wang, C.; Shi, Z.; Johns, V. K.; Ma, L.; Oyer, J.; Copik, A.; Igarashi, R.; Liao, Y. Visible-light activatable organic CO-releasing molecules (PhotoCORMs) that simultaneously generate fluorophores. *Org. Biomol. Chem.* **2013**, *11*, 6671–6674.

(17) Antony, L. A. P.; Slanina, T.; Sebej, P.; Solomek, T.; Klan, P. Fluorescein analogue xanthen-9-carboxylic acid: a transition-metal-free CO releasing molecule activated by green light. *Org. Lett.* **2013**, *15*, 4552–4555.

(18) Palao, E.; Slanina, T.; Muchová, L.; Solomek, T.; Vitek, L.; Klan, P. Transition-metal-free CO-releasing BODIPY derivatives activatable by visible to NIR light as promising bioactive molecules. *J. Am. Chem. Soc.* **2016**, *138*, 126–133.

(19) Pietta, P.-G. Flavonoids as antioxidants. *J. Nat. Prod.* **2000**, *63*, 1035–1042.

(20) Studer, S. L.; Brewer, W. E.; Martinez, M. L.; Chou, P. T. Time-resolved study of the photooxygenation of 3-hydroxyflavone. *J. Am. Chem. Soc.* **1989**, *111*, 7643–7644.

(21) Chou, P.-T.; Martinez, M. L. Photooxygenation of 3-hydroxyflavone and molecular design of the radiation-hard scintillator based on the excited-state proton transfer. *Radiat. Phys. Chem.* **1993**, *41*, 373–378.

(22) Anderson, S. N.; Richards, J. M.; Esquer, H. J.; Benninghoff, A. D.; Arif, A. M.; Berreau, L. M. A structurally-tunable 3-hydroxyflavone motif for visible light-induced carbon monoxide-releasing molecules (CORMs). *ChemistryOpen* **2015**, *4*, 590–594.

(23) Lazarus, L. S.; Esquer, H. J.; Benninghoff, A. D.; Berreau, L. M. Sense and release: a thiol-responsive flavonol-based photonically driven carbon monoxide-releasing molecule that operates via a multiple-input AND logic gate. *J. Am. Chem. Soc.* **2017**, *139*, 9435–9438.

(24) Russo, M.; Orel, V.; Stacko, P.; Srankova, M.; Muchova, L.; Vitek, L.; Klan, P. Structure-Photoreactivity Relationship of 3-Hydroxyflavone-Based CO-Releasing Molecules. *J. Org. Chem.* **2022**, *87*, 4750–4763.

(25) Russo, M.; Stacko, P.; Nachtigalova, D.; Klan, P. Mechanisms of orthogonal photodecarbonylation reactions of 3-hydroxyflavone-based acid-base forms. *J. Org. Chem.* **2020**, *85*, 3527–3537.

(26) Stackova, L.; Russo, M.; Muchova, L.; Orel, V.; Vitek, L.; Stacko, P.; Klan, P. Cyanine-Flavonol Hybrids for Near-Infrared Light-Activated Delivery of Carbon Monoxide. *Chem.—Eur. J.* **2020**, *26*, 13184–13190.

(27) Yang, Q.; Muchova, L.; Stackova, L.; Stacko, P.; Šindelář, V.; Vitek, L.; Klan, P. Cyanine-flavonol hybrids as NIR-light activatable carbon monoxide donors in methanol and aqueous solutions. *Chem. Commun.* **2022**, *58*, 8958–8961.

(28) Cheng, J.; Gan, G.; Shen, Z.; Gao, L.; Zhang, G.; Hu, J. Red Light-Triggered Intracellular Carbon Monoxide Release Enables Selective Eradication of MRSA Infection. *Angew. Chem.* **2021**, *133*, 13625–13632.

(29) Sekhar, A. R.; Chitose, Y.; Janos, J.; Dangoor, S. I.; Ramundo, A.; Satchi-Fainaro, R.; Slavicek, P.; Klan, P.; Weinstain, R. Porphyrin as a versatile visible-light-activatable organic/metal hybrid photo-removable protecting group. *Nat. Commun.* **2022**, *13*, 3614.

(30) Alabugin, A. Near-IR Photochemistry for Biology: Exploiting the Optical Window of Tissue. *Photochem. Photobiol.* **2019**, *95*, 722–732.

(31) Gouterman, M.; Wagnière, G. H.; Snyder, L. C. Spectra of porphyrins: Part II. Four orbital model. *J. Mol. Spectrosc.* **1963**, *11*, 108–127.

(32) Mandal, A. K.; Taniguchi, M.; Diers, J. R.; Niedzwiedzki, D. M.; Kirmaier, C.; Lindsey, J. S.; Bocian, D. F.; Holten, D. Photophysical properties and electronic structure of porphyrins bearing zero to four meso-phenyl substituents: New insights into seemingly well understood tetrapyrroles. *J. Phys. Chem. A* **2016**, *120*, 9719–9731.

(33) Dąbrowski, J. M.; Pucelik, B.; Regiel-Futyr, A.; Brindell, M.; Mazuryk, O.; Kyzioł, A.; Stochel, G.; Macyk, W.; Arnaut, L. G. Engineering of relevant photodynamic processes through structural modifications of metallotetrapyrrolic photosensitizers. *Coord. Chem. Rev.* **2016**, *325*, 67–101.

(34) Dávila, Y. A.; Sancho, M. I.; Almandoz, M. C.; Blanco, S. E. Solvent Effects on the Dissociation Constants of Hydroxyflavones in Organic-Water Mixtures. Determination of the Thermodynamic pKa Values by UV-Visible Spectroscopy and DFT Calculations. *J. Chem. Eng. Data* **2013**, *58*, 1706–1716.

(35) Avani, R.; Busa, K. P.; Mahetar, J. G.; Shah, M. K. A facile synthetic approach for the syntheses of 7-hydroxyflavonol derivatives. *Der Pharma Chem.* **2015**, *7*, 142–146.

(36) Lindsey, J. S.; Hsu, H. C.; Schreiman, I. C. Synthesis of tetraphenylporphyrins under very mild conditions. *Tetrahedron Lett.* **1986**, *27*, 4969–4970.

(37) Gouterman, M. Spectra of porphyrins. *J. Mol. Spectrosc.* **1961**, *6*, 138–163.

(38) Cavaleiro, J. A. S.; Neves, M. G. P. S.; Hewlins, M. J. E.; Jackson, A. H. The photo-oxidation of meso-tetraphenylporphyrins. *J. Chem. Soc., Perkin Trans. 1* **1990**, 1937–1943.

(39) Kalyanasundaram, K.; Neumann-Spallart, M. Photophysical and redox properties of water-soluble porphyrins in aqueous media. *J. Phys. Chem.* **1982**, *86*, 5163–5169.

(40) Dick, B. AM1 and INDO/S calculations on electronic singlet and triplet states involved in excited-state intramolecular proton transfer of 3-hydroxyflavone. *J. Phys. Chem.* **1990**, *94*, 5752–5756.

(41) Sengupta, P. K.; Kasha, M. Excited state proton-transfer spectroscopy of 3-hydroxyflavone and quercetin. *Chem. Phys. Lett.* **1979**, *68*, 382–385.

(42) Schwartz, B. J.; Peteanu, L. A.; Harris, C. B. Direct observation of fast proton transfer: femtosecond photophysics of 3-hydroxyflavone. *J. Phys. Chem.* **1992**, *96*, 3591–3598.

(43) Hammes-Schiffer, S.; Stuchebrukhov, A. A. Theory of Coupled Electron and Proton Transfer Reactions. *Chem. Rev.* **2010**, *110*, 6939–6960.

(44) Migliore, A.; Polizzi, N. F.; Therien, M. J.; Beratan, D. N. Biochemistry and Theory of Proton-Coupled Electron Transfer. *Chem. Rev.* **2014**, *114*, 3381–3465.

(45) Warren, J. J.; Tronic, T. A.; Mayer, J. M. Thermochemistry of Proton-Coupled Electron Transfer Reagents and its Implications. *Chem. Rev.* **2010**, *110*, 6961–7001.

(46) Murray, P. R. D.; Cox, J. H.; Chiappini, N. D.; Roos, C. B.; McLoughlin, E. A.; Hejna, B. G.; Nguyen, S. T.; Ripberger, H. H.; Ganley, J. M.; Tsui, E.; Shin, N. Y.; Koronkiewicz, B.; Qiu, G.; Knowles, R. R. Photochemical and Electrochemical Applications of Proton-Coupled Electron Transfer in Organic Synthesis. *Chem. Rev.* **2022**, *122*, 2017–2291.

(47) Amoretti, M.; Amsler, C.; Bonomi, G.; Bouchta, A.; Bowe, P.; Carraro, C.; Cesar, C. L.; Charlton, M.; Collier, M. J. T.; Doser, M.;

Filippini, V.; Fine, K. S.; Fontana, A.; Fujiwara, M. C.; Funakoshi, R.; Genova, P.; Hangst, J. S.; Hayano, R. S.; Holzscheiter, M. H.; Jørgensen, L. V.; Lagomarsino, V.; Landua, R.; Lindelöf, D.; Rizzini, E. L.; Macri, M.; Madsen, N.; Manuzio, G.; Marchesotti, M.; Montagna, P.; Pruys, H.; Regenfus, C.; Riedler, P.; Rochet, J.; Rotondi, A.; Rouleau, G.; Testera, G.; Variola, A.; Watson, T. L.; van der Werf, D. P. Production and detection of cold antihydrogen atoms. *Nature* **2002**, *419*, 456–459.

(48) Parada, G. A.; Goldsmith, Z. K.; Kolmar, S.; Pettersson Rimgard, B.; Mercado, B. Q.; Hammarström, L.; Hammes-Schiffer, S.; Mayer, J. M. Concerted proton-electron transfer reactions in the Marcus inverted region. *Science* **2019**, *364*, 471–475.

(49) Pettersson Rimgard, B.; Tao, Z.; Parada, G. A.; Cotter, L. F.; Hammes-Schiffer, S.; Mayer, J. M.; Hammarström, L. Proton-coupled energy transfer in molecular triads. *Science* **2022**, *377*, 742–747.

(50) Wu, P. G.; Brand, L. Resonance Energy Transfer: Methods and Applications. *Anal. Biochem.* **1994**, *218*, 1–13.

(51) Fang, B.; Shen, Y.; Peng, B.; Bai, H.; Wang, L.; Zhang, J.; Hu, W.; Fu, L.; Zhang, W.; Li, L.; Huang, W. Small-Molecule Quenchers for Förster Resonance Energy Transfer: Structure, Mechanism, and Applications. *Angew. Chem., Int. Ed.* **2022**, *61*, No. e202207188.

(52) Klan, P.; Wirz, J. *Photochemistry of Organic Compounds: From Concepts to Practice*; John Wiley & Sons: Chichester, UK, 2009.

(53) Lai, R.; Liu, Y.; Luo, X.; Chen, L.; Han, Y.; Lv, M.; Liang, G.; Chen, J.; Zhang, C.; Di, D.; Scholes, G. D.; Castellano, F. N.; Wu, K. Shallow distance-dependent triplet energy migration mediated by endothermic charge-transfer. *Nat. Commun.* **2021**, *12*, 1532.

(54) Cheng, Y. Y.; Fückel, B.; Khoury, T.; Clady, R. G. C. R.; Ekins-Daukes, N. J.; Crossley, M. J.; Schmidt, T. W. Entropically Driven Photochemical Upconversion. *J. Phys. Chem. A* **2011**, *115*, 1047–1053.

(55) Isokuortti, J.; Kuntze, K.; Virkki, M.; Ahmed, Z.; Vuorimaa-Laukkanen, E.; Filatov, M. A.; Turshatov, A.; Laaksonen, T.; Priimagi, A.; Durandin, N. A. Expanding excitation wavelengths for azobenzene photoswitching into the near-infrared range via endothermic triplet energy transfer. *Chem. Sci.* **2021**, *12*, 7504–7509.

(56) Tang, Q.; Zhang, H.-L.; Wang, Y.; Liu, J.; Yang, S.-P.; Liu, J.-G. Mitochondria-targeted carbon monoxide delivery combined with singlet oxygen production from a single nanoplatfrom under 808 nm light irradiation for synergistic anticancer therapy. *J. Mater. Chem. B* **2021**, *9*, 4241–4248.

(57) Queiroga, C. S. F.; Almeida, A. S.; Vieira, H. L. A. Carbon Monoxide Targeting Mitochondria. *Biochem. Res. Int.* **2012**, *2012*, 1–9.

(58) Motterlini, R.; Foresti, R. Biological signaling by carbon monoxide and carbon monoxide-releasing molecules. *Am. J. Physiol.: Cell Physiol.* **2017**, *312*, C302–C313.

(59) Lazarus, L. S.; Simons, C. R.; Arcidiacono, A.; Benninghoff, A. D.; Berreau, L. M. Extracellular vs Intracellular Delivery of CO: Does It Matter for a Stable, Diffusible Gasotransmitter? *J. Med. Chem.* **2019**, *62*, 9990–9995.

(60) Gomes, A.; Neves, M.; Cavaleiro, J. A. S. Cancer, Photo-dynamic Therapy and Porphyrin-Type Derivatives. *An. Acad. Bras. Cienc.* **2018**, *90*, 993–1026.

(61) Vitek, L.; Gbelcová, H.; Muchová, L.; Váňová, K.; Zelenka, J.; Koničková, R.; Šuk, J.; Zadinova, M.; Knejzlík, Z.; Ahmad, S.; Fujisawa, T.; Ahmed, A.; Ruml, T. Antiproliferative effects of carbon monoxide on pancreatic cancer. *Dig. Liver Dis.* **2014**, *46*, 369–375.