Folding-induced Fluorescence Enhancement in a Series of Merocyanine Hetero-Folda-Trimers

Alexander Schulz and Frank Würthner*

Abstract: Many dyes suffer from fast non-radiative decay pathways, thereby showing only short-lived excited states and weak photoluminescence. Here we show a pronounced fluorescence enhancement for a weakly fluorescent merocyanine (MC) dye by being co-facially stacked to other dyes in hetero-folda-trimer architectures. By means of fluorescence spectroscopy (lifetime, quantum yield) the fluorescence enhancement was explained by the rigidification of the emitting chromophore in the defined foldamer architecture and the presence of a non-forbidden lowest excited state in H-coupled hetero-aggregates. This folding-induced fluorescence enhancement (FIFE) for specific sequences of π-stacked dyes points at a viable strategy toward improved fluorophores that relates to the approach used by nature in the green fluorescent protein (GFP).

Fluorescence is one of the most useful properties of dyes with widespread applications in biomolecular imaging.[1–3] Different from absorption properties that are influenced only to a minor extent by the environment,[4] fluorescence is highly sensitive to the fluorophore’s surrounding. This is exemplified by common fluorescence quenching pathways such as internal conversion, often associated with conical intersections,[5,4] excited state proton or electron transfer,[7,2] and aggregation-caused quenching.[9] All of these processes enable fluorescence on/off switching as desired for sensing and imaging applications. Whilst the environmental impact has been considered for many of the most successful fluorophores for the respective applications, for example, DNA intercalation-induced emission enhancement for thiazole dyes such as TOTO (thiazole orange dimer),[10] there are only few examples where chemists have tried to engineer the dyes’ environment for functional control.[11]

Such functional control is accordingly still illustrated by the natural example of the green fluorescent protein (GFP) from jellyfish.[25] Here an essentially non-luminescent p-hydroxybenzylidene-imidazolidinone merocyanine (MC) dye[13] is rigidly embedded within a protein barrel, affording a bright fluorescent dye-protein construct. Genetic engineering of GFP afforded mutants that cover a large spectral range and were applied widely during the last two decades in biomolecular fluorescence imaging.[14] In contrast, the recovery of the luminescence of this particular chromophore in artificial supramolecular constructs was less successful.[15]

MC dyes[10] appear indeed well-suited to elucidate environmental effects on fluorescence properties. Despite being strong absorbers these dyes exhibit with few exceptions[17,18] only very weak fluorescence which has been rationalized by torsional motions leading to conical intersections in the excited state.[9] Accordingly, like in GFP, chromophore embedding within a rigidifying environment may lead to the highly desirable fluorescence enhancement. First evidence for such behavior has been observed in previous research on self-assembled MC oligomers[19–21] and on a series of MC dye foldamers bearing up to six identical chromophores of type R (symbol R for the red-colored solutions of this dye, for definition see Figure 1).[22,23]

However, these dye oligomers suffer strongly from the H-type exciton coupling in the co-facially stacked folded state, thereby enabling only modest fluorescence enhancements with fluorescence quantum yields up to 2% in chloroform/methylcyclohexane (MCH).[23] Because the best results in our previous work were accomplished for trimers, we herein explore a series of trimers composed of the previously applied dimethine MC dye R (abbreviation related to the red color) and a newly synthesized elongated tetramethine analogue B (abbreviation related to the blue color) (Figure 1a). Whilst fluorescence properties of such hetero-foldamers were hitherto never investigated, our previous analyses of exciton coupling in hetero MC aggregates suggested a more beneficial preservation of oscillator strength for the lowest exciton state of these co-facially stacked dyes, thereby enabling higher radiative rates compared to their homo-foldamer congeners.[24,25]

Figure 1a shows the molecular structures of the herein investigated folda-trimers (for synthetic details, see supporting information).[26] Figure 1b a schematic illustration of their dipole-dipole interaction[27] driven folding behavior and Figure 1c our important observation that the hetero-trimers show significantly increased fluorescence quantum yields (ΦF) compared to the monomeric reference dyes and the homo-trimer BBB in the partially folded state. To understand this interesting fluorescence behavior in hetero-trimmers, we first studied the folding behavior of all dyes by NMR and UV/Vis absorption spectroscopy. In general, the folding behavior of all dyes resembles the one of the previously investigated RRR homo-trimer.[23] Thus, the combination of dipole-dipole

[A. Schulz, Prof. F. Würthner
Institut für Organische Chemie and Center for Nanosystems Chemistry, Universität Würzburg Am Hubland, 97074 Würzburg (Germany)
E-mail: wuethner@uni-wuerzburg.de
】Supporting information and the ORCID identification number(s) for the author(s) of this article can be found under:
https://doi.org/10.1002/anie.202114667.
© 2021 The Authors. Angewandte Chemie International Edition published by Wiley-VCH GmbH. This is an open access article under the terms of the Creative Commons Attribution License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited.
interactions between these highly dipolar dyes with the pyridinedicarboxamide turn unit favors the formation of co-facially stacked dyes with antiparallel oriented dipole moments (Figure 1b).

Figure 1. a) Homo- and hetero-folda-trimers investigated in this study, b) schematic illustration for the solvent-dependent folding process and c) fluorescence quantum yields determined for the partially folded form in CHCl₃ (black) and the completely folded stack in CHCl₃/MCH (red) of the MC trimers at 293 K. Arrows indicate dipole moments in these donor-acceptor dyes.

Figure 2. UV/Vis absorption spectra in CHCl₃ (blue) and CHCl₃/MCH 1:3 (red) at 293 K for a) BRR (solid) and RRB (dashed), b) RRB and c) BBB (c = 5-1 μM). The insets show schematic illustrations of the excitation states originating from Coulomb coupling between the neighboring dyes in the fully folded co-facial stacks and the derived transition dipole moment for the low energy transition.

A decrease of the solvent permittivity by the addition of methycyclohexane (MCH) is needed to shift the equilibria towards the completely folded state for the other oligomers (Figure 2a and 2b, red lines). In the case of the hetero-trimmers BRR and RRB with the blue dye at an end position this process competes with intermolecular aggregation into extended H-aggregates as confirmed by additional concentration-dependent studies (Figure S7). For the latter two hetero-trimers it accordingly appears that the shorter red
chromophore in the middle does not perfectly comply with the geometrical requirements of the turn unit to perfectly accommodate the longer blue dye in a close co-facial π-π-stacking arrangement.

Owing to their unsymmetrical structure the hetero-trimers BRR and RRB retain non-negligible oscillator strength of the lowest energy transition even in the completely folded stack, leading to the pronounced absorption over 650 nm. In contrast, for RBB and BBB this transition becomes almost completely forbidden in the completely folded stack, indicating that the absorption around 650 nm in chloroform originates solely from the partially folded structure. This observation can be explained by the interaction of the transition dipole moments according to the molecular exciton theory[28] (Figure 2 insets) and is also nicely reproduced by TDDFT calculations (Figure S10).

Subsequently we studied the emission properties of the trimers in chloroform, in which partially folded and completely folded structures should co-exist according to the absorption spectra. Indeed, whilst the emission of BRR and RRB is dominated by a B-monomer-like band at around 690 nm, an additional broad and red-shifted emission above 750 nm is visible upon excitation at the sharp, high energy band originating from the completely folded stack (Figure 3a and Figure S8b). We therefore attribute the sharp emission to the partially folded state and the exciplex-like broader emission to the completely folded stack. In case of the trimers RBR and BBB, for which the transition to the lowest excited state becomes forbidden for the fully stacked ensemble, only the monomer-like emission can be observed regardless of the excitation wavelength (Figure 3b and Figure S7a), indicating that the fluorescence of the completely folded stack is quenched due to the forbidden nature of the transition from the lowest exciton state to the ground state (Figure 2b,c). This further suggests that the folding-unfolding process is much slower than the deactivation of the excited states. We adjusted our determination method of $\Phi_t$ (Figure S1 and Table S1) and the fluorescence lifetime ($\tau_f$; see Table S2) to accommodate this behavior. The obtained $\Phi_t$ for the partially folded structure shows that while BBB (1.3%) is quenched in comparison to B-ref (3%), the hetero-trimers BRR, RRR and RRB all show pronounced folding-induced fluorescence enhancement (FIFE), resulting in a quantum yield of up to 21% for RRB and 10% for the two other trimers (Figure 1c).

To gain further insight into the photophysical processes in the excited state we carried out time-dependent fluorescence measurements (Figure 4a). For the folda-trimers $\tau_f$-values of 0.6 (BBB) to 1.5 ns (RRR) were determined in chloroform, which are significantly longer compared to the ones of the monomeric references (ca. 0.1 ns). From these values the radiative ($k_r$) and non-radiative ($k_{nr}$) rate constants for all trimers were calculated (Figure 4b and Table S3).

![Figure 3.](image-url) Absorption (black), excitation (red, blue; solid) and emission (colored; dashed) spectra of a) RRB and b) RBR in CHCl$_3$ at 293 K. The excitation and emission wavelengths are indicated by arrows in the respective color. The absorption (dark blue area) and emission (light blue area) spectra of B-ref are included in b) for reference.

![Figure 4.](image-url) a) Fluorescence decays for RBR (black), RRR (red), BRR (blue), BBB (green), B-ref (blue, dashed) and R-ref (red, dashed) in chloroform at 293 K and (b) the radiative (black) and non-radiative (red) rate constants calculated from $\tau_f$ and $\Phi_t$. 

© 2021 The Authors. Angewandte Chemie International Edition published by Wiley-VCH GmbH
As expected for co-facially stacked structures with H-type coupling, the forbidden nature of the radiative transition from the lowest exciton state leads to a reduction of $k_r$ compared to the monomeric references. This effect is most pronounced for the homo-trimer BBB, while the expected preservation of $k_r$ could be observed for the hetero-trimers in accordance with the model shown in the insets of Figure 2. Furthermore, a reduction of $k_{nr}$ can be observed for all trimers, which we attribute to the rigidification of the emitting type B chromophore by the adjacent chromophores, hindering the common deactivation pathway for merocyanines via a conical intersection.[5,6] This rigidification seems to be least effective in the case of BBB, which we attribute to the existence of one weakly associated B chromophore in the partially folded structures, while for the hetero-trimers a more effective rigidification can be observed, indicating close proximity of the B chromophore to at least one adjacent R chromophore in the partially folded structure. In the symmetrical trimer RRR both rates show the strongest reduction, which can be rationalized by the position of the B chromophore in the center of the structure leading to an efficient rigidification but also a more forbidden lowest exciton H-state (Figure 2b, inset). RRR shows a similar rigidification but more beneficial preservation of $k_r$ leading to the increased $\Phi_l$. Surprisingly, the other unsymmetrical congener BRR shows a lower $k_r$ and higher $k_{nr}$ compared to RRR, indicating less effective rigidification and stronger H-coupling.

To investigate the reason for this unexpected result we measured excitation spectra for all trimers at their monomeric-like emission band at 690 nm (Figure 5a). The different intensity and spectral location of the maxima observed for the long wavelength band in the excitation spectra (blue shaded area) indicate a different environment of the B chromophore in all three hetero-trimmers. Notably, the trends observed for the shift and relative intensity coincide with the trends observed for $k_{nr}$ and $k_r$, respectively. This suggests that the observed shift is primarily attributable to a different polarization of the chromophore by the environment in the foldamer structure, while the decrease in intensity is attributable to H-type coupling as illustrated in the insets of Figure 2.[29] The different environmental influence on the blue dye at the end position of the unsymmetrical trimers can also be observed in the DFT optimized structure, which shows a favorable non-covalent contact between the B chromophore and the adjacent turn unit in RRR providing additional rigidification (Figure 5b). This interaction is absent in BRR which is in agreement with the higher $k_{nr}$ value for this trimer compared to RRR. Further, the calculated exciton coupling $J$ between adjacent chromophores[30] in the BR substructure is significantly higher in RRR (2363 cm$^{-1}$) compared to RRB (1629 cm$^{-1}$) resulting in a stronger contribution of the center R dye to the lowest excited state and therefore a reduction of the transition dipole moment in BRR (Figure 2 inset), which is in good agreement with the obtained $k_r$ (Table S5).

In accordance to the measurements in chloroform, only RRB and BRR show notable emission from the completely folded stack in CHCl$_3$/MCH 1:3 (Figure 1c and S9). While for both trimers a superposition of the monomer-like emission (vide supra) and the exciplex-like emission could be observed, isolated emission from the completely folded stack could only be obtained for RRR upon excitation at higher energies. The acquired emission spectrum was found to be rather broad with a red-shifted maximum of 750 nm (Figure S10). While the $\Phi_l$ of B-ref was almost entirely quenched in this solvent system due to the formation of self-assembled dimers,[27] the $\Phi_l$ of the exciplex-like emission of RRR was determined to be around 10% (Table S4). The lifetime of the latter was furthermore found to be significantly increased compared to the partially folded state. The derived rate constants $k_{nr}$ and $k_r$ for this fully folded structure are indeed further reduced compared to the partially folded one which can be rationalized by a reinforced rigidification in the completely folded stack and a more pronounced H-coupling, respectively. In this case the reduction in $k_r$ due to the more forbidden character of the low energy transition is more pronounced than the additional rigidification, thereby leading to the reduction of $\Phi_l$ (Table S4). By using the more viscous liquid paraffin solvent to create a more rigid environment, the $\Phi_l$ of the exciplex-like emission of RRR could be further enhanced to about 14% compared to 2% for B-ref in the same solvent mixture.

In conclusion, the absorption and emission properties of a series of foldamers composed of three merocyanine dyes have been investigated. We were able to discern the beneficial and detrimental effects on the emission of these typically only weakly luminescent chromophores upon organization in β-sheet-like foldamer structures. We found that rigidification
and H-coupling strongly depend on the position of the dye with the smallest band gap (most bathochromic absorption) in the stack and its surrounding environment. Whilst the used monomeric dyes had fluorescence quantum yields of < 3% that were even reduced in their homo-trimers, the best hetero-trimer ensemble showed a significant folding-induced fluorescence enhancement with a fluorescence quantum yield of up to 21% in the far-red to NIR spectral region. We envision that this approach of chromophore rigidification by an embedding environment can be applied to a large variety of dyes to afford GFP-like fluorescence enhancements.

Conflict of Interest

The authors declare no conflict of interest.

Keywords: aggregation · dyes/pigments · fluorescence · folding · merocyanines

[1] J. R. Lakowitz, Principles of fluorescence Spectroscopy, 3rd Ed., Springer, 2006.
[2] L. D. Lavis, R. T. Raines, ACS Chem. Biol. 2008, 3, 142 – 155.
[3] R. Micura, C. Hobartner, Chem. Soc. Rev. 2020, 49, 7331 – 7353.
[4] Strongly solvatochromic dyes may be considered here as an exception. But even for these dyes despite of strong batho- or hypsochromic shifts with increasing solvent polarity the intensity of the absorption band and related oscillator strength change only slightly.
[5] J. Shi, M. A. Izquierdo, S. Oh, S. Y. Park, B. Milián-Medina, D. Roca-Sanjuán, J. Gierschner, Org. Chem. Front. 2019, 6, 1948 – 1954.
[6] J. Hoche, A. Schulz, L. Dietrich, A. Humeniuk, M. Stolte, D. Schmidt, T. Brixner, F. Würthner, R. Mitric, Chem. Sci. 2019, 10, 11013 – 11022.
[7] J. Gierschner, J. Shi, B. Milián-Medina, D. Roca-Sanjuán, S. Varghese, S. Y. Park, Adv. Opt. Mater. 2021, 9, 2002251.
[8] S. Doose, H. Neuwiler, M. Sauer, ChemPhysChem 2009, 10, 1389 – 1398.
[9] J. Mei, N. L. C. Leung, R. T. K. Kwok, J. W. Y. Lam, B. Z. Tang, Chem. Rev. 2015, 115, 11718 – 11940.
[10] A. N. Glazer, H. S. Rye, Nature 1992, 359, 859 – 861.
[11] A. Fin, A. V. Jentzsch, N. Sakai, S. Matile, Angew. Chem. Int. Ed. 2012, 51, 12736 – 12739; Angew. Chem. 2012, 124, 12908 – 12911.

Manuscript received: October 29, 2021
Accepted manuscript online: November 16, 2021
Version of record online: November 29, 2021