Fluorofluorophores: Fluorescent Fluorous Chemical Tools Spanning the Visible Spectrum

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Supporting Information

ABSTRACT: “Fluoro” refers to both fluorescent and fluorinated compounds. Despite the shared prefix, there are very few fluorinated molecules that are soluble in perfluorinated solvents. This paucity is surprising, given that optical microscopy is a ubiquitous strategy throughout the physical sciences and the orthogonality of fluoruous materials is a commonly exploited strategy in synthetic chemistry, materials science, and chemical biology. We have addressed this shortage by synthesizing a panel of “fluoro” fluorophores, fluoruous molecules containing high weight percent fluorine with optical properties spanning the visible spectrum. We demonstrate the utility of these fluoruous fluorophores by preparing fluoruous perfluorocarbon nanoemulsions.

Highly fluorinated, or fluoruous, compounds have gained considerable popularity due to their orthogonality to aqueous and organic species. Fluoruous molecules, often defined as those that contain ≥60 weight percent fluorine (wt% F) in C(sp3)−F bonds, phase-separate from polar and nonpolar compounds due to their disinclination to participate in van der Waals interactions. This phenomenon, discovered by atomic scientists in the World War II era, did not gain widespread recognition by chemists until 1994, when Rábai and Horváth coined the term “fluorous.” Today, the unique properties of fluoruous compounds are exploited in both basic science and commercial applications. Fluorous materials are routinely employed for non-stick, self-cleaning, and anti-fouling coatings. Fluorinated compounds have been reported as components in organic electronics. Perfluorocarbon nanoemulsions have been used in vivo for oxygen delivery, MRI cell tracking, and ultrasound imaging and therapy. Unique nano- and microstructures can be obtained from semifluorinated surfactants and block copolymers. Fluorous phase synthesis facilitates simple structures can be obtained from semi-aqueous and organic species. Fluorous molecules, often defined as both fluorinated compounds due to their disinclination to participate in van der Waals interactions. This shortage extends to fluoruous phase, Matsui and coworkers synthesized fluorinated coumarin dyes for lasers (up to 52 wt% F) and found that fluorination resulted in increased photostability as well as reduced absorption and photoluminescence spectra. Recently, Sun et al. prepared highly fluorinated polyaromatics (60−70 wt% F), which also display improved photostability along with enhanced quantum yields of luminescence. The latter fluorinated polyaromatics are the only organic fluorophores reported to date that have been fully characterized in perfluorocarbons.

Here, we report a panel of highly fluorinated (54−61 wt% F) fluorous compounds, termed “fluoro” fluorophores, in which “fluoro” refers to both fluorescence and fluorination. We employed aminophenol 1 as a common building block to access six fluorophores traversing the visible spectrum in two or three steps from commercial materials. The array of absorption and emission properties of the fluorophores will allow Förster resonance energy transfer (FRET)-based analyses and multicolor optical microscopy experiments to be performed in the fluoruous phase. We demonstrate the utility and necessity of the fluorophores by preparing highly luminescent perfluorocarbon nanoemulsions (Figure 1).

Figure 1. Aminophenol 1 as a building block for fluorous fluorophores and preparation of fluorescent perfluorocarbon emulsions.

We synthesized 1 through dialkylation of 3-aminophenol with (3-perfluoroctyl)propyl iodide and subjected 1 to aromatic substitution reactions with a variety of electrophiles (Scheme 1, SI). Coumarins 2 and 3 were prepared by Lewis acid-mediated Pechmann condensation of 1 and β-ketoesters. A three-component reaction between 1, malononitrile, and benzaldehyde followed by oxidation with 2,3-dichloro-5,6-dicyanobenzo-
Scheme 1. Synthesis of Fluorofluorophores 2–7 from Aminophenol 1

![Scheme 1](image)

Quinone yielded chromene 4. Fluorinated rhodamine 5 was synthesized through a Friedel–Crafts acylation and condensation with phthalic anhydride followed by esterification. Squaraine 6 and oxazine 7 chromophores were obtained by subjecting 1 to squaric acid or N,N-dimethyl-4-nitrosoaniline, respectively. Moderate to low yields of the fluorofluorophores are primarily due to loss of product during the final purification. Despite the sub-optimal yields, the fluorofluorophores are prepared in two or three steps with no specialized techniques, facilitating their accessibility to many scientists.

The absorption (Abs) and photoluminescence (PL) of fluorofluorophores 2–7 extend across the visible spectrum (Figure 2A, Table 1). Coumarin 2 has the highest energy Abs and PL, displaying no color in solution and blue fluorescence upon excitation (Figure 2B). Coumarin 3 and chromene 4 have similar PL spectra, but 3 displays a more hypsochromically shifted Abs and has the largest Stokes shift of the reported fluorofluorophores. Similar extinction coefficients (ε) are observed for 2–4, but the quantum yield (Φ) of 4 is very low. Compound 5 has characteristics consistent with rhodamine dyes, including a large ε and fluorescence Φ in the middle of the visible spectrum. Squaraine dye 6 is deep blue in solution, displays a brilliant red PL when excited, and has the highest ε and Φ of the fluorofluorophores. Oxazine 7 spans a similar region of the electromagnetic spectrum as 6, although its Φ is lower.

Next, we analyzed the solubility of fluorofluorophores 2–7 in solvents with a range of polarities and fluorophilicities. Standard solvents such as acetone, tetrahydrofuran, and dichloromethane solubilized most of the fluorofluorophores, with the exception being squaraine 6, which has very limited solubility overall (Figure S1, Table S1). The best universal solvent for 2–7 was trifluorotoluene (Figure 2B), a solvent often employed in fluororganic synthesis. The affinity of the fluorofluorophores for fluorine-containing solvents was immediately evident when comparing their solubility in trifluorotoluene and toluene, where in the latter solvent the PL of 4–7 is quenched by aggregation (Figure 2B/C). We assayed the solubility of 2–7 in six fluorous solvents: methoxyperfluorobutane, perfluorotripropylamine, perfluoroctyl iodide, perfluoromethylcyclohexane, perfluorodecane, and perfluorohexanes (Figure S2, Table S2). The fluorofluorophores were moderately soluble in fluorous solvents containing a single heteroatom (O, N, or I); however, despite their significant wt% F, they were not readily dissolved in perfluoroalkanes, suggesting that the dispersion interactions of the π system greatly impact the fluorophilicity.

![Figure 2](image)

Table 1. Photophysical Characterization of 2–7

<table>
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<th>compd</th>
<th>wt% F</th>
<th>$\lambda_{max}$ (nm)</th>
<th>ε (cm$^{-1}$ M$^{-1}$)</th>
<th>$\lambda_{max}$ (nm)</th>
<th>Φ$\text{F}$</th>
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<td>641</td>
<td>71 900</td>
<td>664</td>
<td>0.16</td>
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</table>

$^a$Wt% F calculations exclude counterion. $^b$Squaraine 6 data in acetone.
Although the affinity of 2–7 for perfluoroalkanes is low, they can be rendered soluble in mixtures of fluoruous solvents that contain >50% perfluoroalkane (vidi infra).

We envision these fluorophores will be valuable chemical tools for all scientists working with the fluoruous phase. To showcase the utility of our new fluorophores, we prepared fluorescent perfluorocarbon (PFC) nanoemulsions. PFC emulsions were first developed in the 1960s as blood replacements, due to the high oxygen content in fluoruous solvents. Optical probes have only recently been added, as researchers strive to employ PFC nanoemulsions as scaffolds for multimodal imaging and therapy. The absence of fluoruous soluble fluorophores, particularly those with red emission, is evident when the details of optical imaging experiments with these emulsions are analyzed. Most strategies have involved introducing fluorescent molecules to the droplet surfaces either covalently or non-covalently, which results in low loadings of fluorophore that are not protected within the droplets. The panel of fluorophores reported herein should overcome these limitations and help facilitate the transition of PFC emulsions from oxygen delivery agents to multimodal theranostics.

We prepared PFC nanoemulsions using a simplified version of Fluosol-DA, containing a 3:7 mixture of perfluorotripropylamine (PFTPA, 8)/perfluorodecalin (PFD, 9) (20 wt%) and Pluronic-F68 (10, 2.8 wt%) in phosphate-buffered saline (PBS). Nanoemulsions were formed by ultrasonication and had a radius of ~120 nm, as determined by dynamic light scattering (Figure S3). An array of fluorescent emulsions was prepared by predissolving 2–7 in PFD/PFTPA prior to ultrasonication (Figures 3A and S4). The rhodamine 5-containing nanoemulsion was further analyzed by confocal microscopy (Figure 3B,C).

PFC nanoemulsions are kinetically stabilized, self-assembled colloids. To gain insight into their solution dynamics, we prepared emulsions containing 3 and/or 5. Fluorophores 3 and 5 have appropriate spectral overlap such that they undergo FRET, and thus their presence in the same emulsion droplet should be evident through PL spectroscopy. A 1:1 mixture of nanoemulsions containing 3 or 5 excited at 375 nm did not exhibit energy transfer (black dashed line) until subjected to ultrasonication (blue solid line) (Figure 3D). In contrast, nanoemulsions containing both 3 and 5 in each droplet displayed emission from 5 when excited at 375 nm (orange solid line). Even after multiple days and in the presence of serum, no significant FRET was observed in a mixture of emulsions where droplets contained only 3 or 5 (Figure S5). Collectively, these experiments indicate that the fluorophores are confined inside individual droplets and exchange between the droplets is minimal. Thus, despite the dynamic nature of nanoemulsions, adding perfluoroalkyl chains stabilizes the molecules inside the emulsions even when serum is present.

To further demonstrate that fluorination is essential for the long-term residence of fluorophores in the emulsions, we synthesized coumarin 11, which contains fewer fluorine atoms than 3, and coumarin 12, which contains only a CF3 group (Figure 4A, Schemes S2 and S3). Upon examining the photophysical properties of 11 and 12, the effect of the perfluoroalkyl chains was evident. Coumarins 3 and 11 have very similar photophysics, but coumarin 12 displays bathochromically shifted spectra and a ~2-fold reduction in quantum yield. These data are consistent with previous reports that indicate the electron-withdrawing effects of perfluoroalkyl groups can be felt through more than three methylene units and the rigidity of fluoruous moieties provides fewer opportunities for vibrational quenching.

Solutions of coumarins 3, 11, or 12 in PFD/PFTPA (400 μM) were prepared and emulsified with Pluronic-F68. The resulting nanoemulsions, and a control without fluorophore, were agitated continually in the presence of octanol. The PL of the octanol, which contained coumarin expelled from the PFC nanoemulsion droplets, was measured periodically over 2 weeks. The data indicate (Figure 4B) that the fluoruous chain has a large impact on the stability of the coumarin dyes inside the emulsions. The minimally fluorinated coumarin 12 readily displayed a large
amount of PL in the octanol (red line), while coumarins 3 and 11 were both well retained in the nanoemulsions (blue and green lines, respectively). As anticipated, the more fluorinated 3 displayed the best stability inside the droplets, demonstrating the utility of highly fluorinated fluorescent molecules. These results also suggest that wt% F can be correlated to release times, which, if coupled with a cleavable linker, could provide an avenue for controlled drug delivery.

We have synthesized a panel of fluorinated fluorescent molecules, termed fluorous fluorophores, which display photoluminescence spanning the visible spectrum. These compounds have the highest wt% F reported to date for their respective fluorine atoms. Synthesis of the fluorous fluorophores enabled preparation of highly fluorescent perfluorocarbon nanoemulsions, which are promising candidates for bright, non-toxic, in vivo imaging agents.27 We demonstrated that fluorination was essential for the fluorophores to be retained in the nanoemulsion droplets. More broadly, the fluorous fluorophores reported herein will allow scientists working with the fluoruous phase to add multicolor fluorescence microscopy to their experimental toolbox.

**ASSOCIATED CONTENT**

1 Supporting Information
Figures S1–S6, Tables S1 and S2, and Schemes S1–S3; experimental procedures; and characterization of 1–7, 11, and 12. This material is available free of charge via the Internet at http://pubs.acs.org.

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Notes
The authors declare no competing financial interest.

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**REFERENCES**