Supporting Information

Anthryl-Substituted Heterocycles as Acid-Sensitive Fluorescence Probes

Heiko Ihmels,^{a,*} Andreas Meiswinkel,^b Christian J. Mohrschladt,^b Daniela Otto,^a Michael Waidelich,^a

Michael Towler,^c Rick White,^c Martin Albrecht,^d Alexander Schnurpfeil^d

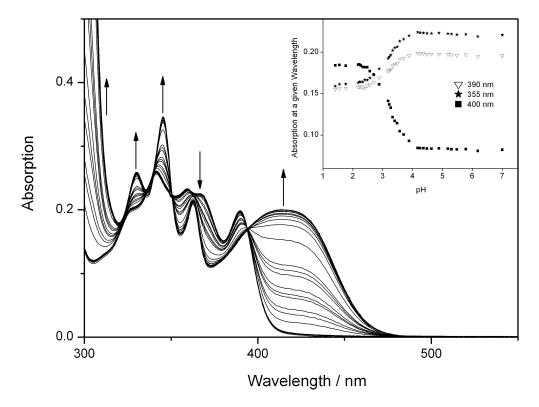


Figure 2. Spectrophotometric titration of **2a** (10⁻⁴ M in MeOH) with aqueous HCl; arrows indicate the development of band maxima during titration; inset: change of absorption intensity at 355, 390, and 400 nm at varying acid concentrations.

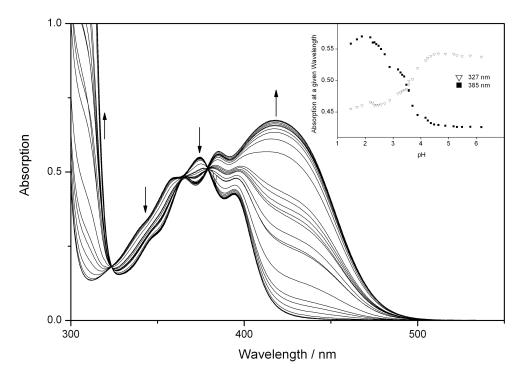


Figure 3. Spectrophotometric titration of **2b** (10⁻⁴ M in MeOH) with aqueous HCl; arrows indicate the development of band maxima during titration; inset: change of absorption intensity at 372 and 385 nm at varying acid concentrations.

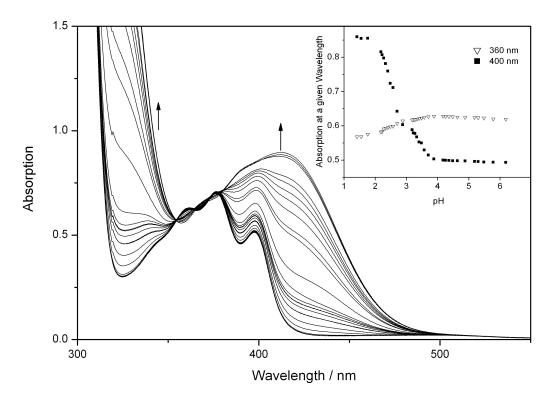


Figure 4. Spectrophotometric titration of **7** (10⁻⁴ M in MeOH) with aqueous HCl; arrows indicate the development of band maxima during titration; inset: change of absorption intensity at 360 and 400 nm at varying acid concentrations.

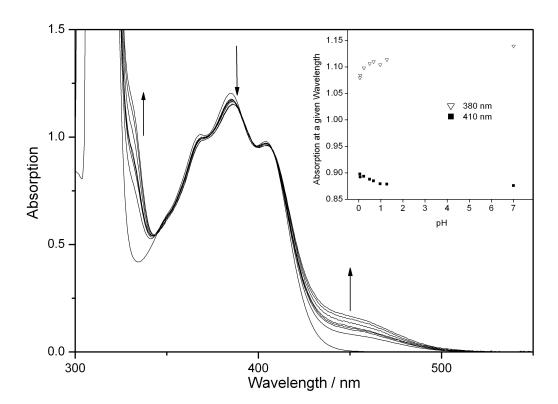


Figure 5. Spectrophotometric titration of **2c** (10⁻⁴ M in MeOH) with aqueous HCl; arrows indicate the development of band maxima during titration; inset: change of absorption intensity at 380 and 410 nm at varying acid concentrations.

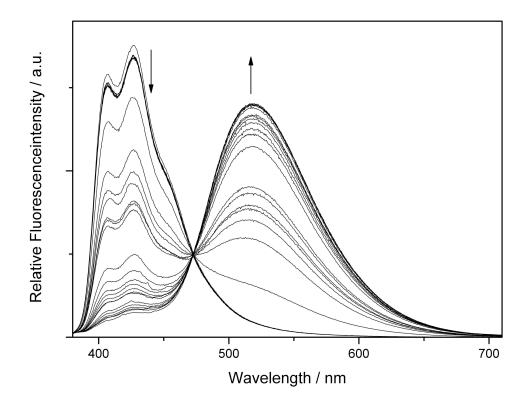


Figure 6. Spectrofluorimetric titration of **2a** (10^{-5} M in MeOH) with aqueous HCl; arrows indicate the development of band maxima during titration; $\lambda_{ex} = 363$ nm; before titration: c(HCl) = 0 M, titration end: $c(HCl) = 3.2 \times 10^{-2}$ M.

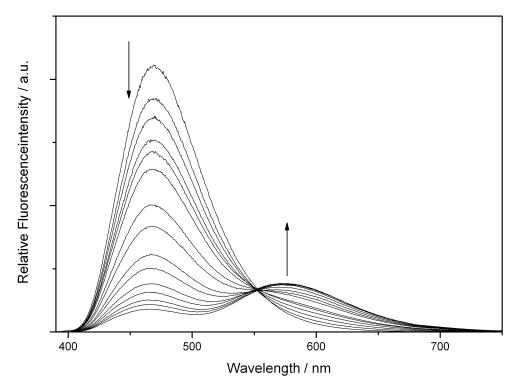
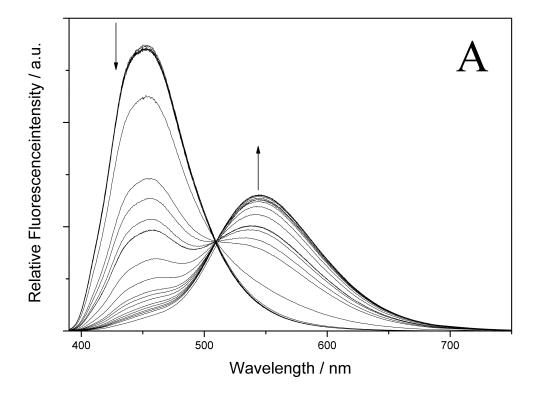


Figure 9. Spectrofluorimetric titration of 2c (10^{-5} M in MeOH) with aqueous HCl; arrows indicate the development of band maxima during titration; $\lambda_{ex} = 380$ nm before titration: c(HCl) = 0 M, titration end: $c(HCl) = 3.2 \times 10^{-1}$ M.



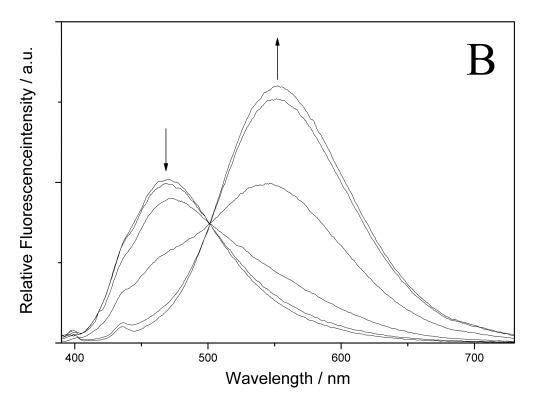
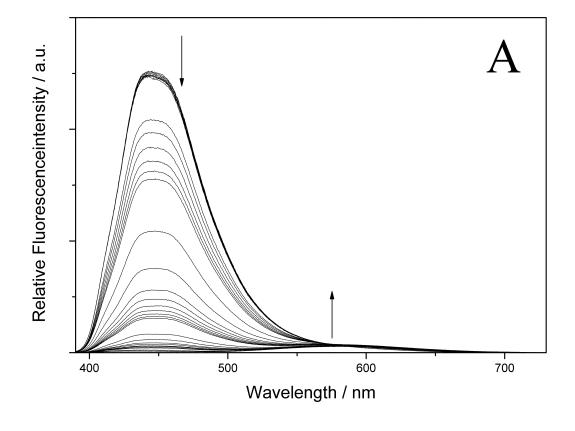


Figure 7. A: Spectrofluorimetric titration of **2b** (10^{-5} M in MeOH) with aqueous HCl; arrows indicate the development of band maxima during titration; before titration: c(HCl) = 0 M, titration end: $c(HCl) = 3.2 \times 10^{-2}$ M. B: Fluoresc ence spectra of **2b** (10^{-6} M in Britton-Robinson buffer at pH 2, 3, 4, 5, 6, 7; cont. 1% DMSO); $\lambda_{ex} = 380$ nm, arrows indicate the development of band maxima upon decreasing pH



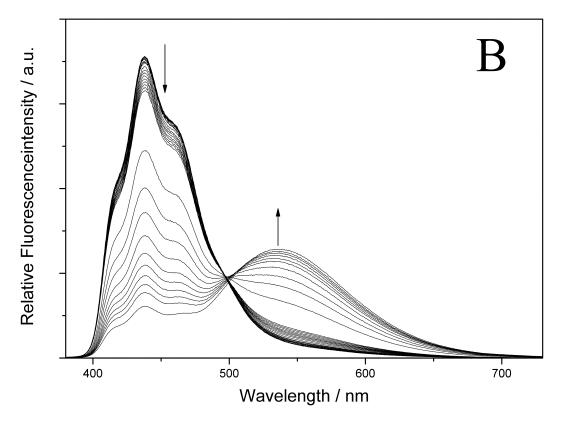


Figure 10. A: Spectrofluorimetric titration of **7** (10^{-5} M in MeOH) with aqueous HCl; arrows indicate the development of band maxima during titration; $\lambda_{ex} = 370$ nm; before titration: c(HCl) = 0 M, titration end: $c(HCl) = 5.0 \times 10^{-2}$ M. B: Spectrofluorimetric titration of **7** (10^{-5} M in CHCl₃) with trifluoroacetic acid; arrows indicate the development of band maxima during titration; $\lambda_{ex} = 370$ nm; before titration: c(HCl) = 0 M, titration end: $c(HCl) = 1.6 \times 10^{-2}$ M.