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NIR optical carbon dioxide sensors based on highly photostable dihydroxy-aza-BODIPY dyes†

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A new class of pH-sensitive indicator dyes for optical carbon dioxide sensors based on di-OH-aza-BODIPYs is presented. These colorimetric indicators show absorption maxima in the near infrared range (λ_{max} 670–700 nm for the neutral form, λ_{max} 725–760 nm for the mono-anionic form, λ_{max} 785–830 nm for the di-anionic form), high molar absorption coefficients of up to 77 000 M⁻¹ cm⁻¹ and unmatched photostability. Depending on the electron-withdrawing or electron-donating effect of the substituents the pK_a values are tunable (8.7–10.7). Therefore, optical carbon dioxide sensors based on the presented dyes cover diverse dynamic ranges (0.007–2 kPa; 0.18–20 kPa and 0.2–100 kPa), which enables different applications varying from marine science and environmental monitoring to food packaging. The sensors are outstandingly photostable in the absence and presence of carbon dioxide and can be read out *via* absorption or *via* the luminescence-based ratiometric scheme using the absorption-modulated inner-filter effect. Monitoring of the carbon dioxide production/consumption of a Hebe plant is demonstrated.

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Introduction

Carbon dioxide is one of the most important parameters in many scientific and industrial fields, such as medicine,^{1–3} marine science,^{4–7} food packaging,⁸ bio processing^{9,10} or environmental and industrial monitoring.¹¹ Routine techniques of CO₂ quantification like infrared (IR) spectroscopy, gas chromatography (GC) or the Severinghaus electrode are well established, but suffer from different drawbacks.^{12,13} For instance IR spectroscopy is mainly used for gas samples,^{14,15} because of strong interferences of water and the Severinghaus electrode is strongly affected by osmotic effects. Carbon dioxide chemosensors are a promising alternative,^{16–23} among which the so-called “plastic type” sensors are the most common ones.^{24–31} The core of this type of sensor is a pH-sensitive indicator dye. It is embedded in a polymer matrix together with a base (mainly a quaternary ammonium base) and responds to carbon dioxide by changing its spectral properties according to the degree of protonation. Read-out of these sensors can be carried out *via* absorption or luminescence intensity. In the case of a fluorescent

indicator dye, referencing of the fluorescence intensity is necessary to achieve reliable results. This can be realized by using an analyte-insensitive reference dye with different spectral properties from the indicator, either a different emission spectrum or a different luminescence decay time. Although necessary in many cases, the addition of reference materials can cause dramatic ratio changes when photobleaching of one dye (reference or indicator) or both occurs. Therefore, self-referencing indicator dyes are highly desired for ratiometric read-out. However, only a few self-referencing dyes (absorption and fluorescence intensity) have been published so far.^{32–38} Optical carbon dioxide sensors are also of great interest in biological applications.

Here, indicators with absorption/fluorescence intensity maxima in the infrared (IR) or near-infrared (NIR) region are preferred, because of several advantages, *e.g.* low light scattering, dramatically reduced autofluorescence and availability of low-cost excitation sources and photodetectors. Recently our group has presented pH-sensitive BF₂-chelated tetraarylazadi-pyrromethene indicators (aza-BODIPYs).³⁸ This indicator class represents an interesting alternative to the state-of-the-art dyes for biological applications as SNARF indicators³⁹ or cyanine dyes.^{40,41} The aza-BODIPYs showed absorption/fluorescence intensity spectra in the near-infrared region and were highly photostable. Although pH-sensitive, these dyes could not be used for optical carbon dioxide sensors. Here, the formation of the ion pair was irreversible and the indicator could not be protonated anymore, even at 100% CO₂. In this study we present a new class of pH-sensitive indicator dyes suitable for

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optical carbon dioxide sensors, the di-OH-aza-BODIPYs containing two de-/protonatable hydroxyl groups. It will be shown that the pK_a values of the di-OH-aza-BODIPYs can be tuned by using different substituents with either the electron-withdrawing or the electron-donating effect. Luminescence-based ratiometric read-out can be realized *via* the absorption-modulated inner-filter effect.^{42–45} An application example demonstrating the carbon dioxide production/consumption of a Hebe plant will also be presented.

Experimental

Materials

Ethyl cellulose (EC49, ethoxyl content 49%), thymol-blue (A.C.S. reagent), *m*-cresol-purple (indicator grade), poly(styrene-*co*-divinylbenzene)microspheres (8 μ m mean particle size; PS-microparticles), 3'-chloro-4'-hydroxyacetophenone, ammonium acetate, benzaldehyde, *N,N*-diisopropylethylamine (DIPEA), dry dichloromethane, boron trifluoride diethyl etherate, MOPS buffer salt, sodium sulfate (anhydrous) and tetraoctylammonium hydroxide solution (TOAOH, 20% in methanol) were obtained from Sigma-Aldrich (www.sigmaaldrich.com). Deuterated dimethyl sulfoxide (DMSO- d_6) was purchased from Eurisotop (www.eurisotop.com). Perfluorodecalin (98%; *cis* and *trans*), 1-butanol (99%, BuOH) and nitromethane were received from ABCR (Germany, www.abcr.de), Hyflon AD 60 from Solvay (www.solvay.com). Nitrogen, 2% oxygen in nitrogen, 5% carbon dioxide in nitrogen, 0.2% carbon dioxide in nitrogen, argon and carbon dioxide (all of 99.999% purity) were obtained from Air Liquide (Austria, www.airliquide.at). Toluene, ethanol (EtOH), tetrahydrofuran (THF), hydrochloric acid (37%), dichloromethane (DCM) and hexane were purchased from VWR (Austria, www.vwr.com). 3'-methyl-4-hydroxyacetophenone, 4'-butoxyacetophenone and 2-fluoro-4-hydroxyacetophenone were from TCI Europe (www.tcichemicals.com). Poly(ethylene terephthalate) (PET) support Melinex 505 was obtained from Pütz (Germany, www.puetz-folien.com). Potassium chloride, potassium carbonate (pro analysi), potassium hydroxide and silica gel 60 (0.063–0.200 mm) were received from Merck (www.merck.at). Sodium hydroxide, ethyl acetate, the buffer salts CHES, MES and CAPS were purchased from Roth (www.carlroth.com). De-ionized water was filtered *via* a Barnstead NANOpure ultrapure water system. 1-(4-Hydroxyphenyl)-4-nitro-3-phenylbutan-1-one (compound **1b**) was synthesized according to Jokic *et al.*³⁸ Silanized Egyptian blue particles were prepared according to the literature procedure.⁴⁶

Synthesis of 3,7-bis(4-butoxyphenyl)-5,5-difluoro-1,9-diphenyl-5H-4 λ^4 ,5 λ^4 -dipyrrolo[1,2-*c*:2',1'-*f*][1,3,5,2]triazaborinine (di-butoxy-complex)

(*E*)-1-(4-Butoxyphenyl)-3-phenylprop-2-en-1-one (compound **a**). 4'-Butoxyacetophenone (1 eq., 2.00 g, 15.5 mmol) was dissolved in absolute ethanol (15 ml). Benzaldehyde (1 eq., 1580 μ l, 15.5 mmol) and potassium hydroxide (3 eq., 2.61 g, 46.4 mmol in 5 ml H₂O) were added to the solution and the reaction was stirred at room temperature for 12 h. Then, the reaction

solution/suspension was acidified with 0.1 M HCl and the resulting precipitate was collected by filtration and was washed with water three times (3 \times 100 ml). The precipitate was dried on a rotary evaporator and was used for the next step without further purification (3.87 g, 89.2%).

1-(4-Butoxyphenyl)-4-nitro-3-phenylbutan-1-one (compound **b**).

A solution of (*E*)-1-(4-butoxyphenyl)-3-phenylprop-2-en-1-one (1 eq., 3.87 g, 13.8 mmol), nitromethane (20 eq., 14.8 ml, 276 mmol) and KOH (0.3 eq., 0.232 g, 4.1 mmol) in 30 ml of absolute ethanol was heated under reflux for 12 h. After cooling to room temperature the solvent was removed on a rotary evaporator. The resulting oily residue was acidified with 0.1 M HCl and was partitioned with ethyl acetate and water in a separating funnel. The organic layer was separated, dried over sodium sulfate and the solvent was removed under reduced pressure (4.035 g, 85.7%).

(5*Z*)-5-(4-Butoxyphenyl)-N-(5-(4-butoxyphenyl)-3-phenyl-2H-pyrrolo-2-ylidene)-3-phenyl-1H-pyrrolo-2-amine (**c**). Compound **b** (1 eq., 2.5 g, 7.33 mmol) and ammonium acetate (35 eq., 19.77 g, 256 mmol) were dissolved in 50 ml of 1-butanol and the reaction solution was heated under reflux for 24 h. The reaction was cooled to room temperature and the solvent was removed under reduced pressure. Then, the solid was redissolved in DCM and washed with water three times (3 \times 100 ml). The crude solid was purified by column chromatography on silica gel, eluting with DCM/cyclohexane (1 : 1 v/v). The product was recrystallized from a hexane-THF mixture to give metallic blue crystals (983 mg, 45.4%). ¹H NMR (300 MHz, CDCl₃) δ 8.06 (d, *J* = 7.1 Hz, 4H), 7.88–7.83 (m, 4H), 7.44–7.31 (m, 6H), 7.11 (s, 2H), 7.02 (d, *J* = 8.8 Hz, 4H), 4.06 (t, *J* = 6.5 Hz, 4H), 1.91–1.72 (q, *J* = 8.5 Hz, 4H), 1.6–1.48 (m, 4H), 1.01 (t, *J* = 7.4 Hz, 6H).

3,7-Bis(4-butoxyphenyl)-5,5-difluoro-1,9-diphenyl-5H-4 λ^4 ,5 λ^4 -dipyrrolo[1,2-*c*:2',1'-*f*][1,3,5,2]triazaborinine. Compound **c** (1 eq., 300 mg, 0.51 mmol) was dissolved in 200 ml of dry DCM. *N,N*-Diisopropylethylamine (DIPEA, 10 eq., 839 μ l, 5.06 mmol) and boron trifluoride diethyl etherate (15 eq., 953 μ l, 7.58 mmol) were added and the reaction solution was stirred under nitrogen for 12 h. The green solution was washed with water three times (3 \times 200 ml) and dried over anhydrous sodium sulfate. The crude product was purified by column chromatography on silica gel, eluting with DCM/cyclohexane (1 : 1 v/v). The product was recrystallized from a hexane-THF mixture to give metallic red needles (179 mg, 55.2%). ¹H NMR (300 MHz, CDCl₃) δ 8.09–8.05 (m, 8H), 7.49–7.38 (m, 6H), 7.04–6.98 (m, 6H), 4.07–4.02 (t, *J* = 6.4 Hz, 4H), 1.58–1.45 (q, *J* = 14.4 Hz, 4H), 1.62–1.42 (m, 4H), 1.02–0.97 (t, *J* = 7.4 Hz, 6H). Electron impact-direct insertion-time of flight (DI-EI-TOF): *m/z* of [M]⁺ was found to be 641.3007, calc. 641.3032.

Synthesis of 4,4'-(5,5-difluoro-1,9-diphenyl-5H-4 λ^4 ,5 λ^4 -dipyrrolo[1,2-*c*:2',1'-*f*][1,3,5,2]triazaborinine-3,7-diyl)bis(2-chlorophenol) (di-Cl-di-OH-complex)

1-(3-Chloro-4-hydroxyphenyl)-3-phenylpropenone (compound **1a**). 3'-Chloro-4'-hydroxyacetophenone (1 eq., 2 g, 11.7 mmol) and



benzaldehyde (1 eq., 1.24 g, 11.7 mmol) were dissolved in 10 ml of ethanol absolute. 10 ml of aqueous potassium hydroxide solution (3 eq., 1.96 g, 35.1 mmol) were added dropwise. The resulting solution was stirred for 8–12 h, during which the product precipitated as a potassium salt. The solution/suspension was poured into 10 ml of hydrochloric acid (1 M) and further concentrated hydrochloric acid was added until the solution became acidic. The obtained yellow solid was washed with water and used for further synthesis without purification (2.33 g, 77%).

1-(3-Chloro-4-hydroxyphenyl)-4-nitro-3-phenylbutan-1-one (compound 1b). A solution of compound **1a** (1 eq., 2 g, 7.7 mmol), nitromethane (20 eq., 8.35 ml, 154.7 mmol) and potassium hydroxide (1.2 eq., 0.52 g, 9.28 mmol) in 10 ml of ethanol was heated at 60 °C under reflux for 12 h. After cooling to room temperature, the solvent was removed *in vacuo* and the oily residue obtained was acidified with hydrochloric acid (4 M) and partitioned between ethyl acetate (50 ml) and de-ionized water (50 ml). The organic layer was separated, dried over anhydrous sodium sulfate and evaporated under reduced pressure. The obtained product was used for further synthesis without purification (2.4 g, 97.2%).

[5-(3-Chloro-4-hydroxyphenyl)-3-phenyl-1H-pyrrol-2-yl]-[5-(3-chloro-4-hydroxyphenyl)-3-phenylpyrrol-2-ylidene]amine (compound 1c). Compound **1b** (1 eq., 2.02 g, 8.76 mmol) and ammonium acetate (35 eq., 19.03 g, 307 mmol) in 50 ml of 1-butanol were heated under reflux for 24 h. The reaction was cooled to room temperature, the salt was removed by extraction with de-ionized water/DCM and the product was purified by column chromatography on silica gel, eluting with 5% ethyl acetate/DCM (after eluting impurities with hexane, 20% DCM/hexane, 1% ethyl acetate/DCM and 2% ethyl acetate/DCM) to yield **1c** as a blue-black solid. The product was recrystallized from a hexane–THF mixture to give metallic green crystals (0.097 g, 5.77%). ¹H NMR (300 MHz, DMSO-*d*₆) δ 8.09 (d, *J* = 7.6 Hz, 6H), 7.90 (d, *J* = 8.5 Hz, 2H), 7.62 (s, 2H), 7.54–7.32 (m, 6H), 7.19 (d, *J* = 8.5 Hz, 2H). EI-DI-TOF: *m/z* [MH]⁺ found 549.1007, calc. 549.1011.

4,4'-(5,5-Difluoro-1,9-diphenyl-5H-4λ⁴,5λ⁴-dipyrrolo[1,2-c:2',1'-ff]-[1,3,5,2]triazaborinine-3,7-diyl)bis(2-chlorophenol) (di-Cl-di-OH-complex). Compound **1c** (0.017 g, 0.07 mmol) was dissolved in 50 ml of dry DCM, treated with DIPEA (10 eq., 0.053 ml, 0.32 mmol) and boron trifluoride diethyletherate (15 eq., 0.061 ml, 0.48 mmol) and stirred under argon for 24 h. Purification was carried out *via* column chromatography on silica gel, eluting with 2% ethanol/DCM (after eluting impurities with DCM). Recrystallization from hexane–THF gave the final product di-Cl-di-OH-complex as a metallic red solid (0.008 g, 44%). ¹H NMR (300 MHz, DMSO-*d*₆) δ 8.29 (s, 2H), 8.16 (d, *J* = 7.0 Hz, 4H), 8.03 (d, *J* = 8.7 Hz, 2H), 7.65 (s, 2H), 7.60–7.41 (m, 6H), 7.13 (d, *J* = 8.7 Hz, 2H). EI-DI-TOF: *m/z* [MH]⁺ found 597.0968, calc. 597.0999.

Synthesis of 4,4'-(5,5-difluoro-1,9-diphenyl-5H-4λ⁴,5λ⁴-dipyrrolo[1,2-c:2',1'-ff]-[1,3,5,2]triazaborinine-3,7-diyl)bis(3-fluorophenol) (di-F-di-OH-complex). The synthesis was performed analogously to that of the di-Cl-di-OH-complex, but starting from

2'-fluoro-4'-hydroxyacetophenone (728 mg). Steps 1 and 2 yielded 1.033 g (90.2%) and 1.28 g (99.4%) of the crude product, respectively. After synthesis step 3, the crude product was purified by column chromatography on silica gel, eluting with 4% THF/DCM to yield [5-(2-fluoro-4-hydroxyphenyl)-3-phenyl-1H-pyrrol-2-yl]-[5-(2-fluoro-4-hydroxyphenyl)-3-phenylpyrrol-2-ylidene]amine **2c** as a blue-black solid. The product was recrystallized from a hexane–THF mixture to give metallic green crystals (0.23 g, 20.9%). ¹H NMR (300 MHz, DMSO-*d*₆) δ 8.04–8.06 (d, *J* = 7.0 Hz, 6H), 7.35–7.47 (m, 8H), 6.78–6.86 (m, 4H). DI-EI-TOF: *m/z* of [M]⁺ found 517.1626, calc. 517.1602.

4,4'-(5,5-Difluoro-1,9-diphenyl-5H-4λ⁴,5λ⁴-dipyrrolo[1,2-c:2',1'-ff]-[1,3,5,2]triazaborinine-3,7-diyl)bis(3-fluorophenol) (di-F-di-OH-complex). 4,4'-(5,5-Difluoro-1,9-diphenyl-5H-4λ⁴,5λ⁴-dipyrrolo[1,2-c:2',1'-ff]-[1,3,5,2]triazaborinine-3,7-diyl)bis(3-fluorophenol) (di-F-di-OH-complex) was synthesized using the same procedure. Purification was carried out *via* column chromatography on silica gel, eluting with 4% THF/DCM. Recrystallization from hexane–THF gave the final product di-F-di-OH-complex as a metallic red solid (0.044 g, 19.6%). ¹H NMR (300 MHz, DMSO-*d*₆) δ 8.12 (d, *J* = 7.0 Hz, 4H), 7.82 (t, *J* = 8.7 Hz, 2H), 7.56–7.48 (q, *J* = 9.4, 7.9 Hz, 6H), 7.31 (d, *J* = 3.4 Hz, 2H), 6.77 (d, *J* = 10.7 Hz, 4H). DI-EI-TOF: *m/z* of [M]⁺ found 565.1615, calc. 565.1591.

Synthesis of 4,4'-(5,5-difluoro-1,9-diphenyl-5H-4λ⁴,5λ⁴-dipyrrolo[1,2-c:2',1'-ff]-[1,3,5,2]triazaborinine-3,7-diyl)diphenol (di-OH-complex). Compound **3a** 1-(4-hydroxyphenyl)-3-phenylpropanone was commercially available. Further synthesis steps were performed analogously to the synthesis steps of the di-Cl-di-OH-complex. The synthesis of compound **3b** gave 2.06 g (80%) of the crude product, starting from 2.02 g of 4'-hydroxychalcone. The crude **3c** was purified by column chromatography on silica gel, eluting with 4% THF/DCM to yield [5-(4-hydroxyphenyl)-3-phenyl-1H-pyrrol-2-yl]-[5-(4-hydroxyphenyl)-3-phenylpyrrol-2-ylidene]amine as a blue-black solid. The product was recrystallized from a hexane–THF mixture to give metallic green crystals (0.41 g, 23.5%). ¹H NMR (300 MHz, DMSO-*d*₆) δ 8.09 (d, *J* = 7.3 Hz, 4H), 7.93 (d, *J* = 8.6 Hz, 4H), 7.53 (s, 2H), 7.42 (m, 6H), 7.01 (d, *J* = 8.6 Hz, 4H). EI-DI-TOF: *m/z* [MH]⁺ found 481.1773, calc. 481.179.

Purification of 4,4'-(5,5-difluoro-1,9-diphenyl-5H-4λ⁴,5λ⁴-dipyrrolo[1,2-c:2',1'-ff]-[1,3,5,2]triazaborinine-3,7-diyl)diphenol. Purification of 4,4'-(5,5-difluoro-1,9-diphenyl-5H-4λ⁴,5λ⁴-dipyrrolo[1,2-c:2',1'-ff]-[1,3,5,2]triazaborinine-3,7-diyl)diphenol was carried out *via* column chromatography on silica gel, eluting with DCM/ethyl acetate (4 : 1). Recrystallization from hexane–THF gave the final product di-OH-complex as a metallic red solid (0.26 g, 57.8%). ¹H NMR (300 MHz, DMSO-*d*₆) δ 8.13 (dd, *J* = 18.6, 8.0 Hz, 8H), 7.65–7.39 (m, 8H), 6.95 (d, *J* = 8.8 Hz, 4H). EI-DI-TOF: *m/z* [MH]⁺ found 529.1771, calc. 529.1779.

Synthesis of the 4,4'-(5,5-difluoro-1,9-diphenyl-5H-4λ⁴,5λ⁴-dipyrrolo[1,2-c:2',1'-ff]-[1,3,5,2]triazaborinine-3,7-diyl)bis(2-methylphenol) (di-CH₃-di-OH-complex). The synthesis was carried out analogously to that of the di-Cl-di-OH-complex, starting from 3'-methyl-4'-hydroxyacetophenone (1.88 g). Steps 1 and 2 yielded 2.77 g (93%) and 1.84 g (53%) of the crude



product, respectively. After synthesis step 3, the crude product was purified by column chromatography on silica gel, eluting with 5% ethyl acetate/DCM (after eluting impurities with hexane/DCM 1 : 1, DCM) to yield [5-(3-methyl-4-hydroxyphenyl)-3-phenyl-1*H*-pyrrol-2-yl]-[5-(3-methyl-4-hydroxyphenyl)-3-phenylpyrrol-2-ylidene]amine **4c** as a blue-black solid. The product was recrystallized from a hexane–THF mixture to give metallic green crystals (0.091 g, 5.8%). ¹H NMR (300 MHz, DMSO-*d*₆) δ 8.10 (d, *J* = 7.3 Hz, 4H), 7.91–7.75 (m, 4H), 7.57–7.33 (m, 8H), 7.00 (d, *J* = 8.4 Hz, 2H), 2.29 (s, 6H). EI-DI-TOF: *m/z* [MH]⁺ found 509.2086, calc. 509.2103.

*Purification of 4,4'-(5,5-difluoro-1,9-diphenyl-5*H*-4λ⁴,5λ⁴-dipyrrolo-[1,2-*c*:2',1'-*f*][1,3,5,2]triazaborinine-3,7-diyl)bis(2-methylphenol).* Purification of 4,4'-(5,5-difluoro-1,9-diphenyl-5*H*-4λ⁴,5λ⁴-dipyrrolo-[1,2-*c*:2',1'-*f*][1,3,5,2]triazaborinine-3,7-diyl)bis(2-methylphenol) was carried out *via* column chromatography on silica gel, eluting with 5% ethyl acetate/DCM (after eluting impurities with DCM, 1% ethyl acetate/DCM and 2% ethyl acetate/DCM). Recrystallization from hexane–THF gave the final product di-CH₃-di-OH-complex as a metallic red solid (0.04 g, 40%). ¹H NMR (300 MHz, DMSO-*d*₆) δ 8.16 (d, *J* = 7.2 Hz, 4H), 8.08–7.90 (m, 4H), 7.52 (m, 8H), 6.95 (d, *J* = 8.5 Hz, 2H), 2.22 (s, 6H). EI-DI-TOF: *m/z* [MH]⁺ found 557.2082, calc. 557.2092.

Staining of PS-microparticles

0.50 g of PS-particles were dispersed in a solution of 5 mg of (1% w/w in respect to the PS-particles) the di-butoxy-complex in 10 ml of tetrahydrofuran (THF) and stirred for 1.5 h. Then 8 ml of de-ionized water were added dropwise. After 10 min of stirring the dispersion was transferred rapidly into a beaker with 50 ml of de-ionized water. The dispersion was again stirred for 10 min. Then the particles were filtered through a cellulose filter and rewashed with 20 ml of ethanol. Afterwards the particles were transferred into a milling cup and were overlaid with a mixture of ethanol/de-ionized water (1:1) to reduce friction and to enhance heat dissipation during grinding in the planet mill. The particles were washed with ethanol and dried in the oven at 70 °C.

Sensor preparation

Absorption: a “cocktail” containing 100 mg of ethyl cellulose and 1 mg of dye (1% w/w with respect to the polymer) dissolved in a toluene: ethanol mixture (6:4 v/v, 1.9 g) was purged with carbon dioxide. Afterwards 100 μl of tetraoctylammonium hydroxide solution (20% w/w TOAOH in MeOH) were added. The “cocktail” was knife-coated on a dust-free PET support. A sensing film of ~7 μm thickness was obtained after evaporation of the solvent. For luminescence-based measurements a second layer was added to the absorption-based sensing foils. To prepare the “cocktail” for the second layer, 0.180 g of Hyflon AD 60 were dissolved in 2.820 g of perfluorodecalin, which was washed prior to use with 1 mol l⁻¹ potassium carbonate solution. Then 0.360 g of Egyptian blue powder (200% w/w with respect to the polymer) and 0.054 g of stained PS-particles (30% w/w with respect to the polymer) were dispersed homogeneously in

the Hyflon solution. The thickness of this layer after evaporation of the solvent was estimated to be ~4.5 μm.

Methods

¹H NMR spectra were recorded on a 300 MHz instrument (Bruker) in DMSO-*d*₆ with TMS as standard. Absorption spectra were recorded on a Cary 50 UV-Vis spectrophotometer (Varian). The determination of the molar absorption coefficients was carried out as an average of three independent measurements. Fluorescence spectra were recorded on a Fluorolog3 fluorescence spectrometer (Horiba) equipped with a NIR-sensitive photomultiplier R2658 from Hamamatsu (300–1050 nm). Photobleaching experiments were performed by irradiating the sensor foils in a glass cuvette with the light of a high-power 10 W LED array (λ_{max} 458 nm, 3 LEDs, www.LED-TECH.de) operated at 6 W input power. A lens (Edmund Optics) was used to focus the light of the LED array on the glass cuvette (photon flux: ~3900 μmol s⁻¹ m⁻² as determined using a Li-250A light meter from Li-COR). The photodegradation profiles were obtained by monitoring the absorption spectra of a sensor foil based on the respective dye. The cuvette was flushed with either carbon dioxide gas for the mono-anionic form, or with nitrogen for the di-anionic form and sealed. Thymol-blue and *m*-cresol-purple were used for comparison. In the case of the anionic form of thymol-blue, *m*-cresol-purple, the di-OH-complex and the di-CH₃-di-OH-complex (di-anionic forms) sodium hydroxide was placed at the bottom of the cuvette to capture carbon dioxide traces. Gas calibration mixtures were obtained using a gas mixing device from MKS (www.mksinst.com). The gas mixture was humidified to about 85% relative humidity, using silica gel soaked with a saturated potassium chloride solution, prior to entering the calibration chamber. Temperature was controlled by a cryostat ThermoHaake DC50. Dyed particles were filtered through cellulose filters type 113A from Roth (www.carlroth.com). Particle milling was carried out using an 80 ml milling cup, zirconia spheres (Ø 5 mm) and a Pulverisette 6 planet mill from Fritsch (www.fritsch.de). For the determination of the p*K*_a values titration curves in aqueous buffer/ethanol mixtures (1:1) were measured and the average value of the point of inflection obtained was used. For the production/consumption measurements of carbon dioxide two Firesting-Mini devices and an oxygen trace sensor from PyroScience (www.pyro-science.com) were used. Illumination of the sample was performed using 3 commercially available halogen bulb lamps with an averaged photon flux of ~217 μmol s⁻¹ m⁻² per lamp. The plant was positioned in a glass desiccator, which was flushed with a gas mixture of 2% oxygen in nitrogen.

Results and discussion

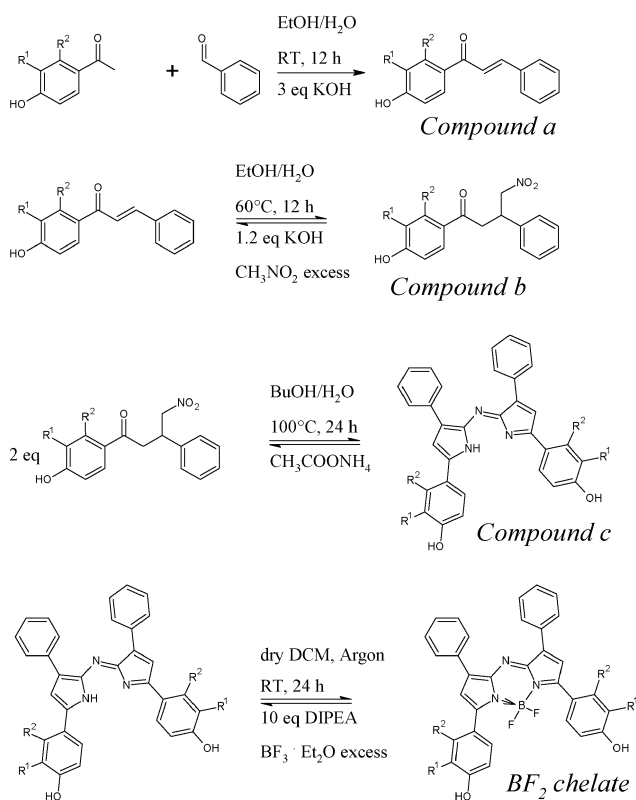
Synthesis

As reported by Jokic *et al.*³⁸ there are two ways for the preparation of azadipyrromethenes to obtain either asymmetrical⁴⁷ or symmetrical⁴⁸ chromophores. The previously published



asymmetrical dyes bearing one hydroxyl group were proved to be promising pH indicators.³⁸ Unfortunately, they were found to be unsuitable for optical carbon dioxide sensors. The ion pair built between the hydroxyl group and the tetraoctylammonium base was comparatively strong. Even after exposure of the sensors to pure carbon dioxide, the amount of CO₂ was not enough to achieve re-protonation of the indicators and only after exposure to strong acid vapors (e.g. hydrochloric acid) the indicators were irreversibly re-protonatable.

Dissociation of both hydroxyl groups in symmetrical azadi-pyrromethenes was expected to be more difficult compared to the mono-hydroxy derivatives since two ion pairs with bulky cations would be built upon deprotonation. Therefore, this new class of pH indicators potentially suitable for CO₂ sensing was investigated. The starting point of our synthesis was diaryl- α,β -unsaturated ketones (chalcones) either commercially available or readily made by an aldol condensation of the corresponding aldehyde and acetophenone (Scheme 1). The Michael addition of nitromethane to the chalcones, using KOH as a base, yields 1,3-diaryl-4-nitrobutan-1-ones in essentially quantitative yields. Their condensation with ammonium acetate in refluxing butanol gave azadipyrromethenes. Finally, the azadipyrromethenes were



di-OH-complex: R¹ = R² = H

di-F-di-OH-complex: R¹ = H, R² = F

di-Cl-di-OH-complex: R¹ = Cl, R² = H

di-CH₃-di-OH-complex: R¹ = CH₃, R² = H

Scheme 1 4-Step-synthesis of the di-OH-aza-BODIPY chelates.

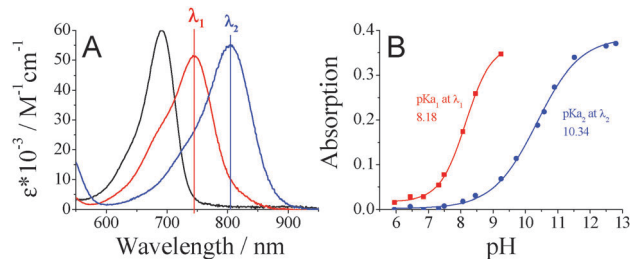


Fig. 1 (A) Absorption spectra of the neutral (black line, pH 6.4), the mono-anionic (red line, pH 9.3) and the di-anionic (blue line, pH 12.8) forms of the di-OH-complex dissolved in the ethanol/aqueous buffer mixture (1 : 1) at 25 °C and (B) the titration curves for pK_{a1} determined at 745 nm (red) and pK_{a2} at 805 nm (blue), respectively.

converted to the corresponding BF₂ chelates *via* reaction with BF₂-etherate using diisopropylethylamine as a base at room temperature for 24 h, giving yields ranging from 20 to 70%.

Photophysical properties

Dissolved in an ethanol/aqueous buffer mixture (1 : 1) all di-OH-aza-BODIPY dyes showed absorption spectra corresponding to the two protonation steps of the hydroxyl groups (Fig. 1). The neutral forms showed maxima at 670–700 nm and the mono-anionic forms at 725–760 nm. The di-anionic forms were again bathochromically shifted for approximately 60 nm (λ_{\max} 785–830 nm). Generally, the di-CH₃-di-OH-complex bearing electron-donating methyl groups showed absorption maxima shifted to higher wavelengths compared to the non-substituted di-OH-complex (Table 1). In contrast, complexes bearing electron-withdrawing groups displayed absorption maxima at shorter wavelengths.

An overview of the spectroscopic properties of the presented dyes is given in Table 1. Molar absorption coefficients are rather high: 55 000–77 000 M⁻¹ cm⁻¹. pK_a determination was carried out in a mixture of ethanol/aqueous buffer solution (1 : 1). For all di-OH-aza-BODIPY dyes two protonation steps can be observed (Table 1). Here, the complex with electron-donating methyl groups in the *ortho*-position to the hydroxyl groups showed the highest pK_a values, followed by the non-substituted di-OH-complex. Electron-withdrawing chlorine atoms located in the proximity of the hydroxyl groups (*ortho*-position) have the strongest impact on the pK_a values, whereas the electron-withdrawing effect of fluorine atoms in the *meta*-position is significantly lower.

Carbon dioxide sensors

The pH-sensitive di-OH-aza-BODIPY dyes were embedded in an ethyl cellulose matrix (EC49) along with tetraoctylammonium hydroxide as a base to obtain optical carbon dioxide sensors. Clearly, the sensors showed well-observable spectral changes in the near infrared (NIR) range according to the protonation of the di-anionic form giving the mono-anionic form (Fig. 2). The neutral forms of the di-OH-complexes showed slight fluorescence in solution, and the sensors based on the indicators did not show detectable fluorescence, neither for the mono-anionic



Table 1 Absorption maxima, molar absorption coefficients (ϵ) and pK_a values of the di-OH-aza-BODIPY-complexes dissolved in the ethanol/aqueous buffer mixture (1 : 1), as well as the absorption maxima for optical CO₂ sensors (EC49) based on the respective aza-BODIPY-complexes

Chelate	Neutral form		Mono-anionic form		Di-anionic form		
	λ_{max} ($\epsilon \times 10^{-3}$) buffer 1 : 1 [nm (M ⁻¹ cm ⁻¹)]	pK_{a1} EtOH/ aq. buffer 1 : 1	λ_{max} ($\epsilon \times 10^{-3}$) EtOH/aq. buffer 1 : 1 [nm (M ⁻¹ cm ⁻¹)]	EC49 [nm]	pK_{a2} EtOH/aq. buffer 1 : 1	λ_{max} ($\epsilon \times 10^{-3}$) EtOH/ aq. buffer 1 : 1 [nm (M ⁻¹ cm ⁻¹)]	EC49 [nm]
Di-CH ₃ -di-OH	701 (76.8 ± 1.9)	8.40	762 (65.5 ± 0.4)	775	10.68	827 (70.7 ± 0.6)	908
Di-OH	692 (59.9 ± 2.0)	8.18	745 (51.3 ± 1.2)	772	10.34	805 (55.2 ± 1.5)	876
Di-F-di-OH	669 (51.1 ± 2.0)	7.32	725 (46.7 ± 2.4)	760	9.35	784 (52.8 ± 0.8)	860
Di-Cl-di-OH	688 (61.4 ± 2.5)	6.52	752 (52.6 ± 2.2)	781	8.72	808 (57.9 ± 0.7)	890

nor for the di-anionic form. Compared to the measurements in solution the CO₂ sensors based on the di-OH-aza-BODIPY dyes showed significantly bathochromically shifted absorption spectra with maxima at 760–780 nm for their mono-anionic forms and maxima at 860–910 nm for their di-anionic forms. Generally, a similar trend for the absorption maxima of the CO₂ sensors based on the respective complexes was observed (Table 1). Here, the shift between the mono-anionic and the di-anionic form was enlarged to over 100 nm. This fact is advantageous for optical sensors, because peak separation becomes easier.

Despite the fact that all di-OH-aza-BODIPY dyes showed two protonation steps in aqueous solution, only one protonation step was observed in optical sensors based on these indicators. In fact, in the absence of carbon dioxide both hydroxyl groups build ion pairs with the quaternary ammonium base and the spectra of the di-anionic forms are observable. The mono-anionic form is built in the presence of carbon dioxide and is stable even at 100% CO₂. Spectra of the neutral forms are not observable anymore (Fig. 2). Therefore, the pK_{a2} values measured in solution are most relevant for optical carbon dioxide sensors based on the di-OH-aza-BODIPY dyes. Indeed, these values correlate very well with the sensitivities of the optical CO₂ sensors (Table 1 and Fig. 3). The sensitivity increases in the following order: di-Cl-di-OH < di-F-di-OH < di-OH < di-CH₃-di-OH. Notably, for the most sensitive sensor based on di-CH₃-di-OH about 25% of the overall signal change is observed already at atmospheric $p\text{CO}_2$ which is, to the best of our knowledge, one of the highest sensitivities reported so far. Di-F-di-OH and di-Cl-di-OH complexes bearing electron-withdrawing groups displayed diminished pK_{a2} values of 9.35 and 8.72, respectively,

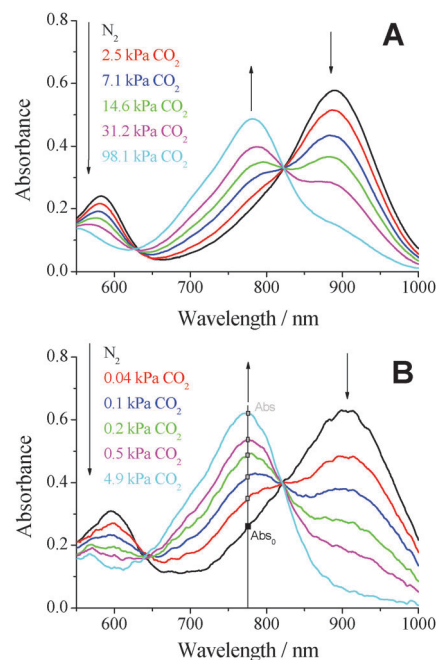


Fig. 2 Absorption spectra of optical carbon dioxide sensors based on (A) the di-Cl-di-OH-complex and (B) the di-CH₃-di-OH-complex in EC49 (base: TOAOH) at 25 °C at various $p\text{CO}_2$ values.



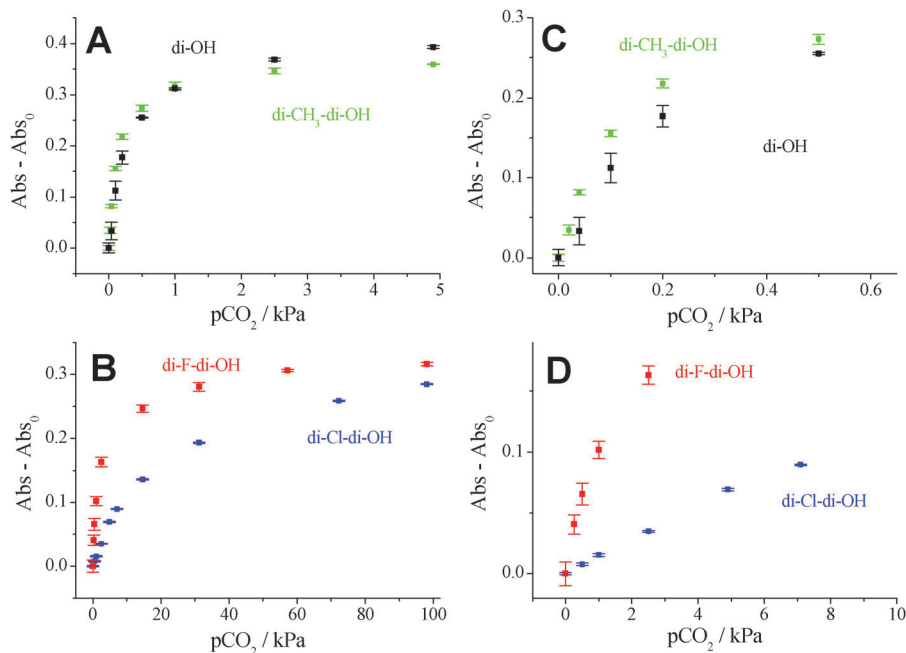


Fig. 3 (A) Calibration curves (absorption–absorption at 0 kPa of the mono-anionic form) for the carbon dioxide sensors based on the di-OH-complex (black) and the di-CH₃-di-OH-complex (green) and (B) di-Cl-di-OH-complex (blue) and the di-F-di-OH-complex (red) 25 °C under humid conditions with the respective “zoom-in” sections (C and D).

and enabled measurements up to 100% CO₂. These great differences in sensitivity lead to a broad range of applications, varying from food packaging and capnography for di-F-di-OH and di-Cl-di-OH complexes to environmental monitoring for di-OH and di-CH₃-di-OH complexes.

Fig. 3 shows the increasing absorption of the mono-anionic form in relation to the absorption at 0 kPa (Abs–Abs₀) carbon dioxide (see Fig. 2B). The higher the sensitivity of the sensor, the steeper the respective calibration curves, as shown in Fig. 3, and the lower the amount of carbon dioxide necessary to fully protonate the sensor. Sensors based on the di-Cl-di-OH-complex and the di-F-di-OH-complex showed limits of detection (LOD) of 0.19 kPa and 0.18 kPa, respectively. However, determining the LOD value for very sensitive sensors can be very challenging. For measurements of low levels of carbon dioxide the measuring system (gas mixer, gas lines, flow cell, *etc.*) has to be completely decarbonized, which is very difficult to achieve in reality. Especially for atmospheric levels of *p*CO₂ (0.04 kPa ≈ 400 ppm in the gas phase ≈ 13.6 μmol l⁻¹ in water at 298.15 K) and below, traces of environmental carbon dioxide disturbed the measurements for sensors based on the di-CH₃-di-OH-complex and the di-OH-complex. Hence, the determined LODs for these two complexes (0.011 kPa for di-OH and 0.007 kPa for di-CH₃-di-OH) were only rough estimations.

Photodegradation profiles of the carbon dioxide sensors were obtained from the absorption spectra after illuminating the sensor foils with a high-power LED array (λ_{max} 458 nm). For comparison, the state-of-the-art indicator dyes such as *m*-cresol-purple, thymol-blue and HPTS were used (ESI,† Fig. S1). The photostability of both the mono-anionic and the di-anionic form were investigated. Therefore, the cuvette was filled either

with pure carbon dioxide (mono-anionic form) or with pure nitrogen (di-anionic form). Clearly, all of the di-OH-aza-BODIPY dyes showed outstanding photostability, much better than the reference indicators embedded in the same ethyl cellulose matrix. After 1.5 h of illumination the presented dyes showed hardly any photobleaching effects for both the mono-anionic and di-anionic forms, whereas *m*-cresol-purple (neutral form), thymol-blue (neutral form) and HPTS (only anionic form) were degraded to less than 80%. The anionic forms of *m*-cresol-purple and thymol-blue were even less photostable than their neutral forms (ESI,† Fig. S1).

Luminescence-based ratiometric read-out using IFE (inner-filter effect) based sensors

Read-out of the planar optodes and fibre-optic sensors based on colorimetric systems is significantly more challenging than that of the luminescent systems. The inner-filter effect (IFE) was made use of in order to enable read-out *via* luminescence. Additionally to the first layer containing the absorption-based indicator dye along with TOAOH in ethyl cellulose, a second layer containing the secondary emitters was used.⁴² It included the pH-insensitive di-butoxy-complex entrapped in PS-particles as a fluorophore and Egyptian blue (EB) as a phosphor, both embedded in Hyflon AD 60. The absorption spectra of the di-OH-complex and the emission spectra of the secondary emitters are shown in Fig. 4A. Here, the broad emission band of the phosphor (EB) overlaps with the absorption spectrum of the di-anionic form of the indicator and the emission band of the fluorophore (di-butoxy-aza-BODIPY-complex) overlaps with the absorption spectrum of the mono-anionic form of the indicator. The isosbestic point of the di-OH-indicator is located at



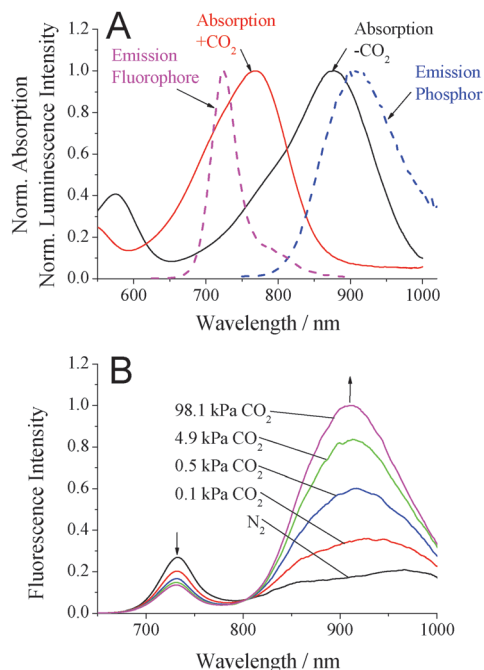


Fig. 4 (A) Emission spectra (λ_{exc} 610 nm) of Egyptian blue (EB; blue dashed line; "Emission Phosphor") and the di-butoxy-complex (dissolved in tetrahydrofuran; magenta dashed line; "Emission Fluorophore"); absorption spectra of the sensor based on the di-OH-complex in the absence (black line) and presence (red line) of carbon dioxide at 25 °C. (B) Emission spectra (λ_{exc} 610 nm) of a carbon dioxide sensor based on the di-OH complex and inner-filter effect read-out at 25 °C under humidified conditions.

610 nm and represents an ideal wavelength for exciting the secondary emitters matching the maxima of the red LEDs available (605, 617 nm). The fluorescence spectra of the sensor based on the combination of EB, the di-butoxy-complex and the non-substituted di-OH indicator are shown in Fig. 4B. Emission peaks in the absence (λ_{max} 913 nm) and presence (λ_{max} 738 nm) of carbon dioxide were well observable. Hence, luminescence-based ratiometric read-out becomes possible. This can be realized either by using two emission filters isolating the respective components or by measuring the luminescence phase shift. In fact, the phase shift for luminescent Egyptian blue is 55.8° at 2000 Hz and the phase shift of the fluorophore is 0 at this modulation frequency.

Carbon dioxide production/consumption of a Hebe plant

The applicability of the presented carbon dioxide sensors is demonstrated by showing the respiration of a Hebe plant. The Hebe plant in soil, a carbon dioxide sensor based on the di-CH₃-di-OH-complex and an oxygen trace sensor were placed in a desiccator. The desiccator was purged for 15 min with a gas mixture of 2% oxygen in nitrogen (in order to achieve better dynamics when measuring with an optical oxygen sensor compared to air saturation) and then closed tightly. During the measurement the setup was alternately kept in darkness and illuminated for 30 min using three halogen lamps. According to these illumination sequences the production/consumption of

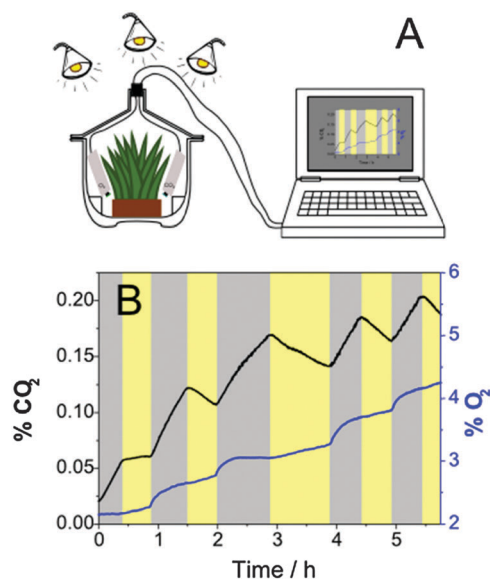


Fig. 5 (A) Experimental set-up and (B) carbon dioxide and oxygen dynamics in a desiccator containing a Hebe plant during illumination (yellow zones) and in darkness (gray zones).

carbon dioxide was observed (Fig. 5). In darkness an increase of carbon dioxide occurred due to respiration, whereas during illumination the concentration of CO₂ decreased. Over the whole measurement more CO₂ was produced than consumed which may be due to stress-induced respiration of the Hebe plant. Corresponding to the applied light sequences also the oxygen concentration was affected. Note that over the whole experiment the oxygen concentration was increasing which can be explained by slow diffusion of oxygen into the desiccator. Surprisingly, oxygen concentration did not increase significantly during the light phase, but it increased abruptly immediately after the light was switched off, reaching a plateau after about 30 min (Fig. 5). This effect may be attributed to the storage of generated oxygen in the plant and its release in the beginning of the dark phase. After this time the equilibrium between oxygen consumption during respiration and oxygen diffusion from outside is reached. As can be seen, the same phenomenon is observed if both the light and dark cycles are extended to 60 min (see hours 2 to 4 in Fig. 5). The above plant behavior is beyond the scope of the paper and calls for more detailed investigation.

Conclusion

A new class of colorimetric pH-sensitive indicators for carbon dioxide sensors has been presented. The di-OH-aza-BODIPY dyes show characteristic CO₂-dependent absorption spectra in the near-infrared region. In addition to the remarkable photostability of the indicators and the high molar absorption coefficients, the dynamic ranges of the sensors can be tuned *via* electron-donating and electron-withdrawing substituents. This enables a broad range of applications from environmental monitoring to food packaging or capnography. The sensors



based on di-CH₃-di-OH-aza-BODIPYs are the most sensitive sensors ever reported and resolve well below atmospheric CO₂ levels. The absorption-modulated inner-filter effect was used to enable referenced luminescence-based ratiometric read-out. As an example the production/consumption of carbon dioxide of a Hebe plant was demonstrated.

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