

Full Paper

## Multi-step Redox Systems with NIR-Fluorescence Based on 4*H*-Imidazoles

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**Abstract:** A new class of 4*H*-imidazoles was synthesized starting from fused-ring aromatic dinitriles. Strong bathochromic shifts of the longest wavelength absorptions were observed in the corresponding UV/vis spectra due to a conversion of the merocyanine chromophores into cyanines/(aza)oxonoles upon protonation/deprotonation of the 4*H*-imidazoles. Novel boratetraazapentalenes were synthesized via a cyclization reaction with boron trifluoride. These mesoionic species bearing a cyanine chromophore not only show NIR-fluorescence, they also participate as part of a quasi-reversible multi-step redox system. Large calculated semiquinone formation constants  $K_{SEM}$  ( $3 \times 10^{10}$  to  $5 \times 10^{11}$ ) indicate a high thermodynamic stability of the corresponding radical anions (SEM).

**Keywords:** 4*H*-Imidazoles, boratetraazapentalenes, multi-step redoxsystems, NIR-fluorescence.

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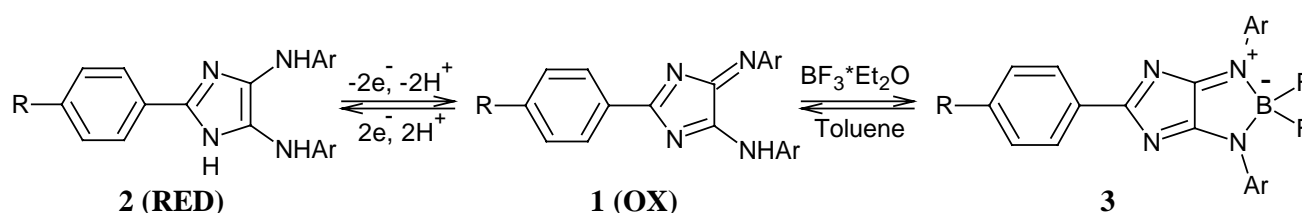
### Introduction

Over the last several years a great number of organic substances with extraordinary absorption, emission or electrochemical properties have been developed. In conjunction with materials science, many of these substances were employed in new materials such as displays, organic LED's or other,

mostly electronic devices. Only a few organic systems combine all the desirable properties for all the mentioned application fields, thus making research on polyfunctional dyes very popular.

4*H*-Imidazoles **1** are well known for their long wavelength UV/vis absorptions and high extinction coefficients ( $\log \epsilon > 4$ ) [1]. As part of a two-electron redox system, the reduction of the deeply coloured 4*H*-imidazoles **1** (OX) leads to nearly colourless, fluorescent 1*H*-imidazoles **2** (RED) via radical-ion intermediates (SEM) (Scheme 1). Cyclization of derivatives **1** with boron compounds yields red fluorescent boratetraazapentalenes **3**, which on the other hand may be regarded as electron-poor heterocycles. The radical anions of this new class of electrophores show unusual high  $K_{SEM}$  values (up to  $10^{15}$ ) [2].

**Scheme 1.** Redox behaviour and cyclization of 4*H*-imidazoles.



As part of a four-electron redox system, in which the radical state is replaced by more stable closed-shell moieties, *bis*-4*H*-imidazoles show almost identical properties as their monofunctional parent compounds [3]. Due to their long wavelength absorptions in the UV/vis spectra, correlated with high extinction coefficients, together with their interesting redox behaviour, 4*H*-imidazoles are promising candidates for the synthesis of functional dyes [4].

Therefore we sought to prepare 4*H*-imidazoles which are connected to other chromophores. The involvement of ring-fused aromatic systems such as anthracene or naphthalene, should lead to bathochromic shifts of the absorptions/emissions of the corresponding boratetraazapentalenes. Due to the switch between merocyanine and cyanine upon protonation of 4*H*-imidazoles, their application as acidochromic dyes also should be of interest.

## Results and Discussion

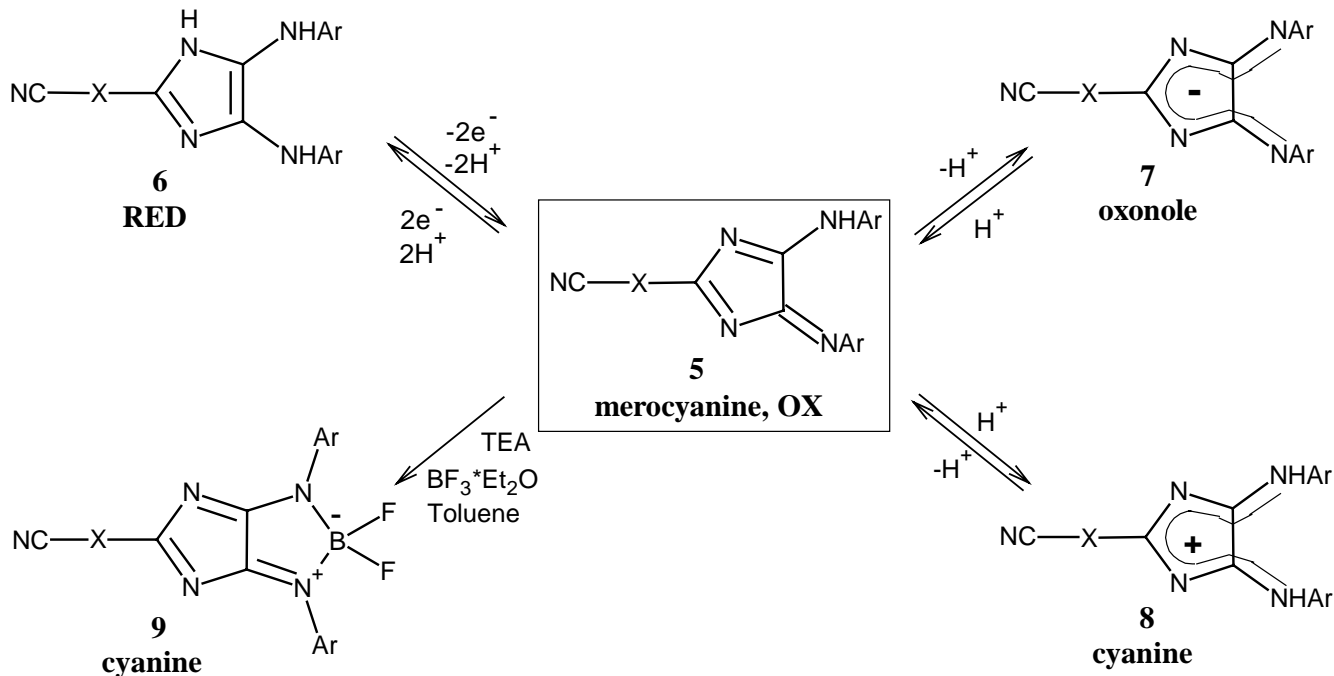
As the starting material required for *bis*-4*H*-imidazoles, dinitriles of type **4** were synthesized via a condensation reaction of *o*-xylylenedicyanide with the corresponding 1,2-diketones in the presence of sodium ethoxide [5,6]. The naphthalene derivative **4d** was obtained starting from 1,5-naphthalenediol [7,8]. All reported dinitriles are yellow-greenish fluorescent solids and, with the exception of derivative **4d**, sparingly soluble in organic solvents. Reactions of the dinitriles **4** with lithium hexamethyldisilazide (LiHMDS) yielded the corresponding amidinates, which were converted into the persilylated amidines by heating with chlorotrimethylsilane under reflux. As depicted in Scheme 2, all attempts to interconvert both nitrile groups failed. This failure can be attributed to considerable delocalization of the negative charge over the aromatic system which significantly lowers the electrophilicity of the second cyano group. The observed decrease of reactivity is in accordance with



**Table 1.** Redox potentials and semiquinone formation constants of 4*H*-imidazoles **5**.

Comp.	Ar	$E^1_{\text{Red}}, E^2_{\text{Red}}$	$K_{\text{SEM}}$
<b>5a</b>	4-tolyl	-0.72V, -1.05V	$3.9 \cdot 10^5$
<b>5a</b>	4- <i>n</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	-0.85V, -1.16V	$1.8 \cdot 10^5$
<b>5a</b>	4- <i>t</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	-0.87V, -1.17V	$1.2 \cdot 10^5$
<b>5b</b>	4- <i>n</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	-1.15V, -1.48V	$3.9 \cdot 10^5$
<b>5b</b>	4- <i>t</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	-1.02V, -1.34V	$2.7 \cdot 10^5$
<b>5c</b>	4-tolyl	-0.84V, -1.12V	$5.6 \cdot 10^4$
<b>5c</b>	4- <i>n</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	-0.75V, -1.12V	$1.9 \cdot 10^6$
<b>5c</b>	4- <i>t</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	-0.79V, -1.13V	$5.8 \cdot 10^5$
<b>5d</b>	4-tolyl	-0.90V, -1.31V	$8.9 \cdot 10^6$
<b>5d</b>	4- <i>n</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	-0.94V, -1.39V	$4.2 \cdot 10^7$

Chemically induced reduction could be achieved by treating a solution of the 4*H*-imidazole **5** with sodium dithionite in THF. The colour of the reaction mixture changed from red to light yellow, which indicates the formation of 1*H*-imidazoles **6**. These nearly colourless, but fluorescent products ( $\lambda_{\text{max,em}}$ : 439 up to 665nm), immediately reoxidize when exposed to air (Scheme 3).

**Scheme 3.** Modifications of 4*H*-imidazoles.

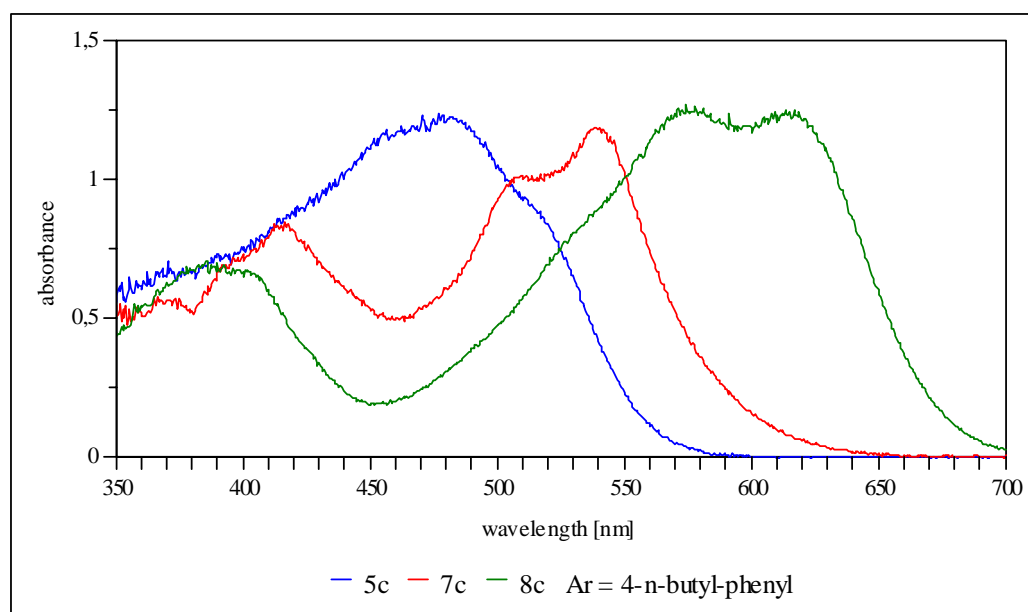
A further interesting feature of 4*H*-imidazoles **5** is the protonation-deprotonation reaction with acids or bases (Scheme 3). While the neutral species is based on a merocyanine system, the formation of the protonated or deprotonated 4*H*-imidazole causes an alteration of their chromophoric systems and consequently of their long wavelength absorption in UV/vis spectra. Compared to the neutral species, both the protonated (cyanine) and the deprotonated 4*H*-imidazole (azaaxonole) display a bathochromic shift in their UV/vis spectra (Table 2).

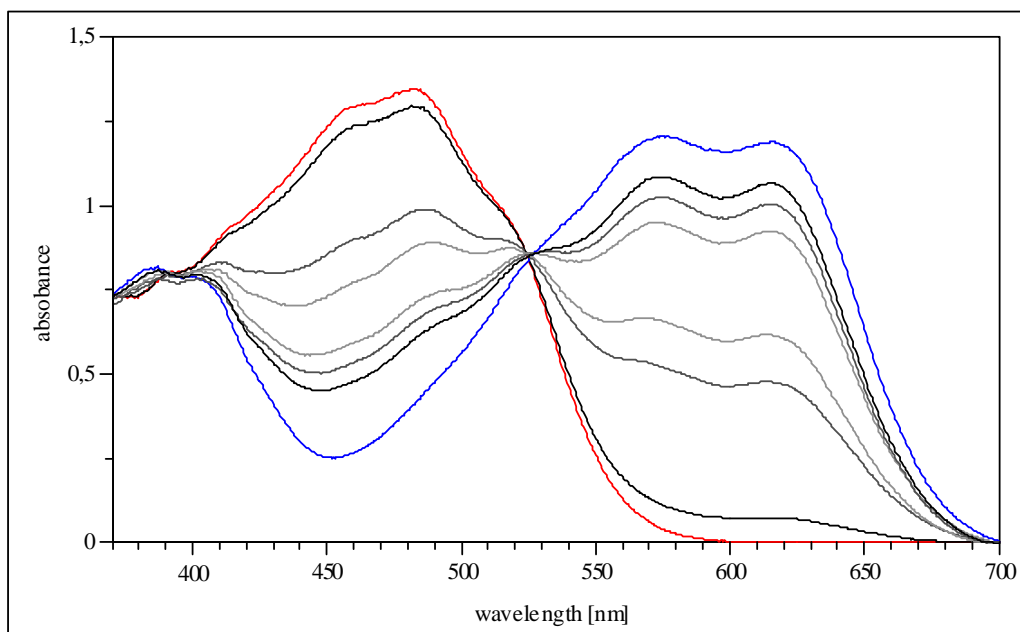
**Table 2:** Absorption and emission maxima of reduced **6**, deprotonated **7** and protonated **8**.

Comp.	Ar	$\lambda_{\max}/\text{type } 5$ (log $\epsilon$ )	$\lambda_{\max}/\text{type } 6$ (log $\epsilon$ )	$\lambda_{\max,\text{em}}/\text{type } 6$ ( $\lambda_{\max,\text{exc.}}$ )	$\lambda_{\max}/\text{type } 7$ (log $\epsilon$ )	$\lambda_{\max}/\text{type } 8$ (log $\epsilon$ )
<b>5-8a</b>	4-tolyl	519 (4.1)	409 (4.2)	439 (401)	533 (4.1)	619 (4.2)
<b>5-8a</b>	4- <i>n</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	524 (4.0)	411 (4.2)	455 (418)	540 (4.1)	623 (4.2)
<b>5-8a</b>	4- <i>t</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	521 (4.1)	412 (4.2)	475 (429)	542 (4.1)	625 (4.2)
<b>5-8b</b>	4- <i>n</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	523 (4.3)	406 (4.2)	535 (457)	535 (4.1)	620 (4.2)
<b>5-8b</b>	4- <i>t</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	517 (4.3)	412 (4.2)	570 (461)	538 (4.1)	622 (4.2)
<b>5-8c</b>	4-tolyl	520 (4.2)	409 (4.2)	448 (408)	535 (4.1)	614 (4.2)
<b>5-8c</b>	4- <i>n</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	525 (4.2)	409 (4.2)	471 (427)	538 (4.1)	617 (4.3)
<b>5-8c</b>	4- <i>t</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	526 (4.1)	410 (4.2)	470 (428)	540 (4.1)	618 (4.2)
<b>5-8d</b>	4-tolyl	536 (4.1)	415 (4.2)	651 (432)	549 (4.1)	631 (4.2)
<b>5-8d</b>	4- <i>n</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	540 (4.2)	421 (4.2)	665 (451)	555 (4.1)	637 (4.2)

UV/vis and fluorescence spectra were recorded in THF-solution.

Usually the cyanines **8** and the oxonoles **7** display bathochromic shifts of approximately 100 nm and 20 nm, respectively, with regards to the merocyanine **5** (Figure 1). The isosbestic point in the UV/vis spectra during protonation or deprotonation of the 4*H*-imidazole **5c** indicates a first order reaction (Figure 2). Thus, 4*H*-imidazoles are well suited for the application as acidochromic dyes, especially in sensor systems.

**Figure 1.** UV/vis spectra (THF) of anthracene derivative **5c**, deprotonated **7c** and protonated **8c** species.

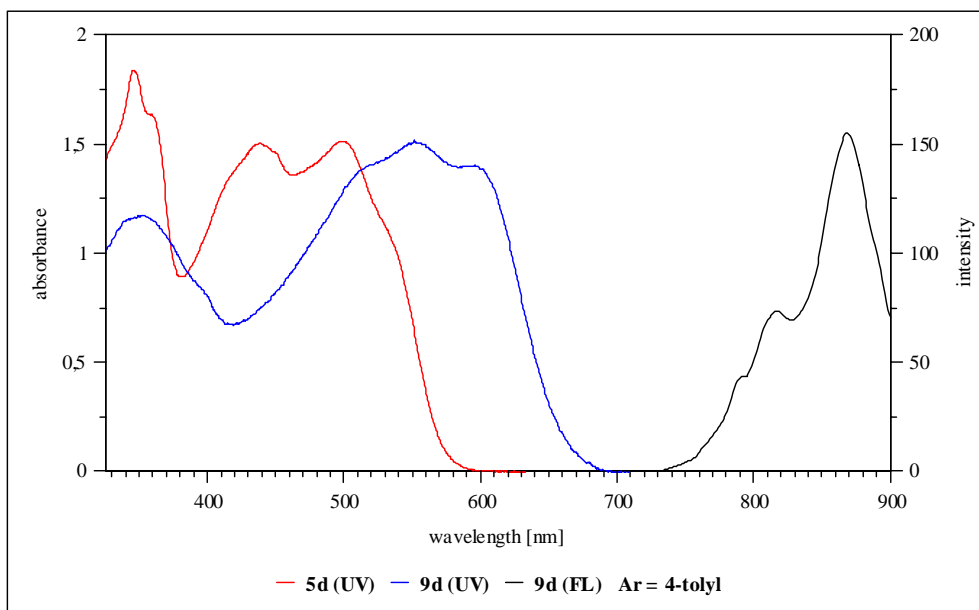
**Figure 2:** Isosbestic point (at 524nm, THF) during protonation of **5c**.

The mesoionic boracycles **9** were obtained in about 50% yield by cyclization of 4*H*-imidazoles **5** with boron trifluoride diethyl ether complex in the presence of triethylamine. These deep blue reaction products were purified by column chromatography and identified by MS-, IR-, NMR- and UV/vis spectroscopy. Cyclic voltammetric measurements showed the quasi-reversibility of the reduction step. The very large calculated semiquinone formation constants  $K_{SEM}$  ( $3 \times 10^{10}$  to  $5 \times 10^{11}$ ) indicate a high thermodynamic stability of the radicalanions (SEM) (Table 3).

**Table 3:** Electrochemical properties of the 1,3,2-diazaborolidines **9**.

Comp.	Ar	$E^1_{Red}, E^2_{Red}$	$K_{SEM}$
<b>9a</b>	4- <i>n</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	-0.45V, -1.14V	$5.0 \cdot 10^{11}$
<b>9a</b>	4- <i>t</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	-0.48V, -1.16V	$3.4 \cdot 10^{11}$
<b>9b</b>	4- <i>t</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	-0.88V, -1.50V	$3.2 \cdot 10^{10}$
<b>9c</b>	4- <i>n</i> -C <sub>4</sub> H <sub>9</sub> -C <sub>6</sub> H <sub>4</sub> -	-0.42V, -1.05V	$4.8 \cdot 10^{10}$
<b>9d</b>	4-tolyl	-0.82V, -1.47V	$1.0 \cdot 10^{11}$

Compared with the phenylogous boracycles **3**, the absorptions of 1,3,2-diazaborolidines **9** are shifted bathochromically (between 600 and 620nm). An unusually high stokes-shift of about 250 to 300 nm leads to the longest wavelength emissions in the area of NIR ( $\Phi < 20\%$ ), exemplified by derivative **9d** which is emitting at 870 nm (Figure 3).

**Figure 3.** UV/vis- and fluorescence spectra of derivative **9d** (solvent THF).

## Conclusions

We have prepared the new 4*H*-imidazoles **5** and mesoionic 1,3,2-diazaborolidines **9**, and have characterized them by elemental analysis, MS and NMR. Strong bathochromic shifts of the longest wavelength absorptions were observed in the corresponding UV/vis spectra due to a conversion of the merocyanine chromophores into cyanines/(aza)oxonoles upon protonation/deprotonation. The mesoionic boracycles **9** bearing a cyanine chromophore not only show NIR-fluorescence with an unusually high Stokes-shift, but they also participate as part of a quasi-reversible multi-step redox system. Large calculated semiquinone formation constants indicate a high thermodynamic stability of the corresponding radical anions. Thus, derivatives of 4*H*-imidazole are well suited for the applications in the functional dyes field.

## Experimental

### General

All reactions were monitored by TLC, carried out on 0.25 mm Merck silica gel plates (60F<sub>254</sub>) using UV light for visualization. <sup>1</sup>H- and <sup>13</sup>C-NMR spectra were recorded with a Bruker DRX 400 or Bruker AC 250 spectrometer. Melting points are measured with a Galen TM 3 apparatus and are uncorrected. UV/vis spectra were recorded on a Perkin-Elmer Lambda 19 spectrophotometer. MS spectra were taken from measurements on a Finnigan MAT SAQ 710 mass spectrometer. Elemental analyses were carried out in-house with an automatic analyzer LECO CHNS 932. Electrochemical measurements were carried out in dichloromethane with a Metrohm 663VA Stand using mercury or platinum electrodes (reference electrode SCE) and tetrabutylammonium hexafluorophosphate as conductive salt. The fluorescence spectra were recorded with a JASCO P-6500 instrument. Dinitriles **4a**, **4b** [5], **4c** [6] and **4d** [7,8] were prepared according to the published literature procedures.

*9,14-Dicyanobenz[b]triphenylene (4a)*. Yield 91%, yellow solid; mp 244-246°C (acetic acid); IR  $\text{cm}^{-1}$ : 2216 (CN); MS (EI):  $m/z$  (%): 328 ( $M^+$ , 100), 299 (20), 164 (15), 150 (18), 136 (20); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 256 nm (4.6), 307 (4.6), 320 (4.4), 395 (4.0), 417 (3.9); fluorescence (THF, 325 nm):  $\lambda_{\text{max,em}}$ : 487 nm.

*7,12-Dicyanobenz[k]fluoranthene (4b)*. Yield 85%, light yellow solid; mp 357-359 °C (tetrachloroethane); Anal. calc. for  $\text{C}_{22}\text{H}_{10}\text{N}_2$ : 87.4% C, 3.3% H, 9.3% N; found 87.2% C, 3.4% H, 9.4% N; IR  $\text{cm}^{-1}$ : 2220 (CN); MS (EI):  $m/z$  (%): 302 ( $M^+$ , 100), 273 (10), 151 (23), 123 (13), 111 (5), 28 (5); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 246 nm (3.7), 302 (3.3), 315 (3.6), 329 (3.8), 371 (4.1), 407 (3.8); fluorescence (THF, 363 nm):  $\lambda_{\text{max,em}}$ : 440 nm.

*9,10-Anthracenedicarbonitrile (4c)*. Yield 70%, yellow needles; mp 340-341°C (*p*-xylene); IR  $\text{cm}^{-1}$ : 2215 (CN); MS (DEI):  $m/z$  (%): 228 ( $M^+$ , 100), 201 (6), 174 (4); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 261 nm (4.5), 359 (3.6), 377 (4.0), 401 (4.0), 425 (4.0); fluorescence (THF, 333 nm):  $\lambda_{\text{max,em}}$ : 458 nm; CV (DMF):  $E_{\text{RED}} = -0.81\text{V}$ .

*2,6-Dicyano-1,5-di-*n*-hexyloxynaphthalene (4d)*. Yield 60% (three steps), orange solid; mp 87°C (lit. [8] mp 88°C); IR  $\text{cm}^{-1}$ : 2227 (CN); MS (DEI):  $m/z$  (%): 378 ( $M^+$ , 10), 294 (27), 223 (18), 209 (100), 85 (20), 43 (84); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 285 nm (3.7), 295 (3.8), 308 (3.7), 347 (3.7), 363 (3.7), 440 (3.9), 469 (3.7);  $^1\text{H-NMR}$  (250 MHz,  $\text{CDCl}_3$ )  $\delta$  (ppm): 0.85 (t,  $^3J = 6.3$  Hz, 6H), 1.28-1.51 (m, 12H), 1.87 (m, 4H), 4.40 (t,  $^3J = 6.3$  Hz, 4H), 7.45 (d,  $^3J = 10.0$  Hz, 2H), 7.65 (d,  $^3J = 10.0$  Hz, 2H); fluorescence (THF, 470 nm):  $\lambda_{\text{max,em}}$ : 511 nm.

#### General procedure for the synthesis of 4*H*-imidazoles 5:

Dinitrile **4** (1 mmol) was dissolved or suspended in dry THF (25 mL). After the addition of 1.0M Li-HMDS solution in THF or *n*-hexane (5 mL, 5 mmol), the reaction mixture was stirred at room temperature for 4 days and then evaporated to dryness *in vacuo*. The residue was dissolved in dry toluene (20 mL) and of chlorotrimethylsilane (0.75 mL, 6.0 mmol) was added. The reaction mixture was heated to 110 °C for 12 h and then the solvent was removed *in vacuo*. The silylated amidine was dissolved in dry THF (20 mL) and cyclized by addition of KF (0.35 g, 6.0 mmol), 18-crown-6-ether (1.58 g, 6.0 mmol) and the appropriate *bis*-imidoylchloride (1.5 mmol; Ar = 4-tol, 4-*n*- $\text{C}_4\text{H}_9$ - $\text{C}_6\text{H}_4$ -, 4-*t*- $\text{C}_4\text{H}_9$ - $\text{C}_6\text{H}_4$ - [11]), whereby the colour changed from yellow to deep red. The mixture was stirred for 3-4 hours at about 40 to 50 °C to ensure completion of the reaction. The 4*H*-imidazoles **5** were obtained by filtration and column chromatography of the crude product ( $\text{SiO}_2$ , toluene/acetone 50:1).

*9-{5-*p*-Tolylamino-4-[*p*-tolylimino]-4*H*-imidazol-2-yl}-benzo[b]triphenylene-14-carbonitrile (5a, Ar = 4-tolyl)*: Yield 21%, red solid, practically insoluble in all organic solvents tested; IR  $\text{cm}^{-1}$ : 3283 (NH), 2210 (CN); MS (DEI):  $m/z$  (%): 577 ( $M^+$ , 100), 486 (28), 459 (88), 326 (20), 215 (12), 106 (6), 91 (9); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 408 nm (4.2), 453 (4.2), 482 (4.3), 519 (4.1); CV:  $E_{\text{RED}}^1 = -0.42\text{V}$  (CN),  $E_{\text{RED}}^2 = -0.72\text{V}$ ,  $E_{\text{RED}}^3 = -1.05\text{V}$ .

9-{5-*p-n*-Butylphenylamino-4-[*p-n*-butylphenylimino]-4*H*-imidazol-2-yl}-benzo[*b*]triphenylene-14-carbonitrile (**5a**, Ar = 4-*n*-butylphenyl): Yield 28%, red solid; IR  $\text{cm}^{-1}$ : 3480 (NH), 2210 (CN); MS (DEI):  $m/z$  (%): 661 ( $M^+$ , 20), 351 (75), 175 (68), 106 (100); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 423 nm (4.1), 455 (4.1), 485 (4.2), 524 (4.0);  $^1\text{H-NMR}$  (250 MHz, THF- $d_8$ )  $\delta$  (ppm): 0.89 (t, 6H), 1.55 (m, 4H), 2.56 (t, 4H), 7.31 (t, 2H), 7.52 (t, 2H), 7.71 (m, 6H), 7.88 (d,  $^3J = 8.8$  Hz, 4H), 8.52 (d,  $^3J = 8.8$  Hz, 2H), 8.61 (d,  $^3J = 8.8$  Hz, 2H), 10.07 (s, 1H); CV:  $E_{\text{RED}}^1 = -0.47$  V (CN),  $E_{\text{RED}}^2 = -0.85$  V,  $E_{\text{RED}}^3 = -1.16$  V.

9-{5-*p-t*-Butylphenylamino-4-[*p-t*-butylphenylimino]-4*H*-imidazol-2-yl}-benzo[*b*]triphenylene-14-carbonitrile (**5a**, Ar = 4-*t*-butylphenyl): Yield 30%, red solid; IR  $\text{cm}^{-1}$ : 3338 (NH), 2212 (CN); MS (EI):  $m/z$  (%): 661 ( $M^+$ , 10), 352 (60), 337 (95), 328 (12), 175 (46), 160 (100), 134 (92); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 211 nm (4.8), 298 (4.7), 386 (4.0), 481 (4.2), 521 (4.1);  $^1\text{H-NMR}$  (250 MHz, THF- $d_8$ )  $\delta$  (ppm): 1.31 (s, 18H), 7.37 (d,  $^3J = 8.8$  Hz, 4H), 7.3-7.4 (m, 6H), 7.76 (d,  $^3J = 8.8$  Hz, 4H), 7.91 (d,  $^3J = 8.8$  Hz, 2H), 8.52 (t, 2H), 8.60 (t, 2H), 10.12 (s, 1H); CV:  $E_{\text{RED}}^1 = -0.51$  V (CN),  $E_{\text{RED}}^2 = -0.87$  V,  $E_{\text{RED}}^3 = -1.17$  V.

12-{5-*p-n*-Butylphenylamino-4-[*p-n*-butylphenylimino]-4*H*-imidazol-2-yl}-benzo[*k*]fluoranthene-7-carbonitrile (**5b**, Ar = 4-*n*-butylphenyl): Yield 15%, red solid; IR  $\text{cm}^{-1}$ : 3298 (NH), 2224 (CN); MS (DEI):  $m/z$  (%): 635 ( $M^+$ , 8), 578 (10), 508 (40), 352 (25), 306 (50), 263 (83), 175 (59), 132 (100), 106 (66); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 315 nm (4.2), 403 (4.3), 487 (4.4), 523 (4.3);  $^1\text{H-NMR}$  (250 MHz, THF- $d_8$ )  $\delta$  (ppm): 0.95 (t, 6H), 1.35 (m, 4H), 1.58 (m, 4H), 2.62 (m, 4H), 6.98-7.33 (m, 14H), 7.73 (d,  $^3J = 8.3$  Hz, 4H);  $^{13}\text{C-NMR}$  (100 MHz, THF- $d_8$ )  $\delta$  (ppm): 13.8, 21.8, 31.5, 31.9, 34.8, 35.0, 101.4, 116.8, 117.6, 120.9, 121.2, 122.5, 125.5, 126.1, 126.3, 126.4, 126.8, 127.2, 128.1, 128.2, 128.6, 129.3, 131.2, 132.0, 132.5, 132.8, 132.9, 133.1, 134.0, 135.4, 136.2, 138.4, 141.0, 148.0, 158.9; CV:  $E_{\text{RED}}^1 = -1.15$  V,  $E_{\text{RED}}^2 = -1.48$  V.

12-{5-*p-t*-Butylphenylamino-4-[*p-t*-butylphenylimino]-4*H*-imidazol-2-yl}-benzo[*k*]fluoranthene-7-carbonitrile (**5b**, Ar = 4-*t*-butylphenyl): Yield 18%, red solid; IR  $\text{cm}^{-1}$ : 3315 (NH), 2221 (CN); MS (DEI):  $m/z$  (%): 635 ( $M^+$ , 30), 578 (35), 477 (58), 352 (100), 337 (94), 302 (26), 160 (39), 132 (28); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 286 nm (4.2), 406 (4.2), 490 (4.4), 517 (4.3);  $^1\text{H-NMR}$  (250 MHz, THF- $d_8$ )  $\delta$  (ppm): 1.28 (s, 18H), 7.07-7.18 (m, 6H), 7.36-7.40 (m, 8H), 7.74 (d,  $^3J = 8.3$  Hz, 4H);  $^{13}\text{C-NMR}$  (100 MHz, THF- $d_8$ )  $\delta$  (ppm): 31.4, 31.6, 34.8, 35.0, 101.4, 116.8, 117.6, 120.9, 121.2, 122.5, 125.5, 126.1, 126.3, 126.4, 126.8, 127.2, 128.1, 128.2, 128.6, 129.3, 131.2, 132.0, 132.5, 132.8, 132.9, 133.1, 134.0, 135.4, 136.2, 138.4, 141.0, 148.0, 158.9; CV:  $E_{\text{RED}}^1 = -1.02$  V,  $E_{\text{RED}}^2 = -1.50$  V.

10-{5-*p*-Tolylamino-4-[*p*-tolylimino]-4*H*-imidazol-2-yl}-anthracene-9-carbonitrile (**5c**, Ar = 4-tolyl): Yield 19%, red solid; IR  $\text{cm}^{-1}$ : 3545 (NH), 2222 (CN); MS (DEI):  $m/z$  (%): 477 ( $M^+$ , 100), 462 (10), 359 (4), 228 (10), 221 (2), 106 (2); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 390 nm (4.1), 410 (4.1), 452 (4.2), 480 (4.3), 520 (4.2);  $^1\text{H-NMR}$  could not be recorded (almost insoluble in organic solvents); CV:  $E_{\text{RED}}^1 = -0.44$  V (CN),  $E_{\text{RED}}^2 = -0.84$  V,  $E_{\text{RED}}^3 = -1.12$  V.

*10-{5-*p-n*-Butylphenylamino-4-[*p-n*-butylphenylimino]-4*H*-imidazol-2-yl}-anthracene-9-carbonitrile (5c, Ar = 4-*n*-butylphenyl):* Yield 27%, red solid; IR  $\text{cm}^{-1}$ : 3257 (NH), 2221 (CN); MS (DEI):  $m/z$  (%): 561 ( $M^+$ , 100), 504 (68), 402 (20), 228 (10); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 393 nm (4.1), 411 (4.2), 455 (4.2), 482 (4.3), 525 (4.2);  $^1\text{H-NMR}$  (250 MHz,  $\text{CD}_2\text{Cl}_2$ )  $\delta$  (ppm): 0.82 (t,  $^3J = 7.5$  Hz, 6H), 1.24 (m, 4H), 1.47 (m, 4H), 2.48 (q,  $^3J = 7.5$  Hz, 4H), 7.12 (d,  $^3J = 8.3$  Hz, 4H), 7.52 (d,  $^3J = 7.8$  Hz, 2H), 7.68 (d,  $^3J = 7.8$  Hz, 2H), 7.77 (d,  $^3J = 8.3$  Hz, 4H), 8.42 (t,  $^3J = 7.3$  Hz, 2H), 8.52 (t,  $^3J = 7.3$  Hz, 2H); CV:  $E_{\text{RED}}^1 = -0.44$  V (CN),  $E_{\text{RED}}^2 = -0.75$  V,  $E_{\text{RED}}^3 = -1.12$  V.

*10-{5-*p-t*-Butylphenylamino-4-[*p-t*-butylphenylimino]-4*H*-imidazol-2-yl}-anthracene-9-carbonitrile (5c, 4-*t*-butylphenyl):* Yield 29%, red solid; IR  $\text{cm}^{-1}$ : 3456 (NH), 2220 (CN); MS (DEI):  $m/z$  (%): 561 ( $M^+$ , 8), 504 (1), 436 (3), 379 (4), 351 (100), 336 (19), 296 (16), 175 (21), 160 (21), 134 (17), 91 (23); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 453 nm (4.3), 484 (4.3), 526 (4.1);  $^1\text{H-NMR}$  (250 MHz, THF- $d_8$ )  $\delta$  (ppm): 1.30 (s, 18H), 7.36 (d,  $^3J = 8.5$  Hz, 2H), 7.41 (d,  $^3J = 8.5$  Hz, 4H), 7.61 (t, 2H), 7.69 (d,  $^3J = 8.5$  Hz, 2H), 7.80 (t, 2H), 7.99 (d,  $^3J = 8.5$  Hz, 4H), 10.01 (s, 1H); CV:  $E_{\text{RED}}^1 = -0.47$  V (CN),  $E_{\text{RED}}^2 = -0.79$  V,  $E_{\text{RED}}^3 = -1.13$  V.

*1,5-Di-*n*-hexyloxy-6-{5-*p*-tolylamino-4-[*p*-tolylimino]-4*H*-imidazol-2-yl}-naphthalene-2-carbonitrile (5d, Ar = 4-tolyl):* Yield 20%, deep red solid; IR  $\text{cm}^{-1}$ : 3278 (NH), 2227 (CN); MS (DEI):  $m/z$  (%): 627 ( $M^+$ , 16), 268 (100), 205 (20), 177 (18), 133 (25), 106 (72); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 345 nm (4.2), 440 (4.4), 500 (4.3), 536 (4.1);  $^1\text{H-NMR}$  (250 MHz, THF- $d_8$ )  $\delta$  (ppm): 0.89 (t, 6H), 1.29 (m, 8H), 1.40 (m, 4H), 1.96 (m, 4H), 2.37 (s, 6H), 4.46 (t, 4H), 7.24 (d,  $^3J = 8.8$  Hz, 4H), 7.60 (d,  $^3J = 8.8$  Hz, 1H), 7.70 (d,  $^3J = 8.8$  Hz, 2H), 7.98 (d,  $^3J = 8.8$  Hz, 4H), 8.70 (d,  $^3J = 8.8$  Hz, 1H); CV:  $E_{\text{RED}}^1 = -0.90$  V,  $E_{\text{RED}}^2 = -1.31$  V.

*1,5-Di-*n*-hexyloxy-6-{5-*p-n*-butylphenylamino-4-[*p-n*-butylphenylimino]-4*H*-imidazol-2-yl}-naphthalene-2-carbonitrile (5d, Ar = 4-*n*-butylphenyl):* Yield 27%, red solid; IR  $\text{cm}^{-1}$ : 3296 (NH), 2227 (CN); MS (DEI):  $m/z$  (%): 711 ( $M^+$ , 20), 483 (24), 378 (28), 352 (20), 210 (100), 149 (10), 106 (21); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 345 nm (4.2), 415 (4.3), 503 (4.3), 540 (4.2);  $^1\text{H-NMR}$  (250 MHz, THF- $d_8$ )  $\delta$  (ppm): 0.93 (t,  $^3J = 7.0$  Hz, 12H), 1.29 (m, 4H), 1.40 (m, 8H), 1.60 (m, 8H), 1.95 (m, 4H), 2.59 (t,  $^3J = 7.5$  Hz, 4H), 4.46 (t,  $^3J = 6.5$  Hz, 4H), 7.15 (d,  $^3J = 8.5$  Hz, 2H), 7.69 (m, 6H), 8.02 (d,  $^3J = 8.5$  Hz, 4H), 10.08 (s, broad, 1H); CV:  $E_{\text{RED}}^1 = -0.94$  V,  $E_{\text{RED}}^2 = -1.39$  V.

Boratetraazapentalenes **9** were prepared according to literature [2].

*9-[2-Difluoro-1,3-bis-(4-*t*-butylphenyl)-1,2-dihydro-1,3,4,6-tetraaza-2-borapentalen-5-yl]-benzo[*b*]-triphenylene-14-carbonitrile (9a, Ar = 4-*t*-butylphenyl):* Yield 52%, dark violet solid; IR  $\text{cm}^{-1}$ : 2212 (CN); MS (DEI):  $m/z$  (%): 709 ( $M^+$ , 18), 661 (6), 488 (24), 462 (26), 328 (100), 253 (14); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 372 nm (4.1), 529 (4.2), 572 (4.2), 614 (4.2);  $^1\text{H-NMR}$  (400 MHz, THF- $d_8$ )  $\delta$  (ppm): 1.33 (s, 18H), 7.42 (d,  $^3J = 8.0$  Hz, 4H), 7.40-7.65 (m, 6H), 7.85 (d,  $^3J = 8.0$  Hz, 4H), 8.02 (d,  $^3J = 8.0$  Hz, 2H), 8.62 (t, 2H), 8.67 (t, 2H); Fluorescence (THF, 475 nm):  $\lambda_{\text{max,em}}$ : 681 nm; CV:  $E_{\text{RED}}^1 = -0.48$  V,  $E_{\text{RED}}^2 = -1.16$  V,  $K_{\text{SEM}} = 3.4 \times 10^{11}$ .

9-[2-Difluoro-1,3-bis-(4-*n*-butylphenyl)-1,2-dihydro-1,3,4,6-tetraaza-2-borapentalen-5-yl]-benzo[*b*]-triphenylene-14-carbonitrile (**9a**, Ar = 4-*n*-butylphenyl): Yield 54%, dark violet solid; IR  $\text{cm}^{-1}$ : 2209 (CN); MS (DEI):  $m/z$  (%): 709 ( $M^+$ , 20), 661 (2), 451 (28), 369 (20), 328 (56), 221 (24), 111 (23), 97 (34), 85 (64), 71 (87), 57 (100); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 375 nm (4.1), 531 (4.1), 573 (4.3), 616 (4.2);  $^1\text{H-NMR}$  (400 MHz, THF- $d_8$ )  $\delta$  (ppm): 0.97 (t, 6H), 1.38 (m, 4H), 1.62 (m, 4H), 2.66 (m, 4H), 7.02-7.39 (m, 14H), 7.79 (d,  $^3J = 8.0$  Hz, 4H); Fluorescence (THF, 441 nm):  $\lambda_{\text{max,em}}$ : 699 nm, (toluene, 440nm):  $\lambda_{\text{max,em}}$ : 706 nm; CV:  $E_{\text{RED}}^1 = -0.45$  V,  $E_{\text{RED}}^2 = -1.14$  V,  $K_{\text{SEM}} = 5.0 \cdot 10^{11}$ .

12-[2-Difluoro-1,3-bis-(4-*t*-butylphenyl)-1,2-dihydro-1,3,4,6-tetraaza-2-borapentalen-5-yl]-benzo[*k*]-fluoranthene-7-carbonitrile (**9b**, Ar = 4-*t*-butylphenyl): Yield 48%, blue-violet solid; IR  $\text{cm}^{-1}$ : 2220 (CN); MS (DEI):  $m/z$  (%): 683 ( $M^+$ , 18), 635 (36), 578 (64), 302 (100), 134 (20), 91 (25), 57 (48); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 407 nm (4.3), 524 (4.4), 571 (4.3), 608 (4.1);  $^1\text{H-NMR}$  (400 MHz, THF- $d_8$ )  $\delta$  (ppm): 1.31 (s, 18H), 7.10-7.22 (m, 6H), 7.38-7.43 (m, 8H), 7.76 (d,  $J = 8.0$  Hz, 4H); Fluorescence (toluene, 405 nm):  $\lambda_{\text{max,em}}$ : 885 nm; CV:  $E_{\text{RED}}^1 = -0.88$  V,  $E_{\text{RED}}^2 = -1.50$  V,  $K_{\text{SEM}} = 3.2 \cdot 10^{10}$ .

10-[2-Difluoro-1,3-bis-(4-*t*-butylphenyl)-1,2-dihydro-1,3,4,6-tetraaza-2-borapentalen-5-yl]-anthracene-9-carbonitrile (**9c**, Ar = 4-*t*-butylphenyl): Yield 40%, dark violet solid; IR  $\text{cm}^{-1}$ : 2219 (CN); MS (EI):  $m/z$  (%): 609 ( $M^+$ , 20), 561 (84), 552 (100), 504 (78), 228 (69), 134 (71); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 388 nm (4.1), 403 (4.1), 528 (4.2), 567 (4.2), 602 (4.2);  $^1\text{H-NMR}$  (400 MHz, THF- $d_8$ )  $\delta$  (ppm): 1.30 (s, 18H), 7.41 (t, 2H), 7.54 (d,  $^3J = 8.0$  Hz, 4H), 7.76 (t, 2H), 7.80-7.95 (m, 4H), 8.08 (d,  $^3J = 8.0$  Hz, 4H); Fluorescence (THF, 462 nm):  $\lambda_{\text{max,em}}$ : 676 nm; CV:  $E_{\text{RED}}^1 = -0.42$  V,  $E_{\text{RED}}^2 = -1.05$  V,  $K_{\text{SEM}} = 4.8 \cdot 10^{10}$ .

1,5-Di-*n*-hexyloxy-6-[2-Difluoro-1,3-bis-(4-tolyl)-1,2-dihydro-1,3,4,6-tetraaza-2-borapentalen-5-yl]-naphthalene-2-carbonitrile (**9d**, Ar = 4-tolyl): Yield 52%, dark violet solid; IR  $\text{cm}^{-1}$ : 2227 (CN); MS (DEI):  $m/z$  (%): 678 ( $M^+$ , 21), 627 (38), 268 (100), 205 (20), 177 (18), 133 (25), 106 (72); UV/vis (THF)  $\lambda_{\text{max}}$  (log  $\epsilon$ ): 352 nm (4.1), 511 (4.2), 553 (4.3), 596 (4.2);  $^1\text{H-NMR}$  (400 MHz, THF- $d_8$ )  $\delta$  (ppm): 0.91 (t, 6H), 1.32 (m, 8H), 1.44 (m, 4H), 1.99 (m, 4H), 2.42 (s, 6H), 4.50 (t, 4H), 7.36 (d,  $^3J = 8.0$  Hz, 4H), 7.81 (d,  $^3J = 8.0$  Hz, 1H), 7.93 (d,  $^3J = 8.0$  Hz, 2H), 8.18 (d,  $^3J = 8.0$  Hz, 4H), 8.90 (d,  $^3J = 8.0$  Hz, 1H); Fluorescence (THF, 394 nm):  $\lambda_{\text{max,em}}$ : 870 nm; CV:  $E_{\text{RED}}^1 = -0.82$  V,  $E_{\text{RED}}^2 = -1.47$  V,  $K_{\text{SEM}} = 1.0 \cdot 10^{11}$ .

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*Sample availability:* Most of the described substances are available from the authors.

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