

RESEARCH ARTICLE

View Article Online

View Journal | View Issue



Cite this: *Org. Chem. Front.*, 2017, 4, 178

Copper-assisted azide–alkyne cycloaddition chemistry as a tool for the production of emissive boron difluoride 3-cyanoformazanates†

Stephanie M. Barbon,^{a,b} Samantha Novoa,^{a,b} Desiree Bender,^{a,b} Hilary Groom,^c Leonard G. Luyt^{*a,d,c} and Joe B. Gilroy^{*a,b}

The synthesis and characterization of emissive boron difluoride (BF₂) complexes of 3-cyanoformazanate ligands produced using copper-assisted azide–alkyne cycloaddition (CuAAC) chemistry is described. Detailed spectroscopic and electrochemical characterization of benzyl-functionalized complexes served as models and demonstrated that triazole formation at the *N*-aryl substituents of the formazanate ligand scaffold led to red-shifted absorption and emission maxima and more difficult electrochemical reduction compared to alkyne-substituted precursors. CuAAC chemistry was also used to append ferrocene and tetraethylene glycol substituents to the formazanate backbone. In the case of the ferrocene-substituted complexes, fluorescence was quenched and a reversible oxidation feature (in addition to the reduction features associated with formazanate complexes) was observed using cyclic voltammetry. Treatment with NOBF₄ oxidized ferrocene to ferrocenium and resulted in the reestablishment of fluorescence. Tetraethylene glycol substitution produced the first water soluble BF₂ formazanate, which was shown to distribute throughout the cytoplasm and nucleus of mouse fibroblast cells when studied as a fluorescence imaging agent.

Received 18th October 2016,
Accepted 2nd November 2016

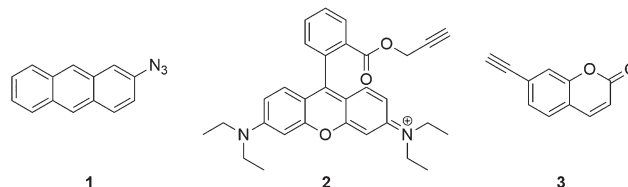
DOI: 10.1039/c6qo00640j

rsc.li/frontiers-organic

Introduction

Fluorophores are a very useful class of molecules as they provide a handle that allows for the visualization of chemical phenomena at the molecular, macromolecular and bulk size regimes.^{1–6} For this and other reasons, the development of new classes of fluorophores has been the focus of a significant amount of research.^{7,8} One of the major hurdles to be overcome in the field was the facile synthetic modification of fluorophores in order to utilize them for an expanded range of applications.^{9–11} Significant impact was made in this area with the reports on copper-assisted azide–alkyne cycloaddition

(CuAAC) chemistry.^{12,13} Since the original reports appeared, the number of alkyne- and azide-modified fluorophores such as those based on anthracene,^{14,15} pyrene,^{16,17} boron dipyrromethenes (BODIPYs),^{18–20} rhodamine^{21,22} and coumarin^{23,24} has expanded dramatically (*e.g.*, 1–3). The onward CuAAC chemistry of these and other fluorophores has led to their use, for example, as protein labels,²⁵ in photosensitizers,²⁶ as molecular switches,²⁷ as probes of DNA hybridization,²⁸ and in radiolabeling.²⁹ CuAAC chemistry has also been used to create a variety of fluorescence sensors for anions,³⁰ acids and bases,^{31,32} nitroxide radicals,³³ hydroxyl radicals,³⁴ and various metal ions including mercury^{35,36} and zinc.^{37,38}



In addition, CuAAC chemistry has been utilized to modify fluorophores for biological imaging applications. This strategy has been used both to impart water solubility (*e.g.*, 4),^{39,40} and functionalities that interact with specific biological targets (*e.g.*, 5).^{41–44} More recently, this strategy has been extended to include copper-free, strain-promoted azide–alkyne cyclo-

^aDepartment of Chemistry, The University of Western Ontario, 1151 Richmond St. N., London, Ontario, Canada, N6A 5B7. E-mail: lluyt@uwo.ca, joe.gilroy@uwo.ca; Tel: +1-519-661-2111 ext. 81561 (JBG), +1-519-685-8600 ext. 53302 (LGL)

^bThe Centre for Advanced Materials and Biomaterials Research (CAMBR), The University of Western Ontario, 1151 Richmond St. N., London, Ontario, Canada, N6A 5B7

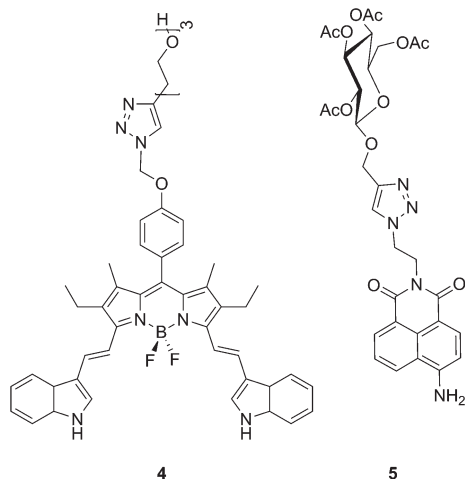
^cDepartment of Oncology, London Regional Cancer Program, 790 Commissioners Rd. E., London, Ontario, Canada, N6A 4L6

^dDepartment of Medical Imaging, The University of Western Ontario, 1151 Richmond St. N., London, Ontario, Canada, N6A 5B7

†Electronic supplementary information (ESI) available. CCDC 1500180. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c6qo00640j



addition chemistry with specific utility in biological systems.^{45,46}



Transition metal^{47–53} and boron^{54–56} complexes of formazanate ligands have received significant attention due to their rich optical and electrochemical properties. More specifically, boron difluoride (BF_2) complexes of 3-cyanoformazanates have recently been shown to have many of the desirable properties associated with common fluorophores, including high fluorescence quantum yields, large Stokes shifts, facile high-yielding syntheses, and perhaps most importantly, emission in the red to near-IR region of the electromagnetic spectrum.^{57–59} These complexes have shown promise as fluorescence cell-imaging agents,⁵⁸ as precursors to $\text{B}(1)$ -carbenoid intermediates⁶⁰ and stable radicals,⁶¹ as efficient electrochemiluminescence emitters,⁶² aggregation-induced emission luminogens (AIEgens),⁶³ and building blocks for functional polymeric materials.^{64,65} Herein, we expand the scope of BF_2 formazanate

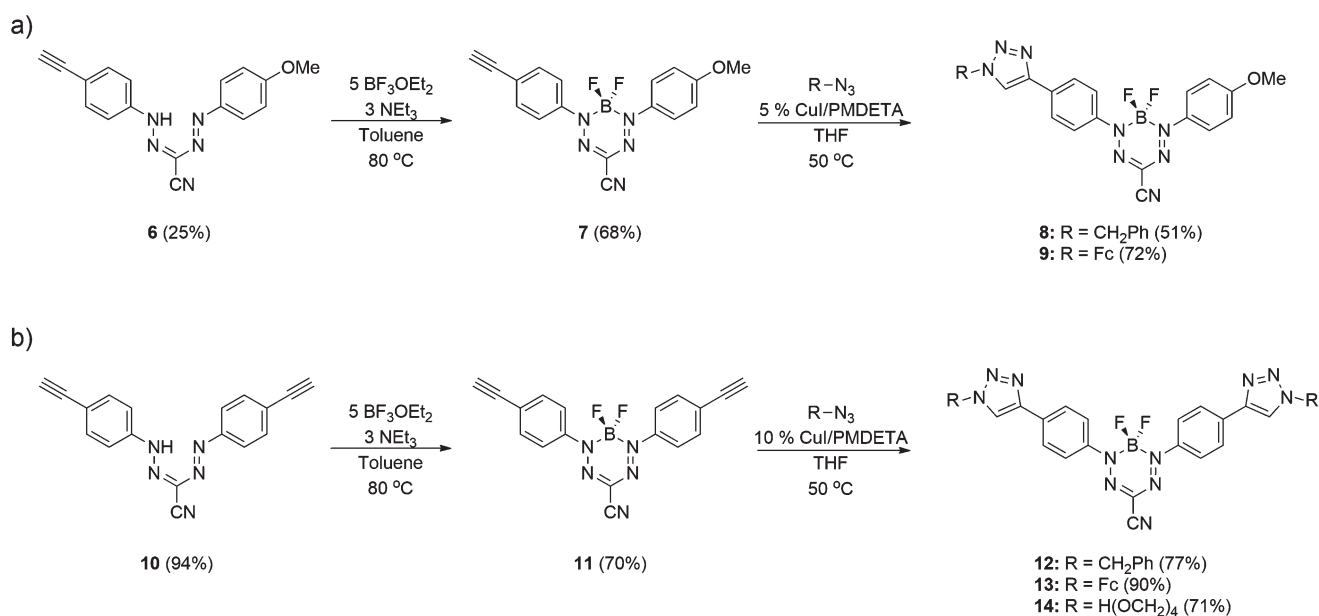
chemistry further by producing a variety of emissive materials through the use of CuAAC chemistry that cannot be readily synthesized by other methods.

Results and discussion

Benzyl-functionalized BF_2 complexes of 3-cyanoformazanates

Formazans **6** and **10** were prepared *via* aryl diazonium coupling reactions in basic solutions of cyanoacetic acid by adapting a reported method.⁶⁶ Formazan **6** was prepared from a mixture of alkyne- and methoxy-substituted aryl diazonium salts and separated from the symmetrically-substituted byproducts using column chromatography. In order to study the effect of triazole formation on the photophysical properties of BF_2 complexes of 3-cyanoformazanate ligands, formazans **6** and **10** were converted to mono-alkyne-substituted complex **7** and bis-alkyne-substituted complex **11** by adaptation of established methods (Scheme 1).⁵⁷ Complex **7** was first characterized by single crystal X-ray diffraction (Fig. 1), and crystallized on a two-fold rotation axis. As a result, the alkyne and methoxy substituents were disordered. However, the structure does confirm the expected connectivity and planarity throughout the molecule (dihedral angle between N_4 plane and aryl substituents: 7.2°). All other structural metrics were consistent with the solid-state structures of previously reported BF_2 adducts of 3-cyanoformazanate ligands.⁵⁷

The alkyne-substituted complexes **7** and **11** underwent CuAAC chemistry with benzyl azide when combined in the presence of copper(I) iodide and N,N,N',N'',N''' -pentamethyldiethylenetriamine (PMDETA) in THF to form mono- and bis-benzyl-functionalized compounds **8** and **12** (Scheme 1). The successful conversion of alkyne-substituted complexes **7** and



Scheme 1 Synthesis of (a) mono- and (b) bis-functionalized BF_2 complexes of 3-cyanoformazanates.



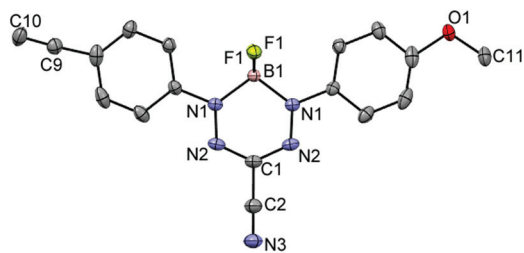


Fig. 1 Solid-state structure of **7**. Anisotropic displacement ellipsoids are shown at 50% probability and hydrogen atoms have been removed for clarity. Selected bond lengths (Å): B1–N1 1.5759(15), N1–N2 1.3023(15), N2–C1 1.3351(13), O1–C11 1.418(5), C9–C10 1.191(7). Selected bond angles (°): B1–N1–N2 124.57(10), N1–N2–C1 117.24(11).

11 to benzyl-functionalized complexes **8** and **12** was confirmed by ^1H , ^{11}B , $^{13}\text{C}\{^1\text{H}\}$ and ^{19}F NMR spectroscopy (Fig. 2 and S1–S17[†]), UV-vis absorption/emission and IR spectroscopy, and high resolution mass spectrometry. CuAAC chemistry was also used to synthesize BF_2 formazanate complexes with appended ferrocene substituents (**9** and **13**) and water-solubil-

izing tetraethylene glycol substituents (**14**). Initial attempts to synthesize the water-soluble complex **14** were unsuccessful, likely due to the TEG chains outcompeting PMDETA for the ligation of copper(I) when all of the reagents were combined simultaneously. We circumvented this issue by stirring CuI and PMDETA in THF for 30 min before the TEG azide was added.

The electrochemical properties of complexes **7**, **8**, **11** and **12** were studied using cyclic voltammetry in THF (Fig. 3 and Table 1). Each voltammogram showed two reversible one-electron reduction waves within the electrochemical window of the solvent. Considering the first reduction wave, which corresponds to the formation of a ligand-centered radical anion, bis-alkyne-substituted complex **11** is significantly easier to reduce than mono-alkyne-substituted complex **7** (−0.50 V and −0.67 V relative to the ferrocene/ferrocenium redox couple, respectively) due to the presence of the electron donating methoxy substituent in **7**. The benzyl-functionalized complexes were more difficult to reduce than their alkyne-substituted precursors (**8**: −0.68 V; **12**: −0.62 V). We have previously demonstrated that extending electronic conjugation in BF_2 complexes of 3-cyanoformazanate ligands results in complexes that are easier to reduce due to a narrowing of the HOMO–LUMO gap.⁵⁹ However, in the case of complexes **8** and **12**, the more negative reduction potentials observed are likely due to the electron donating nature of the benzyl-substituted triazoles, which appear to be better electron donors than alkyne substituents.

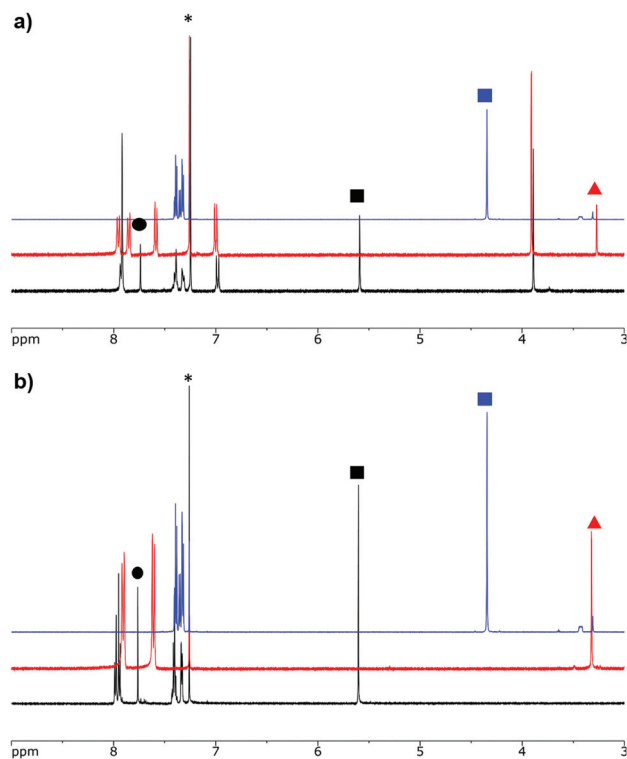


Fig. 2 ^1H NMR spectra of (a) benzyl azide (blue), mono-alkyne-substituted complex **7** (red) and mono-benzyl-functionalized complex **8** (black) and (b) benzyl azide (blue), bis-alkyne-substituted complex **11** (red) and bis-benzyl-functionalized complex **12** (black) in CDCl_3 . The asterisks denote residual solvent signals. The blue squares denote the benzyl azide CH_2 signals, the red triangles denote the alkyne CH signals, the black circles denote the triazole CH signals, and the black squares denote the CH_2 signal from the benzyl azide moieties in complexes **8** and **12**.

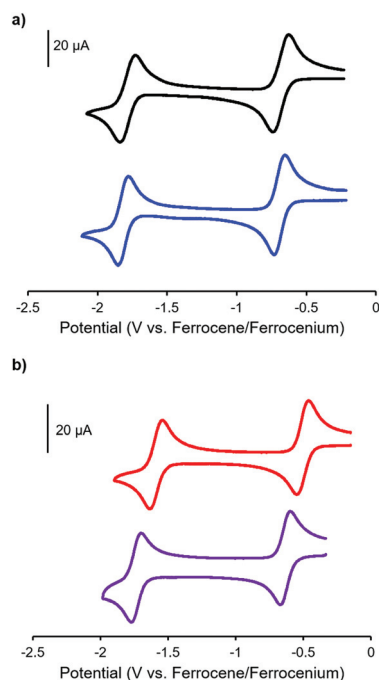


Fig. 3 Cyclic voltammograms of (a) mono-alkyne-substituted complex **7** (black) and mono-benzyl-functionalized complex **8** (blue) and (b) bis-alkyne-substituted complex **11** (red) and bis-benzyl-functionalized complex **12** (purple) recorded at 100 mV s^{-1} in 1 mM THF solutions containing 0.1 M $[\text{nBu}_4\text{N}][\text{PF}_6]$ as supporting electrolyte.



Table 1 Optical and electrochemical properties of BF₂ formazanate complexes **7–9** and **11–14**

	Solvent	λ_{\max} (nm)	ϵ (M ⁻¹ cm ⁻¹)	λ_{em} (nm)	Φ_{F}^a (%)	ν_{ST} (nm)	ν_{ST} (cm ⁻¹)	E_{ox1}^b (V)	E_{red1}^b (V)	E_{red2}^b (V)
7	CH ₂ Cl ₂	552	30 400	647	30	95	2660	—	-0.67	-1.78
	THF	550	32 500	650	27	100	2797			
	Toluene	569	26 800	648	36	79	2143			
8	CH ₂ Cl ₂	560	31 000	660	30	100	2706	—	-0.68	-1.81
	THF	560	33 700	672	25	112	2976			
	Toluene	579	37 400	661	37	82	2143			
9^c	CH ₂ Cl ₂	562	28 600	—	—	—	—	0.20	-0.67	-1.80
	THF	562	31 700	—	—	—	—			
	Toluene	578	23 900	—	—	—	—			
11	CH ₂ Cl ₂	530	32 700	635	18	105	3120	—	-0.50	-1.58
	THF	529	29 600	638	19	109	3230			
	Toluene	547	41 700	637	36	90	2583			
12	CH ₂ Cl ₂	558	39 600	679	46	121	3194	—	-0.62	-1.72
	THF	563	34 100	696	34	133	3394			
	Toluene	577	33 700	680	72	103	2625			
13^c	CH ₂ Cl ₂	569	30 900	—	—	—	—	0.21	-0.56	-1.67
	THF	569	26 100	—	—	—	—			
	Toluene	582	30 700	—	—	—	—			
14	CH ₂ Cl ₂	542	16 500	680	53	138	3744	—	-0.58	-1.68
	THF	563	11 900	697	20	134	3415			
	Toluene	—	—	—	—	—	—			

^aQuantum yields were measured according to published protocols using [Ru(bpy)₃][PF₆]₂ as a relative standard and corrected for wavelength-dependent detector sensitivity (Fig. S18).^{68,69} ^bCyclic voltammetry experiments were conducted in THF containing 1 mM analyte and 0.1 M [nBu₄N][PF₆] as supporting electrolyte at a scan rate of 100 mV s⁻¹. All voltammograms were referenced internally against the ferrocene/ferrocenium redox couple. ^cComplexes **9** and **13** were non-emissive in the range of organic solvents employed.

The second reduction waves, which correspond to the formation of ligand-centred dianions, followed the same trend.

In both the UV-vis absorption and emission spectra, the wavelength of maximum absorption (λ_{\max}) and emission (λ_{em}) red-shift by approximately 10 nm when comparing mono-benzyl-functionalized complex **8** and mono-alkyne-substituted complex **7**. As may be expected, a more drastic red-shift was observed when the spectra of bis-benzyl-functionalized complex **12** and bis-alkyne-substituted complex **11** were compared. In this case, red-shifts of approximately 30 nm (absorption) and 40 nm (emission) were observed (Fig. 4 and Table 1).

Complexes **7**, **8**, **11** and **12** exhibit moderate fluorescence quantum yields in THF, CH₂Cl₂, and toluene (Table 1). Mono-alkyne-substituted complex **7** has a quantum yield of 30% in CH₂Cl₂, which is almost double that of bis-alkyne-substituted complex **11** (Φ_{F} = 18% in CH₂Cl₂), likely due to its asymmetric structure.⁶⁷ Mono-benzyl-functionalized complex **8** has the same quantum yield as complex **7**, while bis-benzyl-functionalized complex **12** showed the most intense emission at a quantum yield of 46%. The Stokes shifts observed (79–133 nm; 2143–3394 cm⁻¹) for these complexes were typical of other BF₂ complexes of 3-cyanoformazanates.⁵⁷

Ferrocene-functionalized BF₂ complexes of 3-cyanoformazanates

Complexes **9** and **13**, which bear one or two ferrocene substituents, are dark purple/blue (Fig. S19 and S20†) and non-

emissive in solution. The lack of emission is likely due to excited-state electron transfer from the BF₂ formazanate moieties to the ferrocenes.^{70–73} In order to further explore the apparent fluorescence quenching, we sequentially oxidized the ferrocene units in **9** and **13** and studied their absorption and emission properties in CH₂Cl₂ (Fig. 5). NOBF₄ was chosen as an oxidant as the gaseous byproducts were assumed not to contribute to the spectra collected upon oxidation and the oxidation potential (*ca.* 1.0 V relative to the ferrocene/ferrocenium redox couple)⁷⁴ was not in a range that would oxidize the 3-cyanoformazanate ligand backbone. In CH₂Cl₂, the absorption maxima associated with the $\pi \rightarrow \pi^*$ transition of the BF₂ formazanate backbone of complexes **9** and **13** were blue-shifted, by up to 20 nm in complex **13**, upon sequential addition of NOBF₄. The blue-shifts appear to arise due to the removal of electron density from the ferrocene moieties upon oxidation, rendering the complexes less electron rich overall. This effect has been observed previously for similar complexes, whereby the introduction of electron withdrawing groups blue-shifted absorption maxima.⁵⁷ Perhaps more significant, was the observation that, upon conversion of ferrocene to ferrocenium, the solutions of **9** and **13** that had been treated with NOBF₄ became emissive. The emission intensity gradually increased with the subsequent addition of oxidizing agent until a maximum intensity (Φ_{F} = 7% for **9** and 14% for **13**) was reached when a full equivalent of NOBF₄ per ferrocene unit



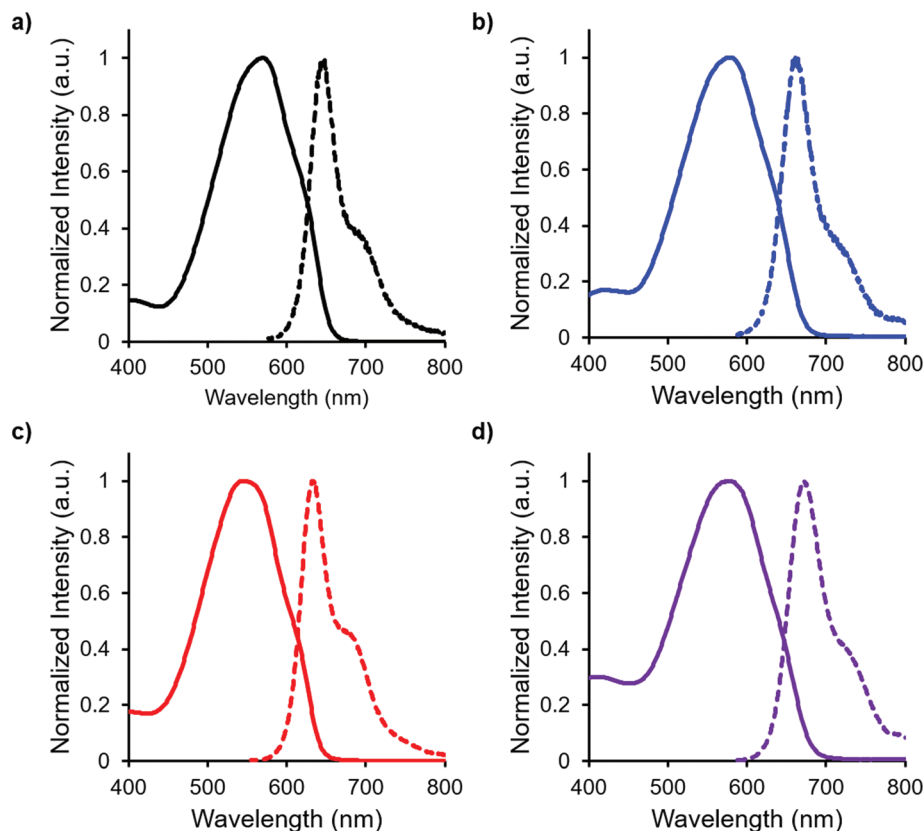


Fig. 4 UV-vis absorption spectra (solid line) and emission spectra (dashed line) of (a) mono-alkyne-substituted complex **7**, (b) mono-benzyl-functionalized complex **8**, (c) bis-alkyne-substituted complex **11** and (d) bis-benzyl-functionalized complex **12**, recorded for 10^{-5} M toluene solutions.

had been added, while λ_{em} was unchanged. These observations corroborate our hypothesis that the emission of these complexes was quenched as a result of excited-state electron transfer from the formazanate backbone to ferrocene.

BF_2 complexes of formazanate ligands generally exhibit very well-behaved reduction chemistry.^{56–61} However, their oxidation is rarely observed within the electrochemical window of most organic solvents. Ferrocene can be reversibly oxidized, and so the redox properties of compounds that contain both ferrocene and formazanate moieties may have unusual properties, including charge-transfer characteristics. Similar compounds based on different fluorophores have been previously used as viscosity probes,⁷⁵ redox-active fluorescent switches,^{70,76} and ion pair recognition receptors.⁷⁷

Cyclic voltammetry studies of complexes **9** and **13** revealed two characteristic BF_2 formazanate reduction waves, and indicated that each complex was more difficult to reduce than the alkyne-substituted precursors (**7** and **11**), due to the electron-rich nature of ferrocene (Table 1). Oxidation waves corresponding to one electron for **9** and two electrons for **13** were observed at potentials of 0.20 V and 0.21 V, which corresponds to the oxidation of the ferrocene groups (Fig. 6). The coincident appearance of the ferrocene waves in the cyclic voltammogram of **13** confirmed that there was little to no electronic communication between the ferrocene units *via* the BF_2

formazanate spacer. This observation was consistent with those noted for polymers and model compounds derived from BF_2 complexes of triarylformazanate ligands, whereby triazole groups were shown to limit the overall degree of electronic conjugation along the polymer backbone.⁶⁵

Tetraethylene glycol-functionalized BF_2 complexes of 3-cyanoformazanates

Complex **14**, which was targeted in an effort to maximize hydrophilicity, is soluble in both polar organic solvents and water. Interestingly, the optical properties were highly dependent on the polarity of the solvent used. In relatively non-polar solvents such as CH_2Cl_2 and THF, **14** was moderately fluorescent (Φ_F : 53% and 20%, respectively). However, the fluorescence intensity dropped off significantly in more polar solvents (Φ_F : 5% in H_2O), and there was also a blue-shift in the wavelength of maximum emission (Table 2, Fig. 7), as is often observed for fluorescent dyes in highly polar solvents.^{78–82}

While the emission intensity of **14** was relatively low in water, many other fluorophores with relatively low quantum yields have shown promise as cell-imaging agents, especially dyes which emit in the red to near-IR region.^{83–87} With these factors in mind, complex **14** was introduced into mouse fibroblast cells and its utility as an imaging agent studied using fluorescence confocal microscopy.



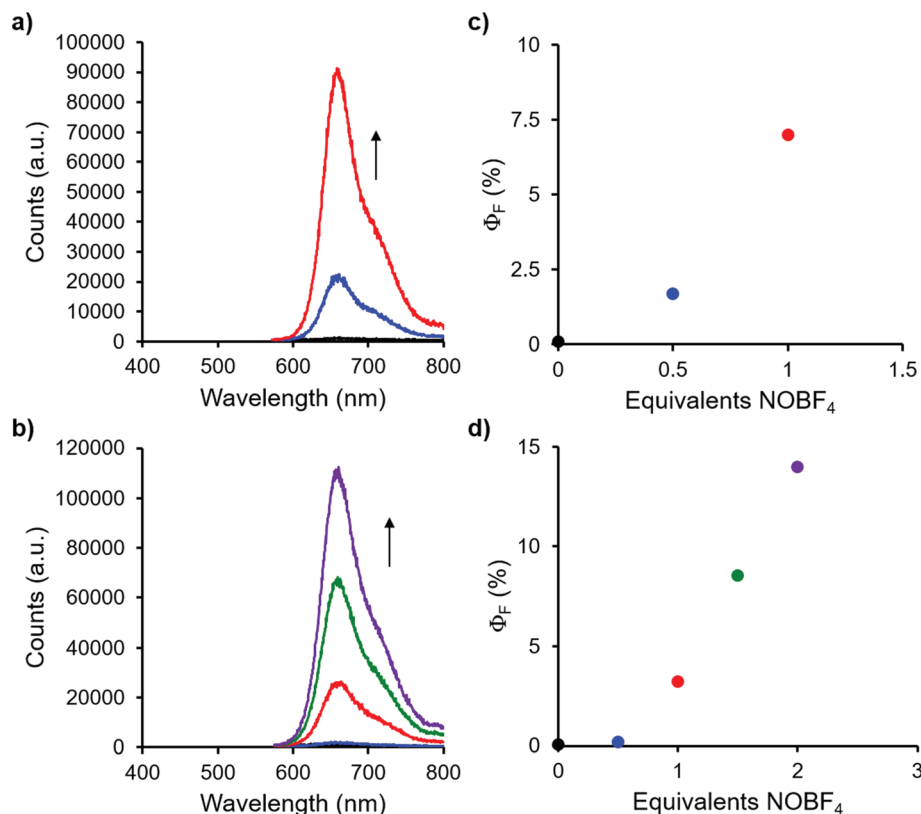


Fig. 5 Emission spectra of solutions of ferrocene-substituted complexes (a) **9** and (c) **13** treated with 0 equiv. (black), 0.5 equiv. (blue), 1.0 equiv. (red), 1.5 equiv. (green) and 2.0 equiv. (purple) of NOBF₄, recorded in degassed CH₂Cl₂ at a concentration of 10^{−5} M. The arrows indicate the trends upon addition of NOBF₄. The corresponding quantum yields for each solution are plotted in panels (b) **9** and (d) **13**.

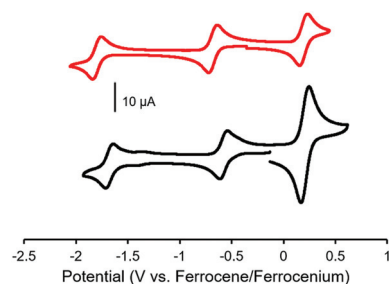


Fig. 6 Cyclic voltammograms of mono-ferrocene substituted BF₂ complex **9** (red) and bis-ferrocene substituted complex **13** (black) recorded at 100 mV s^{−1} in 1 mM THF solutions containing 0.1 M [nBu₄N][PF₆] as supporting electrolyte.

Table 2 Optical properties of complex **14** in different solvents

Solvent	λ _{max} (nm)	ε (M ^{−1} cm ^{−1})	λ _{em} (nm)	Φ _F ^a (%)	ν _{ST} (nm)	ν _{ST} (cm ^{−1})
CH ₂ Cl ₂	542	16 500	680	53	138	3744
THF	563	11 900	697	20	134	3415
MeOH	521	19 900	681	6	160	4510
MeCN	524	18 200	697	10	173	4737
DMSO	526	17 600	716	<1	190	5045
H ₂ O	527	15 100	698	5	171	4649

^aQuantum yields were measured according to published protocols using [Ru(bpy)₃][PF₆]₂ as a relative standard and corrected for wavelength-dependent detector sensitivity (Fig. S18).^{68,69}

Complex **14** was successfully incorporated into the cells, as evidenced by the fluorescence micrographs shown in Fig. 8. Specifically, complex **14** was distributed throughout the cell structure and clearly penetrated the cell nucleus. The dark spots visualized in Fig. 8c are believed to be DNA-free nucleoli, and appear to be the only features of the cells that were not stained by **14**. Despite the widespread incorporation of complex **14** throughout the cell, Fig. 8b and d demonstrate our ability to differentiate between cytoplasm and nucleus when 4',6-diamidino-2-phenylindole (DAPI), a selective nuclear stain,

was employed concurrently. These results differed significantly when compared to previous studies involving BF₂ complexes of 3-cyanoformazanates bearing *p*-anisole substituents.⁵⁸ The hydrophobic nature of the anisole-based dye resulted in selective staining of the cell cytoplasm and required the use of DMSO to achieve cellular uptake.⁵⁸ While we have not produced a site-specific imaging agent as part of this study, we have demonstrated our ability to create hydrophilic analogs of the parent complexes and shown that, in doing so, we can drastically alter their cellular uptake. This work sets the stage for future iterations whereby site-specific staining will be



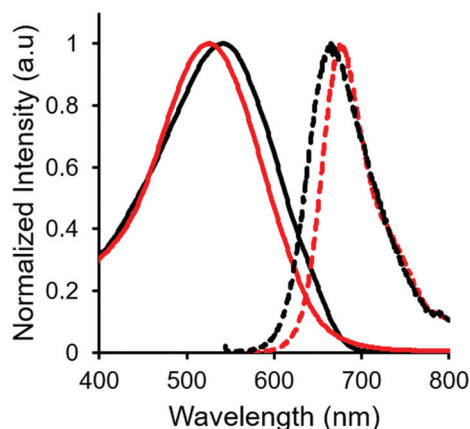


Fig. 7 UV-vis absorption (solid line) and emission (dashed line) spectra of TEG-functionalized complex **14** in CH_2Cl_2 (black) and H_2O (red), recorded for degassed solutions at a concentration of 10^{-5} M.

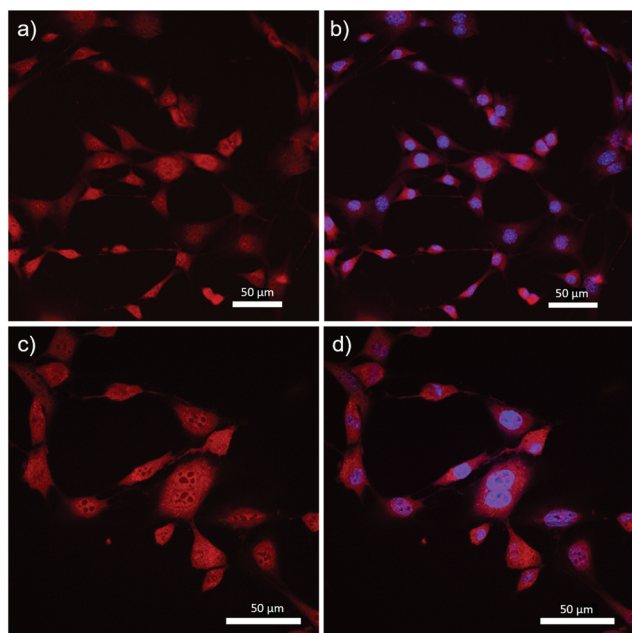


Fig. 8 Confocal fluorescence micrographs of mouse fibroblast cells stained with TEG-functionalized complex **14** and DAPI. Images (a) and (c) were visualized with excitation at 579 nm and emission collected between 620–720 nm. Images (b) and (d) are an overlay of images (a) and (c) with those obtained from excitation at 405 nm and emission collected between 425–475 nm.

achieved through the introduction of hydrophilic, tailor-made peptide chains.

Conclusions

As a result of this work, we have demonstrated the effect of triazole formation on the spectroscopic and electrochemical properties of BF_2 complexes of 3-cyanoformazanates. Complexes

functionalized with benzyl groups showed that the formation of triazole rings resulted in red-shifted wavelengths of maximum absorption and emission, as well as increased fluorescence quantum yields. Triazole formation also resulted in BF_2 complexes that were more difficult to reduce electrochemically. We further demonstrated how CuAAC chemistry could be used to expand the scope of BF_2 formazanate chemistry. The introduction of ferrocene into the BF_2 formazanate scaffold *via* CuAAC resulted in the formation of a non-emissive complex with rich electrochemistry. Specifically, oxidation waves corresponding to one electron per ferrocene and two reduction waves (one electron each) originating from the formazanate backbones were observed by cyclic voltammetry. We also demonstrated that the stepwise conversion of ferrocene to ferrocenium was accompanied by an increase in emission intensity, confirming our hypothesis that quenching arose due to excited-state electron transfer from the formazanate backbone to ferrocene. Finally, CuAAC chemistry was used to append water-solubilizing TEG chains to a BF_2 formazanate complex. The resulting complex was used to image mouse fibroblast cells, and demonstrated our ability to control cellular uptake *via* the modification of hydrophobicity/hydrophilicity.

Experimental section

General considerations

Reactions and manipulations were carried out under a nitrogen atmosphere using standard Schlenk techniques unless otherwise stated. Solvents were obtained from Caledon Laboratories, dried using an Innovative Technologies Inc. solvent purification system, collected under vacuum and stored under a nitrogen atmosphere over 4 Å molecular sieves. Reagents were purchased from Sigma-Aldrich or Alfa Aesar and used as received. TEG-N_3 ⁸⁸ and Fc-N_3 ⁸⁹ were prepared according to literature procedures.

NMR spectra were recorded on 400 MHz (^1H : 399.8 MHz, ^{11}B : 128.3 MHz, ^{19}F : 376.1 MHz) or 600 MHz (^1H : 599.5 MHz, $^{13}\text{C}\{^1\text{H}\}$: 150.8 MHz) Varian INOVA instruments. ^1H NMR spectra were referenced to residual CHCl_3 (7.26 ppm) or $\text{DMSO-}d_6$ (2.50 ppm) and $^{13}\text{C}\{^1\text{H}\}$ NMR spectra were referenced to CDCl_3 (77.2 ppm) or $\text{DMSO-}d_6$ (39.5 ppm). ^{11}B spectra were referenced to $\text{BF}_3\cdot\text{OEt}_2$ at 0 ppm and ^{19}F spectra were referenced to CFCl_3 at 0 ppm. Mass spectrometry data were recorded in positive-ion mode using a high-resolution Finnigan MAT 8200 spectrometer using electron impact ionization or a Micromass LCT electrospray time-of-flight mass spectrometer. UV-vis absorption spectra were recorded using a Cary 5000 instrument. Four separate concentrations were run for each sample and molar extinction coefficients were determined from the slope of a plot of absorbance against concentration. FT-IR spectra were recorded on a KBr disk or using an attenuated total reflectance (ATR) attachment using a Bruker Vector 33 FT-IR spectrometer. Emission spectra were obtained using a Photon Technology International QM-4 SE spectrofluorometer. Excitation wavelengths were chosen based on



λ_{\max} from the respective UV-vis absorption spectrum in the same solvent. Emission quantum yields were estimated relative to $[\text{Ru}(\text{bpy})_3][\text{PF}_6]_2$ and corrected for wavelength dependent detector sensitivity (Fig. S18†).^{68,69}

Electrochemical methods

Cyclic voltammetry experiments were performed with a Bioanalytical Systems Inc. (BASi) Epsilon potentiostat and analyzed using BASi Epsilon software. Electrochemical cells consisted of a three-electrode setup including a glassy carbon working electrode, platinum wire counter electrode and silver wire *pseudo* reference electrode. Experiments were run at scan rates of 100 mV s⁻¹ in degassed THF solutions of the analyte (~1 mM) and supporting electrolyte (0.1 M *n*Bu₄PF₆). Cyclic voltammograms were referenced against an internal standard (~1 mM ferrocene) and corrected for internal cell resistance using the BASi Epsilon software.

X-ray crystallography details

Single crystals of mono-alkyne-substituted complex 7 suitable for X-ray diffraction studies were grown by slow evaporation of a concentrated CH₂Cl₂ solution. The sample was mounted on a MiTeGen polyimide micromount with a small amount of Paratone N oil. X-ray measurements were made on a Bruker Kappa Axis Apex2 diffractometer at a temperature of 110 K. The data collection strategy included a number of ω and φ scans which collected data over a range of angles, 2θ . The frame integration was performed using SAINT.⁹⁰ The resulting raw data was scaled and absorption corrected using a multi-scan averaging of symmetry equivalent data using SADABS.⁹¹ The structure was solved by using a dual space methodology using the SHELXT program.⁹² All non-hydrogen atoms were obtained from the initial solution. The hydrogen atoms were introduced at idealized positions and were allowed to refine isotropically. The structural model was fit to the data using full matrix least-squares based on F^2 . The calculated structure factors included corrections for anomalous dispersion from the usual tabulation. The structure was refined using the SHELXL-2014 program from the SHELXT suite of crystallographic software.⁹³ See Table 3 and CCDC 1500180 for additional crystallographic data.

Cell imaging studies

Mouse fibroblast C3H/10T1/2 cells (ATCC) were released from the tissue culture flask and seeded onto cover slips in a 12-well tissue culture plate at approximately 50 000 cells per well. The cells were incubated overnight in Dulbecco's modified Eagle's medium (DMEM, Sigma-Aldrich) containing 10% fetal bovine serum (FBS, Sigma-Aldrich) with penicillin streptomycin at 37 °C in a 5% CO₂ atmosphere. Cell media was aspirated and the cells were washed once with serum free media and twice with phosphate buffer saline (PBS, Sigma-Aldrich). Water-soluble complex 14 (8 mg) was dissolved into 5 mL DMEM (serum free), filter sterilized with a 0.2 μm filter and diluted to 100 μM with serum free DMEM. The dye stock was incubated at 37 °C for 1 h with the cells. The cells were washed twice

Table 3 X-ray diffraction data collection and refinement details for mono-alkyne-substituted complex 7

	7
Chemical formula	C ₁₇ H ₁₂ BF ₂ N ₅ O
FW (g mol ⁻¹)	351.13
Crystal habit	Purple prism
Crystal system	Monoclinic
Space group	C2/c
<i>T</i> (K)	110
λ (Å)	0.71073
<i>a</i> (Å)	10.248(6)
<i>b</i> (Å)	14.634(7)
<i>c</i> (Å)	11.146(5)
α (°)	90
β (°)	107.352(18)
γ (°)	90
<i>V</i> (Å ³)	1595.5(14)
<i>Z</i>	4
ρ (g cm ⁻³)	1.462
μ (cm ⁻¹)	0.111
<i>R</i> ₁ ^a , <i>wR</i> ₂ ^b [<i>I</i> > 2 σ]	0.0428, 0.0963
<i>R</i> ₁ , <i>wR</i> ₂ (all data)	0.0770, 0.1107
GOF ^c	1.038

$$^a R_1 = \sum(|F_o| - |F_c|)/\sum F_o, ^b wR_2 = [\sum(w(F_o^2 - F_c^2)^2)/\sum(wF_o^4)]^{1/2}, ^c GOF = [\sum(w(F_o^2 - F_c^2)^2)/(\text{no. of reflns.} - \text{no. of params.})]^{1/2}.$$

with PBS, fixed with 4% paraformaldehyde and mounted onto slides containing Pro-Long Antifade mounting medium with DAPI (ThermoFisher Scientific). Fluorescence microscopy images were obtained using an Olympus FluoView FV 1000 confocal microscope. Images based on the fluorescence of DAPI were obtained using 2% laser strength, with excitation at 405 nm and emission collected between 425 and 475 nm. Images based on the fluorescence of complex 14 were obtained using 10% laser strength, with excitation at 579 nm and emission collected between 620 and 720 nm.

Mono-alkyne-substituted formazan 6

In air, cyanoacetic acid (1.00 g, 11.7 mmol) and NaOH (4.70 g, 117 mmol) were mixed with deionized H₂O (60 mL) and the solution was stirred in an ice bath for 20 min. Meanwhile, in a separate flask, 4-ethynylaniline (1.10 g, 9.36 mmol) was mixed with 12 M HCl (2.3 mL, 28 mmol) in deionized H₂O (2.3 mL). The solution was cooled in an ice bath for 15 min before a cooled solution of sodium nitrite (0.75 g, 10 mmol) in deionized H₂O (7 mL) was added dropwise. The resulting reaction mixture, which contained the corresponding diazonium salt, was stirred in an ice bath for an additional 20 min. In a separate flask, *p*-anisidine (1.4 g, 12 mmol) was mixed with 12 M HCl (2.9 mL, 35 mmol) in deionized H₂O (2.9 mL). The solution was cooled in an ice bath for 15 min before a cooled solution of sodium nitrite (0.93 g, 13 mmol) in deionized H₂O (8 mL) was added dropwise. The resulting reaction mixture, which contained diazonium salt, was stirred in an ice bath for an additional 20 min. The diazonium salt solutions were then mixed together and stirred in an ice bath for 10 min. This solution was then added dropwise to the cyanoacetic acid solution described above. The solution turned dark red after approxi-



mately 2 min. After complete addition, the mixture was stirred in an ice bath for an additional 60 min before it was neutralized with 1 M HCl. The resulting red-brown solid was isolated by vacuum filtration and purified by flash chromatography using a gradient strategy (starting at 1:1 *n*-hexanes:CH₂Cl₂ and ending with 2:8 *n*-hexanes:CH₂Cl₂) where the second coloured fraction contained the desired product. Removal of the solvent *in vacuo* afforded mono-alkyne-substituted formazan **6** as a dark red microcrystalline solid. Yield = 0.89 g, 25%. M.p. 212–213 °C. ¹H NMR (400.1 MHz, DMSO-*d*₆): δ 12.49 (br s, 1H, NH), 8.08 (d, ³J_{HH} = 9 Hz, 2H, aryl CH), 7.54 (d, 4H, aryl CH), 7.18 (d, ³J_{HH} = 9 Hz, 2H, aryl CH), 4.19 (s, 1H, alkyne CH), 3.91 (s, 3H, OCH₃). ¹³C{¹H} NMR (100.6 MHz, DMSO-*d*₆): δ 163.2, 145.9, 142.5, 132.8, 126.9, 126.2, 116.6, 115.5, 114.8, 112.6, 83.3, 80.5, 55.7. FT-IR(ATR): 3308 (m), 3230 (m), 2942 (s), 2837 (s), 2224 (m), 2099 (m), 1605 (m), 1579 (m), 1514 (s), 1249 (s), 1183 (s), 1164 (s), 1140 (s), 1110 (m), 1028 (m) cm⁻¹. UV-Vis (CH₂Cl₂): λ_{max} 444 nm (ε = 26 400 M⁻¹ cm⁻¹). Mass Spec. (EI, +ve mode): exact mass calculated for [C₁₇H₁₃N₅O]⁺: 303.1120; exact mass found: 303.1111; difference: -3.0 ppm.

Mono-alkyne-substituted BF₂ complex 7

Formazan **6** (0.40 g, 1.3 mmol) was dissolved in dry toluene (75 mL). NEt₃ (0.40 g, 0.55 mL, 3.9 mmol) was then added slowly, and the solution was stirred for 10 min before BF₃·OEt₂ (0.94 g, 0.81 mL, 6.6 mmol) was added and the solution was heated to 80 °C with stirring for 16 h. The solution became dark purple during this time and after cooling to 22 °C, deionized H₂O (10 mL) was added to quench any excess boron-containing compounds. The toluene solution was then washed with deionized H₂O (3 × 20 mL), dried over MgSO₄, gravity filtered, and concentrated *in vacuo*. The resulting compound was purified by flash chromatography (THF, neutral alumina) to afford mono-alkyne-substituted complex **7** as a dark-purple solid. Yield = 0.31 g, 68%. M.p. 199–201 °C. ¹H NMR (399.8 MHz, CDCl₃): δ 7.96 (d, ³J_{HH} = 9 Hz, 2H, aryl CH), 7.85 (d, ³J_{HH} = 9 Hz, 2H, aryl CH), 7.59 (d, ³J_{HH} = 9 Hz, 2H, aryl CH), 7.00 (d, ³J_{HH} = 9 Hz, 2H, aryl CH), 3.91 (s, 3H, OCH₃), 3.27 (s, 1H, alkyne CH). ¹³C{¹H} NMR (150.7 MHz, CDCl₃): δ 163.0, 142.9, 136.8, 133.0, 125.3, 124.5, 122.5, 115.0, 114.0, 82.5, 80.7, 55.8. ¹¹B NMR (128.3 MHz, CDCl₃): δ -0.7 (t, ¹J_{BF} = 31 Hz). ¹⁹F NMR (376.1 MHz, CDCl₃): δ -133.7 (q, ¹J_{FB} = 31 Hz). FT-IR (ATR): 3282 (m), 2928 (s), 2840 (s), 2240 (m), 1593 (s), 1505 (m), 1407 (s), 1343 (s), 1328 (s), 1307 (s), 1262 (s), 1166 (s), 1138 (s) cm⁻¹. UV-Vis (CH₂Cl₂): λ_{max} 552 nm (ε = 30 400 M⁻¹ cm⁻¹). Mass Spec. (EI, +ve mode): exact mass calculated for [C₁₇H₁₂BF₂N₅O]⁺: 351.1103; exact mass found: 351.1108; difference: +1.4 ppm.

Mono-benzyl-functionalized complex 8

A solution of PMDETA (0.002 g, 0.003 mL, 0.01 mmol) in dry THF (2 mL) was degassed *via* 3 freeze-pump-thaw cycles before CuI (0.003 g, 0.014 mmol) was added and the resulting mixture was stirred for 15 min at 22 °C. Benzyl azide (0.04 g, 0.04 mL, 0.3 mmol) and mono-alkyne-substituted complex **7** (0.1 g, 0.3 mmol) were then added and the reaction mixture

was stirred at 22 °C for 16 h. The THF solution was then purified by flash chromatography (THF, neutral alumina) and recrystallized from MeOH to afford mono-benzyl-functionalized complex **8** as a dark-purple microcrystalline solid. Yield = 0.14 g, 51%. M.p. 188–189 °C. ¹H NMR (400.1 MHz, CDCl₃): δ 7.95–7.93 (m, 6H, aryl CH), 7.75 (s, 1H, =CH), 7.42–7.40 (m, 3H, aryl CH), 7.34–7.32 (m, 2H, aryl CH), 7.00 (d, ³J_{HH} = 9 Hz, 2H, aryl CH) 5.60 (s, 2H, CH₂), 3.90 (s, 3H, CH₂). ¹³C{¹H} NMR (150.7 MHz, CDCl₃): δ 162.7, 146.6, 142.6, 136.8, 134.4, 132.9, 129.2, 128.9, 128.1, 126.4, 125.2, 123.3, 120.5, 114.9, 114.3, 55.8, 54.3. ¹¹B NMR (128.3 MHz, CDCl₃): δ -0.7 (t, ¹J_{BF} = 29 Hz). ¹⁹F NMR (376.1 MHz, CDCl₃): δ -134.2 (q, ¹J_{FB} = 29 Hz). FT-IR(ATR): 3123 (m), 2849 (m), 2250 (m), 1603 (s), 1509 (m), 1460 (m), 1375 (s), 1344 (s), 1326 (s), 1308 (s) cm⁻¹. UV-Vis (CH₂Cl₂): λ_{max} 560 nm (ε = 31 000 M⁻¹ cm⁻¹). Mass Spec. (EI, +ve mode): exact mass calculated for [C₂₄H₁₉BF₂N₈O]⁺: 484.1743; exact mass found: 484.1759; difference: +3.3 ppm.

Mono-ferrocene-functionalized BF₂ complex 9

A solution of PMDETA (0.002 g, 0.002 mL, 0.01 mmol) in THF (3 mL) was degassed *via* 3 freeze-pump-thaw cycles before CuI (0.002 g, 0.01 mmol) was added and the resulting mixture was stirred for 30 min. Ferrocenyl azide (0.024 g, 0.11 mmol) and mono-alkyne-substituted complex **7** (0.037 g, 0.11 mmol) were then added and the reaction mixture was stirred at 50 °C for 18 h. Upon cooling to 22 °C, the resulting dark purple solution was filtered through a plug of celite and purified by flash chromatography (CH₂Cl₂, silica gel) to afford ferrocene-functionalized complex **9** as a dark purple powder. Yield = 0.04 g, 72%. Melting point not observed (m.p. >250 °C). ¹H NMR (400.1 MHz, CDCl₃): δ 8.06 (s, 1H, triazole CH), 8.02–7.95 (m, 6H, aryl CH), 7.01–7.00 (m, 2H, aryl CH), 4.90 (s, 2H, ferrocene CH), 4.32 (s, 2H, ferrocene CH), 4.26 (s, 6H, ferrocene CH), 3.91 (s, 3H, OCH₃). ¹³C{¹H} NMR (150.7 MHz, CDCl₃): δ 162.9, 146.3, 142.9, 137.0, 132.9, 126.7, 125.4, 123.6, 120.0, 115.1, 114.4, 93.6, 70.4, 67.1, 62.4, 62.3, 56.0. ¹¹B NMR (128.3 MHz, CDCl₃): δ -0.7 (t, ¹J_{BF} = 30 Hz). ¹⁹F NMR (376.1 MHz, CDCl₃): δ -134.1 (q, ¹J_{FB} = 30 Hz). FT-IR(ATR): 3092 (w), 2926 (m), 2842 (m), 2241 (m), 1597 (s), 1505 (m), 1342 (m), 1261 (s), 1167 (s), 1029 (m), 965 (m), 832 (m), 733 (s) cm⁻¹. UV-Vis (CH₂Cl₂): λ_{max} 562 nm (ε = 28 600 M⁻¹ cm⁻¹). Mass Spec. (EI, +ve mode): exact mass calculated for [C₂₇H₂₁BF₂FeN₈O]⁺: 578.1249; exact mass found: 578.1261; difference: +2.1 ppm.

Bis-alkyne-substituted formazan **10**

In air, cyanoacetic acid (0.47 g, 5.5 mmol) was dissolved in deionized H₂O (75 mL) containing NaOH (2.20 g, 55 mmol). This colourless solution was stirred for 45 min in an ice bath. Meanwhile, 4-ethynylaniline (1.30 g, 12 mmol) was mixed with concentrated HCl (2.85 mL) in deionized H₂O (30 mL). This solution was cooled in an ice bath for 10 min before a solution of sodium nitrite (1.01 g, 15 mmol) in deionized H₂O (15 mL) was cooled in an ice bath, and then added slowly to the 4-ethynylaniline solution over a 10 min period. This mixture was stirred in an ice bath for 30 min, and then added slowly to the basic cyanoacetic acid solution. A dark red/orange colour per-



sisted almost immediately, and a dark red/orange precipitate formed after a few min. The mixture was stirred in an ice bath for an additional 16 h before ethyl acetate (250 mL) was added and the organic layer was isolated, washed with deionized H₂O (3 × 100 mL), dried over MgSO₄, gravity filtered and concentrated *in vacuo*. The resulting residue was purified by flash chromatography (CH₂Cl₂, neutral alumina) to afford bis-alkyne-substituted formazan **7** as a dark red solid. Yield = 1.55 g, 94%. Melting point not observed (>250 °C). ¹H NMR (599.5 MHz, CDCl₃) δ 12.60 (s, 1H, NH), 7.60 (s, 8H, aryl CH), 3.24 (s, 2H, alkyne CH). ¹³C{¹H} NMR (100.6 MHz, DMSO-*d*₆): δ 146.8, 133.0, 126.9, 121.7, 120.2, 112.5, 83.2, 82.8. FT-IR (KBr): 3397 (br, s), 3261 (m), 2915 (m), 2850 (m), 2221 (s), 1653 (s), 1540 (m), 1268 (m), 1193 (m), 1142 (m) cm⁻¹. UV-vis (CH₂Cl₂): λ_{max} = 434 nm (ε = 22 800 M⁻¹ cm⁻¹). Mass Spec. (EI, +ve mode): exact mass calculated for [C₁₈H₁₁N₅]⁺: 297.1014; exact mass found: 297.1018; difference: +1.3 ppm.

Bis-alkyne-substituted BF₂ complex **11**

Formazan **10** (1.00 g, 3.40 mmol) was dissolved in dry toluene (100 mL). NEt₃ (1.03 g, 1.42 mL, 10.2 mmol) was then added slowly and the solution was stirred for 10 min before BF₃·OEt₂ (2.39 g, 2.08 mL, 16.8 mmol) was added and the solution was heated with stirring at 80 °C for 18 h. The solution gradually turned from dark red to dark purple during this time. After cooling to 22 °C, deionized H₂O (10 mL) was added to quench any excess reactive boron-containing compounds. The purple toluene solution was then washed with deionized H₂O (3 × 50 mL), dried over MgSO₄, gravity filtered and concentrated *in vacuo*. The resulting residue was purified by flash chromatography (CH₂Cl₂, neutral alumina) to afford bis-alkyne-substituted complex **11** as a dark purple microcrystalline solid. Yield = 0.82 g, 70%. Melting point = 218 °C (decomp.). ¹H NMR (599.5 MHz, CDCl₃) δ 7.91 (d, ³J_{HH} = 9 Hz, 4H, aryl CH), 7.61 (d, ³J_{HH} = 9 Hz, 4H, aryl CH), 3.32 (s, 2H, alkyne CH). ¹³C{¹H} NMR (100.6 MHz, CDCl₃): δ 143.0, 133.4, 126.0, 123.2, 113.8, 82.6, 81.8. ¹¹B NMR (128.3 MHz, CDCl₃): δ -0.8 (t, ¹J_{BF} = 31 Hz). ¹⁹F NMR (376.1 Hz, CDCl₃): δ -132.0 (q, ¹J_{FB} = 31 Hz). FT-IR (KBr): 3290 (s), 3189 (m), 3108 (m), 3092 (m), 2244 (s), 1589 (s), 1391 (m), 1331 (s), 1187 (m), 1147 (s), 1033 (m), 958 (m), 838 (s) cm⁻¹. UV-vis (CH₂Cl₂): λ_{max} = 530 nm (ε = 32 700 M⁻¹ cm⁻¹). Mass Spec. (EI, +ve mode): exact mass calculated for [C₁₈H₁₀N₅BF₂]⁺: 345.0997; exact mass found: 345.0999; difference: +0.6 ppm.

Bis-benzyl-functionalized BF₂ complex **12**

A solution PMDETA (0.0075 g, 0.0091 mL, 0.044 mmol) in THF (3 mL) was degassed *via* 3 freeze-pump-thaw cycles before CuI (0.0083 g, 0.044 mmol) was added and the resulting mixture was stirred for 30 min. Benzyl azide (0.174 g, 1.30 mmol) and bis-alkyne-substituted complex **11** (0.150 g, 0.44 mmol) were then added and the reaction mixture was stirred at 50 °C for 18 h. Upon cooling to 22 °C, the resulting dark purple solution was filtered through a plug of celite and purified by flash chromatography (CH₂Cl₂, silica gel) to afford bis-benzyl-functionalized complex **12** as a dark-purple solid. Yield = 0.21 g,

77%. Melting point not observed (>250 °C). ¹H NMR (599.5 MHz, CDCl₃) δ 7.99–7.94 (m, 8H, aryl CH), 7.76 (s, 2H, triazole CH), 7.43–7.38 (m, 6H, aryl CH), 7.34–7.33 (m, 4H, aryl CH), 5.60 (s, 4H, benzyl CH₂). ¹³C{¹H} NMR (150.7 MHz, CDCl₃): δ 146.7, 142.8, 134.4, 133.8, 129.4, 129.2, 128.3, 126.7, 123.8, 120.7, 114.2, 54.6. ¹¹B NMR (128.3 MHz, CDCl₃): δ -0.7 (t, ¹J_{BF} = 29 Hz). ¹⁹F NMR (376.1 Hz, CDCl₃): δ -133.1 (q, ¹J_{FB} = 29 Hz). FT-IR (KBr): 3109 (m), 3100 (s), 3035 (m), 2915 (m), 2246 (s), 1608 (s), 1497 (m), 1456 (s), 1326 (s), 1226 (m), 1074 (s), 1027 (s), 976 (m), 826 (m) cm⁻¹. UV-vis (CH₂Cl₂): λ_{max} = 558 nm (ε = 39 600 M⁻¹ cm⁻¹). Mass Spec. (EI, +ve mode): exact mass calculated for [C₃₂H₂₅N₁₁BF₂]⁺: 612.2356; exact mass found: 612.2362; difference: +1.0 ppm.

Bis-ferrocene-functionalized BF₂ complex **13**

A solution of PMDETA (0.0019 g, 0.0023 mL, 0.011 mmol) in THF (4 mL) was degassed *via* 3 freeze-pump-thaw cycles before CuI (0.0021 g, 0.011 mmol) was added and the resulting mixture was stirred for 30 min. Ferrocenyl azide (0.050 g, 0.22 mmol) and bis-alkyne-substituted complex **11** (0.038 g, 0.11 mmol) were then added and the reaction mixture was stirred at 50 °C for 18 h. The resulting dark purple solution was filtered through a plug of celite and purified by flash chromatography (CH₂Cl₂, silica gel) to afford ferrocene-functionalized complex **13** as a dark purple powder. Yield = 0.080 g, 90%. Melting point not observed (>250 °C). ¹H NMR (599.5 MHz, CDCl₃) δ 8.08 (s, 2H, triazole CH), 8.05 (s, 8H, aryl CH), 4.91 (m, 4H, ferrocene CH), 4.34–4.33 (m, 4H, ferrocene CH), 4.28–4.26 (m, 10H, ferrocene CH). Due to the poor solubility of **13**, and despite a saturated solution in THF-*d*₈ being subjected to 10 000 scans (8 h) on a 600 MHz spectrometer, a publication quality ¹³C{¹H} NMR spectrum was not obtained. ¹¹B NMR (128.3 MHz, CDCl₃): δ -0.6 (t, ¹J_{BF} = 28 Hz). ¹⁹F NMR (376.1 Hz, CDCl₃): δ -132.8 (q, ¹J_{FB} = 28 Hz). FT-IR (KBr): 3131 (w), 3092 (m), 2929 (m), 2246 (m), 1605 (s), 1381 (m), 1341 (s), 1226 (m), 1181 (s), 1027 (m), 966 (s), 816 (s) cm⁻¹. UV-vis (CH₂Cl₂): λ_{max} = 569 nm (ε = 30 900 M⁻¹ cm⁻¹). Mass Spec. (ESI, +ve mode): exact mass calculated for [C₃₈H₂₉N₁₁BF₂Fe₂]⁺: 800.1367; exact mass found: 800.1380; difference: +1.6 ppm.

Bis-TEG-functionalized BF₂ complex **14**

A solution of PMDETA (0.008 g, 0.009 mL, 0.04 mmol) in THF (3 mL) was degassed *via* 3 freeze-pump-thaw cycles before CuI (0.008 g, 0.04 mmol) was added and the resulting solution was stirred for 30 min. TEG-azide (0.29 g, 1.3 mmol) and bis-alkyne-substituted complex **11** (0.15 g, 0.44 mmol) were then added and the reaction mixture was stirred at 50 °C for 18 h. The resulting dark purple solution was filtered through a plug of celite and purified by precipitation of a concentrated THF solution into *n*-hexanes thrice to yield a dark purple solid. Yield = 0.24 g, 71%. ¹H NMR (599.5 MHz, CDCl₃) δ 8.20 (s, 2H, triazole CH), 8.00 (s, 8H, aryl CH), 4.63–4.61 (m, 4H, ethylene glycol CH₂), 3.93–3.90 (m, 4H, ethylene glycol CH₂), 3.71–3.57 (m, 24H, ethylene glycol CH₂). ¹³C{¹H} NMR (150.7 MHz, CDCl₃): 145.9, 142.4, 134.0, 126.5, 123.6, 122.6, 114.1, 72.4, 70.5, 70.4, 70.3, 70.1, 69.4, 61.5, 50.5 ppm. ¹¹B NMR



(128.3 MHz, CDCl₃): δ -0.7 (t, $^1J_{\text{BF}} = 29$ Hz). ^{19}F NMR (376.1 Hz, CDCl₃): δ -133.1 (q, $^1J_{\text{FB}} = 29$ Hz). FT-IR (KBr): 3132 (br), 3317 (w), 2915 (m), 2868 (m), 2242 (w), 1604 (s), 1527 (w), 1461 (m), 1360 (m), 1333 (s), 1227 (m), 1179 (s), 1121 (s), 1069 (s), 1029 (s), 967 (s), 845 (m), 765 (w) cm⁻¹. UV-vis (CH₂Cl₂): $\lambda_{\text{max}} = 542$ nm ($\epsilon = 16\,500$ M⁻¹ cm⁻¹). UV-vis (H₂O): $\lambda_{\text{max}} = 527$ nm ($\epsilon = 15\,100$ M⁻¹ cm⁻¹). Mass Spec. (ESI, +ve mode): exact mass calculated for [C₃₄H₄₅N₁₁O₈BF₂]⁺: 784.3514; exact mass found: 784.3540; difference: +3.3 ppm.

Acknowledgements

We would like to thank the University of Western Ontario, the Natural Science and Engineering Research Council (NSERC) of Canada (J. B. G.: DG, 435675, L. G. L.: DG, 326972 and S. M. B.: CGS D scholarship), the Ontario Ministry of Research and Innovation (J. B. G.: ERA, ER14-10-147) and the Canada Foundation for Innovation (J. B. G.: JELF, 33977) for funding this work. Finally we thank Prof. Elizabeth R. Gillies and Prof. Mark S. Workentin for access to instrumentation in their labs.

Notes and references

- 1 S. Weiss, *Science*, 1999, **283**, 1676–1683.
- 2 J. Eid, A. Fehr, J. Gray, K. Luong, J. Lyle, G. Otto, P. Peluso, D. Rank, P. Baybayan, B. Bettman, A. Bibillo, K. Bjornson, B. Chaudhuri, F. Christians, R. Cicero, S. Clark, R. Dalal, A. deWinter, J. Dixon, M. Foquet, A. Gaertner, P. Hardenbol, C. Heiner, K. Hester, D. Holden, G. Kearns, X. Kong, R. Kuse, Y. Lacroix, S. Lin, P. Lundquist, C. Ma, P. Marks, M. Maxham, D. Murphy, I. Park, T. Pham, M. Phillips, J. Roy, R. Sebra, G. Shen, J. Sorenson, A. Tomaney, K. Travers, M. Trulson, J. Vieceli, J. Wegener, D. Wu, A. Yang, D. Zaccarin, P. Zhao, F. Zhong, J. Korlach and S. Turner, *Science*, 2009, **323**, 133–138.
- 3 B. Huang, W. Wang, M. Bates and X. Zhuang, *Science*, 2008, **319**, 810–813.
- 4 L. D. Lavis and R. T. Raines, *ACS Chem. Biol.*, 2008, **3**, 142–155.
- 5 E. Kim, Y. Lee, S. Lee and S. B. Park, *Acc. Chem. Res.*, 2015, **48**, 538–547.
- 6 S. T. Laughlin, J. M. Baskin, S. L. Amacher and C. R. Bertozzi, *Science*, 2008, **320**, 664–667.
- 7 U. Resch-Genger, M. Grabolle, S. Cavaliere-Jaricot, R. Nitschke and T. Nann, *Nat. Methods*, 2008, **5**, 763–775.
- 8 D. Ding, K. Li, B. Liu and B. Z. Tang, *Acc. Chem. Res.*, 2013, **46**, 2441–2453.
- 9 Z. Yang, A. Sharma, J. Qi, X. Peng, D. Y. Lee, R. Hu, D. Lin, J. Qu and J. S. Kim, *Chem. Soc. Rev.*, 2016, **45**, 4651–4667.
- 10 H. Kobayashi, M. Ogawa, R. Alford, P. L. Choyke and Y. Urano, *Chem. Rev.*, 2010, **110**, 2620–2640.
- 11 L. D. Lavis and R. T. Raines, *ACS Chem. Biol.*, 2014, **9**, 855–866.
- 12 V. V. Rostovtsev, L. G. Green, V. V. Fokin and K. B. Sharpless, *Angew. Chem., Int. Ed.*, 2002, **41**, 2596–2599.
- 13 C. W. Tornøe, C. Christensen and M. Meldal, *J. Org. Chem.*, 2002, **67**, 3057–3064.
- 14 Q. Liu and Y. Tor, *Org. Lett.*, 2003, **5**, 2571–2572.
- 15 B. Khanna, S. R. Parkin and K. D. Revell, *Tetrahedron Lett.*, 2012, **53**, 6383–6387.
- 16 O. Karagollu, M. Gorur, F. Gode, B. Sennik and F. Yilmaz, *Macromol. Chem. Phys.*, 2015, **216**, 939–949.
- 17 R. Chopra, P. Kaur and K. Singh, *Anal. Chim. Acta*, 2015, **864**, 55–63.
- 18 B. Brizet, V. Goncalves, C. Bernhard, P. D. Harvey, F. Denat and C. Goze, *Chem. – Eur. J.*, 2014, **20**, 12933–12944.
- 19 B. Dhokale, T. Jadhav, S. M. Mobin and R. Misra, *Dalton Trans.*, 2015, **44**, 15803–15812.
- 20 H. R. A. Golf, H.-U. Reissig and A. Wiehe, *Org. Lett.*, 2015, **17**, 982–985.
- 21 X. Chen, Q. Wu, L. Henschke, G. Weber and T. Weil, *Dyes Pigm.*, 2012, **94**, 296–303.
- 22 P. Gobbo, P. Gunawardene, W. Luo and M. S. Workentin, *Synlett*, 2015, 1169–1174.
- 23 Y. Su, L. Li, H. Wang, X. Wang and Z. Zhang, *Chem. Commun.*, 2016, **52**, 2185–2188.
- 24 V. Evans, P. Duncanson, M. Motevalli and M. Watkinson, *Dyes Pigm.*, 2016, **135**, 36–40.
- 25 M. Albrecht, A. Lippach, M. P. Exner, J. Jerbi, M. Springborg, N. Budisa and G. Wenz, *Org. Biomol. Chem.*, 2015, **13**, 6728–6736.
- 26 S. Guo, L. Ma, J. Zhao, B. Küçüköz, A. Karatay, M. Hayvali, H. G. Yaglioglu and A. Elmali, *Chem. Sci.*, 2014, **5**, 489–500.
- 27 C. Beyer and H.-A. Wagenknecht, *J. Org. Chem.*, 2010, **75**, 2752–2755.
- 28 S. P. Sau and P. J. Hrdlicka, *J. Org. Chem.*, 2012, **77**, 5–16.
- 29 Z. Liu, Y. Li, J. Lozada, P. Schaffer, M. J. Adam, T. J. Ruth and D. M. Perrin, *Angew. Chem., Int. Ed.*, 2013, **52**, 2303–2307.
- 30 V. Martí-Centelles and P. D. Beer, *Chem. – Eur. J.*, 2015, **21**, 9397–9404.
- 31 R. Arumugaperumal, V. Srinivasadesikan, M. V. Ramakrishnam Raju, M.-C. Lin, T. Shukla, R. Singh and H.-C. Lin, *ACS Appl. Mater. Interfaces*, 2015, **7**, 26491–26503.
- 32 X. Ma, J. Zhang, J. Cao, X. Yao, T. Cao, Y. Gong, C. Zhao and H. Tian, *Chem. Sci.*, 2016, **7**, 4582–4588.
- 33 J. Rosenthal and S. J. Lippard, *J. Am. Chem. Soc.*, 2010, **132**, 5536–5537.
- 34 N. B. Yapici, S. Jockusch, A. Moscatelli, S. R. Mandalapu, Y. Itagaki, D. K. Bates, S. Wiseman, K. M. Gibson, N. J. Turro and L. Bi, *Org. Lett.*, 2012, **14**, 50–53.
- 35 K.-B. Li, H. Wang, Y. Zang, X.-P. He, J. Li, G.-R. Chen and H. Tian, *ACS Appl. Mater. Interfaces*, 2014, **6**, 19600–19605.
- 36 R. S. Singh, R. K. Gupta, R. P. Paitandi, A. Misra and D. S. Pandey, *New J. Chem.*, 2015, **39**, 2233–2239.
- 37 E. Manandhar, J. H. Broome, J. Myrick, W. Lagrone, P. J. Cragg and K. J. Wallace, *Chem. Commun.*, 2011, **47**, 8796–8798.



- 38 S. A. Ingale and F. Seela, *J. Org. Chem.*, 2012, **77**, 9352–9356.
- 39 T. Uppal, N. V. S. D. K. Bhupathiraju and M. G. H. Vicente, *Tetrahedron*, 2013, **69**, 4687–4693.
- 40 M. Sun, K. Müllen and M. Yin, *Chem. Soc. Rev.*, 2016, **45**, 1513–1528.
- 41 P. Kele, X. Li, M. Link, K. Nagy, A. Herner, K. Lőrincz, S. Béni and O. S. Wolfbeis, *Org. Biomol. Chem.*, 2009, **7**, 3486–3490.
- 42 M. Zhou, X. Zhang, M. Bai, D. Shen, B. Xu, J. Kao, X. Ge and S. Achilefu, *RSC Adv.*, 2013, **3**, 6756–6758.
- 43 J. H. Kwak, Y. He, B. Yoon, S. Koo, Z. Yang, E. J. Kang, B. H. Lee, S.-Y. Han, Y. C. Yoo, K. B. Lee and J. S. Kim, *Chem. Commun.*, 2014, **50**, 13045–13048.
- 44 X.-t. Zhang, Z.-y. Gu, L. Liu, S. Wang and G.-w. Xing, *Chem. Commun.*, 2015, **51**, 8606–8609.
- 45 E. M. Sletten and C. R. Bertozzi, *Acc. Chem. Res.*, 2011, **44**, 666–676.
- 46 J. C. Jewett and C. R. Bertozzi, *Chem. Soc. Rev.*, 2010, **39**, 1272–1279.
- 47 R. Travieso-Puente, M.-C. Chang and E. Otten, *Dalton Trans.*, 2014, **43**, 18035–18041.
- 48 M.-C. Chang, T. Dann, D. P. Day, M. Lutz, G. G. Wildgoose and E. Otten, *Angew. Chem., Int. Ed.*, 2014, **53**, 4118–4122.
- 49 E. Kabir, C.-H. Wu, J. I.-C. Wu and T. S. Teets, *Inorg. Chem.*, 2016, **55**, 956–963.
- 50 W. Schorn, D. Grosse-Hagenbrock, B. Oelkers and J. Sundermeyer, *Dalton Trans.*, 2016, **45**, 1201–1207.
- 51 A. Mandal, B. Schwederski, J. Fiedler, W. Kaim and G. K. Lahiri, *Inorg. Chem.*, 2015, **54**, 8126–8135.
- 52 R. Travieso-Puente, J. O. P. Broekman, M.-C. Chang, S. Demeshko, F. Meyer and E. Otten, *J. Am. Chem. Soc.*, 2016, **138**, 5503–5506.
- 53 N. A. Protasenko, A. I. Poddelsky, A. S. Bogomyakov, G. K. Fukin and V. K. Cherkasov, *Inorg. Chem.*, 2015, **54**, 6078–6080.
- 54 M.-C. Chang and E. Otten, *Organometallics*, 2016, **35**, 534–542.
- 55 J. B. Gilroy, M. J. Ferguson, R. McDonald, B. O. Patrick and R. G. Hicks, *Chem. Commun.*, 2007, 126–128.
- 56 M.-C. Chang and E. Otten, *Chem. Commun.*, 2014, **50**, 7431–7433.
- 57 S. M. Barbon, P. A. Reinkeluers, J. T. Price, V. N. Staroverov and J. B. Gilroy, *Chem. – Eur. J.*, 2014, **20**, 11340–11344.
- 58 R. R. Maar, S. M. Barbon, N. Sharma, H. Groom, L. G. Luyt and J. B. Gilroy, *Chem. – Eur. J.*, 2015, **21**, 15589–15599.
- 59 S. M. Barbon, V. N. Staroverov and J. B. Gilroy, *J. Org. Chem.*, 2015, **80**, 5226–5235.
- 60 M.-C. Chang and E. Otten, *Inorg. Chem.*, 2015, **54**, 8656–8664.
- 61 M.-C. Chang, A. Chantzis, D. Jacquemin and E. Otten, *Dalton Trans.*, 2016, **45**, 9477–9484.
- 62 M. Hesari, S. M. Barbon, V. N. Staroverov, Z. Ding and J. B. Gilroy, *Chem. Commun.*, 2015, **51**, 3766–3769.
- 63 R. R. Maar and J. B. Gilroy, *J. Mater. Chem. C*, 2016, **4**, 6478–6482.
- 64 S. Novoa, J. A. Paquette, S. M. Barbon, R. R. Maar and J. B. Gilroy, *J. Mater. Chem. C*, 2016, **4**, 3987–3994.
- 65 S. M. Barbon and J. B. Gilroy, *Polym. Chem.*, 2016, **7**, 3589–3598.
- 66 J. B. Gilroy, P. O. Otieno, M. J. Ferguson, R. McDonald and R. G. Hicks, *Inorg. Chem.*, 2008, **47**, 1279–1286.
- 67 S. M. Barbon, J. T. Price, P. A. Reinkeluers and J. B. Gilroy, *Inorg. Chem.*, 2014, **53**, 10585–10593.
- 68 S. Fery-Forgues and D. Lavabre, *J. Chem. Educ.*, 1999, **76**, 1260–1264.
- 69 K. Suzuki, A. Kobayashi, S. Kaneko, K. Takehira, T. Yoshihara, H. Ishida, Y. Shiina, S. Oishi and S. Tobita, *Phys. Chem. Chem. Phys.*, 2009, **11**, 9850–9860.
- 70 M. Li, Z. Guo, W. Zhu, F. Marken and T. D. James, *Chem. Commun.*, 2015, **51**, 1293–1296.
- 71 M. R. Rao, K. V. P. Kumar and M. Ravikanth, *J. Organomet. Chem.*, 2010, **695**, 863–869.
- 72 B. Dhokale, P. Gautam, S. M. Mobin and R. Misra, *Dalton Trans.*, 2013, **42**, 1512–1518.
- 73 R. Misra, B. Dhokale, T. Jadhav and S. M. Mobin, *Dalton Trans.*, 2013, **42**, 13658–13666.
- 74 N. G. Connelly and W. E. Geiger, *Chem. Rev.*, 1996, **96**, 877–910.
- 75 X. Yin, Y. Li, Y. Zhu, X. Jing, Y. Li and D. Zhu, *Dalton Trans.*, 2010, **39**, 9929–9935.
- 76 J. Rochford, A. D. Rooney and M. T. Pryce, *Inorg. Chem.*, 2007, **46**, 7247–7249.
- 77 F. Otón, M. d. C. González, A. Espinosa, C. Ramírez de Arellano, A. Tárraga and P. Molina, *J. Org. Chem.*, 2012, **77**, 10083–10092.
- 78 S. Tobita, K. Ida and S. Shiobara, *Res. Chem. Intermed.*, 2001, **27**, 205–218.
- 79 P. J. Sadkowski and G. R. Fleming, *Chem. Phys.*, 1980, **54**, 79–89.
- 80 H. Nakamura and J. Tanaka, *Chem. Phys. Lett.*, 1981, **78**, 57–60.
- 81 L. Quan, Y. Chen, X.-J. Lv and W.-F. Fu, *Chem. – Eur. J.*, 2012, **18**, 14599–14604.
- 82 J. Oshima, T. Yoshihara and S. Tobita, *Chem. Phys. Lett.*, 2006, **423**, 306–311.
- 83 C. Patra, A. K. Bhanja, A. Mahapatra, S. Mishra, K. D. Saha and C. Sinha, *RSC Adv.*, 2016, **6**, 76505–76513.
- 84 B. Zhou, W. Liu, H. Zhang, J. Wu, S. Liu, H. Xu and P. Wang, *Biosens. Bioelectron.*, 2015, **68**, 189–196.
- 85 W. Liu, B. Zhou, G. Niu, J. Ge, J. Wu, H. Zhang, H. Xu and P. Wang, *ACS Appl. Mater. Interfaces*, 2015, **7**, 7421–7427.
- 86 M. V. Werrett, P. J. Wright, P. V. Simpson, P. Raiteri, B. W. Skelton, S. Stagni, A. G. Buckley, P. J. Rigby and M. Massi, *Dalton Trans.*, 2015, **44**, 20636–20647.
- 87 Z. Yang, C. Yan, Y. Chen, C. Zhu, C. Zhang, X. Dong, W. Yang, Z. Guo, Y. Lu and W. He, *Dalton Trans.*, 2011, **40**, 2173–2176.
- 88 P. Gobbo, W. Luo, S. J. Cho, X. Wang, M. C. Biesinger, R. H. E. Hudson and M. S. Workentin, *Org. Biomol. Chem.*, 2015, **13**, 4605–4612.
- 89 A. C. Pauly, C. D. Varnado, C. W. Bielawski and P. Theato, *Macromol. Rapid Commun.*, 2014, **35**, 210–213.



- 90 Bruker-AXS, *SAINT version 2013.8*, Bruker-AXS, Madison, WI 53711, USA, 2013.
- 91 Bruker-AXS, *SADABS version 2012.1*, Bruker-Nonius, Madison, WI 53711, USA, 2012.
- 92 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Fundam. Crystallogr.*, 2015, **71**, 3–8.
- 93 G. M. Sheldrick, *Acta Crystallogr., Sect. C: Cryst. Struct. Commun.*, 2015, **71**, 3–8.

