A fluorescent activatable probe for imaging intracellular Mg$^{2+}$

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An activatable BODIPY probe for in vitro detection and fluorescence cell imaging of free Mg$^{2+}$ without interference from Ca$^{2+}$ is described. Fluorescence amplification of the probe is observed upon detection of physiological concentrations of Mg$^{2+}$ due to reduced rotation of the fluorophore and effective chelation by a quinolizine-based core.

Introduction

Intracellular magnesium (Mg$^{2+}$) plays a crucial role in the physiological activity of live cells. At concentrations below 30 mM, Mg$^{2+}$ is known to regulate and participate in a range of fundamental biological processes, including metabolic pathways, DNA synthesis and being a co-factor for more than 600 enzymes. Consequently, the concentration of free unbound intracellular Mg$^{2+}$ falls to around 0.5–1.2 mM.1 A number of pathological disorders have been correlated to abnormal levels of Mg$^{2+}$ within the body.2 For instance, low levels of Mg$^{2+}$ are associated with the progression of Parkinson’s disease, characterised by a depletion in the number of dopaminergic neurons in the brain.3 Therefore, the importance of Mg$^{2+}$ in multiple physiological conditions demands new chemical tools to investigate the storage, concentration and homeostasis of Mg$^{2+}$ in both health and disease.

Small organic fluorophores offer a cheap, reliable and non-destructive method of cellular analysis, and have been widely reported to detect a plethora of biomolecules and enzymes, as well as to monitor biological processes and discriminate cell types.4–9 Advances in fluorophore development have led to the generation of smart or activatable probes, which are chemically designed so that their photophysical properties depend on an external stimulus. The fine-tuning of these chemical architectures not only allows spatiotemporal control of the readout of the probes but also provides enhanced signal-to-noise ratios, which boost their sensitivity in imaging studies under fluorescence microscopy. Furthermore, when significant differences are achieved between the readouts of activated and non-activated fluorophores, quantitative measurements of specific analytes or biological events can be performed. In the context of divalent cation sensing, since the pioneering work from Tsien et al.,10 multiple fluorescent probes have been reported for in vivo and in vitro tracking of divalent cations.11–13 Numerous chemical probes have been developed to monitor the levels of Ca$^{2+}$, the most abundant cation, with only a modest number of probes being available for sensing Mg$^{2+}$. A major challenge in the development of Mg$^{2+}$-responding probes is to ensure that the fluorescence signal is elicited only in response to Mg$^{2+}$ and no other divalent intracellular cations, with Ca$^{2+}$ being a major interference due to its high intracellular levels. Mag-indo-1 and Mag-fura-2 are commercially available probes based on the O-aminophenol-N,N,O-triacetic acid (APTRA) chelating group, which can strongly bind to Mg$^{2+}$ as well as to other intracellular divalent cations.14 Remarkable work has been also recently undertaken by the group of Buccella with the adaptation of APTRA to a range of chemical scaffolds (e.g. HaloTag-binding molecules, ratiometric and red-emitting fluorophores) to improve its capabilities as imaging reagents.15–17 Other chelating groups have been reported for enhanced recognition of Mg$^{2+}$ and minimal cross-reactivity with Ca$^{2+}$. Among these, the 4-oxo-4H-quinolizine-3-carboxylic acid scaffold reported by Levy et al.18 offers good discrimination between the two major divalent ions and can be effectively conjugated to fluorophores (e.g. fluorescein) to produce probes that respond to Mg$^{2+}$ chelation by photo-induced electron transfer (PeT).19 However, the limited cell permeability of fluorescein hinders the use of these dyes for visualising intracellular Mg$^{2+}$ levels in live cells. In addition to high cell permeability, the 4,4-difluoro-4-bora-3a,4a-diaza-s-indacene (BODIPY) scaffold presents excellent photophysical...
features for live cell imaging, such as narrow excitation and emission and excellent photostability. Furthermore, the fluorescence emission of BODIPY can be fine-tuned via incorporation of chelating groups at the meso position, which affect intramolecular PeT and the overall emission of the fluorophores. Therefore, we envisioned that the introduction of the 4-oxo-4H-quinolizine-3-carboxylic acid into the cell-permeable BODIPY scaffold might render novel activatable fluorophores for imaging intracellular levels of Mg$^{2+}$.

Results and discussion

Rational design and chemical synthesis of Mg$^{2+}$ probes

First, we synthesized the quinolizine chelating moiety 1 (Fig. 1), which incorporates a β-diketone moiety to selectively chelate the Mg$^{2+}$ ion and an aldehyde functional group for adaptation to the synthesis of symmetric BODIPY fluorophores. The 4-oxo-4H-quinolizine-3-carboxylic acid moiety shows enhanced selectivity for Mg$^{2+}$ over Ca$^{2+}$ when compared to other chelating groups (e.g. APTRA). The synthesis of the BODIPY fluorophore 2 was achieved in a one-pot single reaction by conjugation of the aldehyde 1 to pyrrole using slightly modified reported procedures to isolate the BODIPY compound 2 (Fig. 1) and examine its properties as a Mg$^{2+}$-sensitive fluorophore.

The low fluorescence quantum yield (i.e. around 2%) of compound 2 after incubation with Mg$^{2+}$ proved ineffective for the detection of Mg$^{2+}$ levels within the physiological range.

![Fig. 1](image)

**Fig. 1** Simplified synthetic scheme for the preparation of Mg-sensitive BODIPY fluorophores based on the 4-oxo-4H-quinolizine-3-carboxylic acid chelating scaffold. Below are represented the chemical structures of BODIPY fluorophores 2, 3 and 4 and their corresponding activation modes upon Mg$^{2+}$ recognition. Fluorescence quantum yields were determined relative to fluorescein in basic ethanol.

Such weak fluorescence emission can be explained due to the free rotation of the quinolizine group conjugated to the meso position, which might result in the energy of the excited state being lost as rotational energy rather than emitted as fluorescence. We assessed this hypothesis by examining the fluorescence emission of compound 2 in environments of increasing viscosity (Fig. S1 in ESI†). The enhancement of fluorescence intensity in glycerol-containing media confirmed the free rotation of the quinolizine group around the meso position as a major cause for the low fluorescence quantum yield of compound 2.

We envisioned that BODIPY probes where the fluorescence enhancements derived from Mg$^{2+}$ chelation and restricted rotation were combined in a single molecular framework might display suitable fluorescent quantum yields for Mg$^{2+}$ detection in cells. As such, we designed and synthesized two additional BODIPY compounds with two or four methyl substituents (3 and 4 respectively, Fig. 1). Whereas compound 3 would retain similar rotational freedom to compound 2, the steric hindrance around the quinolizine core in compound 4 would minimise its free rotation. Compounds 3 and 4 were synthesized by condensation of the aldehyde 1 to 2-methylpyrrole or 2,4-dimethylpyrrole respectively, and isolated with reasonable synthetic yields. The excitation and emission wavelengths of all quinolizine-BODIPY derivatives (2-4) are similar (Fig. 1 and Fig. S2 in ESI†) as well as their photostability properties (Fig. S3 in ESI†). However, their fluorescence quantum yields were significantly different, with the tetramethyl-substituted BODIPY 4 showing the highest fluorescence quantum yield after binding to Mg$^{2+}$ (11%, Fig. 1 and Fig. S4 in ESI†). This observation supports our rational design of BODIPY probes combining the chelating 4-oxo-4H-quinolizine-3-carboxylic acid with restricted rotational features to maximise the fluorescence response upon Mg$^{2+}$ recognition.

**In vitro spectral characterisation of Mg$^{2+}$ binding**

In view of the spectral properties of compound 4, we examined its application for the detection and visualisation of intracellular Mg$^{2+}$ levels. As shown in Fig. 2A, compound 4 showed a dose-dependent fluorescence increase upon incubation with concentrations of Mg$^{2+}$ within its physiological range (i.e. up to 1 mM). Furthermore, compound 4 showed excellent discrimination of Mg$^{2+}$ vs. other divalent cations, with around 4-fold selectivity over Ca$^{2+}$—the most abundant divalent cation found in cells and tissues—marginal cross-reactivity with Mn$^{2+}$ and Zn$^{2+}$ at their physiological concentrations (Fig. 2B and Fig. S5 in ESI†). Mn$^{2+}$ is present in cells in a relatively broad range of concentrations (i.e. between 0.04 and 2 mM) whereas Zn$^{2+}$ is predominantly bound as a co-factor in protein complexes and found as a free cation in cells at very low concentrations (i.e. in the pM range). Notably, the co-incubation with Mg$^{2+}$ and the chelating agent ethylenediaminetetraacetic acid (EDTA) did not lead to any fluorescence enhancement (Fig. 2B). This result confirms that, together with the lack of rotational energy loss, the blockage of the intramolecular PeT due to Mg$^{2+}$ chelation is likely to be the main mechanism
Fig. 2 (A) Fluorescence emission of compound 4 (10 μM, H₂O:EtOH 1:1) upon incubation with increasing concentrations (0, 0.1, 0.25, 0.5, 0.75 and 1 mM) of Mg²⁺; λexc.: 450 nm. (B) Comparative fluorescence of compound 4 upon incubation with different divalent cations at their respective physiological concentrations (Mg²⁺, Ca²⁺, Mn²⁺: 1 mM; Zn²⁺: 1 μM; EDTA: 5 mM). Inset: Pictograms of compound 4 under 365 nm excitation in solutions without (left) or with Mg²⁺ (right). *** for p < 0.001, n.s. for p > 0.05.

Further detailed characterisation studies showed that the Mg²⁺ adduct presented a K_d value of around 100 μM (Fig. 3A), in the range of DCHQ-based quinoline fluorophores and an order of magnitude lower than APTRA-based Mg²⁺-responding fluorophores. We also analysed the fluorescence response of the Mg²⁺-4 adduct to different pH values and observed no major dependence on pH (Fig. S7 in ESI†).

Fig. 3 (A) Determination of the dissociation constant of the Mg²⁺-4 adduct by measuring the fluorescence intensity of compound 4 (10 μM, H₂O:EtOH 1:1) at increasing concentrations of Mg²⁺. Values represented as means ± s.e.m. (n = 3). K_d: 92 ± 30 μM. (B) Emission spectra of compound 4 (50 μM) upon incubation with 2 mM free Mg²⁺ (green) and 2 mM Mg-ATP (grey); λexc.: 450 nm. (Inset) Quantification of the fluorescence intensities for both conditions. Values represented as means ± s.d. (n = 3).

Fig. 4 Fluorescence confocal microscopy images of A549 cells upon incubation with compound 4. Cells were pre-incubated or not with 1 μM FCCP, and then treated with compound 4 (3 μM) for 15 min. Fluorescence images are shown in the left panels and brightfield images on the right. Quantitative analysis of the fluorescence intensity in both conditions is shown in Fig. S8.† Scale bar: 10 μm.

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In view of the excellent properties of compound 4 as a Mg²⁺-activatable fluorescent probe, we examined its capabilities for cell imaging using A549 human adenocarcinoma cells. A549 cells were incubated or not with carbonyl cyanide p-(trifluoromethoxy)phenylhydrazone (FCCP), a molecule that acts as a mitochondria uncoupler and leads to an increase in Mg²⁺ concentration resulting from ATP hydrolysis and/or Mg²⁺ efflux from mitochondria. We acquired fluorescence confocal microscopy images of both untreated and FCCP-treated cells that had been incubated with the same concentration of compound 4, and observed brighter intracellular emission in the latter due to significantly higher intracellular Mg²⁺ levels (Fig. 4 and Fig. S8 in ESI† for quantitative image analysis). We also examined the intracellular localisation of compound 4 by co-staining with different subcellular markers (i.e. LysoTracker Red and MitoTracker Red). Dual-colour images suggested that compound 4 mainly localised in the cytoplasm of A549 cells, with slightly preferential accumulation in the mitochondria over lysosomal compartments (Fig. S9 in ESI†). Furthermore, we performed cell viability experiments and confirmed that the incubation of A549 cells with compound 4 did not induce any significant cytotoxicity at the concentrations used for comparison to the same concentration of Mg-ATP, asserting its utility to monitor changes in the levels of free Mg²⁺ within the intracellular physiological range.
Experimental section

General methods

Commercially available reagents were used without further purification. Thin-layer chromatography was conducted on Merck silica gel 60 F254 sheets and visualized by UV (254 and 365 nm). Silica gel (particle size 35–70 µm) was used for column chromatography. 1H and 13C spectra were recorded in a Bruker Avance 500 spectrometer (at 500 and 125 MHz, respectively). Data for 1H NMR spectra reported as chemical shift δ (ppm), multiplicity, coupling constant (Hz) and integration. Data for 13C NMR spectra reported as chemical shifts relative to the solvent peak. HPLC-MS analysis was performed on a Waters Alliance 2695 separation module connected to a Waters PDA2996 photodiode array detector and a ZQ Micromass mass spectrometer (ESI-MS) with a Phenomenex® column (C18, 5 µm, 4.6 × 150 mm). Compounds were purified using a Waters semi-preparative HPLC system using a Phenomenex® column (C18 Axial, 10 µm, 21.2 × 150 mm) and UV detection.

Synthesis of ethyl 1-formyl-4-oxo-4H-quinolizine-3-carboxylic acid (1)

Ethyl 1-formyl-4-oxo-4H-quinolizine-3-carboxylate (1.04 g, 0.1 mmol, 1 eq.) was dissolved in a mixture of THF (40 mL) and EtOH (40 mL). 1 M aqueous NaOH (5 mL, 1.3 eq.) was added and the reaction was left to stir for 2 h at r.t. Removal of the solvents under vacuum rendered the product (1.11 g) in quantitative yield. 1H NMR (500 MHz, D2O) δ 9.86 (d, J = 1.1 Hz, 1H), 9.37 (ddt, J = 7.3, 1.5, 0.8 Hz, 1H), 9.19 (dd, J = 9.0, 0.8 Hz, 1H), 8.58 (d, J = 0.9 Hz, 1H), 8.13–8.08 (m, 1H), 7.59 (tt, J = 7.1, 1.2 Hz, 1H) ppm. 13C NMR (125 MHz, D2O) δ 191.1, 172.9, 157.4, 146.0, 138.7, 130.0, 122.4, 119.6, 109.4 ppm.

Synthesis and characterisation of BODIPY derivatives (2–4)

Compounds 2, 3 and 4 were synthesized using modified reported procedures30,31 using the aldehyde 1 as the starting material. For the synthesis of compound 2, compound 1 (150 mg, 0.63 mmol, 1 eq.) was dissolved in CH2Cl2 (50 mL) and pyrrole (437 μL, 6.3 mmol, 10 eq.) was then added followed by 1 drop of TFA. The reaction was left to stir overnight under N2. Additional pyrrole (437 μL, 6.3 mmol, 10 eq.) was then added followed by 1 more drop of TFA and the reaction was stirred under N2 for further 7 h. Afterwards, DDQ (143 mg, 0.63 mmol, 1 eq.) in CH2Cl2 (10 mL) was added dropwise and the reaction was left to stir for 20 min. Triethylamine (2.2 mL, 15.75 mmol, 25 eq.) followed by 48% BF3.OEt2 in Et2O (1.2 mL, 9.45 mmol, 15 eq.) were added and the reaction was stirred for 3 h at r.t. under N2. Volatiles were removed under vacuum and flash column chromatography (CH2Cl2/MeOH, 92 : 8) followed by semi-preparative HPLC (0–100% ACN/H2O + 0.1% FA over 25 min) yielded compound 2 (2 mg) as an orange solid. 1H NMR (500 MHz, MeOD) δ 9.54 (d, J = 7.8 Hz, 1H), 8.53–6.37 (m, 1H), 8.03 (d, J = 10.1 Hz, 1H), 8.01–7.85 (m, 2H), 7.80 (s, 1H), 7.63 (dt, J = 13.5, 6.3 Hz, 2H), 6.95 (t, J = 5.0 Hz, 1H), 6.71–6.52 (m, 2H) ppm. 13C NMR (125 MHz, MeOD) δ 167.6, 157.4, 146.0, 138.7, 130.0, 122.4, 119.6, 109.4 ppm.

Conclusions

In summary, we developed a novel BODIPY activatable fluorophore for the detection and imaging of physiological concentrations of Mg2+ and high selectivity over other divalent cations, such as Ca2+. The combined lack of rotational freedom and effective quinolizine-based Mg2+ chelation function as the activation mechanisms for enhanced fluorescence amplification, enabling the imaging of free intracellular Mg2+ ions in live cells and tissues.
2H), 2.68 (s, 6H) ppm. 13C NMR (125 MHz, CD2Cl2) 
160.1, 159.6, 144.8, 141.5, 135.3, 134.6, 129.3, 128.9, 125.6, 123.8, 122.1, 118.7, 110.9, 14.9, 14.7 ppm. HRMS (ESI+) (m/z): [M + H]+ calced for C23H21BF2N3O3 436.1644; found 436.1643.

For the synthesis of compound 3, compound 1 (150 mg, 0.63 mmol, 1 eq.) was dissolved in CH2Cl2 (50 mL), and 2-methylpyrrole (525 µL, 6.3 mmol, 10 eq.) was then added followed by 1 drop of TFA. The reaction was left to stir overnight under N2. Additional 2-methylpyrrole (263 µL, 3.15 mmol, 5 eq.) was then added followed by 1 more drop of TFA and the reaction was stirred for further 7 h. Afterwards, DDQ (0.1 mmol, 1 eq.) was added and the reaction was stirred under N2 for further 20 min. Triethylamine (3.5 mL) was added dropwise and the reaction was left to stir for 20 min. Triethylamine (2.2 mL, 15.75 mmol, 25 eq.) followed by 48% BF3OEt2 in Et2O (1.2 mL, 9.45 mmol, 15 eq.) were added and the reaction was stirred for 3 h at r.t. under N2. Volatiles were removed under vacuum and flash column chromatography (CH2Cl2 : MeOH, 96 : 4) followed by semi-preparative HPLC (0–100% ACN/H2O + 0.1% FA over 25 min) yielded compound 3 (17 mg) as an orange solid. 1H NMR (500 MHz, CDCl3) δ 9.48 (d, J = 7.3 Hz, 1H), 8.63 (s, 1H), 7.90 (d, J = 8.9 Hz, 1H), 7.82–7.73 (m, 1H), 7.56–7.47 (m, 1H), 6.59 (d, J = 4.1 Hz, 2H), 6.32 (d, J = 4.2 Hz, 2H), 2.68 (s, 6H) ppm. 13C NMR (125 MHz, CDCl3) δ 165.4, 160.1, 159.6, 144.8, 141.5, 135.3, 134.6, 129.3, 128.9, 125.6, 125.1, 124.7, 120.3, 118.9, 110.0, 14.7 ppm. HRMS (ESI+) (m/z): [M]+ calced for C23H18BF2N2O3 407.1253; found 407.1251.

For the synthesis of compound 4, compound 2 (26 mg, 0.1 mmol, 1 eq.) was dissolved in CH2Cl2 (5 mL), and 2,4-dimethylpyrrole (20 µL, 0.2 mmol, 2 eq.) was then added followed by 1 drop of TFA. The reaction was left to stir overnight under N2. Afterwards, DDQ (23 mg, 0.1 mmol, 1 eq.) in CH2Cl2 (5 mL) was added dropwise and the reaction was left to stir for 20 min. Triethylamine (350 µL, 2.5 mmol, 25 eq.) followed by 48% BF3OEt2 in diethyl ether (185 µL, 1.5 mmol, 15 eq.) were added and the reaction was stirred for 3 h at r.t. under N2. Volatiles were removed under vacuum and flash column chromatography (CH2Cl2 : MeOH, 95 : 5) followed by semi-preparative HPLC (0–100% ACN/H2O + 0.1% FA over 25 min) yielded compound 4 (13 mg) as an orange solid in a 25% yield over the 3 steps. 1H NMR (500 MHz, CDCl3) δ 9.50 (t, J = 8.8 Hz, 1H), 8.58 (d, J = 8.1 Hz, 1H), 7.88–7.76 (m, 2H), 7.55–7.45 (m, 1H), 6.04 (s, 2H), 2.62 (s, 6H), 1.38 (s, 6H) ppm. 13C NMR (125 MHz, CDCl3) δ 165.6, 160.0, 159.7, 157.6, 144.8, 143.7, 142.0, 141.0, 135.8, 133.1, 131.9, 129.2, 123.8, 122.4, 118.7, 110.9, 14.9, 14.7 ppm. HRMS (ESI+) (m/z): [M + H]+ calced for C23H23BF2N3O4 436.1644; found 436.1643.

**In vitro spectral characterisation**

Spectroscopic and quantum yield data were recorded on a Synergy HT spectrophotometer (Biotek). Compounds were dissolved at the indicated concentrations and spectra were recorded at r.t. Spectra are represented as means from at least two independent experiments with n = 3. When applicable, quantum yields were calculated by measuring the integrated emission area of the fluorescent spectra and comparing it to the area measured for fluorescein in basic ethanol as reference.41

**Fluorescence imaging in A549 cells**

Human lung A549 adenocarcinoma cells (ATCC CCL-185) were grown using Dulbecco’s Modified Eagle Medium (DMEM) supplemented with 10% fetal bovine serum (FBS), antibiotics (100 U mL−1 penicillin and 100 mg mL−1 streptomycin) and 2 mM l-glutamine in a humidified atmosphere at 37 °C with 5% CO2. A549 cells were regularly passaged in T-75 cell culture flasks. For microscopy experiments, cells were plated on glass chamber slides Lab-Tek™ II (Nunc) the day before imaging, reaching 75% to 90% confluence on the day of the experiment. For imaging experiments, cells were incubated for 15 min at 37 °C with compound 4, washed once with PBS and imaged under a Zeiss LSM 510 META fluorescence confocal microscope equipped with a live cell imaging stage. Fluorescence and brightfield images were acquired using a 40× oil objectives. Fluorescent probes were excited with 488 nm (for compound 4) or 543 nm (for MitoTracker Red and LysoTracker Red) lasers. Confocal microscopy images were analysed and processed with ImageJ.

**Fluorescence imaging in Arabidopsis**

*Arabidopsis thaliana* seeds of the Col-0 ecotype were surface sterilised using 10% (v/v) bleach for 10 min, followed by one wash in 70% EtOH, and 5 rinses in sterile ddH2O. Seeds were then plated individually on 0.5× Murashige-Skoog media, solidified with 2% (w/v) Phytoagar. Seeds were stratified at 4 °C for 3 days before being transferred to a vertical position in a controlled growth chamber under long day conditions (16 h light: 8 h dark), with 100 µmol m−2 s−1 white light and a constant temperature of 21 °C. At 11 days after germination, plates were placed flat to encourage the cotyledons to grow away from the media. On day 12, the cotyledons were treated with 0.3 µL of 2.5% (v/v) Adigor for 1 h to improve cotyledon permeability. A 0.3 µL drop of 10 mM compound 4 was then applied to each cotyledon and the seedlings were incubated for around 4 h. The true leaves were then carefully removed, avoiding any contact with the probe, and mounted on glass slides using ddH2O. The leaves were imaged using Leica 5× HC PL Fluotar or 20× HCX APO w objectives on a Leica TCS SP8 Confocal. Samples were excited at 488 nm and emission collected at 500–540 nm for compound 4 and 650–705 nm for chlorophyll.

**Conflicts of interest**

There are no conflicts to declare.

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Notes and references