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Endoplasmic reticulum targeting fluorescent probes to image mobile Zn^{2+} †

Le Fang,^a Giuseppe Trigiante,^b Rachel Crespo-Otero,^a Chris S. Hawes,^c Michael P. Philpott,^b Christopher R. Jones^a and Michael Watkinson^{*c}

Zn^{2+} plays an important role in the normal function of the endoplasmic reticulum (ER) and its deficiency can cause ER stress, which is related to a wide range of diseases. In order to provide tools to better understand the role of mobile Zn^{2+} in ER processes, the first custom designed ER-localised fluorescent Zn^{2+} probes have been developed through the introduction of a cyclohexyl sulfonylurea as an ER-targeting unit with different Zn^{2+} receptors. Experiments *in vitro* and *in cellulo* show that both probes have a good fluorescence switch on response to Zn^{2+} , high selectivity over other cations, low toxicity, ER-specific targeting ability and are efficacious imaging agents for mobile Zn^{2+} in four different cell lines. Probe **9** has been used to detect mobile Zn^{2+} changes under ER stress induced by both tunicamycin or thapsigargin, which indicates that the new probes should allow a better understanding of the mechanisms cells use to respond to dysfunction of zinc homeostasis in the ER and its role in the initiation and progression of diseases to be developed.

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Introduction

The endoplasmic reticulum (ER), an organelle in eukaryotic cells, serves a number of important cellular roles, such as protein synthesis and transport, protein folding, carbohydrate metabolism, and lipid and steroid synthesis.¹ Proteins synthesized in the ER are normally properly folded and transported to the Golgi apparatus, however, when there are changes to ER function, resulting from factors such as ageing, genetic mutations, or the environment, unfolded or misfolded proteins are synthesized and accumulate in the ER,² causing ER stress, which activates the unfolded protein response (UPR).³ It has been found that ER stress can result in a wide range of diseases, such as inflammation,^{4,5} diabetes^{6,7} and neurodegenerative disorders,⁸ including Parkinson's and Alzheimer's diseases.^{9,10} Additionally, the ER acts as an intracellular store for biological mediators, including zinc, which it requires for normal function.^{11,12} For example, it has been found that zinc can be released from thapsigargin- and inositol 1,4,5-trisphosphate (IP3)-sensitive ER storage in cortical neurons.¹³ In light of this, it is unsurprising that a large number of reports demonstrate that

the depletion of zinc transporters and zinc deficiency can cause ER stress and upregulate the UPR,^{14–18} however, the role of 'free' or 'mobile' zinc in this process is little understood due to the lack of suitable molecular tools to image this subcellular region that exist.

Small molecule fluorescent probes, which have great advantages of high sensitivity and selectivity, low toxicity, and good photophysical properties, are increasingly important tools for the investigation of biological events^{19–25} and there have been extensive efforts in the development of a range of probes to detect mobile zinc.^{26–30} However, a failure to control the probe's subcellular location or indeed their different behaviour across cell lines³¹ limits their utility somewhat.^{32,33} Though there is a clear need for ER localised probes to monitor mobile Zn^{2+} flux, to our knowledge there are only two such reports that have appeared in the last decade and in both cases ER localisation was adventitious.^{34,35} Thus, there remains a need for effective and reliable methods to localize small molecule Zn^{2+} probes to target the ER.

The most effective and reliable strategy employed to date to provide organelle targeting capability in small molecule probes has been to incorporate specific organelle targeting vectors into them. For example, the triphenylphosphonium salt (TPP) has been widely demonstrated to be an effective mitochondrial targeting group,^{36–38} whilst basic ethylenediamine³⁹ or morpholine^{40–42} groups have been successfully applied to target the lysosome. However, this strategy has not yet been widely explored in ER localized probes.

Previously we have reported a modular 'click' synthetic methodology to produce an array of biologically targeted Zn^{2+}

^aSchool of Biological and Chemical Science, Queen Mary University of London, The Joseph Priestley Building, Mile End Road, London, E1 4NS, UK

^bCentre for Cutaneous Research, Institute of Cell and Molecular Science, Barts and The London School of Medicine and Dentistry, Queen Mary University of London, London E1 2AT, UK

^cThe Lennard-Jones Laboratories, School of Chemical and Physical Science, Keele University, ST5 5BG, UK. E-mail: m.watkinson@keele.ac.uk

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probes for both extracellular and intracellular imaging of zinc through the incorporation of a range of targeting units.^{43,44} We were therefore hopeful that this approach could be further extended towards the development of ER specific probes, provided that a suitable targeting vector could be identified and incorporated in a modular manner utilising our established chemistry. Herein we present our efforts to this end.

Results and discussion

Probe design and synthesis

Recently a number of reports have appeared in which the methyl sulfonamide group has been used as a targeting unit to visualize hydrogen peroxide,⁴⁵ methylglyoxal,⁴⁶ hypochlorite⁴⁷ and hydrogen sulfide⁴⁸ in the ER. Inspired by this, we first synthesized fluorescent probe **1** (ref. 49) using our modular synthetic protocol that incorporated the methyl sulfonamide unit (Fig. 1) and whilst the fluorescence response of the probe to Zn^{2+} was efficacious *in vitro* and it co-localised to the ER with ER tracker red, it was unfortunately also widely dispersed in other organelles in HeLa cells.

We therefore decided to employ a revised strategy based on the incorporation of glibenclamide as the targeting unit. Glibenclamide, an antidiabetic drug, is employed in the commercial dyes ER tracker red and ER tracker green because it binds and inhibits ATP-sensitive K^+ channels that are abundant in the ER membrane.^{50,51} We were further encouraged by a report of the use of glibenclamide as a targeting group in a small molecule chemical probe which was shown to selectively accumulate in the ER and image glutathione.⁵² We therefore

sought to develop a route to a 'clickable' variant of glibenclamide that we could incorporate into our modular synthetic route and prepared 'click' glibenclamide **2** through six steps in moderate yield (see Scheme S1, ESI†). Gratifyingly the CuAAC reaction with alkyne **5** proceeded, as expected, in good yield (Scheme S2, ESI†), however, to our bemusement, when we came to install the Zn^{2+} binding unit **8**, the second 'click' reaction to give compound **3** failed completely despite a range of Cu(I) sources being tested. Given the success of the first 'click' reaction we decided to investigate the development of a 'reverse' methodology in which the metal-binding unit was first installed into anhydride **9** before addition of the targeting unit *via* a second 'click' process was attempted (see ESI, Scheme S3†). Although the first 'click' reaction again worked well, the second 'click' reaction to incorporate the glibenclamide azide completely failed.

At this point we decided to review our synthetic strategy and were attracted by a number of recent reports that reveal the cyclohexyl sulfonylurea moiety of glibenclamide is an important element in binding with ATP-sensitive K^+ channels, which are present in the ER membrane.^{53–55} We therefore designed an alternative route in which this ER targeting unit was directly incorporated into the naphthalimide, which also makes the synthesis route shorter and easier.

To synthesize the probes, we started from 4-(2-aminoethyl) benzenesulfonamide to give ER-targeting group **6** through the route depicted (see Scheme S4, ESI†) and directly introduced this into the fluorophore through reaction with commercially available 4-bromo-1,8-naphthalic anhydride **5** (Scheme 1). This was then conveniently converted to azide **7** using standard conditions for such $\text{S}_{\text{N}}\text{Ar}$ reactions, which then underwent smooth and high yielding 'click' reactions with alkynes **8** (ref. 56) or **10**,⁴⁴ to yield probes **9** and **12**, the latter after deprotection of **11** with TFA. All products were satisfactorily characterized by ^1H , ^{13}C NMR and IR spectroscopies as well as high-resolution mass spectrometry.

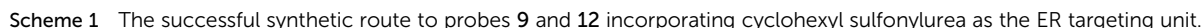
Photophysical properties in solution

Fluorescence titrations with Zn^{2+} were first undertaken to show that both probes have a switch-on response to Zn^{2+} . From Fig. 2a and S2 (ESI†), it can clearly be seen that with increasing Zn^{2+} equivalents, the fluorescence intensity of probes **9** and **12** increased gradually with about 16-fold and 4-fold maximal increase, respectively. Subsequent treatment with TPEN (N,N,N',N' -tetrakis(2-pyridinylmethyl)-1,2-ethanediamine), a strong Zn^{2+} chelator, resulted in the expected decrease in fluorescence with the fluorescence of **9** being completely quenched while that of **12** was partly quenched (Fig. S3, ESI†), which is consistent with previous reports.⁵⁷ The Job's plot (Fig. S4, ESI†) revealed the expected 1:1 binding stoichiometry with Zn^{2+} . The dissociation constants K_{d} for probes **9** and **12** were obtained through non-linear curve fitting using reported methods^{58,59} at a number of probe concentrations (Fig. S5 and Table S1, ESI†). However, because of the very low K_{d} values for these compounds and to mimic the buffered nature of mobile zinc *in cellulo*, EGTA (ethylene glycol-bis(β -aminoethyl ether)-



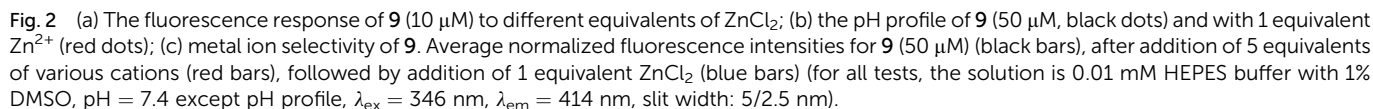
Fig. 1 The structure of probe **1**, glibenclamide azide **2**, unsuccessfully synthesized probe **3** and non-targeting probe **4**.





The quantum yields of both probes were measured using anthracene as the standard ($\Phi = 0.27$ in ethanol). It was found

The pH-dependent fluorescence response of both probes was tested to confirm that they have fluorescence responses to Zn^{2+} in the biologically relevant pH range. Fig. 2b shows that **9** has good switch on fluorescence response to Zn^{2+} over a wide pH



range of 3.0–10.0, and the fluorescence of both probe and complex is increased in an acidic environment. In contrast, **12** (Fig. S11, ESI†) shows a poor response in acid conditions, presumably due to the basic nature of the polyamine ligand, but works well in the pH range 6.0–12.0; given that the reported pH of the ER is the same as the cytoplasm, which is about 7.2,⁶³ both probes should have a response to mobile Zn^{2+} *in cellulo*. Through non-linear curve fitting (applied with the eqn S(4) and S(5)),† the pK_a values of both probes can be obtained, which are $\text{pK}_{a1} = 0.81 \pm 0.19$, $\text{pK}_{a2} = 4.34 \pm 0.11$, $\text{pK}_{a3} = 6.83 \pm 0.53$, $\text{pK}_{a4} = 11.63 \pm 0.41$ for **9**, and $\text{pK}_{a1} = 3.20 \pm 0.14$, $\text{pK}_{a2} = 8.07 \pm 0.11$, $\text{pK}_{a3} = 11.07 \pm 0.58$ for **12** (Fig. S10 and S12, ESI†). These values are comparable to those reported for related compounds.⁴⁴

The selectivity of the two probes over a range of other competing cations was also investigated. As shown in Fig. 2c, beside Zn^{2+} , the fluorescence of **9** was not switched on obviously after addition of 5 equivalents of other cations, except for the stereoelectronic isostere Cd^{2+} , which is a common issue for many Zn^{2+} probes,⁶⁴ but is not a problem in biological milieu due to its negligibly low concentration. Subsequent addition of Zn^{2+} resulted in recovery of a fluorescence response in most cases, however for Co^{2+} , Cu^{2+} and Ni^{2+} fluorescence was not recovered, but as they mostly exist in bound forms in biology, rather than the free cations tested here this is un concerning. For **12**, results were similar, as shown in Fig. S13 (see ESI†), with only Cu^{2+} , Fe^{2+} and Fe^{3+} continuing to quench fluorescence after the addition of Zn^{2+} . Importantly, the metal ions Na^+ , K^+ , Ca^{2+} and Mg^{2+} , which are abundant in cells, show no effect on either probe. Overall these results indicate that they should show a selective response to ‘mobile’ Zn^{2+} *in cellulo*.

DFT simulations

DFT and TDDFT calculations were undertaken to understand the binding behaviour of **9** and **12** with Zn^{2+} and their excited

states. The optimised structure of probe **9** with 1 equivalent of Zn^{2+} (Fig. S14, ESI†) shows the Zn^{2+} is bound to the nitrogen in ligand *N,N*-di-(2-picoly)ethylenediamine (DPEN) and a nitrogen atom of the triazole. For **12** (Fig. S15, ESI†), the triazole nitrogen is involved in binding with Zn^{2+} in addition to the cyclam nitrogen donors, which is consistent with previous observations in closely related analogues.^{65,66}

The calculated vertical excitations and emissions are in relatively good agreement with experimental data (see Table S3 and S4, ESI†). The excited state calculations also show that positions of the absorption maximum and the emission energies are not significantly affected by the formation of the complexes with Zn^{2+} as observed experimentally. This behaviour can be understood by visualising the $\text{S}_1\text{--}\text{S}_0$ electron density plots (Fig. S16 for **9**, Fig. S17 for **12**, ESI†). The electronic transition is localised on the naphthalimide moiety and the vicinal triazole, and that there is no major difference between probes and their complexes with Zn^{2+} ; only a slight decrease of electron density on the nitrogen from the triazole is observed when it is involved in complex formation and the oscillator strength is not significantly affected. These calculations indicate that the enhancement of emissive behaviour of the complexes should be related to a reduced decay through nonradiative pathways after



Fig. 3 The colocalization images of HeLa cells incubated with **9** (20 μM , GFP filter: $\lambda_{\text{ex}} = 470/30$ nm, $\lambda_{\text{em}} = 530/50$ nm) and commercial red organelle tracker dyes (RFP filter: $\lambda_{\text{ex}} = 530/40$ nm, $\lambda_{\text{em}} = 605/55$ nm). (Scale bars = 20 μm).



Fig. 4 (a) Fluorescence microscopy images of HeLa cells treated with **9** (20 μM), **9** (20 μM) with zinc pyrithione (100 μM), and **9** (20 μM) with TPEN (100 μM) (Scale bars = 20 μm); (b) the fluorescence intensity (F) in different cell lines in the presence of zinc pyrithione (red bars) or TPEN (blue bars) relative to the intensity of the probe alone (F_0 , black bars, normalised to 1).





Fig. 5 (a) Fluorescence microscopy images and (b) fluorescence intensity changes of probe 9 (20 μM) in HepG2 cells at ER stress state induced by tunicamycin (5 μg mL⁻¹) or thapsigargin (1 μM). (Scale bars = 20 μm).

complexation with Zn²⁺ in contrast with the typical PET mechanism. The role of alternative mechanisms to PET in enhancing the fluorescence of organometallic complexes has been recently highlighted.^{67,68} The formation of the complex hinders large amplitude vibrations in the vicinity of the fluorophore hampering access to nonradiative mechanisms and increasing the quantum yield of emission. The restriction of intramolecular rotations can also hinder the access to low energy conical intersections associated with ultrafast decay to the ground state. Tang *et al.* have recently shown how restriction of intramolecular motions in a Zn²⁺ complex can hamper the access to a dark state enhancing the fluorescence response.⁶⁹

ER-targeting ability

Encouraged by the promising properties of **9** and **12** *in vitro*, we attempted to assess their suitability for imaging Zn²⁺ in cells. Firstly, the innate toxicity of them was measured by alamarBlue assays with HeLa cells (Fig. S18 and S19, ESI†).²² The cell viabilities were over 90% after 24 h incubation with both probes in a concentration range 10–50 μM compared to the control group, demonstrating both have very low cytotoxicity.⁷⁰

The targeting behaviour of **9** and **12** was then studied using co-staining assays. HeLa cells were co-incubated with **9** and ER-tracker red, and as shown in Fig. 3, the fluorescence of **9** from the green channel overlays well with that of ER-tracker red, which was obtained from the red channel, giving an excellent Pearson's correlation coefficient of 0.92. The negative controls of **9** co-incubated with Mito-tracker red and Lyso-tracker red, did not reveal a good overlay and Pearson's correlation coefficients were 0.52 and 0.58, respectively. Similar results were observed for **12** (see Fig. S22, ESI†) with Pearson's coefficients of 0.85, 0.52, 0.45 being measured for the ER, mitochondria and lysosome respectively. A control probe that did not contain a targeting unit **4** (ref. 49) was observed to be widely distributed in all three organelles (Fig. S23, ESI†) and this is also consistent with our original study.⁶⁵

To confirm the probes have ER-targeting ability in different cell types, a human breast cancer cell line MCF-7, a sweat gland tissue cell line EC23 and a human liver cancer cell line HepG2 were used to study the ER-targeting behaviour. The results (see Fig. S24–S26 for probe **9**, Fig. S27–S29 for probe **12**, ESI†) show that probe localisation is similar to that observed in HeLa cells demonstrating the general applicability of the probes. Based on these data, we can conclude that both **9** and **12** show excellent, and general, ER-targeting capability.

Zn²⁺ fluorescence response in cells

As **9** and **12** can localise to the ER, their fluorescence response to 'mobile' Zn²⁺ *in cellulo* was tested to demonstrate that they have the ability to image Zn²⁺ in the ER. As shown in Fig. 4a, the fluorescence response of probe **9** in HeLa cells could be clearly observed in the ER. Upon the addition of zinc pyrithione, a membrane permeable zinc source, the fluorescence intensity increased considerably; the addition of TPEN resulted in almost complete quenching of fluorescence, as expected from *in vitro* control experiments. The fluorescence intensity read from these images (Fig. 4b) clearly reveals these changes. For example, in HeLa cells the intensity of the fluorescence response increases 3.6 times after the addition of zinc pyrithione and decreases by 30% on addition of TPEN compared to that observed for **9** alone. Similar results were observed in other cell lines and also for **12** (Fig. 4b and S30–S36, ESI†) demonstrating that both probes are tractable in their ability to image mobile Zn²⁺ *in cellulo*.

Fluorescence change under ER stress

As **9** shows excellent ER-localisation and Zn²⁺ response, we sought to explore whether it could be applied to monitor changes in mobile Zn²⁺ levels. It is reported that ER stress can be induced by tunicamycin and thapsigargin with tunicamycin causing unfolded protein accumulation and thapsigargin



reducing Ca^{2+} levels. As shown in Fig. 5, the fluorescence intensity of **9** in HepG2 cells decreased significantly after treatment with both ER stress inducers, indicating that the mobile Zn^{2+} level in the ER was significantly depleted under ER stress within 1 hour. As a control, no fluorescence intensity change was observed in their absence (Fig. S37, ESI†).

Conclusions

In conclusion, the ER localised fluorescent probes **9** and **12** have been developed to image mobile Zn^{2+} in the ER through the introduction of a simple sulfonylurea targeting group. They show a good fluorescence response to Zn^{2+} , high selectivity over biologically relevant competing cations, and can be used to detect Zn^{2+} over a wide range of pH *in vitro*. Their low toxicity, ER-targeting and ability to image mobile Zn^{2+} in different cell lines should allow them to be applied in imaging Zn^{2+} related biological processes in the ER. The fluorescence intensity of **9** was also shown to decrease under ER stress inducers, indicating that mobile Zn^{2+} levels are reduced. The ER localised probes may therefore provide a platform to understand the mechanisms cells use to respond to dysfunction of zinc homeostasis in the ER, and to develop a better understanding of the role of Zn^{2+} in the initiation and progression of related diseases.

Conflicts of interest

There are no conflicts to declare.

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