Fluorescent probes for the detection of magnesium ions (Mg$^{2+}$): from design to application

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Magnesium ions (Mg$^{2+}$) play essential roles in various physiological and pathological processes, its abnormal homeostasis in cells is related to many diseases, such as diabetes, neuromuscular disorders, hypertension and other cardiovascular disorders. Investigation on the regulation of magnesium in cellular processes has attracted considerable interest in the past several decades. Among those reported strategies, fluorescent imaging technology has become a powerful and cost-effective tool for the real-time monitoring of magnesium distribution, uptake and trafficking, due to its superior features of high sensitivity and non-invasiveness, as well as excellent spatial and temporal fidelity. Herein, we critically summarize the progresses in the intracellular magnesium detection with fluorescent imaging probes. Our discussion focuses on the recent contributions concerning fluorescent imaging probes for mapping magnesium in biological processes. All the candidates are organized according to their acceptor structures. The sensing mechanisms of fluorescent probes are also highly taken into account. Challenges, trends and prospects of fluorescent imaging technology in magnesium detection are also set forth.

1. Introduction

The magnesium ion (Mg$^{2+}$) is one of the most abundant essential elements in cells and plays a critical role in numerous cellular processes such as cell proliferation, cell death, enzyme-driven biochemical reactions, channel regulation, stabilization of DNA conformation and signal transduction.$^{1-3}$ There are approximately 300 enzymatic reactions mediated by Mg$^{2+}$ in cells, and the concentration change of intracellular Mg$^{2+}$ is believed to regulate the metabolism of cells.$^4$ In the majority of mammalian cells, intracellular free Mg$^{2+}$ usually ranges from 0.5 mM to 0.7 mM, maintained by Mg$^{2+}$ transport across the cell membrane, intracellular buffering and compartmentation into organelles.$^5$ In medical settings, hypomagnesemia can result from various causes including hypertension, diabetes, osteoporosis, metabolic syndrome, neuronal injury, cell senescence and antitumor efficacy of chemotherapeutics. By contrast, hypermagnesemia occurs less frequently than other common diseases, however, it associates significantly with chronic renal failure and even results in cardiac arrest in extreme cases.$^6$ In addition, magnesium sulfate is presently employed to treat numerous conditions such as arrhythmias, myocardial infarction and eclampsia.$^8$ Therefore, it is of great value and urgency to develop simple, rapid and efficient analytical methods to detect magnesium mobilization, distribution and concentration changes in cells.

Over the last few decades, several conventional analytical methods have been put forward to measure Mg$^{2+}$ in either fixed or live samples. These methods have respective benefits and drawbacks and can be exploited with variable levels of confidence in studies of specific issues. Atomic absorption spectroscopy (AAS) developed by Australian chemist Walsh in the 1950s (ref. 9) has been historically the first and most frequently used technique to detect magnesium in acidic extracts of biological samples. The introduction of AAS partially obviated methodological limitations: the system made it possible to measure the level of magnesium in all biological samples, with a lower detection limit of 10 mM. However, AAS could only better our understanding of magnesium metabolism to some extent. In fact, investigating the physiological dynamics of Mg$^{2+}$ at the cellular level needs techniques that ascendant detection of Mg$^{2+}$ movements across the diversiform compartments of a living cell. Radioactive isotopes ($^{25}$Mg and $^{26}$Mg) were originally used to measure magnesium fluxes but have been abandoned on account of biohazard, short half-life and very high production costs. They are currently replaced by their stable counterparts ($^{24}$Mg or $^{25}$Mg), which can be used in assessment of exchangeable magnesium pool masses as well as the evaluation of magnesium status and fluxes in animals and humans.$^10$ However, the method is costly and demands of detection by mass spectrometry (or better, ICP-MS), which is not easily accessible.
Another sophisticated technique is phosphorus nuclear magnetic resonance spectroscopy ($^{31}$P-NMR), which depends on the fact that most magnesium in the cytosolic matrix is bound to ATP (adenosine triphosphate). When magnesium is bounded to ATP, it shifts the resonance frequencies of signals (chemical shift) coming from the three phosphoric groups of the molecule. Due to the chemical equilibrium between bound and free magnesium, the chemical shift of ATP signals is a function of free Mg$^{2+}$ concentration. $^{31}$P-NMR offers the unique opportunity to measure in vivo cytosolic Mg$^{2+}$ concentration in several tissues and has provided new hints on magnesium homeostasis and its involvement in the cellular bioenergetics of human brain and skeletal muscle. However, this method requires cumbersome sample pretreatment, time-consuming and complicated instruments. What’s more, no information on intracellular magnesium distribution can be derived.

Due to the weaknesses of these methods, it inspires the creation of new analytical methods to help further our understanding of the distribution, uptake, and trafficking of Mg$^{2+}$ in living systems. Hence, fluorescent imaging technologies were developed and stand out as the most effective approach because of its simplicity, high sensitivity, excellent spatial–temporal resolution, fast response and non-invasiveness. Over the last decades, a boom of fluorescent imaging techniques have been witnessed among the most powerful platforms for dynamic behaviour determination and real-time observation in a single living cell, which were also expected to be used in magnesium research. As a result, a variety of novel fluorescent probes for detection of Mg$^{2+}$ have been reported. Together with the rapid development of confocal laser scanning and two-photon microscopy (TPM) techniques, fluorescence methods now provide a robust tool to visualize and measure Mg$^{2+}$ in three dimensions, even deep into tissues.

To date, a plethora of fluorescent probes for Mg$^{2+}$ have been developed and this research field has become very active and exuberant. Therefore, a systematic survey of progress in the rapid development of fluorescent probes for Mg$^{2+}$ is highly demanded. Herein, we will focus on the recent progress in the creation of novel fluorescent imaging probes for Mg$^{2+}$ and their applications in cell biology. Firstly, to gain a better understanding of the rational design approaches and discuss how the fluorescence probes react with the analytes, we will present and outline the general principles of designing effective fluorescent imaging probes for biological applications. Then, we will discuss advances in Mg$^{2+}$ fluorescent imaging probes, which are classified according to different types of Mg$^{2+}$-selective binding sites, for highly specific sensing and detecting Mg$^{2+}$ in cellular systems. In the end, we will conclude with a discussion of the reality and future challenges in this field from the perspective of available probes for fluorescence imaging detection of cellular Mg$^{2+}$.

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2. The general design principles of functional fluorescent probes

Many photophysical processes (quenching via collision, photoinduced electron transfer, exciplex formation, photoinduced proton transfer, energy transfer and etc) may be responsible for the fluorescence changes (excitation and emission spectra, quantum yield, fluorescent lifetime and etc) upon binding of the analyte by the probes. The most widely utilized fluorescence signaling mechanisms in the design of probes will be briefly discussed in this section.

2.1 Photoinduced electron transfer (PET)

Photoinduced electron transfer (PET) is possible the cause of fluorescence quenching, when the PET process is followed by a nonluminescent process back to the normal state. The PET process can be illuminated picturesquely based on the molecular orbital theory, therefore becomes a powerful tool to expound the fluorescence "on-off" switching of fluorescent probes. Fluorescent probes on the basis of PET are often structured as fluorophore-spacer-chelator constructs. In this pattern, an aliphatic spacer separates the fluorophore and chelator, thus electronically disconnect the π-electron systems of them. According to the electron accepting or donating relationships between fluorophore and chelator in one fluorescent probe, PET can be subdivided into the reductive PET and the oxidative PET. This review will focus on the reductive PET due to its frequent use in the design of fluorescent probes (as depicted in Fig. 1). In reductive PET, the fluorophore is reduced while the chelator is oxidized, that is the chelator acts as the electron donor (e.g., the amine group) and the fluorophore serves as the electron acceptor. During the reductive PET process, photo-excitation promotes an electron from the HOMO (the highest occupied molecular orbital) of the fluorophore to its LUMO (the lowest unoccupied molecular orbital). Subsequently, the reaction between the fluorophore and the analytes induces either the appearance or removal of the "nearby" orbital between HOMO and LUMO of the fluorophore, leading to fluorescence quenching or enhancement, respectively.

2.2 Intramolecular charge transfer (ICT)

The intramolecular charge transfer (ICT) strategy for cation sensing was introduced by Valeur. Probes based on ICT are featured by conjugation of an electron-donating unit to an electron-accepting unit in one molecule to rise a "push–pull" π-electron system in the excited state, which have been extensively used for cation sensing. While the electron-donating part interacts with an analyte, the electron-donating character of the probe decreases, generating a blue shift in the absorption spectrum. In contrast, an evident red shift is observed when the ICT becomes more developed owing to the interaction of an analyte with the electron-accepting part (as described in Fig. 2). Besides, changes in the fluorescence lifetimes and quantum yields are also observed. Up to now, a great number of fluorescent imaging molecules are derived from this ICT mechanism by changing either the electron-donating, electron-withdrawing ability or π-conjugation degree of the fluorophores to interact with the target analyte.

2.3 Excited-state intramolecular proton transfer (ESIPT)

Excited-state intramolecular proton transfer (ESIPT) has also been adopted for designing fluorescent imaging probes because...
of its unique and excellent spectral sensitivity to the environmental medium. ESIPT process typically involves a fast proton transfer from a proton donor (often hydroxyl or amino unit) to an acceptor group (usually either carbonyl oxygen or imine nitrogen) atom in the excited state of a fluorophore mediated by an intramolecular hydrogen bond. A pictorial representation of ESIPT process is depicted in Fig. 3.

Owing to the transformation between the enol and keto form in the ESIPT-based system, as such, the ESIPT process not only radically reduces the photochemical reactivity of the excited molecules, but also significantly enhances their photostability. On top of this, a large manifest Stokes shift can also be observed. Therefore, the ESIPT process is quite suitable for designing fluorescent imaging probes requiring spectral shift for selective detection.

2.4 Fluorescent resonance energy transfer (FRET)

Fluorescence resonance energy transfer (FRET) mechanism comprises energy transfer between a pair of fluorophores that serve respectively as energy donor and acceptor. FRET is a distance-dependent interaction between the electronic excited states of two chromophores in which excitation energy is non-radiatively transferred from a donor chromophore to an acceptor chromophore via nonradiative dipole–dipole coupling (as represented in Fig. 4). In those transfer processes, several vibronic transitions in the donor have approximately equivalent energy to those in the acceptor. Molecules in which FRET is operative are quintessentially used to artificially increase the Stokes shift. As applied in sensing applications, the emission of the donor at relatively short wavelengths serves to activate emission of the acceptor at longer wavelengths with the ratio of the fluorescence intensities of the donor and acceptor emissions modulated by the target analytes. To obtain a high FRET efficiency, it is required that spectral overlap between the donor emission and the acceptor absorption spectrum is substantial. Many FRET-based fluorescent probes have been created subsequent to these pioneering efforts for the ratiometric detection of metal ions, such as Hg²⁺, Zn²⁺. So far, no FRET-based fluorescent probes for Mg²⁺ have been reported, which suggests that it is a hard nut to crack in this field.

3. Fluorescent probes for magnesium ions based on different binding sites

Besides the fluorescence signaling mechanisms, there are several important characteristics of fluorescent imaging probes must to be tuned when selecting or designing a probe for imaging the analyte in living cellular systems, such as the excitation/emission wavelength, photostability, fluorescence quantum yield, fluorescence lifetime and so on. Conceivably, the most significant factor to consider when choosing a fluorescent probe is its binding affinity, which mainly depends on the acceptor. The acceptor, in other words, is the selective binding sites and dictates the sensitivity to molecular/ion recognition. To be applied in living cellular systems, a high-affinity Mg²⁺ binding probe should be selective for Mg²⁺ over...
other biologically abundant ions, which exist at much higher cellular concentrations. In the further section, we will try to provide a critical overview of Mg\textsuperscript{2+} fluorescent probes based on different binding sites known by now, with particular regard to the latest advances.

3.1 Fluorescent probes for Mg\textsuperscript{2+} based on o-aminophenol-\(N,N,O\)-triacetic acid (APTRA)

The \(o\)-aminophenol-\(N,N,O\)-triacetic acid (APTRA) structure, which is the most common acceptor, is generally deemed to be the Mg\textsuperscript{2+}-selective binding site (its structure is shown in Fig. 5). The first widely available fluorescent Mg\textsuperscript{2+} probe was anchored in the chemical design by Roger Tsien, who rationally devised fluorescent Ca\textsuperscript{2+} probes for intracellular use in the late 1970s.\textsuperscript{27,28} The prototype of such Ca\textsuperscript{2+} probes was modified as fluorescent probes for measuring cytosolic free Mg\textsuperscript{2+} by Robert London.\textsuperscript{29,30} One modification method is that fluorinated derivatives of the chelator APTRA have been designed and synthesized for use as \(19^F\)-NMR indicators of free cytosolic Mg\textsuperscript{2+} concentration. Another way is that the APTRA structure is modified around the furan ring, resulting in the fluorescent probe \textit{Mag-fura-2} (as shown in Fig. 5).\textsuperscript{31} From then on, many other dyes purported to be selective for Mg\textsuperscript{2+} have been put forward, and most of them are simply analogues of the corresponding Ca\textsuperscript{2+} probes: for example, \textit{Mag-indo-1} (analogue of indo-1) and \textit{Magnesium Green} (analogue of calcium green) (see Fig. 5 for their chemical structures). It’s obvious that the chemical structure of those probes contain a common APTRA unit as Mg\textsuperscript{2+}-selective binding site.

\textit{Mag-fura-2}, the first fluorescent Mg\textsuperscript{2+} probe, was designed to visualize and understand intracellular Mg\textsuperscript{2+} distributions.\textsuperscript{32} \textit{Mag-fura-2} is excited by ultraviolet (UV) light (\(\lambda\text{max} = 369\) nm) showing a blue shift to 330 nm in absorption spectra upon Mg\textsuperscript{2+} binding, and also a shift in emission spectrum in the visible region from 511 nm in Mg\textsuperscript{2+}-free medium to 491 nm at saturating Mg\textsuperscript{2+} concentrations. The binding dissociation constant for Mg\textsuperscript{2+} is 1.9 mM. \textit{Mag-fura-2} has been exploited to measure the intracellular Mg\textsuperscript{2+} concentration in cells from the muscle, heart, liver, and nervous system.\textsuperscript{33,35}

Thereafter, \textit{Mag-indo-1} has been synthesized on the strength of identical molecular design as \textit{Mag-fura-2}.\textsuperscript{36} \textit{Mag-indo-1} has a dissociation constant (\(K_d\)) of 2.7 mM for Mg\textsuperscript{2+} and is excited by UV light to emit fluorescence in the visible region. \textit{Mag-fura-2} undergoes a palpable shift in excitation wavelength after binding to Mg\textsuperscript{2+}, whereas \textit{Mag-indo-1} presents a shift in both its excitation and emission wavelengths. The \(K_d\) of \textit{Mag-indo-1} for Mg\textsuperscript{2+} is lower than that of \textit{Mag-fura-2} (the parameter is shown in Table 1), which is conducive for the measurement of high spikes in intracellular Mg\textsuperscript{2+}, for example the temporal analysis of Ca\textsuperscript{2+}-induced Mg\textsuperscript{2+} mobilization in neurons or the glutamate-stimulated Mg\textsuperscript{2+} concentration change.\textsuperscript{37-39}

Later, several other fluorescent probes which possess the APTRA group as the binding site for Mg\textsuperscript{2+} have been developed, such as \textit{Mag-fluo-4} and \textit{Magnesium Green}.\textsuperscript{40,41} The remarkable feature of those probes compared to the aforementioned ones is that they are excited by visible light (see Table 1) rather than UV irradiation which is regarded to be more cytotoxic.\textsuperscript{42} The maximum excitation and emission of \textit{Magnesium Green} are 490 nm and 520 nm, respectively. Upon binding to Mg\textsuperscript{2+}, the fluorescence intensity of \textit{Magnesium Green} increases without any shift in the wavelength. Moreover, \textit{Magnesium Green} presents a higher affinity for Mg\textsuperscript{2+} compared to both \textit{Mag-fura-2} and \textit{Mag-indo-1}. In biological applications, \textit{Magnesium Green} has been exploited to probe intracellular Mg\textsuperscript{2+} for the exploration of the binding of free Mg\textsuperscript{2+} by the bacterial SecA protein and the investigation of ATP hydrolysis in spontaneously contracting cardiomyocytes.\textsuperscript{43,44}

Unfortunately, Mg\textsuperscript{2+} fluorescent probes possessing APTRA as binding site lack the appropriate selectivity for Mg\textsuperscript{2+} over Ca\textsuperscript{2+}, which implies that Ca\textsuperscript{2+} become a confounding factor in Mg\textsuperscript{2+} detection when Ca\textsuperscript{2+} concentrations exceed about 1 \(\mu\)M.\textsuperscript{45} Therefore, Ca\textsuperscript{2+} disturbs the accuracy of Mg\textsuperscript{2+} measurement and limits the interpretation of the read-out eventually. To deal with this problem, Bong Rae Cho and co-workers developed efficient two-photon fluorescent probes \textit{FMg1} and \textit{FMg2} (structures are shown in Fig. 5) to simultaneously monitor Mg\textsuperscript{2+}/Ca\textsuperscript{2+} activities.\textsuperscript{46} Through using BCaM and \textit{FMg2}, one can simultaneously detect Ca\textsuperscript{2+}/Mg\textsuperscript{2+} activities in live cells and even in live tissues at a depth of more than 100 \(\mu\)m for long periods of time without photobleaching artifacts. What is worth mentioning is that BCaM, a two-photon probe for near membrane Ca\textsuperscript{2+} was also designed by the same research team.\textsuperscript{47} In \textit{FMg1} and \textit{FMg2}, the

| Probes       | Fluorophore   | \(\lambda_{\text{max,abs}}\) (nm) | \(\lambda_{\text{max,em}}\) (nm) | \(K_d\) (Mg\textsuperscript{2+})\textsuperscript{a} | \(K_d\) (Ca\textsuperscript{2+}) | Ref. |
|--------------|---------------|---------------------------------|---------------------------------|---------------------------------|--------------------------------|-----|
| Mag-fura-2   | Benzofuran    | 369/330                        | 511/491                         | 1.9 mM                          | 25 \(\mu\)M                    | 31  |
| Mag-indo-1   | Indole        | 349/330                        | 480/417                         | 2.7 mM                          | 35 \(\mu\)M                    | 36  |
| Mag-fluo-4   | Fluorescein   | 490/493                        | None/517                        | 4.7 mM                          | 22 \(\mu\)M                    | 40  |
| Magnesium-Green | Fluorescein | 506/none                       | None/531                        | 5.0 mM                          | 6.0 \(\mu\)M                   | 41  |
| FMg1         | Fluorene      | 362/none                       | 540/none                        | 1.5 mM                          | 8.8 \(\mu\)M                   | 46  |
| FMg2         | Fluorene      | 368/none                       | 555/none                        | 1.7 mM                          | 9.8 \(\mu\)M                   | 46  |
| AMg1         | Naphthalene   | 365/none                       | 485/none                        | 1.4 mM                          | 9.0 \(\mu\)M                   | 48  |
| Mag-S        | Furan         | 396/330                        | 572/547                         | 3.2 mM                          | 48 \(\mu\)M                    | 49  |
| Mag-Se       | Furan         | 412/360                        | 584/562                         | 3.3 mM                          | 41 \(\mu\)M                    | 49  |

\(\lambda\text{max,abs}\) excitation wavelength; \(\lambda\text{max,em}\) emission wavelength; \(K_d\) dissociation constant; \(K_d\) values were measured in aqueous buffers; for details please see the cited references.\textsuperscript{a} None represents the spectral no change.
APTRA group serves as the receptor for Mg\(^{2+}\), while 2-acetyl-7-diethylamino-9, 9-dimethyl-9H-fluorene and 2-diethylamino-7-nanonoyl-9,9-dimethyl-9H-fluorene acts as the fluorophore, respectively. The acetoxymethyl (AM) ester form of them (FMg1-AM and FMg2-AM) can passively diffuse through the plasma membrane and readily stand enzymatic hydrolysis to reform the metal-ion probe inside the cell. FMg1 and FMg2 display absorption maxima at 362 nm and 368 nm with fluorescence maxima at 540 nm and 555 nm, respectively. When Mg\(^{2+}\) is added to FMg1 and FMg2 in MOPS buffer, the fluorescence intensity increases remarkably without impact on absorption spectra, which may be a result of the blocking of the PET from APTRA to fluorophore upon chelation with Mg\(^{2+}\). The \(K_d\) values of FMg1 and FMg2 for Mg\(^{2+}\) are 1.5 mM and 1.7 mM, respectively (see Table 1). FMg2-AM in combination with BCaM are utilized as two-photon probes to investigate Mg\(^{2+}\)/Ca\(^{2+}\) activities in HepG2 cells by dual-color imaging.

As noted above, the two-photon microscopy (TPM) is able to visualize the distribution of intracellular metal ions in living cells and tissues. TPM utilizes two photons with lower energy as the excitation source, is getting increasingly prevalent among biologists owing to several distinct advantages. Two-photon probes offer numerous unique benefits over one-photon probes, from increased penetration depth (>500 mm), lower binding site. The maximum excitation wavelength of metal-ion complexation. The increase, a dramatic enhancement in the fluorescence is getting increasingly prevalent among biologists owing to several distinct advantages. Two-photon fluorescence is

3.2 Fluorescent probes for Mg\(^{2+}\) based on charged β-diketone binding site

There is a great demand for the development of optimized Mg\(^{2+}\) fluorescent probes to achieve the dynamic and three-dimensional imaging of intracellular Mg\(^{2+}\). Alternative acceptor featuring a charged β-diketone or related β-dicarbonyl-bidentate binding motifs has been efficiently incorporated into fluorescent probes for the measurement of Mg\(^{2+}\) (as shown in Fig. 6).

As early as 2002, Suzuki et al. reported the development of two novel Mg\(^{2+}\) fluorescent imaging probes KMG-20-AM and KMG-27-AM (as shown in Fig. 6), in which AM is an acetoxymethyl group, grounded on a coumarin possessing a charged β-diketone substructure. Both probes have a β-hydroxycarboxylic acid group and an aromatic amino group combined

Live cell fluorescence imaging of Mg\(^{2+}\). Mag-S and Mag-Se show a promising result in reality. Mag-S and Mag-Se display excitation maximum at 392 nm and 410 nm in the free-state, respectively, while the two probes show emission maximum at 572 nm and 584 nm in unbound form, respectively. Hence, a single replacement of O by S or Se in the structure of the fluorophore generates a bathochromic shift greater than 60 nm in the emission of the probe. Both Mag-S and Mag-Se's responses to Mg\(^{2+}\) show a significant hypochromic shift in the fluorescence excitation and emission spectra. The separation between excitation bands for the metal-free and bound probe anchored to the electron pooper thiazole and selenazole is dramatically larger than that of the oxazole derivative. This feature is desirable for ratiometric imaging and the test for responding of Mag-S for the ratiometric measurement of Mg\(^{2+}\) in HeLa cells was performed. Recently, the same research group using Mag-S as the parent probe introduced a clever two-step strategy for in situ anchoring and activation of fluorescent sensing Mg\(^{2+}\) within intracellular organelles of choice. Step one, they choose append a tetrazine moiety to Mag-S through a fluoreogenic reaction; step two, as trained bicycloonynone in the structure of fluorophore covalently linked to a genetically encoded HaloTag fusion protein of known cellular localization. Experiments proved that the sensing system could realize the aim for the ratiometric detection of Mg\(^{2+}\) in target organelles in HEK293T cells. The labelling strategy is, furthermore, fully compatible with live cell imaging and enables the real time visualization of changes in metal ion distribution involved in cellular processes, providing a valuable tool for tracking changes in magnesium distribution that to date have been an unsolved mystery in the cell biology of Mg\(^{2+}\).

However, the other photophysical properties of the probes do not improve significantly. For example, the interference of Ca\(^{2+}\) or other biologically relevant metal ions is not negligible. Moreover, the fluorescence quantum yield is lower. On top of these, the dissociation constants of Mag-S and Mag-Se decrease slightly in comparison to Mag-fura-2. To sum up, the chalcogen replacement strategy may open the doors to enhance red-shifted ratiometric detection for other biologically relevant ions.
measurement of intracellular Mg\(^{2+}\) concentration imaging with fluorescent microscopy in living cells. Take PC12 cells as an example, the addition of KMG-20-AM and KMG-27-AM into PC12 cells led to a strong fluorescence in the cytoplasm and a weak fluorescence in the nuclear region. After treatment with high-K\(^+\) medium, the fluorescence intensity increases owing to increasing intracellular Mg\(^{2+}\) concentrations. High K\(^+\) can depolarize the cell membrane and activates several ion channels, therefore, Mg\(^{2+}\) is released from intracellular stores, making it possible for KMG-20-AM and KMG-27-AM to image Mg\(^{2+}\) distribution successfully. As another application of KMG-20-AM, Park and co-workers developed Mg\(^{2+}\)-sensitive nanoparticles for intracellular use by PEBBLE (probe encapsulated by biologically localized embedding) technology.\(^{22}\) The particles are stable but show affinities and spectroscopic limitations analogous with those of KMG-20AM.

Fluorescein and rhodamine are known to be excitable at long wavelength, and have a high fluorescence quantum yield in an aqueous medium, as is generally used in biological experiments.\(^{23}\) Nagano and co-workers developed four novel fluorescent Mg\(^{2+}\) probes, 2’-CF, 2’-CF-Cl, 2’-CF-F and 2’-CFR (these probes are shown in Fig. 6). All of them contain 2’-carboxy-fluorescein or its derivatives as fluorophore, and a carboxyl group at the 2’-position to make up a \(\beta\)-keto acid moiety as the Mg\(^{2+}\) chelating site.\(^{24}\) Those probes could be used in the confocal microscope, however, they showed a lower affinity for Mg\(^{2+}\) (\(K_d = 15.8\) mM of 2’-CF, other parameters are shown in Table 2) and suffered from strong pH interference, therefore, the project were forced to be abandoned. Then, the focus shifts to the KMG series. Similar to KMG-20-AM, these compounds possess a charged \(\beta\)-diketone as a specific binding site for Mg\(^{2+}\), which is attached to the 9’-position of xanthene (as presented in Fig. 6), resulting in the novel Mg\(^{2+}\) fluorescent probes, KMG-101, -103, -104, and KMG-301.\(^{53,26}\) This molecular design results in an intensive off-on-type fluorescent response toward Mg\(^{2+}\), showing both the highly sensitive photoinduced electron transfer-type (PET) response and Mg\(^{2+}\) affinity modification, as well as the allowance of Ar\(^+\) laser excitation. Among the four probes mentioned above, KMG-104 possesses the most desirable properties, for it can be excited by the 488 nm line of Ar lasers and its fluorescence emission rises along with Mg\(^{2+}\) concentrations, while the absorption spectrum only displays minor changes. In addition, its \(K_d\) is about 2.1 mM. Therefore, comparable to that of APTRA-based probes, the new dye has a better affinity for Mg\(^{2+}\) and is suitable to detect free intracellular Mg\(^{2+}\). No response of it to alkali metals is observed yet. More interestingly, KMG-104 displays a \(K_d\) for Ca\(^{2+}\) of 7.5 mM, which is much higher than the resting physiological free Ca\(^{2+}\) concentration. Therefore, the selectivity of this probe for Mg\(^{2+}\) over Ca\(^{2+}\) is much superior to that exhibited by Mag-fura-2 and Mag-fluo-4. Moreover, the fluorescence of KMG-104 appears to be insusceptible by pH changes in the range from 6.0 to 7.6. In order to simplify cell loading, KMG-104-AM, a membrane-permeable probe of AM ester form of KMG-104 has also been synthesized. The probe was successfully incorporated into PC12 cells through microinjection method. By using a confocal microscope, the intracellular three-dimensional Mg\(^{2+}\)
distributions were successfully imaged with the KMG-104-AM probe. KMG-104 was adopted to detect Mg\(^{2+}\) transients upon mitochondrial depolarisation, and this led to the hypothesis that mitochondria could be intracellular magnesium stores. Features of KMG-104 in terms of excitation spectrum, fluorescence enhancement and ion selectivity make this probe become one of the most promising alternatives to detect free Mg\(^{2+}\) by live cell imaging.

In order to combine the fluorescent probe KMG-104 with a protein and explore the mobilization as well as underlying mechanisms of Mg\(^{2+}\), KMG-104-AsH has been designed and developed (as shown in Fig. 6). KMG-104-AsH consists of the highly selective fluorescent Mg\(^{2+}\) probe and a tetracysteine peptide tag (TCTag), which can be genetically incorporated into any protein. After the combination, KMG-104-AsH has a highly selective affinity for Mg\(^{2+}\) (K\(_d\) for Mg\(^{2+}\) = 1.7 mM, K\(_d\) for Ca\(^{2+}\) = 100 mM), with its fluorescence intensity increases by more than 10-fold. Furthermore, fluorescent imaging of intracellular Mg\(^{2+}\) in HeLa cells reveals that this Flash-type Mg\(^{2+}\) sensing probe is membrane-permeable and bound specifically to tagged proteins, such as TCTag-actin and mKeima-TCTag targeted to the cytoplasm and the mitochondrial intermembrane space. This probe is a promising tool for elucidating the dynamics and mechanisms of intracellular localization of Mg\(^{2+}\).

Developing an efficient two-photon probe for the detection of Mg\(^{2+}\) ions in living cells and live tissues is a very useful strategy. Bong Rae Cho and his research group designed another two-photon probe CMG1. In contrast to the APTRA binding site in AMG1, CMG1 contains a strong donor-acceptor pair in the cyclic planar framework to facilitate the intramolecular charge transfer (ICT) for high two-photon cross-section and an integral \(\beta\)-keto acid as the Mg\(^{2+}\)-binding site to increase water solubility, while maintaining a small molecular weight for optimal cell permeability. CMG1 can be excited at 880 nm and its spectral properties was dramatically influenced by the lipophilicity of the solvent, deciphered as a better metal ion binding in a more hydrophobic environment. Indeed, the binding affinity of CMG1 for Mg\(^{2+}\) increases gradually with sodium dodecyl sulfate (SDS) concentration up to the critical micellar concentration, and then remains constant at 1.3 mM. Interestingly, the titration curves displayed the complexation between CMG1 and Mg\(^{2+}\) in a 10 mM SDS solution and were found to be similar to that in ionophore-treated cells. Thus SDS micelles seem to represent a suitable model for the intracellular environment. While the K\(_d\) of CMG1 for Mg\(^{2+}\) is comparable to that of probes whose binding site is APTARA such as Magfura-2 and Mag-fluo-4, its K\(_d\) for Ca\(^{2+}\) is only 3.6 mM, showing obvious selectivity for Mg\(^{2+}\) over Ca\(^{2+}\). However, the selectivity is still not as good as that of KMG-104. Fortunately, thanks to the order of magnitude of typical free Mg\(^{2+}\) and Ca\(^{2+}\) intracellular concentrations, CMG1 can effectively measure free intracellular Mg\(^{2+}\) with no interference from Ca\(^{2+}\). Apart from this, no response toward either alkali metal ions (Na\(^+\), K\(^+\)) or transition-metal ions (Zn\(^{2+}\), Fe\(^{2+}\), Mn\(^{2+}\), Cu\(^{2+}\), Co\(^{2+}\)) has been discovered. In addition, CMG1 is capable of selectively detecting intracellular Mg\(^{2+}\) in living Hep3B cells and imaging endogenous stores of free Mg\(^{2+}\) at 100–300 \(\mu\)M depth in live tissues using two-photon microscopy.

Coomarin and its derivatives have been extensively used as organic dyes to design fluorescent probes. CS was a good example, Gupta et al. using coomarin derivatives, 4-methyl-7-hydroxy-8-formyl coomarin, a simple molecule served as a selective probe for Mg\(^{2+}\). As presented in Fig. 6) CS shows a remarkable enhancement and a small red shift of 12 nm from 473 to 485 nm in fluorescence emission upon complexation with Mg\(^{2+}\). The phenolic OH and the oxygen atom of carbonyl group have been proved to be the binding sites of CS to Mg\(^{2+}\) to form a rigid \(\beta\)-dicarbonyl bidentate binding system. The limit of detection of Mg\(^{2+}\) is found to be much lower as 0.43 \(\mu\)M and CS may be a practical probe for detection of Mg\(^{2+}\) in analytical as well as environmental and biological samples.

### 3.3 Fluorescent probes for Mg\(^{2+}\) based on crown ether binding site

Crown ether has been a popular receptor for alkaline or alkaline earth metal ions owing to various sizes of cavities available as...
well as the effective chelation ability of the oxygen lone pairs for these “hard” metal ions. The common types of crown ether include 15-crown-5 ether, 18-crown-6 ether and the diaza-18-crown-6 ether, in which two nitrogen atoms are inserted into the structure of 18-crown-6 ether. All of them possess the outstanding properties of metal ion coordination.

In 2007, Hama et al. published a report on the probe B15C5 (as shown in Fig. 7), which possesses benzo-15-crown-5 as recognition moiety and 1-naphthaleneacetamide as chromophore moiety. Its fluorescence emission showed a significant enhancement after the addition of Mg\(^{2+}\), and a detectable, yet not as prominent, increase in the existence of other alkaline earth metal ions (Ca\(^{2+}\), Sr\(^{2+}\), Ba\(^{2+}\)). Nevertheless, at present this compound does not show a promising future, for the observation of its excitation and emission spectra in the UV range and spectroscopic characterisation in organic solvents, which are incompatible with biological samples.

8-Hydroxyquinoline (8-HQ) is deemed as the second Mg\(^{2+}\) chelation agent in significance after ethylene diaminetetra-acetic acid (EDTA). The most noteworthy characteristic of 8-HQ is its extremely low quantum yield in aqueous or organic solutions and the fluorescence enhancement arising from cation binding, which results in an intense yellow-green fluorescence. Furthermore, the fluorescence of the complex species varies with the environment and becomes brighter as lipophilicity increases.

A significant and promising work had been accomplished by an Italian research group, who designed and developed a new family of fluorescent molecules for Mg\(^{2+}\) obtained by conjugation of a diaza-18-crown-6 ether with 8-hydroxyquinolines bearing various substituents (DCHQ1-6, Fig. 7).

Among the panel of the novel synthesized DCHQ probes, various of groups are inserted into the 5-position of the 8-HQ side arms to obtain diverse target probes. DCHQ1 and DCHQ2 were obtained by substitution of hydrogen atom and chlorine atom, respectively, while DCHQ3 bears two terminal acetoxymethyl ester groups instead with the goal of enhancing cellular uptake and/or trapping efficiently. Furthermore, by bearing long alkyl chains on the 8-hydroxyquinoline side arms, DCHQ4 obtains higher lipophilicity and, consequently, better affinity for membranes. Finally, one aromatic group each is inserted on the 8-HQ side arms to give DCHQ5 and DCHQ6, conducing to the overall charge delocalization, in an attempt to bathochromic-shift the absorption and emission spectra with respect to the parent probe DCHQ1.

The photophysical properties of these probes have been extensively studied. Several probes exhibit a remarkable increase in fluorescence intensity in the existence of Mg\(^{2+}\) while other alkaline-earth ions induce no noticeable fluorescence increase. Probes in this family display an intense band at around 240–250 nm and a smaller and broader band at around 310–330 nm in free-state. The former can be ascribed to a \(\pi-\pi^*\) transition, while the latter shows a main charge transfer feature with charge density shifting from the hydroxyl oxygen atom to the quinolone. All the compounds present a weak emission band at around 465–510 nm with low values of fluorescence quantum yields in free form, due to the presence of the nitrogen atom in the macrocycle and the 8-HQ derivative, which can be elucidated by the photoinduced electron transfer (PET) mechanisms. The increasement of Mg\(^{2+}\) can be observed in the absorption spectra of all the DCHQ derivatives, featuring the decrease of the bands centered at 240–250 and 310–330 nm as well as the appearance of new bands around 260–265 and 370–390 nm. These new bands are characterized by 8-HQ derivatives when the complexation process is accompanied by the deprotonation of the hydroxyl group in the 8 position. Apart from this, Mg\(^{2+}\) binding causes a tremendous enhancement of the emission spectra for all the studied species, up to 20 times higher than the fluorescence intensity of the free ligand, which can be explained by the fact that PET is inhibited by complexation.

In order to investigate their affinity and photochemical behaviors in the presence of Mg\(^{2+}\) and other possible competitive ions, a systematic study of the probes have been performed. Interestingly, all the DCHQ derivatives presented identical trends with regard to complexation ability: no changes were observed either in absorption or emission spectra after the addition of Na\(^{+}\), K\(^{+}\), Ca\(^{2+}\), Fe\(^{2+}\) and Fe\(^{3+}\), indicating that these species do not form any complex with the DCHQ ligands. On the contrary, Hg\(^{2+}\), Cu\(^{2+}\) and Ni\(^{2+}\) were efficiently bound to the molecules but constituted nonluminescent complexes. Among all the tested metal ions, only Mg\(^{2+}\), Zn\(^{2+}\) and Cd\(^{2+}\) induced a prominent enhancement of the fluorescence in the system (See Table 3). However, Zn\(^{2+}\) and Cd\(^{2+}\) ions are not anticipated to give rise to substantial interference in total intracellular Mg\(^{2+}\) determination for the following reasons: (1) the intracellular concentration of Zn\(^{2+}\) is approximately 100 times lower than that of Mg\(^{2+}\), not to mention that most of Zn\(^{2+}\) is bound to zinc fingers with relatively high association constants; (2) Cd\(^{2+}\) is not present in non-poisoned cells.

Loading cell, and the distribution of the probes between cells and medium as well as the detection of Mg\(^{2+}\) by live cell imaging were further investigated. In analogy to KMG-104, the experiments were conducted by Kubota et al. While KMG-104 showed cytosolic free Mg\(^{2+}\) transients upon mitochondrial depolarization, an astonishingly similar enhancement in fluorescence in DCHQ1-loaded cells was found, despite differences of the binding affinities between them. In addition, Chiara Marraccini et al. performed experiments with the DCHQ compounds by two-photon imaging in the study of intracellular
Table 3 Basic photophysical properties of the fluorescent Mg$^{2+}$ probes based on crown ether binding site

| Probes | Chromophore                  | $\lambda_{\text{max,abs}}$ (nm) | $\lambda_{\text{max,em}}$ (nm) | $K_d$ (Mg$^{2+}$) | $K_d$ (Zn$^{2+}$) | $K_d$ (Cd$^{2+}$) | Ref. |
|--------|-----------------------------|---------------------------------|---------------------------------|------------------|------------------|------------------|------|
| DCHQ1  | Hydroxyl-quinoline          | 244/244                         | 505/505                         | 5.02 ± 0.08$^a$  | 5.85 ± 0.06$^a$  | 9.39 ± 0.05$^b$  | 66   |
| DCHQ2  |                             | 245/245                         | 510/510                         | ND$^c$           | ND$^c$           | ND$^c$           | 66   |
| DCHQ3  |                             | 246/246                         | 510/513                         | 10.1 ± 0.1$^b$   | 6.12 ± 0.07$^a$  | 9.53 ± 0.05$^b$  | 68   |
| DCHQ4  |                             | 247/247                         | 463/513                         | 11.2 ± 0.4$^b$   | 6.50 ± 0.2$^c$   | 9.30 ± 0.06$^b$  | 68   |
| DCHQ5  |                             | 249/249                         | 512/517                         | 5.08 ± 0.06$^a$  | 6.60 ± 0.2$^c$   | 9.4 ± 0.2$^b$   | 69   |
| DCHQ6  |                             | 245/245                         | 504/514                         | 4.8 ± 0.4$^b$    | 5.80 ± 0.09$^a$  | 8.91 ± 0.06$^b$  | 68   |
| B1SC5  | Naphthaleneacetamide        | 258/250                         | 338/338                         | ND$^c$           | ND$^c$           | ND$^c$           | 61   |

$^a$ Value obtained for 1 : 1 metal-to-ligand stoichiometry. $^b$ Value obtained for 1 : 2 metal-to-ligand stoichiometry. $^c$ ND, the data of $K_d$ is not determined by reference; $K_d$ values of DCHQ1 and DCHQ2 for Mg$^{2+}$ are 44 μM and 73 μM, respectively. $^d$ If not otherwise stated, $K_d$ values were measured in aqueous buffers; for details please see the cited references.

total Mg$^{2+}$ distribution.$^{66}$ According to the results, the aromatic probe DCHQ5 presented unique characteristics and appears to be a valuable and efficient candidate tool for the evaluation of total intracellular Mg$^{2+}$ and understanding the homeostasis of this important element better. As an example of the sensitivity of DCHQ5, the participation of Mg$^{2+}$ in multidrug resistance in human colon adenocarcinoma LoVo cells sensitive (LoVo-S) and resistant (LoVo-R) to doxorubicin was studied. The results revealed a higher Mg$^{2+}$ concentration in LoVo-R cells. The intriguing results indicate that the photochemical characteristics of DCHQ probes might be applied to detect Mg$^{2+}$ movements across intracellular compartments, which is worth further investigation. What’s amazing, in 2017 Lvova et al. developed novel all-solid-state optodes via inclusion inside PVC-based polymeric films of different DCHQ based receptors to perform fast monitoring of Mg$^{2+}$.$^{72}$

### 3.4 Fluorescent probes for Mg$^{2+}$ based on a Schiff base ligand

A Schiff base is a compound with the general structure R$_2$C = NR’. It derives from condensation of an aliphatic or aromatic amine and a carbonyl compound. In the realm of coordination chemistry, the Schiff base plays an important role and is commonly used as ligands to form coordination complexes with metal ions.$^{73}$ Recently, moderating the corresponding groups of C=N structure to construct fluorescent probes for Mg$^{2+}$ were based on the combination action of fluorescence signaling mechanisms. For example, probe SBL-1 (as shown in Fig. 8, take the first letter of each word in phrase: Schiff base ligand, named SBL) selectively recognizes Mg$^{2+}$ due to rapid isomerization around the C=N bond,$^{74}$ meanwhile, the introduction of a hydroxyl to the adjacent nitrogen can induce excited state intramolecular proton transfer (ESIPT) to generate a keto tautomer with proficient Mg$^{2+}$ binding capability, such as probe SBL-2, -3, -4, and SBL-5. They are also identified as the photo-induced electron transfer (PET) action from the lone pair of the sp$^2$ nitrogen to the fluorophore when the fluorescence could be quenched in coordination without Mg$^{2+}$, however, when probes bind to Mg$^{2+}$, these processes will be blocked.

SBL-1, a novel probe, which was designed and developed by Ray and co-workers,$^{74}$ connected two different coumarin derivatives through an imine linkage to form a metal binding site in the middle of fluorescence signaling system. The molecule possesses satisfying photophysical properties, including a large Stokes shift, visible excitation and emission wavelengths as well as little or no interference from other metal ions. Disappointingly, this indicator is completely useless for...
biological applications, because its emission dramatically decreases upon the addition of water, and entirely vanishes in 10% v/v water, which makes this indicator completely futile for biological applications.

**SBL-2**, which was designed and developed by Narinder Singh and colleagues, has been constructed for the ratiometric recognition of Mg$^{2+}$ in semi-aqueous solution at pH 7.0. This probe with a Schiff base ligand in structure undergoes ESIPT process which can provide dual channel emission to realize ratiometric determination of Mg$^{2+}$, along with the keto tautomer as an efficient receptor for Mg$^{2+}$. The fluorophore exists preferentially in the enol form and is stabilized in the ground state by an intramolecular six-membered ring H-bonding motif. Upon excitation at 275 nm, the excited enol form ($E^*$) is transformed into the excited keto tautomer ($K^*$) arisen from ESIPT. This process results in an emission band (band at 355 nm) with a large Stokes shift. The keto form ($K$) returns to the enol form via a reverse proton transfer after decaying to the ground state (the proposed responding model with Mg$^{2+}$ is showed in Fig. 8). Perturbations to the intramolecular arrangement by means of intermolecular H-bonding interactions or variations in solvent polarity have been extensively studied. **SBL-2** displays good selectivity of Mg$^{2+}$ over other metal ions and does not suffer from interference by Ca$^{2+}$ in particular and can measure Mg$^{2+}$ ion concentration ranging from 2.0 and 30.0 $\mu$M.

Driven by the situation that few fluorescent probes for Mg$^{2+}$ were capable of applying in both animal cells and plant tissues, Zhou et al. developed **SBL-3**.**SBL-3** shows high selectivity and sensitivity for the main group Mg$^{2+}$ through fluorescence “turn-on” response in ethanol solution with no interference from Ca$^{2+}$ in particular. Detection limit of probe **SBL-3** is 1.47 x 10^{-6} M and the rapid response could reach about 15–20 s. What’s more, probe **SBL-3** presents great photostability, low toxicity, and cellular permeability and can successfully sense and detect Mg$^{2+}$ in HeLa cells. Furthermore, it is successfully applied as a Mg$^{2+}$ developer in plant tissues, indicating that it not only can successfully track the transport of Mg$^{2+}$ but also is able to make a corresponding fluorescence response to different Mg$^{2+}$ concentrations.

**SBL-4** is an aggregation-induced emission (AIE)-active fluorescence probe. It can be used as a sensitive probe to detect Mg$^{2+}$ ion in acetonitrile and the other common ions shown no noticeable interference. Particularly, upon complexation with Mg$^{2+}$, the colour of the **SBL-4** solution changes from colourless to yellow under hand-held ultraviolet lamp, which can be observed by naked-eye. However, the biological application of probe **SBL-4** is limited by its inferior characteristics.

Interestingly, **SBL-5**, as described by Liu and colleagues, exhibits a solvent-dependent “turn-on” fluorescence recognition for Mg$^{2+}$ anchored in combination of C=N isomerization as well as inhibition of ESIPT (the probable binding mode and action between **SBL-5** and Mg$^{2+}$ are showed in Fig. 8). The unique property of **SBL-5** is that it shows turn-on fluorescence response for Mg$^{2+}$ only in alcohol solvent with sensitivity and negligible fluorescent response in other nonpolar or polar solvents. Upon binding with Mg$^{2+}$, a 148-fold fluorescence enhancement was observed at 458 nm when 20 $\mu$M Mg$^{2+}$ was added, along with the colour changing from colourless to blue. However, due to these photophysical changes only happen in alcohol solvent, which is not close to the physiological environment, thus the probe might not be applied in colorometric and ratiometric sensing of Mg$^{2+}$ in living cells.

**SBL-6**, which was reported by Zhanget al. in 2017, is a novel “off-on” fluorescent probe based on 1, 8-naphthalimide derivative for the detection of Mg$^{2+}$ in ethanol solution. The o-hydroxyl Schiff base, a very important unit in the structure of **SBL-6**, which not only plays the role of connecting 1,8-naphthalimide chromophore and 2-hydrazinylbenzo[\(n\)] thiazole group, but also acts as recognizer to realize “off-on” fluorescence recognition. The process and proposed sensing process of **SBL-6** are showed in Fig. 8). Its complexation towards Mg$^{2+}$ is a 1:2 binding mode in ethanol. **SBL-6** displays responses to Mg$^{2+}$ with a fluorescence enhancement at 523 nm, accompanying with a distinct fluorescence change from nearly colorless to bright yellow-green. In addition, biological application experiments revealed that the probe is nearly non-toxic and the fluorescence scanning microscopic experiments demonstrated that the probe can be qualified for monitoring at the intracellular Mg$^{2+}$ level successfully.

Recently, Li et al. designed, synthesized and evaluated **SBL-7** as a novel Mg$^{2+}$ “turn-on” fluorescent probe, which connected fluorescein-derived Schiff-base ligand to a chromone moiety.**SBL-7** binds Mg$^{2+}$ with good selectivity and exhibits excellent sensitivity towards Mg$^{2+}$ over other important metal ions investigated. Upon binding with Mg$^{2+}$, the fluorescence emission intensity at 504 nm increases by about 50-fold. The remarkable enhancement in fluorescence emission is due to the fact that the ring opening process of the fluorescein fluorophore in **SBL-7** upon complexation with Mg$^{2+}$. **SBL-7** responses to Mg$^{2+}$ within 3 min, and can be utilized to sense and monitor Mg$^{2+}$ for real-time detection.

**SBL-8** serves as a dual analyte probe and quantifies Mg$^{2+}$ and Zn$^{2+}$ ions, and that the binding constant of Mg$^{2+}$ is lower. It was designed and synthesized by Maity and his workmates using coumarin derivative 3-amino-7-(diethylamino)-2H-chromen-2-one as fluorophore and it was conjugated to 2,6-diformyl-4-methylphenol via a C=N bond as well. The proposed binding mode of **SBL-8** with Mg$^{2+}$ and Zn$^{2+}$ is based on C=N bond isomerisation mechanism and the binding sites maybe include the carbonyl oxygen of coumarin ring, imine nitrogen as well as phenolic oxygen (as shown in Fig. 8). However, owing to lack of selectivity towards Mg$^{2+}$, thus the application of **SBL-8** will be limited to some extent.

### 3.5 Miscellanea

As discussed in the preceding part of this review, four families of Mg$^{2+}$-responsive fluorescent probes have been reported and put into biological applications, *i.e.* the receptor groups based upon an APTRA, a charged β-diketone, a crown ether structure and a Schiff base ligand. However, there are still other types of Mg$^{2+}$-responsive fluorescent probes, which do not belong to the above classification, have also been developed. There-in-after, we will give a brief introduction of these probes.
Calixarenes are cyclic oligomeric host molecules formed through a condensation reaction between phenol and formaldehyde. They exist in different ring sizes, each bearing a hydrophobic interior, among which the four-monomer ring calix[4]arene is the most studied one. The usage of calix[4]arene derivative as a probe for the detection of Mg$^{2+}$ ions was described by Song and colleagues. The probe exhibits a large Mg$^{2+}$-induced red-shifted emission. The selective sensing of Mg$^{2+}$ over other potentially interfering ions could be realised by analysing the ratio between fluorescence emission at 493 and 409 nm, and the value was significantly higher for Mg$^{2+}$. Such discriminating behaviour suggested that this molecule could be utilized in ratiometric sensing of magnesium in semi-aqueous environments.

Analogously, porphyrins and their derivatives, which contain both a fluorophore and a coordination platform, are promising candidates for the development of cation-responsive probes. These molecules have ideal properties such as a cell-penetrating far-red or near-infrared fluorescent emission, as well as large Stokes shifts which minimize the influence of the background fluorescence. In accordance with these superior features, a porphyrin-related macrocycle probe for Mg$^{2+}$ recognition was proposed by Ishida and co-workers, which also shows fluorescence enhancement and bathochromic shift upon Mg$^{2+}$ complexation, while displaying inappreciable interference from other physiologically relevant cations (Na$^+$, K$^+$, Ca$^{2+}$). Thus it is recommended as a cell-permeable fluorescent sensor for Mg$^{2+}$ in semi-aqueous solution enabling ratiometric measurements.

To the best of our knowledge, almost all fluorescent probes devised for the detection of Mg$^{2+}$ meet a common problem of nonnegligible interference from either alkali metal ions or transition-metal ions. Fortunately, the use of a single-molecular multi-analyte, namely, a single probe that allows the sensing of multiple analytes with different spectral responses is promising for overcoming the poor selectivity of probe for Mg$^{2+}$. Inspired by this idea and towards this objective, two novel fluorescent probes, KCM-1 and DFC-8-AQ (see in Fig. 9), were successfully designed, synthesized and characterized. KCM-1, one of single-molecular multi-analyte fluorescent probe for the simultaneous sensing of both Ca$^{2+}$ and Mg$^{2+}$, was developed by Koji Suzuki and co-workers. The molecule of novel probe KCM-1 includes a coumarin moiety as a stable fluorophore which can be excited with visible light, BAPTA (O,O′-bis(2-aminophenyl)ethyl- eneglycol-N,N,N′,N′-tetraacetic acid) as the Ca$^{2+}$-selective binding site and a charged β-diketone as the Mg$^{2+}$-selective binding site. Consequently, fluorescence emission of KCM-1 shows a spectral hypochromatic shift in the presence of Ca$^{2+}$ and a bathochromic shift upon complexation to Mg$^{2+}$.

Furthermore, Ca$^{2+}$ and Mg$^{2+}$ concentrations can be measured by a calibration performed on data acquired by a combination of three different excitation/emission wavelengths. However, the binding characteristics of KCM-1 are quite disillusionary, as $K_d$ values for Ca$^{2+}$ and Mg$^{2+}$ are reported to be 14 μM and 26 mM, respectively, which are obviously too high to effectively measure common intracellular concentrations of these ions. Nevertheless, KCM-1 has been applied to detect distinct Ca$^{2+}$ and Mg$^{2+}$ transients upon mitochondrial depolarisation.

The second probe in Fig. 9, DFC-8-AQ possesses a diformyl-p-cresol (DFC)-8-aminoquinoline moiety as the chromophore, with the Schiff base linked to the benzene ring and the quinoline ring coupled with the phenolic and aldehyde groups to constitute the receptor moiety of the probe, which is very close to the charged β-diketone binding site. DFC-8-AQ displays colorimetric and fluorescence properties upon selective and sensitive binding towards Mg$^{2+}$ and Zn$^{2+}$. DFC-8-AQ turns on fluorescence enhancement as high as 40 folds and 53 folds upon binding of Mg$^{2+}$ and Zn$^{2+}$, respectively. Additionally, this probe could be selective towards Mg$^{2+}$ over Zn$^{2+}$ in presence of the specific cell-permeable heavy metal chelator TPEN ($N,N,N',N'$-tetrakis(2-pyridylmethyl)ethylenediamine), under both intra- and extracellular conditions. Furthermore the dissociation constant of the Mg$^{2+}$-(DFC-8-AQ) complex ($K_d = 6.60 \mu M$) is far lower than a large proportion of reported Mg$^{2+}$ probes which fall in the mM range. However, despite the encouraging results, the project hasn’t been pursued further yet.

A controlled supramolecular approach using allosteric effect in combination with conformation restriction in a fluorescence-based ring-off/open system was less investigated in the field of fluorescent probes. Yu et al. adopted this strategy to design and synthesize a novel fluorescent and colorimetric probe FL-1 for Mg$^{2+}$ or Ca$^{2+}$ over other competing ions. FL-1 displays turn on fluorescent and colour changes upon addition of Mg$^{2+}$ and Ca$^{2+}$. Through the experiments of fluorescence emission, UV-visible titrations and $^1$H-NMR titrations revealed that the cation-driven can construct conformation of FL-1. And that Mg$^{2+}$ and Ca$^{2+}$ could be selectively sense by the allosteric effect. Even though this strategy is novel-innovative, one of its fatal drawbacks is that it cannot exclude the interference from Ca$^{2+}$.

4. Conclusions and outlook

In the present review, we attempt to cover the advances in fluorescent probes for detection of Mg$^{2+}$ in past several decades, with the goal of providing some valuable clues to find the safe exit of such a colorful maze and direct future developments. Firstly, we placed emphasis on the discussion of general design principles of fluorescent probes for ion detection, which would contribute to the understanding of the operational mechanisms of fluorescent probes for both the users and the inventors. Simultaneously, we classified numerous fluorescent probes for sensing Mg$^{2+}$ based on the structure of their recognition units. The appropriate combination of the ion acceptor unit and the fluorophore reporter unit can offer a potent tool for the visualisation of Mg$^{2+}$ in vitro and in vivo to clarify the roles of Mg$^{2+}$ in living organisms. Since the pioneering work of

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**Fig. 9** Chemical structures of other types of Mg$^{2+}$ probes.
fluorescent probes for sensing intracellular Ca^{2+}, a multitude of fluorescent probes for the detection of Mg^{2+} have sprung up that operate according to various mechanisms and are constructed based on different chromophores in conjunction with multifarious selective-Mg^{2+} acceptors. Despite the rapid progress of Mg^{2+}-selective fluorescent probes, a diverse array of problems and challenges are shrouded in this research field at present. The current major knotty problems and challenges include:

4.1 Concerning the acceptor

A large proportion of Mg^{2+} probes suffer from the lack of selectivity over Ca^{2+} or other metal ions and the $K_d$ value is relatively high. In addition, Mg^{2+} fluorescent probes which involve AM (acetoxyethyl) esters also have multiple problems, including compartmentalisation within intracellular organelles, incomplete hydrolysis resulting in the formation of fluorescent ion-insensitive species as well as leakage from cytosol to extracellular medium. Therefore, active search and creative development of acceptors with extraordinary affinity for Mg^{2+} without interference from competitive ions is in urgent demand. Meanwhile, further improvement of the acceptors would also contribute to their utility in cells.

4.2 In terms of the fluorophore

There is no doubt that the sensitivity and signal-to-noise ratio of a probe mainly depend on the brightness and stability of the probe’s fluorophore(s) as well as the features of the instrumentation. In general, an ideal fluorophore should meet certain requirements, such as, (a) absorption and emission wavelengths in the visible region, or more preferably in near-infrared or infrared range; (b) good photo-stability and very low light scattering; (c) low or no toxicity to living cells; (d) excellent equilibrium between water soluble and fat-soluble property; (e) easy ways to produce and modify probes without high costs. However, the vast majority of fluorophores in Mg^{2+} fluorescent probes don’t possess most of these elements, which drastically reduces the potential biological applications. Therefore, it is of highly priority to develop brighter fluorophores with better photo-physical nature.

4.3 Designing ratiometric probes or functional probes based on multi-sensing mechanisms

It is well-known that ratiometric fluorescent recognition prevails over conventional monitoring at a single wavelength since the method is free from the errors related to receptor concentration, photobleaching and environmental effects. Although it is an important step forward to develop control probes, ratiometric probes act as their own internal controls to correct potential dye-dependent localization and sample thickness effects. Such probes are allowed to evaluate the concentrations more quantitatively for Mg^{2+} buffered in the cell. The FRET probe for the detection of Mg^{2+} is currently blank, which is likely to be a future direction for effort. Besides, the combination of multi-responding mechanisms to construct functional probes will possibly be an ideal choice, too. Looking forward, it is obvious that the performance of probes designed by simultaneous multi-sensing mechanisms to elucidate interactions between Mg^{2+} pools and the living systems would be an attractive target in future.

4.4 Continued innovations in analytical methods and imaging techniques

Generally speaking, the possibility to monitor Mg^{2+} in a living organism would contribute significantly to magnesium research. Accordingly, the ultimate challenge is not only to design advanced probes, but also to develop satisfactory tools for in vivo investigations. Besides, an additional Frontier is to enlarge the toolkit for imaging Mg^{2+} in living organisms, which would undoubtedly enhance our comprehension of Mg^{2+} homeostasis at the organismal level. In the last decade, multitudes of different imaging techniques and analytical methods have been employed, ranging from wide-field fluorescence microscopy to confocal microscopy, from single photon fluorescence microscopy to two-photon fluorescence microscopy, and so on. It hopes that the spur that has inspired imaging techniques will also be passed on to the realm of magnesium research.

To sum up, we do believe that with on-limits attitude and extensive collaboration of researchers in chemistry, physics, medicine and biology, the design and synthesis of novel probes as well as the development of advanced imaging techniques and instrumentation will facilitate our understanding of the comprehensive role of Mg^{2+} in living cells.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

1. H. Rubin, BioEssays, 2005, 27, 311–320.
2. A. M. Alexandra, Sci. Signaling, 2016, 9, ec269.
3. M. H. Pontes, J. Yeom and E. A. Groisman, Mol. Cell, 2016, 64, 480–492.
4. B. O’Rourke, P. H. Backx and E. Marban, Science, 1992, 257, 245–248.
5. A. Roman, Arch. Biochem. Biophys., 2007, 458, 90–102.
6. F. I. Wolf and A. Cittadini, Mol. Aspects Med., 2003, 24, 3–9.
7. W. Jahnen-Dechent and M. Ketteler, Clin. Kidney J., 2012, 5, i3–i14.
8. J. F. Committee, British National Formulary, The Pharmaceutical Press, London, 2015, ISBN: 978085711562, p. 463.
9. A. Walsh, Spectrochim. Acta, 1955, 7, 108–117.
10 C. Feillet-Coudray, A. Trzebiakiewicz, C. Coudray, M. Rambeau, A. Chanson, Y. Rayssiguier, A. Opolski, F. I. Wolf and A. Mazur, *Eur. J. Nutr.*, 2006, 45, 171–177.

11 H. R. Halvorson, A. M. Vande-Linde, J. A. Helpern and K. M. Welch, *NMR Biomed.*, 1992, 5, 53–58.

12 S. Iotti and E. Malucelli, *Magnetism Res.*, 2008, 21, 157–162.

13 H. Ishikawa, K. Suzuki, M. Nakanishi and A. Inokai, *Bioimaging for Molecular Dynamics and Cellular Functions*, Kyoritsuushuppan Co., Ltd., Tokyo, 1998.

14 J. A. Thomas, *Chem. Soc. Rev.*, 2015, 44, 4494–4500.

15 S. Lee, K. K. Yuen, K. A. Jolliffe and J. Yoon, *Chem. Soc. Rev.*, 2015, 44, 1749–1762.

16 G. G. Guilbault, *Practical Fluorescence*, 2nd edn, Marcel Dekker Inc, New York, 1990, ISBN 0824783506.

17 A. P. de Silva, H. Q. Gunaratne, T. Gunnalugsson, A. J. Huxley, C. P. McCoy, J. T. Rademacher and T. E. Rice, *Chem. Rev.*, 1997, 97, 1515–1566.

18 N. Boens, V. Leen and W. Dehaen, *Chem. Soc. Rev.*, 2012, 41, 1130–1172.

19 B. Valeur and I. Leray, *Coord. Chem. Rev.*, 2000, 205, 3–40.

20 Z. R. Grabowski and J. Dobkowski, *Pure Appl. Chem.*, 1983, 55, 245–252.

21 Y. M. Yang, Q. Zhao, W. Feng and F. Y. Li, *Chem. Rev.*, 2013, 113, 192–270.

22 J. S. Kim and D. T. Quang, *Chem. Rev.*, 2007, 107, 3780–3799.

23 J. Fan, M. Hu, P. Zhan and X. Peng, *Chem. Soc. Rev.*, 2013, 42, 29–43.

24 Y. Feng, J. Cheng, L. Zhou, X. Zhou and H. Xiang, *Analyst*, 2012, 137, 4885–4901.

25 A. Coskun and E. U. Akkaya, *J. Am. Chem. Soc.*, 2006, 128, 14474–14475.

26 Y. Kurishita, T. Kohira, A. Ojida and I. Hamachi, *J. Am. Chem. Soc.*, 2010, 132, 13290–13299.

27 R. Y. Tsien, *Biochemistry*, 1980, 19, 2396–2404.

28 G. Grynkiewicz, M. Poenie and R. Y. Tsien, *J. Biol. Chem.*, 1985, 260, 3440–3450.

29 L. A. Levy, E. Murphy, B. Raju and R. E. London, *Biochemistry*, 1988, 27, 4041–4048.

30 B. Raju, E. Murphy, L. A. Levy, R. D. Hall and R. E. London, *Am. J. Physiol.*, 1989, 256, C540–C548.

31 R. E. London, *Annu. Rev. Physiol.*, 1991, 53, 241–258.

32 E. Murphy, C. C. Freudenberg, L. A. Levy, R. E. London and M. Lieberman, *Proc. Natl. Acad. Sci. U. S. A.*, 1989, 86, 2981–2984.

33 M. T. Touyz and E. L. Schiffrin, *J. Biol. Chem.*, 1996, 271, 24353–24358.

34 M. Tashiro, H. Inoue and M. Konishi, *Biophys. J.*, 2014, 107, 2049–2058.

35 E. Gouafon, F. Leercf and M. German-Fattal, *J. Pharm. Pharmacol. Sci.*, 2012, 15, 389–398.

36 L. Csernoch, J. C. Bernengo, P. Szentesi and V. Jacquemond, *Biophys. J.*, 1998, 75, 957–967.

37 C. Cheng and I. J., *Neurosciences*, 2000, 95, 973–979.

38 L. S. Meena, S. R. Dhakate and P. D. Sahare, *Biotechnol. Appl. Biochem.*, 2012, 59, 429–436.
