A low-molecular-weight ditopic MRI probe for ratiometric sensing of zwitterionic amino acid neurotransmitters†

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We report a novel ditopic Gd(III)-based probe selective to zwitterionic amino acid neurotransmitters (ZNTs) crafted for ratiometric MRI imaging. The probe displayed increased binding affinity to ZNTs and non-synchronized concentration-dependent changes of the $r_1$- and $r_2$-relaxity. Through the application of a $T_2/T_1$ weighted MRI strategy, we demonstrated signal enhancement for cooperatively bound glutamate and $\gamma$-aminobutyric acid ZNTs over competitive hydrogen carbonate, which remained MR silent.

A wide variety of neuronal diseases are characterized by the overexpression of zwitterionic amino acid neurotransmitters (ZNTs) into the synaptic cleft, which in abnormal amounts trigger a plethora of processes resulting in the partial impairment or death of nerve cells. Excess secretion of the major excitatory glutamate (Glu) or inhibitory $\gamma$-aminobutyric acid (GABA) ZNTs causes an imbalance in the excitation level of the brain, leading to various diseases, such as epileptic seizures that result in the degeneration of neuronal tissue. Thus, decoding the spatiotemporal patterns of brain chemodynamics could imply connection with related diseases. For this reason, ZNTs are recognized as irreplaceable biomarkers for monitoring neuronal activity. Currently, non-invasive imaging techniques based on magnetic resonance (MR), i.e. diamagnetic chemical exchange saturation transfer (diaCEST) or $^{13}$C MR spectroscopy, are preferably used due to good spatiotemporal resolution and depth of penetration. However, they do not distinguish between intra- and extracellular concentration of ZNTs, and have insufficient chemical resolution and low sensitivity. A promising way to address these issues involves the introduction of a paramagnetic sensor capable of altering image contrast upon interaction with the target metabolite. Therefore, the development of MR imaging (MRI) probes responsive to extracellular ZNTs is of emerging importance.

Molecular recognition between magnetic host sensors and guest molecules occurs via the formation of a ternary adduct, and is evaluated via its binding affinity. In the case of a ‘turn-off’ response, the interaction of biomarkers with the coordination cage of an MRI probe restricts inner-sphere water accessibility to the paramagnetic center resulting in a signal decrease. So far, the direct sensing of NTs has been approached with two distinct designs, via genetically engineered metalloproteins and by small-molecule ditopic paramagnetic probes. Although large molecule CAs tailored for monoamine neurotransmitters were shown to be capable of detecting target metabolites in the µM range, further improvement of small-sized probes is imperative due to issues associated with translation into biological systems. In terms of host-guest interaction, the low-molecular-weight sensors are bismacro cyclic Gd(III)-based complexes with a 1,4,7,10-tetraazacyclododecane-1,7-dicarboxylic acid (DO2A) chelator, whose geometry allows for easy access to small anions. In particular, the selectivity of the current sensors is greatly hindered by hydrogen carbonate (HCO$_3^-$), which binds in a similar fashion as the ZNTs. Owing to high extracellular concentration (∼25 mM), small molecular size and consequently low steric hindrances, HCO$_3^-$ predominantly binds to the paramagnetic label, thus preventing selective interaction with ZNTs. On the other hand, the advantageous feature of ZNTs is their existence in the form of an ion-pair entity at physiological pH, which opens the possibility of exploiting a ditopic platform to strengthen binding through cooperativity. In accordance with this, different binding modes may unequally affect parameters such as inner-sphere water exchange rate or molecular tumbling ($r_m$ and $r_t$, respectively), which determine both longitudinal ($r_1$) and transverse ($r_2$) relaxivities. In turn, the occurrence of non-synchronized changes of $r_1$ and $r_2$ would allow for the utilization of the ratiometric $r_2/r_1$ MRI approach.

In order to develop such a sensor, it is necessary to evoke non-synchronized stimuli-induced changes of parameters that determine $r_1$ and $r_2$. Therefore, we set out to exploit the cooperative binding potential of a ditopic low-molecular weight Gd-chelate to improve the sensing of ZNTs. The probe design consisted of a bismacro cyclic framework, providing receptor sites to favour cooperative ion-pair binding of ZNTs.
interactions involve simultaneous coordination of the carboxylate to Gd(III) and multiple hydrogen bonds established between the 18-crown-6 derived moiety (18C6) and the NH₃⁺ terminal of ZNTs (Fig. 1). The rationale for choosing 1,4,7,10-tetraazacyclododecane-1,4,7-tricarboxylic acid (DO3A) as the MR responsive unit over the coordinatively less saturated DO2A macrocycle is due to its higher selectivity towards carboxylates from ZNTs over HCO₃⁻/CO₃²⁻, owing to stronger electronic repulsions from the acetate pendant arms. To reinforce the affinity towards the ammonium cation, we incorporated a formanilide pendant that bears two amide groups, one proximate and one distant to the 18C6 ring-frame, expecting to engage as the H-bond donor and additionally stabilize the binding of the ammonium cation. The role of the flexible amide linker is to preserve the high degree of conformational mobility of the binding sites, providing an adaptable geometry for cooperative binding of the zwitterions (Fig. 2).

The proposed sensor GdL was synthesized by bridging two macrocyclic components, followed by complexation with Gd(III) (Scheme 1). The 18C6 fragment 1 was obtained by direct mono-N-alkylation of the 1,10-diaza-18-crown-6 ether with 2-bromo-N-(4-nitrophenethyl)acetamide. Next, the macrocyclic precursors 1 and 2 were covalently linked to form bismacroyclic derivative 3. As the preparation of bismacrocylics of this type is often limited by a bridging step, the yield of 62% can be considered as excellent. In the following step, the aromatic nitro group of 3 was subjected to Pd(OH)₂/C-catalysed hydrogenation to afford the amine 4. Simultaneous hydrolysis of the tert-butyl esters and conversion of the amine into formamide was conducted in formic acid to obtain the final ligand L with a yield of 64%, and a ratio of isomers of 72:28. The complexation was performed in water media at neutral pH with GdCl₃ to give the corresponding complex GdL. To avoid entrapment of a Gd(III) ion within the 18C6 component, the complexation was performed using a ligand to Gd(III) ratio of 1.0:0.8 and the excess of ligand was effectively removed by the HPLC. The absence of free Gd(III) ions was additionally confirmed by the xylenol orange test.

The bulk ¹H T₁ and T₂-relaxation time dependencies of GdL on the concentration of the ZNTs: Glu, glycine (Gly), GABA, the non-zwitterionic transmitter acetylcholine (ACh) and competitive metabolite HCO₃⁻ were assessed in relaxometric titrations (Fig. 3). The initial values calculated for r₁ and r₂ were 10.14 and 13.46 mM⁻¹ s⁻¹, respectively. The displayed downward trends revealed that GdL is ACh-insensitive while HCO₃⁻ exhibited a comparable drop in r₁ (69%) to the ZNTs that induced changes, with r₂ ranging from 61 to 69%. However, notably diverse r₂

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**Fig. 1** The chemical structure of the ditopic Gd(III)-chelate, GdL.

**Fig. 2** Binding modes of structurally different guests with GdL. (a) Monotopically bound HCO₃⁻ and (b) ditopically bound ZNT.

**Scheme 1** Synthesis of the crown ether-containing ligand L. Reagents and conditions: (i) 2-bromo-N-(4-nitrophenethyl)acetamide, NaHCO₃, MeCN, 45 °C, 52%; (ii) DMF, NaHCO₃, RT, 3 days, 62%; (iii) H₂, Pd(OH)₂/C (20 wt%), EtOH, 3.1 bar, 84%; (iv) HCOOH, 65 °C, 16 h, 64%.

**Fig. 3** ¹H relaxometric study of GdL (a and b) the experimentally measured (symbols) and fitted data (curves) for (a) r₁ and (b) r₂ relaxivities for GdL (2.5 mM) with different analytes (0–100 equiv.) in 50 mM HEPES (pH 7.4, 7.05 T, 298 K). Lines in (a) correspond to the fit described in the ESI† while the dashed lines in (b) show the mismatch of the experimental to those fitted in the same way, while assuming Ka values obtained from fitting the r₁ data. (c) Decreases in r₁ and r₂ for the host–guest molar ratio of 1:60. (d) The corresponding r₂/r₁ ratio.
changes between HCO₃⁻ and ZNTs were observed. Upon saturation with the guests (60 equiv.), the r₂ of ternary adduct GdL-HCO₃⁻ was 3.69 mM⁻¹ s⁻¹, while the values for the Gly, -Glu and -GABA adducts were 5.64, 5.78 and 5.89 mM⁻¹ s⁻¹, respectively. The gathered results are in agreement with the simulations performed for low-molecular-weight Gd(III)-based CAs at high magnetic fields (>1.5 T). Namely, T₁ is relatively constant over a broad range of values for the rₐ and rᵢ parameters. On the contrary, for the same parameters, T₂ undergoes a rather drastic change. Hence, the occurrence of non-synchronized changes of T₁ and T₂ within the same MR reporter, opens the possibility to implement a rᵢ/r₁ ratiometric MRI strategy. When presented as the rᵢ/r₁ ratio, there is an increase from the initial value of 1.32 for GdL to 1.7, 1.9 and 2.1 for Gly, Glu and GABA, respectively. In the case of HCO₃⁻, the decrease in rᵢ is followed by a proportional decrease in r₂, resulting in insignificant fluctuations for the rᵢ/r₁ values over the entire titration range.

The binding affinities (Kᵢ) were obtained by fitting the r₁ data to eqn (S2) (ESI†) (Table 1). Comparison of Kᵢ values revealed higher affinities of 116 and 106 M⁻¹ for GABA and Glu, respectively, over HCO₃⁻ and Gly, which were 86 and 64 M⁻¹, respectively. The observed trends suggested greater structural complementarity of the receptor-binding pocket to the long-chain ZNTs, Glu and GABA. Comparing HCO₃⁻ and Gly, the former binds stronger due to its potential to interact in the bidentate binding mode. The decreased affinity of Gly compared to GABA and Glu is likely due to the absence of cooperative host-guest binding as the carboxylate–amine terminals in GABA and Glu are more suited to the spatial disposition of binding sites within the host. The ambient concentrations of Glu, GABA, Gly and Gln in physiological brain states of ~6 μM, 0.23 μM, 16.7 μM and 1.738 mM, respectively, are still below the detection level of the 1H NMR spectra (see above), which led us to exclude the involvement of the formanilide moiety in binding.

The conformational mobility of the binding sites in the host molecule is affected by the interactions with guest molecules. Therefore, to gain insight into the host–guest binding modes, various NMR studies were performed on the diamagnetic yttrium(III) analogue, YL. High-resolution ¹H NMR spectra were recorded with YL in the presence of 20 equiv. of Glu or HCO₃⁻ (Fig. 4). The resonances of both ternary complexes shifted upfield, with a more pronounced effect for Glu, which are on the level of ~10 mM, still require slight improvements to allow for the detection of the relevant ZNTs, whose concentrations are closer to low mM values.

![Fig. 4 Overlapped 2D EXSY NMR spectra of YL (blue) and YL-Glu (red) in D₂O (pD 7.8, 800 MHz).](image)

Anion binding to the paramagnetic label coordinated by DO3A-based chelates is well documented, and thus, we focused on investigating interactions that involve the 18C6 binding site. 2D EXSY NMR spectra displayed conformational diversity for the 18C6 moiety within the ternary adducts (Fig. 4 and Fig. S6–S9 in ESI†). The higher density of cross-peaks in YL-Glu pointed towards a binding mode that involves multiple interactions between the 18C6 and the ammonium cation. Conversely, the more defined cross-resonances for 18C6 in YL-HCO₃⁻ were assigned to Na⁺ inclusion (bicarbonate was added as the monosodium salt). Moreover, no change was observed in the ratio of the isomers in the ¹H NMR spectra (see above), which led us to exclude the involvement of the formanilide moiety in binding.

Another parameter sensitive to molecular motions of bound protons is the T₁ relaxation time. Thus, to quantify the observed conformational flipping, we measured T₁ of protons in the NH₂CH₂CH₂O groups within the 18C6 fragment and the meta-H in the remote pendant (Table S2 and Fig. S10 in ESI†). The acquired T₁ values for the 18C6 moiety in YL-Glu and YL-HCO₃⁻ were 692 ms and 651 ms, respectively, whereas the one of binary YL was measured to be 628 ms. The elevated T₁ value in the presence of Glu indicated a higher rate of conformational exchange of the 18C6 binding site. This was rationalized as the impact of reversible interactions of multiple hydrogen bonds between NH₃⁺ of the zwitterion and free electronic pairs on the heteroatoms in 18C6. Slightly increased mobility of 18C6 in the monotopically bound YL-HCO₃⁻ is likely related to stronger interactions with entrapped Na⁺. The identical trend of mobility was observed for the 18C6 pendant, which is in alignment with the higher degree of freedom for linear compounds.

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Table 1 The ¹H r₁ relaxation decrease and affinity constants (Kᵢ) of the ternary complexes GdL-analyte in 50 mM HEPES (298 K, pH 7.4, 7.05 T)

<table>
<thead>
<tr>
<th>Analyte</th>
<th>Gly</th>
<th>Glu</th>
<th>GABA</th>
<th>ACh</th>
<th>HCO₃⁻</th>
</tr>
</thead>
<tbody>
<tr>
<td>Δr₁ (%)</td>
<td>68</td>
<td>69</td>
<td>71</td>
<td>N/A</td>
<td>75</td>
</tr>
<tr>
<td>Kᵢ (M⁻¹)</td>
<td>64</td>
<td>106</td>
<td>116</td>
<td>N/A</td>
<td>86</td>
</tr>
</tbody>
</table>

a The values are calculated for the complex to analyte ratio of 1:60.
b Binding affinities were obtained from the fit of r₁ curves to eqn (S2) (ESI).
adducts, YL-Glu and YL-HCO₃⁻, compared to the binary YL (Fig. S11 and Table S3 in ESIT). The corresponding hydrodynamic radii shrank from 1.54 nm to 1.29 nm upon moving from the binary complex to the ternary species, suggesting a different spatial arrangement of YL prior to its interaction with guests. These values excluded the formation of aggregates or intermolecular self-assembly. The rotational correlation times (τᵣ), calculated according to the Stokes–Einstein-Debye model (see ESIT), were identical for the ternary complexes (234 ps), which ruled out molecular tumbling as the explanation for the observed diversity in T₂ trends.²¹

To capitalize on this effect, we tested the selectivity of the Gdl probe by comparing the signal enhancements of T₁w and T₂w with T₁/T₂w images of MRI tube phantoms in a 7.05 T MRI scanner (Fig. 5 and Fig. S12, S13 and Tables S4, S5 in ESI†). The contrast enhancements were calculated as differences in obtained signal-to-noise ratio (SNR). The T₁w images gave negligible ΔSNRs for Gly, GABA and Glu with respect to the signal enhancement produced by HCO₃⁻, which are in the range 1–4%. On the other hand, HCO₃⁻ caused greater MRI signal alteration in the T₂w images, thus masking the effects of other ZNTs. Specifically, Gly, GABA and Glu only reached 70, 47 and 57% of the signal enhancement observed for HCO₃⁻, respectively. In contrast to the results obtained using T₁w and T₂w MRI protocols, the T₂/T₁w phantoms gave more intense signals for Gly, GABA and Glu compared to HCO₃⁻ (4, 11 and 10%, respectively). Moreover, HCO₃⁻ remained completely MR silent with the ΔSNR equal to that of Gdl. In addition, comparing the greater potential of the T₂/T₁w MRI protocol to the commonly used T₁w analogue to produce images with greater SNR,¹⁴ it should be noted that the effective discrimination of HCO₃⁻ was successfully achieved under physiologically-relevant conditions.

In summary, we have developed a novel ditopic bismacrocyclic MRI probe with an increased binding affinity towards the major inhibitory (GABA) and excitatory (Glu) neurotransmitters and employed an r₂/r₁ imaging strategy to discriminate from their physiological competitor, HCO₃⁻. We demonstrated that the bound guests induced uneven changes of r₁ and r₂ in the supramolecular adducts. Unlike the long-chain ZNTs, the monotopically bound HCO₃⁻ induced a proportional decrease in both r₁ and r₂, whose ratio remained equal to that of unbound Gdl. Hence, greater SNR changes were observed for GABA and Glu in MRI phantoms, and were less emphasized for Gly. Overall, these findings establish an efficient strategy to sense polydentate metabolites by employing low-molecular-weight polytopic MR probes. We anticipate that the introduced concept of utilizing a polytopic host with a ratiometric imaging methodology will be beneficial in the future development of MR probes for sensing polycharged guest molecules.

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Conflicts of interest
There are no conflicts to declare.

Notes and references