Selenoglycosides as Lectin Ligands: $^{77}$Se-Edited CPMG-HSQMBC NMR Spectroscopy To Monitor Biomedically Relevant Interactions

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The fundamental importance of protein–glycan recognition calls for specific and high-resolution techniques for their detailed analysis. After the introduction of $^{19}$F NMR spectroscopy to study the recognition of fluorinated glycans, a new $^{77}$Se NMR spectroscopy method is presented for complementary studies of selenoglycans with optimised resolution and sensitivity, in which direct NMR spectroscopy detection on $^{77}$Se is replaced by its indirect observation in a 2D $^1$H,$^{77}$Se HSQMBC spectrum. In contrast to OH/F substitution, O/Se exchange allows the glycosidic bond to be targeted. As an example, selenodigalactoside recognition by three human galectins and a plant toxin is readily indicated by signal attenuation and line broadening in the 2D $^1$H,$^{77}$Se HSQMBC spectrum, in which CP/MG-INEPT long-range transfer ensures maximal detection sensitivity, clean signal phases, and reliable ligand ranking. By monitoring competitive displacement of a selenated spy ligand, the selective $^{77}$Se NMR spectroscopy approach may also be used to screen non-selenated compounds. Finally, $^1$H,$^{77}$Se CPMG-INEPT transfer allows further NMR sensors of molecular interaction to be combined with the specificity and resolution of $^{77}$Se NMR spectroscopy.

Glycans are increasingly recognised for their capacity to encode biological information at high density, thus attracting interest in understanding how it is specifically read and transduced into a plethora of physiological effects.[1] For instance, the recognition of cellular glycoconjugates by various endogenous sugar receptors[2] underlies specific bridging in the cell–cell/matrix adhesion and initiation of signalling in diverse regulatory processes such as the induction of anoikis/apoptosis, or mediator release in degenerative and inflammatory diseases like (osteo)arthritis.[3] The analysis of such clinically relevant molecular recognition processes and screening for potent antagonists with maximal specificity, sensitivity, and resolution are still great experimental challenges on the way to cracking the sugar code and its biomedical implications.[4,5]

Solution-state NMR spectroscopy can probe molecular interactions with atomic resolution over a wide range of affinities,[6] and NMR screening is now widely used in pharmaceutical research. It may be implemented with detection of suitable proteins, which requires their isotopic labelling, evinces binding sites, and includes irreversible binding. Alternatively, the unlabelled small ligands are observed, which allows faster throughput by pooling and spectrally separating compounds, and thus, is particularly suited to detect weak binding, but often overlooks strong binding because only the dissociated state is detected. Inherent problems of $^1$H NMR spectroscopy, where signals often show poor dispersion, complex fine structure (from extensive homonuclear J($^1$H,$^1$H) coupling), and overlap (aggravated by a vast spectral background and intense solvent signals), further complicate ligand identification, especially in mixtures of chemically related compounds. These shortcomings are avoided upon selecting another NMR-active nucleus with superior signal dispersion and no natural background, which may be introduced by chemical derivation. For glycans with their particularly complex and similar $^1$H NMR spectra,[6] the substitution of hydroxyl groups for fluorine or oxygen for selenium atoms are established modifications, for example, to inhibit enzymatic degradation. This also introduces the extremely well dispersed spin-$^{1/2}$ NMR isotopes $^{19}$F and $^{77}$Se, with chemical shift ranges of $\delta \approx 300$ and 3000 ppm, respectively. For $^{19}$F, high natural abundance (100%) and sensitivity (83% of $^1$H) facilitate direct NMR detection and have boosted the development and broad application of diverse $^{19}$F NMR spectroscopy techniques for efficient screening and detailed analysis of molecular interactions.[7] For the rare (7.63% abundance) and insensitive (0.7% of $^1$H) $^{77}$Se isotope,[6] in contrast, direct $^{77}$Se NMR detection[9] is unfavourable. To enhance the sensitivity substantially and enable reliable ligand ranking by affinities, we propose indirect $^{77}$Se detection and initial $^1$H→$^{77}$Se polarisation.
sation transfer, as implemented in the 2D $^1$H/$^{77}$Se HSQMB experiment with CPMG-INEPT out-and-back transfer.

Despite great success, broad applications, and the advanced technical state of $^1$H NMR based interaction studies, analogous $^{77}$Se NMR approaches may become an important complementary technique because the required OH/F and O/Se substitutions label different molecular sites for NMR observation and with distinct biochemical impacts. In particular, whereas OH/F substitution removes a potential hydrogen-bond donor that may impair or even abolish molecular recognition, O/Se exchange in the glycosidic bond has no known adverse effects, but increases its stability against hydrolysis. Several previous findings indicate that selenoglycosides are excellent biomimetics of natural O-glycosides and exhibit similar dynamic and conformational properties to those of thioglycosides, whereas molecular docking analyses predict full biocompatibility with O-glycosides. Methyl selenoglycosides co-crystallise readily with their cognate bacterial, fungal, and human lectins to enable multi-wavelength anomalous diffraction (MAD) phasing of their complex crystal structures. The diselenodiglycoside of β-GlcNac was shown to be a ligand for wheat germ agglutinin by means of $^1$H saturation transfer difference (STD) NMR spectroscopy, whereas its presence in serum was traced by means of $^{77}$Se NMR spectroscopy. Similar to their thioglycosides, selenodigalactoside (SeDGal; Scheme 1) and diselenodigalactoside (DSeDGal) were recognised by human galectins and a ricin-like plant toxin, in contrast to diselenodi-glucoside (DSeDGlc). At the same time, selenoglycoside synthesis has advanced greatly, and $^{77}$Se NMR signal shifts were shown to be sensitive indicators for phenylselenyl acetate binding to α-chymotrypsin and selenomethyl glycoside binding to plant lectins. All studies, however, employed very insensitive direct $^{77}$Se NMR detection.

![Scheme 1. Structures of the selenodiglycosides used in this study.](image)

To further explore the NMR spectroscopy potential of selenated compounds far more efficiently, we herein propose indirect $^{77}$Se detection through $^1$H and initial $^1$H→$^{77}$Se polarisation transfer to increase the sensitivity by $12 = (\gamma_H/\gamma_{^{77}Se})^2$ and $5.23 = (\gamma_H/\gamma_{^{77}Se})^2$, respectively, that is, by a factor of 63 overall. Yet, the approach relies on moderate scalar two- and three-bond $^2$J($^1$H/$^{77}$Se) couplings (10–30 Hz) for out-and-back transfer of $^1$H coherence to a $^{77}$Se spin in the glycosidic bond that require rather long transfer periods $2.3 \approx (3^1J(\nu,^{77}Se))^{-1} = 30–100$ ms. Transfer efficiency is then degraded by transverse $^1$H $R_t$ relaxation and by competing evolution of homonuclear $^2$J($H,H$) coupling that causes periodic signal phase and amplitude modulation. The use of CPMG-INEPT for $^1$H→$^{77}$Se coherence transfer can alleviate these deleterious effects of COSY-type $^1$H($H$,$H$) evolution by enforcing TOCSY-type evolution instead, which is purely in-phase and eventually leads to equipartition of magnetisation in the coupled $^1$H spin system. This removes signal phase modulation and greatly reduces intensity losses from co-evolving $^1$H($H$,$H$) coupling. Moreover, CPMG-INEPT can suppress line broadening from chemical exchange ($R_{\text{exch}}$) that often increases the net $^1$H $R_2^*$ relaxation rate in cases of transient interactions provided that the CPMG echo delay ($\tau$ in Figure 1) is shorter than the lifetime of interchanging states ($\tau_{\text{exch}}$). If this condition is maintained for all ligands binding to the same site of a protein, relaxation losses are minimised and $R_2^*$ is reduced to a population-weighted sum of slow $R_{2,b}$ and fast $R_{2,f}$ relaxation rates for the free (f) and bound (b) ligand state, respectively [Eq. (1)]:

$$R_2^* = p_0 R_{2,b} + p_0 R_{2,f} = R_{2,b} + p_0 (R_{2,\text{protein}} - R_{2,b})$$  \hspace{1cm} (1)

For small ligands ($R_{2,f}$|$R_{2,\text{protein}}$) in the weak binding regime ($p_0 \approx K_a$), $R_2^*$ then becomes proportional to the affinity constant, $K_a$ [Eq. (2)]:

$$R_2^* \approx R_{2,b} + K_a R_{2,\text{protein}}$$  \hspace{1cm} (2)

Thus, the signal intensity ($I$) of a ligand in the presence of a weakly binding protein is attenuated according to Equa-

![Figure 1. Pulse sequence of the 2D $^1$H/$^{77}$Se CPMG-HSQMB experiment.](image)
is not suppressed during chemical shift evolution, thus agglutinin (VAA).

...with Equation (6) suggests $I/D$ pulses $R_1$ and $I/D$ relaxation [Eq. (4)]:

$$I_0 \propto \exp(-2\Delta R_{1D})$$

The $R_{1D}$ dependence then cancels out in the ratio of the signal intensity of a ligand in the presence and absence of a cognate protein [Eq. (5)]:

$$I/I_0 \approx \exp(-2\Delta K_{2D\text{protein}})$$

Thus, relative signal attenuations from binding allow a ranking of two ligands, a and b, that bind to the same site of a protein, and an estimation of the ratio of their affinity constants [Eq. (6)]:

$$K_{a,b}/K_{AB} \propto \ln(I/I_0)/\ln(I/I_0)$$

This approximation holds for small ligands in the weak binding regime [Eq. (2)] if line broadening from chemical exchange ($R_{\text{exch}}$) is suppressed. By ensuring this as well undistorted signal shapes, CPMG-INEPT enhances sensitivity and is critical for reliable ligand ranking by relative affinities. Nevertheless, $R_{\text{exch}}$ is not suppressed during chemical shift evolution, thus adding to $R_{1s}$ relaxation [Eq. (2)] of $^{77}$Se ($t_1$) and $^1H$ ($t_2$) as extra line broadening. Although this enhances indication sensitivity for molecular interactions, it biases signal intensities, but not their integrals. Thus, Equations (4) to (6) strictly apply to signal integrals, but are also approximately valid for signal intensities.

The 2D $^1H$/$^{77}$Se CPMG-HSQMBC (Figure 1) is derived from our previously presented 2D CPMG-HSQMBC experiment with composite $90^\circ-180^\circ_x-90^\circ_y$ inversion pulses for increased CPMG bands at reduced power and clean coherence selection by echo/antiecho gradients ($G_y$, $G_z$). Contrary to the original experiment, which was developed to accurately measure long-range coupling constants, a refocusing second CPMG-INEPT module is inserted prior to FID acquisition to enable $^{77}$Se decoupling in order to obtain sharp and purely in-phase signals for easy quantification of signal intensities or integrals.

A 2D $^1H$/$^{77}$Se HSQMB spectrum of SeDGal and DSeeDGal (Figure 2) illustrates the significant phase distortions (in the $^1H$ dimension) and sensitivity losses due to amplitude modulation from COSY-type co-evolution of homonuclear $^2J(H,H)$ couplings during conventional INEPT transfer. In contrast, CPMG-INEPT suppresses these deleterious effects and allows a quantitative analysis of the substantially enhanced signal intensities.

We then recorded 2D $^1H$/$^{77}$Se CPMG-HSQMBC spectra to analyse binding of SeDGal, DSeeDGal, and SeDGlc (Scheme 1) to three human galectins and to Viscum album agglutinin (VAA). Both selenogalactosides show distinct signal attenuation after addition to Gal-3, an anti-apoptotic and pro-inflammatory effector and Gal-1 antagonist, at a molar excess of 20:1 (Figure 3, right, top) or 60:1 (Figure 3, right, bottom). The extracted $^1H$/$^{77}$Se traces readily show strong signal attenuation for SeDGal (blue: $I/I_0 \approx 8\%$ at 20:1 molar ratio) and weak attenuation for DSeeDGal (red: $I/I_0 \approx 84\%$ at 20:1 molar ratio), whereas the SeDGlc signals (black) remain unaffected. The extent of binding-induced signal attenuations agree well with previously reported IC$_{50}$ values and Equation (6) suggests about 15-fold higher affinity for SeDGal than for DSeeDGal. In the presence of Gal-1 (Figure S1 in the Supporting Information), the signal attenuations for SeDGal (at 20:1 molar ratio: $I/I_0 \approx 10\%$) and DSeeDGal ($I/I_0 \approx 90\%$) are very similar to those caused by Gal-3, thus confirming a similar differential, albeit somewhat weaker, recognition by this related human galectin. In contrast, Gal-7 (Figure S2) provokes less attenuation of SeDGal signals (at 20:1 molar ratio: $I/I_0 \approx 36\%$), whereas DSeeDGal signals are again attenuated only weakly ($I/I_0 \approx 92\%$), thus suggesting that Gal-7 discriminates less between SeDGal and DSeeDGal than Gal-1 or Gal-3. Finally, VAA causes comparable signal attenuation for SeDGal and DSeeDGal (Figure S3: $I/I_0 \approx 22$ and 27\%, respectively), which confirms a very similar affinity for VAA for these two ligands, in contrast to the galectins. As a negative control, none of the four lectins affected SeDGlc signal intensities, in line with their specific galactose, but not glucose, recognition. In summary, induced signal attenuations observed in 2D $^1H$/$^{77}$Se CPMG-HSQMBC spectra appear to be a valid indicator of ligand binding and allow their reliable ranking by relative affinities.

Our experiments with mixtures of three selenoglycosides implicitly register their competitive displacement if they share a common binding site. To verify this assumption, we also recorded signal attenuations in the 2D $^1H$/$^{77}$Se CPMG-HSQMBC spectrum after addition of only DSeeDGal (2.5 mM) to a lectin.
(125 µM; 30 µM for Gal-3 to avoid its precipitation), and after subsequent coaddition of SeDGal (2.5 mM). The latter caused a significant rebound of the DSeDGal signal attenuation ($I/I_0$) with all four test lectins, that is, from 55 to 91% for Gal-1 (Figure S4), 67 to 86% for Gal-3 (Figure 4), 89 to 97% for Gal-7 (Figure S5), and 12 to 37% for VAA (Figure S4). Thus, SeDGal displaces DSeDGal from all four lectins, which confirms their competitive binding at the same recognition site. This makes DSeDGal a suitable spy ligand to indirectly monitor galectin binding by means of 2D $^1$H,$^{77}$Se CPMG-HSQMBC through its competitive displacement, which allows application of this specific and sensitive detection method to be extended to non-selenated compounds as well. In such assays with a single selenated spy ligand, the 1D version of the experiment, that is, a $^1$H($^2$F)-filtered CPMG-INEPT, may easily be combined with other sensors of molecular interaction than the signal attenuation (from increased $R_2$ relaxation) exploited here. For instance, intermolecular $^1$H$_{\text{pmm}}$—$^1$H$_{\text{ligand}}$ saturation transfer can be combined with the extreme spectral resolution, clarity, and specificity of $^{77}$Se editing in a 2D STD-$^1$H,$^{77}$Se-HSQMBC experiment in order to study transient molecular interactions without spectral background and artefacts. For analyses of stable protein–ligand complexes, the same

Figure 3. Selenoglycoside binding to Gal-3 monitored by 2D $^1$H,$^{77}$Se CPMG-HSQMBC. The 2D spectra of the ligand mixture containing DSeDGal (red), SeDGal (blue), and SeDGlc (black) at 2.5 mM concentrations each were recorded before (left) and after (right) adding Gal-3 to a protein/ligand ratio of 1:20 (top) and 1:60 (bottom). $^1$H($^2$F) traces extracted at the three $^{77}$Se signals are shown next to the contour plots. The indicated attenuated signal intensities from binding are relative to the free ligand intensity. Each 2D $^1$H,$^{77}$Se CPMG-HSQMBC spectrum was measured and plotted with identical parameters. Further experimental details can be found in the Supporting Information.

Figure 4. Competitive displacement of diselenodigalactoside (DSeDGal) binding to Gal-3, monitored by 2D $^1$H,$^{77}$Se CPMG-HSQMBC. A) $^1$H($^2$F) traces of DSeDGal (red, top) and SeDGal (blue, bottom) in the absence of Gal-3, yielding the reference signal intensities, $I_0 = 100\%$. B) The $^1$H($^2$F) trace of DSeDGal (2.5 mM) after adding Gal-3 (30 µM, that is, molar ratio $= 0.75:60$) reveals a binding-induced signal attenuation to $I/I_0 = 67\%$. C) Equimolar addition of SeDGal (2.5 mM) causes a rebound of the attenuated DSeDGal signal to 86% (red, top), thus indicating its competitive displacement. The SeDGal spectrum is conversely attenuated to 57% (blue, bottom), which confirms its preferred binding by Gal-3.
CPMG-HSQMBC building block may be used, for example, in a diagonal free 2D $^{75}$Se HSQC experiment to substantially increased sensitivity and resolution compared with that of the previously proposed direct 1D $^{75}$Se detection method.\(^\text{[19]}\) The proposed 2D $^{1}H$/$^{75}$Se CPMG-HSQMBC experiment instead detects $^{75}$Se indirectly in a second spectral dimension, by employing $^{1}H$/$^{75}$Se CPMG-INEPt out-and-back transfer through $^{1}J(H,Se)$ long-range coupling. As shown for the example of selenoglycoside recognition by three clinically relevant human galectins and the plant toxin VAA, this experiment enables efficient screening of selenated ligands, sensitive detection of molecular binding, and reliable ligand ranking by relative affinities. The range of applications is vast and similar to that of $^{19}$F NMR-based techniques, whereas selenium introduction through O/Se exchange has far less potential impact on molecular interactions and can target other sites than OH/F exchange. This may be particularly important and beneficial in glycoscines, where O/Se exchange in the glycosidic bond is now well established and allows glycan/protein recognition to be studied directly at this often critical linkage, and without risk of impairing molecular interactions, in contrast to OH/F exchange.

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**Conflict of Interest**

The authors declare no conflict of interest.

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[1] a) P. J. Winterburn, C. F. Phelps, *Nature* 1972, 236, 147; b) R. A. Laine in *Glycosciences: Status and Perspectives* (Eds.: H.-J. Gabius, S. Gabius), Chapman & Hall, London, 1997, p. 1; c) H.-J. Gabius, H.-C. Siebert, S. Andry, J. Jiménez-Barbero, H. Rüdiger, *ChemBioChem* 2004, 5, 740; d) H.-J. Gabius, J. Roth, *Histochem. Cell Biol.* 2017, 147, 111.

[2] a) H. Lü, N. Sharon, *Chem. Rev.* 1998, 98, 637; b) J. C. Manning, A. Romero, F. A. Habermann, G. García Caballero, H. Kaltner, H.-J. Gabius, *Histochem. Cell Biol.* 2017, 147, 199; c) H. Kaltner, G. García Caballero, A.-K. Ludwig, J. C. Manning, H.-J. Gabius, *Histochem. Cell Biol.* 2018, 149, 547.

[3] a) F. L. Harrison, C. J. Chesterton, *FEBS Lett.* 1980, 122, 157; b) D. Solis, N. V. Bovin, A. P. Davis, J. Jiménez-Barbero, A. Romero, R. Roy, K. Smetana, Jr., H.-J. Gabius, *Biochim. Biophys. Acta* 2015, 1850, 186; c) H. Kaltner, S. Tögel, G. García Caballero, J. C. Manning, R. W. Ledeen, H.-J. Gabius, *Histochem. Cell Biol.* 2017, 147, 239; d) S. Mayer, M. K. Rauf, B. Lepenies, *Histochem. Cell Biol.* 2017, 147, 223; e) D. Weimann, M. K. Sen, H. Schmidt, K. Schmidt, S. M. Walzer, B. Kubista, R. Windhager, W. Schreiner, S. Tögel, H.-J. Gabius, *Cell. Mol. Life Sci.* 2018, 75, 4187; f) Q. Xiao, A.-K. Ludwig, C. Romano, I.Buzzacchera, S. E. Sherman, M. Vetro, S. Vertesy, H. Kaltner, E. H. Reed, M. Möller, C. J. Wilson, D. A. Hammer, S. Oscarson, M. L. Klein, H.-J. Gabius, V. Percece, *Proc. Natl. Acad. Sci. USA* 2019, 115, E2509; g) A.-K. Ludwig, M. Michaal, Q. Xiao, U. Gille, F. J. Medrano, H. Ma, F. G. Fitzgerald, W. D. Hasley, A. Melendez-Davila, M. Liu, K. Rahimi, N. Yu Kostina, C. Rodriguez-Emmenegger, M. Möller, I. Lindner, H. Kaltner, M. Cudic, D. Reusch, J. Kopitz, A. Romero, S. Oscarson, M. L. Klein, H.-J. Gabius, V. Percece, *Proc. Natl. Acad. Sci. USA* 2019, 116, 2837.