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Design strategies for molecular thermometers by magnetic resonance are essential for enabling new non-invasive means of temperature mapping for disease diagnoses and treatments. Herein we demonstrate a new design strategy for thermometry based on chemical control of the vibrational partition function. To do so, we performed variable-temperature 59 Co NMR investigations of four air-stable Co(III) complexes: Co(accp)₃ (1), Co(bzac)₃ (2), Co(tBu₂-acac)₃ (3), and Co(acac)₃ (4) (accp = 2-acetylcyclopentanonate; bzac = benzoylacetonate; tBu₂-acac = 2,2,6,6-tetramethyl-3,5-heptanedionate and acac = acetylacetonate). We discovered 59 Co chemical shift temperature sensitivity ($\Delta\delta/\Delta T$) values of 3.50(2), 3.39(3), 1.63(3), and 2.83(1) ppm °C⁻¹ for 1–4, respectively, at 100 mM concentration. The values observed for 1 and 2 are new records for sensitivity for low-spin Co(III) complexes. We propose that the observed heightened sensitivities for 1 and 2 are intimately tied to the asymmetry of the accp and bzac ligands *versus* the acac and tBu₂-acac ligands, which enables a larger number of low-energy Raman-active vibrational modes to contribute to the observed $\Delta\delta/\Delta T$ values.

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Introduction

Thermometry by magnetic resonance imaging, which utilizes nuclear magnetic resonance (NMR) technology, is a powerful technique for noninvasive monitoring of temperature. This capability is important in biomedical treatments, *e.g.* monitoring the progression of thermal ablation of tumors. Moreover, the technique is also important for simply achieving a deeper fundamental understanding of internal temperature regulation in the body, which is poorly understood and still yielding new insights to this day. Designing molecular probes that enable high temperature/spatial resolution *in vivo* is an essential component of improving thermometer capabilities.

One method for temperature detection via NMR utilizes a temperature-dependent chemical shift. This technique can use protons (^{1}H) , $^{5-7}$ ^{19}F , 8,9 ^{13}C , 10,11 and ^{31}P . 12,13 Yet, the temperature sensitivities of these systems are often low, <1 ppm $^{\circ}\text{C}^{-1}$ (e.g., 0.01 ppm $^{\circ}\text{C}^{-1}$ for ^{1}H in H_{2}O , 6 or 0.012 ppm $^{\circ}\text{C}^{-1}$ in ^{19}F as perfluorotributylamine 9). A related technique, PARACEST (PARAmagnetic Chemical Exchange Saturation Transfer), can imbue a temperature dependence to the ^{1}H chemical shift of bulk water through hyperfine coupling in open-shell lantha-

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nide complexes. ^{14,15} These strategies also generally produce temperature sensitivities of ca. 1 ppm ${}^{\circ}C^{-1}$.

The ⁵⁹Co nucleus in low-spin Co(III) complexes is a particularly attractive alternative system for chemical-shift-based thermometry applications. ¹⁶ The ⁵⁹Co nucleus is 100% naturally abundant, exhibits a nuclear spin of $I = ^7/_2$, has a high receptivity (*ca.* 30% that of ¹H), and displays a wide reported chemical shift (δ) window (20 000 ppm), which reflects an exquisite sensitivity of δ to electronic structure and the ligand field splitting ($\Delta_{\rm o}$). ^{17,18} Importantly, the ⁵⁹Co nucleus also typically yields a large temperature-dependent chemical shift, $\Delta \delta/\Delta T$, of 1–3 ppm °C⁻¹ in magnitude, which stems from the temperature dependence of $\Delta_{\rm o}$. In 1980, Levy and co-workers set the record for $\Delta \delta/\Delta T$ with Co(acac)₃ (acac = acetylacetonate), which displayed a 3.15 ppm °C⁻¹ $\Delta \delta/\Delta T$ for the ⁵⁹Co nucleus. ¹⁹

A target sensitivity of ca. 10 ppm °C⁻¹ for ⁵⁹Co is ideal to balance the typical wide linewidths of the peaks, and the gradient range in an MRI scanner, and avoid the need for retuning the instrument mid-measurement. A molecule to reach this target parameter should ideally also be air stable to facilitate biomedical imaging applications. In light of these requirements, the recently reported²⁰ record (>100 ppm °C⁻¹) sensitivities in [(CpCo(OP(OR)₂)₃)₂]Co⁺ (R = Me, Et, tBu) are too large. The complexes are also air sensitive, and these two properties together limit applicability. Separately, one complex that is closer to the 10 ppm °C⁻¹ goal, [tBu (PNP)Fe–H], exhibits thermal sensitivities of 14 ppm °C⁻¹ (1 H)²¹ and 34 ppm °C⁻¹ (31 P)²² though these molecules are also, unfortunately, air sensitive. Hence, design strategies to maximize $\Delta \delta/\Delta T$ in an air-

stable compound remains a key target of fundamental understanding.

We recently proposed a route to a higher $\Delta \delta/\Delta T$ through designing the vibrational partition function. In this strategy, the target parameter for higher $\Delta \delta/\Delta T$ is a larger number of low-energy symmetric (Raman-active) vibrational modes.²³ A group-theory analysis of the CoO6 coordination shell of Co $(acac)_3$ reveals 15 possible vibrations, but local D_3 symmetry ensures that only 12 of these are Raman active (Fig. 1). We hypothesized that the 3 lost vibrations could be diminishing the partition function, and, hence, lowering $\Delta \delta / \Delta T$. We hypothesized that a system with asymmetric bidentate ligands would lower local point group symmetry to C_3 (or lower), symmetries where all vibrations are Raman active, and hence ensure a higher partition function and thus higher $\Delta \delta / \Delta T$.

Herein we report the synthesis and ⁵⁹Co NMR characterization of a set of Co(III) complexes with asymmetric analogues of the acac ligand: Co(accp)₃ (1, accp = 2-acetylcyclopentanonate), $Co(bzac)_3$ (2, bzac = benzoylacetonate), $Co(tBu_2-acac)_3$ (3, tBu_2 -acac = 2,2,6,6-tetramethyl-3,5-heptanedionate) and Co $(acac)_3$ (4, acac = acetylacetonate), Fig. 2. We hypothesized that the ligands of 1 and 2 would enforce a higher $\Delta \delta/\Delta T$ than the original report of Co(acac)₃. ¹⁹ We show that this asymmetryfocused design strategy based on maximizing the vibrational partition function can indeed be harnessed to raise $\Delta \delta / \Delta T$.

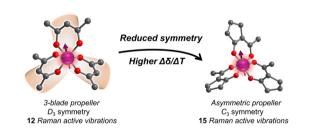


Fig. 1 The CoO_6 coordination geometry in D_3 symmetry (like a threebladed propeller) has 15 possible vibrations, only 12 are active for temperature sensitivity. In contrast, the molecule on the right, with asymmetric ligands, could exploit all 15 for temperature sensitivity.

Results and discussion

Syntheses and molecular structures

Compounds 1-4 are all pseudooctahedral, low-spin Co(III) complexes. Single crystals of 1 were grown by slow evaporation from acetonitrile (MeCN), and the structures of 2-4 were previously reported²⁴⁻²⁶ - all are shown in Fig. 2. All complexes exhibit six-coordinate inner coordination shells with 6 donor O atoms. Average Co-O bond distances are 1.88(2) Å for 1, which are in the range of reported average distances of 1.81(2), 1.86(1), and 1.90(2) for 2-4. The average O-Co-O bite angle for 1 is 89.2(2)°, close to 90.04(13), 92.3(2), and 92.9(3)° for 2-4, respectively. Continuous-shape-measurement analyses were also performed to check for geometric structural trends. 27,28 For an idealized octahedral O_h geometry, SHAPE scores are 0. The scores for 1-4 are 0.34, 0.28, 0.08 and 0.37, respectively, indicating that all four complexes closely resemble perfect octahedral geometry.

The ligands in 1-4 direct the molecular point group of the complex. In 3 and 4, which have C_{2v} -symmetric acac and tBu_2 acac ligands, the local coordination environments of the Co(III) ions are D_3 symmetry. In contrast, 1 and 2, which have asymmetric ligands, have two possible forms. If the ligands in these complexes are all oriented similarly, then local C_3 symmetry is possible, resulting in a fac isomer. If one of the ligands is flipped in orientation relative to the other two, the local symmetry is C_1 (the *mer* isomer). In the single-crystal diffraction data, only the mer isomer is observed for 1. However, in the structure of 2 one of the bzac ligands is disordered and hence the mer and fac isomers are both present in the single crystal.24

Electronic structures. We first studied the electronic structures of the cobalt(III) ions in 1-4 by electronic absorption spectroscopy. UV-vis absorption spectra of compounds 1-4 measured in CH₂Cl₂ at room temperature (Fig. 3a and Fig. S1†) revealed two sets of absorption bands. Absorptions with molar absorptivity coefficients, ε , of ca. 30-80 M⁻¹ cm⁻¹ around 16 700 cm⁻¹ (600 nm) give 1-4 deep green colors in solution, and these are likely d-d transitions, specifically the 1 A_{1g} to 1 T_{2g} transition. $^{29-31}$ All molecules also give more

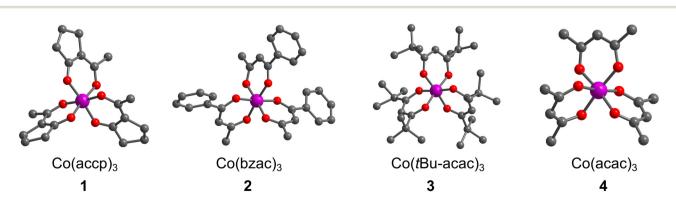
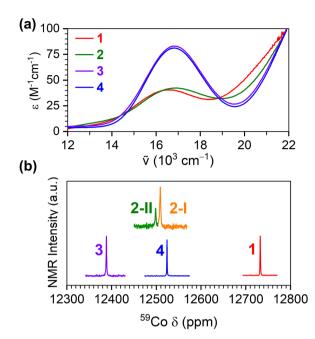


Fig. 2 The crystal structures of the Co(III) complexes studied in this manuscript. Structures of 2-4 are taken from previous reports. 24-26 H atoms are omitted for clarity. Color scheme; Co: violet, O: red, C: gray.





Research Article

Fig. 3 Characterization of Co(III) electronic structure in 1-4. (a) UV-vis spectra for 1-4 in CH₂Cl₂ at room temperature, focused on the d-d transition. (b) 59 Co NMR spectra (ca. 118.3 MHz) for 1-4 in CDCl₃ at room temperature. Spectra were collected in a 500 MHz ¹H magnet (11.7 T).

intense (ca. 250 M⁻¹ cm⁻¹) peaks at much higher energies; 29 000 and 30 600 cm⁻¹ for 1 and 4. These same high-energy transitions are very broad for 3, spanning 28 571 to 33 333 cm⁻¹. Lastly, two transitions of this type are observed for 2 at 34 965 and 38 610 cm⁻¹. These high-energy transitions are likely ligand π - π * transitions.³²

The energies of the low-energy peaks and the d⁶ Tanabe-Sugano diagram³³ allowed us to quantify Δ_0 , which increases in order from 1 (16502 cm⁻¹) to 4 (16850 cm⁻¹) to 2 (16 852 cm⁻¹) to 3 (16 885 cm⁻¹). The value of 4 is consistent with the literature value, 29 whereas the values for the other three have not been previously reported. In all, these values suggest generally weak ligand fields for the Co(III) ions and control for the possibility of substantial ligand field differences causing the observed trends in $\Delta \delta / \Delta T$.

For all four complexes, the 59Co NMR chemical shift varied within 12 400 to 12 800 ppm (Fig. 3b). These values are within the range of ⁵⁹Co NMR peaks of previously reported all-oxygenbound octahedral Co(III) complexes. 16 An important feature of the ⁵⁹Co NMR spectra of 2 is the presence of two peaks. We note that prior DFT calculations and ¹H NMR analyses of 2 reveal both fac and mer isomers to be stable, 34 and disorder in the crystal structure of 2 indicates the presence of both isomers. We tentatively assign the two ⁵⁹Co NMR peaks to the respective isomers as 2-I (for mer) and 2-II (for fac). This assignment is based on literature references indicating that the fac isomer exhibits greater shielding, narrower linewidth and lower signal intensity. 34,35 We note that only one peak is observed for 1, suggesting that only one isomer is present,

which is consistent with the lack of any disorder in the crystal structure. Finally, the known correlation of 59Co chemical shifts to Δ_o^{17} indicates the magnitude of Δ_o increases in the order of 1 < 4 < 2 < 3, in agreement with the lowest-energy peak in the UV-vis data above.

We also conducted diffuse-reflectance spectroscopy (DRS) on these compounds for comparison between solution and solid-state electronic structures and, by proxy, physical structure. The diffuse reflectance spectra (Fig. S2†), transformed from reflectance to the Kubelka-Munk parameter F(R), have generally sharper features than the solution phase UV-vis spectra. For 1-4, two to three peaks noticeable below 25 000 cm⁻¹. The energies of the lowest-energy peaks increase in order from 2 (13 626 cm⁻¹) to 4 (13 633 cm⁻¹) to 3 (13 645 cm⁻¹) to 1 (14 004 cm⁻¹) in the solid-state, but are generally close in energy, like the solution-phase UV-vis data. This general similarity leads us to conclude there are no large differences structurally from compound to compound in solution, at least with respect to the CoO6 coordination shell. However, the lowest-energy peaks are also generally lower than solution-phase values. This distinction, and other differences in the data, suggests the possibility of small structural changes in the solution phase, though we note the colors of the compounds in the solid state and the solutions are strikingly similar (Fig. S2†), suggesting a lack of gross changes to coordination geometry/structure/oxidation state, consistent with the general chemical stabilities of low-spin Co(III) ion bound by chelating ligands.36

Variable-temperature, solvent, and concentration ⁵⁹Co NMR analyses. We collected variable-temperature ⁵⁹Co NMR spectra to determine the temperature sensitivity of the ⁵⁹Co nuclei in 1-4. Spectra were first collected on 10 mM solutions in CDCl₃ from 5 to 35 °C in the 11.74 T magnet of a 500 MHz (1H) NMR (see Fig. S3-S6†). With increasing temperature, the ⁵⁹Co peaks for all complexes shift downfield to higher δ . Linear regression of the temperature dependence of the ⁵⁹Co chemical shifts yields the values of 2.71(1), 2.67(2), 1.55(4), and 2.78(2) ppm °C⁻¹ for 1-4, respectively. We note that prior variabletemperature 59Co NMR analysis of 4 set the prior record for $\Delta \delta / \Delta T$ for ⁵⁹Co NMR in air-stable Co(III) complexes at 3.15 ppm °C⁻¹. But, that original set was recorded at 3.52 T with a 100 mM solution in CDCl3. We further note that Kanakubo later reported³⁷ a $\Delta \delta / \Delta T$ for 4 in CHCl₃ as 2.79(2) ppm, which aligns with our findings.

We tested the matrix sensitivity of the 59Co thermal responses of 1-4 through variable-solvent (see Tables S3-S7†) and variable-concentration studies (see Tables S13, S14 and Fig. S8, S12, S16 and S25†). We selected a variety of solvents: CHCl₃, CH₂Cl₂, CD₂Cl₂, MeOH, DMSO and 1,2,4-trichlorobenzene (TCB), based on variance in physical properties such as polarity (from 2.2 to 47 for solvent TCB and DMSO, respectively) and viscosity (from 0.43 cP for CH2Cl2 to 32.9 cP for TCB).³⁸ Notably, 1-4 are not water-soluble and, hence, were not tested in aqueous solutions. The 59Co NMR peak and its temperature sensitivity differed in each solvent (see Table S2†). The highest ⁵⁹Co $\Delta \delta/\Delta T$ values were observed for 1, 2-II, 2-III,

and 4 in CDCl₃, 2.71(1), 2.69(1), 2.64(2), and 2.78(1) ppm °C⁻¹ and 3, 2.00(3) ppm °C⁻¹, in MeOH (see Tables S3–S7 and Fig. S7, S11, S15, S19 and S22†). Among these, the sensitivity of 3 was found to be the lowest of 1–4 in each solvent. In 2–4, only slight changes (*ca.* 0.4 ppm °C⁻¹) were observed in $\Delta \delta/\Delta T$ with deuteration and no effect for 1.

The origin of the solvent sensitivity likely stems from an impact from the solvent cage, though it is unclear precisely how that mechanism functions. In Co(III) amine complexes, solvent dependence of $\Delta \delta/\Delta T$ has been suggested to result from hydrogen-bonding interaction with donor solvents.³⁹ In 1-4, there are no hydrogen bonding possibilities, which precludes that interaction as one of importance for $\Delta \delta/\Delta T$. There are also no apparent direct relationships between $\Delta \delta/\Delta T$ and polarity or viscosity of the solvent (Fig. S28 and S29†). Finally, we recognize that the solvent dependence is counterintuitive on the basis of sterics because of the variation of the chemical shift of 3 with solvent, which is wider than 1, 2, and 4. One might expect, given the steric congestion afforded by the tBu functional groups, that 3 should have the weakest solvent dependence, yet it does not, in agreement with some previous observations by Kanakubo.37

Variable-temperature ⁵⁹Co NMR spectra were collected on **1–4** at 1, 10, and 100 mM, revealing changes in the chemical shift and its temperature sensitivity with concentration (Fig. 4

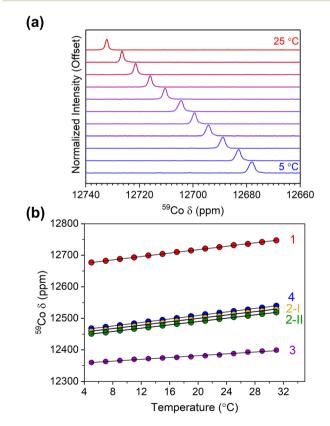
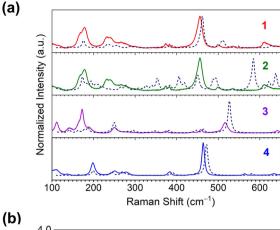


Fig. 4 (a) Variable-temperature 59 Co NMR for 1 in CDCl₃ at 100 mM concentration. (b) Peak positions for 1–4 as a function of temperature at 100 mM concentration in CDCl₃.

and Fig. S7, S11, S15†). The ⁵⁹Co peak for compounds 1-4 in a 100 mM solution exhibited a downfield shift in the range of approximately 1-3 ppm compared to a 1 mM solution. Notably, the value of $\Delta \delta / \Delta T$ changed significantly, particularly for compound 1, increasing to 3.50(2) ppm $^{\circ}\text{C}^{-1}$, at the increased concentration (Fig. 4). A similar trend is observed for 2, where the $\Delta \delta / \Delta T$ value increased from 2.70(2) to 3.23(2) ppm ${}^{\circ}\text{C}^{-1}$ for 2-I, and from 2.64(2) to 3.26(4) ppm ${}^{\circ}\text{C}^{-1}$ for 2-II. Similarly, the $\Delta \delta/\Delta T$ value increased from 1.55(4) to 1.63(3) ppm °C⁻¹ for 3. Lastly, in 4 the ⁵⁹Co peak position appears significantly less dependent on concentration; only a small increase was observed in the sensitivity, from 2.78(2) to 2.83(2) ppm ${}^{\circ}C^{-1}$. To the best of our knowledge, the observed value of 3.50 ppm ${}^{\circ}C^{-1}$ for 1 establishes a record for temperature sensitivity not only among closed-shell Co(III) complexes, but for all closed-shell species.

The origin of the concentration dependence of $\Delta \delta/\Delta T$ needs further investigation. It is reasonable to propose aggregation effects enhance the temperature sensitivity on account of the observed concentration dependence. However, there is no clear chemical difference in 1-4 that presents the possibility of aggregation in an intuitive way, e.g. functional groups hydrogen bonding or open coordination sites. Furthermore, if such interactions exist, they are likely not magnetic in nature because they would be too weak. For instance, the dipole-dipole interaction between two ⁵⁹Co nuclei held at 7.47(1) Å apart (from the structure of 1, the closest Co···Co distance in the crystal structures of 1-4 and likely closest possible distance in solution) is calculated to be only ca. 0.3 Hz (~6 ppm), significantly smaller than the total ranges observed (ca. 80 ppm, see Fig. 4). An alternative explanation of the concentration dependence is that aggregation is modifying how the individual 59Co nuclei interact with the solvent cage. Changes in this interaction may induce changes in the electric field gradient at the cobalt nucleus, 40-43 which might also contribute to the concentration dependence of the chemical shift. Dynamic light scattering experiments as a function of concentration and solvent suggest that some degree of aggregation may be happening (see ESI†), though these changes do not trend with $\Delta \delta/\Delta T$, or any other magnetic resonance characteristic, across all studied concentrations and solvent choice. Note, finally, that the test of viscosity dependence of $\Delta \delta/\Delta T$ through the variable-solvent studies appears to disprove a mechanism stemming directly from increased viscosity of the higher-concentration solvents, or simply slower molecular motion in solution from aggregation.

Vibrational analyses. We anticipated that a descent in symmetry in the coordination environment would amplify the number of low-energy Raman-active modes in the complex, and therefore increase $\Delta\delta/\Delta T$. Hence, we applied Raman spectroscopy to evaluate the effect of the descending ligand symmetry on the vibrational spectra of **1–4** (Fig. 5). The selected spectral window (<650 cm⁻¹) targets vibrations that are appreciably populated at room temperature and the expected energy range of Co–L vibrations. As the probed molecules decrease in symmetry, the Raman spectra increase in complex-



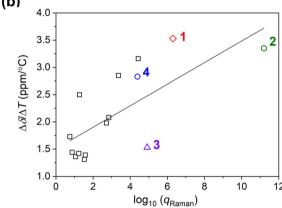


Fig. 5 (a) Room-temperature Raman spectra collected on microcrystal-line samples of 1–4. All spectra are baseline corrected. The dotted lines indicate the calculated spectra for the complexes. (b) The relationship between the $^{59}\mathrm{Co}$ temperature sensitivity $(\Delta\delta/\Delta T)$ and total vibrational partition function of Raman modes, q_{Raman} , for 1–4 (colored data points), and other Co(III) complexes (square data points). The linear correlation by R^2 is 0.49. Data for the other complexes are taken from ref. 15.

ity: asymmetric 1 and 2 display 12 and 17 bands below 650 cm⁻¹, respectively, whereas symmetric 3 and 4 exhibit 9 and 10 bands respectively. An exact correlation between the observed numbers of peaks and the simple group-theory analysis in the introduction (Fig. 1) does not occur. We suspect that this imperfect correlation is due to 1–4 having far more atoms than the local CoO₆ moiety and that there are potentially different isomers in 1 and 2. Note, however, that the crystal structure of 1 seems to indicate only a single *mer* isomer, while the crystallographic data for 2 suggest that both *fac* and *mer* isomers are present. Hence, for 2 some of the additional peaks may stem from different isomers, whereas for 1 new peak (relative to 4) are likely attributable to the descent in symmetry and splitting of *E*-symmetry Raman modes. There are additional considerations described further below.

Computational analysis of the vibrational spectra enabled assignment of the observed modes in the Raman spectra and comparison with $\Delta\delta/\Delta T$. The experimental spectra match the predicted spectra closely (see Fig. 5a). Experimentally, the most intense modes in all species are symmetric Co–O bond

stretching modes ("breathing" modes). These modes are observed at 456 cm⁻¹ for 1, and in the region of 563–568 cm⁻¹ for 2. For the symmetric complexes, this band is observed at 517 and 463 cm⁻¹ for 3 and 4, respectively. Bands below 200 cm⁻¹ are mainly related to torsional modes of methyl or alkyl groups and the chelating ring.⁴⁴

Role of ligand asymmetry in $^{59}\text{Co}\ \Delta\delta/\Delta T$. We compared $\Delta\delta/\Delta T$ to the vibrational partition functions, q_{Raman} ; of 1–4 using the computed vibrations from the frequency calculations concerning only the Raman active modes (see ESI for the calculation details and Tables S8–S11†). We compared these values against the aggregate of $q_{\text{Raman}} \nu$. $\Delta\delta/\Delta T$ data compiled in previous works^{23,45} (Fig. 5). In general, the higher q_{Raman} values trend with higher $\Delta\delta/\Delta T$. Together, the comparison therefore suggests that a model for predicting temperature sensitivity based on a higher density of low-energy Raman vibrations is mostly correct, and that a ligand-asymmetry design strategy is one way to enhance $\Delta\delta/\Delta T$ through that model.

However, we note that the observed correlation between $q_{\rm Raman}$ and $\Delta \delta / \Delta T$ is weak, as indicated by the low correlation coefficient ($R^2 = 0.49$), and suggests the opportunity for significant future insight. We propose three important initial hypotheses behind the weak correlation for future study. First, we propose the need to factor in the extent to which a vibration is focused on the inner coordination shell into the partition function analyses. One may surmise a ligand-only vibration should have little impact on the 59 Co δ versus a Co-O-based mode, but that intuition needs quantitative testing, which could be available by local mode analyses. 46 Second, we propose that couplings between the solvent cage (or bulk solvent) and individual vibrational modes also need to be investigated in detail, because these interactions may ultimately alter which modes should be weighted for temperature sensitivity. This latter point may ultimately explain the concentration dependence if aggregation affects the couplings between specific vibrations and the solvent cage. Finally, third, we propose that understanding the exact physical structure of the species in solution is important, as the diffuse reflectance suggests some small alteration of the coordination shell. Such dynamic structural changes likely affect the vibrational spectra, and how this relates to the temperature sensitivity is a key future target.

⁵⁹Co NMR linewidth analyses. As one final experimental characterization, we performed linewidth analyses of the observed ⁵⁹Co peaks for **1–4** in all solvents and as a function of temperature (Fig. 6a). The linewidths ultimately help us describe the thermometer resolution (see below). The ⁵⁹Co NMR peaks for **1–4** in CDCl₃ at 25 °C all exhibit relatively sharp peaks with full width at half maximum values ($\nu_{1/2}$) of 1.12, 2.37, 1.38, and 0.85 ppm, respectively. These already narrow linewidths sharpen further with increasing tempera-

[‡] q_{Raman} values are calculated from the product of each normal mode by $q_{\text{Raman}} = \prod_n \left(\frac{1}{1-e^{-E_n}/k_{\text{B}}T}\right)$ where E_n is the energy of the nth normal mode (cm^{-1}) , k_{B} is Boltzmann's constant, and T is the temperature (K).

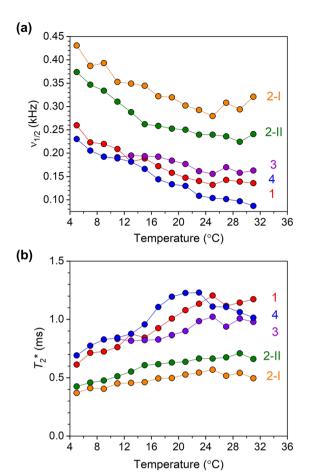


Fig. 6 (a) 59 Co NMR linewidth values for 10 mM solutions of 1-4 in CDCl₃ as a function of temperature. (b) Temperature dependence of T_2^* values for 1-4, calculated using ⁵⁹Co NMR peak linewidths.

ture for 1-4 (Fig. 6a). In all complexes, only slight changes (ca. ~0.1 ppm in magnitude) were observed in linewidth with solvent (see Fig. S9, S13, S17, S20 and S23†), but in general, the narrowest peaks were observed in CH2Cl2, CDCl3, and MeOH. Finally, for 2, the linewidth of isomer I is found to be slightly broader than isomer II in all measured solvents. In all complexes, the broadest 59Co NMR peaks are observed in DMSO.

The ⁵⁹Co NMR linewidths contain information about the relaxation time T_2^* , the dephasing time of the nuclear spin (analogous to the coherence time, T_2). The values of T_2^* were extracted from the temperature-dependent NMR linewidths through the relationship $T_2^* = 1/(2\pi\nu_{1/2})$ where $\nu_{1/2}$ (kHz) is the linewidth (at FWHM) of the 59Co NMR peak. The temperaturedependent trends in T_2^* for all complexes over the 5-30 °C range are shown in Fig. 6b (see also Fig. S10, S14, S18, S21, and S24†) Complexes 1-4 all show generally increasing T_2^* with increasing temperature, where the temperature dependence is stronger for 1, 3 and 4 than both isomers of 2. Among all four complexes, 1 and 4 show the longest values of T_2^* . For both 1 and 4, the longest T_2^* values are found in MeOH; 1.2 and 1.6 ms for 1 and 4, respectively. While deuteration

increases T_2^* in CDCl₃ for 1, only a small effect on T_2^* was observed in CD_2Cl_2 . On the other hand, an increase in T_2^* in 4 upon deuteration was observed. For both isomers of 2, T_2^* values are found to be similar. In all complexes, except 2-II and 3 where the latter is insoluble, noticeably shorter T_2^* values are observed in DMSO. The T_2^* values were also found to decrease for all four complexes as the concentration increased from 10 mM to 100 mM. We also measured the spin-spin relaxations (T_2) for 1-4 via Carr-Purcell-Meiboom-Gill ("CPMG") experiments at room temperature (Fig. S26†). The T_2 values are 2.38(3), 0.99(2), 1.98(3) and 3.47(4) ms for 1-4, respectively, all generally higher than the T_2^* values observed, but only a factor of 2. The T_2^* value is only determined for isomer 2-II due to limitations in the data collections for 2-I.

The T_2^* value of a quadrupolar nucleus like ⁵⁹Co is often dictated by spin-lattice relaxation (T_1) , if T_1 is short enough that it limits the magnitude of T_2^* . We measured T_1 relaxation times via inversion recovery of 1-4 at room temperature (295 K) in 100 mM CDCl₃ solution (Fig. S27†). The T_1 values are 4.15(4), 1.23(2), 1.10(2), 2.24(3) and 4.01(4) ms for 1-4, respectively, all on the order of the T_2 values obtained from CPMG and T_2^* from simple linewidth analyses. The effects of the T_1 times are observed in the ⁵⁹Co peak linewidths. Complexes 1 and 4, with the longest T_1 values, have the narrowest 59 Co NMR peaks, whereas 2 and 3 exhibit shorter T_1 times. Hence, the linewidths here appear to follow spin-lattice relaxation times for the ⁵⁹Co nuclei in 1-4.

There is additional importance of the linewidths and relaxation times in the context of the concentration dependence. Aggregation should be expected to slow down molecular rotation, potentially inducing larger linewidths through lengthening the correlation time.⁴⁷ If true, then the measured T_1 values would shorten, as would T_2 , and all spectra would increase in linewidth with the increased aggregation observed from the DLS measurements. That we do not see these trends for relaxation time measurements from 1 to 100 mM concentrations (see Table S12† for DLS results and Tables S13 and S14 for variable-concentration relaxation data) suggests that whatever the nature of the observed aggregation, it is not having conventional impacts on the ⁵⁹Co NMR characteristics. Further, expecting the T_1 and T_2 data to follow the aggregation through correlation time variation does not acknowledge the substantial quadrupolar coupling for ⁵⁹Co, which is important for spin relaxation of this nucleus. 48-50 If aggregation were noticeably distorting the structure of the molecules, then these two competing systems ensure this system is far from trivial. Understanding these effects is key to sharper linewidths in future systems, and hence a key target for our future study.

⁵⁹Co NMR temperature sensing resolution. A key aspect of a sensor's utility is the resolution afforded in the sensing, which has not been described in the literature for 59Co NMR systems to the best of our knowledge. We can parameterize this resolution through a ratio that compares the temperature sensitivity $(\Delta \delta/\Delta T)$ with the full width half maximum of the peak $(\nu_{1/2})$: $(\Delta \delta/\Delta T)/\nu_{1/2}$. A higher ratio (in units of °C⁻¹) will enable

higher resolution of temperature sensing. The ratio points out an intuitive point: a sharp peak with high $\Delta\delta/\Delta T$ is advantageous for higher resolution thermometry.

We calculated the temperature resolution for 1-4 and contrasted them against values for other prominent NMR thermometer systems (Table 1). In complex 1, the highest temperature sensitivity and the narrowest peak were observed in CDCl₃, which led to the ratio of 3.18, which is higher than Co(acac)₃ under the same conditions that we measured 1-4 in this paper. We also note that both isomers of complex 2 show lower resolution than 1 and 4 despite the high $\Delta \delta/\Delta T$, owing to the relatively wide linewidths of the two isomers. Among ⁵⁹Co NMR thermometers, these are the highest resolutions, eclipsing the recent spin-crossover 59Co results.20 They are also notably higher than the record temperature sensitivities for ¹H and ³¹P. The sole measurement that eclipses our results in this manuscript is the original measurement of Co(acac)3. Again, we stress the difference in the measurement technique as an essential part of the comparison, specifically the field of measurement which is likely affecting linewidth (11.7 T vs. ca. 3.2 T in the original Levy report). The resolution here is likely limited by the field of the measurement and the resolution for 1 would likely eclipse 4 if measured under the same conditions.

Conclusion

Tuning the number of allowed and forbidden IR and Ramanactive modes in a molecule is a common exercise in inorganic and graduate level inorganic chemistry courses. Here we show that applying the same course-work level analyses to conventional Co(III) complexes can yield new design strategies for $\Delta\delta/\Delta T$. Indeed, we present two new records for the temperature sensitivities of the ⁵⁹Co chemical shift in the asymmetric complexes Co(accp)₃ and Co(bzac)₃, the first in the over-40-year period since Co(acac)₃ was initially identified as an exciting NMR thermometer. ¹⁹

Table 1 Linewidths, $\nu_{1/2}$, and calculated temperature resolutions, $(\Delta\delta/\Delta T)/\nu_{1/2}$, for **1–4** and other notable NMR thermometers

Compound	$\nu_{1/2}^{a}$ (ppm)	$(\Delta \delta/\Delta T)/\nu_{1/2} (^{\circ}\text{C}^{-1})$
Co(acac) ₃ ^b	0.86	3.66
$Co(accp)_3 (1)^c$	1.10	3.18
$Co(bzac)_3 (2)^c$	2.47	1.37
$Co(tBu_2-acac)_3 (3)^c$	1.34	1.22
$Co(acac)_3 (4)^c$	0.91	3.11
$K_3[Co(CN)_6]^d$	0.40	2.98
[Co(diNOsar)]Cl ₃ ^d	12	0.17
[(CpCo(OP(OtBu) ₂) ₃) ₂]Co ^{+ e} H-[tBuPNPFe-H] ^f	128	1.18
¹ H-[tBuPNPFe-H] ^f	50	0.28
31 P-[t Bu(PNP)Fe-H] g	121	0.28

^a From room temperature values. ^b From original ref. 19 by Levy *et al.* ^c From this work. ^d Collected in H₂O at 33.3 mM concentration, 500 MHz magnet, see ref. 38. ^e Collected in CH₂Cl₂ at 400 mM, 500 MHz magnet, see ref. 20. ^f For ¹H NMR signal, from ref. 21. ^g For ³¹P NMR signal, from ref. 22.

While our study has predominantly focused on the design strategies to enhance temperature sensitivity in ⁵⁹Co NMR thermometry using air-stable Co(III) complexes, it is imperative to recognize that the translation of these findings into practical applications, particularly in biological or clinical settings, requires addressing issues related to toxicity and biocompatibility. Future investigations should explore encapsulation techniques and other strategies to increase stability and to mitigate potential toxicity concerns when applying these molecular thermometers *in vivo*. ^{52–55}

In a similar vein, it is important to note that new molecular imaging agents do not simply appear, but follow years of fundamental studies of magnetic properties in molecules. 52,56,57 It is in this context that this work should be appreciated for testing a new mechanism to increase temperature sensitivity in an ⁵⁹Co nucleus by molecular design. Yet, we acknowledge that further work is still necessary. For example, the trend in $q_{\rm Raman}$ with $\Delta \delta/\Delta T$ is not a perfect correlation, implying future experiments are needed, and while the measurements here focus on high-field analyses, conventional MRI is often performed at lower fields, suggesting low-field studies are also imperative. Finally, we need to understand what mechanisms are important for governing linewidth, as this knowledge could lead to higher resolution. Two parameters, quadrupolar coupling of the $I = \frac{7}{2}$ ⁵⁹Co nucleus and chemical shift anisotropy, are both proposed to dictate ⁵⁹Co NMR linewidth.⁵⁸ Studies of all of these effects will be reported in due course.

Author contributions

Ö. Ü. – methodology, investigation, formal analysis, validation, visualization, writing – original draft. S. S. – investigation, formal analysis. T. M. O – investigation, formal analysis. J. M. Z. – conceptualization, methodology, formal analysis, validation, funding acquisition, supervision, writing – review & editing.

Conflicts of interest

There are no conflicts to declare.

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