Magnetic Properties

Single-Molecule Magnets DyM₂N@C₈₀ and Dy₂MN@C₈₀ (M = Sc, Lu): The Impact of Diamagnetic Metals on Dy³⁺ Magnetic Anisotropy, Dy···Dy Coupling, and Mixing of Molecular and Lattice Vibrations

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Abstract: The substitution of scandium in fullerene single-molecule magnets (SMMs) DyScN@C₈₀ and Dy₂ScN@C₈₀ by lutetium has been studied to explore the influence of the diamagnetic metal on the SMM performance of dysprosium nitride clusterfullerenes. The use of lutetium led to an improved SMM performance of DyLuN@C₈₀, which shows a higher blocking temperature of magnetization (T_BLK = 9.5 K), longer relaxation times, and broader hysteresis than DyScN@C₈₀ (T_BLK = 6.9 K). At the same time, Dy₂LuN@C₈₀ was found to have a similar blocking temperature of magnetization to Dy₂ScN@C₈₀ (T_BLK = 8 K), but substantially different interactions between the magnetic moments of the dysprosium ions in the Dy₂MN clusters. Surprisingly, although the intramolecular dipolar interactions in Dy₂LuN@C₈₀ and Dy₂ScN@C₈₀ are of similar strength, the exchange interactions in Dy₂LuN@C₈₀ are close to zero. Analysis of the low-frequency molecular and lattice vibrations showed strong mixing of the lattice modes and endohedral cluster vibrations in k-space. This mixing simplifies the spin–lattice relaxation by conserving the momentum during the spin flip and helping to distribute the moment and energy further into the lattice.

Introduction

Enclosing lanthanide ions within the fullerene cage is a versatile route to a variety of molecular magnets.[1] In particular, when non-metal atoms (C, N, O, S) are captured by the carbon cage together with lanthanides, the strong ionic interactions emerging in such endohedral clusterfullerenes may lead to a large magnetic anisotropy.[2] In addition, different magnetic states can emerge from the intramolecular interactions of lanthanide ions in clusterfullerenes.[2a, e, 3] This combination of properties made lanthanide-clusterfullerenes promising candidates for single-molecule magnets (SMMs). Single-molecule magnetism is a phenomenon involving the slow relaxation of magnetization in molecules with a bistable magnetic ground state and has been a hot topic in the field of molecular magnetism during the last decades.[4] Indeed, some lanthanide-clusterfullerenes exhibit single-molecule magnetism,[5] which is most robust in dysprosium-clusterfullerenes.[6, 7, 8] Nitride clusterfullerenes (NCFs) with the composition (M³⁺)ₙN³⁻@Cₙ₈₀, and in particular the species with C₈₀ = C₈₀⁻Ịₙ (Ịₙ denotes the symmetry of the carbon cage) and M = Sc, Y, or heavy lanthanides (Gd–Lu), show the largest synthetic yield and are therefore the most studied clusterfullerenes to date.[2a, c, 6b, 8] The structure of the trimetal nitride cluster, with N⁺ in its center and three M³⁺ ions located at the vertices of the triangle with rather short M–N bonds, offers a convenient platform to create a variety of molecular magnets with divergent properties. First of all, the nitride ion at a distance of only 2.0–2.2 Å from the lanthanides[2a, 6b, 8] generates a strong axial ligand field (LF), which leads to an easy-axis magnetic anisotropy for cerium, praseodymium, neodymium, terbium, dysprosium, and holmium ions, and an easy-plane anisotropy for erbium and thulium ions[26] (note that we use the term “ligand field” instead of the more commonly used “crystal field” to avoid unnecessary connotations to intermolecular interactions in crystals). The strong axial LF also ensures that mₗ is a good quantum number, at least for several lowest-energy LF states. In particular, the lowest-energy Kramers doublets for Dy³⁺ in dysprosium-scandium NCFs are essentially pure mₗ states, with the high-spin Jₗ = ± 15/2 states as the ground Kramers doublet with the magnetic moment oriented along the Dy–N bond.[26a–c]
The purity of the $J_z$ states in terms of $m_j$ composition ensures that the magnetic system is weakly susceptible to external perturbations such as those introduced by dipolar magnetic fields from neighboring molecules or molecular and lattice vibrations.

Another advantage that lanthanide NCFs offer for tuning magnetic properties is the possibility of combining different metals within one molecule to give the so-called mixed-metal NCFs.\[5a, 8c, 9\] The importance of this can be best illustrated with the dysprosium-scandium NCFs Dy$_x$Sc$_{1-x}$N@C$_8$I$_h$ ($x = 1–3$) as an example.\[3a] As $Sc^{3+}$ is diamagnetic, this series essentially allows analysis of how two or three dysprosium ions interact magnetically and the effect of these interactions on SMM behavior. Dy$_x$Sc$_{1-x}$N@C$_8$I$_h$ was the first endohedral metallofullerene (EMF) proven to be a SMM.\[3b, d]\ At temperatures below 7 K, the molecule exhibits magnetic hysteresis with the abrupt drop of the magnetization in zero magnetic field ascribed to the quantum tunneling of magnetization (QTM), which is typical for single-ion magnets. Dy$_x$Sc$_{1-x}$N@C$_8$I$_h$ shows magnetic hysteresis below 8 K without fast QTM relaxation in zero field, which is explained by ferromagnetic exchange and dipolar coupling between the non-collinear magnetic moments of the two dysprosium ions in the Dy$_2$ScN cluster (Figure 1), thus creating an additional barrier and preventing zero-field QTM.\[1a]\ The temperature dependence of the magnetization relaxation times in Dy$_x$Sc$_{1-x}$N@C$_8$I$_h$ revealed a high barrier of 1735 K, because of the Orbach relaxation via the fifth Kramers doublet, in good agreement with CASSCF calculations.\[1b]\ Similar ferromagnetic interactions between dysprosium ions are also present in Dy$_x$Lu$_{1-x}$N@C$_8$I$_h$, but the triangular arrangement of the dysprosium ions forbids simultaneous realization of ferromagnetic coupling for all three Dy–Dy contacts resulting in a frustrated ground state (Figure 1) with faster relaxation of magnetization than in the mono- and dinuclear analogues.\[2a, 3a]\ Thus, due to different cluster compositions and intramolecular Dy–Dy interactions, Dy$_x$Sc$_{1-x}$N@C$_8$I$_h$, Dy$_x$ScN@C$_8$I$_h$, and Dy$_x$N@C$_8$I$_h$ exhibit substantially different SMM behavior at low temperatures.

It is natural to consider that the magnetic properties of the NCFs in the Dy$_x$Sc$_{1-x}$N@C$_8$I$_h$ ($x = 1–3$) series discussed above are determined by the Dy$^{3+}$ ions, with the diamagnetic Sc$^{3+}$ ions acting just as placeholders, which are needed to keep the trimetallic cluster composition. However, in this work, we shed more light on the role that the diamagnetic metal can play in the SMM properties of NCFs. For this, we chose lutetium as a diamagnetic lanthanide, the ionic radius ($R^{3+} = 0.86\ A$) of which is noticeably larger than that of scandium ($R^{3+} = 0.75\ A$). Yet, as we reported recently, Dy$_x$Lu$_{1-x}$N@C$_8$I$_h$ and Dy$_x$LuN@C$_8$I$_h$ can still be synthesized, albeit in lower yields than the dysprosium-scandium analogues.\[6c]\ The larger size of Lu$^{3+}$ leads to changes in the internal structure of the trimetallic nitride cluster, and in particular results in shorter Dy–N bonds than in the dysprosium-scandium NCFs. The shortening of the Dy–N bonds may affect the magnetic anisotropy and the strength of the intramolecular Dy–Dy coupling. Furthermore, lutetium is much heavier than scandium, which changes the low-frequency part of the vibrational spectrum. Because the relaxation of magnetization involves spin–phonon interactions, alterations of molecular vibrations may also result in changes in the SMM properties. Thus, the goal of this work was to study a possible influence of these factors on the SMM properties of the Dy$_x$M@C$_8$I$_h$ and Dy$_x$M$_2$@C$_8$I$_h$ NCFs.

**Results and Discussion**

**Magnetization behavior of Dy$_x$Lu$_{1-x}$N@C$_8$ and Dy$_x$LuN@C$_8$**

The mixed-metal dysprosium-lutetium NCFs with a C$_8$I$_h$ fullerene cage (for clarity, the symmetry designation will be omitted hereafter) were obtained by arc-discharge synthesis and separated by recycling HPLC as reported earlier.\[3b]\ Similar retention behavior of Dy$_x$N@C$_8$, Dy$_x$LuN@C$_8$, Dy$_x$LuN@C$_{10}$, and Lu$_x$N@C$_{10}$ substantially complicated the separation of the individual compounds. However, a compositional purity exceeding 90–95% could be achieved for Dy$_x$LuN@C$_{10}$ as verified by MS analysis. Due to the similar retention behavior, the separation of Dy$_x$LuN@C$_{10}$ from Lu$_x$N@C$_{10}$ was not possible, and the relative content of the two NCFs in the studied sample was around 1:3.5. Because Lu$_x$N@C$_{10}$ is diamagnetic, its presence in the sample does not lead to strong changes in the magnetic properties of Dy$_x$LuN@C$_{10}$. A possible influence of dilution on the quantum tunneling of magnetization will be specifically considered below.

The magnetization curves of Dy$_x$LuN@C$_{10}$ and Dy$_x$LuN@C$_{10}$ measured by SQUID (superconducting quantum interference device) magnetometry, are shown in Figures 2 and 3, respectively. Dy$_x$LuN@C$_{10}$ exhibits hysteresis up to 9 K at a sweep rate of 3 mT s$^{-1}$ (Figure 2a). The “butterfly” shape of the hysteresis curves points to efficient zero-field relaxation by QTM similar to that observed in Dy$_x$ScN@C$_{10}$ (Figure 2b).\[6b\] However, in the latter, the opening of the hysteresis is narrower and the QTM induces complete loss of magnetization at zero field, whereas in Dy$_x$LuN@C$_{10}$ upon crossing zero-field, the magnetization drops to around 30% of the saturation magnetization value, resulting in a coercivity of 0.9 T at 2 K. Recently, we showed that the QTM in Dy$_x$ScN@C$_{10}$ is strongly affected by dilution in the diamagnetic matrix, including dilution with Lu$_x$N@C$_{10}$.\[6b\] To...
ensure that the difference in the QTM is intrinsic for the two NCFs and is not caused by the dilution of DyLu$_2$N@C$_{80}$ with Lu$_3$N@C$_{80}$. In Figure 2b we also show the magnetization curve of DySc$_2$N@C$_{80}$ diluted with Lu$_3$N@C$_{80}$ in a 1:1 ratio. The inset in (b) shows the determination of the blocking temperature, $T_B$, from the temperature dependence of the magnetic susceptibility, $\chi$ (temperature sweep rate of 5 K min$^{-1}$).

Similarly to Dy$_2$ScN@C$_{80}$, Dy$_2$LuN@C$_{80}$ exhibits open hysteresis without a pronounced QTM step (Figure 3a). Its blocking temperature of magnetization, $T_B = 8$ K, is the same as that of Dy$_2$ScN@C$_{80}$ [24]. However, at 2 K, the magnetic hysteresis of Dy$_2$LuN@C$_{80}$ is narrower (Figure 3b) and the coercive field of 0.4 T is smaller than that of Dy$_2$ScN@C$_{80}$ (0.7 T). Thus, the substitution of scandium by lutetium narrows the magnetic hysteresis but does not change the temperature scale of the slow relaxation. Nevertheless, as we show below, the temperature dependence of relaxation times reveals that the mechanisms of the relaxation of magnetization in Dy$_2$LuN@C$_{80}$ and Dy$_2$ScN@C$_{80}$ are different.

Relaxation times of magnetization in Dy$_2$M@C$_{80}$ and Dy$_2$MN@C$_{80}$

The opening of magnetic hysteresis in the magnetization curves of molecular magnets indicates that the magnetization attains its equilibrium value slower than the rate of the magnetic field sweep. The slow relaxation of magnetization is a key characteristic of SMMs and needs to be understood in detail. The relaxation of magnetization requires energy exchange between the spin system and a thermal bath, which is mediated...
by the phonon system. The spin–phonon interaction is crucial for such an energy exchange, and the role of vibrational degrees of freedom becomes paramount. Several mechanisms of phonon-mediated spin relaxation have been recognized in studies of paramagnetic salts and adapted for SMMs.

A direct mechanism implies a single-phonon process in which phonon frequency matches the energy difference between two opposite spins. Because a magnetic field increases the energy gap between the opposite spins, and the phonon density at near-zero frequency is low and increases with frequency, the relaxation due to a direct mechanism accelerates in a magnetic field according to Equation (1)

\[ \tau_{\text{M,dir}}^{-1}(H) = A_1 H^2 T + A_2 H^4 T \]

in which \( \tau_{\text{dir}} \) is the magnetization relaxation time, \( H \) is the magnetic field, with two terms describing the relaxation of a Kramers ion in the absence (ca. \( H^0 \)) and in the presence (ca. \( H^2 \)) of hyperfine interactions, and \( A_1 \) and \( A_2 \) are fitting parameters. Thus, the relaxation rate scales linearly with temperature, \( \tau_{\text{M,dir}}^{-1} \sim T \). However, when at low temperature the phonon density may be very low, the energy exchange between the phonon system and the bath can become the limiting step. This effect, known as a phonon bottleneck, can change the temperature dependence to \( \tau_{\text{M,dir}}^{-1} \sim T^2 \).

The Raman mechanism implies a spin flip through the absorption and emission of two phonons, with the frequency difference being equal to the energy gap between the opposite spins. As the frequencies can be much higher than the Zeeman energy, and the phonon density increases with frequency, the mechanism becomes more efficient than the direct mechanism once the temperature is sufficiently high to ensure sufficient phonon population. The original consideration with only acoustic phonons in the Debye model gave the power-law temperature dependence expressed by Equation (2)

\[ \tau_{\text{M,Ram}}^{-1}(T) = C T^n \]

in which \( n = 9 \) for Kramers ions and \( n = 7 \) for non-Kramers ions, and \( C \) is the fitting parameter. However, if optical phonons are also included in the model, powers of 6, 5, and even lower can be expected.

A special case of the Raman mechanism is the energy matching of the opposite spin levels, and therefore the application of a magnetic field can quench the QTM when Zeeman splitting becomes large enough.

It has become common practice to analyze the relaxation of magnetization in SMMs as a combination of these processes. Characteristic temperature and field dependencies of relaxation times allow identification of the prevailing relaxation mechanism. Usually, the direct mechanism is the most important at the lowest temperatures of a few K, at somewhat higher temperatures the Raman mechanism becomes dominant, and with further increases in temperature, the Orbach mechanism involving LF excited states takes over. Neither the direct nor Raman mechanism would show an Arrhenius temperature dependence, so the latter usually serves as an indication of the Orbach mechanism. However, back in the 1960s, Klemens and others argued that a localized vibrational mode can cause an Arrhenius temperature dependence with \( U^\text{eff} \) corresponding to the frequency of the mode. Furthermore, it was shown that the direct mechanism can also demonstrate Arrhenius behavior in some conditions.

The relaxation times of magnetization in DyLu\(_3\)N@C\(_{80}\) and DyLuN@C\(_{80}\) in this work were determined at different temperatures by magnetizing the sample to saturation, quickly sweeping the magnetic field to zero or any other required field value, and then following the decay of magnetization with subsequent fitting of the measured decay curves with a stretched exponential (see the Supporting Information for the decay curves and Tables containing all the fitted parameters). Below, the values obtained here are compared with those of DyScN@C\(_{80}\) from ref. (6b), whereas the relaxation times in DyScN@C\(_{80}\) were re-measured in this work for better consistency.

The zero-field measurements for DyLu\(_3\)N@C\(_{80}\) are complicated by the relatively long stabilization of the field and fast zero-field QTM. In addition, the decay curves showed two types of behavior: A fast drop of magnetization for around 90% of the sample, followed by a much slower relaxation of the remaining magnetization. The fit of the decay curves with two stretched exponents gave values ranging from 54 ± 1 s at 1.8 K to 21 ± 6 s at 5 K for the fast process (Figure 4a), which we assigned to QTM. The abrupt change in the relaxation rate may be caused by the redistribution of dipolar fields in the sample when a significant number of spins flip, the dilution effect of Lu\(_3\)N@C\(_{80}\) and may also reflect different relaxation of the molecules with different dysprosium isotopes. In addition, the slow process may also be caused by slow relaxation of the remnant magnetization in the magnet, or the deviation of the real magnetic field from zero. For comparison, \( \tau_{\text{QTM}} \) in non-diluted DyScN@C\(_{80}\) determined by AC magnetometry in the same temperature range is 1–3 s. Overall, we can conclude that the rate of relaxation by the QTM mechanism in DyLu\(_3\)N@C\(_{80}\) is slower than in DyScN@C\(_{80}\). Note that both compounds show a temperature dependence of the relaxation rate, even in the QTM regime, which may reflect the temperature dependence of the phonon collision rate, as suggested by Chilton and co-workers, or result from dipolar intermolecular interactions.
in a field of 0.2 T, applied to 10-

reasons. In addition, the $T_s$ values are many orders of magnitude longer than are usually found for the Orbach mechanism. We tentatively propose that the relaxation of magnetization in SMM EMFs in this temperature range may follow the Raman mechanism with involvement of local vibrations that would also follow this temperature range may follow the Raman mechanism with involvement of local vibrations that would also follow

Short relaxation times cannot be measured very reliably by DC magnetometry, and further relaxation measurements were performed for DyLuN@C$_{60}$ in a field of 0.2 T, applied to quench the QTM. Over a range of only 5 K, the in-field relaxation times in DyLuN@C$_{60}$ vary by almost four orders of magnitude, from $4.7 \times 10^2$ s at 2 K to 58 s at 7 K (Figure 4a). When plotted in Arrhenius coordinates, the temperature dependence of the relaxation times has a linear form below 5 K. For comparison, the $r_{\text{arr}}$ values of DySc,N@C$_{60}$ are systematically shorter than those of DyLu,N@C$_{60}$ by a factor of six, but they also show a linear temperature dependence with a very similar inclination (Figure 4a). Fitting the relaxation times of DyLuN@C$_{60}$ with Equation (3) gave a $U_{\text{eff}}$ value of 24.2 ± 0.7 K and $r_0$ of 2.8 ± 0.5 s. The analogous fit for DySc,N@C$_{60}$ gave $U_{\text{eff}} = 23.6 ± 1$ K and $r_0 = 0.6 ± 0.2$ s.$^{[6d]}$ Thus, both NCFs have essentially identical $U_{\text{eff}}$ values and differ only in their attempt times. The reason for the low-temperature $U_{\text{eff}}$ barriers of 24 K in both DyM,N@C$_{60}$ molecules is not clear. Due to the very strong LF splitting of dysprosium in the NCFs, the energies of the lowest-energy excited states exceed hundreds of K (see the discussion of the ab initio calculations below).$^{[2a–d, f, g]}$ In addition, the $r_s$ values are many orders of magnitude longer than are usually found for the Orbach mechanism. We tentatively propose that the relaxation of magnetization in SMM EMFs in this temperature range may follow the Raman mechanism with involvement of local vibrations that would also follow

$$r_{\text{arr}}(T) = C T^{n_2} + r_0^\text{eff} \exp(-U_{\text{eff}}/T)$$

(4)

The fit of the experimental zero-field relaxation times of DyLuN@C$_{60}$ with Equation (4) gives $C = (1.14 ± 0.28) \times 10^{-6}$ s$^{-1}$ K$^{-5.45}$, $n_1 = 5.45 ± 0.15$, $r_0 = 435 ± 50$ s, and $U_{\text{eff}} = 4.3 ± 0.2$ K (Table 1). The Raman mechanism dominates above 4 K, whereas Arrhenius behavior is predominant below 2.5 K, and both mechanisms make comparable contributions in between. As for DyLuN@C$_{60}$, neither $r_0$ nor $U_{\text{eff}}$ of the Arrhenius part is typical for the standard Orbach relaxation process via ligand-field excited states. Spin relaxation in DySc,N@C$_{60}$ below 8 K can be also described by Equation (4) with $C = (0.51 ± 0.26) \times 10^{-6}$ s$^{-1}$ K$^{-5.91}$, $n_1 = 5.99 ± 0.33$, $r_0 = 56 ± 4$ s, and $U_{\text{eff}} = 8.0 ± 0.1$ K (somewhat different values, $r_s = 11.9 ± 1.5$ s and $U_{\text{eff}} = 10.7 ± 0.3$ K, reported by us in ref. [2d], were obtained with a smaller

![Figure 4](https://example.com/figure4.png)

Figure 4. (a) Magnetization relaxation times of DyLuN@C$_{60}$ measured in zero field and in a field of 0.2 T and compared with those of DySc,N@C$_{60}$ from ref. [6] (the zero-field values for DySc,N@C$_{60}$ were measured by AC magnetometry). The lines are fits of the Arrhenius behavior. (b) Magnetization relaxation times of DyLuN@C$_{60}$ measured in zero field and in a field of 0.2 T and compared with the relaxation times of DySc,N@C$_{60}$ in zero field. The solid lines are total fits with Equations (4) and (5) (0 and 0.2 T, respectively). The contributions of the Arrhenius, Raman, and direct (for 0.2 T) processes are also shown. (c) Field dependence of magnetization relaxation times of DyLuN@C$_{60}$ and DySc,N@C$_{60}$ measured at 2.5 K. The green line is the fit of DyLuN@C$_{60}$ data with Equation (1). (d) Comparison of the magnetization relaxation times of DyLuN@C$_{60}$ and DySc,N@C$_{60}$.

| Table 1. Low-temperature relaxation parameters for DyM,N@C$_{60}$ and DyM,N@C$_{60}$ (M = Sc, Lu). |
|--------------------------------------------------------------------------------------------------|
| **QTM** | **Raman** | **Arrhenius** | **Direct** |
| $r_{\text{arr}}$ | $T^n$ | $n_1$ | $r_0$ | $U_{\text{eff}}$ [K] | $A(T)^{[7]}$ [s$^{-1}$ K$^{-5}$] |
| DyLu,N, 0 T | 20–50 | – | – | 2.8 ± 0.5 | 24.2 ± 0.7 | – |
| DyLu,N, 0.2 T | – | – | – | 0.6 ± 0.2 | 23.6 ± 1.0 | – |
| DySc,N, 0 T | 1–3 | – | – | 2.8 ± 0.5 | 24.2 ± 0.7 | – |
| DySc,N, 0.2 T | – | – | – | 0.6 ± 0.2 | 23.6 ± 1.0 | – |
| DyLu,N, 0 T | 1.14 ± 0.28 | 5.45 ± 0.15 | 435 ± 50 | 4.3 ± 0.2 | – |
| DyLu,N, 0.2 T | 1.14 ± 0.28 | 5.45 ± 0.15 | 435 ± 50 | 4.3 ± 0.2 | 7.59 ± 0.57 | 1.67 ± 0.09 |
| DySc,N, 0 T | 0.51 ± 0.26 | 5.99 ± 0.33 | 56 ± 4 | 8.0 ± 0.1 | – |
| DySc,N, 0.2 T | – | – | – | – | – |

[a] Below $T_g$. | www.chemeurj.org | 2440 | © 2019 The Authors. Published by Wiley-VCH Verlag GmbH & Co. KGaA, Weinheim | Chem. Eur. J. 2020, 26, 2436 – 2449 | 2440
and Dy have similar parameters in the Raman process, but are significantly different in their Arrhenius behavior, as DyScN@C\textsubscript{80} has a two-fold higher $U^{\text{eff}}$ and an eight-fold shorter $\tau_0$. As a result, although the zero-field relaxation times of DyLuN@C\textsubscript{80} and DyScN@C\textsubscript{80} are very similar at 1.8 K and virtually identical by 7 K, between 2 and 6 K the relaxation of magnetization in DyLuN@C\textsubscript{80} is considerably slower.

Measurements of magnetization relaxation times in DyLuN@C\textsubscript{80} at 2.5 K in different magnetic fields (Figure 4c) revealed considerable acceleration of the relaxation with increasing field. The temperature dependence of the relaxation times measured in a field of 0.2 T also showed noticeable deviations from zero-field values below 5 K (Figure 4b). Such a dependence of $\tau_\text{rel}$ on the magnetic field is a characteristic of the direct relaxation mechanism [Eq. (1)]. To describe the temperature dependence of the relaxation times measured in a field of 0.2 T, we used Equation (4) with the addition of a term describing the direct process, given by Equation (5)

$$
\tau_\text{rel}(T) = CT^n + \tau_0^{-1} \exp(-U^{\text{eff}}/T) + A(H) T^n,
$$

where $C$, $n$, $\tau_0$, and $U^{\text{eff}}$ were fixed to the values determined for zero-field relaxation.

The fit of the experimental data measured in a field of 0.2 T with Equation (5) gives $A(H) = (7.59 \pm 0.57) \times 10^{-5}$ s$^{-1}$ K$^{-1.67}$ and $n_1 = 1.67 \pm 0.009$ (Table 1). The exponent of approximately 1.7 lies between the values expected for normal (n = 1) and bottleneck (n = 2) direct processes, and indicates that both are likely to take place. If instead of using the fitting procedure, the $n_2$ value is fixed to 1 and $A(H)$ is determined from the field dependence at 2.5 K by using Equation (1) (Figure 4c), then Equation (5) describes well the temperature dependence above 2.5 K, but shows increasing deviations at lower temperature. This indicates that the bottleneck process has higher impact at low temperatures, when the number of excited phonons is not sufficient for efficient energy transfer. Note that the direct and Arhenius processes in DyLuN@C\textsubscript{80} have similar rates at 0.2 T, and hence in-field relaxation rates are around twice as fast as up to 3 K, when the Raman process starts to dominate, and the field dependence eventually vanishes by 5 K.

In striking contrast to DyLuN@C\textsubscript{80} the relaxation times of DyScN@C\textsubscript{80} at 2.5 K do not depend on the external magnetic field until it exceeds 0.4 T (Figure 4c). The relaxation times of DyScN@C\textsubscript{80} measured at different temperatures in a field of 0.2 T almost coincide with the zero-field values and start to show small deviations only below 2 K. This shows that the direct mechanism contributes to the spin relaxation in DyScN@C\textsubscript{80} at considerably higher fields and lower temperatures than in DyLuN@C\textsubscript{80}, which explains why the coercive field in the magnetic hysteresis of DyScN@C\textsubscript{80} is larger than in DyLuN@C\textsubscript{80} (Figure 3b).

The temperature dependencies of the relaxation times of DyLuN@C\textsubscript{80} and DyScN@C\textsubscript{80} measured in zero field and in a field of 0.2 T are compared in Figure 4d. Once the QTM in DyLuN@C\textsubscript{80} is quenched by the application of a finite field, its relaxation rate is much slower than in DyLuN@C\textsubscript{80} and at 2 K the difference between mono- and di-dysprosium NCFs exceeds two orders of magnitude. Intramolecular interactions between dysprosium spins in DyM@C\textsubscript{80} block zero-field QTM and create a manifold of new low-energy coupled spin states. Apparently, spin relaxation in DyLuN@C\textsubscript{80} at low temperature proceeds via such coupled states and is therefore much faster than in DyLuN@C\textsubscript{80}, which has only single-ion excited spin states. A similar difference in the low-temperature relaxation mechanisms was also observed for DyScN@C\textsubscript{80} and DyScN@C\textsubscript{80}\textsubscript{C\textsubscript{80}}\textsubscript{C\textsubscript{80}}. If indeed this is the case, the $U^{\text{eff}}$ of the Arrhenius process in DyM@C\textsubscript{80} may be related to the energy difference between the ground and the first excited state of the coupled spin system. At higher temperature, the relaxation rates of DyM@C\textsubscript{80} and DyM@C\textsubscript{80} tend to be more similar, which indicates that relaxation via single-ion states becomes equally efficient for both types of NCFs.

### Single-ion anisotropy of dysprosium ions in DyM@C\textsubscript{80} and DyM@C\textsubscript{80} (M = Sc, Lu)

The central nitride ion is the main source of the magnetic anisotropy in lanthanide NCFs, and the LF is expected to become stronger with decreasing distance between N$^+$ and Dy$^{3+}$. DFT calculations at the PBE level with the 4f-in-core effective potential showed that the increase in the ionic radius from Sc$^{3+}$ to Lu$^{3+}$ shortens the Dy–N bond from 2.156 Å in DyScN@C\textsubscript{80} to 2.090 Å in DyLuN@C\textsubscript{80}. Likewise, the DFT-optimized Dy–N bonds in DyLuN@C\textsubscript{80} (2.073 and 2.074 Å) are shorter than those in DyScN@C\textsubscript{80} (2.105 and 2.108 Å). These geometrical changes may substantially affect the LF acting on the dysprosium ions.

To explore the influence of this effect, we performed ab initio CASSCF/RASSI calculations on the LF splittings in DyLuN@C\textsubscript{80} and DyLuN@C\textsubscript{80} molecules and compared the results with those for the dysprosium-scandium analogues. For DyM@C\textsubscript{80} molecules, only one dysprosium ion was treated ab initio at a time, and the other dysprosium was replaced by yttrium. The energies of the Kramers doublets (KDs) in DyM@C\textsubscript{80} molecules, the pseudo-spin g-tensor of the ground state KD, and the transition probability in the first KD are presented in Figure 5, Table 2, and Table S7 in the Supporting Information. The calculations show that the Dy$^{3+}$ ions in DyLuN@C\textsubscript{80} and DyLuN@C\textsubscript{80} exhibit high magnetic anisotropy with overall LF splitting of 1340–1360 cm$^{-1}$. The quantization axis is aligned along the Dy–N bond, and the ground Kramers doublet is described as an essentially pure state with $|m_z| = 15/2$. The energy of the second Kramers doublet is predicted to be close to 400 cm$^{-1}$, which ensures that the low-temperature magnetic properties of both DyLuN@C\textsubscript{80} and DyLuN@C\textsubscript{80} are determined solely by the ground state of Dy$^{3+}$, as in the previously studied dysprosium-scandium NCFs. A comparison of DyScN@C\textsubscript{80} and DyLuN@C\textsubscript{80} shows that the increase in metal size from scandium to lutetium leads to an increase in LF splitting from 1284 cm$^{-1}$ (DyScN) to 1348 cm$^{-1}$ (DyLuN). The energy of the second KD (relative to the first KD) also shows an increase from 356 cm$^{-1}$ in DyScN@C\textsubscript{80} to 391 cm$^{-1}$ in DyLuN@C\textsubscript{80}. As the Dy–N bond in DyLuN@C\textsubscript{80}
are somewhat shorter than that in DyLuN@C_{60}, the LF splitting for the dysprosium ion in the former is also slightly higher. Likewise, the LF splitting in Dy$_2$LuN@C$_{60}$ is somewhat higher than in Dy$_2$ScN@C$_{60}$, which also correlates with the shorter Dy–N bond lengths (see Tables S8 and S9).

For the relaxation of magnetization, not only the energies of the KD states, but also the transition probabilities between them as well as the composition of the wave functions in the $|J, m_J\rangle$ basis are very important (Figure 5, see also Table S7 in the Supporting Information). The first KD with $g_i$ close to 19.8 and infinitesimally small $g_j$ and $g_i$ values has more than 99% contribution from the $|m_J\rangle = 15/2$ function for all the discussed NCFs. The probability of a QTM transition within the first KD is only $1.1 \times 10^{-9} \mu_B^2$ in Dy$_2$ScN@C$_{60}$ and $1.7 \times 10^{-9} \mu_B^2$ in Dy$_2$LuN@C$_{60}$. The QTM transition probabilities within one KD remain low up to the fourth KD (Figure S5), and similar values are also found for Dy$_2$MN@C$_{60}$ molecules. Likewise, transitions between the states of different $m_i$ and opposite spin are also not efficient until KD4. The reasons for this situation are rooted in the composition of the KD wave functions, which can be described as essentially pure $m_i$ states up to KD4–KD5 (Figure 5).

Thus, ab initio calculations predict that the relaxation of magnetization in all dysprosium NCFs should proceed via the KDs, as indeed was observed experimentally in Dy$_2$ScN@C$_{60}$.[2d] There is no considerable difference between dysprosium-scandium and dysprosium-lutetium nitride clusterfullerenes in this regard.

To summarize, the replacement of scandium by lutetium in mixed-metal nitride clusterfullerenes shortens the Dy–N bonds and increases the LF splitting by 5–10%. Otherwise, there is no significant difference in terms of KD composition and the expected relaxation pathways via excited KDS.

**Intramolecular interactions of dysprosium magnetic moments in Dy$_2$LuN@C$_{60}$**

The system of two weakly interacting dysprosium centers with magnetic moments $J_{i,2}$ can be described by the effective spin Hamiltonian given by Equation (6)

$$\hat{H}_{\text{spin}} = \hat{H}_{\text{eff}} + \hat{H}_{\text{dip}} - 2j_1J_1 \hat{J}_2$$  (6)

in which $\hat{H}_{\text{eff}}$ is the single-ion LF Hamiltonian for the $i$th dysprosium site, dysprosium moments $\hat{J}_i$ are treated in the $|J, m_J\rangle$ basis, and $j_{1,2}$ is the coupling constant between the localized dysprosium moments. Here, $j_{1,2}$ is treated isotropically in the spirit of the lines model[2d] and includes both exchange and dipolar interactions.

To determine the $j_{1,2}$ constant for Dy$_2$LuN@C$_{60}$, we simulated magnetization curves with different values of $j_{1,2}$ and compared them with the experimental data. In these simulations the angle, $\alpha$, between the single-ion anisotropy axes of the dysprosium ions is set to $61.7^\circ$, as determined by ab initio calculations. The best agreement is achieved for $j_{1,2} = 0.02$ - 0.03 cm$^{-1}$ (Figure 6). As follows from Equation (6), for the two dysprosium spins oriented at an angle of $\alpha = 61.7^\circ$, the energy difference between the states with ferromagnetic (FM) and antiferromagnetic (AF) coupling of the dysprosium ions ($U = 15/2$) can be calculated from Equation (7)

$$\Delta E_{\text{FM-AF}} = 4j_{1,2}^2\cos(\alpha) - 225j_{1,2}\cos(\alpha)$$  (7)

which gives an estimation of 2.1 – 3.2 cm$^{-1}$ (3.1–4.6 K) for Dy$_2$LuN@C$_{60}$. This energy difference is close to the $U_{\text{eff}}$ value of 4.3 K determined for the low-temperature relaxation process with Arrhenius behavior (see above). The assumption that the relaxation proceeds by excitation to the antiferromagnetically coupled state (i.e., $\Delta E_{\text{FM-AF}} = U_{\text{aff}}^{[2d,3d]}$) allows a more precise estimation of $j_{1,2} = 0.028 \pm 0.001$ cm$^{-1}$. This value also gives a reasonable agreement between the experimental and simulated $\chi T$ curves (see Figure S8 in the Supporting Information).

Magnetic Dy–Dy interactions have two components, dipolar and exchange. The energy difference between the dipolar interactions in the FM and AF states, $\Delta E_{\text{Dip}}^{\text{FM-AF}}$, can be calculated by using the well-known formula for the energy of dipolar interactions between two magnetic moments [Eq. 8],

$$\Delta E_{\text{Dip}}^{\text{FM-AF}} = \frac{1}{2} \frac{4\pi^2}{3} \mu_B^2 \frac{1}{d^3}$$

where $d$ is the distance between the dysprosium ions.
Table 2. Ligand-field splitting of the Dy\(^{4+}\) states and the ground-state g-tensor in Dy\(_2\)Ni@C\(_{88}\) and Dy\(_2\)MN@C\(_{88}\) (M = Sc, Lu) molecules as predicted by ab initio calculations.

|          | DySc\(_2\)N | DyLu\(_2\)N         | Dy1         | Dy2         | Dy1         | Dy2         |
|----------|-------------|---------------------|-------------|-------------|-------------|-------------|
| KD1 [cm\(^{-1}\)] | 0           | 0                   | 0           | 0           | 0           | 0           |
| KD2 [cm\(^{-1}\)] | 356         | 391                 | 399         | 378         | 408         | 403         |
| KD3 [cm\(^{-1}\)] | 666         | 715                 | 704         | 698         | 726         | 726         |
| KD4 [cm\(^{-1}\)] | 906         | 960                 | 938         | 945         | 967         | 971         |
| KD5 [cm\(^{-1}\)] | 1050        | 1104                | 1081        | 1084        | 1109        | 1112        |
| KD6 [cm\(^{-1}\)] | 1140        | 1194                | 1176        | 1174        | 1203        | 1202        |
| KD7 [cm\(^{-1}\)] | 1222        | 1276                | 1253        | 1263        | 1283        | 1286        |
| KD8 [cm\(^{-1}\)] | 1284        | 1348                | 1336        | 1333        | 1365        | 1366        |

Table 2. Ligand-field splitting of the Dy\(^{4+}\) states and the ground-state g-tensor in Dy\(_2\)Ni@C\(_{88}\) and Dy\(_2\)MN@C\(_{88}\) (M = Sc, Lu) molecules as predicted by ab initio calculations.

\[
\Delta E_{\text{FM-AF}} = 4.7 \, \text{K}, \quad \text{but as the } \Delta E_{\text{FM-AF}} \text{ energy in Dy}_2\text{ScN@C}_{88} \text{ is twice as large as in Dy}_2\text{LuN@C}_{80}, \text{ one can conclude that the exchange term for Dy}_2\text{ScN@C}_{88} \text{ is of a similar size to the dipolar term (Table 3). Thus, substitution of scandium by lutetium in the Dy}_2\text{MN cluster results in considerable variation of the coupling constant, mainly because of the negligible exchange interactions in Dy}_2\text{LuN@C}_{80}, \text{ which also leads to a smaller energy difference between the ferromagnetically and antiferromagnetically coupled states, and through this difference has a strong influence on the relaxation of magnetization at low temperature.}
\]

Table 3. Exchange and dipolar coupling parameters for di-Dy EMFs.

|                  | \(\Delta E_{\text{FM-AF}}\) [K] | \(\Delta E_{\text{FM-AF}}\) [K] | \(j_{12} [\text{cm}^{-1}]\) | \(\varepsilon_{12}^\alpha [\text{cm}^{-1}]\) | \(R_{ij}^\alpha [\text{Å}]\) |
|------------------|---------------------------------|---------------------------------|-----------------------------|---------------------------------|-----------------------------|
| Dy\(_2\)LuN@C\(_{80}\) | 4.3 \pm 0.2                     | 4.8                             | 0.028                       | 61.7                            | 3.559                        |
| Dy\(_2\)ScN@C\(_{80}\) | 8.0 \pm 0.1\(^{10}\)           | 4.7                             | 0.055                       | 63.4                            | 3.576                        |

[a] \(\Delta E_{\text{FM-AF}}\) is considered to be equal to \(U^\alpha\) of the low-temperature Arrenius process. [b] \(\alpha\) is the angle between the quantization axes of two Dy ions, determined from ab initio calculations. [c] \(R_{ij}\) is the distance between two Dy ions in DFT-optimized molecules. [d] The more precise measurements in this work give a somewhat different value from our previous report in ref. [2d] and are closer to the data reported earlier in ref. [3a].

Low-frequency molecular and lattice vibrations in dysprosium-metal NCFs

As follows from the ab initio calculations discussed above, dysprosium-scandium and dysprosium-lutetium NCFs have very similar single-ion magnetic anisotropy and ground-state properties. In addition, the LF splitting in these NCFs is very large and is not relevant for the low-temperature relaxation of magnetization. Therefore, the difference in their relaxation behavior
cannot be explained by the LF splitting and it is necessary to take into account other factors. In addition, the observation of Arrhenius behavior in DySc$_2$Ni@C$_{80}$ and DyLu$_2$Ni@C$_{80}$ with a $U^{\text{eff}}$ of 24 K and unconventional parameters also cannot be explained by LF splitting. A deeper insight into the spin–phonon interactions may be needed to explain these phenomena.

The drawbacks of the commonly applied phenomenological approaches to spin–phonon interactions developed in the 1960s are that they are based on the Debye model for vibrations in the crystal and do not provide a clear connection with the microscopic parameters of the molecules (except for the LF splitting, when the Orbach process is involved). For instance, although the relaxation times measured in this work can be well fitted by Equations (1)-(3) and their combinations, the fitted parameters do not provide sufficient insight into the relation between molecular structure and the relaxation of magnetization.

Recently, Lunghi et al.\cite{17} analyzed spin–lattice relaxation in molecular magnets by using quantum spin dynamics, considering explicit vibrations of the molecule and deriving the spin–phonon coupling parameters from ab initio calculations. They found that spin–phonon relaxation via anharmonic phonons may also result in Arrhenius behavior, but with the $U^{\text{eff}}$ corresponding to half of the vibrational frequency. Furthermore, other research teams analyzed the locality of the spin–phonon interaction and concluded that molecular vibrations spatially localized close to the metal center usually have the strongest contribution to the spin relaxation\cite{17, 19}. These findings show that the Debye model is oversimplified for the analysis of spin–phonon relaxation in molecule magnets\cite{20} and an analysis of the real vibrational spectra may give better insight into the relaxation mechanism.\cite{21} With this in mind, we decided to analyze the low-frequency part of the vibrational spectra of the NCFs. The experimental Raman spectra of DyM$_2$Ni@C$_{80}$ and DyM$_2$Ni@C$_{80}$ (M = Sc, Lu; limited to frequencies above 50 cm$^{-1}$ due to instrument limitations) are compared in Figure 7 with the vibrational density of states (VDOS), computed by DFT for isolated molecules. In addition to the total VDOS, Figure 7 also shows contributions to the VDOS of the whole metal nitride cluster and of only the dysprosium atoms. The computed and experimental frequencies of the cluster-based modes are presented in Table 4.

EMFs have rather peculiar vibrational spectra as their molecules consist of two semi-independent units, the vibrations of which show almost no overlap in the energy scale, as can be

**Table 4.** Experimental and DFT-computed frequencies of cluster-based vibrations in DyM$_2$Ni@C$_{80}$ and DyM$_2$Ni@C$_{80}$ (M = Sc, Lu).\cite{24}

|          | DySc$_2$Ni@C$_{80}$ | DySc$_2$Ni@C$_{80}$ | DyLu$_2$Ni@C$_{80}$ | DyLu$_2$Ni@C$_{80}$ |
|----------|--------------------|--------------------|--------------------|--------------------|
| Mode     | Calc Ex | Calc Exp | Calc Ex | Calc Exp | Calc Ex | Calc Exp | Calc Ex |
|          | R      | R       | R       | R       | R       | R       | R       |
| 41       | 38     | 27      | 32      | 32      | 32      | 32      | 32      |
| 44       | 45     | 33      | 39      | 39      | 39      | 39      | 39      |
| 62       | 53     | 40      | 42      | 42      | 42      | 42      | 42      |
| 89       | 91     | 75      | 79      | 79      | 79      | 79      | 79      |
| 105      | 111    | 78      | 80      | 80      | 80      | 80      | 80      |
| 125      | 122    | 79      | 82      | 82      | 82      | 82      | 82      |
| 168      | 172    | 99      | 93      | 93      | 93      | 93      | 93      |
| 210      | 198    | 157     | 161     | 161     | 161     | 161     | 161     |
| 223      | 218    | 163     | 163     | 163     | 163     | 163     | 163     |

[a] Designation of cluster modes: R, rotation (libration); T, translation; $\delta_{\text{M-N}}$, $\delta_{\text{M-N}}$, $\delta_{\text{M-N}}$; M-N-M bending; $\gamma_{\text{N}}$, nitrogen out-of-plane displacement; $\nu_{\text{M-N}}$, M-N stretching vibration. [b] Experimental Raman spectrum of DySc$_2$Ni@C$_{80}$ has a peak at 145 cm$^{-1}$ that cannot be assigned on the basis of calculation results.

**Figure 7.** Raman spectra ($T = 77 \text{ K}$, $\lambda_{\text{ex}} = 514$ or 647 nm) and DFT-calculated VDOS for (a) DyM$_2$Ni@C$_{80}$ and (b) DyM$_2$Ni@C$_{80}$ (M = Sc, Lu). The total VDOS is shown in gray, and the contributions to the VDOS of the whole M,N cluster and of the Dy atoms are shown in pink and green, respectively.

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well seen in Figure 7. Thus, the vibrations of the relatively rigid carbon cage occur at frequencies exceeding 240 cm\(^{-1}\), whereas the frequencies of metal-involving modes rarely exceed 230 cm\(^{-1}\), because the metal atoms are much heavier than carbon atoms. Only in the border range of 220–260 cm\(^{-1}\) do the squashing cage modes partially mix with the “breathing” mode of the nitride cluster, in which all three metal atoms move radially in one phase along the M–N bond. Other metal-based vibrations occur at frequencies below 200 cm\(^{-1}\). When the nitride cluster is encapsulated inside the fullerene, its external degrees of freedom (i.e., translations and rotations) are transformed into internal ones (i.e., molecular vibrations). The frustrated rotations (i.e., librations) are the lowest-frequency intramolecular modes predicted to be close to 30–40 cm\(^{-1}\) in dysprosium-lutetium NCFs and at 40–60 cm\(^{-1}\) in dysprosium-scandium NCFs. The frustrated translations are mixed with deformations of the cluster (such as in-plane oscillations of the M–N–M angles). In DyLu,N@C\(_{60}\) and Dy, LuN@C\(_{60}\) these modes are clustered into two groups close to 80 and 160 cm\(^{-1}\) (because dysprosium and lutetium have similar atomic masses, both dysprosium-lutetium NCFs have very similar VDOS). In dysprosium-scandium NCFs, such modes are more uniformly spread in the 80–200 cm\(^{-1}\) range. Finally, the nitrogen out-of-plane mode also falls in the range of 90–210 cm\(^{-1}\).

The results of the computations agree reasonably well with the experimental Raman spectra. Above 220 cm\(^{-1}\), the spectra of all the NCFs are quasi-continuous because of densely spaced cage vibrations. The calculations seem to overestimate the cage frequencies by around 10%. In the cluster frequency range, DyLu,N@C\(_{60}\) and Dy, LuN@C\(_{60}\) exhibit only two Raman peaks, at around 80 and 162 cm\(^{-1}\), close to the predicted frequencies of the mixed translation/deformation modes. The dysprosium-scandium NCFs exhibit richer spectral patterns, and most of the observed peaks can be reliably assigned to the computed modes, as listed in Table 4 (see refs. [22] for a more detailed discussion of the vibrational spectra of MSc,N@C\(_{60}\) NCFs). Due to technical limitations, we cannot record the spectra below 50 cm\(^{-1}\), which precludes experimental observation of the cluster librations in dysprosium-lutetium NCFs. However, good agreement between experiment and theory for the cluster modes above 50 cm\(^{-1}\) ensures that the calculated frequencies are not far from reality. In addition, in some of the earlier Raman studies of the NCFs, peaks at around 30–40 cm\(^{-1}\) were reported for Dy,N@C\(_{60}\), Lu,N@C\(_{60}\), and some other M,N@C\(_{60}\) molecules.

**Librations of the cluster and lattice phonons in spin relaxation**

The low-frequency vibrations localized on the metal nitride cluster of the M,N@C\(_{60}\) molecules are expected to mediate energy transfer between the spin and a thermal bath. The rotational motions of the cluster are especially of interest from the point of view of low-temperature spin relaxation because they not only happen at low frequencies (i.e., in the relevant energy range), but also because they may help to conserve the total angular momentum when the spin flips. The relevance of the Einstein–de Haas effect on the single-molecule level was demonstrated by Wemdsorfer and co-workers for TbPC\(_3\) (PC = phthalocyanine) grafted on a carbon nanotube (CNT)\,[24] To conserve the total momentum, the spin reversal of terbium had to result in rotation of the TbPC\(_3\) molecule around the terbium quantization axis. However, because the molecule was rigidly bonded to the CNT, to fulfill the rotational invariance, the rotational momentum had to be transferred to the momentum of the phonon propagating along the nanotube. As a result, the efficient spin reversal proceeded by the direct mechanism when the external magnetic field created a Zeeman splitting matching the frequency of the longitudinal nanotube phonon. In a dysprosium nitride cluster, the reversal of the spin aligned along the Dy–N bond should induce rotation of the cluster around this bond. But as discussed above, interaction with the fullerene cage restricts the rotational motion of the cluster and turns it into a vibration, albeit retaining its rotational character. In DyLu,N@C\(_{60}\) and DySc,N@C\(_{60}\), the frequencies of the corresponding cluster librations are predicted at 27 and 62 cm\(^{-1}\), respectively (Table 4). However, localized molecule vibrations at the \(\Gamma\) point do not transfer moment either, and hence their dispersions and interactions with lattice phonons should be studied further.

Consideration of the lattice phonons in EMF solids is also necessary because their frequency range is likely to overlap with low-frequency intramolecular vibrations. Unfortunately, experimental information on lattice phonons in EMFs is very limited. To the best of our knowledge, there has been only one study of monometallofullerenes by inelastic neutron scattering, which showed almost featureless VDOS in the low-frequency range\,[25] Far-IR\,[25, 26] as well as the aforementioned Raman studies proved the presence of some low-frequency modes in EMFs, but metal-based intramolecular vibrations could not be distinguished from the lattice modes, and, in addition, the selection rules limit the optical activity only to the \(\Gamma\) point. Lattice vibrations of empty fullerenes, and especially C\(_{60}\) are much better studied. According to inelastic neutron scattering and ab initio computations, the lattice phonons of C\(_{60}\) exhibit rather strong dispersion and cover the range up to 60–70 cm\(^{-1}\), whereas optical spectroscopic studies revealed the bands of librational modes at 7 and 18 cm\(^{-1}\), and those of translational modes at 28, 41, and 59 cm\(^{-1}\).\,[28] Thus, we can tentatively suggest that the frequency ranges of lattice modes and the intramolecular cluster vibrations of EMFs do overlap, which may lead to significant mode mixing.

In the absence of experimental information, we performed computational modeling of the lattice phonons of M,N@C\(_{60}\) to analyze their possible interaction with intramolecular vibrations. Complete calculations of the phonons of fullerene crystals at the DFT level are hardly feasible at this moment, and therefore simulations were performed by using the less demanding density-functional based tight-binding (DFTB) approach.\,[29] The pair-atomic interaction potentials available for Sc,N@C\(_{60}\) showed a reasonable prediction of the molecular geometry and vibrations.\,[10] As potentials for other lanthanide atoms of interest are not known, we used the Sc,N@C\(_{60}\) model to determine the Hessian and then computed the dispersion
spectra and vibrational eigenvectors for different lanthanide-containing NCFs by using a proper mass correction in a dynamic matrix.

The model lattice of $\text{M}_3\text{N}@\text{C}_{80}$ molecules was simulated with face-centered cubic (FCC) packing with the optimized unit cell parameter $a$ of 15.5 Å and the distance between the centers of fullerene molecules of 11 Å (see Figure S11 in the Supporting Information). The vibrational spectra computed for isolated DySc$_2$N@C$_{80}$ and DyLu$_2$N@C$_{80}$ molecules are compared in Figure 8 with those of the crystal phase, and dispersion of the phonons along the high-symmetry line $\Gamma$-X can be seen. In both cases, the DFTB-computed spectra of the isolated molecules start above 50 cm$^{-1}$, which is just on the borderline for the acoustic bands of the crystals. The molecular modes away from the frequency range of the acoustic modes show noticeable Davydov splitting but negligible $k$-dispersion. But the local modes close to the acoustic bands intertwine with the dispersed lattice bands giving rise to a dense phonon structure starting from zero frequency on. Three clear acoustic modes are perturbed by a set of what appear to be local modes with fluid character across $k$-space. To follow the possible mode mixing in $k$-space, we chose $\Gamma$-point vibrational eigenvectors as a basis space, in which eigenvectors computed at different $k$ values were projected. The results of this projection analysis for one pure acoustic mode and for one with cluster libration character are presented in Figure 8a,b using a color code (blue for the acoustic and red for the libration) and a scatter plot, for which the size of the dots is proportional to the magnitude of the projection (Figure 8). In this representation we can see and quantify how strongly the acoustic mode couples to more localized modes as a function of $k$. This redistribution is also reflected in the projected DOS in Figure 8, with the acoustic band showing a steady increase of cluster contributions as the energy increases. This model computation clearly shows that substantial mode mixing indeed takes place in $k$-space. However, the 3D model is challenging to grasp due to extensive mixing and a high density of states. For illustrative purposes, we simplified the model to one dimension and considered a linear chain of $\text{M}_3\text{N}@\text{C}_{80}$ molecules with $a = 10.75$ Å (see Figure S11 in the Supporting Information).

Figure 8c–e shows dispersion relationships computed for 1D chains of three NCF molecules, namely Sc$_2$N@C$_{80}$, DySc$_2$N@C$_{80}$, and DyLu$_2$N@C$_{80}$. The spectra contain only one prominent acoustic band with a large dispersion of 50 cm$^{-1}$ and two bands with a smaller dispersion of 10 cm$^{-1}$ produced by on-site rotational degrees of freedom. In 1D Sc$_2$N@C$_{80}$, the lattice and cluster modes are high in frequency, the cluster libration band is flat, and a weak mixing with the acoustic mode can be detected only around the X symmetry point. With the increase in mass in DySc$_2$N@C$_{80}$, the cluster-based frequencies decrease faster than those of the lattice phonons, and the degree of mixing increases as manifested in the considerable dispersion of the intramolecular mode propelled by the acoustic band. Furthermore, the local and acoustic mode frequencies are even closer in 1D DyLu$_2$N@C$_{80}$ and therefore the mode mixing is much more pronounced.

The mixing of cluster libration modes with the lattice phonons outside of the $\Gamma$ point has profound consequences for momentum transfer. These results show how in the first instance the librations, still being local modes, can assist the total conservation of momentum during spin flip and would be able to redistribute the excess angular momentum onto the lattice at some $k$ values. Mixing of the cluster rotations with the lattice phonons thus facilitates spin reversal by interaction of the local and lattice modes in $k$-space. Based on this conclusion, we tentatively suggest that the Arrhenius behavior with a $U_{\text{eff}}$ of 24 K (17 cm$^{-1}$) observed in the temperature dependence of magnetization relaxation times of DySc$_2$N@C$_{80}$ and DyLu$_2$N@C$_{80}$ (Figure 4a) may be caused by spin reversal assisted by the rotational modes of the $\text{M}_3\text{N}$ cluster. $U_{\text{eff}}$ in this case should correspond not to the frequency in the $\Gamma$ point, but rather to the frequencies in the high density of states of the phonons with strong mixed character. In addition, the difference in the masses of the dysprosium-scandium and dysprosium-lutetium nitride clusters will result in a different degree of mixing between the cluster librations and lattice phonons, thus leading to a different efficiency of the spin–lattice relaxation.

**Conclusions**

In this work we have studied the magnetic properties of the nitride clusterfullerenes DyLu$_2$N@C$_{80}$ and DyLu$_2$N@C$_{80}$ and ana...
alyzed how substitution of scandium by lutetium in the mixed-metal clusterfullerenes DyM₆N@C₈₀ and DyMN@C₈₀ affects the single-molecule magnetism thereof. DyLuN@C₈₀ and DyLuN@C₈₀ have been found to be SMMs with blocking temperatures of 9.5 and 8 K, respectively. DyLuN@C₈₀ exhibits a higher blocking temperature, longer relaxation times, and broader hysteresis than the dysprosium-scandum analogue DyScN@C₈₀. Both DyScN@C₈₀ and DyLuN@C₈₀ feature zero-field QTM, and when the QTM is quenched in a finite field of 0.2 T, the magnetization relaxation times of both compounds show Arrhenius behavior with an effective barrier of 24 K.

DyLuN@C₈₀ and DyScN@C₈₀ have identical blocking temperatures, but show different temperature and field dependence of the relaxation times. In particular, a direct relaxation mechanism with enhanced field dependence is observed for DyLuN@C₈₀ below 5 K, whereas the relaxation times of DyScN@C₈₀ remain independent of field until the field exceeds 0.4 T. The magnetization relaxation times of DyLuN@C₈₀ show Arrhenius behavior with an effective barrier of 4.3 K, which has been assigned to the energy of the excited state with antiferromagnetic coupling of the dysprosium moments, ΔE₁₈₇₆₁₆₆. In DyScN@C₈₀, the energy of this state is around two-fold higher, at 8.0 K. Because DyLuN@C₈₀ and DyScN@C₈₀ have almost identical energies of intramolecular dipolar interactions of 4.7 K, the considerable difference in their ΔE₁₈₇₆₁₆₆ values is attributed to the strong variation in the exchange coupling when scandium is substituted by lutetium. Essentially, the magnetic moments in DyLuN@C₈₀ show only dipolar interactions and their exchange coupling vanishes.

To aid the understanding of possible spin–phonon energy exchange, the low-frequency vibrational spectra of the dysprosium-lutetium and dysprosium-scandum NCFs were analyzed experimentally and with the help of DFT calculations. Enclosing the M₆N cluster inside the fullerene cage transforms its rotational degrees of freedom into molecular vibrations, which retain rotational character and are dubbed as librations of the cluster. The low frequencies of these modes lead to overlap with the frequency range of the lattice phonons. Furthermore, projection analysis also revealed the strong mixing of the local cluster librations with acoustic phonons of the fullerene lattice in the k-space away from the Γ point. As a result, these modes are predicted to facilitate the relaxation of magnetization by helping to conserve momentum during the spin reversal. Thus, the results of our study emphasize that the mixing of local and lattice modes in k-space may be an important mechanism of the spin–lattice relaxation and should be considered for other molecular magnets.

**Experimental Section**

Powder samples of fullerenes for magnetometry studies were prepared by drop-casting from toluene or C₅₋₇ solutions. The magnetic properties were studied with a Quantum Design MPMS3 Vibrating Sample Magnetometer (VSM). Modeling of the magnetization curves and the spin Hamiltonian solution was accomplished with the PHI program.[31]

Raman spectra were recorded at 77 K with a T 64000 triple spectrometer (Jobin Yvon) using an excitation wavelength of λₑ = 514 nm (Ar⁺ laser) or λₑ = 647 nm (Kr⁺ laser). The samples for Raman measurements were drop-cast onto single-crystal KBr disks. DFT calculations on isolated M₆N@C₈₀ molecules were performed at the PBE-D level with a plane-wave basis set and the corresponding projector augmented-wave potentials, treating 4f electrons as a part of the core as implemented in the VASP 5.0 package.[32] These calculations employed a cubic unit cell with a lattice parameter of 25 Å and the atomic cut-off energy was set to 400 eV. The precision was set to be "accurate" with real-space projector operators optimized down to 10⁻⁶ eV per atom. The Γ-point Hessian matrix and then the vibration frequencies (or Γ-point phonons) were determined by using density functional perturbation theory as implemented in VASP.

The ab initio energies and wave functions of LF (ligand field) multiplets for the dysprosium-lutetium NCF molecules were calculated at the CASSCF/SO-RASSI level of theory with atomic natural orbital extended relativistic basis set (ANO-RCC)[33] of the valence double-zeta (VDZ) quality using the quantum chemistry package MOLCAS 8.0.[34] The active space of the CASSCF calculations included the 4f shell, that is, 11 active electrons and 7 active orbitals. All 21 sextet states and 108 quartets and only 100 doublets were included in the state-averaged CASSCF procedure and further used in the RASSI procedure with a spin-orbit Hamiltonian. The single-configuration properties and LF parameters were calculated on the basis of the ab initio data with the use of a SINGLE ANISO module.[35]

Phonon spectra modeling and analysis were performed with in-house Python scripts based on ASE libraries and with the improved version of some functions.[36] The DFTB⁺ software[37] was used as a force derivation code in the Calculator class of ASE. The finite differences method was used in phonon calculations with atomic displacement of 0.03 Å along each Cartesian axis. Prior to phonon calculations, 1D and 3D systems (see Figure S11 in the Supporting Information) were optimized with a quasi-Newton algorithm down to 0.0002 eV Å⁻¹ with 2 k-point sampling in each periodic direction. In all calculations, the non-charge-self-consistent model was employed with the Slater–Koster parameters developed for similar types of system.[38]

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