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Hyperfine collisional excitation of ammonia by molecular hydrogen

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ABSTRACT

Ammonia is one of the most widely observed molecules in space, and many observations are able to resolve the hyperfine structure due to the electric quadrupole moment of the nitrogen nucleus. The observed spectra often display anomalies in the satellite components of the lines, which indicate substantial deviations from the local thermodynamic equilibrium. The interpretation of the spectra thus requires the knowledge of the rate coefficients for the hyperfine excitation of NH₃ induced by collisions with H₂ molecules, the dominant collider in the cold interstellar medium. In this paper we present the first such calculations using a recoupling approach. The rate coefficients are obtained for all hyperfine levels within rotation-inversion levels up to j = 4 and temperatures up to 100 K by means of quantum scattering close-coupling calculations on an accurate, five-dimensional, potential energy surface. We show that the rate coefficients depart significantly from those obtained with the statistical approach and that they do not conform to any simple propensity rules. Finally, we perform radiative transfer calculations to illustrate the impact of our new rate coefficients by modelling the hyperfine line intensities of the inversion transition in ground state *para*-NH₃ ($j_k = 1_1$) and of the rotational transition $1_0 \rightarrow 0_0$ in *ortho*-NH₃.

Key words: radiative transfer - molecular data - molecular processes - scattering - ISM: molecules - ISM: abundances

1 INTRODUCTION

Since its first detection in the interstellar medium (Cheung et al. 1968), ammonia (NH₃) has often been used to probe the physical properties of molecular clouds due to the favourable structure of its energy levels and its high abundance in a variety of astrophysical sources. The rotational levels of NH_3 are denoted as j_k (where j is the angular momentum and k its projection on the molecular axis), and the presence of metastable states for i = k that can only decay through molecular collisions makes NH₃ an excellent probe of the local temperature. NH3 exists in two forms, namely ortho-NH3 (for k = 3n, *n* integer) and *para*-NH₃ (for $k = 3n \pm 1$) that cannot be interconverted through radiative or inelastic collisional processes. In addition, the inversion (or umbrella) motion causes a splitting of each j_k level into a doublet (with the exception of k = 0 due to spin statistics) whose components, denoted as j_k^+ and j_k^- , are separated by 0.79 cm⁻¹ (23.68 GHz). NH₃ can thus be detected in different frequency ranges corresponding to pure inversion, rotational, or vibrational transitions.

In most astrophysical observations, the hyperfine structure due to the electric quadrupole moment of nitrogen is resolved. N nuclei possess a spin of I = 1, so that each rotation-inversion level is further split into three hyperfine components F (with the exception of ground state *ortho*-NH₃), with a separation of the order of a MHz. A rotational line is thus split into five lines: the main line ($\Delta F = 0$), as well as two inner and two outer satellite lines characterized by $\Delta F = \pm 1$. The lines are further split by the spins of the H nuclei, but this interaction is much weaker and will be neglected here.

The relative intensity of hyperfine lines can be used to derive opacities if the various hyperfine components have the same excitation temperature and if the opacity is not too large (Barrett et al. 1977; Ho & Townes 1983). However, NH₃ hyperfine-resolved observations commonly display anomalies in the spectra, where the intensities within a pair of satellite lines are not equal. This indicates different excitation temperatures, meaning that the satellite components do not obey the local thermodynamic equilibrium (LTE) assumption in the interstellar medium. Such anomalies were first observed by Matsakis et al. (1977) in source DR21 and subsequently by various authors (see e.g., Matsakis et al. (1980); Gaume et al. (1996); Rathborne et al. (2008); Rosolowsky et al. (2008); Camarata et al. (2015); Caselli et al. (2017); Zhou et al. (2020)) in diverse sources. If the hyperfine levels are not populated thermally, the interpretation of observations requires radiative transfer models to account for non-LTE effects. These occur mainly because of collisional excitation by H₂ molecules, and a knowledge of accurate rate coefficients for the hyperfine excitation of NH₃ by H₂ is thus necessary. In addition, modelling the strength of the hyperfine anomalies can be used to deduce the local H₂ density.

Models to interpret anomalies were developed by Stutzki & Winnewisser (1985a,b) based on rate coefficients calculated for NH_3 -He collisions as a proxy for NH_3 -H₂ collisions. Beside this work, Chen et al. (1998) also computed hyperfine-resolved rate coefficients for NH_3 -He collisions. To this day there are thus no accurate collisional data for the hyperfine excitation of NH_3 by H_2 molecules.

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Various theoretical techniques exist to compute such rate coefficients. Among these, the simplest is the statistical approach, which assumes that the rate coefficients for hyperfine excitation are proportional to the hyperfine degeneracy of the final state. In this case, rate coefficients for pure rotational transitions, which are available for molecules such as NH₃, are the only ingredient needed to obtain hyperfine-resolved rate coefficients. This method was used to interpret the observations of Caselli et al. (2017) for the $1_0 - 0_0$ transition in the pre-stellar core L1544. The infinite-order-sudden limit approximation, used by Stutzki & Winnewisser (1985a), ignores the rotational energy spacing and allows again to obtain the hyperfineresolved rate coefficients on the basis of the pure rotational ones. Given the nature of its assumptions, this approximation is however not expected to be valid at low temperatures, i.e. at temperatures close or lower than the rotational spacings. Finally, the method used in the present work is the recoupling approach, which was also adopted by Chen et al. (1998) but for NH₃-He collisions and combined with the coupled-states approximation¹. The only approximation in our version of the recoupling approach is the assumption that the hyperfine levels are degenerate, i.e. their splitting is negligible compared to the temperature (Alexander & Dagdigian 1985). While computationally more demanding, it is also expected to be more accurate than the statistical or sudden methods. A summary and comparison of these different approaches can be found in the paper by Faure & Lique (2012) for the linear molecules CN and HCN.

In this work we present rate coefficients for the hyperfine excitation of *ortho*- and *para*-NH₃ in collisions with *para*-H₂ for all levels up to j = 4 and for temperatures up to 100 K. While the rotationinversion collisional excitation of NH₃ by H₂ has been widely studied theoretically over the past decades (Danby et al. (1987); Offer & Flower (1990); Maret et al. (2009); Bouhafs et al. (2017); Ma et al. (2015); Demes et al. (2023)) and experimentally with crossed beams (Schleipen & ter Meulen 1991; Tkac et al. 2015; Gao et al. 2019), to the best of our knowledge this represents the first effort to provide hyperfine collisional excitation rate coefficients of NH₃ (or any nonlinear molecule) by H₂ with the accurate recoupling approach.

The paper is structured as follows: in Section 2 we discuss the theoretical methods employed. In Section 3 we discuss the features of the hyperfine-resolved rate coefficients, and in Section 4 we present radiative transfer calculations to investigate the impact of these new data on the modelling of ammonia spectra. Finally, we summarize our findings in Section 5.

2 THEORY

The scattering calculations were performed in two steps. First, we computed the scattering matrix for rotation-inversion transitions by means of the fully quantum-mechanical close-coupling (CC) method. This method has been used to obtain accurate cross sections for rotational excitation of NH₃ induced by collisions with H₂ by several authors (see *e.g.*, Danby et al. (1987); Rist et al. (1993)) and we refer to these papers for further methodological details. Similarly to recent theoretical works (Ma et al. 2015; Bouhafs et al. 2017; Demes et al. 2023), we employ the accurate five-dimensional rigid-rotor potential energy surface of Maret et al. (2009), which was constructed by fitting *ab initio* energies obtained with the coupled cluster method with single, double, and perturbative triple excitations.

The cross sections for rotation-inversion transitions can be obtained from the *T*-matrix elements $T_{j_1kvj_2j_{12}l,j'_1k'v'j'_2j'_{12}l'}^{I}$, where *v* denoted the inversion level, *J* is the quantum number associated with the total angular momentum of the collision complex without nuclear spin ($J = j_{12} + L$), $j_{12} = j_1 + j_2$, where j_1 and j_2 are the angular momenta of NH₃ and H₂, respectively, and *L* is the relative orbital angular momentum of the collision. We do not discuss here the properties of the cross sections for rotation-inversion transitions as these are identical to those reported recently by Bouhafs et al. (2017) and Demes et al. (2023).

In a second step, the recoupling approach is used to compute hyperfine-resolved cross sections. When the nuclear spin of the N atom (I = 1) is included, each level of NH₃ is split into three components, and the total angular momentum J_t is obtained by the coupling $J_t = J + I$. However, the hyperfine splittings are of the order of the MHz (i.e. about $10^{-5} - 10^{-4}$ K), which is much smaller than the rotational spacings and collisional energies considered here. We thus assume that the hyperfine levels of NH₃ are degenerate. In this case the nuclear spin wavefunctions can be decoupled from the rotational wavefunctions and, as shown by Offer et al. (1994), the angular momenta can also be recoupled as $F = j_1 + I$, $j_R = j_2 + L$, $J_t = j_R + F$. The *T*-matrix elements including nuclear spin, T^{J_t} , can then be obtained from the spin-free *T*-matrix elements T^J as

$$T_{j_{1}\nu F j_{2}j_{R}l, j_{1}'\nu' F' j_{2}' j_{R}'l'}^{J_{t}} = \sum_{J, j_{12}, j_{12}'} (-1)^{j_{R}+j_{R}'+l+l'+j_{2}+j_{2}'} \\ ([F][F'][j_{12}][j_{12}][j_{R}][j_{R}']]^{1/2}[J] \left\{ \begin{array}{cc} j_{1} & j_{2} & j_{12} \\ l & J & j_{R} \end{array} \right\} \left\{ \begin{array}{cc} j_{1}' & j_{2}' & j_{12}' \\ l' & J & j_{R}' \end{array} \right\} \\ \left\{ \begin{array}{cc} j_{R} & j_{1} & J \\ I & J_{t} & F \end{array} \right\} \left\{ \begin{array}{cc} j_{R}' & j_{1}' & J \\ I & J_{t} & F' \end{array} \right\} T_{j_{1}k\nu j_{2}j_{12}l, j_{1}'k'\nu' j_{2}'j_{12}'l'}^{J}$$
(1)

where $\{\cdots\}$ denotes a 6-j symbol and [x] = 2x + 1.

The state-to-state hyperfine cross sections for NH_3 - H_2 can be then computed from the *T*-matrix according to the expression (Offer et al. 1994):

$$\sigma_{j_1kvFj_2,j'_1k'v'F'j'_2} = \frac{\pi}{k_i^2(2F+1)(2j_2+1)} \sum_{J_T} (2J_T+1) \sum_{\substack{II'j_Rj'_R}} |T_{j_1kvFj_2j_Rl,j'_1k'v'F'j'_2j'_Rl}|^2$$
(2)

The scattering calculations were carried out with the Hibridon program (Alexander et al. 2023) for total energies between 0 and 600 cm⁻¹ on a grid of 700 energies for *ortho*-NH₃ and 660 energies for para-NH₃. These calculations were performed independently for ortho-NH₃ and para-NH₃ as ortho-para conversion is forbidden in inelastic collisions. The energy step was 0.5 cm^{-1} at the lowest energies and increased gradually up to 4 cm^{-1} at the highest energies. We used the log-derivative propagator in the short range, and the Airy propagator in the long range $(R > 25a_0)$. Transitions between levels up to $j_1 = 6$ with $j_2 = 0$ were converged with a rotational basis including all levels up to $j_1 = 8$ and $j_2 = 0, 2$. The rotational constants of NH₃ were taken as $A = B = 9.9402 \text{ cm}^{-1}$, C = 6.3044 cm^{-1} with an inversion splitting of 0.7903 cm^{-1} , while the rotational constant of H₂ was taken as 59.3801 cm⁻¹ as in Bouhafs et al. (2017). Tests were performed to ensure convergence of the cross sections with respect to J_t in Eq. (2) at all energies, as well with respect to the parameters of the radial grid.

Rate coefficients were then calculated by averaging cross sections for initial and final states *i* and *f*, $\sigma_{if}(E_c)$, over the Maxwell-

¹ We note that the results of Chen et al. (1998) are questionable and should not be used because some of their rate coefficients are unphysically large (up to 10^{-7} cm³s⁻¹).

$$k_{if}(T) = \left(\frac{8}{\pi\mu\beta}\right)^{\frac{1}{2}}\beta^2 \int_0^\infty E_c \sigma_{if}(E_c) e^{-\beta E_c} dE_c \tag{3}$$

Where $\beta = \frac{1}{k_B T}$ and k_B , T and μ denote the Boltzmann constant, the kinetic temperature and the NH₃-H₂ reduced mass, respectively.

3 HYPERFINE RATE COEFFICIENTS

Hyperfine excitation rate coefficients were calculated for temperatures up to 100 K and for transitions involving all levels up to $j_1 = 4$ for both *ortho*-NH₃ and *para*-NH₃, corresponding to a total of 625 transitions for *ortho*-NH₃ and 2304 transitions for *para*-NH₃. Only excitation due to p-H₂ ($j_2 = 0$) is considered here since excited H₂ levels are not significantly populated at the temperatures of molecular clouds where the hyperfine structure is resolved.

Rate coefficients for hyperfine transitions from the rotational state 2_1^+ down to level 1_1^- are presented in Fig. 1 as a function of temperature up to 100 K. The rate coefficients are seen to be weakly dependent on *T*, a trend that follows directly from the behaviour of the rate coefficients for pure rotational transitions. On the same figure, we also present for comparison the statistical hyperfine rate coefficients. The statistical approach, also known as the M_J randomizing limit (Alexander & Dagdigian 1985), is a commonly used approximation to compute hyperfine-resolved excitation rate coefficients for hyperfine excitation are directly proportional to the degeneracy of the final hyperfine state and independent of the initial hyperfine state. For NH₃-H₂ the statistical rate coefficients are expressed as

$$k_{jk\nu F,j'k'\nu'F'}^{\text{stat}} = \frac{2F'+1}{(2I+1)(2j'+1)}k_{jk\nu,j'k'\nu'}$$
(4)

where $k_{jk\nu,j'k'\nu'}$ are the nuclear spin-free rate coefficients calculated with the CC method and I = 1. To simplify the notation, in the remainder of the paper, we use *j* to denote the rotational quantum number of NH₃ instead of j_1 without ambiguity since $j_2 = 0$.

As can be seen from Fig. 1, for all three final hyperfine states of the transition $2_1^+F \rightarrow 1_1^-F'$ the statistical rate coefficients fall between the recoupling values, but a strong dependence on F' can be observed and large deviations from the M_J randomizing limit are seen to occur. It is thus clear that the statistical method cannot be used to obtain accurate hyperfine rate coefficients in the case of NH₃-H₂ collisions.

From Fig. 1, it can be seen that the known propensity rule $\Delta F = \Delta j$ (Alexander & Dagdigian 1985) holds for the transitions within $2_1^+ \rightarrow 1_1^-$. However, this is not a general trend. In Table 1 we report the rate coefficients for transitions from levels 2_1^+F down to levels 1_1^+F and 1_1^-F at a fixed temperature of 30 K. For the case of 2_1^+F to 1_1^+F transitions (as well as for a large number of transitions involving higher lying rotational levels), the $\Delta F = \Delta j$ propensity rule is not respected. In addition, Table 1 also shows that the largest rate coefficients are not necessarily those with the largest F', as would be expected from a statistical behavior.

Of interest are also the quasi-elastic rate coefficients, i.e. rate coefficients for hyperfine excitation within a single rotational level. We present in Table 2 the quasi-elastic rate coefficients $j_k vF \rightarrow j_k vF'$ at 25 K for the first three rotational states of *para*-NH₃, and for the antisymmetric inversion level (v = v' = -). The corresponding F - F' transitions involving the symmetric inversion levels (v = v' = +) are similar although not identical, with differences of up to a few



Figure 1. Rate coefficients for the transitions $2_1^+ F \rightarrow 1_1^- F'$ of NH₃ induced by collisions with H₂ as a function of temperature. Panel (a), F' = 0; panel (b), F' = 1; panel (c), F' = 2, and comparison to the statistical rate coefficients given by Eq. (4).

percent (not shown). In addition, we also report in the same table the rate coefficients corresponding to a change of inversion level $(- \rightarrow +;$ the reverse rate coefficients can be obtained by detailed balance). It should be noted that those are not strictly quasi-elastic, since the $- \rightarrow +$ pure inversion transition correspond to an energy transfer of 0.79 cm^{-1} . It can be noticed that the rate coefficients strongly depend on the final inversion level, with differences that reach a factor 5 between v' = - and v' = +. Interestingly, the rate coefficients for inversion-conserving transitions $j_k^- F \rightarrow j_k^- F'$, which would be elastic if the nuclear spin were neglected, are in many cases smaller than those of the corresponding inversion-changing transitions $j_{k}^{-}F \rightarrow j_{k}^{+}F'$. Finally, the last column of Table 2 compares our values to the results of Stutzki & Winnewisser (1985a), which are independent of the inversion level, and show important discrepancies of up to an order of magnitude for some quasi-elastic transitions. It is worth emphasizing that those results were obtained using a different method, namely the infinite order sudden (IOS) approximation,

Table 1. Rate coefficients for the transitions $2_1^+F \rightarrow 1_1^+F'$ and $2_1^+F \rightarrow 1_1^-F'$ of NH₃ induced by H₂ at a temperature of 30 K. Brackets indicate powers of 10.

$j_k v - j'_{k'} v'$		F' = 0	F' = 1	F' = 2
$2_1^+ - 1_1^+$	F = 1	5.16[-13]	1.98[-11]	7.27[-12]
	F = 2	8.62[-12]	3.11[-12]	1.58[-11]
	F = 3	1.85[-13]	8.99[-12]	1.84[-11]
$2_1^+ - 1_1^-$	F = 1	3.31[-11]	2.57[-11]	4.97[-12]
	F = 2	3.96[-13]	4.58[-11]	1.76[-11]
	F = 3	7.19[-13]	1.83[-12]	6.12[-11]

Table 2. Quasi-elastic rate coefficients for the excitation $j_k vF \rightarrow j_k v'F'$ for rotational levels 1_1 , 2_2 , and 2_1 of NH₃ by H₂ at 25 K. Comparison with the data of Stutzki & Winnewisser (1985a) (last column). Brackets indicate powers of 10.

j _k	F - F'	<i>v</i> = -, <i>v</i> ′ = -	<i>v</i> = -, <i>v</i> ′ = +	Ref.
1_{1}	0 - 2	2.43[-11]	1.22[-11]	2.49[-11]
1_{1}	0 - 1	1.83[-11]	8.96[-11]	8.66[-11]
1_{1}	2 - 1	1.55[-11]	2.79[-11]	3.28[-11]
22	1 – 3	2.58[-12]	1.42[-12]	1.92[-11]
2_{2}	1 – 2	1.44[-11]	3.05[-11]	7.02[-11]
22	3 - 2	7.42[-12]	1.41[-11]	3.88[-11]
2_{1}	2 – 3	5.31[-12]	8.29[-12]	5.26[-11]
2_{1}	2 - 1	4.13[-12]	6.84[-12]	4.14[-11]
2_{1}	3 – 1	7.93[-13]	9.17[-13]	7.41[-12]

which is not expected to be accurate at low temperatures. In addition, Stutzki & Winnewisser (1985a) used a potential for NH_3 -He instead of NH_3 -H₂, which leads to an additional error. Finally, we note that our quasi-elastic rate coefficients are of the same order of magnitude as those for inelastic transitions.

From their IOS calculations, Stutzki & Winnewisser (1985a) compute relative factors, defined as the ratio of hyperfine rate coefficients divided by the rotational excitation rate coefficient,

$$g_{FF'} = \frac{k_{jk\nu F, j'k'\nu'F'}}{k_{jk\nu, j'k'\nu'}}$$
(5)

While the absolute IOS rate coefficients are not expected to be accurate at low temperature, the method should provide relative factors that are better than the statistical ones. These relative factors obtained with the recoupling approach are presented in Table 3 and compared to the IOS results of Stutzki & Winnewisser (1985a) for the $1_1^+ - 2_1^+$ and $1_1^+ - 2_1^-$ excitation transitions. We observe that in most cases the IOS method does predict the correct propensities, and performs better than the statistical approach, as expected. On the other hand, it is clear that IOS is not sufficient to obtain accurate rate coefficients, since discrepancies of up to 50% are seen for the dominant rate coefficients and up to a factor 10 for the smallest rate coefficients. Moreover, both the IOS and recoupling relative factors are almost independent of the temperature in the range explored here (see Fig. 1), which shows that the discrepancies discussed above remain up to temperatures of at least 100 K.

4 RADIATIVE TRANSFER CALCULATIONS

In order to investigate the impact of the hyperfine rate coefficients on the non-LTE modelling of NH_3 spectra, we present below simple

Table 3. Relative factors (Eq. (5)) for the transitions $1_1^+F \rightarrow 2_1^+F'$ and $1_1^+F \rightarrow 2_1^-F'$ of NH₃ induced by H₂ at a temperature of 25 K. The values in parentheses are from Stutzki & Winnewisser (1985a).

		F' = 1	F' = 2	F' = 3
$1_1^+F - 2_1^+F'$	F = 0	0.03 (0.08)	0.95 (0.65)	0.02 (0.27)
	F = 1	0.43 (0.31)	0.11 (0.20)	0.46 (0.48)
	F = 2	0.09 (0.16)	0.35 (0.35)	0.56 (0.50)
$1^+_1F - 2^1F'$	F = 0	0.93 (0.57)	0.02 (0.28)	0.05 (0.15)
	F = 1	0.24 (0.27)	0.72 (0.50)	0.04 (0.23)
	F = 2	0.03 (0.09)	0.17 (0.24)	0.81 (0.67)

radiative transfer calculations using two different sets of hyperfineresolved collisional data: (1) the precise set of *recoupling* rate coefficients and (2) the approximate set of *statistical* rate coefficients, as defined above in Eq. (4).

Radiative transfer calculations were performed with the non-LTE RADEX program (van der Tak et al. 2007) using the large velocity gradient (LVG) formalism for an expanding sphere. The code was employed to compute the radiation temperature T_R of low-lying hyperfine transitions in both para- and ortho-NH3. The hyperfine energy levels and radiative rates were taken from the CDMS catalog (Müller et al. 2005). The kinetic temperature was fixed at 10 K, which is typical of cold molecular clouds, while the hydrogen density was varied in the range $10^2 - 10^9$ cm⁻³. The *para*-NH₃ (or *ortho*-NH₃) column densities were selected in the range $10^{12} - 10^{15}$ cm⁻² with a line width (full width at half maximum) of 1 km s⁻¹. No radiation field was considered except the cosmic microwave background (CMB) at 2.73 K. We have also checked with the MOLPOP code (Asensio Ramos & Elitzur 2018) that line overlap effects, which are ignored by RADEX, are mostly negligible with the above parameters, allowing us to focus on the impact of the collisional data.

We first consider the inversion transition $1_1^- \rightarrow 1_1^+$ of para-NH₃ at 23.68 GHz. This transition has been widely observed in the interstellar medium since its first detection by Cheung et al. (1968). Anomalies in the hyperfine intensities of this line were reported in the 1970s (Matsakis et al. 1977) and these anomalies are now known to be very common in star forming regions. In the anomalous spectra, the intensities of the outer satellite lines are not equal: the $F = 0 \rightarrow 1$ component is stronger and the $F = 1 \rightarrow 0$ component is weaker than expected. Radiative transfer models have shown that these anomalies can be reproduced by non-LTE effects induced by photon trapping of selected hyperfine transitions in the rotational transition $2_1^{+/-} \rightarrow 1_1^{-/+}$, as explained by Stutzki & Winnewisser (1985b) (see in particular their Fig. 2). Systematic motions, such as expansion and collapse, are also expected to play a role and the relative contribution of these effects remains a matter of debate (see Zhou et al. 2020, and references therein). In any case, because non-LTE populations depend on the competition between the radiative and collisional rates, the accuracy of hyperfine selective collisional rate coefficients should be critical for a correct interpretation of the anomalies in the $1_1^- \rightarrow 1_1^+$ inversion transition.

In Fig. 2, we have plotted the intensity ratio of the outer hyperfine satellite line $F = 0 \rightarrow 1$ relative to the strongest component $F = 2 \rightarrow 2$, for the two sets of collisional rates (recoupling and statistical). At low hydrogen density, the ratio is found to increase above the "natural" intensity ratio (0.266) as soon as the column density exceeds ~ 10^{14} cm⁻². The ratios then increase with increasing density and plateaus are reached when $n(H_2)$ is larger than about 10^5 cm⁻³, where LTE conditions are fullfilled. For column densi-





Figure 2. The intensity ratio of the hyperfine satellite line $F = 0 \rightarrow 1$ of the *para*-NH₃ $j_k = 1_1$ inversion transition relative to the strongest component $F = 2 \rightarrow 2$ as function of the hydrogen density for a set of column densities. The kinetic temperature is fixed at 10 K and the line width at 1 km s⁻¹.

ties above ~ 10^{14} cm⁻², we can notice the impact of the collisional dataset: the intensity ratios are decreased by up to 20% when the statistical dataset is employed. This demonstrates the importance of the hyperfine propensity rules when the total opacity of the line is larger than ~ 1. A similar result was actually observed by Stutzki & Winnewisser (1985a) using hyperfine collision rates computed with the IOS approximation and using Helium instead of H₂ as the collider. As concluded by these authors, a precise knowledge of the hyperfine-resolved rate coefficients is thus required for a reliable modelling of observational data.

A second example is the ortho-NH3 ground-state rotational transition $1_0^+ \rightarrow 0_0^-$ at 572.5 GHz. The electric quadrupole hyperfine structure of this line was resolved for the first time in space thanks to Herschel observations towards the cold pre-stellar core L1544 (Caselli et al. 2017). The three hyperfine components were found to be heavily self-absorbed and a sophisticated radiative transfer model was employed to reproduce the observed profile and to deduce the NH3 radial abundance in the core. In that work, hyperfine selective collisional data were generated from the rotational rate coefficients of Bouhafs et al. (2017) by applying the statistical approach to the electric quadrupole and magnetic hyperfine splittings. In Fig. 3, we have plotted the intensity ratio of the hyperfine component $F = 1 \rightarrow 1$ relative to the strongest component $F = 2 \rightarrow 1$, again for the two sets of collisional rates (recoupling and statistical). We note that our statistical set is identical to that used by Caselli et al. (2017) except that the magnetic splittings (< 100 kHz) are neglected here. At low hydrogen density, the ratio is found to lie slightly below the natural intensity ratio (0.6) for column density above ~ 10^{12} cm⁻². When the H₂ density exceeds ~ 10^5 cm⁻³, however, the intensity ratios rise above 0.6 and the LTE plateaus are reached at H₂ densities larger than $\sim 10^8$ cm⁻³, as expected from the large radiative rates $(A \sim 1.58 \times 10^{-3})$. The impact of the rate coefficients is smaller than in the case of the para-NH₃ transition (Fig. 2) with a maximum difference of $\sim 10\%$ in the intensity ratios. This is not really surprising since the recoupling and statistical rate coefficients for the

Figure 3. The intensity ratio of the hyperfine component $F = 1 \rightarrow 1$ of the *ortho*-NH₃ $j_k = 1_0 \rightarrow 0_0$ rotational transition relative to the strongest component $F = 2 \rightarrow 1$ as function of the hydrogen density for a set of column densities. The kinetic temperature is fixed at 10 K and the line width at 1 km s⁻¹.

hyperfine transitions $1_0^+F \rightarrow 0_0^-F' = 1$ are identical and equal to the pure rotational rate coefficients (by construction since the rotational 0_0^- level has a single hyperfine sub-level F = 1). Thus, only the rate coefficients corresponding to hyperfine transitions within the rotational level 1_0^+ , the so-called quasi-elastic rate coefficients, are different in the two datasets: they are set to zero in the statistical set while they are comparable to the pure rotational rate coefficient in the recoupling set. We can conclude that the quasi-elastic rate coefficients have a significant although moderate impact in the radiative transfer calculation.

Finally, we wish to stress that the above results are only illustrative: more sophisticated radiative transfer calculations are clearly needed to explore a much larger parameter space and to include line overlap as well as systematic motion effects. Preliminary calculations with MOLPOP show, for instance, that for kinetic temperatures larger than 10 K and column densities above 10^{15} cm⁻³, line overlap effects can increase the intensity ratio $F = 0 \rightarrow 1/F = 2 \rightarrow 2$ in the $1^-_1 - 1^+_1$ inversion transition by more than 50%. The combination of the present recoupling rate coefficients with one-dimensional radiative transfer codes able to handle hyperfine overlaps, such as ALICO (Magalhães et al. 2018), will be particularly valuable to revisit the NH₃ hyperfine anomalies and to more firmly establish the role of non-LTE effects.

5 CONCLUSION

We have presented a full set of rate coefficient for the hyperfine excitation of *ortho*- and *para*-NH₃ in collisions with *para*-H₂ for all levels of NH₃ up to j = 4 and for temperatures up to 100 K. The calculations were performed by means of the accurate recoupling technique, which relies on quantum-mechanical close-coupling scattering calculations. In general, the rate coefficients are weakly dependent on the temperature, but depend strongly on all quantum numbers of NH₃. We found no clear propensity rules, and a comparison with statistical rate coefficients (which are independent of the initial hyperfine level) showed important deviations from this approximation. In addition, we also found large differences with previously published results on NH_3 -He collisions.

Using a simple non-LTE model, we illustrated the difference between the accurate recoupling rate coefficients and the approximate statistical rate coefficients on the inversion transition in ground state *para*-NH₃ $(1_1^-F \rightarrow 1_1^+F')$ at 23.68 GHz and on the rotational transition $1_0^+F \rightarrow 0_0^-F'$ in *ortho*-NH₃. These simple calculations demonstrate that the two sets of rate coefficients lead to significant differences. Moreover, non-LTE effects are predicted for column density in excess of ~ 10^{12} cm⁻². An important extension of the present work will be to perform radiative transfer calculations using a model that can treat line overlaps which play a key role in NH₃ hyperfine anomalies. This would allow the investigation of non-LTE effects at higher temperatures, thus involving higher-lying rotational states of ammonia.

As a reminder, the present calculations only considered the hyperfine splitting due to the N nucleus but did not take into account the (much smaller) further hyperfine magnetic splitting due to the three protons, which is sometimes resolved in astronomical spectra (Rydbeck et al. 1977). We suggest that the corresponding rate coefficients can be obtained from those presented here using the statistical approximation.

Finally, it should also be noted that the recoupling calculations presented in the present work are accurate but computationally expensive. At the other end, statistical rate coefficients can be immediately obtained provided the rate coefficients for pure rotational excitation are available. An intermediate approach is the IOS method, which has been used with some success for the hyperfine excitation of linear molecules by H_2 (Goicoechea et al. 2022). Extending the IOS method and recoupling theory to asymmetric polyatomic molecules would be beneficial in order to generate sets of hyperfine-resolved rate coefficients for other molecules such as deuterated ammonia (NH₂D, NHD₂) or H₂CO.

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DATA AVAILABILITY

The data underlying this article are available as supplementary material to the article and the full set of rate coefficients will be made available through the EMAA database (https://emaa.osug.fr and https://dx.doi.org/10.17178/EMAA).

REFERENCES

- Alexander M. H., Dagdigian P. J., 1985, The Journal of Chemical Physics, 83, 2191
- Alexander M., Dagdigian P., Werner H.-J., Kłos J., Desrousseaux B., Raffy G., Lique F., 2023, Computer Physics Communications, 289, 108761 Asensio Ramos A., Elitzur M., 2018, A&A, 616, A131

- Barrett A. H., Ho P. T. P., Myers P. C., 1977, ApJ, 211, L39
- Bouhafs N., Rist C., Daniel F., Dumouchel F., Lique F., Wiesenfeld L., Faure A., 2017, Mon. Notices Royal Astron. Soc., 470, 2204
- Camarata M. A., Jackson J. M., Chambers E., 2015, The Astrophysical Journal, 806, 74
- Caselli P., et al., 2017, A&A, 603, L1
- Chen J., hui Zhang Y., Zeng Q., Pei C.-C., 1998, Journal of Physics B: Atomic, Molecular and Optical Physics, 31, 1259
- Cheung A. C., Rank D. M., Townes C. H., Thornton D. D., Welch W. J., 1968, Phys. Rev. Lett., 21, 1701
- Danby G., Flower D. R., Valiron P., Kochanski E., Kurdi L., Diercksen G. H. F., 1987, Journal of Physics B: Atomic and Molecular Physics, 20, 1039
- Demes S., Lique F., Loreau J., Faure A., 2023, Monthly Notices of the Royal Astronomical Society, 524, 2368
- Faure A., Lique F., 2012, Monthly Notices of the Royal Astronomical Society, 425, 740
- Gao Z., Loreau J., van der Avoird A., van de Meerakker S. Y. T., 2019, Phys. Chem. Chem. Phys., 21, 14033
- Gaume R. A., Wilson T. L., Johnston K. J., 1996, The Astrophysical Journal, 457, L47
- Goicoechea J. R., Lique F., Santa-Maria M. G., 2022, A&A, 658, A28
- Ho P. T. P., Townes C. H., 1983, ARA&A, 21, 239
- Ma Q., van der Avoird A., Loreau J., Alexander M., van de Meerakker S. Y. T., Dagdigian P. J., 2015, J. Chem. Phys., 143, 044312
- Magalhães V. S., Hily-Blant P., Faure A., Hernandez-Vera M., Lique F., 2018, A&A, 615, A52
- Maret S., Faure A., Scifoni E., Wiesenfeld L., 2009, Mon. Not. R. Astron. Soc., 399, 425
- Matsakis D. N., Brandshaft D., Chui M. F., Cheung A. C., Yngvesson K. S., Cardiasmenos A. G., Shanley J. F., Ho P. T. P., 1977, ApJ, 214, L67
- Matsakis D. N., Bologna J. M., Schwartz P. R., Cheung A. C., Townes C. H., 1980, ApJ, 241, 655
- Müller H. S. P., Schlöder F., Stutzki J., Winnewisser G., 2005, J. Mol. Struct., 742, 215
- Offer A., Flower D. R., 1990, J. Chem. Soc., Faraday Trans., 86, 1659
- Offer A. R., Hemert M. C. v., Dishoeck E. F. v., 1994, The Journal of Chemical Physics, 100, 362
- Rathborne J. M., Lada C. J., Muench A. A., Alves J. F., Lombardi M., 2008, The Astrophysical Journal Supplement Series, 174, 396
- Rist C., Alexander M. H., Valiron P., 1993, J. Chem. Phys., 98, 4662
- Rosolowsky E. W., Pineda J. E., Foster J. B., Borkin M. A., Kauffmann J., Caselli P., Myers P. C., Goodman A. A., 2008, The Astrophysical Journal Supplement Series, 175, 509
- Rydbeck O. E. H., Sume A., Hjalmarson A., Ellder J., Ronnang B. O., Kollberg E., 1977, ApJ, 215, L35
- Schleipen J., ter Meulen J., 1991, Chem. Phys., 156, 479
- Stutzki J., Winnewisser G., 1985a, A&A, 144, 1
- Stutzki J., Winnewisser G., 1985b, A&A, 144, 13
- Tkac O., Saha A. K., Loreau J., Ma Q., Dagdigian P. J., Parker D. H., van der Avoird A., Orr-Ewing A. J., 2015, Mol. Phys., 113, 3925
- Zhou D.-d., Wu G., Esimbek J., Henkel C., Zhou J.-j., Li D.-l., Ji W.-g., Zheng X.-w., 2020, A&A, 640, A114
- van der Tak F. F. S., Black J. H., Schöier F. L., Jansen D. J., van Dishoeck E. F., 2007, A&A, 468, 627

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