A simple model for high rotational excitations of molecules in a superfluid

To cite this article: Igor N Cherepanov et al 2022 New J. Phys. 24 075004

View the article online for updates and enhancements.
PAPER

A simple model for high rotational excitations of molecules in a superfluid

Igor N Cherepanov\textsuperscript{1}, Giacomo Bighin\textsuperscript{1,2}, Constant A Schouder\textsuperscript{3}, Adam S Chatterley\textsuperscript{3,\dagger}, Henrik Stapelfeldt\textsuperscript{3,∗} and Mikhail Lemeshko\textsuperscript{1,∗}

\textsuperscript{1} Institute of Science and Technology Austria, Am Campus 1, 3400 Klosterneuburg, Austria
\textsuperscript{2} Institut für Theoretische Physik, Universität Heidelberg, Philosophenweg 19, D-69120 Heidelberg, Germany
\textsuperscript{3} Department of Chemistry, Aarhus University, 8000 Aarhus C, Denmark

∗ Authors to whom any correspondence should be addressed.
E-mail: henriks@chem.au.dk and mikhail.lemeshko@ist.ac.at

Keywords: helium nanodroplets, molecular alignment, molecules in fields

Abstract

Recently it became possible to study highly excited rotational states of molecules in superfluid helium through nonadiabatic alignment experiments (Cherepanov \textit{et al} 2021 \textit{Phys. Rev. A} 104 L061303). This calls for theoretical approaches that go beyond explaining renormalized values of molecular spectroscopic constants, which suffices when only the lowest few rotational states are involved. As the first step in this direction, here we present a basic quantum mechanical model describing highly excited rotational states of molecules in superfluid helium nanodroplets. We show that a linear molecule immersed in a superfluid can be seen as an effective symmetric top, similar to the rotational structure of radicals, such as OH or NO, but with the angular momentum of the superfluid playing the role of the electronic angular momentum in free molecules. The simple theory sheds light onto what happens when the rotational angular momentum of the molecule increases beyond the lowest excited states accessible by infrared spectroscopy. In addition, the model allows to estimate the effective rotational and centrifugal distortion constants for a broad range of species and to explain the crossover between light and heavy molecules in superfluid \textsuperscript{4}He in terms of the many-body wavefunction structure. Some of the above mentioned insights can be acquired by analyzing a simple $2 \times 2$ matrix.

1. Introduction

Interactions of individual molecules with superfluid helium-4 has been extensively studied during the last decades both experimentally and theoretically [1–3]. According to infrared spectroscopy, the rotational motion of most molecules is affected by superfluid helium only quantitatively: while no drastic qualitative changes in rotational spectra is observed, the spectroscopic constants of molecules become ‘renormalized’ due to the molecule–solvent interactions. In particular, the rotational constant, $B$, and the centrifugal distortion constant, $D$, assume different values as compared to gas phase molecules, $B^* < B$ and $D^* > D$. However, for the lowest $J$-levels, the rotational energy, $E_J$, can still be accurately described by the gas-phase expression [4–6]:

$$E_J = B^*J(J+1) - D^*J^2(J+1)^2.$$  \hspace{1cm} \text{(1)}

Although there is little doubt that equation (1) describes the low-energy rotational structure for most molecules in superfluid \textsuperscript{4}He, little is known about the higher excited rotational states. In particular, we are talking about the states that are not initially thermally populated due to the helium environment ($T \approx 0.37$ K in helium nanodroplets). Due to the spectroscopic selection rules, $\Delta J = \pm 1$, conventional infrared and microwave spectroscopies are able to reach as far as only one rotational state above the initial Boltzmann distribution, on the order of $J \sim 5$.

Recently it became possible to experimentally probe highly excited rotational states of molecules in helium nanodroplets using non-adiabatic alignment protocols [7–10]. Namely, analysing the Fourier
transforms of alignment traces allowed to reveal the energies and lifetimes of rotational levels in superfluid 4He, up to \( J \sim 16 \). Moreover, the technique is applicable to molecules, that are non-responsive to infrared spectroscopy, such as I\(_2\). At the moment, theoretical approaches that would explain the behavior of solvated molecules at high angular momenta are lacking, be it simple models or \textit{ab initio} numerical techniques. Apart from understanding the fundamentals of molecular solvation, such theory is required to control molecular rotation in solvents, e.g. using an optical centrifuge [11, 12].

The goal of this paper is to present a simple quantum mechanical model that provides insights into high-\( J \) states of molecules in superfluids and to put it in the context of the experimental data gathered during the years in the low \( J \)-regime, i.e. on renormalization of the effective \( B^* \) and \( D^* \) constants of equation (1). We would like to emphasize that a lot of theoretical work has been previously devoted to explaining renormalization of rotational and centrifugal distortion constants in superfluid helium. Most quantum models and \textit{ab initio} approaches, however, focused on properties of molecules in the ground and the lowest excited rotational states [4, 13–20]. The extension of first principle treatments to highly excited states seems quite challenging [21]. Since at the moment even simple-model approaches to such states are lacking, here we resort to a phenomenological treatment, based on the previously discussed angulon model [22–24], which we simplify further in order to make it more transparent.

The present study builds upon our recent experimental and theoretical work [10], where we briefly sketched the model and showed that it provides a good agreement with experimental data for CS\(_2\) and I\(_2\) molecules up to \( J \sim 16 \). Here, apart from describing the theoretical machinery of our model in more detail, we gain several novel insights into the highly excited states and connect their behavior to the large amount of experimental data available for the states with low angular momentum. In particular, we focus on the following aspects:

(a) The relatively sharp crossover between the spectra of light and heavy molecules in helium. It is considered common knowledge that rotations of light molecules (usually defined as \( B \lesssim 1 \text{ cm}^{-1} \)) are almost not affected by helium, while the rotational constants of heavy molecules (\( B \gtrsim 1 \text{ cm}^{-1} \)) are strongly renormalized [1]. It is, however, clear that the effects of the solvent cannot be reduced to the magnitude of the rotational constant alone (as in ‘heavy’ vs ‘light’), since the strength of the molecule–He interactions and the available excitation channels in the superfluid also have to play a role. This is confirmed e.g. by path integral Monte-Carlo simulations for the LiH molecule in 4He clusters [25]. Despite being a light rotor (LR) (\( B \approx 7.5 \text{ cm}^{-1} \)), LiH features strong anisotropic interactions with helium [26], which results in strong renormalization of its rotational constant, \( B^*/B \sim 0.06 \) [25].

Using our simple model, we show how the interplay between the value of \( B \) and the strength of the anisotropic molecule–He interactions result in a sharp transition between the heavy and light molecules, cf figure 3.

(b) We show that the statement that ‘light molecules are not strongly affected by helium’ does not necessarily hold for highly excited rotational states with \( J \gtrsim 5 \), cf the simulated Raman spectrum in figure 2.

(c) The scaling of the effective centrifugal distortion constant \( D^* \) with \( B \). In reference [27] the following empirical relation of was found by fitting to experimental data:

\[
D^* = 0.031 \times B^{1.818}.
\]

The microscopic picture behind this empirical law is, however, not clear, let alone the origin and dimensionality of the prefactor. Furthermore, assuming that light molecules are almost unaffected by helium, one would expect \( D^* \rightarrow 0 \) for large \( B \), which contradicts equation (2). We show that a similar law follows from our model, with a prefactor dependent on molecule–helium interactions, cf equations (11) and (12) and figure 4.

The points (a)–(c) above allow to integrate the results of our model (which focuses on highly excited states) into the previously gathered knowledge on low molecular excitations. We start by describing the model in section 2 and show that a linear molecule in superfluid helium can be seen as an effective symmetric top. This description is similar to that of open-shell molecules like OH or NO [28, 29], but with the angular momentum of the superfluid playing the role of the angular momentum of the electronic shell. Here we are going to omit a detailed discussion of the angular momentum transfer between the molecule and the superfluid and refer the interested reader to our earlier work [10]. In section 3 we analyze the energy level structure of such an effective symmetric top and gather insights relevant to experiments on solvated molecules in excited rotational states. In section 4 we show how the model can be simplified even further and that important insights can be gathered from solutions of a \( 2 \times 2 \) matrix. Finally, in section 5
we connect the model to the data available in the low-$J$ regime and present analytic estimates for effective spectroscopic constants, $B^*$ and $D^*$, and compare them with experiment. Section 6 provides the conclusions of this study.

2. A solvated linear molecule becomes a symmetric top

2.1. The angulon Hamiltonian

We consider a linear molecule with a rotational constant $B$ revolving in the bath of bosons (collective excitations in $^4$He). To obtain the simplest possible model, we take into into account only a single mode of the bath with energy $\omega$ and angular momentum $\lambda$. In the case of superfluid helium it might be tempting to label these excitations as rotons, however we intentionally keep the treatment as general as possible. In addition, we take into account only the linear molecule–He coupling term. This corresponds to a further simplification of the previously developed angulon model [22, 23]. In the molecular (body-fixed) frame, the system is described by the following Hamiltonian [30]:

$$\hat{H} = B(\hat{L} - \hat{\Lambda})^2 + \omega \sum_n \hat{b}^\dagger_{n\lambda} \hat{b}_{n\lambda} + u(\hat{b}^\dagger_{0\lambda} + \hat{b}_{0\lambda}), \quad (3)$$

where $\hat{b}^\dagger_{n\lambda}$ $(\hat{b}_{n\lambda})$, obeying the commutation relation $[\hat{b}_{n\lambda}, \hat{b}^\dagger_{n'\mu}] = \delta_{\lambda\mu}\delta_{n'n'}$, create (annihilate) a bosonic excitation with angular momentum $\lambda$ and projection onto the molecular (i.e. interatomic) z-axis $n$. The parameter $u$ reflects the strength of the anisotropic molecule-bath interaction. $\hat{L}$ is the total angular momentum of the system and $\hat{\Lambda} = \sum_{n\lambda} \hat{b}^\dagger_{n\lambda} \sigma_{\lambda\mu} \hat{b}_{n\lambda}$ defines the angular momentum acquired by the bath. Here, $\sigma_{\lambda\mu}$ denotes the angular momentum matrices fulfilling the $SO(3)$ algebra in the representation of angular momentum $\lambda$.

In this paper we focus on the weak-coupling theory, that is, we start from a non-interacting case, corresponding to no helium excitations and add excitations one by one. The weak coupling angulon theory accounting for a single excitation of helium was shown to predict renormalization of rotational constants of corresponding rotational states can be expressed through the symmetric-top states $|LNM\rangle \rightarrow |\Lambda\mu\nu\rangle$. This is impossible to do for the case of a rotating molecular impurity, since different components of the angular momentum $\hat{L}$ do not commute with each other and it is therefore impossible to replace $\hat{L}$ in equation (3) by a classical number $L$, as one could do for the total linear momentum operator, $\hat{P} \rightarrow P$. Although the magnitude of the total angular momentum, $L$, is conserved, a general solution is going to be a superposition of states corresponding to different projections $N$, which, in turn, can contain different numbers of bosonic excitations (the $M$-quantum number plays no role in the absence of external fields).

2.2. Basis states and diagonalization

It is worth noting that in the case of a particle linearly moving in a bosonic environment (the so-called ‘polaron problem’), writing the Hamiltonian in the frame co-moving with the particle (by analogy with equation (3)) allows to completely decouple the particle and environment degrees of freedom [33]. This is impossible to do for the case of a rotating molecular impurity, since different components of the angular momentum $\hat{L}$ do not commute with each other and it is therefore impossible to replace $\hat{L}$ in equation (3) by a classical number $L$, as one could do for the total linear momentum operator, $\hat{P} \rightarrow P$. Although the magnitude of the total angular momentum, $L$, is conserved, a general solution is going to be a superposition of states corresponding to different projections $N$, which, in turn, can contain different numbers of bosonic excitations (the $M$-quantum number plays no role in the absence of external fields).
We diagonalize the Hamiltonian, equation (3), in the following basis:

$$\psi_{LM}^{(m)} = \langle LN M \rangle_{\text{mol}} (b_{L M}^{\dagger} b_{L \lambda 1}^{\dagger} \ldots b_{L \lambda n}^{\dagger} |0\rangle_{\text{bos}}).$$

$$N = \sum_{i=1}^{m} n_i$$ and M refer to the total projection of $L$ on the molecular and laboratory z-axis, respectively. $N$ and M take values in the range $[-L, L]$. An additional condition on the total projection $N$ is imposed by the following limitation on $n_i$: $|n_i| \leq \lambda$. As stated above, we restrict our basis set to $m \leq 3$. Note that although equation (3) is a substantial simplification of the original angulon Hamiltonian, the ansatz of equation (4) represents a substantially expanded basis set compared to the previous treatments, where only single excitations ($m = 1$) were taken into account [22]. Including multiple bath excitations allows to describe a broader range of molecules using the weak-coupling theory.

The $m = 0$ case describes a bare (‘gas phase’) molecular state $\psi_{LM}^{(m=0)} = |L, N = 0, M\rangle_{\text{mol}} |0\rangle_{\text{bos}}$. Diagonalization of the Hamiltonian in this basis obviously leads to the $(2L + 1)$-fold degenerate energy spectrum of an isolated rigid rotor, $BL(L + 1)$. The projection $N$ equals to zero in the absence of the excitation since we assume that the molecule is linear. The $m > 0$ cases introduce multiple excitations of the bath $b_{L M}^{\dagger} b_{L \lambda 1}^{\dagger} \ldots b_{L \lambda n}^{\dagger} |0\rangle_{\text{bos}}$.

In our model, the molecule can directly induce only deformations of the boson density that preserve $N = 0$. Thus, they are strongly aligned along the molecular z-axis. This can be seen from the third term in equation (3) and from the corresponding density plot for $L = 0$ in figure 1(a). Nevertheless, the presence of the spin-orbit-like (or Coriolis-like) interaction, the $-2L \cdot \Lambda$ term in equation (3), causes precession of $\Lambda$ about the molecular z-axis, somewhat similar to a spin in a magnetic field. Minimization of the angle between $L$ and $\Lambda$ (which, in turn, minimizes the energy of the system) leads to increase in $N$ and hence to the wider distribution of the bosons density with respect to the molecular z-axis as shown in figure 1(a). As a result, the linear molecule dressed by a cloud of excitations resembles a symmetric top whose non-zero projection $N$ is exclusively provided by the angular momentum of the He atoms in the solvation shell. In the following sections we discuss how the spectrum of such an effective symmetric top differs from the quadratic spectrum of a rigid linear rotor.

3. Highly excited rotational states in a superfluid

We start from exploring the stationary states of the system, previously briefly described in reference [10]. In what follows, we show that through analysing the states of an effective symmetric top (cf section 2.1), one can understand the distribution of angular momentum due to the molecule–helium interaction and how it changes in an external laser field.

Figure 1 shows the possible states of the system for the case of one, two and three bath excitations. Each dot in figures 1(b)–(d) represents a unique configuration, their energies are obtained by diagonalization of the Hamiltonian, equation (3), assuming $n = 0$. To facilitate the visualisation of each contribution, we perform diagonalization in each of three bases, equation (4) with $m = 1–3$, separately and plot the calculated energies in figures 1(b)–(d), respectively. The blue dots correspond to $m = 0$, i.e. to the energies of an isolated gas phase molecule, $BL(L + 1)$. In this case, the molecular angular momentum equals to the total angular momentum $J = L$, no excitations of the bath are present.

The red dots in figures 1(b)–(d) form the band of excited states. For illustrative purposes, figure 1(a) shows the molecular-frame densities of He corresponding to three of these excited states at $L = 0, 14$ and 35, also marked in figure 1(b). In these configurations, the total angular momentum $L$ is shared between the molecule and the helium excitations. For simplicity we begin with the states involving single excitations, $m = 1$, carrying energy $\omega$ and angular momentum $\lambda$ with projection $n$ onto the molecular z-axis, as shown in figure 1(b). Neglecting the off-diagonal $L \Lambda \pm$ terms in equation (3), the energies in the $|LN M\rangle$-basis are given by:

$$E_{\Lambda n}^{\lambda} = BL(L + 1) - 2Bn^2 + BL(\lambda + 1) + \omega,$$

where we introduced an additional shift by the excitation energy $\omega$. Equation (5) corresponds to the energies of an oblate (disk-shaped) symmetric top, shifted by $B(\lambda + 1) + \omega$ from zero. Since the off-diagonal components of the $-2L \cdot \Lambda$ term in equation (3) mix $n$, the resulting state in the most general case corresponds to a superposition of different $n$ projections. From the shape of the band of excited states in figure 1(b) one can see that the energetics remains similar to that of an oblate symmetric top even when the off-diagonal terms are fully taken into account. In particular, the system tends to occupy the states with non-zero $n$.

To provide an intuitive understanding of the perturbations caused by molecular rotation, we plot the distribution of helium density in the molecular frame for selected excited states in figure 1(a). Note that
Figure 1. (a) Boson density distributions in the molecular (body-fixed) frame for selected excited states (marked by the corresponding symbols in (b)). (b)–(d) Energy diagram of the band of excited states involving single \((m = 1)\), double \((m = 2)\), and triple \((m = 3)\) excitations of the bosonic bath (red dots), respectively. The gas-phase rotational spectrum \((m = 0)\) is shown by the blue dots. The green line indicates configurations with projection \(n \approx 0\) onto molecular \(z\)-axis, the black lines denote the excited states with minimum energy and largest possible total projection, \(N = \sum n_i\), for a given \(L\). The absolute energy units correspond to the choice of \(B = 0.04\) cm\(^{-1}\).

these densities are obtained at \(u = 0\) and do not correspond to the density deformations induced by the molecule (as discussed in reference \([10]\)). Instead, these plots are supposed to illustrate how the excited bath states look like in real space in the absence of molecule–helium interactions.

Let us consider a particular excited state at \(L = 0\) with a well-defined projection, \(n = 0\), marked by the green square in figures 1(a) and (b). The energy cost to create such an excitation is \(\omega + B\lambda(\lambda + 1)\). The angular density distribution plotted in figure 1(a) shows that the bosons primarily reside at the poles of the molecule (linear configuration). As \(L\) increases, the states with the dominating zero projection contribution form the upper edge of the band in figure 1(b) coloured in green. Classically, they might be thought of as rigid rotation of the molecule with its solvation shell.

However, the states with \(n \approx 0\) are not the ground state of an effective oblate top described by equation (5). For \(L > 0\), as soon as the excitation is created, the Coriolis coupling \(-2\hat{L} \cdot \hat{\Lambda}\) makes the \(n \neq 0\) configurations energetically more favorable. The bosons density shifts to the waist of the molecule (\(T\)-shape configuration) as \(L\) increases. The states with the maximum \(|n|\) build the lower edge of the band in figure 1(b) coloured in black. Its parabolic shape is defined by the above mentioned restrictions set on \(n\): (i) \(|n| \leq L\) and (ii) \(|n| \leq \lambda\). The minimum energy equals to \(\omega\) and it is reached at \(L = \lambda\). Furthermore, the lowest excited state at \(L = \lambda\) shows a perfectly uniform distribution over \(n\). This state is marked by the
black square in figures 1(a) and (b). Its angular density distribution is delocalized as plotted in figure 1(a). In the classical picture, we interpret these observations as manifestation of non-rigidity of the molecule–bosons coupled rotation.

Further growth of \( L \) beyond \( \lambda \), nevertheless, leads to the bending up of the lower edge of the band since \(|n|\) can no longer increase. In figures 1(a) and (b) we mark one of the states satisfying \( L < \lambda \) by the black square. The boson density moves back towards the poles of the molecule. We would like to stress that these findings are analogous to the resonance behavior of the helium anisotropy found within the semiclassical toy model [17]. In that model, the solvation shell is modelled as a ring of \( N_{16} \) helium atoms. Identically to \( \lambda \) in our model, \( N_{16} \) determines the symmetry of the helium solvation shell. The maximum anisotropy observed at \( L = N_{16} \) draws parallels to the results discussed above.

Qualitatively, similar considerations are valid for double (\( m = 2 \)) and triple (\( m = 3 \)) excitations. The corresponding energy diagrams are shown in figures 1(c) and (d). The only noticeable difference arises from the possibility to sum up individual projections \( n \) to the total projection \( N \). The constraint \(|n| \leq L \) is thereby lifted which substantially expands the Hilbert space of the bath excitations. As a consequence, the lower edge of the band in the range of \( L < \lambda \) becomes flat. It happens due to the fact that the combinations of several excitations having the largest possible projections \(|n| = \lambda \) of the opposite sign are allowed even for small \( L \). The minimum energy therefore reduces to \( m \omega \), the lower edge of the band starts bending upwards at \( L = m \lambda \).

### 4. Even simpler: a two-level model

In the previous section we discussed the possible states of the ‘many-body symmetric top’ without explicitly taking into account the coupling between these states induced by the molecule–helium interactions. A non-zero value of \( u \) results in coupling of the bare molecular state (\( m = 0 \)) to the excited states with \( m > 0 \) discussed above. The deviations of the final energies with respect to the gas-phase spectrum describe the net effect of the surrounding environment on molecular rotation. These perturbations can be detected in experiments as a change of the effective spectroscopic constants and are therefore of particular interest. While it is possible to evaluate them numerically, we would like to focus on the aspects of the model available for analytical treatment at first.

In our model, bare rotational states couple in first order only to the single excitations with \( n = 0 \) (cf the third term in equation (3)). For small \( L \), the gas-phase energies and the band of excited states are separated by the relatively large energy gap, \( \Delta_0 \), as compared to the rotational kinetic energy:

\[
\Delta_0 = \Delta_{L=0} = \omega + B \lambda (\lambda + 1).
\]  

(6)

Note that the gap depends on \( B \) and never closes for small \( L \). In particular, this means that a few well-distinguished rotational levels must be present even for very LR with \( B \) exceeding \( \omega \), as confirmed by experiment [1]. Since it is hard to obtain an accurate analytical expression for the gap, \( \Delta_0 \), for an arbitrary \( L \), we use its numerically calculated values shown in figure 1(b). Nevertheless, one can say that in the linear approximation its slope is approximately given by \( B \lambda \). The ratio \( u/\Delta_0 \) and its dependence on \( L \) define how strong the bath perturbs the molecular energies. If the interaction strength is comparable to or exceeds the kinetic energy of the excitation, \( u \gtrsim \Delta_0 \), the rotational spectrum is subject to strong renormalization. In the opposite case of \( u \ll \Delta_0 \), the molecule does not experience a strong influence from the bath.

In figure 2 we calculate the Raman-like (\( \Delta L = \pm 2 \)) rotational spectrum for a typical light molecule with \( B = 4 \text{ cm}^{-1} \) and \( u/\Delta_0 \approx 0.05 \). Although the energy of the first excited state, \( L = 1 \), exceeds \( \omega \), we see three well-defined spectral lines attributed to the total angular momentum states up to \( L = 6 \). Because of the small \( u/\Delta_0 \) ratio, they exhibit a slight red shift and minor changes in intensity distribution in comparison to the gas phase. This coincides with a few percent change in the rotational constants observed for light molecules [1]. A secondary substructure originating from the perturbed band of the excited states is separated from the main peak by the energy gap of order \( \Delta_0 \). We do not label these spectral features in the figure because of their negligible spectral weight.

The situation changes radically when \( L \) is further increased. As may be seen in figure 1(b), the \( L \)-dependent energy gap \( \Delta_0 \) shrinks and finally closes at

\[
L_0 \approx \left( \frac{\omega + B \lambda (\lambda + 1)}{2B} \right)^{1/2}.
\]  

(7)

Referring back to figure 2, we observe that the lines involving the states \( L \gtrsim L_0 \) \((L_0 = 8 \) in this particular case) develop a rich substructure consisting of multiple secondary peaks. In principle, all of them might be ascribed to the transitions between excited states that preserve \( N \) but change \( L \) according to the selection
rules $\Delta L = \pm 2$. We expect that these lines will be substantially broadened if one goes beyond the single mode approximation and includes a full continuous dispersion of bulk helium $\omega(k)$ into the model. This effect has been demonstrated in reference [34] for symmetric top molecules. It quantitatively explains the anomalous broadening of spectral lines, initially observed in experiments with CH$_3$ [35] and NH$_3$ [36] in helium droplets.

Next, we derive simple analytical formulas for the renormalized spectroscopic constants. In first order, the gas-phase rotational states are coupled only to the states with $n = N = 0$ and $m = 1$, which are, in turn, coupled to states with nonzero $N$ and $m \geq 1$ in higher orders. To simplify the problem, we can assume that all higher-order interactions can be incorporated into an effective energy shift, $\delta_L$, of the single excitations with $N = 0$ with respect to the energy given by equation (5), $E^0_L = \Delta_0 + BL(L + 1)$.

In such a way, we can qualitatively describe the $L$-dependent deformations of the gas phase spectrum as coupling of the bare molecular states to a single ‘dressed’ $N = 0$, $m = 1$ state for a given $L$, which corresponds to an effective two-level system:

$$\hat{H}'_L = \begin{bmatrix} BL(L + 1) & u \\ u & BL(L + 1) + \Delta_0 - \delta_L \end{bmatrix}.$$  \hspace{1cm} (8)

After dropping $L$-independent contributions, the ground state energy of the Hamiltonian (8) reads:

$$E_L = BL(L + 1) - \frac{\delta_L}{2} - \sqrt{(\Delta_0 - \delta_L)^2 + 4u^2}.$$  \hspace{1cm} (9)

Based on equation (5), we set $\delta_L = 2B\gamma L(L + 1)$ with the parameter $\gamma \in [0, 1]$ defining how strong is the effect of high order interactions on the $N = 0$, $m = 1$ states, i.e. how much their energy effectively shifts from $E^0_L$ (located close to the green line in figure 1(b)) towards the lower edge of the band (black line in the same figure). In both limits of light ($B \to \infty$) and heavy ($B \to 0$) rotors, $E_L$ can be expanded in a series:

$$E_L = B^* L(L + 1) - D^* L^2(L + 1)^2 + O(L^3(L + 1)^3),$$  \hspace{1cm} (10)

cf equation (1). For LRs, we make use of the condition $u \ll \Delta_0$ to show that the zero-order term in $u$ cancels out leading to weak renormalization of spectroscopic constants:

$$\frac{B_{LR}^*}{B} \approx 1 - \frac{2\gamma u^2}{\Delta_0^2}; \quad D_{LR}^* \approx \frac{4B^2\gamma^2 u^2}{\Delta_0^2}.$$  \hspace{1cm} (11)

These expressions coincide with the exact analytical results obtained for small $L$ in reference [10]. Both renormalized spectroscopic constants contain the small parameter $u/\Delta_0$ which guarantees that $B^* \to B$ and $D^* \to 0$ in the free-rotor limit.

In the opposite limit of HRs, the expansion of energy in powers of a small parameter $(\Delta_0 - \delta_L)^2$ gives

$$\frac{B_{HR}^*}{B} \approx 1 - \gamma; \quad D_{HR}^* \approx \frac{B^2\gamma^2}{2u}.$$  \hspace{1cm} (12)

Since the average value of the parameter is $\gamma \sim 1/2$, the rotational constant shows non-negligible renormalization in this case. The expression for $D_{HR}^*$ closely resembles the empirical formula equation (2) found in reference [27] by fitting to existing experimental data. In our case, however, the prefactor explicitly depends on the interaction strength and has a clear physical meaning (setting $\gamma = 1/2$ and $u = 10$ gives the prefactor of $\sim 0.01$, i.e. on the same order as in equation (2)). Moreover, equation (12) predicts the same dependence on $B$ as the approximate solutions of the strong coupling model reported in reference [23].
5. Scaling of spectroscopic constants

The model described in section 2 has already been successfully used to describe experimental data on rotational states of CS₂ and I₂ molecules up to \( J \sim 16 \) [10]. The experimental data available on highly excited rotational states is, however, still quite limited and it is important to place the model in the context of the large amount of measurements for the low-\( J \) excitations. Over the past two decades a lot of experimental and theoretical data were collected for effective spectroscopic constants of a broad range of molecular species in superfluid helium (see e.g. reference [1]). In this section, we work with the numerical solutions of the full model developed in section 2, as opposed to the simplified solutions discussed in the previous section. The values of \( B^\prime \) and \( D^\prime \) discussed below were obtained by fitting the energies of the \( L = 0–3 \) states to equation (1). Apart from providing the estimates for spectroscopic constants, we address the transition between the light and heavy species as well as the scaling of \( D^\prime \) with \( B \), see sub. (a)–(c) in the introduction, section 1.

In figure 3(a) we compare the effective rotational constants \( (B/B^\prime \) as a function of \( B \)) estimated using our model (lines) with the results of experiments (black circles) [1, 4–6, 9, 13, 14, 19, 27, 35–56]. The energy of the bosonic mode \( \omega \) was fixed to 6 cm\(^{-1}\), the roton energy of bulk helium [57]. This choice is motivated by the extremely large density of solvent states near the roton excitation energy. The angular momentum of excitation \( \lambda \) was set to 14, which has already been used to explain the experimental data in reference [10]. Although this number seems quite large, it reflects the density profile around the molecule and corresponds to approximately 2\( \lambda \) He atoms forming the solvation shell in the semiclassical picture. Furthermore, it is important to emphasize that the Hamiltonian of equation (3) is only an effective one. It is clear that a microscopic coupling of a molecule with helium is going to involve terms corresponding to excitations of many bosonic modes, such as \( b b^\dagger, b^\dagger b b^\dagger b^\dagger \), in addition to single excitations of equation (3). Therefore, the third term of equation (3) is to be seen as coupling to some effective mode with angular momentum \( \lambda \), which at the microscopic level consists of several excitations with much lower angular momentum.

We would like to emphasize that the results presented here for the spectroscopic constants are merely estimates: we are trying to see whether some general trends, previously obtained by fitting to experimental data, can also be derived from the model. For example, the molecule–helium coupling constant, \( u \), depends on the details of the molecule–He potential energy surface (PES) and is going to be different for each molecule. Moreover, \( u \) does not show any significant correlation with \( B \): according to reference [23], the interaction parameter extracted from the molecule–He PES and expressed in absolute units varies within one order of magnitude for the species whose rotational constant cover more than three orders of magnitude. Since here we focus on the general trend, we present the theoretical curves for three different values of \( u \) and the experimental data on \( B/B^\prime \), without discussing concrete molecular species.

The overall trend seen in figure 3(a) can be explained semiclassically by the ‘adiabatic following’ model [14, 58, 59], revealing the crossover between the heavy and light species. In a simple picture, HRS \((B \lesssim 1 \text{ cm}^{-1})\) rotate slow enough for the helium solvation shell to follow. Such strong coupling leads to a significant reduction of \( B \), up to a factor of 6. LRs \((B \gtrsim 1 \text{ cm}^{-1})\), in contrast, rotate so fast that they decouple from helium and their rotational constant is almost not renormalized.

Note that the ansatz of equation (4) corresponds to the weak-coupling approximation, which breaks down in the limit of \( B \to 0 \), i.e. for very heavy molecules. The results furnished by the model in this regime (dotted lines figure 3(a)) are unphysical. This behavior might also be rationalized within the effective two-level model of section 4. If \( B \to 0 \), the shift \( \delta_1 \) in equation (9) vanishes, thereby eliminating the \( L \)-dependence from the model (or alternatively, the lower edge of the band in figure 1(b) becomes flat).

Although the admixture of bosonic excitations into the total wave function might be dominant, it does not bring any \( L \)-dependent contribution to the energy. The decreasing renormalization in this region is, thus, of a completely different nature than in the case of LRs. Including the excitations with \( m > 3 \) into the basis may substantially improve solutions of the model Hamiltonian, equation (3), in this regime.

Figures 3(b) and (c) break down the contributions of different numbers of helium excitations into the total wavefunction for heavy and light molecules, respectively. The coupling parameter is set to the same value of \( u = 10 \text{ cm}^{-1} \) in both cases. For heavy molecules, \( B = 0.04 \text{ cm}^{-1} \), figure 3(b), the contribution of the bare molecular state, \( m = 0 \), is approximately 50% for \( L = 0 \) and monotonously decreases with \( L \), while higher excitations \( m = 2 \) and 3 get more populated. For light molecules, on the other hand, there is a sharp transition point \( L_o \), such that for \( L < L_o \) only \( m = 0 \) states are populated, while for \( L > L_o \) also the states with nonzero \( m \) are, see the example for \( B = 4 \text{ cm}^{-1} \) in figure 3(c).

This shows an important difference between heavy and light molecules, previously broadly discussed in the literature from other points of view [1]. For heavy molecules, even in the absence of rotation, the molecule–helium interaction distorts the surrounding superfluid and creates He excitations co-rotating...
Figure 3. (a) The reduction factor of rotational constants, $B/B^\ast$, for molecules in helium as a function of the gas phase rotational constant, $B$. The solid lines show the theoretical predictions obtained in the present work for selected values of the interaction strength, $u$. The dotted part of the lines indicates the range of $B$ where the weak coupling theory fails. Experimental data points are shown by the black circles. (b) and (c) The relative contributions of the states involving single ($m = 1$), double ($m = 2$), triple ($m = 3$) excitations of helium as well as bare molecular states ($m = 0$) to the total wave function for a HR ($B = 0.04 \text{ cm}^{-1}$) and a LR ($B = 4 \text{ cm}^{-1}$), respectively. The interaction parameter $u = 10 \text{ cm}^{-1}$ in both cases.

Figure 4. The effective centrifugal distortion constant, $D^\ast$, as a function of the effective rotational constant $B^\ast$ (solid lines) for selected values of the interaction parameter $u$. The dotted parts of the lines indicate the range of $B^\ast$ where the weak-coupling theory fails. Experimental data points are shown by the black circles. The dashed black line corresponds to the empirical formula from reference [27].

with the molecule (a ‘non-superfluid solvation shell’). For light molecules at small $L$ the bath excitations are only virtual (in agreement with the results of reference [23]), resulting in a very small $B$-renormalization. After some critical value of $L \sim L_0$, the bare molecular states cross the excitation threshold and start coupling to the bath strongly, which results in substantial population of $m \neq 0$ states.

Figure 4 shows the effective centrifugal constant $D^\ast$ as a function of $B^\ast$ in comparison with the experimental data listed in references [4–6, 9, 27, 35, 38, 44, 51, 52, 54, 55]. In agreement with the established experimental and theoretical result, $D^\ast$ measured in helium droplets is found to be $10^2$–$10^4$ times larger than the corresponding gas-phase value. The LRs with $B \gtrsim 3 \text{ cm}^{-1}$, whose $B/B^\ast$ ratio is barely distinguishable from 1, show large $D^\ast$ of the order of 0.01 cm$^{-1}$ only if the interaction parameter, $u$, is large (orange and blue lines). Otherwise, $D^\ast$ does not scale with $B^\ast$ (green line) and might be comparable to the gas-phase centrifugal constant for some of the molecules. In the case of HRs this tendency is not apparent,
$D^*$ shows persistent dependence on $B^*$ in a wide range of $u$. In particular, one can see that, similarly to equation (12), the scaling of $D^*$ closely resembles the already mentioned empirical formula, equation (2), found in reference [27].

6. Conclusions

Thus we presented a simple quantum mechanical model describing the rotational level structure of molecules in superfluid helium nanodroplets and, in particular, capturing highly excited states (recent work [10] compared the model calculations with experiment up to $J \gtrsim 16$ for I$_2$ and CS$_2$ molecules). Here we provided details on the theoretical machinery of the model, benchmarked its results against the data on the effective spectroscopic constants $B^*$ and $D^*$ for a broad range of molecules. Although the model is already based on a simplified version of the previously reported angulon Hamiltonian [22], we have substantially simplified it further and have shown that several properties of molecules in superfluids can be understood by analyzing a simple $2 \times 2$ matrix, equation (8).

Among other results, we gathered the following insights:

(a) A linear molecule in superfluid He can be described as an effective symmetric top, with an additional quantum number describing the projection of superfluid angular momentum on the molecular $z$-axis. Coupling between the superfluid and molecular rotational angular momenta is reminiscent of that between the electronic and rotational angular momenta in the gas-phase radicals, such as OH or NO. This provides a means to work with different angular momentum coupling schemes involving the solvent angular momentum and to introduce some kind of ‘many-body Hund’s cases’ describing such coupling in the future.

(b) Analyzing the structure of such a symmetric top, whose states can be mixed by molecule–helium interactions, furnishes a few qualitative insights. For example, the crossover between the rotational behavior of light and heavy molecules in a superfluid (approximately at $B \sim 2–3\,\text{cm}^{-1}$) can be explained in terms of the many-particle wavefunction structure shown in figure 3(c), which, in turn, follows from the $L$-dependent energy gap $\Delta_L$ shown in figure 1(a).

It is important to underline the main differences between the model described here from the simple models previously discussed in the literature, in particular that of reference [17]. First and foremost, the model presented here is capable of describing highly excited rotational states of molecules in a superfluid. Secondly, it considers a rotor immersed in a bosonic bath in three-dimensional space. Thirdly, the bath is treated as a Bose–Einstein condensate of collective excitations of superfluid $^4\text{He}$. For the sake of simplicity, we restricted our consideration to a single mode of the bath. However, further generalization of the treatment is straightforward.

The results presented here and in reference [10] reveal that the structure of the highly excited rotational states can substantially deviate from the gas-phase-like equation (1), in particular for heavier molecules, such as I$_2$ and CS$_2$. This deviation needs to be taken into account while creating molecular superrotors using the optical centrifuge technique [11, 12]. In particular, one might need to redefine the adiabaticity criterion of molecule-laser interactions and to use non-linear ramp pulses in order to account for the threshold of the states as shown in figure 1.

Acknowledgments

IC acknowledges the support by the European Union’s Horizon 2020 research and innovation programme under the Marie Skłodowska-Curie Grant Agreement No. 665385. GB acknowledges support from the Austrian Science Fund (FWF), under Project No. M2461-N27 and from the Deutsche Forschungsgemeinschaft (DFG, German Research Foundation) under Germany’s Excellence Strategy EXC2181/1-390900948 (the Heidelberg STRUCTURES Excellence Cluster). ML acknowledges support by the Austrian Science Fund (FWF), under Project No. P29902-N27, and by the European Research Council (ERC) starting Grant No. 801770 (ANGULON). HS acknowledges support from the Independent Research Fund Denmark (Project No. 8021-00232B) and from the Villum Fonden through a Villum Investigator Grant No. 25886.

Data availability statement

All data that support the findings of this study are included within the article (and any supplementary files).
ORCID iDs

Adam S Chatterley https://orcid.org/0000-0003-3847-5936

References

[1] Toennies J P and Vilesov A F 2004 Angew. Chem., Int. Ed. 43 2622
[2] Ancilotto F, Barranco M, Coppens F, Eloranta J, Halberstadt N, Hernando A, Mateo D and Pt M 2017 Int. Rev. Phys. Chem. 36 621
[3] Verma D, Tanyag R M P, O’Connell S M O and Vilesov A F 2019 Adv. Phys. X 4 1553569
[4] Grebenev S, Hartmann M, Havenith M, Sartakov B, Toennies J P and Vilesov A F 2000 J. Chem. Phys. 112 4485
[5] Nauta K and Miller R E 2001 J. Chem. Phys. 115 10254
[6] Nauta K and Miller R E 2001 J. Chem. Phys. 115 8384
[7] Pentlehner D, Nielsen J H, Slenzka A, Møller K and Stapelfeldt H 2013 Phys. Rev. Lett. 110 93002
[8] Shepperson B, Sondergaard A A, Christiansen L, Kaczmarczyk J, Zillich R E, Lemeshko M and Stapelfeldt H 2017 Phys. Rev. Lett. 118 203203
[9] Chatterley A S et al 2020 Phys. Rev. Lett. 125 013001
[10] Cherepanov I N, Bighin G, Schouder C A, Chatterley A S, Albrechtsen S H, Muñoz A V, Christiansen L, Stapelfeldt H and Lemeshko M 2021 Phys. Rev. A 104 L061303
[11] Karczmarek J, Wiertz J, Corkum P and Ivanov M 1999 Phys. Rev. Lett. 82 3420
[12] Korobenko A, Milner A A and Milner V 2014 Phys. Rev. Lett. 112 113004
[13] Hartmann M, Miller R E, Toennies J P and Vilesov A F 1995 Phys. Rev. Lett. 75 1566
[14] Lee E, Farrell D and Whaley K B 1999 Phys. Rev. Lett. 83 3812
[15] Kwon Y and Whaley K B 1999 Phys. Rev. Lett. 83 4108
[16] Callegari C, Conjusteau A, Reinhard I, Lehmann K K, Scoles G and Dalfovo F 1999 Phys. Rev. Lett. 83 5058
[17] Lehmann K K 2001 J. Chem. Phys. 114 4643
[18] Lehmann K K and Callegari C 2002 J. Chem. Phys. 117 1959
[19] Zillich R E, Kwon Y and Whaley K B 2004 Phys. Rev. Lett. 93 250401
[20] Zillich R E and Whaley K B 2004 Phys. Rev. B 69 104517
[21] Zillich R E, Paesani F, Kwon Y and Whaley K B 2005 J. Chem. Phys. 123 114301
[22] Schmidt R and Lemeshko M 2015 Phys. Rev. Lett. 114 203001
[23] Lemeshko M 2017 Phys. Rev. Lett. 118 095301
[24] Lemeshko M and Schmidt B 2017 Molecular impurities interacting with a many-particle environment: from ultracold gases to helium nanodroplets Low Energy and Low Temperature Molecular Scattering ed O Dulieu and A Osterwalder (Cambridge: Royal Society of Chemistry)
[25] Zillich R E and Whaley K B 2007 J. Phys. Chem. A 111 7489
[26] Taylor B K and Hinde R J 1999 J. Chem. Phys. 111 973
[27] Choi M Y, Douberly G E, Falconer T M, Lewis W K, Lindsay C M, Merritt J M, Stiles P L and Miller R E 2006 Int. Rev. Phys. Chem. 25 15
[28] Lefebvre-Biron H and Field R W 2004 The Spectra and Dynamics of Diatomic Molecules (New York: Elsevier)
[29] Brown J M and Carrington A 2003 Rotational Spectroscopy of Diatomic Molecules (Cambridge: Cambridge University Press)
[30] Schmidt R and Lemeshko M 2016 Phys. Rev. X 6 011012
[31] Bighin G and Lemeshko M 2017 Phys. Rev. B 96 85410
[32] Bighin G, Tschibul T V and Lemeshko M 2018 Phys. Rev. Lett. 121 165301
[33] Devreeze I T 2015 arXiv:1012.4576v6
[34] Cherepanov I N and Lemeshko M 2017 Phys. Rev. Mater. 1 35602
[35] Morrison A M, Raston P L and Douberly G E 2015 J. Phys. Chem. A 117 11640
[36] Slipchenko M N and Vilesov A F 2005 Chem. Phys. Lett. 412 176
[37] Harms J, Hartmann M, Toennies J P, Vilesov A F and Sartakov B 1997 J. Mol. Spectrosc. 185 204
[38] Callegari C, Conjusteau A, Reinhard I, Lehmann K K and Scoles G 2000 J. Chem. Phys. 113 10353
[39] Conjusteau A, Callegari C, Reinhard I, Lehmann K K and Scoles G 2000 J. Chem. Phys. 113 4840
[40] Nauta K and Miller R E 2000 J. Chem. Phys. 113 9466
[41] Callegari C, Lehmann K K, Schmied R and Scoles G 2001 J. Chem. Phys. 115 10090
[42] Madeja F, Markwick P, Havenith M, Nauta K and Miller R E 2002 J. Chem. Phys. 116 2870
[43] Förtnier N, Toennies J P and Vilesov A F 2002 J. Chem. Phys. 117 6054
[44] Lindsay C M and Miller R E 2005 J. Chem. Phys. 122 104306
[45] Paesani F 2005 Phys. Rev. Lett. 94 153401
[46] von Haeften K, Metzeltin A, Rudolph S, Staemmler V and Havenith M 2005 Phys. Rev. Lett. 95 215301
[47] Scheele I, Conjusteau A, Callegari C, Schmied R, Lehmann K K and Scoles G 2005 J. Chem. Phys. 122 104307
[48] Kuyanov K E, Slipchenko M N and Vilesov A F 2006 Chem. Phys. Lett. 427 5
[49] Skvortsov D, Choi M Y and Vilesov A F 2007 J. Phys. Chem. A 111 12711
[50] Hoshina H, Skvortsov D, Sartakov B G and Vilesov A F 2010 J. Chem. Phys. 132 074302
[51] Raston P L, Knapp C J and Jäger W 2011 Phys. Chem. Chem. Phys. 13 16789
[52] Raston P L, Kelloway D D and Jäger W 2012 J. Chem. Phys. 137 041302
[53] Raston P L, Liang T and Douberly G E 2013 J. Phys. Chem. A 117 8103
[54] Raston P L, Douberly G E and Jäger W 2014 J. Chem. Phys. 141 044301
[55] Faulkner T, Miller I and Raston P L 2018 J. Chem. Phys. 148 044308
[56] Raston P L 2021 Phys. Chem. Chem. Phys. 23 25467
[57] Donnelly R J, Donnelly J A and Hills R N 1981 J. Low Temp. Phys. 44 471
[58] Patel M V, Viel A, Paesani F, Huang P and Whaley K B 2003 J. Chem. Phys. 118 5011
[59] Markovskiy N D and Mak C H 2009 J. Phys. Chem. A 113 9165