



Collision-induced Raman scattering from a pair of dissimilar particles: An intriguing mathematical model predicting the suppression of the odd-numbered partial waves

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Résumé en anglais

Relying on a simple analytic two-atom model in which the anisotropy of the interaction dipole polarizability obeys an inverse power law as a function of separation, we offer mathematical and numerical evidence that, in a monoatomic gas, the free-free Raman spectrum for a collisional pair of two different isotopes, a - a' , may vastly differ from that for a - a . This result is obtained even if a and a' are assumed to have the same mass and zero nuclear spin and even if a - a and a - a' are subject to the same interaction polarizability and potential. The mechanism responsible for this effect is inherent in the parity of the partial-wave rotational quantum number J : given that the contribution of each partial wave to the Raman cross section is controlled by a polarizability-transition matrix-element and that each of those matrix-elements has a radial component with a magnitude slightly smaller than that of the preceding partial wave, a deficit which disfavors the odd-numbered waves is accumulated upon summing over J . In the far high-frequency wing, this deficit tends to generate spectral intensities for a - a' about half as great as the a - a ones, a tendency which becomes all the more effective as temperature is decreased. We show for instance that, for the spectral branch $\Delta J = 2$, the fractional difference between the free-free differential cross sections for a - a and a - a' is $12(1-x^2)^{31} + 3x^4 12(1-x^2)^{31} + 3x^4$, with $x = E/E' - \sqrt{x} = E/E'$ (E (E') being the initial (final) state energy of the pair and $E' - E = h\nu$ ($\nu > 0$)). Remarkably, this quantity is zero at $\nu \approx 0$ but goes to 1212 for $\nu \rightarrow 0$. For $\Delta J = 0$, analogous conclusions may be drawn from the expression $(1 + \ln(1+x) - \sqrt{x})^2 \arctan x - 1(1 + \ln(1+x) - \sqrt{x})^2 \arctan x - 1$.

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[1] <http://okina.univ-angers.fr/michel.chrysos/publications>

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