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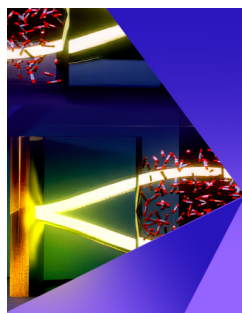
Erratum: "Intramolecular vibrational redistribution in aromatic molecules. I. Eigenstate resolved CH stretch first overtone spectra of benzene" [J. Chem. Phys. 113, 10583 (2000)] **FREE**

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J. Chem. Phys. 114, 3344 (2001)

<https://doi.org/10.1063/1.1347028>



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Erratum: "Intramolecular vibrational redistribution in aromatic molecules. I. Eigenstate resolved CH stretch first overtone spectra of benzene" [J. Chem. Phys. 113, 10583 (2000)]

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[DOI: 10.1063/1.1347028]

Due to a production error, incorrect versions of Figs. 2 (p. 10586) and 7 (p. 10593) appeared in the original paper. AIP apologizes for this error. Corrected versions of Figs. 2 and 7 are reproduced below.

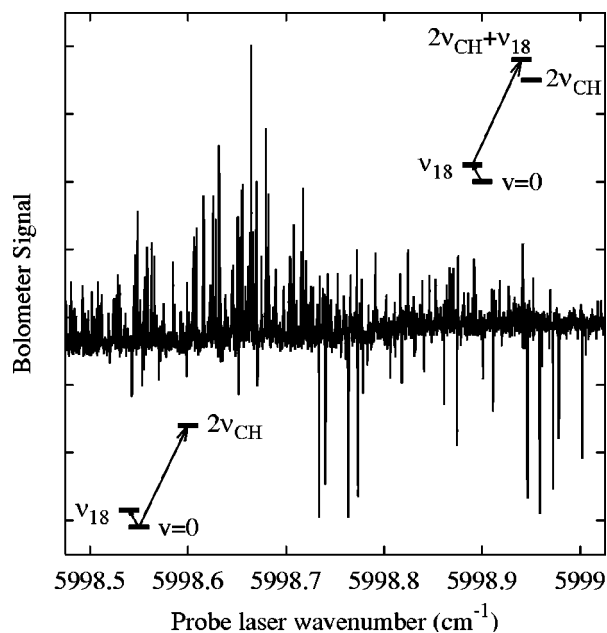


FIG. 2. The double resonance molecular beam spectrum of benzene observed with the probe laser tuned to the region of the CH stretch first overtone. Downward-going peaks correspond to transitions from the ground state, depleted by the pump laser, to the $\nu_{\text{CH}}=2$ pure overtone manifold. Upward-going peaks correspond to transitions from the upper state (ν_{18}) populated by the pump laser, to the $2\nu_{\text{CH}}+\nu_{18}$ manifold.

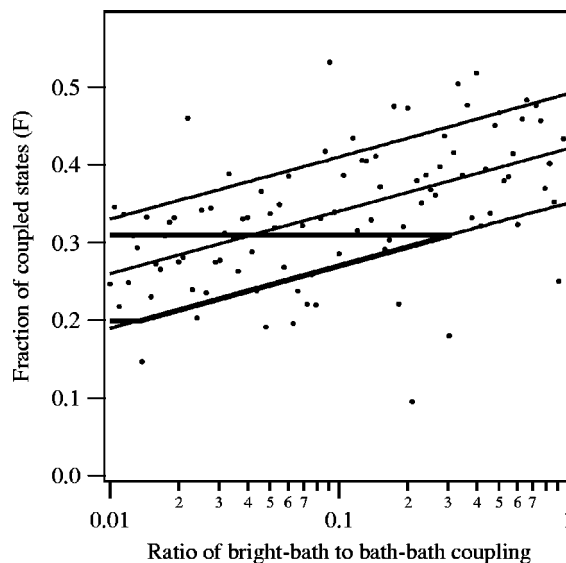


FIG. 7. The calculated fraction (F) of available states accessed by the IVR dynamics as a function of the bright state–bath to bath–bath coupling ratio. Thin lines show the result of a pseudolinear fit and the 1σ confidence bands. The thick lines show the region compatible with the range of values of F observed experimentally.

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