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Quantum state dependent chemistry of ultra-cold <sup>6</sup>Li<sub>2</sub> dimers<sup>1</sup> ERIK FRIELING, GENE POLOVY, DENIS UHLAND, University of British Columbia, JULIAN SCHMIDT, Albert-Ludwigs-Universitt Freiburg, KIRK MADI-SON, University of British Columbia — Starting from an ultra-cold ensemble of <sup>6</sup>Li<sub>2</sub> Feshbach molecules, we produce deeply bound molecules by STIRAP in the lowest energy levels of the v = 0, 5, 8 and 9 vibrational manifolds of the  $a(1^{3}\Sigma_{u}^{+})$  potential. The ensemble lifetime is found to be limited by two-body collisions with a loss rate near the universal rate for three of these states and, remarkably, below universality for the  $|v = 9, N = 0\rangle$  state. In addition, unlike all prior experimental work with ultra-cold molecules, we observe a rotational state dependence of the reaction rate. We observe that molecules in the absolute lowest triplet level are unstable. Because of the suppression of spin-changing collisions and absence of other inelastic collision channels, we conclude this instability is primarily due to trimer formation, consistent with theoretical predictions.

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