

OBSERVATION OF TWO $\text{La}(\text{C}_3\text{H}_2)$ ISOMERS FORMED BY DEHYDROGENATION OF PROPYNE

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C-H bond activation of small hydrocarbons is of importance in chemistry and industrial applications. $\text{La}(\text{C}_3\text{H}_2)$ was formed by the reaction of laser-ablated La atoms and propyne (C_3H_4) in supersonic molecular beams. Two isomers of $\text{La}(\text{C}_3\text{H}_2)$ were detected for the first time by mass-analyzed threshold ionization (MATI) spectroscopy. From the MATI spectra, the two isomers exhibit origin bands at 42953(5) and 43609(5) cm^{-1} and vibrational intervals of 425 and 535 cm^{-1} , respectively. They were identified as $\text{La}(\text{CCCH}_2)$ formed from 1,3-dehydrogenation and $\text{La}(\text{HCCCH})$ formed by 3,3-dehydrogenation and were confirmed by measurements with deuterium substituted propyne (C_3D_4) as the precursor. The 1,3-dehydrogenated complex shows a higher ionization energy and larger metal-ligand stretching frequencies than the 3,3-dehydrogenated species. Based on DFT/B3LYP calculations, the electronic transitions responsible for the observed MATI spectrum of $\text{La}(\text{HCCCH})$ isomer is $^1\text{A} \leftarrow ^2\text{A}$, and that of $\text{La}(\text{CCCH}_2)$ isomer is $^1\text{A}' \leftarrow ^2\text{A}'$.