

INFRARED EMISSION SPECTRUM OF He₂ OBSERVED BY A PULSED DISCHARGE

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The He₂ molecule is known to have many electronic transitions between Rydberg states in visible and ultraviolet regions. However, in infrared region only two bands $b^3\Pi_g - a^3\Sigma_u^+$ and $B^1\Pi_g - A^1\Sigma_u^+$ have been studied by using a DC discharge method so far. Recently we developed a time resolved Fourier transform spectroscopy with high resolution Bruker IFS 120 HR by using micro controller SX^a. In the present study, the FT system was applied to infrared emission spectroscopy of He₂ which was produced by a pulsed discharge in He with pulse width of 20 μ sec and 1 A peak-to-peak current. In the 1800-10000 cm^{-1} region, many electronic transitions have been observed in addition to the previously reported two bands. From observed time profiles of emission spectra, Rydberg states with higher energy than the *b* state are produced efficiently in afterglow plasma after termination of the discharge.

A least-squares analysis was carried out for the $h^3\Sigma_u^+ - g^3\Sigma_g^+$ and $g^3\Sigma_g^+ - d^3\Sigma_u^+$ bands in the 3200 cm^{-1} region to determine the molecular constants. A transition from an un-identified state to the $d^3\Sigma_u^+$ state has been observed with irregular P- and R- branch intensities.

^aK. Kawaguchi, O. Baskakov, Y. Hosaki, Y. Hama, and C. Kugimiya, *Chem. Phys. Lett.* **369**, 293(2003)