

Toward laser spectroscopy of rotationally cooled CaH^+ ions trapped in a cryogenic linear Paul trap

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Synopsis We have generated rotationally cooled CaH^+ ions with a cryogenic linear Paul trap toward laser spectroscopy. The direct observation of cold CaH^+ ions via laser Raman spectroscopy is now undergoing.

The study of possible time variations in fundamental constants plays an important role for the development of a ground unification theory (GUT) [1]. In addition to the fine structure constant α , the proton-to-electron mass ratio ($\beta = m_p/m_e$) is also one of important parameters for GUT, since β is related to $\dot{\alpha}/\alpha$ [1]. The ratio between a molecular vibrational transition frequency and an optical atomic transition yields the information of β . However, compared with the atomic transitions, precision measurements of the molecular transitions have not been progressed almost. In connection with this context, Kajita *et al.* proposed to detect possible time variation of β using a vibrational transition frequency of a single cold CaH^+ ion in a linear Paul trap (LPT) [2]. Then CaH^+ ions have been successfully produced by laser-induced reactions of $\text{Ca}^+(^2P_{1/2}) + \text{H}_2 \rightarrow \text{CaH}^+ + \text{H}$ in a LPT [3].

In this work we have developed a cryogenic LPT (Fig.1 (a)) in order to generate rotationally cooled CaH^+ ions. As shown in Fig.1 (b), the temperature at the LPT is cooled down to 13 K. Figure 1 (c) shows laser-induced fluorescence (LIF) images of Ca^+ Coulomb crystals in the cryogenic LPT before and after the laser-induced reactions. Since the mass of CaH^+ is a little larger than that of Ca^+ , CaH^+ ions are distributed at the outside of the ion crystal. In this example, the number of CaH^+ ions is on the order of 10^3 . In order to evaluate the rotational state distribution in $1^1\Sigma$ ($v = 0$), we have performed the rate-equation analysis including up to $J = 18$ rotational levels. The transition dipole moments and the rotational state constants were taken from the theoretical values [4] in order to include the transitions by black body radiations and spontaneous emissions. Figure 1 (d) shows plots of the time-dependence of the populations in $J = 0-3$ at 13 K, where we assume that the initial populations are the same as those at room temperature. The ro-

tational state population of $1^1\Sigma$ ($v = 0$, $J = 0$) is evaluated to be 55% after 16 minutes.

We have been working on the laser Raman spectroscopy by $1^1\Sigma \rightarrow 2^1\Sigma$ ($\lambda = 405$ nm) excitations of rotationally cooled CaH^+ ions [4]. By inserting a half mirror in the detection optics, the Raman photons ranging from 430–750 nm [5] and the LIF from Ca^+ are simultaneously detected by a photomultiplier tube and a cooled CCD camera, respectively. At present we are searching for the resonance wavelengths of the rovibrational transitions in $1^1\Sigma \rightarrow 2^1\Sigma$, which are still unknown experimentally. The status of the experiment will be presented at the conference.

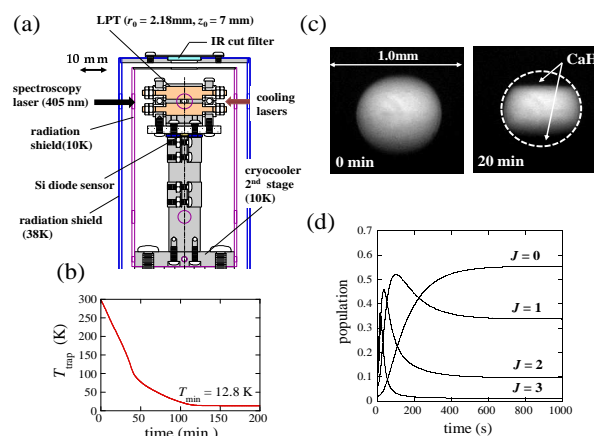


Figure 1 (a) Schematic drawing of the cryogenic LPT. (b) A plot of the wall temperature enclosing the LPT after turning on the cryocooler. (c) LIF images of Ca^+ ion crystals before and after the laser-induced reactions. (d) Simulation results of the rotational state populations of CaH^+ $1^1\Sigma$ ($v = 0$, J) as a function of time at 13 K.

References

- [1] X. Calmet *et al.*, Eur. Phys. J. C **24**, 639
- [2] M. Kajita *et al.*, J. Phys. B **42**, 154022
- [3] N. Kimura *et al.*, Phys. Rev. A **83**, 033422
- [4] M. Abe *et al.*, J. Phys. B **43**, 245102
- [5] M. Abe *et al.*, CPL **521**, 31

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