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Generation of high-repetitive, multi-MeV, pure proton beams via Coulomb explosion of micron-size hydrogen cluster target

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Laser-driven ion acceleration has been one of the most active areas of research over approximately the past decade, because accelerated multi-MeV ion beams have unique properties that can be employed in a broad range of applications. From a view point of practical applications, high-purity proton beams with high reproducibility are quite advantageous. In experiments using thin foil targets, however, protons from surface contaminants along with the high-*z* component materials are accelerated together, making the production of impurity-free proton beams unrealistic.

Here we introduce a micron-size hydrogen cluster (composed of 10⁸⁻¹⁰ hydrogen molecules) as a target to generate impurity-free, highly-reproducible, and robust multi-MeV proton beams [1, 2]. Because of the recent progress in intense laser technology, the advanced PW class lasers can now achieve intense laser fields around 10²² W/cm² [3]; with such fields, all the electrons inside the micron-size hydrogen cluster up to 3.0 μm in diameter can be fully stripped off, resulting in a pure Coulomb explosion with a pronounced increase in accelerated maximum proton energies scaled as $E_{max} = 276(d/2)^2$ MeV, where *d* is a diameter of clusters.

By using the micron-size hydrogen cluster target, we have conducted ion acceleration experiments with the 0.1 Hz PW class J-KAREN laser at QST-KPSI [4]. In order to characterize the accelerated ions, we used nuclear track detector plates (CR-39), nuclear emulsion plates, and a real-time Thomson parabola equipped with a micro-channel plate (MCP), a phosphor screen, and a CCD camera. We found that only protons having the maximum energy of ~12 MeV, consistent with the theoretical prediction, were accelerated in the laser propagation direction at a laser focused intensity of 1×10²⁰ W/cm². Based on the experimental results, the detailed ion acceleration mechanism by Coulomb explosion of clusters is discussed with the help from numerical simulations using a particle-in-cell (PIC) method.

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3. A. S. Pirozhkov *et al.*, Opt. Express **25**, 20486 (2017).
4. H. Kiriya *et al.*, Opt. Lett. **43**, 2595 (2018).

References

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