

Synthesis of halide perovskite quantum dot liquid scintillator for particle detection

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Introduction

Motivation

- We aim to develop a liquid scintillation solution (LS) for particle detection in the low-energy range ($\sim$ keV)
- Utilizing the unique luminescent and scintillation characteristics of perovskite quantum dots.
- LS based on perovskite quantum dots is designed to resolve the limitations of existing scintillators and achieving enhanced light emission at a particular wavelength.
- We try to develop a simple synthesis method that can be expanded at room temperature & pressure without the need for special high costing equipment for the next generation neutrino experiment

Experiment Setup

- MAPbBr₃ perovskite quantum dots were synthesized and coated with Oleic acid (OA) and Oleylamine (OAm), using toluene as the solvent.
- The synthesized sample was centrifuged into upper and lower fractions and a sediment, and then the particle size distribution and luminescence of both fractions were analyzed to assess sample stability.

Development of Perovskite-based LS (PVLS)

1. Fabrication of perovskite precursor
 - 1) Dissolve 2.5 mmol of PbBr₂ in 5 ml of DMF for 4 hours, then pour 0.08 ml of OA and 0.45ml of OAm and mix for about 2 hours.
 - 2) After loading 2.5mmol of MABr, mix.
 2. Fabrication of perovskite-based LS
 - 1) Pour 1 ml of precursor solution into 30 ml of toluene and mix using a stirrer for about 2 hours.
 - 2) Perform centrifugation at 15,000 rpm for 30 minutes to remove the precipitate and obtain the supernatant.
 - 3) To improve transmittance, adjust the concentration of the perovskite-based LS or use PPO to shift the fluorescence.
 - a. If it is necessary to shift the fluorescence additionally, pour 1.8 ml of the solution from step 2-1) into 98.2 ml of toluene, load PPO at a concentration of 3 g/L, and mix using a stirrer for 1–2 hours.
 - b. Adjust the concentration by adding toluene so that there are 3.006×10^{-5} mol of perovskite crystals per 100 ml of toluene.
- If the fluorescent wavelength band is shifted, Br is replaced with Cl using TMCS (TMS-Cl, (CH₃)₃SiCl) in process 2-1) or 2-2).
- An additional centrifugation step was conducted when TMCS was used.

MAPbBr₃ + PPO

Visible

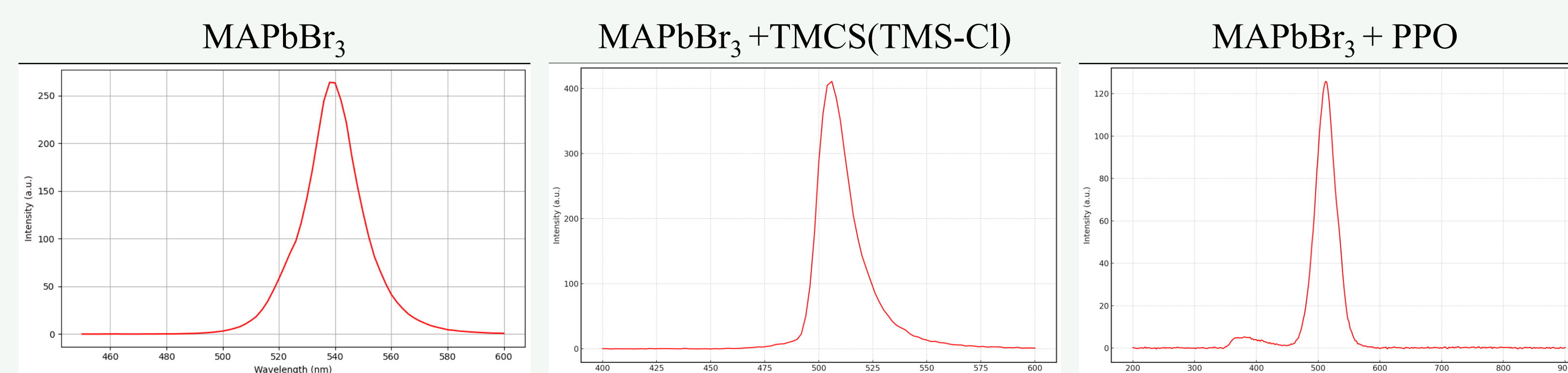


UV365nm



Material Characterization

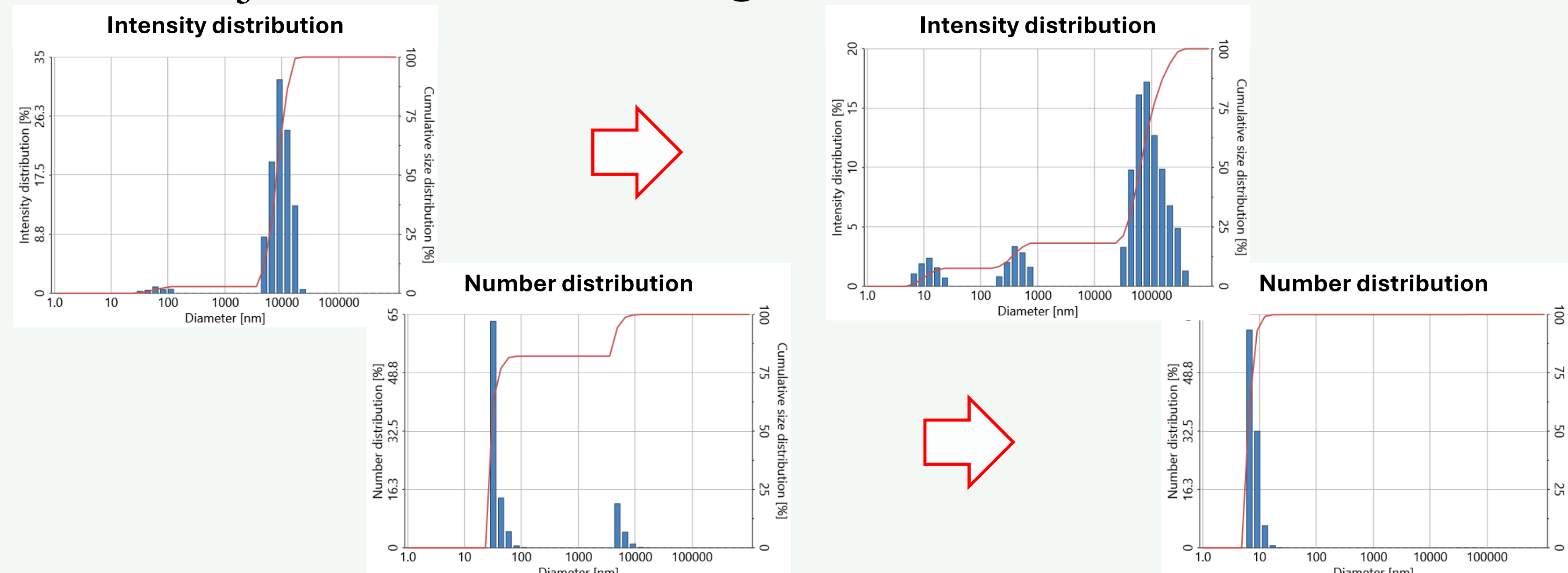
Fluorescence Measurement



- The MAPbBr₃ quantum dots exhibited a strong emission peak near 538 nm in the fluorescence spectrum.
- After the addition of TMCS (TMS-Cl), a similar emission peak was observed around 506 nm, indicating a slight blue shift.
- This suggests that a substitution (Br to Cl) occurred between TMCS and the quantum dots.
- After the addition of PPO, a similar emission peak was observed around 515 nm, indicating a slight blue shift.
- This suggests that energy transfer occurred between PPO and the quantum dots. PPO was added to investigate the energy transfer phenomenon.

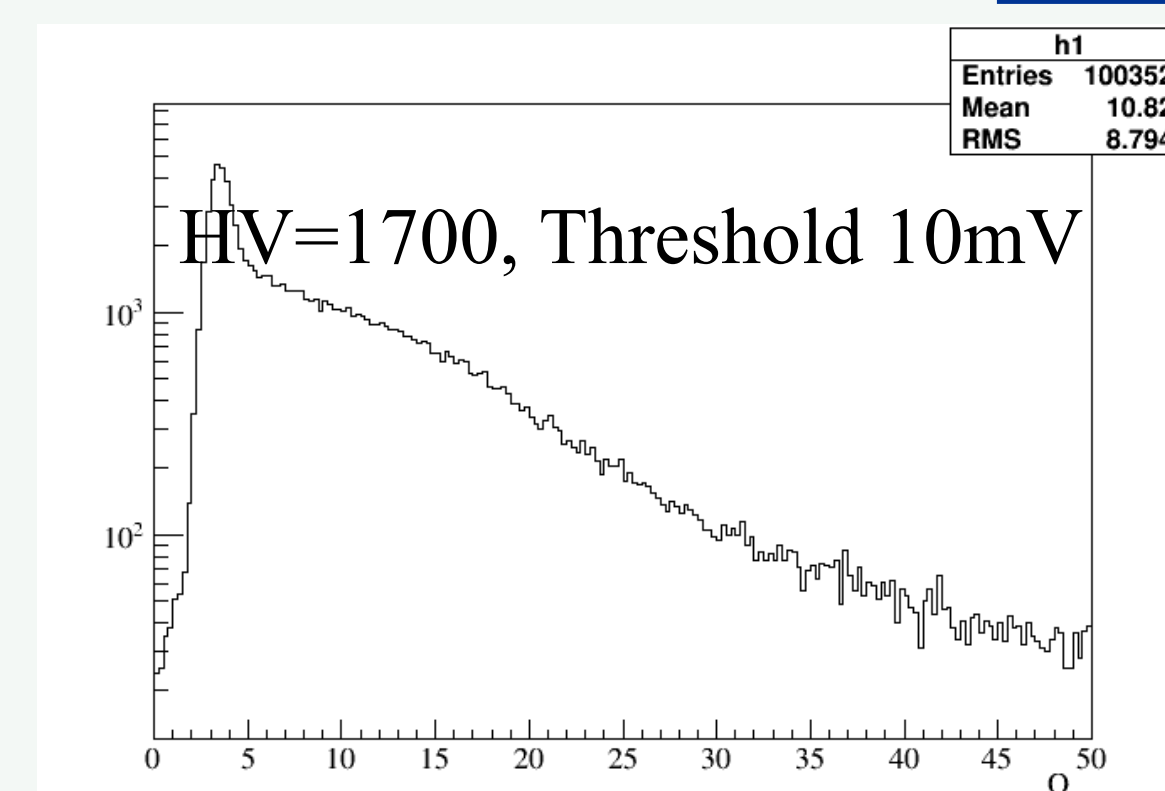
Particle Size Distribution

MAPbBr₃ Before/After Centrifugation



- Before) The particle size is distributed across 50–100 nm and 5000–8000 nm ranges, resulting in relatively large dispersion.
- After) The particle size distribution was reduced to a smaller range (7–15 nm), and the overall dispersion was decreased.

Light Yield Measurement



- ✓ Light yield measurement using a radioactive source(¹³⁷Cs)
- ✓ The Compton edge appears around Q_{tot} values of 18 to 20.

Summary & Future Plan

- ✓ Through the synthesis of PVLS, promising results can be observed as a result of improved stability and consistent light emission.
- ✓ Results showed that the emission wavelength band of PVLS was adjusted using TMCS (TMS-Cl) and PPO (2,5-Diphenyloxazole).
- ✓ Using the centrifugation method, the particle size can be adjusted from 50–100 nm to 7–15 nm to achieve a smaller and more uniform distribution.
- ✓ Subsequent studies will focus on enhancing the scintillation efficiency of PVLS and improving sample stability.

Reference :

William R. LEO, “Techniques for Nuclear and Particle Physics Experiments”, 2 ed, Springer (1994)

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E. M. Kim et al. Poster, KSHEP Fall Meeting, UNIST, 2024.