



Development of a microfluidic device for the study of plasmainduced nucleation

Camille Zindy, C Lafargue, J.-F Audibert, A Salmon, A Brisset, R B Pansu,
Anne Spasojevic - de Biré

► To cite this version:

Camille Zindy, C Lafargue, J.-F Audibert, A Salmon, A Brisset, et al.. Development of a microfluidic device for the study of plasmainduced nucleation. 27th International School on Low Temperature Plasma Physics: Basics and Applications, Oct 2024, Bad Honnef, Germany. hal-04737017

HAL Id: hal-04737017

<https://hal.science/hal-04737017v1>

Submitted on 15 Oct 2024

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Development of a microfluidic device for the study of plasma-induced nucleation

C. Zindy¹, C. Lafargue², J.-F. Audibert³, A. Salmon⁴, A. Brisset⁵, R. B. Pansu², A. Spasojevic¹

¹ Laboratoire SPMS, CNRS, CentraleSupélec, Université Paris-Saclay,

² Laboratoire LuMin, CNRS, ENS Paris-Saclay, CentraleSupélec, Université Paris-Saclay,

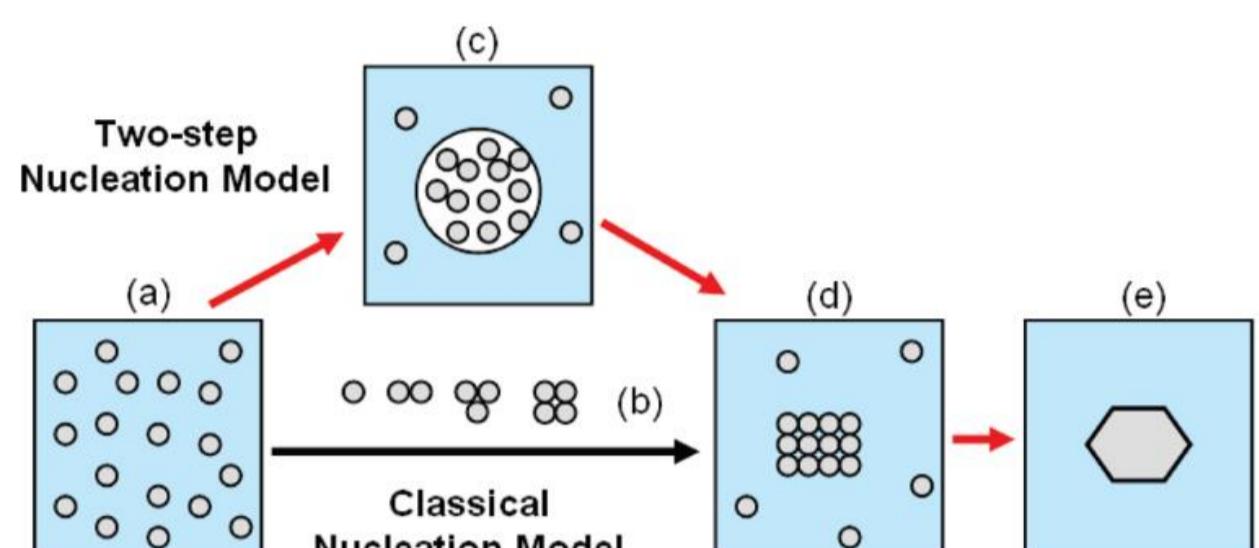
³ Laboratoire PPSM, CNRS, ENS Paris-Saclay, Université Paris-Saclay

⁴ Laboratoire LGPM, CentraleSupélec, Université Paris-Saclay,

⁵ Laboratoire EM2C, CNRS, CentraleSupélec, Université Paris-Saclay,

CONTEXT

Crystallization is a state transition from a liquid or gas phase to an organized solid structure called crystal. It is widely used in industry as separation and purification process. Two main theories explain its mechanism [1].

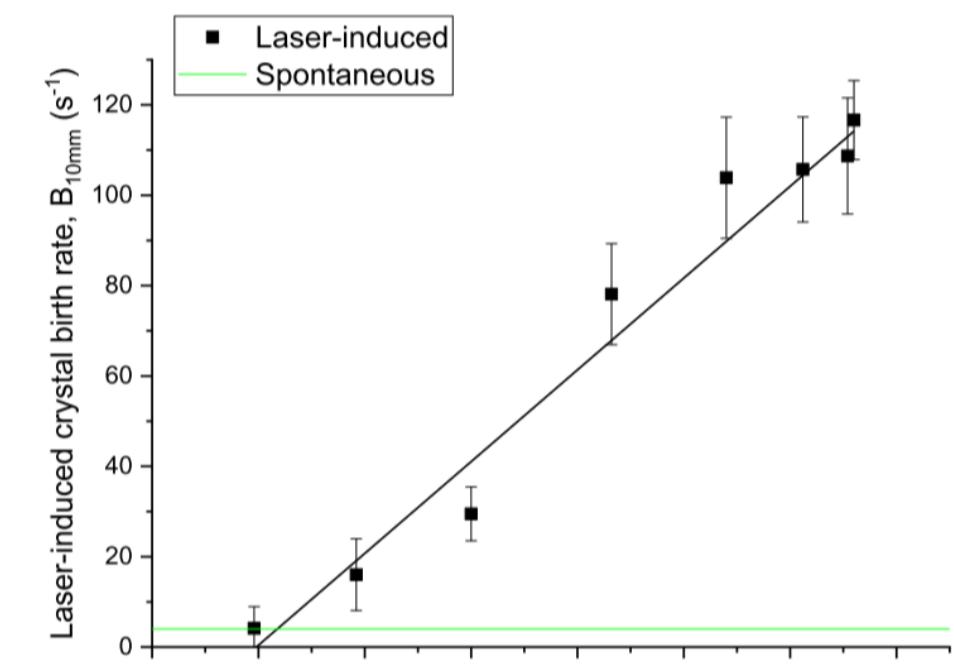
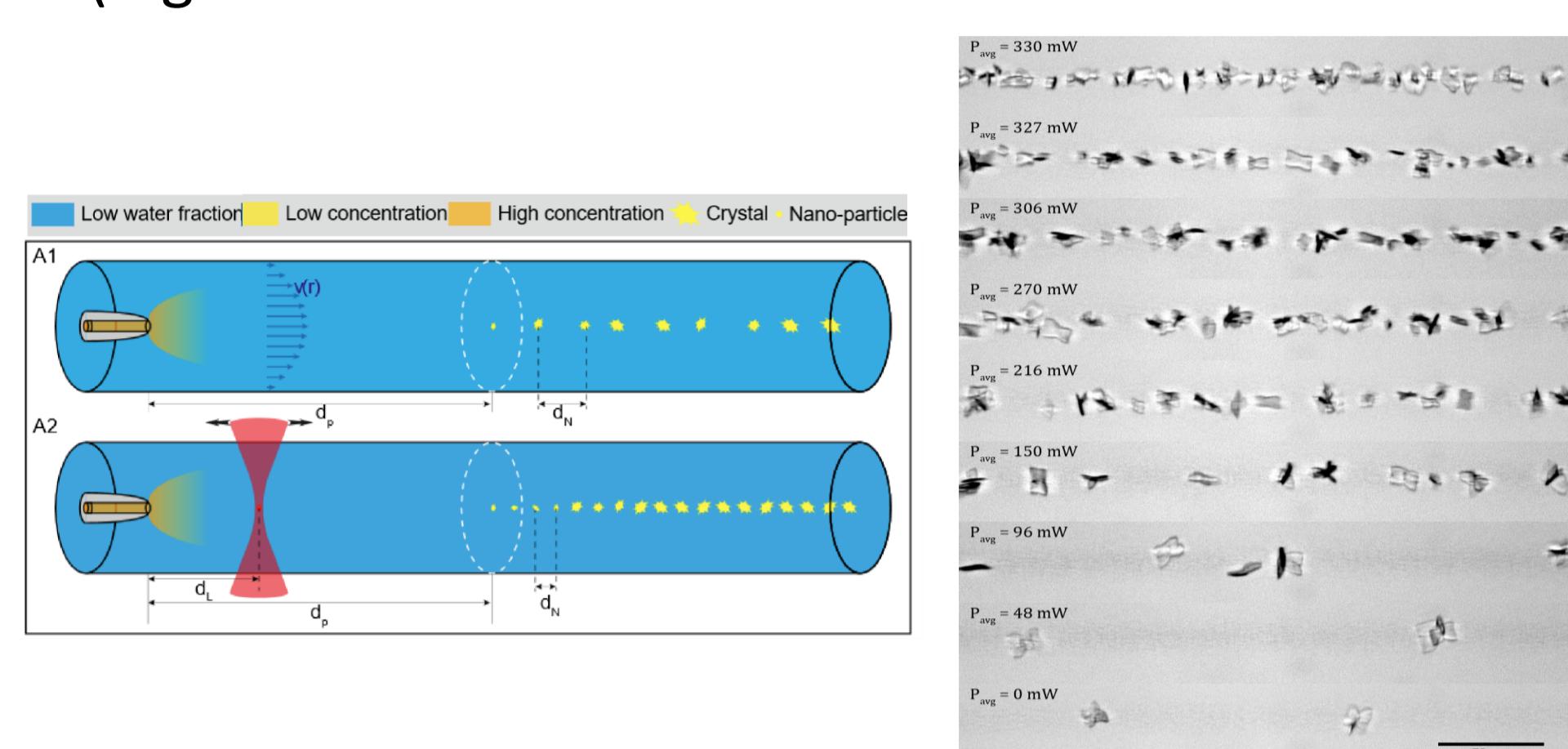


Several crystallization techniques such as precipitation, slow evaporation, grinding exist but those don't allow any control on nucleation and don't give any information about where and when nucleation is initiate.

There is a need for **in situ, on-demand and scalable seeding** techniques allowing a high level of polymorph and size distribution controls.

PREVIOUS WORKS

Nucleation can be **triggered by external fields** with a better control over it. (e.g. laser induced nucleation NPLIN increases the nucleation rate [2])



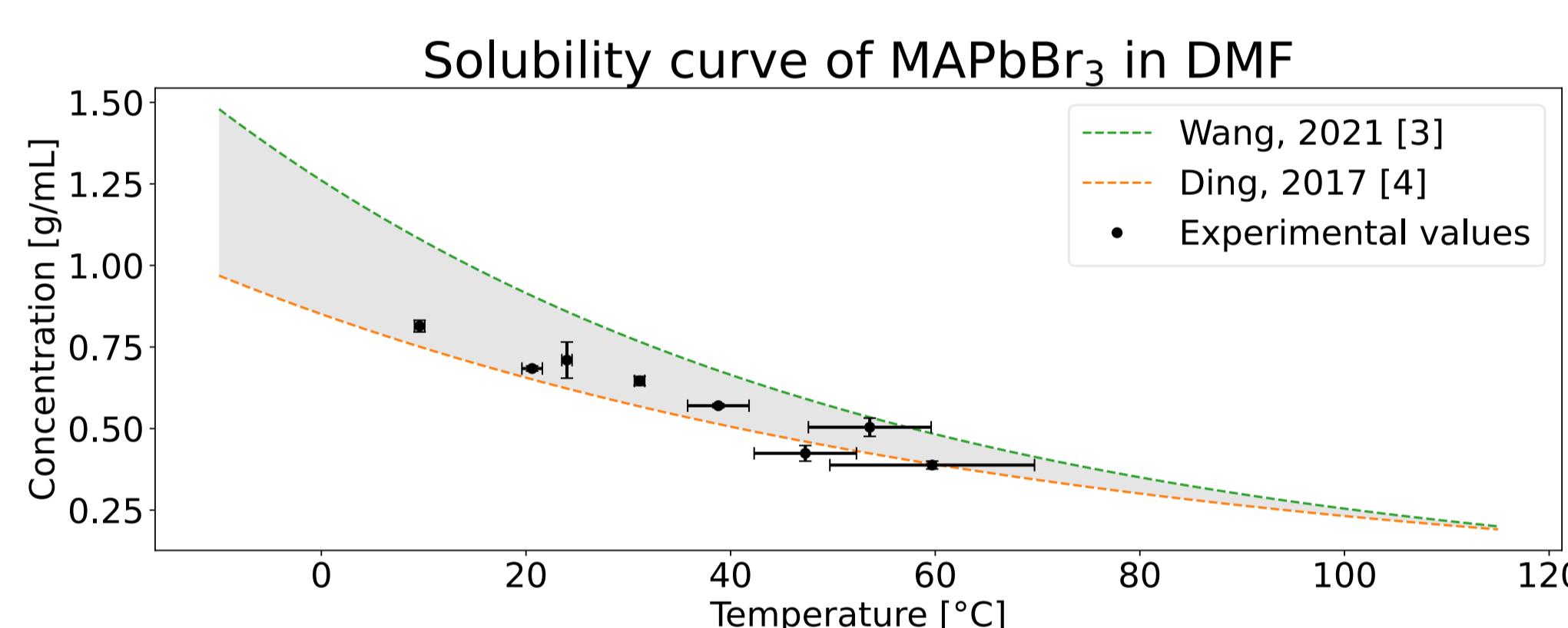
Recent studies in CentraleSupélec show that **cold plasma** can induce and control nucleation with an even greater efficiency.

METHODOLOGY

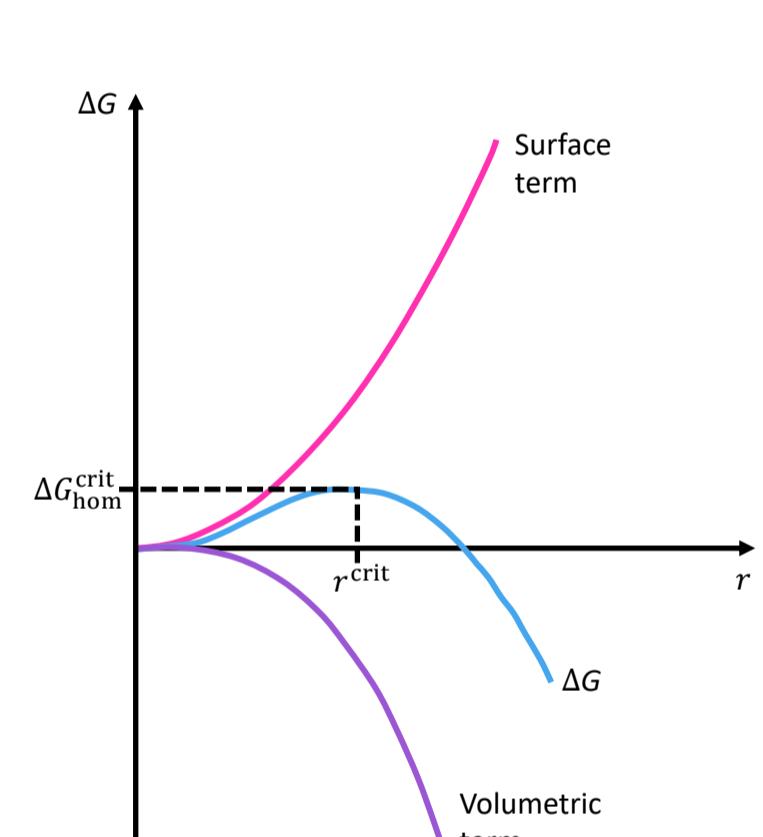
Nanosecond pulsed discharges will be applied on a solution of supersaturated MAPbBr_3 perovskites in N,N-Dimethylformamide (DMF). Nucleation mechanisms will be studied in a microfluidic device by epifluorescence; and several plasma diagnosis. First stages are to characterize the solution and develop the microfluidic system.

SOLUTION CHARACTERIZATION

The **solvability diagram** is necessary to design a crystallization cycle.



Solubility limit is a key information to control the solution's **supersaturation** (concentration on concentration at solubility).



Gibbs free energy :
(energy from supersaturation)

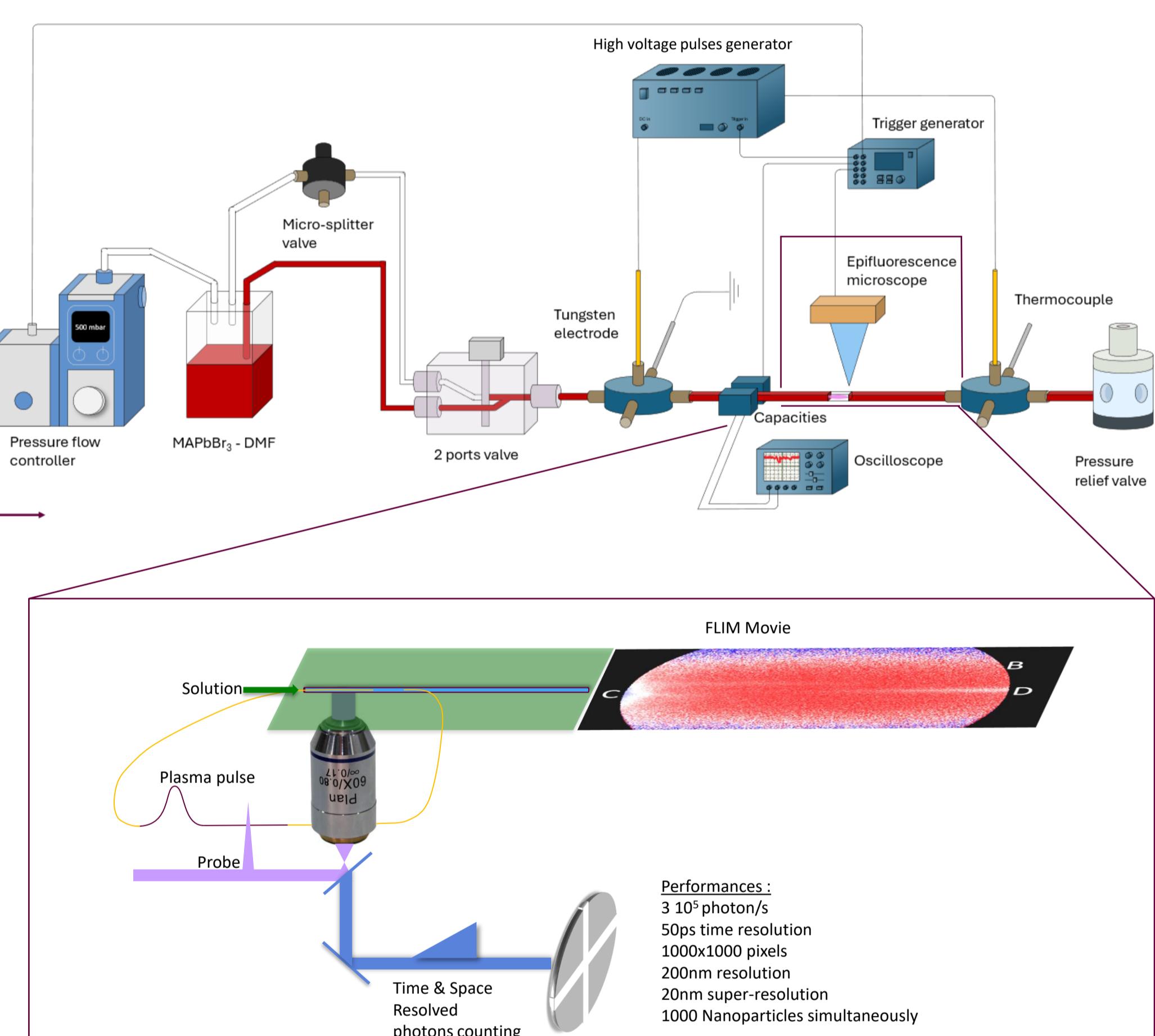
$$\Delta G = -\frac{4\pi r^3}{3V} \Delta\mu + 4\pi r^2 \gamma$$

Volumetric term Surface term

The interfacial energy γ between germ and liquid works as a **potential barrier** preventing the formation of the nucleus.

The critical size of a cluster r_{crit} depends on the **supersaturation** of the solution.

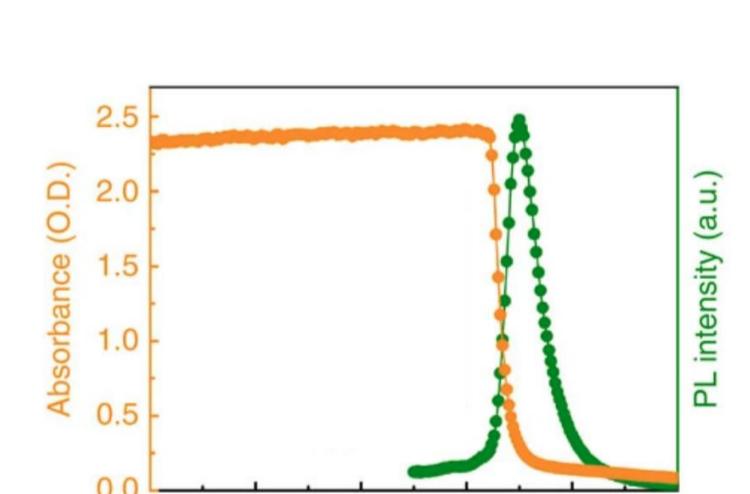
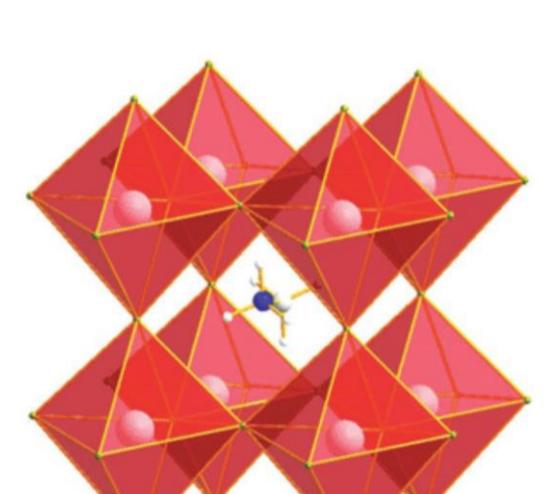
MICROFLUIDIC EXPERIMENTAL SETUP



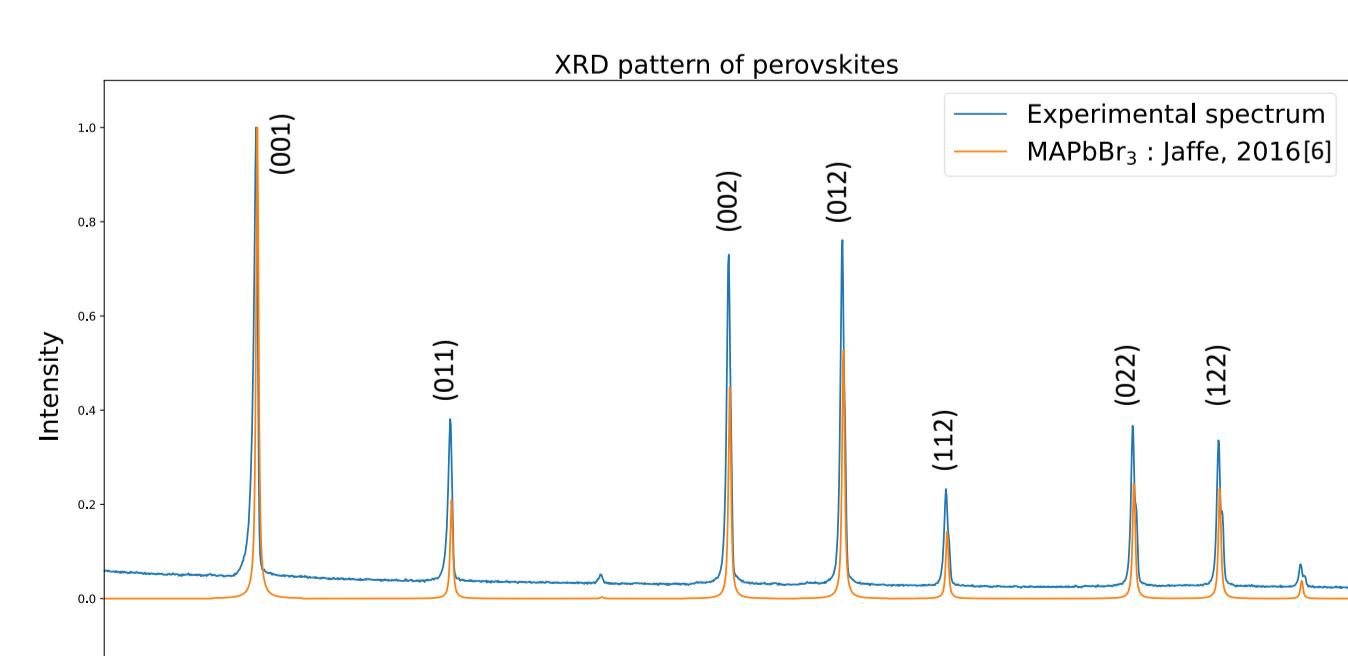
RESULTS

CRYSTALS

MAPbBr_3 ($\text{MA}=\text{CH}_3\text{NH}_3^+$) perovskites are fluorescent in solid form but not when dissolve in DMF.

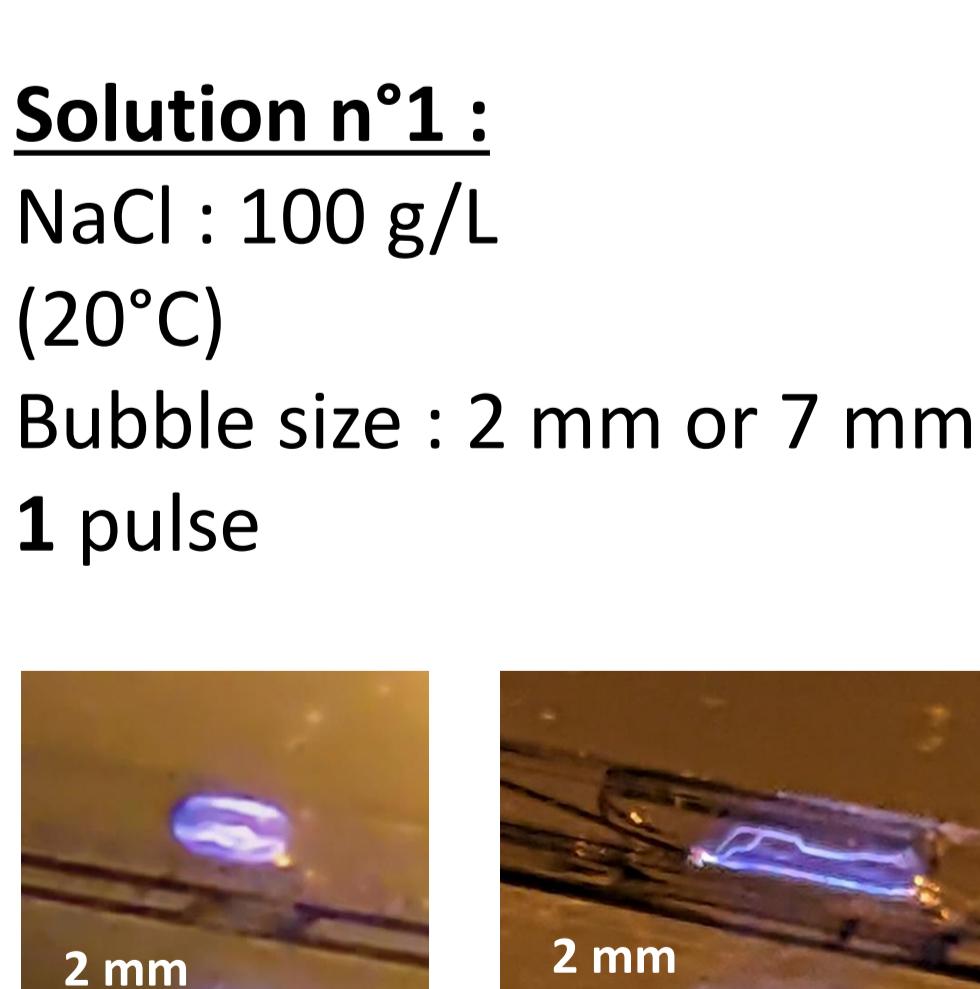


MABr and PbBr_2 powders in a DMF solution crystallize as MAPbBr_3 perovskites.

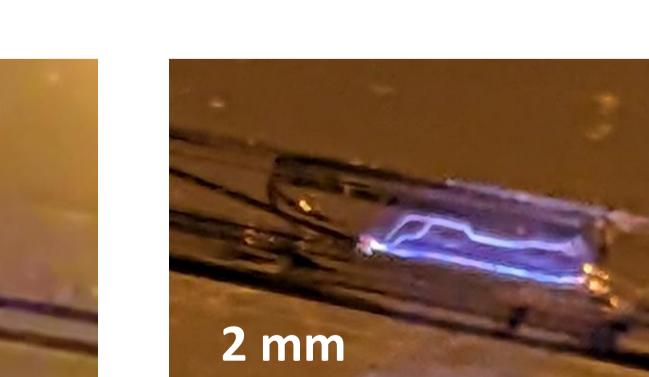


PLASMA

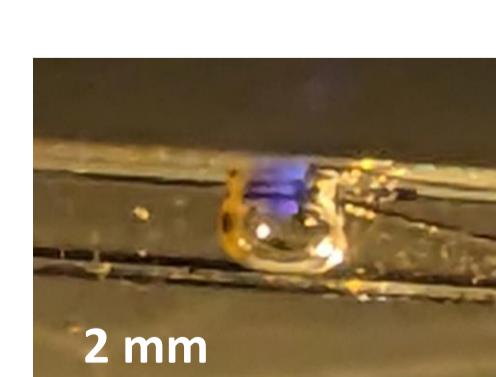
We apply 10 kV nanosecond discharges at 10 kHz in different solutions to trigger plasma discharges.



Solution n°1 :
 $\text{NaCl} : 100 \text{ g/L}$
(20°C)
Bubble size : 2 mm or 7 mm
1 pulse



Solution n°2 :
 $\text{MAPbBr}_3\text{-DMF} : 0,6 \text{ g/mL}$
(S=90%)
Bubble size : 1 mm
30 pulses at 10 kHz



CHALLENGES

- Having a regular bubble train for a future automatization of the process.
- Triggering the plasma discharge with only 1 pulse.
- Reduce electromagnetic plasma radiations.
- We can have a DBD discharge between the electrodes in and outside of the capillary.



ACKNOWLEDGMENTS

ANR INCRYS | ANR-23-CE51-0033

References

- Erdemir, D. et al., Acc. Chem. Res. 42, 5 (2009) 10.1021/ar800217x
- Zhang, Z., Laser-Induced Nucleation in a Coaxial Microfluidic Mixer (2019)
- Wang, H. et al., Mater. Sci. Semicond. Process., 135 (2021) 10.1016/j.mssp.2021.106107
- Ding, J. et al., J Mater Sci. 52 (2017) 10.1007/s10853-017-0995-8
- Saidaminov, M. I., et al., Nature Comm., 6(1) (2015) 10.1038/ncomms8586
- Jaffe, A. et al., ACS Cent. Sci., 2, 4, 201–209 (2016) 10.1021/acscentsci.6b00055