

# New insights into colloidal phase transitions using neutron scattering techniques



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## TECHNIQUES

DLS & SEM, RMMF



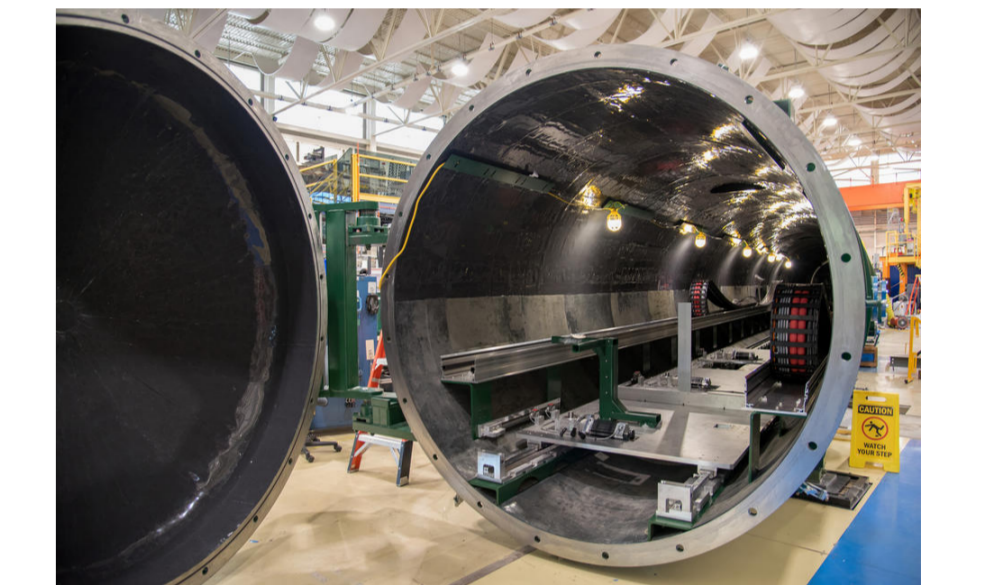
SAXS, RMMF & Australian Synchrotron



U/SANS, ANSTO



VSANS, NIST



## INTRODUCTION

- Fundamentals of **crystallisation** and **glass transition** are not fully understood.
- Binary colloidal suspensions** are a good model.
- Brownian, real-time motion.
- Limited studies on **dynamics** and **structure**. [1-3]

## RESEARCH QUESTIONS

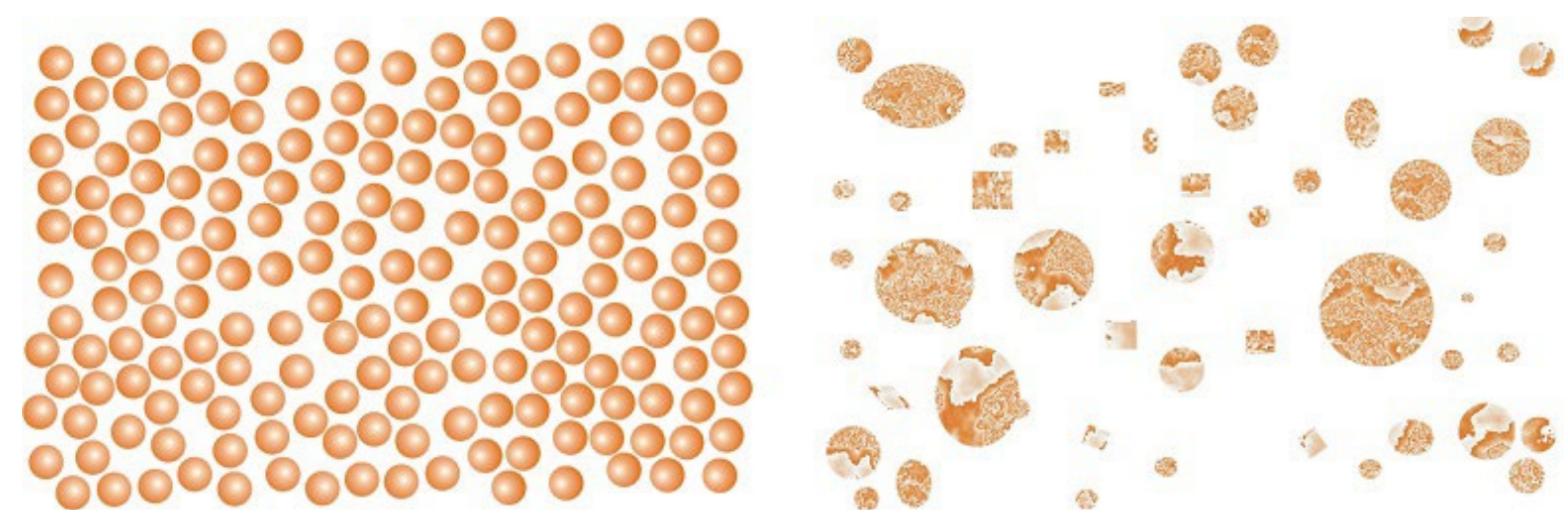
- What **combination of nanoparticle and solvent** will be the ideal colloidal system to study?



- How do **different volume fractions  $\phi$**  affect the crystallisation and phase behaviours of the system?

$$\phi = \frac{\text{Volume}_{\text{particle}}}{\text{Volume}_{\text{total}}}$$

- How does **change in polydispersity** affect the crystallisation and phase behaviours of the system?



- What **resolution** do we need to measure the **full range of scattering vectors  $q$**  during solidification processes in our neutron scattering experiments?



## CHOOSING THE IDEAL SYSTEM

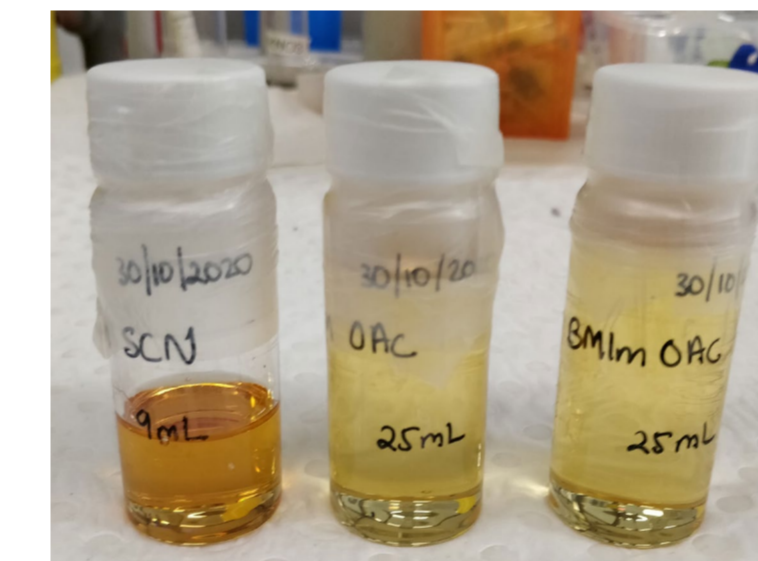
### Particle

- Large particles, relatively cheap, non-volatile. Testing: **Silica** ( $d = 20\text{-}2470$  nm) and **Polystyrene** ( $d = 62\text{-}2500$  nm)

### Solvent

- Refractive index  $n$  match, similar density  $\rho$ , transparent, relatively cheap, reasonable viscosity  $\eta$ . Potentials: ionic liquids and deep eutectic solvents.

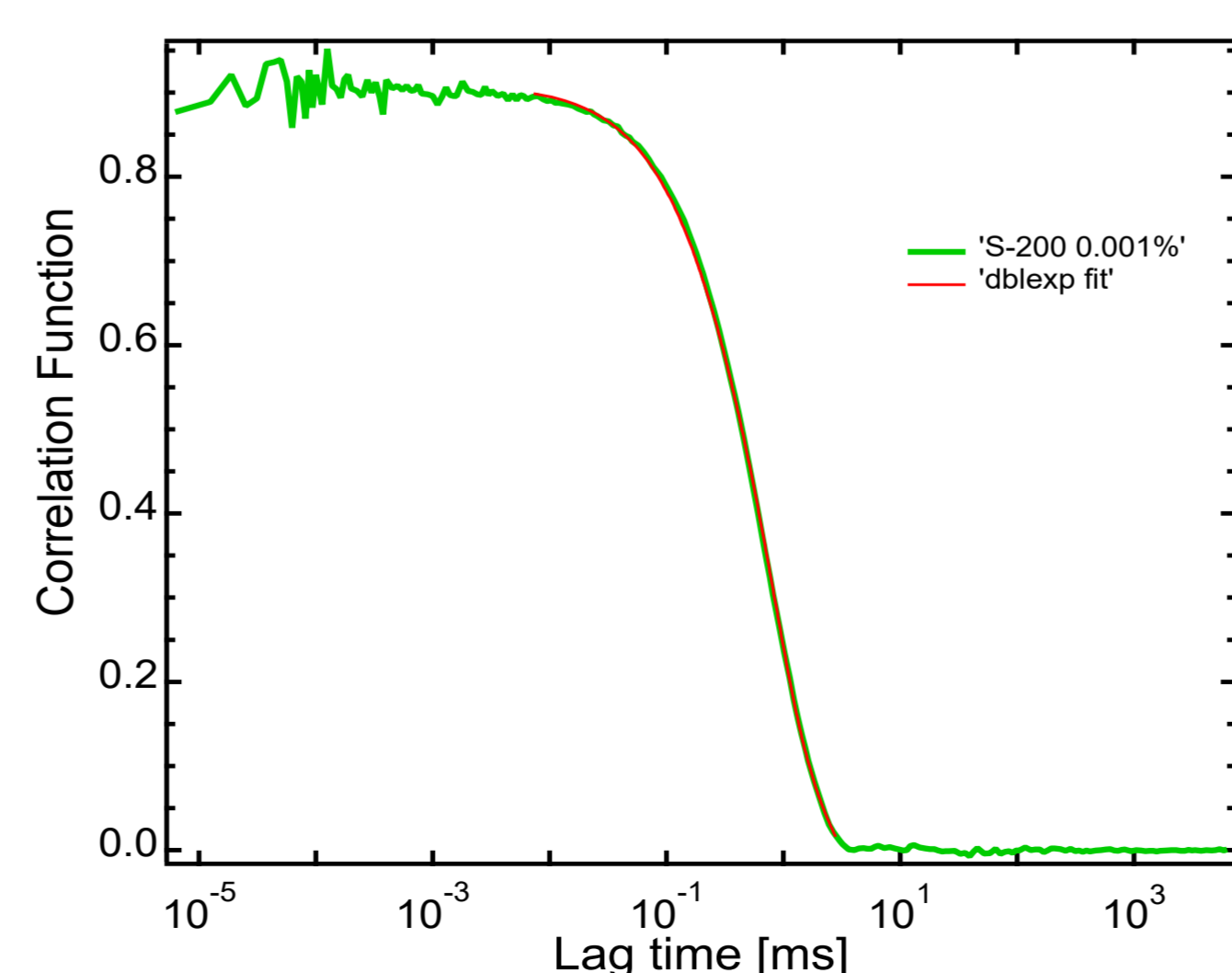
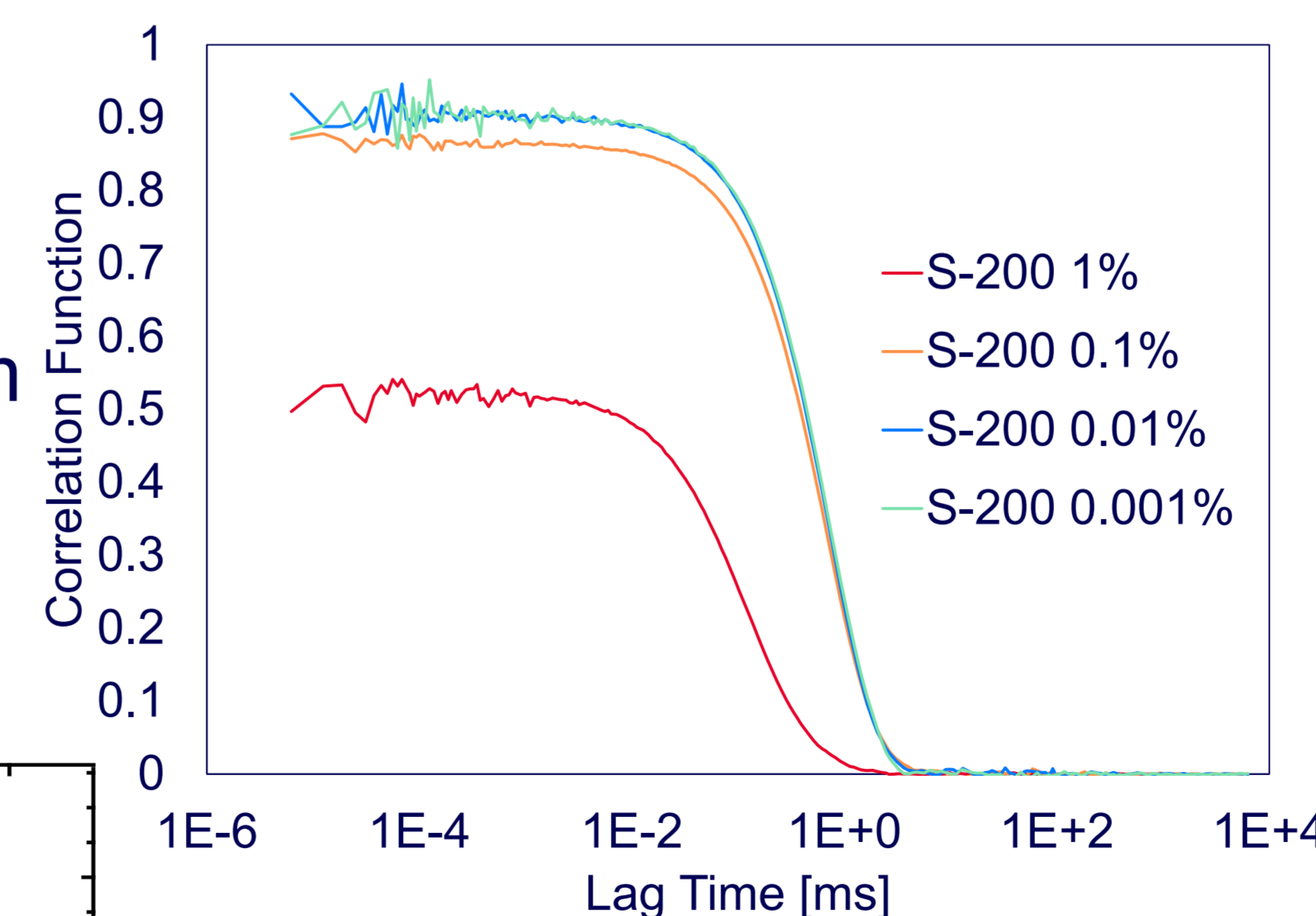
Testing: **BMIm Acetate** and **BMIm Thiocyanate**



## PRELIMINARY RESULTS

- Checking particles are stable with DLS.

**Fig. 1** Light scattering data for silica  $d = 200$  nm at four concentrations.

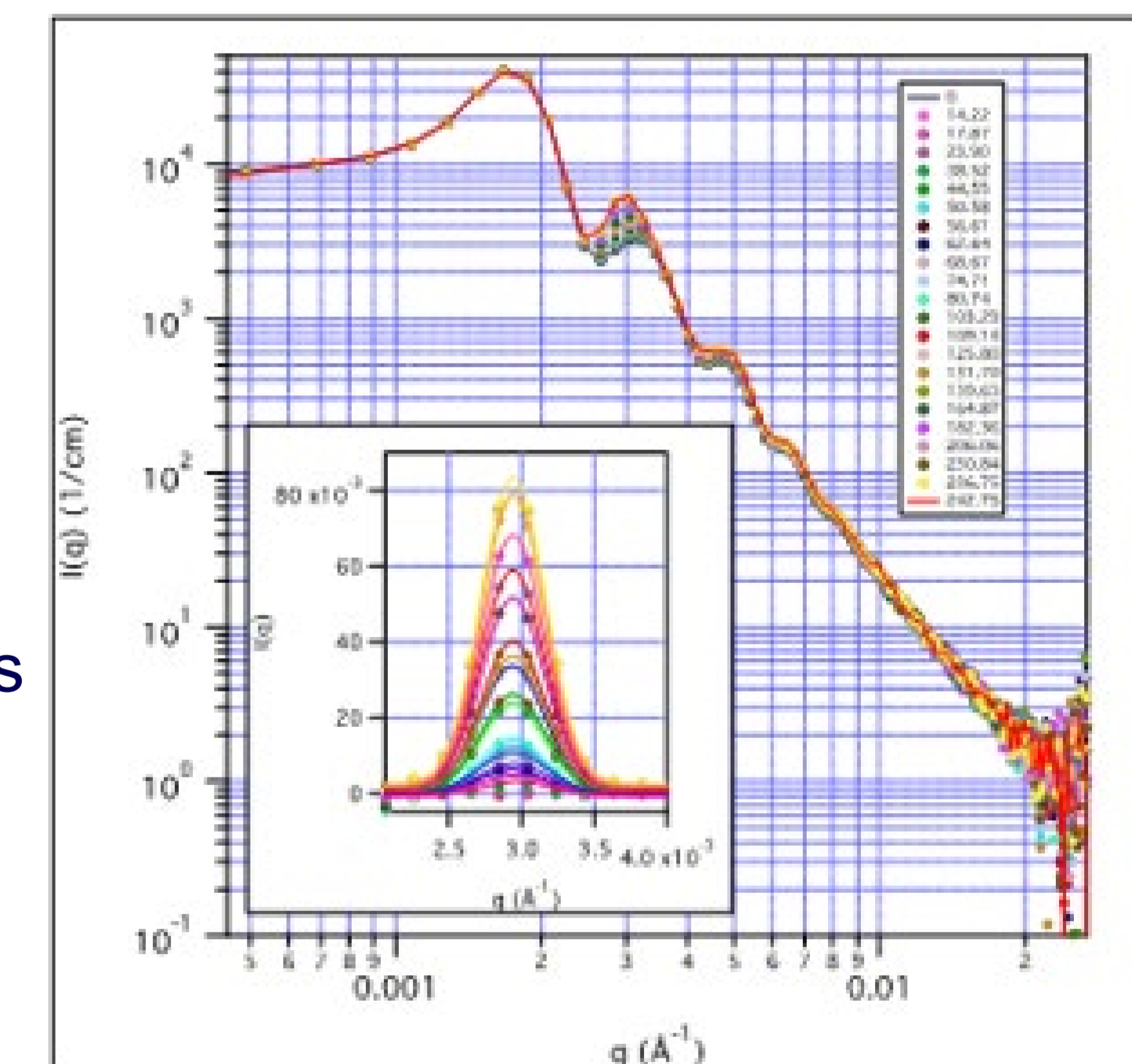


**Fig. 2** Double exponential fit of dataset for 0.001% concentration of silica (S-200 0.001%).

## PREVIOUS WORK [4-8]



**Fig. 3** SANS data for SMU39 at a volume fraction of 0.54 over a period of ~250 minutes following the quench. The inset highlights the growth of the main crystalline peak with Gaussian fits.



## REFERENCES

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