

Newsletter

Combined Imaging and in plate Dynamic Light Scattering

XtalConcepts GmbH Hamburg, Germany

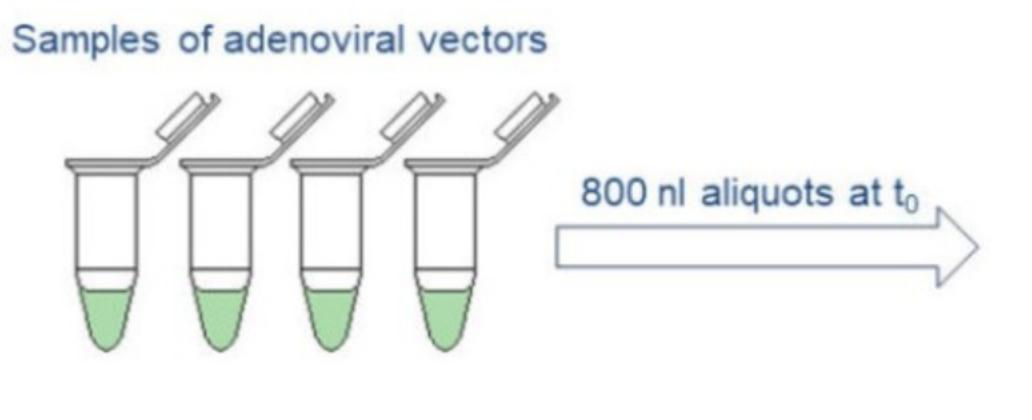


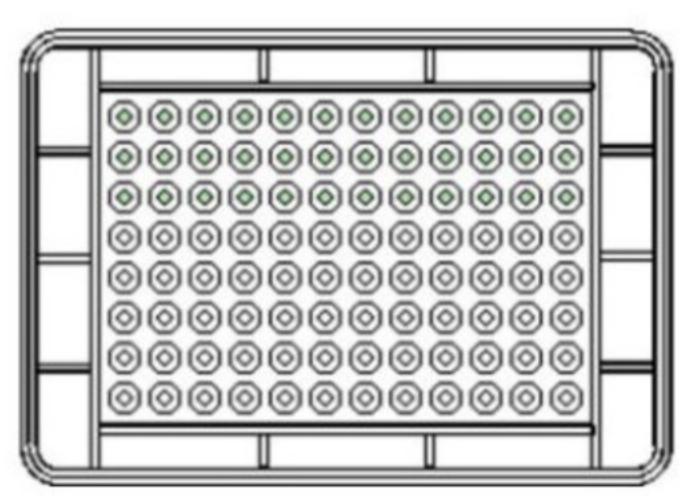
Please find more information about the system on our website: www.xtal-concepts.de

The Sample: Adenoviral mRNA Vectors

Preparation of various Adenoviral Vectors for in plate DLS Analysis

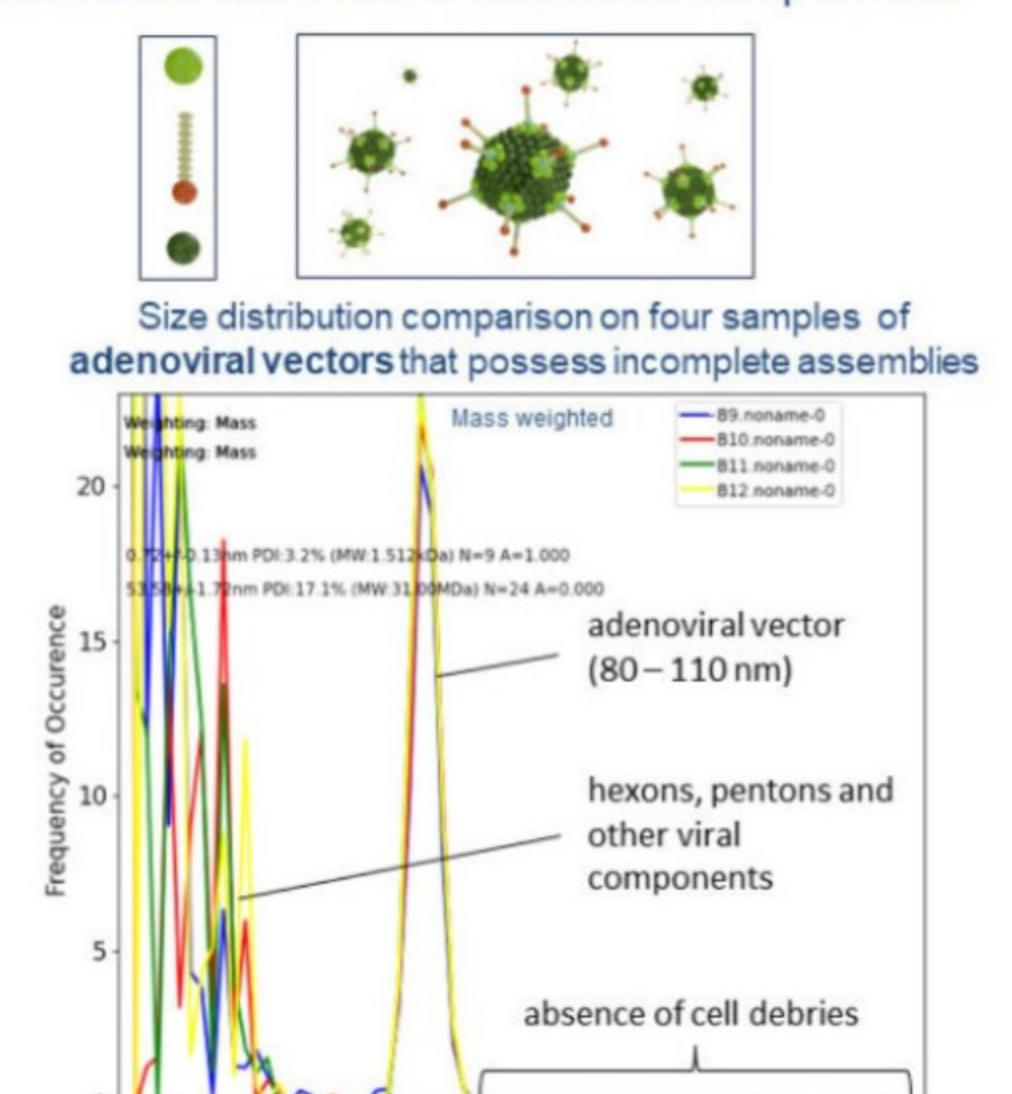
800 nl aliquots from several batches were transferred to a 96-well Douglas Instruments plate and sealed with paraffin oil.





Vector Purity Assessment based on Size Distribution Analysis

Mass-weighted size distribution monitoring of the ratio of fully assembled viral vectors to residual components.

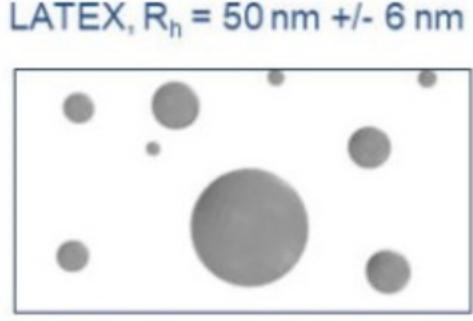


100nm Radius (nm)

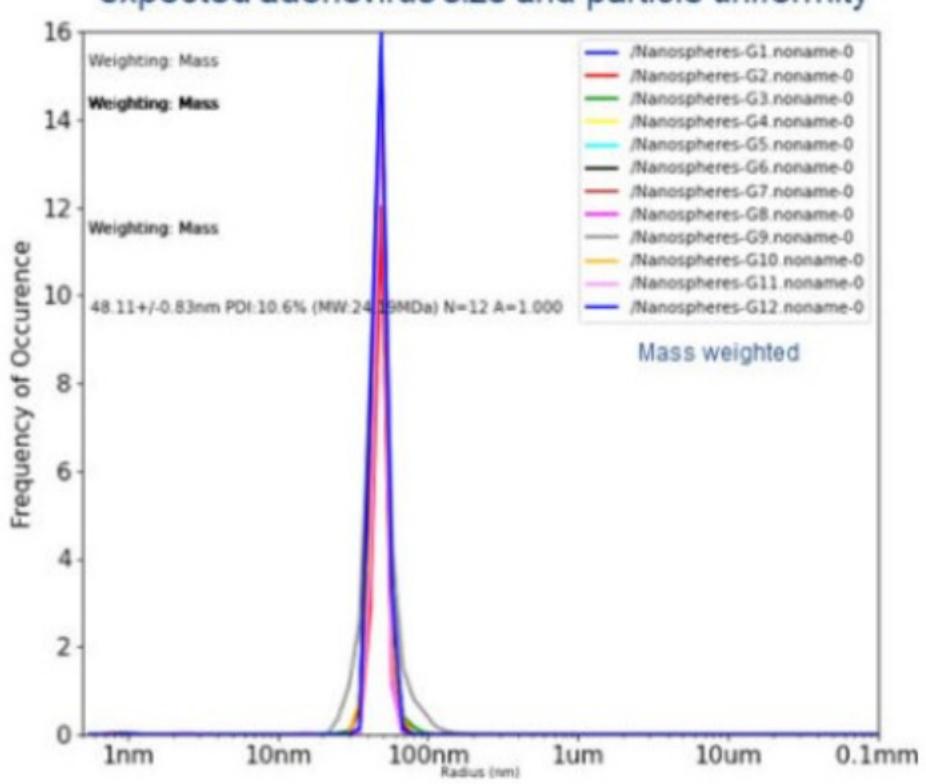
lum

10nm

lnm



Control: LATEX Nanosphere Size Standard of the expected adenovirus size and particle uniformity

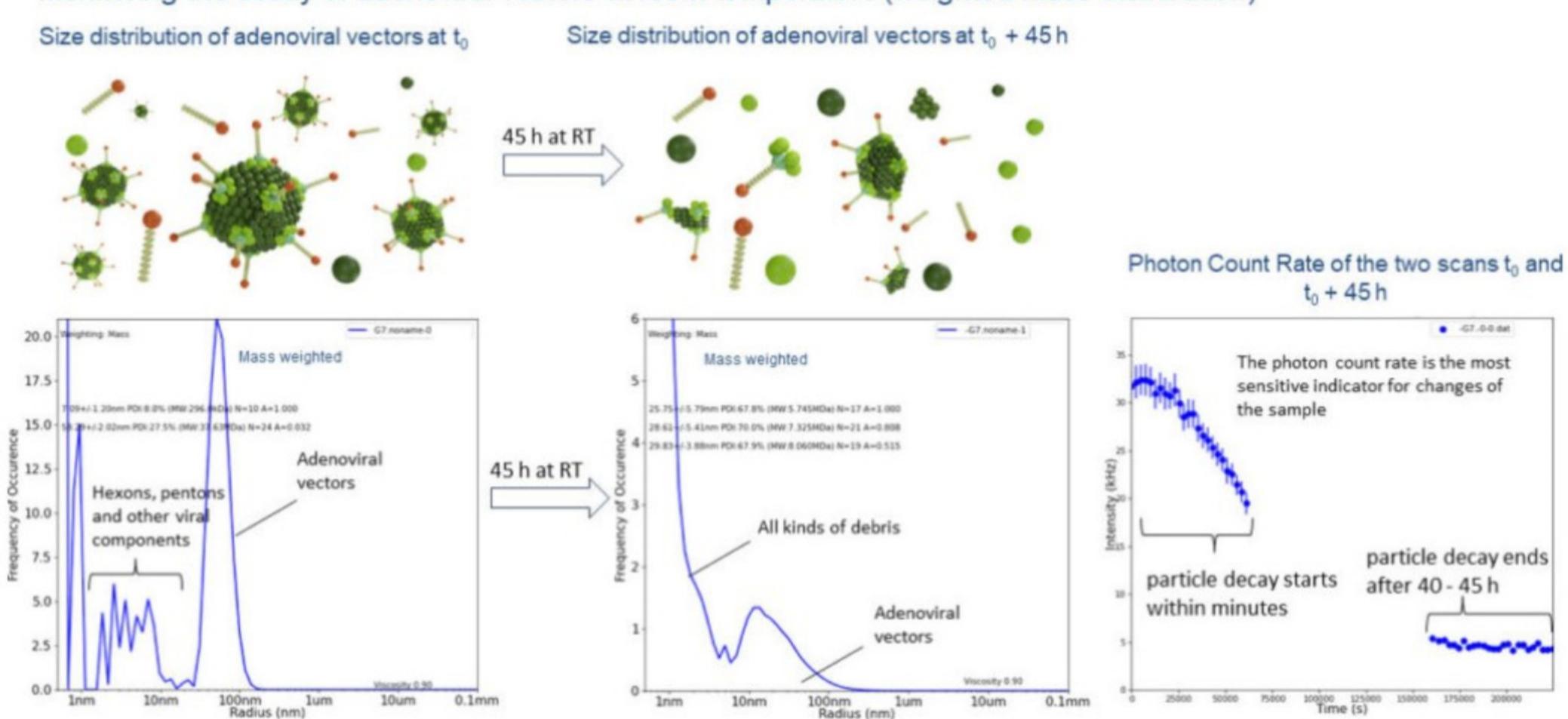


Size Distribution Signature of an instable adenoviral Vector when incubated at Room Temperature

0.1mm

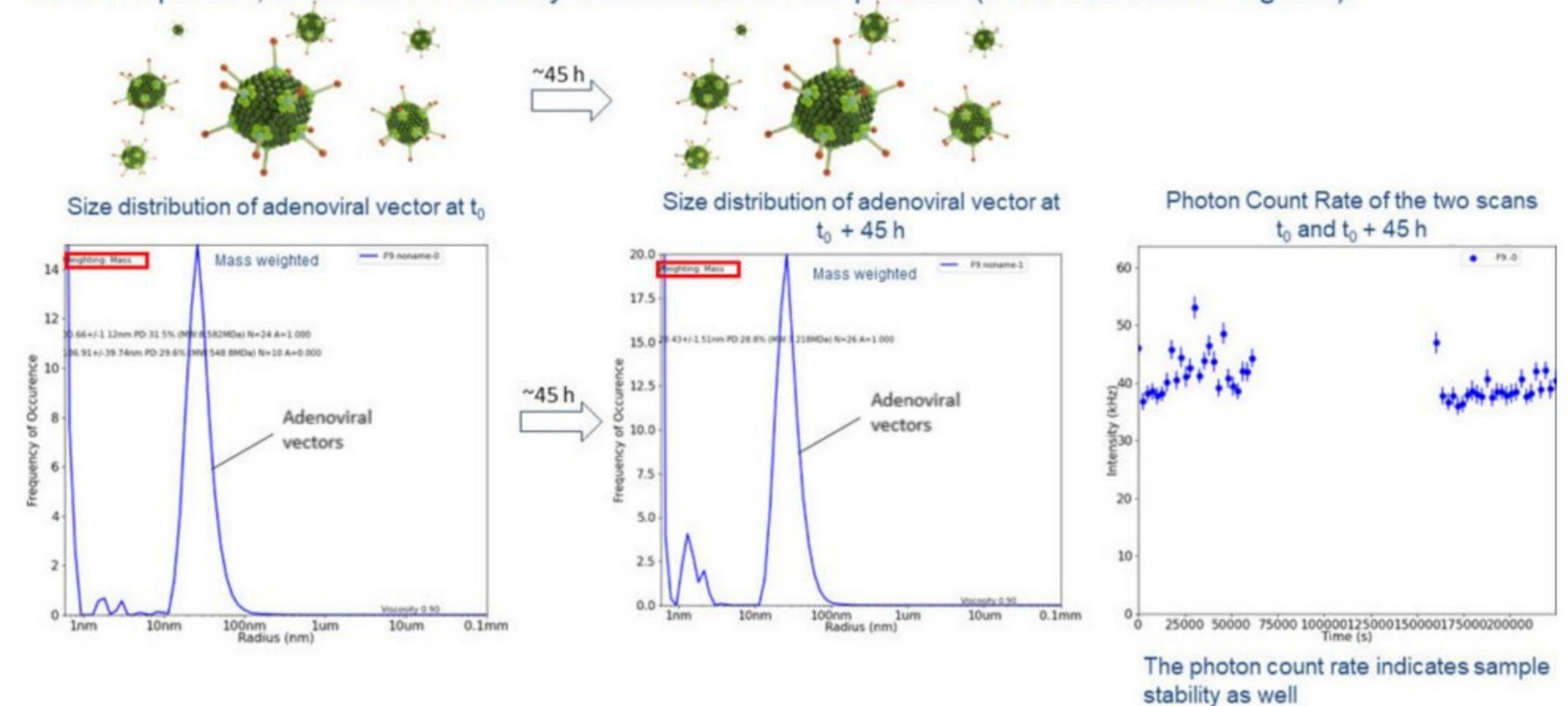
Monitoring the decay of adenoviral vectors at room temperature (weighted mass distribution)

10um



Size Distribution Signature of a stable adenoviral Vector when incubated at Room Temperature

As a comparison, some vectors are very stable at at room temperature (mass distribution weighted)

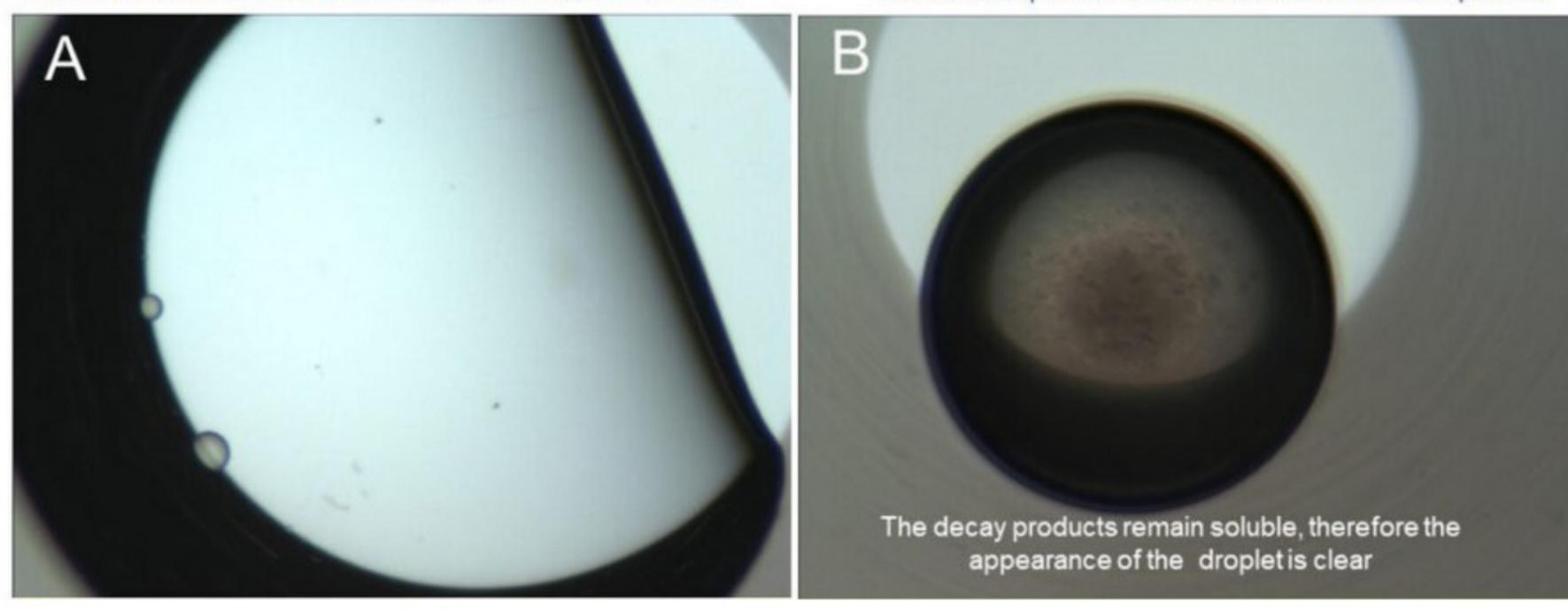


Visual Inspection of two instable adenoviral Vectors

The degradation products remain soluble, so the appearance of the droplet is clear (A). Other adenovirus samples were full of colloidal impurities (B).

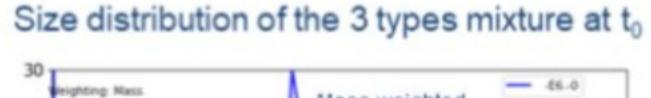
Adenoviral vector sample A, bright light image at to + 45 h

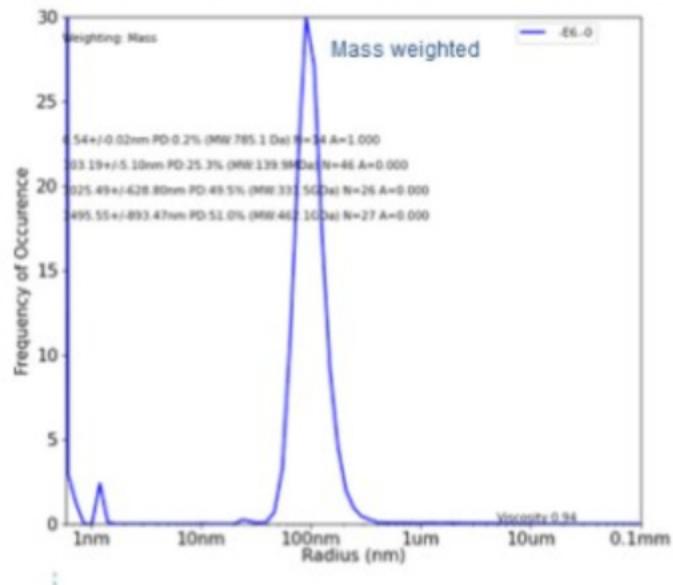
Adenoviral vector sample B: Bright light image of another sample with debris and other colloidal impurities



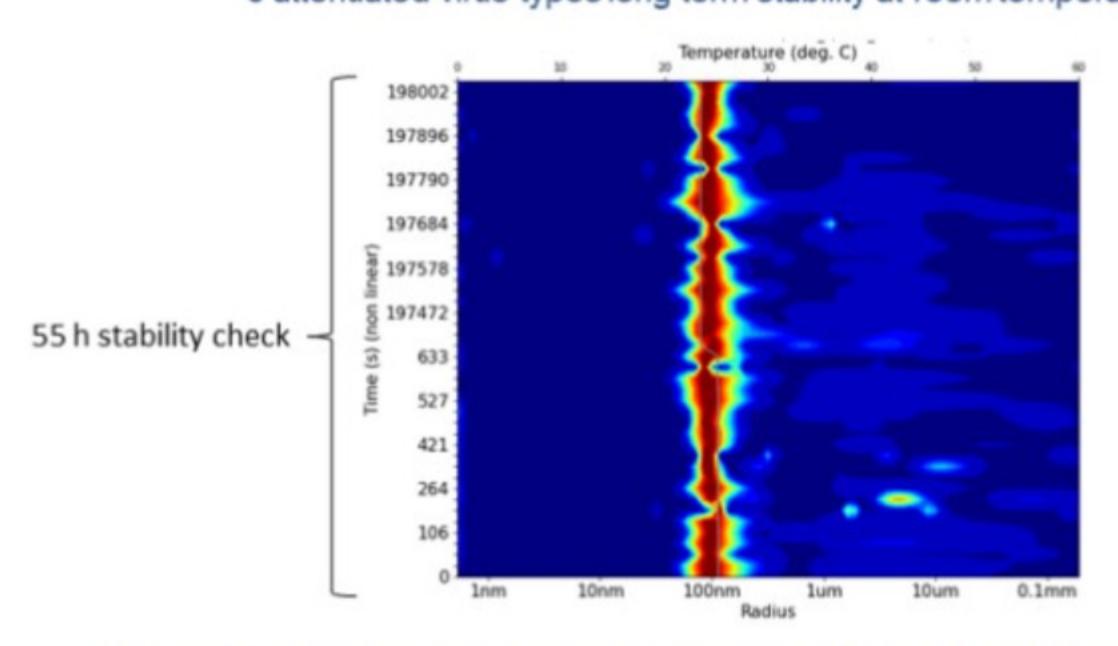
Control, a Mixture of three stable attenuated Viruses also incubated at Room Temperature

- Mumps virion ~150 nm
- Measels virion diameter ~150 nm
- German Measels virion diameter 50-70 nm





3 attenuated virus types long term stability at room temperature



In this plot the particle distribution is not weighted. Instead, the distribution is based on the intensity of the scattered light, which is strongly size dependent. Larger particles are therefore visible. However, their mass fraction is negligible.

Plate scanning was fully automated. The identity of smaller particle impurities was subsequently determined by other methods. They were identified as hexons and pentons as well as other capsid components such as fibres etc.

Please visit our website for more information on the incredible possibilities of in situ and in plate DLS,

Your Xtal-Concepts Team