DynaPro Plate Reader III

Automated characterization of size, stability and molecular weight in industry-standard microwell plates





DynaPro Plate Reader III Dynamic and Static Light Scattering in Microwell Plates

Analyze Size, Screen Stability

Dynamic Light Scattering (DLS) is a core technology for any lab engaged in nanoparticle sizing or biotherapeutic stability studies.

Most DLS instruments require tedious, one-at-atime sample analyses by manually inserting and replacing cuvettes. The DynaPro Plate Reader III eliminates virtually all of the manual labor involved in measuring many different samples; it even enables fully-automated screening of processing or formulation conditions by making measurements in standard well plates.

Applications

- Develop stable biopharmaceuticals
- Optimize protein buffers for crystallization, SAXS or chromatography
- Formulate theranostic nanoparticles
- Discover small-molecule and peptide inhibitors of protein-protein interactions

Easy, automated all-in-one

With automated, in-plate capabilities you can perform experiments and carry out novel studies you wouldn't have imagined before.

- Acquire in one day the data that would otherwise take weeks
- Screen dozens of samples with thousands of formulation and temperature combinations
- Analyze and visualize an entire dataset at once, then zoom in for a detailed study of the most promising conditions
- Export your results with a single click
- Measure non-destructively and transfer plates to other analytical instruments

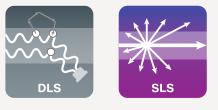
In-plate analysis saves you time and money!

- Measure directly in 96, 384 or 1536 well plates, simply load and walk away
- Microwell plates are disposable and less expensive per sample than disposable cuvettes
- Integrate with plate and liquid-handling robots for even more time savings

Maximize Characterization with DLS and SLS

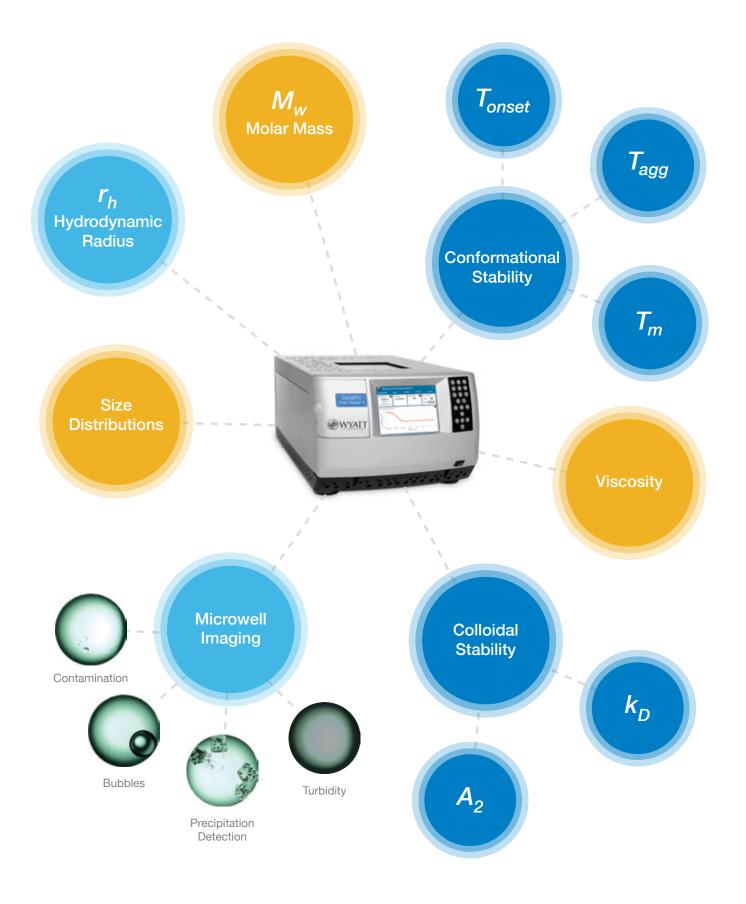
DLS is widely used to characterize proteins, nanoparticles, colloids and macromolecules from subnanometers to several micrometers. Requiring relatively small amounts of material, DLS—along with the new static light scattering (SLS) capabilities helps assess key factors:

- Size (*r_h*) and size distributions
- Molar mass (M_w)
- Aggregation and stability indicators (T_m, T_{agg}, k_D, A₂)
- Purity or contamination, turbidity

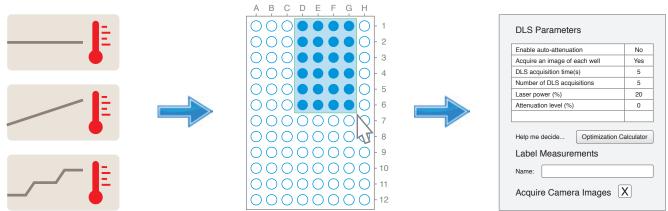




Automated All-in-One



Design High-throughput Experiments



1. Select temperature profiles Combine multiple profiles

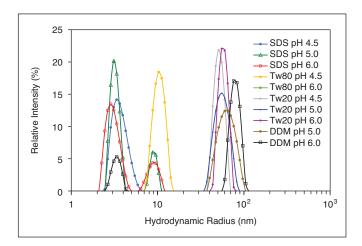
for complex protocols.

2. Select wells Include replicates and control samples.

3. Finalize design

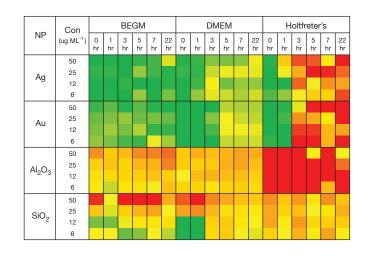
Fine-tune parameters, add camera images.

Identify Optimal Samples



Which process creates the ideal nanoparticle size range?

Overlay size distributions from multiple wells.



In which media are the particles most stable?

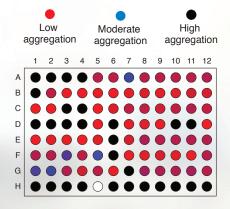
Immediately visualize an aggregation time course.

Advantages

- Compatible with industry-standard 96, 384 or 1536 microwell plates
- > Enhanced thermal isolation prevents condensation
- > First and only plate reader to provide molar mass
- > Sample volumes as low as $4 \mu L$
- > Measure from 4 °C to 85 °C
- > Infrared wavelength not susceptible to fluorescence
- > 21 CFR Part 11 compliant software

70 **F**5 B3 10 -60 8 50 % Intensity 100 nm 3.5 nm % Intensity 40 6 30 4 3.5 nm 20 2 10 0 0 -10² 10⁴ 10² 10⁴ 0.01 0.01 1 1 Radius (nm) Radius (nm)

Aggregation in a 96 well plate



Explore content-rich screening data and tackle comprehensive DLS experiments you never thought possible

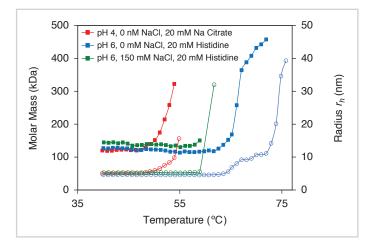
Weeks of Experiments in Just Hours

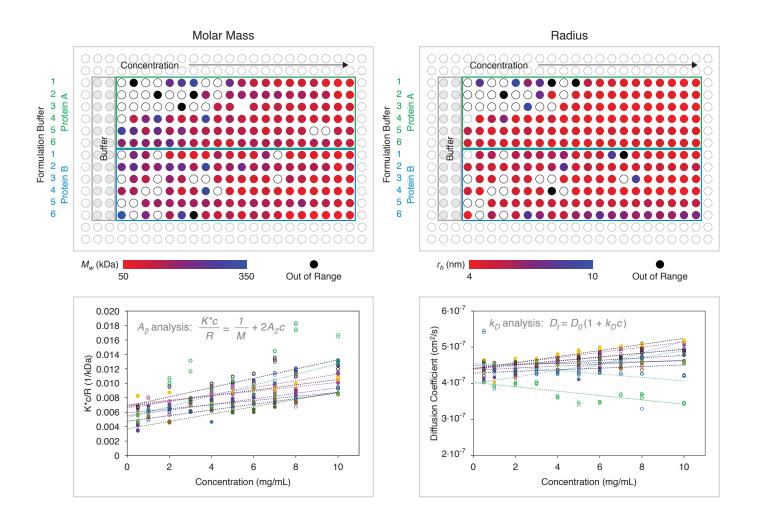
Biologics: Thermal and Colloidal Stability

The DynaPro Plate Reader III performs high-throughput screening of biotherapeutic candidates to determine multiple properties of a plateful of candidates and formulations. Shown to the right and below is an analysis of proteins for thermal and colloidal stability.

Right: The thermal stability of three formulations of an IgG is determined, in parallel, through changes in size (T_{onset} by DLS) and molar mass (T_{agg} by SLS) across a temperature ramp. Filled squares – molar mass; empty circles – hydrodynamic radii. One formulation exhibits multiple transitions.

Bottom: Analysis of aggregation and colloidal stability for two proteins via 10-point concentration series with two replicates at each concentration and condition. Size and molar mass are indicated in the SpectralViewTM heat maps, while plots of second virial coefficient A_2 and diffusion interaction parameter k_D , two measures of colloidal stability, are shown below.

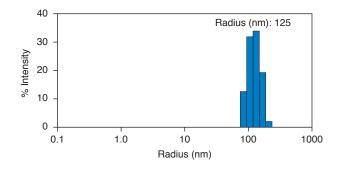




Nanoparticles: Process Optimization

Development of nanoparticle manufacturing processes is greatly accelerated when a matrix of processing conditions is combined with high-throughput particle sizing in the DynaPro Plate Reader III.

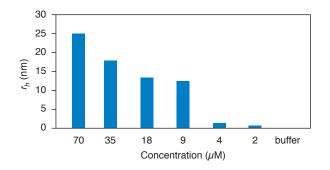
In this example, acoustic resonance milling of drug solids to nanoparticle size was performed in 96 well plates and samples aliquoted periodically to a 384 well plate for analysis. Conditions tested included surfactant type, ratio of surfactant to drug solids and length of milling time. Analysis parameters were median size D50 and degree of polydispersity. The SpectralView heat map indicates D50 size with different sections representing four processing times.

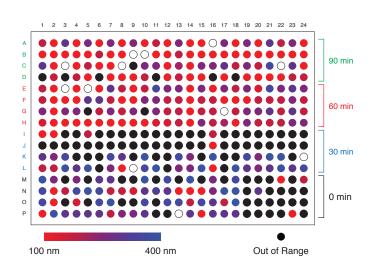


Small Molecules: Compound Library Screening for Drug Discovery

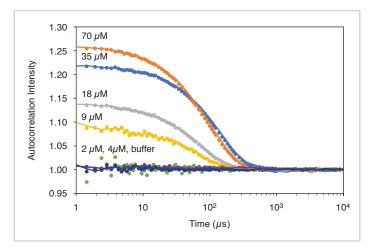
Large aggregates of small-molecule drugs are known to produce false positives through non-specific binding and inhibition of protein targets. Compounds may be screened in different buffers to identify aggregation conditions and optimal, non-aggregating buffers.

These DLS data show autocorrelation and size results for a common small-molecule drug compound. The critical aggregation concentration (CAC) is identified at approximately 4 μ M.





Polymer / surfactant additives	Drug compound nanoparticle measurements	
	D50 (nm)	%Pd
HPMC/SDS	144	23.2
HPC-SL/SDS	127	22.7
PVP K29-32/SDS	127	22.9
Poloxamer 407	228	23.1
Polysorbate 80	129	23.7



Specifications

Dimensions	60 cm (l) x 36 cm (w) x 25 cm (h)	
Digital Communication	Ethernet (TCP/IP)	
Onboard Camera	3 megapixels, operates up to 50 °C	
Correlator	512 channels, 100 ns sampling time in a multi-tau layout	
Electronics		
Read Time per Well	5 to 20 seconds (~1.5 hours for a 384 well plate)	
Temperature Control	4 °C to 85 °C*	
Attenuation Range	1 to 10 ⁵	
Laser Power	Programmable 10% to 100%	
Laser Wavelength	830 nm	
Optics		
Minimum Sample Volume	4 μL (2 mg/mL lysozyme in 1536 well plate)	
	Many industry-standard well plates are supported	
Supported Formats	96, 384, or 1536	
Well Plates		
at 67 kDa	Thight (00 µE BOA III Greiner 304 weil plate)	
Minimum Concentration	1 mg/mL (50 μL BSA in Greiner 384 well plate)	
Molar Mass Range	1000 to 1,000,000 Da	
Static Light Scattering		
Minimum Concentration at 14 kDa	0.125 mg/mL (50 μL lysozyme in Greiner 384 well plates)	
Size Range	0.5 to 1000 nm (hydrodynamic radius, r_h)	
Dynamic Light Scattering		

Warranty: All Wyatt instruments are guaranteed against manufacturing defects for 1 year.

* Absolute accuracy of \pm 0.5 °C from 4 °C to 50 °C, and \pm 1 °C from 50 °C to 85 °C. Minimum temperature of 4 °C requires a laboratory ambient temperature of 24 °C or below.

Wyatt Technology is committed to continual improvement. Specifications are subject to change without notice.

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Left to right: Geofrey K. Wyatt, President Dr. Philip J. Wyatt, Chief Executive Officer Clifford D. Wyatt, Executive Vice President

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The DynaPro Plate Reader III is just one of the many tools in the Light Scattering Toolkit for Essential Biophysical and Nanoparticle Characterization.

Learn more at www.wyatt.com

