

PL2609: Key publications on the characterization of nanoparticle drug and gene delivery systems by FFF-MALS-DLS

Summary

This document lists key publications using AF4- MALS-DLS for the characterization of nanoparticle drug delivery systems (nanoDDS), including nanoparticles for delivering small molecules, proteins, DNA, RNA and other therapeutics.

Characterization of drug products: methodology, validation and cross-comparisons

- Measuring particle size distribution of nanoparticle enabled medicinal products, the joint view of EUNCL and NCI-NCL. A step by step approach combining orthogonal measurements with increasing complexity

Caputo, F. et al., *J. Controlled Release* **299**, 31-43 (2019). <https://doi.org/10.1016/j.jconrel.2019.02.030>

- Measuring particle size distribution by asymmetric flow field flow fractionation: a powerful method for the preclinical characterization of lipid-based nanoparticles

Caputo, F. et al., *Mol. Pharm.* **16** (2), 756-767 (2019). <https://doi.org/10.1021/acs.molpharmaceut.8b01033>

- Are existing standard methods suitable for the evaluation of nanomedicines: some case studies.

Gioria, S. et al., *Nanomedicine (Lond)* **13**(5), 539-554 (2018). <https://doi.org/10.2217/nnm-2017-0338>

- FFF-MALS method development and measurements of size and molecular weight

Mehn, D. from EU Nanomedicine Characterization Laboratory web site, <http://www.euncl.eu/about-us/assay-cascade/PDFs/PCC/EUNCL-PCC-022.pdf?m=1468937868&>

Liposomes and lipid nanoparticles

- Physical characterization of liposomal drug formulations using multi-detector asymmetrical-flow field flow fractionation

Parot, J. et al. *J. Controlled Release* available online 28/1/2020

<https://doi.org/10.1016/j.jconrel.2020.01.049>

- Mechanism and kinetics of the loss of poorly soluble drugs from liposomal carriers studied by a novel flow field-flow fractionation-based drug release-/transfer-assay.

Hinna, A. H. et al., *J. Controlled Release* **232**, 228-37 (2016). <https://doi.org/10.1016/j.jconrel.2016.04.031>

- Size fractionation and size characterization of nanoemulsions of lipid droplets and large unilamellar lipid vesicles by asymmetric-flow field-flow fractionation/multi-angle light scattering and dynamic light scattering.

Vezocnik, V. et al., *J. Chromatography A* **1418**, 185-91 (2015). <https://doi.org/10.1016/j.jconrel.2016.04.031>

- Determination of size distribution and encapsulation efficiency of liposome-encapsulated hemoglobin drug substitutes using asymmetric flow field-flow fractionation coupled with multi-angle light scattering

Arifin, D. and Palmer, A., *Biotechnology Progress* **19**(6), 1798-1811 (2003)

- Application of Asymmetric Flow Field-Flow Fractionation hyphenations for liposome–antimicrobial peptide interaction

Iavicoli, P. et al., *J. Chromatography A* **1422**, 260-269 (2015). <https://doi.org/10.1016/j.jconrel.2016.04.031>

Polymersomes

- Toward Functional Synthetic Cells: In-Depth Study of Nanoparticle and Enzyme Diffusion through a Cross-Linked Polymersome Membrane

Gumz, H. et al., *Advanced Science* (2019). <https://doi.org/10.1002/advs.201801299>

- An alternative route to dye-polymer complexation study using asymmetrical flow field-flow fractionation

Boye s. et al., *J Chromatogr A* **1217**(29), 4841-9 (2010). <https://doi.org/10.1016/j.chroma.2010.05.036>

Exosomes

- Size characterization and quantification of exosomes by asymmetrical-flow field-flow fractionation

Sitar, S. et al., *Anal. Chem.* **87** (18), 9225-33 (2015). doi: 10.1021/acs.analchem.5b01636

- Identification of distinct nanoparticles and subsets of extracellular vesicles by asymmetric flow field-flow fractionation.

Zhang, H. et al. *Nature Cell Biol.* **1** (2018). <https://doi.org/10.1038/s41556-018-0040-4>

- Size Dependent Lipidomic Analysis of Urinary Exosomes from Patients with Prostate Cancer by Flow Field-Flow Fractionation and Nanoflow Liquid Chromatography-Tandem Mass Spectrometry

Yang, J.S. et al., *Anal. Chem.* **89**(4), 2488-2496 (2017). <https://doi.org/10.1021/acs.analchem.6b04634>

- A review of exosome separation techniques and characterization of B16-F10 mouse melanoma exosomes with AF4-UV-MALS-DLS-TEM

Petersen, K.E. et al., *Anal. Bioanal. Chem.* **406**(30), 7855-7866 (2014). <https://doi.org/10.1007/s00216-014-8040-0>

- Asymmetric-flow field-flow fractionation technology for exomere and small extracellular vesicle separation and characterization

Zhang H. and Lydem D., *Nature Protocol* **14**(4), 1 (2019). doi: 10.1038/s41596-019-0126-x

Virus-like particles

- Characterization of virus-like particle assembly for DNA delivery using asymmetrical flow field-flow fractionation and light scattering

Citkowicz, A. and Petry, H. *Anal. Biochem.* **376**, 163-172 (2008). <https://doi.org/10.1016/j.ab.2008.02.011>

- Quantitative analysis of virus-like particle size and distribution by field-flow fractionation

Chuan, Y.P. et al., *Biotechnology and Bioengineering* **99**(6) 1425-1433 (2008). <https://doi.org/10.1002/bit.21710>

- Quantitative characterization of virus-like particles by asymmetrical flow field flow fractionation, electrospray differential mobility analysis, and transmission electron microscopy

Pease, L.F. et al., *Biotechnology and Bioengineering* **102**(3) 845-855 (2009). <https://doi.org/10.1002/bit.22085>

- Quaternary size distribution of soluble aggregates of glutathione-S-transferase-purified viral protein as determined by asymmetrical flow field flow fractionation and dynamic light scattering

Lipin, D.I., Lua, L.H.L., Middelberg, A.P.J., *J. Chromatography A* **1190**(1-2), 204-2012 (2008).
<https://doi.org/10.1016/j.chroma.2008.03.032>

AAVs and other viruses

- Biophysical characterization of influenza virus subpopulations using field flow fractionation and multiangle light scattering: correlation of particle counts, size distribution and infectivity

Z. Wei et al, *J. Virol. Meth.* **144**(1-2), 122-132 (2007). <https://doi.org/10.1016/j.jviromet.2007.04.008>

- Quantitation of influenza virus using field-flow fractionation and multi-angle light scattering for quantifying influenza A particles

Bousse, T. et al., *J. Virological Meth.* **193**(2), 589-596 (2013). <https://doi.org/10.1016/j.jviromet.2013.07.026>



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