Physical interpretation of the size and concentration of extracellular vesicles measured by advanced techniques

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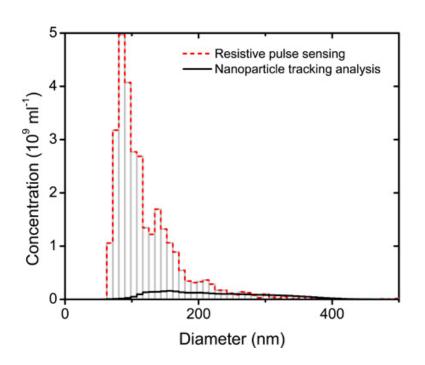


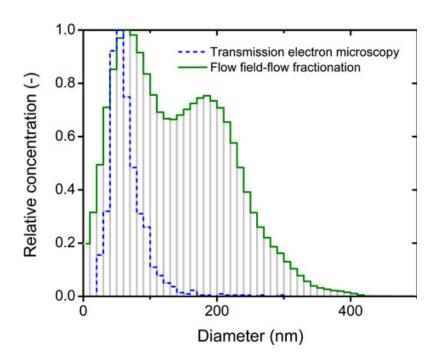
Disclosures of: Edwin van der Pol

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Other	No conflict of interest to disclose

Presentation includes discussion of the following off-label use of a drug or medical device: N/A

Introduction





- vesicles measured by different techniques*
- each technique obtains a different size distribution

Goals

- enable data comparison between techniques
- obtain the size distribution of vesicles

Methods

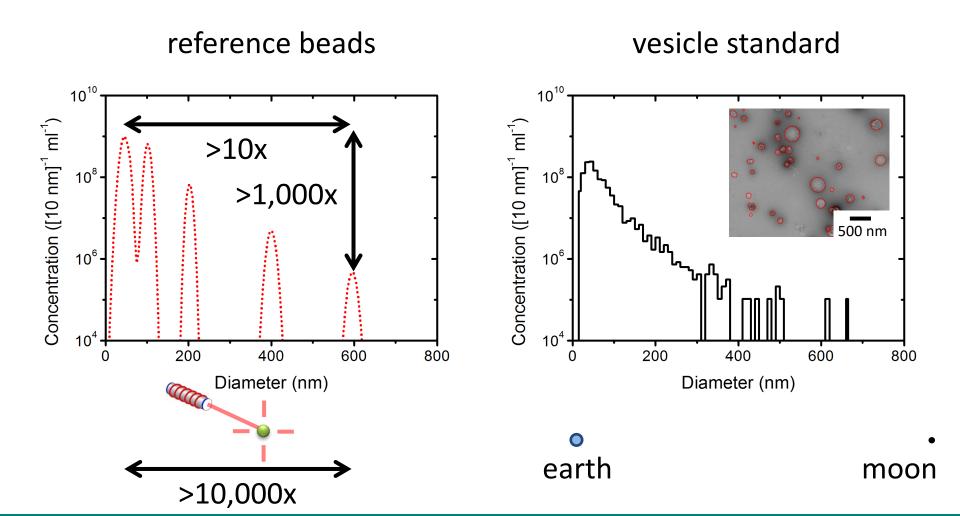






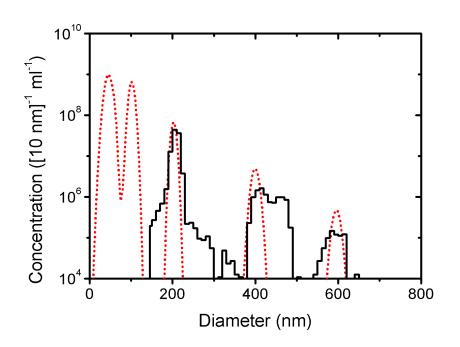
- standard population of
 - polystyrene beads
 - urine vesicles
- analyzed by
 - transmission electron microscopy
 - old flow cytometer (FACSCalibur)
 - new flow cytometer (Apogee A50-Micro)
 - nanoparticle tracking analysis (Nanosight NS500)
 - resistive pulse sensing (*Izon qNano*)

Transmission electron microscopy

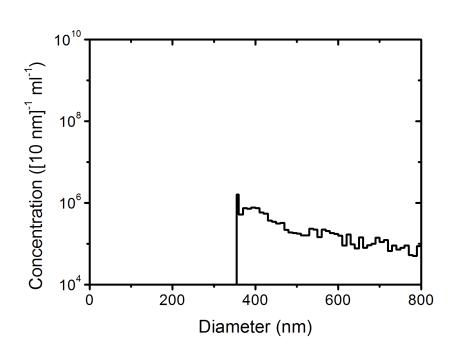


Conventional flow cytometry





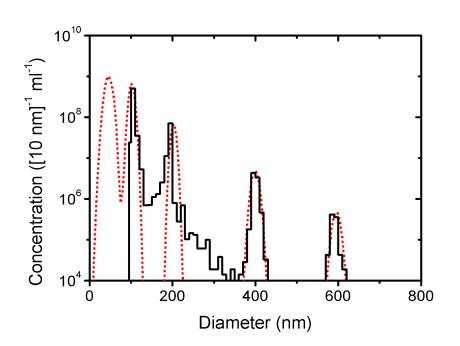
refractive index polystyrene = 1.61



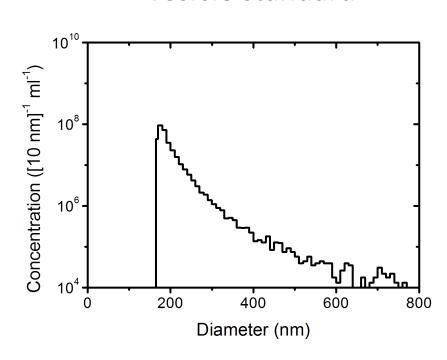
refractive index vesicles* = 1.4

Novel flow cytometry





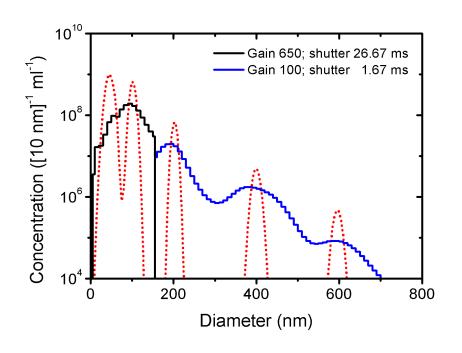
refractive index polystyrene = 1.61

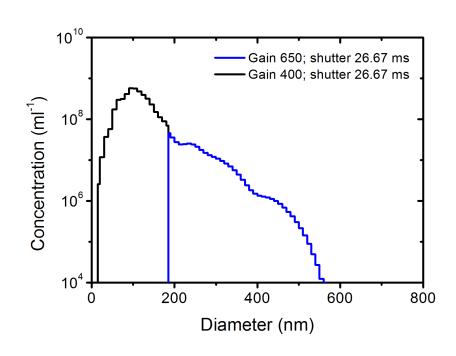


refractive index vesicles* = 1.4

Nanoparticle tracking analysis

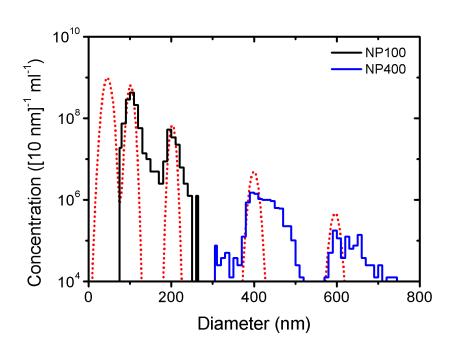
reference beads

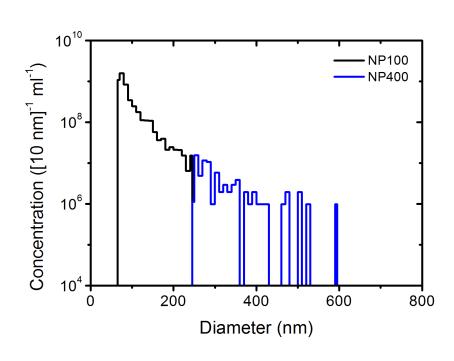




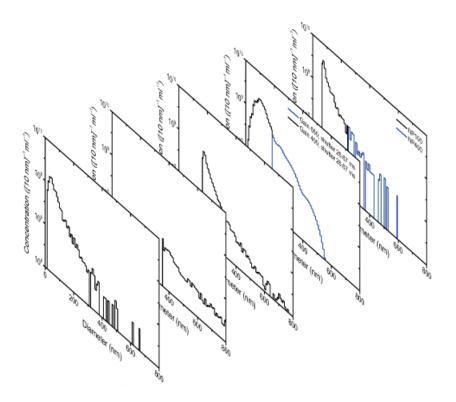
Resistive pulse sensing

reference beads

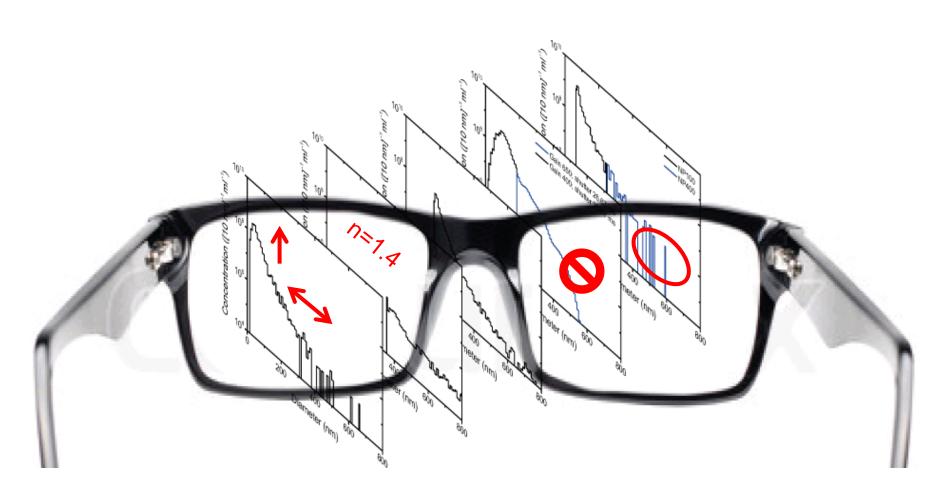




Summary



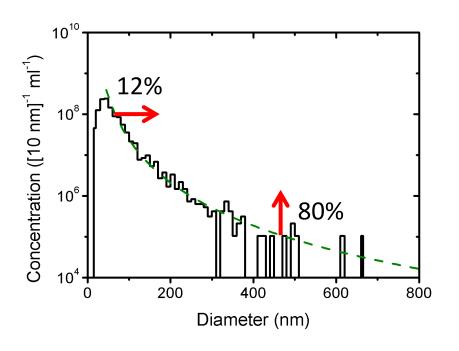
Summary

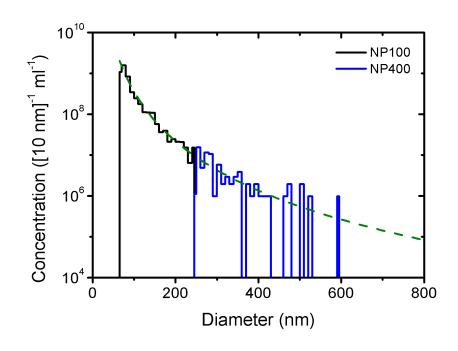


Outlook – data comparison

transmission electron microscopy

resistive pulse sensing





Outlook – fit parameters for clinical studies

transmission electron microscopy

power-law function:

plankton



$$C(d) = \left(k \frac{d}{d_0}\right)^{-m}$$

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More on microparticle detection: edwinvanderpol.com