

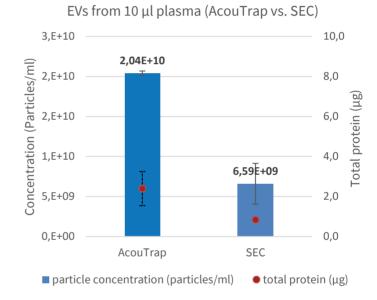
AcouTrap

Isolation of Extracellular Vesicles

HIGH QUALITY PURIFICATION FROM MINUTE SAMPLES

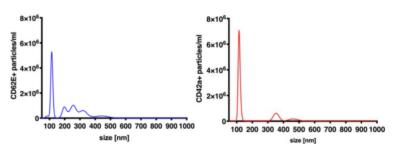
Demonstrated Suitable for Biomarker Discovery

- Rapid and automated isolation
- Reliable results from low-volume biological samples
- **High-quality** samples with demonstrated suitability for
 - Olink proteomics
 - miRNA analysis
 - Mass Spectrometry



High Quality EVs

- fNTA confirms endothelial-(CD62E+) and platelet-derived (CD42a+) extracellular vesicles in isolated plasma
- ELISA confirms presence of CD9/CD81/CD63



Data from:

Top: 10 μ l pooled platelet-poor plasma from healthy donors was processed using AcouTrap (n=2) and a commercially available size exclusion chromatography column (n=3) following the manufacturer's instructions. The resulting samples were measured using NanoSight LM10 and microBCA.

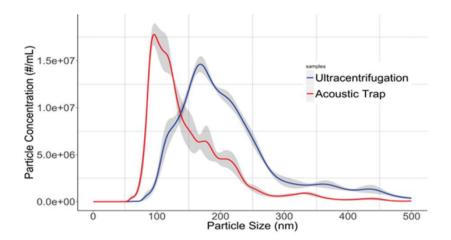
Bottom: Bryl-Górecka et al. Mol. Nutr. Food Res. 2020. <u>Bilberry Supplementation after Myocardial Infarction Decreases</u> <u>Microvesicles in Blood and Affects Endothelial Vesiculation</u>: Post enrichment, EVs were stained with anti-CD62E or anti-CD42a and analyzed in NanoSight equipped with 488 nm blue laser.



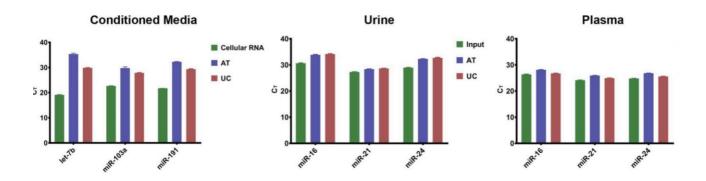


Comparison to Centrifugation

- **Proteomic profile:** LC-MS shows significant overlap (80%) between plasma samples processed using AcouTrap and differential centrifugation
- **Size distribution:** size distribution from NTA of of 300 μl plasma isolated either using AcouTrap or ultracentrifugation



• **miRNA content**: qRT-PCR shows similar miRNA levels in AcouTrap (AT) and ultracentrifugation (UC) samples from different sources



Data from:

Top: Rezeli et al. Anal. Chem. 2016, 88. <u>Comparative Proteomic Analysis of Extracellular Vesicles Isolated by Acoustic Trapping or Differential Centrifugation</u>. Plasma was processed either using AcouTrap or differential centrifugation and label-free LC-MS quantification was used to assess proteomic profiles.

Middle and bottom: Ku et al. Anal. Chem. 90, 2018 <u>Acoustic Enrichment of Extracellular Vesicles from Biological Fluids</u>: Samples were processed and assessed for EV size by nanoparticle tracking analysis, NTA (Top) and Ct values of EV-specific miR markers (Bottom).

