Standardized and fully automated deep plasma profiling of plasma integrated with Evosep One enables large scale clinical cohort analysis

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Complete end-to-end workflow allowing deep plasma profiling in a single analysis.

Standardized and fully automated enrichment of membrane-bound vesicles and depletion of abundant proteins from just 4 µl of plasma.

Highlights

Coupled to SCIEX ZenoTOF, the workflow reproducibly identifies >3,500 (▼) protein groups at a throughput of 30 samples per day by the push of a button.

Validation of enrichment and depletion

Different approaches to plasma preparations

Plasma preparation protocols vary between labs and we tested three procedures to assess their impact on the achievable proteome depth using the 100 samples per day method and the ZenoTOF 7600. The analysis of undepleted plasma samples revealed similar proteomic profiles across all three plasma preparation methods, while the Mag-Net enriched samples showed clear differences. All preparation strategies resulted in similar levels of depletion, but the 1x1200g prep resulted in the highest level of enrichment.

Rapid, fully automated membrane vesicle enrichment

Push-button solution - from plasma to peptides on Evotips

Here we present an end-to-end workflow on the Opentrons (OT-2) for the analysis of undepleted plasma and implementation of the Mag-Net protocol with PAC digestion and automated peptide cleanup for complete hands-free deep plasma proteome profiling. By simultaneously enriching vesicle-cargo proteins and depleting abundant plasma proteins using MagReSyn SAX beads, this workflow significantly increases plasma proteome depth in a standardized easy-to-use workflow. With just 4 µl of plasma input this method presents a cost-effective and sample conserving strategy for robust analysis of large sample cohorts. The workflow is based on this pre-print https://www.biorxiv.org/content/10.1101/2023.06.10.544439v2





depleted or enriched in the Mag-Net protocol using the Enrichr tool within Jensen COMPARTMENTS.

Good enrichment of vesicles across plasma preparations

High-throughput and ease-of-use

After deposition of 4 µl plasma and 1 µl MagResyn SAX beads, the protocol efficiently prepares 96 enriched plasma samples in 10 hours. To facilitate more comprehensive analysis, an additional set of 96 paired undepleted plasma samples can be prepared using just one additional µl of plasma and an additional 2 hours of running time.



Significant enrichment was achieved in Mag-Net 1x1200g detecting 80% of the EV marker proteins listed in Vesiclepedia. This enrichment was accomplished through efficient reduction of the dynamic range within the sample.



Rapid and robust plasma analysis

The reproducibility of the protocol was evaluated by preparing 40 Mag-Net samples and 8 samples of undepleted plasma (1x1200g). The analysis reproducibly resulted in the identification of 2,500 precursors in undepleted plasma and 7,500 precursors in Mag-Net samples with a digestion efficiency of ~85% using the 200 SPD method in combination with the SCIEX ZenoTOF 7600 and analyzed in library-free mode using DIA-NN. This resulted in the identification of 250 and 1,250 protein groups in undepleted plasma and Mag-Net samples, respectively. Finally, Mag-Net samples showed improved precursor precision compared to undepleted plasma.

More than 3,500 proteins identified in plasma

In combination with our standard methods the protocol allows for high-throughput application and in-depth plasma analysis. This resulted in the identification of 1,700 and 2,500 protein groups with the 100 and 60 samples per day methods respectively, when coupled to the SCIEX ZenoTOF 7600. This was increased to 30,000 precursors leading to more than 3,500 protein groups with the 30 samples per day method in library-free mode using DIA-NN. The protocol shows an effective reduction of the dynamic range and a 4-fold increase in the amount of identified protein groups relative to undepleted plasma.







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