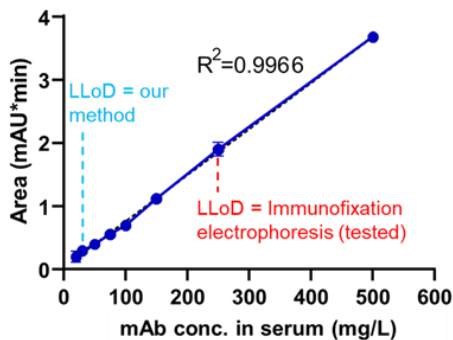




CTBT

Centre for Transfer of
Biomedical Technologies

TECHNOLOGY SUMMARY



Biomedical
Research
Centre

Technology Owner

University Hospital Hradec Králové

Inventors

Rudolf Kupčík
Marie Vajrychová
Ondřej Soukup
Jakub Radocha
Juraj Lenčo
Adéla Tomášová

IPR Status

Know-how

Stage of Development

Proof-of-Concept

Contact

Lucie Bartošová, Ph.D.
lucie.bartosova@fnhk.cz
+420 727 802 314

Blood-based highly sensitive diagnostics of multiple myeloma and monitoring of its progression

Background

- **Multiple myeloma (MM)** = cancer of plasma cells (PC) that produce monoclonal immunoglobulins (mAb, M-protein).
- **Disease complications** = high likelihood of **developing persistent minimal residual disease (MRD)** and **relapse**, both of which are associated with the presence of drug-resistant malignant plasma cell clone(s) - primary or emerged during treatment through clonal heterogeneity and evolution.
- **Detection of minor and persistent clones from blood is currently limited** = low sensitivity of current electrophoretic serum-based methods x more sensitive methods requiring invasive bone marrow biopsy.
- **Main objective** = to develop a simple and widely applicable method for sensitive detection of malignant PC clones based on **antibody** analysis in **human serum**. This method will allow early and sensitive detection of malignant clone progression.

Our Method

- **Isolation** = selective affinity isolation of antibodies from a **blood sample** using ≤ 3 different affinity matrices.
- **Analysis** = separation and detection using conventional HPLC system with UV-VIS detector.
- **Validation samples** = 5 different mAb standards + 28 patient serum samples (pre and post treatment).
- **Possible upgrade** = automation of isolation procedure + sensitivity enhancement via mass spectrometric detection of clonotypic peptides.

Conclusion

This novel approach allows highly sensitive and specific identification of primary myeloma cell clones in blood, those that become resistant and require following treatment as well as novel newly emerged clones. Additionally, the method enables distinguishing between pathological and therapeutic antibodies. The method employs a common analytical instrument, thus facilitating its widespread use. The method has a potential to replace standard immunofixation electrophoresis, avoid invasive and problematic bone marrow aspiration, and by that to facilitate early detection of disease relapse due to possibility of close patient monitoring at brief intervals.