

Proteomics Facility

Operator and User Regulations

1. Terms of Usage

The Proteomics core facility (PF) of the IZKF Aachen provides access for all members of the RWTH Aachen to proteomic technologies in order to facilitate the analysis and investigation of protein function on a molecular level in health and disease. The facility provides advice and support for the generation of such experimental setups and workflows based on the individual scientific question of the customer. This includes advice and suggestions for the process of appropriate sample generation. The acquired data is subsequently discussed with the researchers. Furthermore, the PF contributes in explanation and advice on data interpretation and possible follow-up experiments. The PF also offers experimental support in protein enrichment strategies such as co-IPs and affinity purification as well as PTM-specific protein and peptide enrichment strategies (such as phosphopeptide analysis).

2. Sample analysis requests:

All researchers have to contact the PF via email or phone (see contact details below) if they are interested in any form of proteomic analysis. The PF will then discuss the biological question, the necessary sample preparation, the required analysis procedure and the estimated costs with the respective researchers. After a decision has been made on a particular form of sample analysis, the respective researchers have to submit a signed "sample submission form" describing the nature and state of the individual samples, the planned sample analysis and an agreement to cover the costs for the analysis. Furthermore, a signed certificate of compliance regarding S1 regulations has to be provided. Samples that require S2 handling or contain radioactivity will not be analysed.

The PF is open to the entire RWTH Aachen. However, the PF has the right to refuse sample analysis if the project is ill-defined, too time-intensive or lacks the required funding.

3. Cost of sample analysis:

The individual groups have to carry the expenses for all the reagents required for sample analysis. These include - not exclusively - proteases (mainly trypsin and LysC), labelling reagents (formaldehyde (+/- Deuterium and ^{13}C) and cyanoborohydride (+/- Deuterium)), nanoLC columns, chromatography material (e.g. TiO_2 and/or Ti-IMAC for phosphopeptide enrichment), precasted gradient gels and others. Since the PF is required to employ refinancing measurements, the subsequent sample analysis will be charged at a time dependent rate ("hours of instrument-time"), depending on the number of samples. The current fee is 25 €/hour for members of the RWTH (according to DFG-guidelines), 30€/hour for external users from other universities and 50€/hour for commercial users.

4. Sample submission:

The experimental work required for sample generation is usually carried out in the individual laboratories (including the expression of recombinant proteins in bacteria or eukaryotic host systems and their application in in vitro assays, the growing and potential treatment of cells in cell culture, treatment of transgenic mice vs control animals and isolation of the relevant organs/tissues), not the facility itself.

Samples that are "ready-to-use" (these may include Coomassie stained protein gels, protein lysates/solutions with or w/o sample buffer, organs, tissues or similar) can be submitted to the PF. The samples are then prepared in the PF for their respective analysis.

5. Order of sample analysis:

Samples are usually analysed in a timely order depending on their date of submission. Exceptions to this order (at the discretion of the head of the PF) are made for samples that have to be measured for manuscript revisions or similar deadlines.

6. Working in the laboratory of the PF:

As described above, most of the work for sample generation is carried out in the individual labs of the respective researchers. These experiments cannot be performed in the PF lab. The PF has to operate in a very clean environment in order to minimize potential sources of contamination (such as dust, hair or skin particles) and can therefore not be considered a "standard wet lab". In case participating researchers need to perform specific parts of their sample generation within the lab of the PF, this needs to be discussed in advance. All personnel working in the lab is required to adhere to federal health and safety regulations and obey S1 regulation rules. It is also necessary to participate in the IZKF safety instruction lecture before working in the PF lab.

7. Instrument operation:

The PF harbours expensive equipment that requires significant long-term experience for operation. Therefore, both the chromatography systems as well as the mass spectrometers are exclusively operated by the staff members of the PF.

8. Core facility contribution:

Routine analysis in the PF is performed as a service (such as repeated measurements of a particular protein in a number of plasma samples). All work requiring a significant scientific input is carried out in form of a collaboration between the respective research groups and the PF.

9. Hinweis / please note:

All users of an IZKF core facility have to acknowledge the support of the respective facility in all relevant publications. Please use the following wording:

„Diese Arbeit wurde unterstützt durch die Proteomics core facility eine Core Facility des Interdisziplinären Zentrums für Klinische Forschung (IZKF) Aachen der Medizinischen Fakultät der RWTH.“

"This work was supported by the Proteomics core facility, a core facility of the Interdisciplinary Center for Clinical Research (IZKF) Aachen within the Faculty of Medicine at RWTH Aachen University."

Experimental approaches offered by the facility

Sample preparation (depending on the "state" of the submitted sample)

- Gel electrophoresis of protein samples
- Lysis and proteolytic digestion of protein samples in-solution, in-gel or on-bead (proteases: Trypsin, LysC, GluC,...); the PF uses standard lysis conditions such as urea or the FASP protocol. Novel protocols (such as the SP3 method) will be used for the analysis of samples with low concentrations (sub µg-range, <100000 cells, etc.).
- Sample purification and desalting (C18); small (zip-tips) to large scale (C18 cartridges)
- Sample concentration
- Sample depletion (plasma/serum)
- Sample labelling: If protein quantitation is required within the particular project, peptides can be modified using isobaric tags (TMT or iTRAQ). Furthermore, peptides can also be isotopically labelled using the dimethyl stable isotope labelling method. Cells that have been grown in SILAC medium can be used as an alternative. However, the label-free quantification approach is usually the preferred choice of quantitation for most analyses.

Sample fractionation/enrichment – peptide chromatography:

- The PF provides peptide separation using high pH reversed phase chromatography or strong anion/cation exchange (SAX/SCX) or similar techniques (such as HILIC, WAX).
- The PF also provides phosphopeptide enrichment using either a TiO₂ based chromatographic approach (mainly for peptides phosphorylated on serine and threonine residues) or anti-phosphotyrosine antibody mediated immunoprecipitation of peptides phosphorylated on tyrosine residues, followed by subsequent mass spectrometry analysis and quantification of phosphorylation changes.

Mass spectrometry:

- Analysis of desalted/purified samples by nanoLC-MS/MS using standardized methods and gradients. All samples are analysed on either the Q Exactive Plus or the Exploris 480 system. Both mass spectrometers are coupled to a nano-UPLC system.

Data analysis:

- Analysis of the raw data is performed by the PF in house using designated software packages (MaxQuant (with the built in Andromeda search engine) and Perseus). This includes protein (and PTM) identification and their respective quantification. The resulting data is explained to and discussed with the researchers. Furthermore, the facility provides advice on data interpretation and possible follow-up experiments.
- Storage of raw and analysis data in house (IZKF cloud) on separate hard drives for access at later time points.

Special services:

- The PF can also provide Western Blot analysis for groups that are not equipped with the required instrumentation. The PF also provides technical assistance and expertise for "unusual" biological questions in the field of protein biochemistry, including non-common protein chromatography and in vitro assays.

Operator and User Regulations

Proteomics Facility

2023-01



Contact:

Dr. Christian Preisinger

IZKF Aachen

cpreisinger@ukaachen.de

Tel.: 0241 80 88832

Stefanie Gostek, MTA, PTLA

IZKF Aachen

stgostek@ukaachen.de

Tel.: 0241 80 37295

Bettina Schweikart, CTA

IZKF Aachen

bschweikart@ukaachen.de

Tel.: 0241 80 38682