# CATALOGUE OF TECHNIQUES



### Technique Catalogue for ReMade@ARI

If you are looking at the catalogue to plan for your proposal, we would like to highlight that the Smart Science Cluster is available to help you.

For the discussion of suitable techniques please submit a pre-proposal: <u>https://apply.remade-project.eu/submit-call/remade-pre-proposal-submission</u>

or write your scientific question to: sciencesupport@remade-project.eu

We are looking forward to hearing from you!

All instruments marked in purple are actively involved in the industry access routes. If you are a company and wish to access another of our instruments, please contact us at: industry@remade-project.eu

Instruments in marked in orange are currently not available.

This document was last updated: 25 March 2025

It is a work in progress, which we try to keep up to date to the best of our knowledge. We do not give a guarantee that all listed instruments are available.

If you spot any inaccurate information, please feel free to reach out via <u>sciencesupport@remade-project.eu</u>

Thank you!



UK Research and Innovation



Federal Department of Economic Affairs Education and Research EAER State Secretariat for Education, Research and Innovation SERI

Funded by the European Union as part of the Horizon Europe call HORIZON-INFRA-2021-SERV-01 under grant agreement number 101058414 and co-funded by UK Research and Innovation (UKRI) under the UK government's Horizon Europe funding guarantee (grant number 10039728) and by the Swiss State Secretariat for Education, Research and Innovation (SERI) under contract number 22.00187.

Views and opinions expressed are however those of the author(s) only and do not necessarily reflect those of the European Union or the UK Science and Technology Facilities Council or the Swiss State Secretariat for Education, Research and Innovation (SERI). Neither the European Union nor the granting authorities can be held responsible for them.

#### Contents

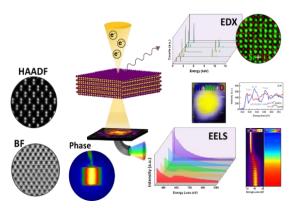
·	Electron Microscopy	3
•	High magnetic fields	8
•	Ion beam materials modification	10
•	Focused Ion Beams for material modification	12
•	<ul> <li>Ion beam analysis: spectrometry</li> </ul>	14
•	<ul> <li>Ion beam composition mapping and imaging</li> </ul>	16
C	Laser photo chemistry & spectroscopy	18
·	Laser processing	21
•	Muon Spectroscopy	23
·	<ul> <li>Neutron-based elemental composition analysis</li> </ul>	24
•	Neutron diffraction	26
•	Neutron imaging	28
·	Neutron reflectometry	30
·	Neutron small angle scattering	32
·	Neutron spectroscopy	33
C	Nuclear Magnetic Resonance	35
C	Positrons	36
c	• X-ray diffraction	37
C	IR- to VUV-beamlines	40
C	• X-ray imaging	41
C	• X-ray small/wide-angle scattering (SAXS-WAXS)	43
C	• X-ray spectroscopy	45
C	• X-ray tomography	48
•	All NanoEnviCz Instruments	50



#### **Electron Microscopy**

Electron Microscopy based techniques employ electrons to visualize and analyse the structure, morphology and composition of a sample at an atomic level achieving a resolution up to 50 pm. Usually, electron transparent samples (thinner than 100 nm) are illuminated with a "parallel" electron beam in conventional transmission electron microscopy (CTEM) or scanned with an electron probe in scanning transmission electron microscopy (STEM). Both techniques are capable of providing an atomic-level understanding of structures, properties and fundamental mechanisms in structural, functional and electronic materials.

In order to obtain information about the elemental composition in **TEM mode**, energy-filtered



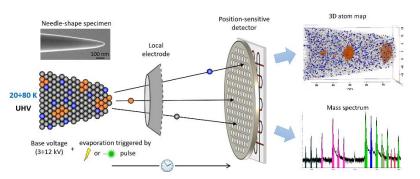
Schematic representation of Electron microscopy techniques. Image credit to Sara Martí-Sánchez.

transmission microscopy can be used. Here, only inelastically scattered electrons with specific, characteristic energies are used for imaging, and only these contribute to the generation of energy-filtered images. In this way, the distribution of chemical elements can be displayed. In **STEM**, energy dispersive X-ray spectroscopy (EDS) is used to determine the elemental composition of a sample at the atomic level. At the same time, Electron Energy Loss Spectroscopy (EELS) can be employed for determining the elemental composition, oxidation state / bonding information and electronic band structure and collective excitations in the material (plasmons, band gap, JDOS, excitons, phonons...).

Ga based <u>FIB</u>s are widespread and are commonly used for the preparation of TEM lamella and in combination with an SEM and EDS or EBSD detectors for tomography, furthermore, cryo stages and nanomanipulators can enhance the versatility of this technique.

Atom probe tomography (APT) is

a powerful technique for quantitative three-dimensional elemental analysis of solids at the near-atomic scale. The method employs the phenomenon of field evaporation, where atoms are ionized and desorbed from the surface of a tiny needle-



shaped specimen (30-200 nm end Schematic representation of APT.

radius) in a high electric field. This field is generated by applying a standing voltage of a few kV, combined with extremely short voltage or laser pulses that trigger ion-by-ion evaporation events. The emitted ions are then accelerated toward a position-sensitive detector.

By analysing detector hit positions and ion time-of-flight, software algorithms reconstruct the 3D atomic distribution within the specimen with sub-nanometer precision. APT essentially combines ion projection microscopy with time-of-flight mass spectrometry, producing a dataset where each detected ion has spatial coordinates (x, y, z) and a mass-to-charge ratio.

Key features: high spatial resolution (~ 1 nm), high mass resolution (m/ $\Delta$ m > 1000, resolving isotopes), quantitative elemental composition without external standards, detection sensitivity down to 10 ppm, equal sensitivity to all elements in the periodic table.

**APT is highly complementary to electron microscopy-based methods**, particularly transmission electron microscopy (TEM). While TEM provides structural information, APT delivers exceptionally high analytical sensitivity in true 3D space but lacks structural details.

#### Techniques

- In situ (S)TEM Temperature (heating, cooling), gas, liquid, bias, strain... induced changes can be analysed at the atomic scale
- Electron Diffraction (ED) Crystal phase identification
- Electron tomography (ET) 3D reconstruction of the materials
- High Resolution Transmission Electron Microscopy (HRTEM) Crystal structure visualization
- (Aberration Corrected) Scanning Transmission Electron Microscopy ((AC-)STEM) Imaging of the samples (with atomic resolution) with different contrast sources including high-angle annular dark field (HAADF), medium-angle annular dark field (MAADF), low-angle annular dark field (LAADF), annular bright field (ABF) and bright field (BF)
- **High-angle annular dark field (HAADF) imaging** imaging of incoherently scattered electrons, Z-contrast image, sensitive to the atomic number, heavier atoms with brighter contrast
- Integrated differential phase contrast (IDPC) phase contrast of projected electrostatic potential, roughly proportional to the atomic number, sensitive to the light elements compared to HAADF
- **4-Dimensional Scanning Transmission Electron Microscopy (4D-STEM)** Phase contrast technique, in-situ strain, phase, electric and magnetic field mapping
- Electron holography (EH) Phase contrast technique, electric and magnetic field mapping
- Electron Energy Loss Spectroscopy (EELS) Compositional mapping, valence state determination and low energy excitations mapping (plasmons, phonons, bandgap...), sensitive to light elements
- Energy Dispersive X-Ray Spectroscopy (EDS) Compositional mapping
- Energy Filtered TEM (EFTEM) Compositional mapping with improved contrast in both image and diffraction pattern
- Lorentz Microscopy magnetic materials, imaging the magnetic domain structure at large defocus
- Scanning Electron Microscopy (SEM) Surface morphology and compositional information
- Electron Paramagnetic Resonance (EPR) Anisotropy and spin states, identification of paramagnetic substances, geometric and electronic structure of a paramagnet, information about distances of radicals
- **TEM lamellae preparation** thanks to the IBL capabilities, it is possible to cut and manipulate small structures to produce samples to be studied in a dedicated TEM (optionally in Cryo conditions).
- Electron Back Scatter Diffraction (EBDS) crystal structure determination through backscattered electrons for grain, defects and plastic deformation analysis
- Atom probe tomography (APT) Analysing elemental segregation at grain boundaries and interfaces, mapping and quantifying dopants in nanostructures, investigating clustering and precipitation in materials, determining local compositions at the nanoscale.

Network	Coun- try	Access pro- vider	Infra- structure	Instrument	Technique	Link
e-DREAM	IT	CNR	CNR	SEM Laboratory	SEM, EDS	<u>learn</u> more
e-DREAM	IT	CNR	CNR	TEM Laboratory	TEM, STEM, EDS	<u>learn</u> more
e-DREAM	DE	FZJ	ER-C	TFS Arctica G2	cryo SEM, TEM	<u>learn</u> more
e-DREAM	DE	FZJ	ER-C	FEI Helios NanoLab 400S FIB-SEM	SEM, EDS, FIB	<u>learn</u> more
e-DREAM	DE	FZJ	ER-C	FEI Helios NanoLab 460F1 FIB-SEM	SEM, EDS, FIB	<u>learn</u> more
e-DREAM	DE	FZJ	ER-C	FEI Tecnai G2 F20	TEM, EELS, EFTEM, STEM with HAADF STEM imaging, Lorentz microscopy, ET, in-situ heating/ cooling/ strain	learn more
e-DREAM	DE	FZJ	ER-C	FEI Titan 80-300 STEM	EELS, ET, HRSTEM with HAADF STEM imaging, monochromator	<u>learn</u> more
e-DREAM	DE	FZJ	ER-C	FEI Titan 80-300 TEM	HRTEM, in-situ heating/ cooling/ strain	<u>learn</u> more
e-DREAM	DE	FZJ	ER-C	FEI Titan G2 60-300 HOLO	HRTEM, EELS, EFTEM, HRSTEM with HAADF STEM imaging, off-axis EH, Lorentz microscopy, ET, in-situ heating/ cooling/ strain	learn more
e-DREAM	DE	FZJ	ER-C	FEI Titan G2 80-200 ChemiSTEM	EELS, ET, HRSTEM with HAADF STEM imaging, 4D-STEM, IDPC, EDS, in-situ heating/ cooling/ strain	learn more

				1 1		
e-DREAM	DE	FZJ	ER-C	Hitachi HF 5000 TEM/STEM	EELS, ET, HRSTEM with HAADF STEM imaging, 4D-STEM, IDPC, EDS, in-situ gas	<u>learn</u> more
e-DREAM	DE	FZJ	ER-C	TESCAN Tensor	STEM, EDS, 4D- STEM	<u>learn</u> more
e-DREAM	DE	FZJ	ER-C	TFS Spectra 300	HRTEM, EELS, EFTEM, HRSTEM with HAADF STEM imaging, Lorentz microscopy, ET, 4D- STEM, IDPC, monochromator	learn more
e-DREAM	DE	FZJ	ER-C	LEAP 4000X HR (Cameca Scientific Instruments, Madison, WI, USA)	APT	
e-DREAM	ES	ICN2	ICN2	FEI MAGELLAN 400L SEM	SEM, EDS, STEM	<u>learn</u> more
e-DREAM	ES	ICN2	ICN2	FEI Quanta 650F ESEM	SEM, EDS, STEM	<u>learn</u> more
e-DREAM	ES	ICN2	ICN2	FEI Tecnai F20 (S)TEM	TEM, EELS, EFTEM, STEM with HAADF STEM imaging, Lorentz microscopy, ET	<u>learn</u> more
e-DREAM	ES	ICN2	ICN2	Thermo Fisher SPECTRA 300	HRTEM, EELS, EFTEM, HRSTEM with HAADF STEM imaging, Lorentz microscopy, ET, 4D- STEM, IDPC, monochromator	learn more
non-ARIE	EU	CERIC- ERIC	CERIC-ERIC	FEI Titan Krios 3Gi@SOLARIS	cryo TEM	<u>learn</u> more
non-ARIE	EU	CERIC- ERIC	CERIC-ERIC	FE-SEM@CUP	SEM	<u>learn</u> more
non-ARIE	EU	CERIC- ERIC	CERIC-ERIC	JEOL JEM ARM 200F@NIMP	HR TEM/STEM, EDS & EELS	<u>learn</u> more
non-ARIE	CZ	NanoE nviCz	NanoEnviCz	HRSEM FEI NanoSEM 450 (UACH4)	HRSEM, STEM	<u>learn</u> more

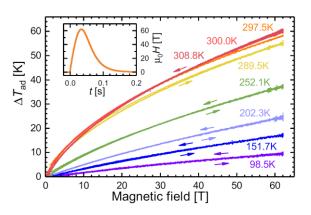
				1		
non-ARIE	CZ	NanoE nviCz	NanoEnviCz	Scanning Electron Microscope, Hitachi (UFCH22)	FESEM, EDS	<u>learn</u> more
non-ARIE	CZ	NanoE nviCz	NanoEnviCz	Scanning Electron Microscope (SEM) Hitachi SU6600 (UPOL10)	SEM, EDS	<u>learn</u> more
non-ARIE	CZ	NanoE nviCz	NanoEnviCz	Transmission Electron Microscope (TEM) JEOL 2100 (UPOL11)	TEM, EDS	<u>learn</u> more
non-ARIE	CZ	NanoE nviCz	NanoEnviCz	Electron-Paramagnetic- Resonance Spectrometer (UPOL13)	EPR	<u>learn</u> more
non-ARIE	CZ	NanoE nviCz	NanoEnviCz	High resolution transmission electron microscope (JEOL) JEM 3010 (UACH10)	HRTEM, EDS, ED, phase/orientation mapping	<u>learn</u> more
non-ARIE	CZ	NanoE nviCz	NanoEnviCz	High resolution transmission electron microscope, HRTEM FEI Talos F200X (UACH16)	HRTEM, STEM- HAADF, EDS	<u>learn</u> more
non-ARIE	CZ	NanoE nviCz	NanoEnviCz	High resolution transmission electron microscope (UFCH21)	HRTEM	<u>learn</u> more
non-ARIE	CZ	NanoE nviCz	NanoEnviCz	High Resolution Transmission Electron Microscope (UPOL5)	HRTEM, EDS, EELS	<u>learn</u> more
non-ARIE	CZ	NanoE nviCz	NanoEnviCz	XPS/ESCA and Auger electron spectroscopy (UJEP3)	XPS, ESCA, AES, SEM	<u>learn</u> more
non-ARIE	CZ	NanoE nviCz	NanoEnviCz	Precision Ion Polishing System (PIPS) Model 691(Gatan) (UACH12)	Sample preparation	learn more
Laserlab- Europe	HR	LLE- AISBL	CALT	SEM	SEM	<u>learn</u> more
Laserlab- Europe	CZ	IP- ASCR	HILASE	SEM	SEM	<u>learn</u> more



#### High magnetic fields

The application of high magnetic fields allows a controlled tuning of the physical properties of bulk and thin film material samples. The electronic or lattice magneto response can be investigated by various experimental techniques. The high magnetic fields are generated either as pulsed or continuous fields with maximum strengths of up to 95 T and up to 38 T, respectively, and at sample temperatures down to about 1 K. For selected techniques, hydrostatic pressures of up to several GPa can be applied to the samples.

High magnetic fields are suitable, for example, for characterising new materials with hard magnetic properties for electric motor technology, or with pronounced magnetocaloric properties for applications in cooling technology.



Magnetocaloric effect of single-crystalline gadolinium: adiabatic temperature change in pulsed magnetic fields up to 62 T. The inset shows the temporal profile of the magnetic field pulse (T. Gottsschall et al., Phys. Rev. B 99, 134429 (2019)).

#### Techniques

- Magnetocaloric effects (MCE) measures an adiabatic temperature change of a material, caused by a pulsed magnetic field
- Magnetotransport measures the electrical resistance and Hall effect in magnetic fields
- Magnetization / Vibrating Sample Magnetometer (VSM) measures the uniform bulk magnetization
- Ultrasound measures the sound velocity and attenuation
- Electric polarization measures the electric bulk polarization of a material
- Electron magnetic resonance (ESR) measures electronic magnetic properties and low-energy magnetic excitations
- Electrical Transport Option (ETO) supports three types of measurements including resistivity, IV curves and differential resistance
- Magnetostriction measures the relative length change of a sample in a magnetic field
- Nuclear magnetic resonance (NMR) measures the internal magnetic fields and crystal electric field gradients at the nuclear sites
- **Magnetic torque** measures the magnetic torque due to the bulk magnetization of a sample in a magnetic field
- Magneto-optical transmission measures the absorption of light in the presence of a magnetic field
- Magnetic Birefringence measures the refractive index of a material in a magnetic field
- Microscopy visual study of organisms, materials or solutions
- Far-infrared Spectroscopy probes low-energy optical excitations
- Ultrafast Spectroscopy probes the dynamics in materials on extremely short time scales

Networ k	Coun- try	Access provider	Infrastru cture	Instrument	Technique	Link
EMFL	DE	HLD	HLD	Various types of pulsed-field magnets, fields up to 95 T and pulse durations of about 100 ms, sev- eral 10 mm^3 of sample space	MCE, Magnetotransport, Magnetization, Ultrasound, Electric polarization, ESR, Magnetostriction, NMR, Magnetic torque, Magneto- optical transmission	<u>learn</u> more
EMFL	NL	HFML	HFML	Various types of con- tinuous-field mag- nets, fields up to 38 T, several 10 mm^3 of sample space	Magnetotransport, Magnetization, Magnetic torque, VSM, Magnetic Birefringence, Microscopy, Far-infrared Spectroscopy, Ultrafast spectroscopy	<u>learn</u> more
LEAPS	NL	SRU	FELIX	DC Magnet	High magnetic field (also possible in combination with IR/THz spectroscopy)	<u>learn</u> more
non- ARIE	CZ	NanoEnvi Cz	NanoEnvi Cz	Physical Properties Measurement Sys- tem - PPMS (UPOL2)	VSM, DC measurements, Electrical transport option (ETO)	<u>learn</u> more
non- ARIE	CZ	NanoEnvi Cz	NanoEnvi Cz	Low temperature in- duction magnetome- ter - PPMS (UPOL14)	VSM, DC and AC, ETO	<u>learn</u> more

11



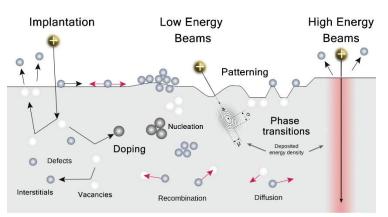
#### Ion beam materials modification

Ion beams have been for the past decades used routinely to **modify and study** the structure and **properties of metals, insulators and semiconductors**. Making use of different ion-target interactions, either by nuclear collisions or electronic excitations, ion beams are used to:

- **introduce dopants and defects** in materials by ion-implantation and hence induce impuritydefect interactions that change the electronic behaviour of the materials (e.g. semiconductors, optoelectronics, stress-strain engineering);
- **create nano-patterns** on surfaces by ion sputtering or in bulk through ion-induced phase-transitions (refer to the following section on Focus Ion Beams);
- grow layered structures by ion-beam assisted deposition;
- perform radiation-related degradation studies;
- **inspect** composition and crystallographic structure of materials, with depth resolution, by studying their response to the impinging ions (see following sections Ion Beam Analysis: Spectrometry and Mapping).

Radiate offers beams of a large variety of ion species across a broad spectrum of kinetic (from a few

eV to several GeV) and potential energies (charge states up to +45). Furthermore, RADIATE provides nanometric ion beams that can be used for maskless lithography in micro and nanofabrication workflows by atomic sputtering of the surface. Additionally, RADIATE beams (including broad, micro- and nano-metric beams) can perform a wide range of Ion Beam Analysis (IBA) experiments. Please refer to the subsequent sections for more details on Focus Ion Beams and IBA.



Schematic of the main techniques and mechanisms involved in the modification of materials by ion beams. Image credit to Stefan Facsko.

#### Techniques

- Ion Implantation keV-MeV beams are used to introduce dopants and defects in materials. Moreover, these beams are employed in IBA, either in- or ex-situ.
- Low-Energy Ion Beams keV irradiation experiments, such as Highly Charged Ions (HCL) or Low Energy Irradiation (LEI) radiations, are used in surface modification and degradation studies. Furthermore, these beams are used for near-surface high-resolution IBA).
- **High-Energy Beams** Such beams typically consist of heavy and highly energetic ions (SHI) that interact with solids mainly through ion-electron excitation processes, contrasting with typical ion implantation, where the impinging ions lose their energy primarily by nuclei collisions. These beams are explored in nanofabrication and to test radiation-hard electronics.

• Focus Ion Beams - Nanometric ion beams used for maskless lithography in micro and nanofabrication workflows by atomic sputtering. They are also used to characterise samples.

The table below provides a general overview of the principal technical contributions delivered by each of the participating laboratories of RADIATE. Note that the list does not reflect the entire range of competencies available at each lab. Please refer to the subsequent sections for the possibilities related to Focus Ion Beams and IBA within ReMade@ARI.

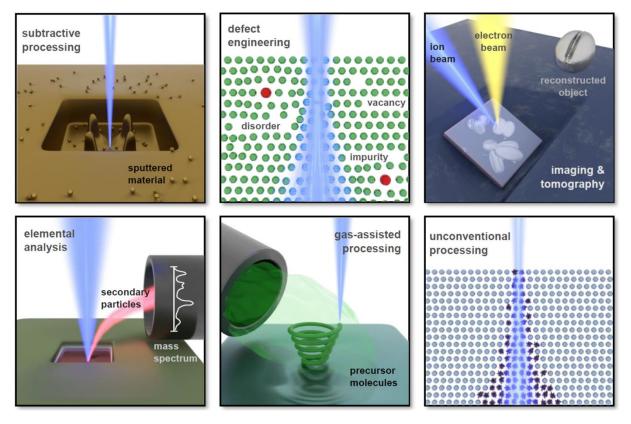
Network	Country	Access provider	Infrastructure	Instrument	Techniques	Link
RADIATE	FR	CNRS	GANIL	GANIL	Swift-heavy ion and	learn
					highly charged ion irradi-	more
					ations	learn
						more
RADIATE	СН	ETHZ	LIP	6 MV Tandem and	Deep implantations in	learn
				1.7 MV Tandetron	the MeV ion energy	more
				Accelerators	range.	
RADIATE	DE	HZDR	IBC	Tandem Accelera-	High/low-energy and	learn
				tors, 500 kV Im-	highly-charged ion im-	more
				planter, low-energy	plantation, focused ion	
				ion irradiation	implantation, clean-room	
					environment	
RADIATE	PT	IST	IBL	high flux 210 kV ion	High flux implantations	learn
				implanter, S1090	at different tempera-	more
				Danfysik	tures.	
RADIATE	РТ	IST	IBL	2.5 MV Van de	proton irradiation, in situ	learn
				Graaff and 2.5 MV	electrical	more
				Tandem	characterization	
				accelerators		
RADIATE	BE	KU	IMBL	High Flux ion Im-	High flux implantations.	learn
		Leuven		planter, Danfysik		more
				S1090		
RADIATE	CZ	NPI	LT	Ion beam lines at	Ion beam lithography	learn
				Tandetron	and implantation, exter-	more
					nal beam.	
RADIATE	HR	RBI	RBI-AF	6 MV Tandem and	Multi-beam radiations,	learn
				1 MV Tandentron	single ion implantation	more
				Accelerators	and deep implantations.	
RADIATE	ES	UAM	CMAM	5MV tandem +	High energy implanta-	learn
				Implantation BL +	tions coupled to optical	more
				internal and	in-situ measurements	
				external	(e.g. temperature moni-	
				microbeam	toring), concurrent ion- laser irradiations.	
	<u>с</u> г	UU	Tandom	2EO W/Implantor	Provides MeV radiations	loarn
RADIATE	SE	00	Tandem Laboratory	350 kV Implanter	for material modifica-	learn more
			Laboratory		tion.	more



#### Focused Ion Beams for material modification

A focused ion beam (FIB) is a class of ion beams that utilizes a nanosized beam of ions with a few keV to a few 10 keV of energy for the modification and analysis of materials (see also sections on <u>Ion Beam</u> <u>Materials Modification</u> and <u>Ion Beam Analysis</u>). It is in particular the ability for on-demand spatially resolved implantation, removal, or addition of material and defect generation which makes the FIB tools so interesting for micro and nanoengineering.

A wide variety of ions can be used in a FIB tool. This includes commercially available ions like He, Li, N, O, Ne, Si, Ar, Ga, Ge, Xe, Cs, Au, Bi as well as academic developments including B, C, Fe, Co, Cu, Rb, Ce, Pr, Dy, Pb and many others. A special variant of the FIB employs a pattern generator and precursor gases for 3D additive manufacturing of metallic and insulating structures. In combination with in-situ methods like  $\mu$ -manipulators, heating stages, specialized detectors (Scanning Transmission Ion Microscopy, Secondary Ion Mass Spectrometry, etc) they become powerful instruments for spatially resolved studies of material modification and analysis. While all ions can be used for imaging, it is Helium Ion Microscopy that stands out due to the capability to also investigate uncoated insulating samples without additional coatings while inducing minimal damage.



Schematic overview on the different FIB techniques. Reused from: K. Höflich et al, Roadmap for focused ion beam technologies. <u>arXiv:2305.19631</u>.

#### Techniques

- Spatially resolved doping using Liquid metal alloy ion sources
- Single Ion Implantation for quantum technology applications
- Defect engineering in oxides, semiconductors, metals and 2D materials
- Radiation hardness for semiconductors and metals
- Direct write material removal and additive manufacturing
- Ion Beam lithography for resist-based processes

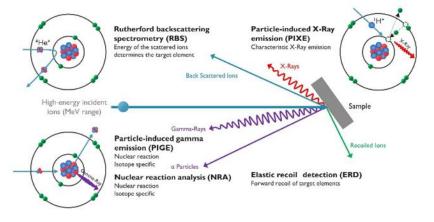
Network	Coun- try	Access pro- vider	Infra- structure	Instrument	Technique	Link
RADIATE	DE	HZDR	IBC	Helium Ion Microscopy	IBL, ET, EBDS	learn more
RADIATE	DE	HZDR	IBC	Ga and non-Ga FIB	IBL	learn more
e-DREAM	DE	FZJ	ER-C	FEI Helios NanoLab 400S FIB- SEM	IBL, SEM, EDS	learn more
e-DREAM	DE	FZJ	ER-C	FEI Helios NanoLab 460F1 FIB-SEM	IBL, SEM, ET, EDS	learn more
e-DREAM	ES	ICN2	ICN2	Thermo Fisher HELIOS 5UX FIB	SEM, ET, EDS	learn more
non-ARIE	CZ	NanoE nviCz	NanoEn- viCz	Precision Ion Polishing Sys- tem (PIPS) Model 691(Ga- tan) (UACH12)		learn more



#### Ion beam analysis: spectrometry

lon beam analysis (IBA) comprises а suite of analytical techniques that explore the interaction between high-energy ions and atoms within а substrate. This interaction gives rise to various outcomes, enabling to obtain information about elemental quantification,

compositional analysis, elemental depth profiling, density analysis and even crystallographic analysis. The



Schematic representation of different ion beam analysis techniques. Image credit to Masoud Dialameh.

incident ions are typically in MeV energy range, with interactions encompassing elastic scattering (RBS), elastic recoil scattering (ERD), nuclear reactions (NRA), X-ray emission (PIXE), etc. Frequently, diverse IBA techniques can be simultaneously applied, offering complementary insights. The utility of these IBA techniques extends beyond the confines of traditional analysis. By precisely characterizing materials and products, these techniques aid in optimizing recycling processes, identifying material degradation, and ensuring the quality and longevity of products throughout their lifecycle. Additional advantages lie in the non-destructive nature of these techniques for certain samples and the possibility of both in-situ and ex-situ experiments during radiation.

#### Techniques

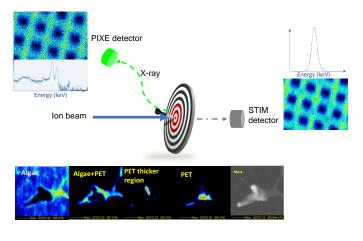
- **Rutherford backscattering spectrometry (RBS) and Ion Channeling** for elemental quantification, depth profiling of elements heavier than the substrate with a sensitivity down to 1E+13 atoms/cm<sup>2</sup>, and defect studying depth profiles.
- Elastic recoil detection (ERD) for compositional analysis and depth profiling for a wide range of elements starting from H up to heavy elements with a sensitivity reaching to 1E+14 atoms/cm<sup>2</sup>.
- Nuclear reaction analysis (NRA) and Particle-induced gamma emission (PIGE) isotopespecific elemental quantification of low Z elements (below Na) and depth profiling with a sensitivity down to 1E+12 atoms/cm<sup>2</sup>.
- **Particle-induced X-ray emission (PIXE)** for elemental identification and compositional analysis from sodium onwards with a sensitivity down to 100 ppm.
- **Proton elastic Scattering Analysis (PESA)** proton beams are used for hydrogen detection in very thin samples in transmission mode.
- Accelerator mass spectrometry (AMS) most famous for radiocarbon (<sup>14</sup>C) dating, uses similar set-ups as IBA, but samples needs to be chemically processed before being introduced in the ion source. Negative ions (e.g., C<sup>-</sup>) are extracted from the sample, further accelerated to MeV, stripped to positively-charged ions and radionuclide-to-stable nuclide ratios are detected.

Network	Coun- try	Access provider	Infra- structure	Instrument	Techniques	Link
RADIATE	сн	ETHZ	LIP	1.7 MV Tandetron and 6 MV	RBS, ERD-ToF,	learn
	en	21112	2	HVEC EN-Tandem	NRA, PIXE	more
				accelerator		
RADIATE	DE	HZDR	IBC	6 MV Tandem Accelerator,	RBS, RBS-C,	learn
				3 MV Tandem Accelerator,	ERD-ToF, NRA,	more
				100 kV Ion Platform	PIGE, PIXE	
RADIATE	IT	INFN	LABEC	3 MeV Tandetron	RBS, PIGE, PIXE	learn
						more
RADIATE	PT	IST	IBL	2.5 MV Van de Graaff and	RBS, ERD, NRA,	learn
				2.5 MV Tandem accelerators	PIGE, PIXE	more
RADIATE	FI	JYU	AL	1.7 MV Tandem Pelletron	RBS, ERD-ToF	learn
						more
RADIATE	BE	KU Leuven	IMBL	1.7 MV Tandem Pelletron	RBS, ERD-ToF,	learn
				and 2.5 MV Van de Graaff	PIXE	more
RADIATE	CZ	NPI	LT	Ion beam lines at Tandetron	RBS, RBS-C,	learn
					ERD, ERD-ToF,	more
					PESA, PIXE	
RADIATE	HR	RBI	RBI-AF	1 MV Tandentron and 6 MV	RBS, ERD, NRA,	learn
				Tandem Van de Graaff	PIGE, PIXE	more
RADIATE	ES	UAM	CMAM	5 MV Tandem	RBS, ERD-ToF,	learn
					NRA, PIGE, PIXE,	more
					microbeam	
RADIATE	AT	UNIVIE	VERA	Accelerator mass spectrom-	AMS	learn
				etry facility		more
				(3 MV tandem)		
RADIATE	ES	USE	CNA	3 MeV Tandem	RBS, ERD, NRA,	learn
					PIGE, PIXE	more
RADIATE	SE	UU	Tandem	5 MV pelletron accelerator,	RBS, ERD, NRA,	learn
			Labaratory	Low-energy Ion Scattering System	PIXE	more



#### Ion beam composition mapping and imaging

The techniques used for ion beam composition mapping and imaging are also included in the IBA techniques mentioned in the section IBA: spectrometry. The IBA technique can be implemented using both defocused and focused ion beams. Using a focused ion beam enables the achievement of composition mapping and imaging. The dimensions of the focused beam are predominantly within the micrometric range, although on occasion, they can become even finer, extending into the submicrometric scale. Furthermore, MeV ions have a greater penetration depth into samples compared to keV electrons, and they also exhibit significantly reduced lateral scattering. There are two methods: the beam is swept over a stationary target,



General scheme of STIM, PIXE, and its simultaneously measurement of a copper grid. *Bottom:* STIM mass and elemental maps of algae sample contaminated with microplastics. Image credit to Noelia Maldonado-Gavilán.

or the sample is moved with the beam fixed. These characteristics prove particularly advantageous when analysing samples with thicknesses spanning a few micrometres, allowing for the visualization of micrometric or submicrometric details. The insights gleaned from the maps are contingent on the technique employed, spanning from elemental composition (PIXE, RBS and PIGE) and mass density distribution (STIM) to the electronic properties inherent in semiconductors (IBIC).

#### Techniques

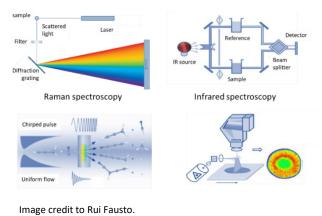
- **Particle Induced X-ray Emission (PIXE)** for elemental identification and compositional analysis from sodium onwards with a sensitivity down to 100 ppm.
- Particle Induce Gamma-Ray Emission (PIGE) for determining and quantifying low-Z elements such as F, Al, Li, ...
- Scanning Transmission Ion Microscopy (STIM) for evaluation of density distribution, via final energy of transmitted ions.
- **Ion Beam Induced Charge (IBIC)** for measuring and imaging the electron transport properties of semiconductor materials and devices.

Network	Coun- try	Access pro- vider	Infra- structure	Instrument	Techniques	Link
RADIATE	DE	HZDR	IBC	Helium Ion Microscopy, 100 kV Ion Platform, 6 MV Tandem Accelerator		<u>learn</u> more
RADIATE	IT	INFN	LABEC	3 MeV Tandetron	PIXE, PIGE, STIM	<u>learn</u> more
RADIATE	РТ	IST	IBL	2.5 MV Van de Graaff with Oxford microprobe	PIXE, RBS, IBIL	<u>learn</u> more
RADIATE	SI	JSI	MIC	Ion beam lines at Tandetron	PIXE, PIGE	learn more
RADIATE	FI	JYU	AL	1.7 MV Tandem Pelletron	PIXE, PIGE, IBIC	<u>learn</u> more
RADIATE	BE	KU Leuven	IMBL	1.7 MV Tandem Pelletron	PIXE, PIGE	<u>learn</u> more
RADIATE	CZ	NPI	LT	Ion beam lines at Tandetron	PIXE, STIM	<u>learn</u> more
RADIATE	HR	RBI	RBI-AF	1 MV Tandentron and 6 MV Tandem Van de Graaff	PIXE, PIGE, STIM	learn more learn more
RADIATE	ES	UAM	CMAM	5MV tandem + internal and external microbeam	PIXE, PIGE, STIM	learn more
RADIATE	SE	UU	Tandem Labarato ry	350 kV Implanter	PIXE	<u>learn</u> more



#### Laser photo chemistry & spectroscopy

Laser photo chemistry and spectroscopy includes a plethora of laser-based techniques to promote and probe chemical and physical events. Several steady-state and time-resolved spectroscopic techniques are available covering wide ranges of energy and time, applying a multitude of specialized sampling techniques.



#### Techniques

- Raman and infrared (IR) spectroscopies for vibrational studies, with micro-sampling techniques available, which include Raman and IR mapping/imaging.
- **Specialized IR spectroscopy** for low temperature matrix-isolation research, for characterization of short-living species, evaluation of photochemical mechanistic aspect of organic chemistry reactions, and studies of intramolecular energy redistribution. The technique is also applicable for direct observations of quantum mechanically-driven processes and their entanglement with vibrationally-induced processes.
- Chirped-pulse Fourier transform microwave (CP-FTMW) spectroscopy coupled with supersonic jet expansions sampling for structural studies, including of complex mixtures.
- **Photoacoustic calorimetry (PAC) and photoacoustic tomography (PAT)** for monitoring non-radiative molecular processes following photo-excitation and imaging.
- Transient absorption spectroscopy (TAS) in fs, ps and ns timescales ns laser flash photolysis and fs pump-probe spectroscopy set-ups for UV, Vis and NIR regions, for detection of transient species in different experimental conditions.
- **Fluorescence and phosphorescence spectroscopy** for studies in different phases, including solid state fluorescence quantum yield determinations.
- Scattering-type Scanning Near-field Optical Microscopy (s-SNOM) for materialcharacteristic maps of chemical and optical properties of the sample surface.
- Hyperspectral microscopy (HSM) based on a birefringent interferometry to acquire hyperspectral images in the visible and NIR ranges.
- Fluorescence Lifetime Imaging Microscopy (FLIM) a molecular fluorescence lifetime-based microscopy technique used to map the spatial distribution of biological cells and materials.
- **Reflectron time-of-flight mass spectrometer (RETOFMS):** for studies of ablation plasma composition, to determine elemental or isotopic signature of a sample and masses of particles and molecules.

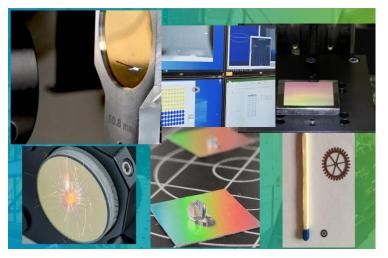
Network	Coun- try	Access provider	Infrastructure	Instrument	Link
Laserlab-	GR	FORTH	IESL-FORTH	time-resolved spectroscopy (UV-VIS-NIR-	learn more
Europe				THz-white light, fs-ps)	
Laserlab-	GR	FORTH	IESL-FORTH	micro-absorption/PL spectroscopy (4K-	learn mor
Europe				380K)	
Laserlab-	CZ	IP-ASCR	Hilase	Raman and AFM	learn mor
Europe					
Laserlab-	CZ	IP-ASCR	Hilase	RETOFMS	learn mor
Europe					
Laserlab-	ES	LLE-AISBL	CLPU	VEGA	learn mor
Europe					
Laserlab-	IT	LLE-AISBL	CUSBO	Hyperspectral imaging VIS-NIR-SWIR	learn mor
Europe					
Laserlab-	IT	LLE-AISBL	CUSBO	Hyperspectral microscope	learn mor
Europe					
Laserlab-	IT	LLE-AISBL	CUSBO	Time resolved fluorescence &	learn mor
Europe				microscopy	
Laserlab-	IT	LLE-AISBL	CUSBO	Ultrafast transient absorption	learn mor
Europe				•	
Laserlab-	ES	LLE-AISBL	ICFO	Attoseconds SXR beamline	learn
Europe	-	_			more,
					learn mor
Laserlab-	HR	LLE-AISBL	CALT	Home-made Raman spectrometer	learn mor
Europe Laserlab-	HR		CALT	Ti-Connhire Comtoscond Locar	loorn mor
	пк	LLE-AISBL	CALI	Ti:Sapphire Femtosecond Laser	learn mor
Europe Laserlab-	HR		CALT	Near field sSNOM	loorn mor
	пк	LLE-AISBL	CALI	Near field SSNOW	learn mor
Europe Laserlab-	NL		LLAMS	Stimulated Daman Coattoring	loorn mor
	INL	LLE-AISBL	LLAIVIS	Stimulated Raman Scattering	learn mor
Europe Laserlab-	NU			microscopy	
	NL	LLE-AISBL	LLAMS	Deep-UV Raman spectroscopy	learn mor
Europe	NU			Law for an an and the second sec	
Laserlab-	NL	LLE-AISBL	LLAMS	Low-frequency Raman spectroscopy	learn mor
Europe					
Laserlab-	PT	UC	CLL	Transient Absorption and	learn mor
Europe	DT			Photoaccoustics	lasur
Laserlab-	PT	UC	CLL	Fluorescence	learn mor
Europe				5118.4	
Laserlab-	PT	UC	CLL	FLIM	learn mor
Europe					
Laserlab-	PT	UC	CLL	Raman Spectroscopy	learn mor
Europe					
Laserlab-	PT	UC	CLL	Matrix-Isolation Infrared spectroscopy	learn mor
Europe					
Laserlab-	PT	UC	CLL	Single Photon Counting	learn mor
Europe					
Laserlab-	PT	UC	CLL	Vibrationally-Induced Photochemistry	learn mor
Europe					
Laserlab-	PT	UC	CLL	Rotational Spectroscopy (MRR and CP-	learn mor
Europe				FTMW)	

				and the second s		
non-ARIE	CZ	NanoEn- viCz	NanoEnvi		escence inverted confocal spinning nicroscope Olympus SpinSR10 12)	learn more
LEAPS	NL	SRU	FELIX	US 11	Ultrafast spectroscopy with IR/THz and table top laser sys- tems	learn more
LEAPS	NL	SRU	FELIX	US 12	IR pump-probe	learn more
LEAPS	NL	SRU	FELIX	US 10	Infrared spectroscopy combined with FTICR and ion mobility	learn more
LEAPS	NL	SRU	FELIX	US 3	Ultrafast spectroscopy with IR/THz and table top laser sys- tems	<u>learn more</u>
LEAPS	NL	SRU	FELIX	US 9	Infrared spectroscopy combined with mass spectrometry	learn more
LEAPS	NL	SRU	FELIX	FELICE beamline	Cluster spectroscopy	learn more



#### Laser processing

Laser processing is primarily focused on surface material modification, assessment. testing and Often nanosecond and shorter laser pulses are utilized for treatment of various materials, while available services are not limited on material processing itself but include damage testing and general laser ablation field and related phenomena. Equipment and facilities thus can be used for variety supporting activities, laser as processing window assessment, ablation rate studies or process optimization and materials qualification for space environment. Also, in combination with ultra-short pulses and high energy lasers, facilities can be used for laser plasma generation and EUV/XUV secondary



Top: laser multibeam processing of substrate, diagnostics of the beam and diffractive structure written on alloy. Bottom: laser damage of thin film, superhydrophobic surface prepared by laser surface processing and miniature composite cogwheels prepared by laser micromachining. Image credit to Jan Vanda.

source development or particle acceleration.

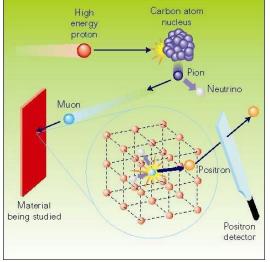
#### Techniques

- Laser Shock Peening (LSP) increase of stress corrosion cracking resistance via plasmainduced deep compressive stresses.
- Laser Micro-Machining (LMM) material structuring for surface functionalization (hydrophobicity/hydrophilicity, tuned friction, bio-compatibility, optical properties).
- Laser Induced Damage Threshold (LIDT) laser resistance testing, laser ablation rate and laser processing window research.
- Pulsed Laser Deposition (PLD) thin film deposition at high vacuum with possibility of assistant gas
- Ultrafast laser ablation (ULA) and micromaterial processing (MMP)
- 2D and 3D sub-micron structures processing for tailored physical and biological properties
- Laser additive manufacturing (LAD) and photochemical modification (LPM) for micro- and nano- scale structures

Network	Country	Access provider	Infrastructure	Instrument	Link
Laserlab-	GR	FORTH	IESL-FORTH	material micro-/nano 2D & 3D pro-	learn
Europe				cessing	<u>more</u>
Laserlab-	GR	FORTH	IESL-FORTH	additive manufacturing and photochem-	learn
Europe				ical modification	<u>more</u>
Laserlab-	CZ	IP-ASCR	HILASE	BIVOJ, nanosecond, 100 J, 1 kW, 1030	learn
Europe				nm	<u>more</u>
Laserlab-	CZ	IP-ASCR	HILASE	Laser shock peening station	<u>learn</u>
Europe					<u>more</u>
Laserlab-	CZ	IP-ASCR	HILASE	LIDT station	<u>learn</u>
Europe					more
Laserlab-	CZ	IP-ASCR	HILASE	Micromachining station	<u>learn</u>
Europe					more
					<u>learn</u>
					more
Laserlab-	CZ	IP-ASCR	HILASE	PERLA B, picosecond, 10 mJ, 100 W,	learn
Europe				1030 nm	more
Laserlab-	CZ	IP-ASCR	HILASE	PERLA C PERLA C, picosecond, 5 mJ, 500	learn
Europe				W, 1030 nm	more
Laserlab-	CZ	IP-ASCR	HILASE	PLD workstation for thin film deposition	learn
Europe				and laser ablation plume analysis	<u>more</u>
Laserlab-	ES	LLE-	CLPU	LAB2 - ULAMP	<u>learn</u>
Europe		AISBL			<u>more</u>
Laserlab-	IT	LLE-	ENEA	ABC Laser Facility	<u>learn</u>
Europe		AISBL			<u>more</u>
Laserlab-	IT	LLE-	ENEA	CETRA Facility	learn
Europe		AISBL			<u>more</u>
Laserlab-	HR	LLE-	CALT	Nd:YAG laser	learn
Europe		AISBL			more
Laserlab-	HR	LLE-	CALT	PLD chamber	learn
Europe		AISBL			more
non-ARIE	CZ	NanoEn-	NanoEnviCz	Industrial femtosecond pulsed laser	learn
		viCz		(TUL13)	more
non-ARIE	CZ	NanoEn-	NanoEnviCz	MicroWriter ML3 Pro (UFCH25)	<u>learn</u>
		viCz			more
non-ARIE	CZ	NanoEn-	NanoEnviCz	Laser scanning confocal microscop	learn
		viCz		(UPOL15)	more
non-ARIE	CZ	NanoEn-	NanoEnviCz	Confocal microscope - LEICA CLSM	learn
		viCz		SP8/DLS (UJEP40)	more

#### Muon Spectroscopy

Muons provide a complementary probe of materials to neutrons. Muons are short-lived heavy versions of the electron. Fully spinpolarised muons are implanted into materials where they sense the local magnetic fields and the polarisation of the muon ensemble responds to these. This makes muons extremely sensitive to magnetism and superconductivity effects. They can also be used to study ionic diffusion, e.g. in battery materials, and can be used as mimics of isolated hydrogen to investigate hydrogen behaviour in materials. Muon studies of materials can be performed at the ISIS Neutron and Muon Source.



#### Life of a Muon. Source: https://nmi3.eu/muon-research/characteristics-ofmuons.html

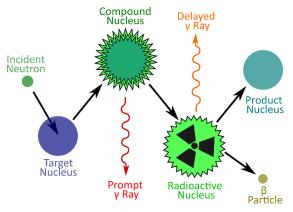
Network	Country	Access provider	Infrastru cture	Instru ment	Technique	Link
LENS	GB	UKRI	ISIS <sup>1</sup>	MuSR	longitudinal and transverse measure- ments	<u>learn</u> more
LENS	GB	UKRI	ISIS <sup>1</sup>	EMU	zero field and longitudinal field measurements, magnetism and ion diffusion in solids	<u>learn</u> more
LENS	GB	UKRI	ISIS <sup>1</sup>	HiFi	applied longitudinal fields up to 5T	<u>learn</u> more

<sup>&</sup>lt;sup>1</sup> LIMITED ACCESS ONLY for proposals requiring environments ONLY available at ISIS. All other requests will be transferred to BNC (Hungary) or SINQ/PSI (Switzerland).



#### Neutron-based elemental composition analysis

Neutron activation analysis methods use the neutron-induced transient radioactivity or nuclear reactions to obtain information on the elemental composition of samples. Two basic types of neutron activation analysis exist from a practical point of view - Neutron Activation Analysis (NAA) and Prompt Gamma Activation Analysis (PGAA, or PGNAA). If the irradiation and the detection of neutron-induced radioactivity are separated in time and space, this is the "traditional", Instrumental NAA (INAA), suitable to trace element analysis. In PGAA, the neutrons are transferred to the sample in form of a guided beam, and the irradiation and the detection of gamma-rays take place simultaneously.



Principle of neutron activation analysis. Image credit to Christina Ossig.

These techniques are non-destructive, bulk-representative, applicable to materials where "exotic" elements (such as light elements (H, B, Cl, S), valuable noble metals, rare-earth elements, environmentally-relevant heavy metals) are to be quantified, if the material is not soluble, or where standard reference material are unavailable. They can be combined for an almost panoramic analysis. A relevant application example is to follow material recovery yields during subsequent stages of production or reprocessing.

#### Techniques

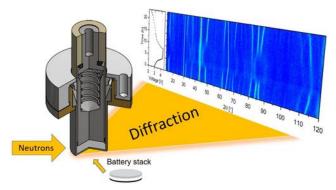
- Neutron Activation Analysis (NAA) For trace elemental analysis
- **Prompt Gamma Activation Analysis (PGAA)** For average elemental composition
- Neutron Depth Profiling (NDP) For near-surface analysis of concentration of light elements
- **Prompt-Gamma Activation Imaging (PGAI)** For determination of the composition and the spatial distribution of traced elements
- **Neutron Radiography (NR)** utilizes the transmission of neutrons and photons to obtain visual information on the structure and/or dynamic processes inside of an object
- Neutron Tomography (NT) For 3D spatial resolution
- Prompt-Gamma Irradiation (PGI) For qualitative and quantitative elemental analysis

Network	Country	Access provider	Infrastructure	Instrument	Technique	Link
LENS	HU	EK	BNC	NAA	NAA	learn more
LENS	HU	EK	BNC	NIPS-NORMA	PGAI, NR, NT, PGAA	<u>learn more</u> learn more
LENS	HU	EK	BNC	PGAA	PGAA	learn more
LENS	DE	TUM	FRM II	FaNGaS	Irradiation of large samples, PGI	<u>learn more</u>
LENS	DE	TUM	FRM II	NAA	NAA	learn more
LENS	DE	TUM	FRM II	PGAA	PGAA	learn more



#### Neutron diffraction

diffraction Neutron reveals structural information on the arrangement of atoms and magnetic moments in condensed matter. Single-crystal diffraction provides the most precise and detailed information but requires crystal samples of suitable quality and size. Otherwise samples exist in a form in which some of the structural information is spatially averaged and the corresponding experimental technique is here referred to generically as 'powder diffraction'. It however includes diffraction on liquids, biological samples (e.g. membranes) and engineering components. In the latter case the measurement is focussed to



Specific sample cells allow the measurement of inoperando neutron diffraction patterns of battery materials. source: https://doi.org/10.1002/cmtd.202200046

determine atomic distances of a well-known structure within a given small gauge volume. Scanning a region of interest enables the determination of stress fields inside the component.

#### Techniques

- Single-crystal neutron diffraction for specific structural information
- Powder neutron diffraction for average structural information
- Zero-field spherical neutron polarimetry (SNP) for magnetic structure determination
- Polarised neutron diffraction (PND) in magnetic field named also Flipping-Ratio method
- Non-polarised diffraction under special conditions (very low temperatures, magnetic and electric fields, high pressures, high temperatures and their combinations) using dedicated sample environments and out-of-plane lifting counter

Network	Coun- try	Access provider	Infrastru cture	ı Instrumen t	Technique	Link
LENS	HU	EK	BNC	ATHOS	phonon dispersion relations, tunneling mode studies, quasielastic scattering studies of rotational and non-local diffu- sion, vibrations of surfaces or molecules adsorbed on surfaces, phonon density of states, neutron scattering to the study of hydrogenous materials	<u>learn</u> more
LENS	HU	EK	BNC	MTEST	material testing diffractometer, powder, liquid and amorphous total diffraction	learn more
LENS	HU	EK	BNC	PSD	atomic structure investigations, amor- phous, liquids and crystalline materials	learn more
LENS	FR	ILL	ILL	Neutron single- crystal diffrac- tion instru- ments D3, D9, D10, D15, D19, D23, VIVALDI		learn more

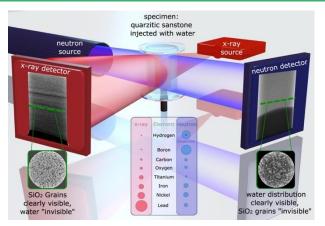
LENS	FR	ILL	ILL	Neutron		learn
				powder		more
				diffraction		
				instru-		
				ments		
				D1B, D2B,		
				D4, D7,		
				D20		
LENS	FR	ILL	ILL	SALSA	Neutron stress / strain diffraction instru-	learn
					ment	more
LENS	СН	PSI	SINQ	HRPT,	High-Resolution Powder Diffractometer	learn
				DMC,	for Thermal Neutrons, Cold Neutron	more
				Zebra,	Powder Diffractometer, Single Crystal	
				POLDI	Neutron Diffractometer, Time-Of-Flight	
					Neutron Diffractometer	
LENS	DE	TUM	FRM II	BioDiff	Protein crystallography, structure deter-	learn
					mination of biological macromolecules	more
LENS	DE	TUM	FRM II	Heidi, Poli	Single crystal diffractometer on hot	learn
					source, SNP, PND	more
						learn
						more
LENS	DE	TUM	FRM II	Spodi	HR- powder diffraction	learn
						more
LENS	DE	TUM	FRM II	StressSpec	Stress and texture measurements	learn
						more
LENS	GB	UKRI	ISIS <sup>2</sup>	Neutron		learn
				diffraction		more
				instrument		
				S		
LENS	GB	UKRI	ISIS <sup>2</sup>	Engin-x	Neutron stress / strain diffraction instru-	learn
					ment	more

<sup>&</sup>lt;sup>2</sup> LIMITED ACCESS ONLY for proposals requiring environments ONLY available at ISIS. All other requests will be transferred to BNC (Hungary) or SINQ/PSI (Switzerland).



#### Neutron imaging

Neutron imaging is a non-destructive technique, highly complementary to X-ray imaging, that can see inside materials and examine processes therein. White beam imaging is based on the attenuation of the neutron beam, due to absorption or scattering, through an object. Grating interferometry is sensitive to materials properties such as porosity down to the micrometer-scale. Polarised neutron imaging reveals magnetic domains and textures. Monochromatic and energy-resolved imaging enhances element specific contrasts or diffraction contrast of materials. Tomography is performed by rotating the sample and reconstructing the 3-dimensional volume from a



Neutron and X-ray imaging can be performed in parallel, e.g., at NeXT/ILL. Source: https://doi.org/10.1016/j.nima.2020.163939

series of images. The high sensitivity to hydrogen containing materials reveals even small contaminations. In some instruments, such as NeXT at ILL, neutrons and X-rays can be used in parallel.

#### Techniques

- Neutron Radiography (NR) / Tomography (NT) are methods to provide 2D projections (radiographs) and 3D reconstructed volumes of attenuation of specimens. NR and NT are tools to investigate internals such as features, structures, cracks, and defects of a sample for spatial resolutions as low as 10 μm.
- Energy resolved neutron imaging (ERNI) provides information about element concentrations (via resonance analysis) or microstructure information such as phase composition and residual strain (via Bragg edge analysis) typically in 2D (although possible in 3D) for spatial resolutions of hundreds of microns.
- Grating interferometry (GI) or Dark Field Imaging (DFI) produces maps of attenuation, phase contrast and ultra-small angle scattering (dark field) signals, providing information about concentrations, porosity, and magnetic domains, for real-space spatial resolutions of tens or hundreds of microns.
- **Combined imaging with X-rays or gamma-rays** bi-modal approach that enhances material contrast by taking advantaging different sensitivities of X-rays and neutrons for different elements and isotopes.
- **Prompt gamma activation imaging (PGAI)** provides spatially resolved element compositions for a large number of elements and isotopes, for resolutions of hundreds of microns.
- Neutron diffraction mapping (ND) provides spatially resolved composition, strain and texture information, typically for 1D or 2D real space resolutions of hundreds of microns up to a few millimetres.

Network	Coun- try	Access provider	Infra- structure	Instrument	Techniques	Link
LENS	HU	EK	BNC	NORMA	PGAI-NT	<u>learn</u> more
LENS	HU	EK	BNC	RAD	Static/dynamic white-beam- neutron and X-ray imaging sta- tion	<u>learn</u> more
LENS	FR	ILL	ILL	Neutron imaging instrument NeXT	Imaging and in-situ X-ray imag- ing	<u>learn</u> more
LENS	СН	PSI	SINQ	ICON, NEUTRA	HR-tomography, Grl, ERI	<u>learn</u> more
LENS	DE	TUM	FRM II	Antares	Cold neutron radiography and tomography facility	<u>learn</u> more
LENS	DE	TUM	FRM II	Nectar	fission neutron radiography and tomography	<u>learn</u> more
LENS	СН	PSI	SINQ	ICON, NEUTRA	HR-tomography, Grl, ERI	<u>learn</u> more
LENS	BG	UKRI	ISIS <sup>3</sup>	Neutron imaging instrument IMAT	neutron radiography, neutron tomography, and energy-re- solved neutron imaging	<u>learn</u> more

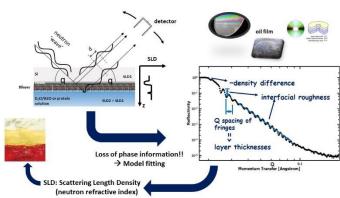
<sup>&</sup>lt;sup>3</sup> LIMITED ACCESS ONLY for proposals requiring environments ONLY available at ISIS. All other requests will be transferred to BNC (Hungary) or SINQ/PSI (Switzerland).



#### Neutron reflectometry

Neutron reflectometry (NR) gives information on the structure of surfaces and interfaces (depth-dependent composition). It is also a powerful technique to study air/solid, solid/solid, solid/liquid, liquid/liquid and liquid/air interfaces.

Reflectivity is the ratio of the reflected intensity to the incident intensity for a beam directed onto an interface or surface. The technique



Neutron Reflectivity - an interference phenomenon

Schematic of neutron reflectivity. Image credit to Philip King.

provides valuable information over a wide variety of scientific and technological applications including chemical aggregation, polymer and surfactant adsorption, structure of thin film magnetic systems, biological membranes, etc.

In the simplest case contrast matching can be employed to isolate the reflected signal from a particular adsorbate within a mixture. The signal is directly proportional to the adsorbed amount and often the NR technique is the only way such quantitative information can be obtained. Building on this idea specific deuteration can be employed to vary the refractive index of components both intra and inter molecular. A set of reflectivity data for the same chemical or biological system is obtained and used to constrain a real space model of the molecular organisation with a resolution of ~0.2nm or better.

#### Techniques

- Reflectometry: chemical composition (depth profile) at interfaces
- Polarised Neutron Reflectometry: magnetic depth profile at interfaces

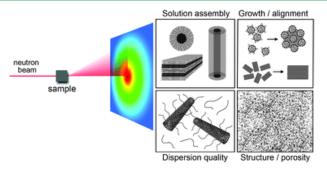
Networ k	Coun- try	Access provider	Infra- structure	Instrument	Techniques	Link
LENS	HU	EK	BNC	GINA	Neutron Reflectometer with Polarized Beam Option	learr more
LENS	HU	EK	BNC	REF	neutron reflectometry	learr more
LENS	FR	ILL	ILL	Neutron reflectome- try instruments D17, Figaro, SuperAdam, D16	neutron reflectometry	<u>learr</u> more
LENS	СН	PSI	SINQ	AMOR	neutron reflectometry	learr more
LENS	DE	TUM	FRM II	Maria, N-ReX, Ref- sans	VR with HIA, Polarized, ver- tical neutron reflectometry, Horizontal ToF	learr more
LENS	GB	UKRI	ISIS <sup>4</sup>	Neutron reflectome- try instruments (In- ter, Offspec, Polref, Surf)	neutron reflectometry	<u>learr</u> more

<sup>&</sup>lt;sup>4</sup> LIMITED ACCESS ONLY for proposals requiring environments ONLY available at ISIS. All other requests will be transferred to BNC (Hungary) or SINQ/PSI (Switzerland).



#### Neutron small angle scattering

Small-angle neutron scattering (SANS) snapshots the collective characteristics of particles and molecules in dispersed or assembled systems, rather than looking at single atoms or molecules. The signal is based on the contrast of the targeted matter and the background. Due to the distinctive scattering length density (SLD) of light elements of similar atomic numbers, neutron scattering will enable differentiation of species abundant in light elements.



Typical information extractable from the data acquired at a SANS instrument.

Source: https://doi.org/10.1039/C3CP50293G

As the scattered neutrons show information as

in the reciprocal space (or q-space) instead of real space, the larger the target objects are, the smaller the scattering angle we get. Since the lower end of q is limited by the beamline setup, typically, the length scale of structures that can be investigated using SANS ranges from a few nanometres to hundreds of nanometres. With a spin-echo SANS (SESANS) setup, one may extend the range to characterise larger structures up to tens of microns. A wide scope of systems can be studies using SANS, including soft matters (gels and colloids), biological materials, and magnetic materials, with controlled environment enabling us to study materials structure under (not limited to) different temperature, pressure, moisture, and mechanically stressed conditions.

Additionally, using isotopic substitution (e.g., hydrogen (H) to deuterium (D), called "deuteration"), one can alter the scattering signals of hydrogenated groups in the molecules. This contrast variation method is based on the significant difference in the SLD of the isotopes, and is widely used in hydrogenrich systems such as soft matters and biological samples (e.g., protein binding). This allows us to render specific parts of the sample with minimal interference from the other parts.

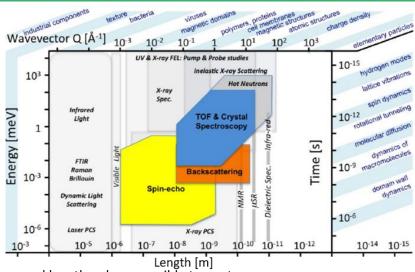
Infrastructures							
Networ k	Coun- try	Access provider	Infra- structure	Instrument	Technique	Link	
LENS	HU	EK	BNC	F-SANS	SANS	learn more	
LENS	HU	EK	BNC	Yellow Submarine	SANS	learn more	
LENS	France	ILL	ILL	Small angle scattering instru- ments (D11, D22, D33, D16)	SANS	learn more	
LENS	СН	PSI	SINQ	SANS-I	SANS	learn more	
LENS	DE	TUM	FRM II	KWS-1, KWS-2, KWS-3, SANS- 1	SANS	learn more	
LENS	GB	UKRI	ISIS <sup>5</sup>	Small angle scattering instru- ments (Zoom, Sans2d, LoQ, Larmor)	SANS	<u>learn more</u>	

### <sup>5</sup> LIMITED ACCESS ONLY for proposals requiring environments ONLY available at ISIS. All other requests will be transferred to BNC (Hungary) or SINQ/PSI (Switzerland).



#### Neutron spectroscopy

Neutron spectroscopy probes the dynamics of magnetic moments, atoms, molecules or atom lattices over length scales ranging from fractions of a nanometer to tens of nanometers, and over timescales from tens of femtoseconds (molecular vibrations) up to the microsecond (motion of large biological molecules). Within neutron spectroscopy, there are 4 main techniques which use different methods to determine the energy of the incident and scattered neutrons and are adapted to different kinds of scientific studies.



Time- and lengthscales accessible to neutrons.

https://europeanspallationsource.se/sites/default/files/downloads/2017/09/TDR\_online\_ver\_all.pdf

#### Techniques

- Time-of-flight (ToF) spectroscopy / quasi-elastic spectroscopy (QENS) Surveys of lattice and magnetic dynamics or atom diffusion on the ps timescale.
- Vibrational spectroscopy for assessing molecular bonds, typically in the fs timescale.
- **Triple-axis spectroscopy (TAS)** More focussed studies of lattice and magnetic dynamics in the ps timescale.
- High resolution neutron backscattering (HR-BS) / neutron spin-echo (NSE) spectroscopy for slow and diffuse motions

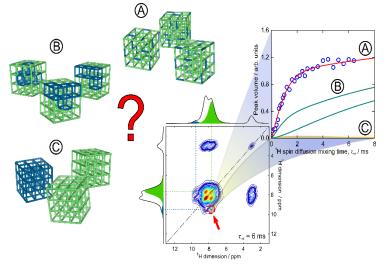
Networ k	Coun- try	Access provider	Infra- structure	Instrument	Technique	Link
LENS	FR	ILL	ILL	IN13, IN15, IN16B, WASP	Neutron spectroscopy – high resolution	<u>learn</u> more
LENS	FR	ILL	ILL	IN5, IN8, IN12, IN20, IN22, SHARP, Thales	Neutron spectroscopy medium resolution	<u>learn</u> more
LENS	FR	ILL	ILL	IN1-Lagrange, Panther	Neutron spectroscopy - vibra- tional spectroscopy	<u>learn</u> more
LENS	СН	PSI	SINQ	FOCUS, TASP, CAMEA, EIGER	ToF for cold neutrons, thermal triple-axis, cold triple-axis	<u>learn</u> more
LENS	DE	TUM	FRM II	Kompass, La- Diff, Puma, Panda, Trisp	lattice excitations and magnetic excitations	<u>learn</u> more
				TOFTOF	ToF	_
				Spheres	HR-BS	_
				NSE, Reseda	Spin-echo spectrometers	
LENS	GB	UKRI	ISIS <sup>6</sup>	MAPS, MARI, LET, MERLIN	Neutron spectroscopy – excita- tions instruments	<u>learn</u> more
LENS	GB	UKRI	ISIS <sup>6</sup>	IRIS, OSIRIS, TOSCA, VESU- VIO	Neutron spectroscopy – molec- ular spectroscopy	<u>learn</u> more

<sup>&</sup>lt;sup>6</sup> LIMITED ACCESS ONLY for proposals requiring environments ONLY available at ISIS. All other requests will be transferred to BNC (Hungary) or SINQ/PSI (Switzerland).



### Nuclear Magnetic Resonance

NMR is an abbreviation for Nuclear Magnetic Resonance. The advantages of NMR for the analysis of molecular structures at the atomic level are that sample measurements are nondestructive and that there is little sample preparation required. NMR spectroscopy is a versatile tool that provides information not only on the structures, but also on the dynamics of various biological and synthetic molecules at an atomic level. The investigated samples are put in a magnetic field that is tens of thousands of times stronger than the earth's magnetic field. The NMR method is very sensitive to the features of molecular structure because the neighboring atoms



Linker molecules in metal-organic frameworks can be distinguished as shown in the work of Krajnc et al. Image adapted from: A. Krajnc et al., (2015), Angew. Chem. Int. Ed., 54: 10535-10538. https://doi.org/10.1002/anie.201504426

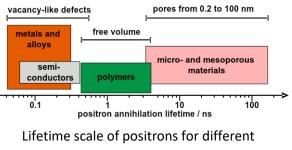
influence the signals from individual nuclei as well and this is important for determining the 3Dstructure of molecules. With NMR spectroscopy one can study liquid, solid and semi-liquid samples. Fields of application include bio, foods, chemistry, as well as new ones such as batteries.

Network	Country	Access provider	Infrastructure	Instrument	Link
non-ARIE	EU	CERIC-ERIC	CERIC-ERIC	ASKA@SloNMR	<u>learn more</u>
non-ARIE	EU	CERIC-ERIC	CERIC-ERIC	DAVID@SloNMR	learn more
non-ARIE	EU	CERIC-ERIC	CERIC-ERIC	LARA@SIoNMR	learn more
non-ARIE	EU	CERIC-ERIC	CERIC-ERIC	MAGIC@SloNMR	learn more
non-ARIE	EU	CERIC-ERIC	CERIC-ERIC	NIKA@SloNMR	learn more
non-ARIE	EU	CERIC-ERIC	CERIC-ERIC	ORO@SloNMR	learn more
non-ARIE	CZ	NanoEnviCz	NanoEnviCz	Solid State NMR Spectrometer	learn more
				Jeol (TUL15)	
EMFL	DE	HZDR	HLD	Pulsed-field magnet	<u>learn more</u>



### Positrons

Being the anti-particle of electrons, positrons are used to probe material defects on the nanometre scale, at low concentrations and with high sensitivity. It is a non-destructive method developed to serve as a proven tool for the study of metals, semiconductors, polymers, and open or closed microporous systems. Positron based techniques are used to study a variety of phenomena and material properties on a nanometre scale, like:



materials. Image credit to Eric Hirschmann.

- performance parameters in semiconductors or alloys by characterization of **atomic defects**, there types (e.g. mono-/ di- vacancies) or concentration.
- optimization of process parameters in e.g. polymer membranes or composites regarding **free/open volume** effects (e.g. interstitial volume in polymer chains due to fatigue).
- determination of **pore size and pore size distribution (up to 100 nm diameter)** in e.g. nano filters or catalysts for high-performance or innovative applications.

It is recommended to discussed sample size, sample preparation as well as in-situ / operando options with the experts.

### Techniques

- **Doppler broadening spectroscopy (DBS)** electron momentum at annihilation site for the investigation of defect decoration and concentration
- Positron annihilation lifetime spectroscopy (PALS) electron density at annihilation site for the investigation of defect size/type and concentration
- **Positron Auger Spectroscopy (PAES)** Positron initiated emission of Auger electrons for chemical analysis of the near-surface area

Netw ork	Coun- try	Access provider	Infra- structure	Instrument	Techniques	Link
ELBE	DE	HZDR	ELBE	Monoenergetic Positron Source (MePS)	PALS, DBS	<u>learn</u> more
ELBE	DE	HZDR	ELBE	Slow-Positron System of Rossendorf (SPONSOR)	DBS for thin films and depth profiling up to 3000 nm	<u>learn</u> more
ELBE	DE	HZDR	ELBE	Conventional Positron Spectroscopy (CoPS)	PALS for bulk solids and pow- ders, in-situ temperature (30 - 510 K) and humidity measure- ments	<u>learn</u> more
LENS	DE	TUM	FRM II	Nepomuc	Pulsed low-energy positron system (PLEPS) Coincident Doppler Broaden- ing Spectrometer (CDBS)	<u>learn</u> more

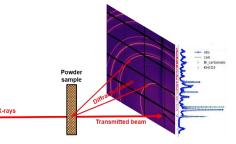


# X-ray diffraction

X-ray diffraction (XRD) enables the identification of crystalline materials by their atomic structure.

The high flux of synchrotron sources allows for fast data acquisition, and the use of high energy photons (hence, high transmittivity through samples) enables operando/in situ studies in real operating systems (e.g., electrochemical cells).

Through the analysis of the XRD data, one can retrieve the different phase(s) in the sample from the peaks position, while the intensities of the peaks provide information about the quantity of each crystalline phase present in the sample.



Scheme of an X-ray beam scattered onto a detector. Image credit to Marta Mirolo.

Finally, the analysis of the peak broadening provides insights into the crystallite size and microstrain in the crystal.

### Techniques

- High-resolution XRD for the identification of an unknown phase
- High space- and time-resolution XRD to identify the onset of a reaction
- Microdiffraction XRD to study the behaviour of single particles
- **High-energy XRD** for operando/in situ or high-throughput materials screening
- Surface XRD for the investigation of layers growth on single crystals or flat surfaces
- Laboratory XRD for ex situ materials screening

Network	Coun- try	Access provider	Infra- structure	Instrument	Technique	Link
LEAPS	ES	ALBA-	ALBA	MSPD	High-angular resolution and high-	learn
		CELLS			throughput Powder Diffraction,	more
					High-pressure Micro Diffraction	
LEAPS	DE	HZB	BESSY II	KMC-2	Diffraction	learn
						<u>more</u>
LEAPS	DE	DESY	PETRA III	P02.1	Powder diffraction, Bragg diffrac-	learn
					tion, PDF analysis	<u>more</u>
LEAPS	DE	DESY	PETRA III	P07	GISAXS, GIWAXS	Learn
						<u>more</u>
		DESY				loorn
LEAPS	DE	(GEMS-	PETRA III	P07 <sup>7</sup>	XRD, 3D-XRD, mirco-tomography	learn
		Hereon)				more
LEAPS	DE	DESY	PETRA III	P08	HRXRD	learn
						more
LEAPS	DE	DESY	PETRA III	P10	XPCS, CDI, Bragg CDI, Holographic	learn
					imaging	more

<sup>&</sup>lt;sup>7</sup> For SME-access only feasibility studies with up to max. 4 hours beamtime are possible.

LEAPSITELETTRAELETTRAMCXNon-single crystal XRD, grazing angle diffraction and reflectivity, residual stress and texture analy- sis, phase analysis, kinetic studiesLEAPSFRESRFESRFID01CDI, GISAXS, XRD, Ptychographylearr moreLEAPSFRESRFESRFID11Diffraction contrast tomography, aD-XRD, Imaging, PDF analysis, Ptychography, SAXS, XRDlearr moreLEAPSFRESRFESRFID13µXRF, µCrystallography, Ptychography, SAXS, XRDlearr moreLEAPSFRESRFESRFID15ADCT, EDD, Imaging, Laminogra- phy, PDF analysis, Pump-probe, SAXS, TR-WAXS, X-ray scatteringlearr moreLEAPSFRESRFESRFID22XRD, Powder diffraction, PDF analysis, anomalous diffraction, more sAXS, GIDlearr more sAXS, GISAXS, PDF analysis, more sAXS, GISAXS, PDF analysis, more sAXS, GIDlearr more sAXS, GIDLEAPSFRESRFESRFID22XRD, Powder diffraction, more sAXS, GIDlearr more sAXS, GIDLEAPSFRESRFESRFID31XRD, Compton scattering, XRR, wAXS, GISAXS, PDF analysis, more sAXS, GIDlearr more sAXS, GIDLEAPSFRSOLEILSOLEILPSICHEEDXRF, ADXRD, Tomography more sAXS, GIDLEAPSFRSOLEILSOLEILPSICHEEDXRF, ADXRD, Tomography more sAXS, GIDLEAPSFRSOLEILSOLEILSOLEILSOLEILCHPSICRELab XRD <th>RADIATE</th> <th>РТ</th> <th>IST</th> <th>IBL</th> <th>Lab XRD (Bruker D8)</th> <th>Powder diffraction</th> <th>learn more</th>	RADIATE	РТ	IST	IBL	Lab XRD (Bruker D8)	Powder diffraction	learn more
LEAPS         FR         ESRF         ESRF         ID01         CDI, GISAXS, XRD, Ptychography more powder diffraction, XRD         learr more more powder diffraction, XRD           LEAPS         FR         ESRF         ESRF         ESRF         ID11         Diffraction contrast tomography, powder diffraction, XRD         learr more powder diffraction, XRD           LEAPS         FR         ESRF         ESRF         ESRF         ID13         µXRF, µCrystallography, Pychography, SAXS, XRD, Purp-probe, SAXS, TR-WAXS, X-ray scattering         learr phy, PDF analysis, powder diffraction, PDF analysis, anomalous diffraction, PDF analysis, anomalous scattering         learr more analysis, anomalous diffraction, PDF analysis, anomalous scattering           LEAPS         FR         ESRF         ESRF         ID31         XRD, Compton scattering, XRR, WAXS, GISAXS, PDF analysis, sAXS, GID         learr more analysis, anomalous diffraction, PDF analysis, anomalous diffraction, MRR, WAXS, GISAXS, VDF), QEXAFS         learr more more sAXS, GID           LEAPS         FR         ESRF         ISLS         Debye         XAS, XRD, SAXS, (PDF), QEXAFS         learr more more sAXS, GID           LEAPS         FR         SOLEIL         SOLEIL         PSICHE         EDXRF, ADXRD, Tomography         learr more more sAXS, GID           LEAPS         FR         SOLEIL         SOLEIL         PSICHE         ZD, D XRD, Tomography         learr more more se	LEAPS	IT	ELETTRA	ELETTRA	· · ·	angle diffraction and reflectivity, residual stress and texture analy-	learn more
LEAPSFRESRFESRFID11Diffraction contrast tomography, powder diffraction, XRDlear yowder diffraction, XRDLEAPSFRESRFESRFID13µXRF, µCrystallography, Ptychography, SAXS, XRDmorLEAPSFRESRFESRFID15ADCT, EDD, Imaging, Laminogra- phy, PDF analysis, Pump-probe, SAXS, TR-WAXS, X-ray scatteringlear phy, PDF analysis, Pump-probe, SAXS, TR-WAXS, X-ray scatteringLEAPSFRESRFESRFID22XRD, Powder diffraction, mor anomalous scatteringLEAPSFRESRFESRFID31XRD, Compton scattering, XRR, uansulous scatteringLEAPSFRSOLEILSOLEILPSICHEEDXRF, ADXRD, Tomography mor SAXS, GIDLEAPSFRSOLEILSOLEILPSICHEEDXRF, ADXRD, Tomography mor SAXS, GIDLEAPSFRSOLEILSOLEILPSICHEEDXRF, ADXRD, Tomography mor sAXS, GIDLEAPSFRSOLEILSOLEILPSICHEEDXRF, ADXRD, Tomography mor saves, latice parameter refine- ment, X-ray reflectivity and SAXSe-DREAMITCNRXRD LabZD, 1D XRD, temperature stage mor fractometer powder diffraction, X-ray powder micro-diffraction, quanti- tative phase analysis, qualitative phase analysisnon-ARIECZNanoEnviNanoEnviX-ray powder diffraction, X-ray powder micro-diffraction, quanti- tative phase analysis, qualitative phase analysisnon-ARIECZNanoEnviX-ray Pow- c cX-ray powder diffraction, phase <b< td=""><td>LEAPS</td><td>FR</td><td>ESRF</td><td>ESRF</td><td>ID01</td><td></td><td>learn more</td></b<>	LEAPS	FR	ESRF	ESRF	ID01		learn more
LEAPSFRESRFESRFID13μXRF, μCrystallography, Ptychography, SAXS, XRD Ptychography, SAXS, XRD phy, PDF analysis, Pump-probe, SAXS, TR-WAXS, X-ray scatteringleart more phy, PDF analysis, Pump-probe, SAXS, TR-WAXS, X-ray scatteringLEAPSFRESRFESRFID22XRD, Powder diffraction, PDFleart more analysis, anomalous diffraction, PDFLEAPSFRESRFESRFID31XRD, Compton scattering, XRR, MAXS, GISAXS, PDF analysis, more sAXS, GIDleart more anomalous scattering, XRR, SAXS, GIDleart more more sAXS, GIDLEAPSFRESRFESRFID31XRD, Compton scattering, XRR, MAXS, GISAXS, PDF analysis, more SAXS, GIDleart more more sAXS, GIDLEAPSFRSOLEILSOLEILPSICHEEDXRF, ADXRD, Tomographyleart more more tallinity, crystallite size and lattice stress, lattice parameter refine- ment, X-ray reflectivity and SAXSe-DREAMITCNRCNRXRD Lab2D, 1D XRD, temperature stage powder dif- 	LEAPS	FR	ESRF	ESRF	ID11	3D-XRD, Imaging, PDF analysis,	learn more
LEAPSFRESRFESRFID15ADCT, EDD, Imaging, Laminogra- phy, PDF analysis, Pump-robe, SAXS, TR-WAXS, X-ray scatteringleart onandous diffraction, PDF 	LEAPS	FR	ESRF	ESRF	ID13	μXRF, μCrystallography,	<u>learn</u> more
LEAPSFRESRFESRFID31XRD, Compton scattering, XRR, lear WAXS, GISAXS, PDF analysis, SAXS, GIDLEAPSCHPSISLSDebyeXAS, XRD, SAXS, (PDF), QEXAFSlear more sAXS, GIDLEAPSCHPSISLSDebyeXAS, XRD, SAXS, (PDF), QEXAFSlear 	LEAPS	FR	ESRF	ESRF	ID15A	DCT, EDD, Imaging, Laminogra- phy, PDF analysis, Pump-probe,	learn more
LEAPSCHPSISLSDebyeXAS, XRD, SAXS, (PDF), QEXAFSlearrLEAPSCHPSISOLEILSOLEILPSICHEEDXRF, ADXRD, TomographylearrLaserlab- EuropeCZIP-ASCRHiLASELab XRDXRD, (RIR) phase analysis, of errys- tallinity, crystallite size and lattice stress, lattice parameter refine- ment, X-ray reflectivity and SAXSlearr more tallinity, crystallite size and lattice 	LEAPS	FR	ESRF	ESRF	ID22	analysis, anomalous diffraction,	<u>learn</u> more
LEAPSFRSOLEILSOLEILPSICHEEDXRF, ADXRD, Tomographylearr moorLaserlab- EuropeCZIP-ASCRHiLASELab XRDXRD, (RIR) phase analysis, reitveld analysis, analysis of crys- 	LEAPS	FR	ESRF	ESRF	ID31	WAXS, GISAXS, PDF analysis,	learn more
Laserlab- EuropeCZIP-ASCRHiLASELab XRDXRD, (RIR) phase analysis, Rietveld analysis, analysis of crys- tallinity, crystallite size and lattice 	LEAPS	СН	PSI	SLS	Debye	XAS, XRD, SAXS, (PDF), QEXAFS	
EuropeRietveld analysis, analysis of crystallinity, crystallite size and lattice stress, lattice parameter refine- ment, X-ray reflectivity and SAXSmore learn moree-DREAMITCNRCNRXRD Lab2D, 1D XRD, temperature stagelearn morenon-ARIECZNanoEnvi CzNanoEnvi CzMultipur- powder dif- fractometer PANalytical XPertPRO MPD (UACH14)X-ray powder diffraction, X-ray powder diffraction, X-ray powder diffraction, X-ray powder diffraction, X-ray powder diffraction, quanti- tative phase analysislearn morenon-ARIECZNanoEnvi CzNanoEnvi CzMultipur- pose X-ray powder diffraction, X-ray powder diffraction, quanti- tative phase analysis, qualitative phase analysislearn morenon-ARIECZNanoEnvi CzCzX-ray powder diffraction, X-ray powder diffraction, quanti- tative phase analysis, qualitative phase analysislearn more morenon-ARIECZNanoEnvi CzX-ray Pow- czX-ray powder diffraction, phase analysis, structure, crystallinity, particle size, solid state transfor- mationslearn morenon-ARIECZNanoEnvi CzX-ray Pow- czX-ray powder diffraction, phase analysis, structure, crystallinity, particle size, solid state transfor- mationslearn more	LEAPS	FR	SOLEIL	SOLEIL	PSICHE	EDXRF, ADXRD, Tomography	
e-DREAMITCNRCNRXRD Lab2D, 1D XRD, temperature stagelearr morenon-ARIECZNanoEnvi CzNanoEnvi CzMultipur- powder dif- fractometer PANalytical XPertPRO MPD (UACH14)X-ray powder diffraction, X-ray powder micro-diffraction, quanti- tative phase analysis, qualitative phase analysislearr morenon-ARIECZNanoEnvi CzNanoEnvi CzMultipur- powder dif- fractometer PANalytical XPertPRO moved diffraction, X-ray powder diffraction, X-ray powder diffraction, X-ray powder diffraction, X-ray powder diffraction, X-ray powder diffraction, Quanti- tative phase analysis, qualitative powder diffraction, quanti- tative phase analysis, qualitative powder diffraction, quanti- tative phase analysis, qualitative powder diffraction, quanti- tative phase analysis, qualitative phase analysis, qualitative phase analysis, qualitative phase analysislearr more morenon-ARIECZNanoEnvi CzNanoEnvi CzX-ray pow- CzX-ray powder diffraction, phase analysis, structure, crystallinity, tion (UPOL7)X-ray powder diffraction, phase analysis, structure, crystallinity, particle size, solid state transfor- mationslearr more		CZ	IP-ASCR	Hilase	Lab XRD	Rietveld analysis, analysis of crys- tallinity, crystallite size and lattice stress, lattice parameter refine-	<u>learn</u> more
non-ARIECZNanoEnvi CzNanoEnvi CzMultipur- pose X-ray powder dif- fractometer PANalytical XPertPRO MPD (UACH14)X-ray powder diffraction, X-ray 	e-DREAM	IT	CNR	CNR	XRD Lab		learn more
CzCzCzpose X-ray powder dif- fractometer (Co tube), 	non-ARIE	CZ			pose X-ray powder dif- fractometer PANalytical XPertPRO MPD	powder micro-diffraction, quanti- tative phase analysis, qualitative	learn more
non-ARIE       CZ       NanoEnvi Cz       NanoEnvi Cz       X-ray Pow- der Diffrac- tion (UPOL7)       X-ray powder diffraction, phase analysis, structure, crystallinity, particle size, solid state transfor- mations       learr more         non-ARIE       CZ       NanoEnvi       X-ray Pow- tion (UPOL7)       X-ray powder diffraction, phase particle size, solid state transfor- mations       learr         non-ARIE       CZ       NanoEnvi       NanoEnvi       X-ray difrac-       Phase analysis of materials, struc-       learr	non-ARIE	CZ			pose X-ray powder dif- fractometer (Co tube), Empyrean, series 3	powder micro-diffraction, quanti- tative phase analysis, qualitative	
non-ARIE CZ NanoEnvi NanoEnvi X-ray difrac- Phase analysis of materials, struc- learn	non-ARIE	CZ			X-ray Pow- der Diffrac-	analysis, structure, crystallinity, particle size, solid state transfor-	
	non-ARIE	CZ			-	Phase analysis of materials, struc-	

Panalytical X lattice parameters, lattice (UJEP5) crystallinity.	strain,	



## IR- to VUV-beamlines

The instruments sorted into this category cover an energy range which also includes that of typical lasers, but their photon source is based at a synchrotron, free-electron laser, or large-scale laser system.

Beamline based IR- to VUV-light sources can allow for high stable beams, with a wide spectral range and potentially very short pulses.

### Techniques

- Fourier-transform infrared (FTIR) spectroscopy for measuring infrared absorption and emission spectra, to determine chemical composition and functional groups
- Synchrotron radiation circular dichroism (SRCD) Very fast acquisition of data about folding and stability of biomacromolecules
- Pump-probe measurements for measuring fast dynamics of systems

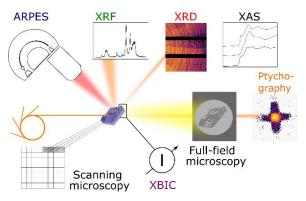
Network	Country	Access provider	Infrastru cture	Instrume nt	Technique	Link
LEAPS	ES	ALBA- CELLS	ALBA	MIRAS Beamline	Fourier Transform Infrared (FTIR) spectroscopy and microscopy	learn more
LEAPS	DE	DESY	FLASH	FLASH	Pump-probe, X-ray spectroscopy, XUV RAMAN, TR-RIXS	<u>learn more</u>
LEAPS	IT	ELETTRA	ELETTRA	SISSI	IR spectroscopy, microspectros- copy and imaging	<u>learn more</u>
Laserlab- Europe	ES	LLE- AISBL	CLPU	VEGA	Pump-probe measurements	learn more
LEAPS	FR	SOLEIL	SOLEIL	DISCO	VUV: SRCD, microspectrofluorim- eter, full-field inverted micro- scope, DUV imaging	learn more



# X-ray imaging

X-ray imaging refers to the set of techniques that provide two-dimensional visualizations of a sample. Images are obtained by recording one or more effects of the interaction between matter and X-rays. Emission of electrons or Xray photons, absorption or diffraction of the impinging beam, creation of free charge carriers are some of these effects and can be used as a contrast mechanism for image formation.

Imaging experiments can be performed in a **full-field** setup, where the sample is illuminated by the whole X-ray beam and the transmitted beam is recorded by a pixelated detector, or in



Schematic representation of multiple possible X-ray imaging modalities. Image credit to Giovanni Fevola and Christina Ossig.

a scanning setup, where the sample is raster-scanned through the highly focused X-ray beam.

Regardless of the setup, synchrotron images can be recorded much faster than with a laboratory source and enable live visualization of physicochemical processes. Furthermore, synchrotrons allow variation of the X-ray energy with few eV resolution. That allows, by scanning the energy over the absorption edge of a specific element, to gain chemical information about an element in a system.

Depending on their energy, X-rays can probe different elements, although with varying penetration depth. Soft X-rays, with energy below 2.5 keV, can probe low-Z elements that are of particular interest for life sciences, but have a low penetration depth. Hard X-rays, with energy above 10 keV, can be used to probe heavy metals and thicker samples.

Resolution is generally linked to the size of the focus spot or to the pixel size of detectors, although methods such as holography and ptychography exploit coherence of X-ray beam to achieve higher resolution (typically 100 to 10 nm).

### Techniques

- X-ray fluorescence measures chemical composition of a material
- X-ray diffraction evaluates the crystal structure of a material
- X-ray beam induced current measures charge collection efficiency in a semiconductor
- X-ray transmission measures transmittance (absorptance) of a material
- X-ray absorption spectroscopy probes chemical state of an element
- Scanning photoemission spectroscopy probes the electronic states of the valence bands

Network	Coun- try	Access provider	Infra- structure	Instrume nt	Technique	Link
LEAPS	ES	ALBA-	ALBA	MISTRAL	FFTXM, cryo-nano tomography, spec-	learn
		CELLS			troscopy imaging, magnetic imaging	more
LEAPS	DE	DESY	PETRA III	P05 <sup>8</sup>	Holography, Tomography	learn
		(GEMS-				more
		Hereon)				
LEAPS	DE	DESY	PETRA III	P06	XRF, XAS, XRD, Ptychography, Tomog-	learn
					raphy	<u>more</u>
LEAPS	DE	DESY	PETRA III	P07 <sup>9</sup>	XRD, 3D-XRD, mirco-tomography	learn
		(GEMS-				more
		Hereon)				
LEAPS	IT	ELETTRA	ELETTRA	ESCA	SPEM	learn
						<u>more</u>
LEAPS	IT	ELETTRA	ELETTRA	NANO-	XPEEm, LEEM, SPLEEM	<u>learn</u>
				SPEC-		more
				TROS-		
				COPY		
LEAPS	IT	ELETTRA	ELETTRA	TwinMic	soft XTM, XEM	learn
						more
LEAPS	FR	ESRF	ESRF	ID13	μXRF, μCrystallography, Ptychography,	learn
					SAXS, XRD	<u>more</u>
LEAPS	FR	ESRF	ESRF	ID19	μTomography	learn
						<u>more</u>
LEAPS	FR	SOLEIL	SOLEIL	ANTARES	HRPES, XAS, ResPES, PhD, XPD, ARPES	<u>learn</u>
						more
LEAPS	FR	SOLEIL	SOLEIL	NANO-	XRF, FF XAS, μTomography	learn
				SCO-		more
				PIUM		
Laserlab-	ES	LLE-	CLPU	VEGA	Pump-probe measurements	learn
Europe		AISBL				more
		(CLPU)				
non-	EU	CERIC-	CERIC-ERIC	DEME-	STXM, XRF, XAS	learn
ARIE		ERIC		TER-		more
				STXM@S		
				OLARIS		

 <sup>&</sup>lt;sup>8</sup> For SME-access only feasibility studies with up to max. 4 hours beamtime are possible.
 <sup>9</sup> For SME-access only feasibility studies with up to max. 4 hours beamtime are possible.

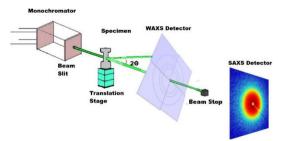


## X-ray small/wide-angle scattering (SAXS-WAXS)

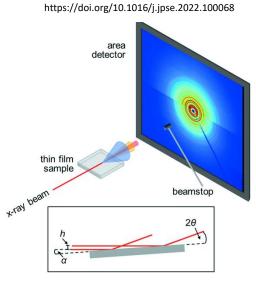
Small- and wide-angle X-ray scattering (SAXS/WAXS) probe the size, shape, orientation, and crystallinity of mesoscale structures on length scales ranging from about one to several hundred nanometers. The techniques can be applied to solids, liquids, and dispersions in both bulk and, with restrictions, thin films.

By focusing the X-ray beam, SAXS/WAXS can be used as an imaging technique in scanning mode. In this case, one can map nanoscale structures within the sample, with the real-space resolution essentially given by the size of the incident X-ray beam. Scanning-based imaging can be carried out in two, three, or six dimensions, depending on the sample.

In the study of surfaces, interfaces and thin films, the structure of a single monolayer up to material thickness of up to tens of nanometers on or within a bulk sample are investigated. To limit the penetration depth of the X-ray beam to the surface-near/interfacenear region grazing incidence (GI) geometry is very effective, where the sample is illuminated under an incident angle smaller or around the critical angle of total external reflection, which amounts to values < 0.1° at high photon energies. To successfully perform GI experiments, the positional alignment of the surface/interface with respect to the X-ray beam requires high precision. To restrict the X-ray footprint resulting from the very shallow incident angle to not exceed the sample dimension, a tightly focused X-ray beam is essential.



Scheme of WAXS and SAXS geometry. From: Connolly et al., J. of Pipeline Science and Engineering, **2** (3), 10068, 1011,



Scheme of a GiSAXS geometry. From: Dippel *et al.* (2019). *IUCrJ*, **6**, 290-298, https://doi.org/10.1107/S2052252519000514

#### Techniques

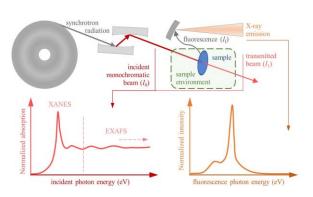
- SAXS/WAXS probing size, shape, orientation, and crystallinity of mesoscale structures
- Scanning SAXS/WAXS imaging for real space spatial resolution
- Grazing-incidence (Gi)SAXS/WAXS for surface layers on flat substrates.

### Infrastructures

Networ k	Country	Access provider	Infrastru cture	Instrument	Technique	Link
LEAPS	ES	ALBA- CELLS	ALBA	NCD-SWEET Beamline	SAXS/WAXS, GISAXS/GIWAXS	learn more
LEAPS	DE	DESY	PETRA III	P03	GISAXS, GIWAXS	learn more
LEAPS	DE	DESY	PETRA III	P07	SAXS, WAXS, GISAXS, GIWAXS	learn more
LEAPS	FR	ESRF	ESRF	ID01	GISAXS	learn more
LEAPS	FR	ESRF	ESRF	ID13	SAXS	learn more
LEAPS	FR	ESRF	ESRF	ID15A	SAXS, TR-WAXS	learn more
LEAPS	FR	ESRF	ESRF	ID31	SAXS, GISAXS, WAXS	learn more
LEAPS	FR	SOLEIL	SOLEIL	SWING	SAXS, WAXS	learn more
LEAPS	SE	ULUND	MAX IV	ForMAX beamline	SAXS, WAXS	learn more
non- ARIE	CZ	NanoEn- viCz	NanoEn- viCz	X-ray powder diffractometer with optics for nanolayers and nanosur- faces Panalyti- cal X Pert PRO (UJEP33)	X-ray diffractometer equipped with optics for structure anal- ysis of polycrystalline thin films and nanosurfaces.	<u>learn more</u>

## X-ray spectroscopy

X-ray spectroscopy (XAS) is sensitive to the local atomic and electronic structure around the element of interest. This element selective technique has versatile applications for solid, liquid and even gaseous materials, including timeresolved in situ studies. X-ray absorption nearedge structure spectroscopy (XANES), which covers the region of 50-100 eV above the absorption edge, probes the transition from the core-level to unoccupied electronic states, being sensitive to the oxidation state, ligand surrounding and local symmetry of the absorbing atom. Extended X-ray absorption fine structure (EXAFS) provide the coordination numbers and interatomic distances for the absorbing atom.



Scheme of X-ray spectroscopy setup and measured data. Image credit to Aram Bugaev.

If the concentration of element of interest is high enough, XAS can measured in transmission geometry. Such measurement can be performed under in situ/operando conditions, high pressures and temperatures and with sub-second time resolution. In fluorescence mode samples with low concentrations of the element of interest down to few ppm can be measured. High time resolution (down to 100 ps) can be also achieved in pump-probe regime. This regime can be applied to liquid samples such as metalloproteins, colloidal nanoparticles and homogenous catalysts, solid samples with low concentration of element of interest or/and the presence of other highly absorbing elements (e.g. led oxide doped with noble metal), and samples that cannot be manipulated to optimize their thickness for transmission geometry (e.g. artefacts of cultural heritage). The energy profile of the fluorescence signal can be also scanned resulting in X-ray emission spectrum (XES) or resonant XES (RXES), if the incident photon energy is tuned to the absorption edge of the element under consideration. The spectra are sensitive to the electronic configuration of the absorbing atom, thus providing important information about chemical bonding. In case both incident and florescence photon energies are scanned, the high-energy-resolution fluorescence-detected (HERFD)-XANES spectra or resonant inelastic X-ray scattering (RIXS) maps are obtained. Finally, in X-ray photoemission spectroscopy (XPS), the kinetic energy of the excited photoelectrons is measured providing information on the binding energies of electrons in materials. The XPS spectra are therefore sensitive to the atomic composition of the sample and the chemical state of each type of atom. Since the photoelectron mean free path is not big, XPS is surface sensitive (1-10 nm).

#### Techniques

- **X-ray absorption spectroscopy** element selective local atomic and electronic structure: oxidation state, ligand surrounding, coordination numbers and interatomic distances.
- Pump-Probe XAS time resolved (down to 100 ps) electronic changes of active site
- X-ray emission spectroscopy electronic configuration of the element of interest
- X-ray photoelectron spectroscopy chemical composition and chemical state of surface atoms

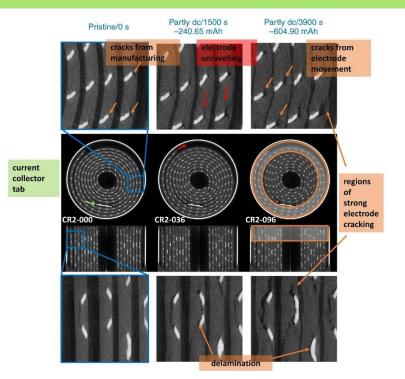
Network	Coun- try	Access pro- vider	Infrastru cture	Instrument	Technique	Link
LEAPS	ES	ALBA-	ALBA	CLAESS Beam-	XAS, XES, XRF, in-situ	learn
		CELLS		line		more
LEAPS	ES	ALBA-	ALBA	CIRCE Beam-	PEEM, NAPP, XPS-PEEM, XMC(L)D-	learn
		CELLS		line	PEEM, IV-LEEM and u-LEED, NEXAFS or XAS	more
LEAPS	DE	DESY	PETRA III	P22	HAXPES, HAXPEEM, high pressure XPS, k-microscopy	<u>learn</u> more
LEAPS	DE	DESY	PETRA III	P64/P65	EXAFS, QEXAFS, RXES, XAFS	learn
				·		more
						learn
						more
LEAPS	DE	DESY	FLASH	FLASH	Pump-probe, X-ray spectroscopy, XUV	learn
					RAMAN, TR-RIXS	more
LEAPS	IT	ELET-	ELETTRA	ESCA	SPEM	learn
		TRA				more
LEAPS	IT	ELET-	ELETTRA	ALOISA	XPS, NEXAFS, PED	learn
		TRA				more
LEAPS	DE	HZB	BESSY II	KMC-2	XANES	learn
						more
LEAPS	СН	PSI	SLS	Debye	XAS, XRD, SAXS, (PDF), QEXAFS	learn
						more
LEAPS	СН	PSI	SLS	SuperXAS	XAS, XES, QEXAFS, pump-probe, TR-	learr
				-	XAFS	more
LEAPS	FR	SOLEIL	SOLEIL	LUCIA	μXRF, μXAS, XANES, EXAFS, Raman	learr
					spectroscopy	more
LEAPS	FR	SOLEIL	SOLEIL	SAMBA	XAS	learn
						more
LEAPS	FR	SOLEIL	SOLEIL	HERMES	STXM, X-PEEM, XMCD, XMLD, XAS,	learn
					XANES, XPS, ARPES	more
LEAPS	FR	SOLEIL	SOLEIL	ANTARES	HRPES, XAS, ResPES, PhD, XPD, ARPES	learn
						more
LEAPS	SE	ULUND	MAX IV	Balder beam-	EXAFS, XANES, XAS, XES, XRF	learn
				line		more
non-ARIE	EU	CERIC-	CERIC-	Envi-	XPS	learn
		ERIC	ERIC	roESCA@CUP		more
non-ARIE	EU	CERIC-	CERIC-	DFEMETER-	XPS, XAS (soft X-rays)	learn
		ERIC	ERIC	PEEM@SO-		more
				LARIS		
non-ARIE	EU	CERIC-	CERIC-	PHELIX@SO-	XAS (soft X-rays), LEED-AES, CDS	learn
		ERIC	ERIC	LARIS		more
non-ARIE	EU	CERIC-	CERIC-	ASTRA@SO-	XAFS	learn
		ERIC	ERIC	LARIS		more
non-ARIE	EU	CERIC-	CERIC-	URANOS@SO-	ARPES, LEED	learn
		ERIC	ERIC	LARIS		more
non-ARIE	EU	CERIC-	CERIC-	PIRX@SO-	XAS (soft X-rays) XMCD	learr
		ERIC	ERIC	LARIS	· · ·	more

non-ARIE	CZ	NanoE nviCz	NanoEn- viCz	X-Ray Photoe- lectron Spec- troscopy (UPOL3)	XPS, Surface chemical composition, Valence state, Chemical quantification	<u>learn</u> more
non-ARIE	CZ	NanoE nviCz	NanoEn- viCz	WDRF spec- trometer Rigaku Primus IV (UJEP 37)	Elemental analysis, WDXRF spectrome- try, Thin layer analysis, Analysis of de- fects	<u>learn</u> more
Laserlab-	ES	LLE-	CLPU	VEGA		learn
Europe		AISBL				more
Laserlab-	ES	LLE-	ICFO	Attoseconds		<u>learn</u>
Europe		AISBL		SXR beamline		more

## X-ray tomography

X-ray tomography is a non-destructive full-field imaging technique applicable to materials from The metals to organic tissue. basic experimental setup is relatively simple: the sample is placed on a rotating stage and aligned in the X-ray beam, with a detector placed some distance after the sample. A tomographic dataset is collected by rotating the sample in the X-ray beam and collecting a series of projections from different angles. The series of projections are fed collected into а reconstruction algorithm to produce a threedimensional volume representation of electron density in the sample.

For many material science samples, preparation is minimal: the sample can simply be mounted securely on the sample stage, making tomography a simple experiment to carry out. Diverse experimental setups allow measurements optimised for different samples approaches, with microtomography or beamlines offering voxel sizes in the range of 0.35-50 µm. Hard X-rays (typically from 20 to over 250 keV) are used to enable penetration



X-ray tomograms of a commercial  $Li/MnO_2$  primary battery. The pristine state and two partly discharged states are presented. The images show the cracking and volume expansion of the  $MnO_2$  electrode during cell discharging.

Source: Nat Commun 11, 777 (2020). https://doi.org/10.1038/s41467-019-13943-3

of large samples. Tomography can be readily combined with sample environments such as a cryostream or a furnace to carry out *in situ* experiments.

The data collected from X-ray tomography provides a 3D view of the internal structure of a sample. Important features such as inclusions or cracks can be observed non-destructively. *In situ* experiments can provide feedback on the real-time evolution of materials under processing or, for example, undergoing tests in a load frame.

### **Techniques**

- Full-field tomography absorption contrast
- Scanning tomography absorption, element specific, or phase contrast
- Ptycho-tomography phase contrast, high resolution

Networ k	Coun- try	Access provider	Infrastructure	Instrume nt	Technique	Link
LEAPS	DE	DESY (GEMS- Hereon)	PETRA III	P05 <sup>10</sup>	Holography, Tomography	<u>learn</u> more
LEAPS	DE	DESY	PETRA III	P06	XRF, XAS, XRD, Ptychography, To- mography	<u>learn</u> more
LEAPS	DE	DESY (GEMS- Hereon)	PETRA III	P07 <sup>11</sup>	XRD, 3D-XRD, mirco-tomography	<u>learn</u> more
LEAPS	FR	ESRF	ESRF	BM05	X-ray diffractometry, reflectome- try, topography, tomography	<u>learn</u> more
LEAPS	FR	ESRF	ESRF	ID19	μTomography	<u>learn</u> more
LEAPS	FR	SOLEIL	SOLEIL	ANATO- MIX	FF radiography, tomography, in ab- sorption and phase contrast	learn more
LEAPS	FR	SOLEIL	SOLEIL	PSICHÉ	EDXRF, ADXRD, tomography	<u>learn</u> more

<sup>&</sup>lt;sup>10</sup> For SME-access only feasibility studies with up to max. 4 hours beamtime are possible. <sup>11</sup> For SME-access only feasibility studies with up to max. 4 hours beamtime are possible.



# All NanoEnviCz Instruments

NanoEnviCz integrates the infrastructure facilities of Czech research organizations for research in nanotechnology sciences.

The program is focused on research in the field of nanomaterials and nanocomposites for environmental and related applications. Our services include controllable syntheses of nanomaterials, their complex chemical, structural, morphological and surface characterization, tuning their functional properties, monitoring their potential toxicity and hazard.

Analysers		
Instrument	Description	Link
Electrokinetic analyser SurPASS (UJEP6)	Measurement of zeta potential of solid samples. Solid surfaces can be measured in the form of flat samples (polymer foils, glass, etc.) and powders or fibres (all size has to be higher than 25 micrometers).	<u>learn</u> <u>more</u>
Laboratory of computational chemistry (UJEP 34)	This laboratory is composed of two computer clusters for high performance computing. The newer cluster consists of 18 nodes "DELL PowerEdge T640 Server", providing a great background, especially for more demanding parallel computations. Each node is equipped with 2 Intel Xeon Gold 6240 processors and 192 GB (16 nodes) or 384 GB (2 nodes) RAM. In addition, 10 nodes are equipped with four GeForce RTX 2080 Ti graphics cards for GPU or GPU/CPU calculations. The older cluster consists of 9 older nodes "Dell PowerEdge R720" (2x Intel Xeon E5-2695 v3, 128 GB RAM) and two newer nodes "TYAN - GPU Server FT48TB7105" (2 x Intel Xeon Gold 6240, 192 GB RAM) which are moreover equipped also with 3 x GeForce RTX 2080 Ti for GPU accelerated calculations.	learn more
Microarray laser scanner - NEW (UJEP27)	Development of novel sensitive optical microarray diagnostic devices (biosensors) in the area of biosensing for environmental or biomedical applications.	<u>learn</u> <u>more</u>

### All of the NanoEnviCz instruments are also available in the industry access routes.

## Centrifuges

Instrument	Description	Link
Refrigerated Centrifuge (UFCH5)	Centrifuge is power-driven machine that separates liquids from solid matter, liquids mixtures, or solid mixtures by centrifugal force. The max. rotation speed is 15000 rpm and max. volume capacity 3200 ml. Refrigerated centrifuge is equipped by temperature controller controller in range of -20 to +40°C.	<u>learn</u> more
Ultracentrifuge (UFCH16)	The ultracentrifuge is a centrifuge optimized for spinning a rotor at very high speeds, capable of generating acceleration as high as 1 000 000 g (approx. 9 800 km/s <sup>2</sup> ). It can also be used for gradient separations, in which the tubes are filled from top to bottom with an increasing concentration of a dense substance in solution.	<u>learn</u> <u>more</u>

## **Electrochemical techniques**

Instrument	Description	Link
FRA -	Integrated electrical and optical measurement system. Set-up for	<u>learn</u>
PhotoEchem	electrochemical measurements including Frequency Response	more
System (UFCH 20)	Analyzer interfaced to electrochemical impedance spectroscopy, Potentiostat/Galvanostat, Solar simulator and IPCE module	

### Chromotographic techniques

Instrument	Description	Link
GC-qMS Agilent	Gas chromatograph Agilent 7980 with electron ionization and simple	<u>learn</u>
(UJEP16)	quadrupole detection (MS 5977E) equipped with autosampler	more
	CombiPAL for liquid, headspace and SPME sample introduction.	
	Agilent MassHunter Workstation Software is used for data	
	acquisition and analysis.	
GC/MS/MS	Gas Chromatograph Thermo Trace 1310 with two injection ports	<u>learn</u>
(TUL11)	(SSL and PTV). Autosampler CTC CombiPal RTC), Mass spectrometer	more
	TSQ 8000 Evo – triple quadrupole with unit mass resolution.	
HPLC/MS/MS	The instrument is an HPLC chromatograph with a triple	learn
(TUL1)	quadrupole/linear ion trap mass spectrometer. The HPLC is a binary	more
	system with two pumps enabling very fast mobile phase gradients.	
lon	Ion Chromatograph DIONEX ICS – 1000	learn
Chromaatograph		more
DIONEX (UJEP14)		
Liquid	Liquid chromatograph with DAD datactor. Marck/Hitachi	loorn
Liquid Chromatograph	Liquid chromatograph with DAD detector, Merck/Hitachi.	<u>learn</u>
(UJEP15)		more
Liquid	Agilent 1290 Infinity UHPLC system consisting of an Agilent 1290	<u>learn</u>
chromatograph	Infinity Binary Pump (G4220A), an Agilent 1290 Infinity High	more
with MS	Performance Autosampler (G4226A), a sample cooler (G1330B), and	
detection	an Agilent 1290 Infinity Thermostatted Column compartment	
(UJEP18)	(G1316C). The UHPLC system is coupled to an Agilent G6495 Triple	
	Quadrupole LC/MS System equipped with an Agilent Jet Stream	
	electrospray ionization source. Agilent MassHunter Workstation	
	Software is used for data acquisition and analysis.	
Liquid	Liquid chromatograph with DAD detector – DIONEX Ultimate 3000	learn
chromatograph	Pump – LPG-3400SD Quarternary Standard Pump	more
with diode-array		
detector Dionex		
(UJEP9)		
Two-Dimensional	Gas Chromatograph (Agilent 7890) equipped with multimode inlet	learn
Gas	(split/splitless/LVI/PTV), GCxGC modulator ZOEX and coupled with	more
Chromatograph	flame ionization detector and mass spectrometry detector (Agilent	
(UJEP36)	7250) quadrupole – time-of-flight (q-ToF). Deans Switch placed blind	
	GCxGC modulator allows two-dimensional chromatography on FID	
	or q-ToF.	

## **Microbiologic techniques**

Instrument	Description	Link
Laboratories for mammalian cell cultivation	Complete infrastructure for mammalian cell cultivation and related experiments equipped by Biohazard box class 2, inverted fluorescence microscope and flow cytometer Attune	<u>learn</u> more
(UJEP21) Laboratory of Nanotoxicology and Model	Laboratory of model organism - Danio Rerio - for toxicity testing	<u>learn</u> more
Organisms (UJEP29) Laboratory of	Complete infrastructure for the design, manufacturing and testing of	learn
biosensors and microfluidics (UJEP22)	biosensors and microfluidic devices for biomedical and environmental applications. Scanning electron microscope with electron lithography module , UV photolithographic instrument , magnetron sputtering device, microabrasive CNC lathe, reactive ion etching station , microfluidic liquid sample delivering system	<u>more</u>
Real-time PCR device (TUL6)	The system features the LightCycler® 480 Instrument, a versatile, plate-based real-time PCR device that supports mono- or multicolor applications, as well as multiplex protocols.	<u>learn</u> <u>more</u>
Respirometer - NEW (TUL12)	Continuous monitoring of metabolic gases concentrations.	<u>learn</u> more
The LightCycler® 480 Real-Time PCR System (IEM11)	The LightCycler <sup>®</sup> 480 System is a plate-based, highly adaptable, and versatile real-time PCR system for gene expression analysis, SNP genotyping, and mutation scanning via high resolution melting (HRM). Key benefits of the LightCycler <sup>®</sup> 480 Thermal Block Cycler: - Run any assay format or application with fast PCR protocols (< 40 minutes for 40 cycles in 384-well plate format) Obtain rapid and accurate temperature adjustment Achieve exceptional data homogeneity across the entire multiwell plate.	<u>learn</u> more

## Micromolecular techniques

Instrument	Description	Link
Metafer Slide	Metafer is an automated multi-purpose slide scanning platform.	<u>learn</u>
Scanning System	Equipped with CometScan software for MSearch, it enables	more
(IEM2)	automatic detection of single cell gel electrophoresis (Comet assay) samples.	
Metafer Slide	The automated scanning system Metafer 4, Version 3.2.1, is a set of	<u>learn</u>
Scanning System	motorized Axio Imager Z1 microscope and software for scoring of	more
(IEM3)	binucleated cells and metaphases.	
MiSeq System	The Illumina MiSeq is a desktop sequencer with integrated computer	<u>learn</u>
(IEM6)	which enables a broad range of applications, from targeted gene	more
	sequencing to metagenomics, small genome sequencing, targeted	
	gene expression analysis, amplicon sequencing starting at 10 ng	
	DNA, and HLA typing. New MiSeq reagents enable up to 15 Gb of	
	output with 25 M sequencing reads and 2x300 bp read lengths.	
SpectraMax	A five-mode microplate reader with three-mode cuvette port for	learn
Multimode Plate	endpoint, kinetic, spectrum, and area-well scanning with PathCheck	more
Reader (IEM1)	sensor and SoftMax Pro Software.	
iScan System	The iScan System is a laser-based, high-resolution optical imaging	<u>learn</u>
(IEM5)	system that can rapidly scan and collect large volumes of data from	more
	Illumina DNA analysis and RNA analysis high-density BeadChips.	

### Microscopic techniques

Instrument	Description	Link
AFM (UACH1)	Atomic Force microscope provides imaging sample topography at high resolution, measuring magnetic structure of the sample surface by MFM and measuring electrical properties by STM.	<u>learn</u> more
DXR Raman mikroscope (UACH9)	Thermo Scientific DXR Raman Microscope for phase identification and determination of the molecular structure of the chemical compounds. Analysis of the organic and inorganic compounds, carbon materials, nanomaterials, etc.	<u>learn</u> more
Fluorescence Microscope (IEM4)	A set of fluorescent microscope and computer equipped with ISIS color fluorescence and FISH imaging system for analysis of chromosomal aberrations and fluorescently stained biological materials.	<u>learn</u> more
Fluorescence inverted confocal spinning disk microscope Olympus SpinSR10 (UEM12)	The Olympus SpinSR10 is a fluorescence inverted confocal spinning disk microscope with super-resolution mode. It is designed for fast 3D super resolution imaging and prolonged cell viability in time- lapse experiments, the IXplore SpinSR microscope system offers XY resolution down to 120 nm without the need for dedicated labeling procedures.	<u>learn</u> more
HRSEM FEI NanoSEM 450 (UACH4)	FEI Nova NanoSEM <sup>™</sup> scanning electron microscopes combine best- in-class imaging with superb analytical performance in one easy-to- use instrument. It is a high-resolution scanning electron microscope, with two modes of measuring and five different detectors.	<u>learn</u> more
High Resolution Transmission Electron Microscope (UPOL5)	High Resolution Transmission Electron Microscope (HRTEM) FEI Titan 60-300 kV an electron source of X-FEG, accelerating voltage from 60–300 kV and a point to point in TEM mode resolution of 0.08 nm. The microscope is equipped with GIF (Gatan Image Filter) and analytic methods EDS and EELS and special holders for reactive samples (vacuum holder, cryo holder, double-tilt holder). The characterization of the nanomaterial's samples (carbon structures, iron oxides, nanotubes, metal nanoparticles, ect.) in the atomic scale is provided.	<u>learn</u> <u>more</u>
High resolution transmission electron microscope (UFCH21)	HRTEM will enable the viewing and imaging of details in nanostructure of nanomaterials down to the dimensions of nanometres with the resolution down to about 0.2 nm.	<u>learn</u> <u>more</u>
High resolution transmission electron	Equipped with EDX detector (Oxford Instruments) and precession diffraction DigiStar (NanoMegas). It is used for high-quality materials	<u>learn</u> <u>more</u>

microscope (JEOL) JEM 3010 (UACH10)	characterization - morphology, phase analysis on nanometer scale, maps of various crystallographic phases and the crystal orientation.	
High resolution transmission electron microscope, HRTEM FEI Talos F200X (UACH16)	High resolution measurement of powder materials in the atomic scale with confirmation of the elemental composition and crystal structure for particle size up to 100nm. The identification of nanoparticles – quality of production, size and shape, determination of d-spacing, projection of atomic structure is studied including chemical composition confirmation (elemental mapping, EDS spectra).	<u>learn</u> more
Infrared imaging microscope with FTIR spectrometer (TUL3)	FTIR spectrometer Nicolet iZ10 - DTGS (room temperature) detector, suitable spectral range 4000 – 400 cm-1, standard resolution 4 cm-1 or more. Infrared imaging microscope Nicolet iN10 MX – DTGS (room temperature) and MCTA (nitrogen cooled) detector, suitable spectral range 4000 – 400 cm-1, standard resolution 4 cm-1 or more	<u>learn</u> more
Laser scanning confocal microscop (UPOL15)	The laser scanning confocal microscopy instrument with Airyscan 2 includes options for fast imaging with improved resolution. Suitable applications include live cell imaging/time courses, colocalization studies, Photo-activation, FRAP, FRET, spectral imaging, stitching of large areas, and imaging of fixed samples. CLSMs, enabling fast multiplexed super-resolution imaging (2x increase in spatial resolution) at 4x faster speed, especially suitable of dynamic live-cell imaging, containing linear scanner and enabling fast imaging 13images/second with resolution (512 x 512 pixels). It contains planapochromat objectives: 10x/0.45 M27 [working distance (WD) 2.1mm], 20x/0.8 M27 (WD 0.55mm) with DIC, 40x/1.2 Imm DIC M27 (WD 0.41mm) immersion: water, silicone oil or glycerol, 63x/1.4 Oil DIC M27 (WD 0.19mm), incubator XL multi S2 Dark premium with incubation set CO2/O2 and temperature heating desk, antivibration table, ZEN 3.3 system with ZEN module FRAP, spectral 32 channel GaAsP PMT and 2 channels MA-PMT detectors. The confocal microscope covers the whole spectra with lasers (405, 445, 488, 514, 543,594, and 639nm).	<u>learn</u> more
Low temperature UHV (UPOL8)	Set of Ultra High Vacuum chambers with Cryostat and Scanning Tunneling Microscope/Atomic Force Microscope for surface analysis.	<u>learn</u> <u>more</u>
Raman microscopy (TUL4)	Raman microscopy with laser 532 nm, Nicolet DXR Raman microscopy.	<u>learn</u> more
Scanning Electron Microscope (SEM) Hitachi SU6600 (UPOL10)	Scanning Electron Microscope (SEM) Hitachi SU6600	learn more

Scanning Probe	Scanning Probe Microscope (SPM) NTEGRA NT-MDT Measuring in	learn
Microscope	different modes: o Atomic force microscopy (AFM) o Magnetic force	<u>more</u>
(UPOL6)	microscopy (MFM) o Scanning tunneling microscopy (STM)	
Transmission	Transmission Electron Microscope	learn
Electron		more
Microscope		more
(TEM) JEOL 2100		
(UPOL11)		
(OPOLII)		
Confocal	Fully motorized confocal laser scanning microscope SP8 from Leica	learn
microscope -	enclosed in an environmental chamber allowing temperature,	more
LEICA CLSM	humidity and CO2 levels control around the sample. Equipped with a	
SP8/DLS (UJEP40)	digital light sheet (DLS) module. Laser lines: 405 nm, argon laser	
	(458, 488, 514 nm), 561 nm, 633 nm.	
Scanning Electron	Field emission scanning electron microscope FESEM model Hitachi S-	learn
Microscope,	4800	more
Hitachi (UFCH22)		
(0.0.22)		
System AFM-	Scanning probe microscope, NTEGRA Spectra which integrates	<u>learn</u>
Raman (UPOL9)	common SPM and micro Raman scattering spectroscopy. AFM-	<u>more</u>
	Raman system delivers nondestructive analysis of the sample	
	surface.	

## Particle size distribution techniques

Instrument	Description	Link
ZetaSizer NanoS (UFCH1)	Non-invasive back scatter (175 degrees) technology takes particles sizing to new levels of sensitivity in the nanometre to micron range size. ZS provides ability to measure three characteristics of particles or molecules in a liquid medium.	<u>learn</u> <u>more</u>
Zetasizer nano ZS (IEM8)	Non-invasive, well-established technique for measuring the size and size distribution of molecules and particles typically in the submicron region - newly purchased within Pro-NanoEnviCz project	<u>learn</u> more

## Physical properties measuring systems

Instrument	Description	Link
Equipment of the laboratory of nanotoxicology in cell cultures (IEM 9)	The set of instruments forms completely new laboratory of nanotoxicology. It includes a MPT-2 Multipurpose titrator (Malvern), a Bugbox Plus (BAKER RUSKINN), a Laminar flow cabinet (HERASAFE KS,) a CO2 incubator (HERACELL VIOS 250i), aThermo Scientific Barnstead Smart2Pure 3 UV/UF Water Purification System. New laboratory equipment was purchased within the Pro-NanoEnviCz project.	<u>learn</u> <u>more</u>
Fragment Analyzer (IEM 10)	The Fragment Analyzer is a parallel capillary electrophoresis instrument for biological effects of manufactured nanoparticles´studies -a new equipment purchased within Pro- NanoEnviCz project	<u>learn</u> more
Low temperature induction magnetometer - PPMS (UPOL14)	The physical properties measurement system (PPMS) is a complex device that allows to operator a broad option of measurements including magnetic properties, electron-transport properties, and thermal properties. PPMS uses a vibrating sample magnetometer (VSM) for the magnetic moment detection and provides both, DC (direct current) and AC (alternative current), types of measurement in a wide range of temperatures from 1.9 K – 400 K and the presence of an external magnetic field ranging from -9 T to +9 T. The Electrical Transport Option (ETO) enables users to make several different types of transport measurements over a wide range of resistance values and sample types. The ETO supports three types of measurements including resistivity, IV curves, and differential resistance. The current source has a minimum precision of 1 nA and a maximum current of 100 mA. It is capable of supplying both DC and AC current with frequencies from 0.1 Hz to 200 Hz. Last, but not least the heat capacity measurement is also possible at PPMS to complete the full magnetic information.	<u>learn</u> more
Physical Properties Measurement System - PPMS (UPOL2)	The physical properties measurement system (PPMS) allows to operator a broad option of measurements including magnetic properties, electro-transport properties and thermal properties. Regarding magnetic properties, PPMS using a vibrating sample magnetometer (VSM) which is less sensitive than SQUID and provide only DC (direct current) measurement option.	<u>learn</u> more
Tester of Liquid Permeability of nanofibrous membranes (UJEP31)	Tester of Liquid Permeability	<u>learn</u> more

Tester of	Tester of air permeability	learn
Membrane Air		more
Permeability of		
nanofibrous		
membranes		
(UJEP30)		
Tester of	Tester of mechanical strength – tensile tests.	learn
mechanical		more
strength of		
nanofiber		
membranes		
(UJEP32)		

Reactors	ł

Instrument	Description	Link
Autoclave for synthesis, catalysts testing and kinetic measurements (UFCH2)	Set of three autoclaves equipped for liquid phase bath synthesis, catalysts testing and kinetic measurement.	<u>learn</u> more
Autoclave for synthesis, catalysts testing and kinetic measurements (UFCH3)	Set of three autoclaves equipped for liquid phase bath synthesis, catalysts testing and kinetic measurement.	learn more
Catalytic flow microreactor B (UFCH10)	The Microactivity-Reference reactor (PCT/ES2005/070079) is an automatic and computerized laboratory catalytic micro-reactor which includes the valves and process layout in a hot box to avoid the possible condensation of volatile products, at the same time that preheats the reactants efficiently.	<u>learn</u> more
Catalytic flow microreactor A (UFCH9)	The Microactivity-Reference reactor (PCT/ES2005/070079) is an automatic and computerized laboratory catalytic micro-reactor which includes the valves and process layout in a hot box to avoid the possible condensation of volatile products, at the same time that preheats the reactants efficiently.	<u>learn</u> more
Fluidized Bed Reactor (UJEP2)	Equipment for plasma treatment of powder materials	<u>learn</u> more
Laboratory reactors (UJEP12)	Laboratory – scale reactors for preparing metal oxide-based sorbents and related materials by homogeneous hydrolysis, sol-gel process, precipitation/calcination and similar techniques.	<u>learn</u> more
Photocatalytic degradation liquide phase (UACH2)	Set of two photoreactors for the photocatalytic degradation of organic pollutants (dyes, cytostatics, pesticides, etc.) conected with UV-VIS Spectrophotometer ColorQuestXE for signal detection of the organic pollutants and kinetic measurements and FL2000 fluorescence detector	<u>learn</u> more
Universal magnetron deposition system (UJEP1)	The magnetron deposition system with variable system up to 3 magnetron with 2inch targets in diameter. Various power supplies are available RF, DC, RF pulsed, DC pulsed for sputtering of metals or metal oxides.	<u>learn</u> more

### Sample preparation techniques

Instrument	Description	Link
Dip Coater 5 (UFCH18)	Dip Coater 5 is design for uniform deposition of layers on plates and similar objects by dipping and dragging into bath with solution.	<u>learn</u> more
Extruder, Multi- Gran (UFCH6)	Granulator is designed to manufacture granules of ceramics, organic materials, polymers/biopolymers in form of cylinders of the diameter 1 to 3 mm adjusted by the selection of a die with proper openings.	<u>learn</u> more
Industrial femtosecond pulsed laser (TUL13)	Industrial femtosecond laser (Onefive Origamy XP, NKT Photonics) with laser scanner head (intelliSCAN 14, SCANLAB). The tool is used for single or multielemental nanoparticle synthesis. It delivers high energy and frequency pulses capable of material ablation (LAL) and material fragmentation (LFL) in both gases and liquids. Further, it is used for laser melting in liquids (LML) and laser-mediated photoreduction approaches of nanoparticle synthesis.	<u>learn</u> more
Laboratory electric superkanthal furnace (UFCH17)	Furnace for preparation and heat treatment of ceramics, glass phases and metals/metal alloys up to 1700°C under air or an inert atmosphere.	<u>learn</u> more
Laboratory of nanofibrous materials (UJEP28)	Device for electrospinning - needle spinning of polymeric nanofibrous membranes (InoCure).	<u>learn</u> more
MicroWriter ML3 Pro (UFCH25)	MicroWriter ML3 Pro (Durham MagnetoOptics Ltd.) is a direct-write photolithography machine for rapid prototyping in R&D laboratories and small clean rooms. It is compatible with most photolithography resists (385 nm), minimum feature size is 400 nm.	<u>learn</u> more
Microarray printer - NEW (UJEP26)	Fabrication of active biosensor surfaces on different substrates	<u>learn</u> more
Precision Ion Polishing System (PIPS) Model 691(Gatan) (UACH12)	Precision ion polishing system is used for thinning of prepared samples by current of ionized argon to the thickness of few nanometers for transmission electron microscopy.	<u>learn</u> more
Reactor Speedwave four (UFCH12)	Microwave digestion system with built-in, non-contact temperature and pressure measurements. The system has been designed to perform chemical digestion procedures under extreme pressure and temperature conditions in chemical laboratories. Digestion is understood to mean the decomposition of a solid material by means	<u>learn</u> more

	of a suitable digestion reagent at increased temperature in a vessel	
	that is permeable with regard to microwaves .	
T2 Glove Box	Manipulating moisture and/or oxygen sensitive products and testing	<u>learn</u>
(UFCH13)	of materials (including long-term stability tests) in inert gas	more
	atmosphere. The glove box is equipped with cables for performing	
	electrochemical measurements.	
Vacuum	This equipment allows for the deposition of in their size and	<u>learn</u>
Apparatus for the	composition well defined subnanometer clusters on flat surfaces.	<u>more</u>
Deposition of	The principle of operation is analog to the equipment described in	
Size- and	"Atomically Precise (Catalytic) Particles Synthesized by a Novel	
Composition	Cluster Deposition Instrument", C. Yin, E. Tyo, K. Kuchta, B. von	
Selected Clusters	Issendorff, and S. Vajda; J. Chem. Phys. 140, 174201 (2014), DOI:	
(UFCH33)	10.1063/1.4871799	
Clean room	The clean room is equipped with spin coater, mask aligner, oxygen	learn
(UFCH11)	plasma etcher, sputtering machine, thermal evaporator.	more

## Spectroscopic techniques

Instrument	Description	Link
FTIR Spectrometer (UFCH8)	FT-IR Nicolet 6700, Infrared Fourier transform spectrometer (FTIR) for qualitative and quantitative analysis for solid and liquid phase in range Mid-IR (4000-400 cm-1).	<u>learn</u> more
Electron- Paramagnetic- Resonance Spectrometer (UPOL13)	Electron Paramagnetic Resonance (EPR) spectroscopy is similar to any other technique that depends on the absorption of electromagnetic radiation.	<u>learn mo</u>
ICP-OES Optical Emission Spectrometer (UJEP17)	Dual-view optical system (axial/radial) with High-dispersion echelle grating, spectral range 165-900 nm with resolution of < 0.009 nm (200 nm)	<u>learn</u> more
Infrared Spectrometer (UFCH23 -new Pro- NanoEnviCzII)	The Nicolet iS50 is the scientific infrared spectrometer for universal material analysis. The spectrometer is primarily used to identify the structure of materials for heterogeneous catalytic and adsorption processes for environmental protection and for catalytic technologies.	<u>learn</u> more
Laboratory of spectroscopy (UFCH27)	The spectroscopic laboratory consists of Horiba Raman spectrometer, WITec Raman spectrometer, Horiba photoluminescence spectrometer. The LabRAM HR Raman microscope is a suitable system for both micro and macro measurements and offers advanced confocal imaging capabilities in 2D and 3D. The true confocal Raman microscope enables detailed images and analysis. WITec microscope system has an exceptional optical throughput, unparalleled signal sensitivity, and outstanding imaging capabilities. Fluorolog 3 system is a state-of- the-art system for measuring excitation and emission spectra of thin-film and liquid samples	<u>learn</u> <u>more</u>
Mass Spectrometer with inductively coupled plasma ICP-MS (TUL8)	ICP-MS NexIOn300D (Perkin Elmer) with autosampler and possible combination with HPLC (Flexar - Perkin Elmer)	<u>learn</u> <u>more</u>
Microplate spectrophotometer (UJEP19)	Universal 96-well microplate UV-VIS spectrophotometer with PathCheck technology for correction of sample volume variations.	<u>learn</u> <u>more</u>
Sciex X500R QTOF HR mass spectrometer - new	A QTOF type HRMS mass spectrometer coupled to an HPLC chromatograph. Suitable for screening and trace determination of pollutants (e.g. pesticides) and their metabolites, pharmaceuticals	<u>learn</u> <u>more</u>

(TUL14) -Pro- NanoEnviCz II	and biomolecules. Use of libraries and in silico fragmentation software allows for the identification of unknowns.	
Solid State NMR	Classic solid state nuclear magnetic resonance instrument for	learn
Spectrometer Jeol	chemical analysis. The instrument produces a high intensity	more
(TUL15)	magnetic field and studies its interaction with magnetic nuclei of	more
(10113)	the exposed sample. It finds application in basic characterization of	
	chemical structure of synthesized nanomaterials and composites.	
	chemical structure of synthesized hanomaterials and composites.	
Spectrophotometer	UV/Vis spectrophotometer Cary 50 with wavelength 190-1100nm	<u>learn</u>
Cary 50 (UJEP13)		more
Thermo Nicolet -	Mid-infrared Fourier transform spectrometer (FTIR) for	<u>learn</u>
FTIR (UACH8)	determination of the molecular structure of the chemical	more
	compounds and in-situ observation of the adsorption, surface	
	chemical and photochemical reactions.	
Thermoanalytical	System SETARAM for thermal analysis is using sophisticated	<u>learn</u>
Complet (UACH15)	system of QMS module connection so-called SuperSonic System in	<u>more</u>
	which gas molecules are accelerated and directed to the mass	
	spectrometer.	
Thermogravimeter	Thermogravimeter STA449F1 (Netzsch) allows is devoted to	learn
STA449F1(Netzsch)	measure: thermogravimetry (TG) and differential scanning	more
connected with	calorimetry (DSC). TG determine sample mass loss during the	
Mass Spectrometer	thermal treatment and DSC determines the heat capacity of the	
(Anamet) (UFCH14)	sample.	
Thermogravimetric	Thermogravimetric analyzer Q500 is suitable for studying material	learn
analyser with FTIR	thermal stability from ambient to 1000 °C. T Evolved gases can be	more
spectrometer	online studied by FTIR spectrometer Nicolet iS10 with MCTA	
(TUL2)	(nitrogen cooled) detector in spectral range 4000 – 650 cm-1 and	
	maximum spectral resolution 1 cm-1.	
X-Ray	Acquired Information are determination and quantification of	learn
Photoelectron	chemical composition of surfaces (max. depth 10 nm),	more
Spectroscopy	determination of valence states of atoms, chemical composition	
(UPOL3)	depending on the depth (depth chemical concentration).	
XPS/ESCA and	The instrument is an electron spectrometer SPECS with an X-Ray	learn
Auger electron	source of achromatic (Al/Mg) and monochromatized (Al/Ag)	more
spectroscopy	photons for electron spectroscopy (XPS/ESCA) analyses with	
(UJEP3)	electron source-based charge compensation. The system also is	
· -	equipped with an electron source (50 eV – 3000 eV) for Auger	
	electron spectroscopy (AES) with scanning options and an SE	
	detector (SEM/SAM). The detection unit is 5 channel channeltron.	
	The base pressure is about 4x10-9 mbar. The solid samples and	
	powder samples can be analyzed. The limitation is mainly in the	
	sample stability under the measurement conditions. A depth	

profiling of elemental composition is possible by Argon ions sputtering from an external ion source.	

### Surface characterization techniques

Instrument	Description	Link
Apparatus for the determination of the texture features and adsorption properties of solid materials - Pro- NanoEnviCz (UFCH19)	A device for determining the surface area, pore size distribution and pore volume by the physical gas sorption	<u>learn</u> <u>more</u>
BET (TUL10)	Surface area and pore size analyzer. The analysis is based on physisorption of either nitrogen or argon on the surface of a sample.	learn more
Nanoindentor (UFCH24) - new Pro-NanoEnviCz II	The Hysitron TI 980 nanoindenter provides rapid, multi-sample, and multi-technique automated testing capabilities for high- throughput characterization. It includes quantitative nanoscale-to- microscale indentation, nano-scratch, nano-wear, high-resolution in-situ scanning probe microscopy (SPM) imaging, dynamic nanoindentation, and high-speed mechanical property mapping; providing a comprehensive understanding of material behavior at the nanoscale. The equipment enables: 1. quantitative determination of localized mechanical properties such as elastic modulus, hardness, creep, stress relaxation, and fracture toughness for a wide variety of materials, 2. continuous measurement of elastic-plastic and viscoelastic properties as a function of indentation depth, frequency, and time, 3. to obtain comprehensive nanomechanical property maps and property distribution statistics in a record amount of time.	<u>learn</u> more
Sensor characterization laboratory (UFCH 26)	The laboratory for characterization of sensors equipped by gas system, electrical parameter measurement unit, and optical excitation unit.	<u>learn</u> <u>more</u>
Surface Area and Pore Size Analyzer (BET) (UACH5)	The Surface Area and Pore Size Analyzer, which uses the static dosing method.	<u>learn</u> more
Surface Characterization System (UFCH32)	Combined ultra-high vacuum apparatus for complex study of thin films, interfaces and surface nanostructures (SPECSR) encompassing: - X-ray photoelectron spectroscopy (XPS) with microfocused (200 $\mu$ m) monochromatic X-ray source (hv=1486.6 eV) - ultraviolet photoelectron spectroscopy with excitation of electrons by monochromatized He I (21.2 eV) and He II (40.8 eV)	<u>learn</u> more

radiation - hemispherical electron energy analyzer with two-	
dimensional electron and ion detector and sample manipulator	
allowing measurement of high resolution spectra from room	
temperature down to liquid Helium temperature at different polar	
and azimutal detection angles, band structure mapping by angle-	
resolved photoemission spectroscopy (ARPES) technique using	
scanning angle lens - low-energy electron diffraction (LEED)	
technique for the determination of the surface structure and	
accurate surface atomic positions of materials - ion gun for	
cleaning of surfaces - scanning probe microscopy (SPM) for	
investigations at atomic scale of a wide variety of materials	

## XRD techniques

Instrument	Description	Link
Multipurpose X-ray powder diffractometer PANalytical XPertPRO MPD (UACH14)	Multipurpose X-ray powder diffractometer PANalytical XPertPRO MPD equipped with Cu K $\alpha$ or Co K $\alpha$ X-ray tube allows analyzing powdered or solid samples and/or micro-samples in reflection or transmission mode. This diffractometer is equipped with programmable divergence slit, focusing mirror and fast linear PSD detector. It allows in-situ analyses at elevated temperatures up to 1200 °C.	<u>learn</u> <u>more</u>
Multipurpose X-ray powder diffractometer (Co tube), Empyrean, series 3 (UACH17)	Non-destructive analysis, qualitative and quantitative phase analysis of crystalline solids, determination of amorphous content by indirect method using an internal standard addition are available. It enables studies of changes in materials connected with their applications, usage, functionality and caused by ageing, fatigue at operation conditions.	<u>learn</u> more
WDRF spectrometer Rigaku Primus IV (UJEP 37)	Tube-above wavelength dispersive X-ray spectrometer for fast elemental analysis of powders, liquids and bulk materials in range F-U. This spectrometer is equipped with micro-mapping utility for analysis of defects and for mapping of chemical composition. It also enables thin layer analysis and analysis of defects.	<u>learn</u> more
X-ray Powder Diffraction (UPOL7)	The instrument is used for identification of crystalline phases, quantitative phase analysis, determination of amorphous phase content, structural analysis of powder samples, determination of particle size, determination of Mean X-ray Coherence Length (MCL), determination of residual stress in (nano)material samples, monitoring and determination of structural/phase transformations in non-ambient conditions, and determination of temperature dependent dilatation.	<u>learn</u> more
X-ray difractometer Panalytical X (UJEP5)	Universal XRD powder diffractometer, with Cu K $\alpha$ x-ray tube. Measurements could be done in reflection and transmission mode. The device is equipped with smart detector, collimator, Göbbel mirror and Euler stage. Identification of unknown crystalline phase and qualitative and quantitative phase analysis of polycrystalline materials are provided.	<u>learn</u> more
X-ray powder diffractometer with optics for nanolayers and nanosurfaces Panalytical X Pert PRO (UJEP33)	X-ray diffractometer equipped with Co tube is usually used in Bragg-Brentano geometry with linear X'Celerator detector. The goniometer is vertical, the sample is placed horizontally. It is designed for measuring powder, bulk and thin layer polycrystal samples. X-ray diffractometer equipped with optics for structure analysis of polycrystalline thin films and nano-surfaces.	<u>learn</u> <u>more</u>