

<b>U41-TXM, HZB</b>	
<b>Microscope type</b>	TXM
<b>Source</b>	Undulator beamline, linear polarization
<b>Condenser/Objective aperture</b>	Partially coherent illumination
<b>Energy range</b>	250 – 1500 eV
<b>Spectral resolution</b>	up to $E/\Delta E = 10^4$
<b>Zone plates for user operation</b>	$dr_n = 25 \text{ nm}, 40 \text{ nm}$
<b>Spatial resolution</b>	2D: $\approx 10 \text{ nm}$ half-period (depends on objective) 3D: $\approx 36 \text{ nm}$ Rayleigh criterion
<b>Sample format</b>	<ol style="list-style-type: none"> <li>1) Special designed rectangular grids (HZB-2 or IFR-1) having 2 mm x 1 mm sample area and open grid slits width <math>50 \mu\text{m} \times 200 \mu\text{m}</math> or <math>150 \mu\text{m} \times 700 \mu\text{m}</math> these grids are cryo- and tomography- compatible</li> <li>2) 3 mm diameter EM grids(cryo possible and tomography) or <math>3.5 \cdot 3.5 \text{ mm}^2</math> wafer (no cryo and no tomography possible)</li> </ol>
<b>Tomography capability</b>	yes
<b>Tilt range</b>	$-80^\circ \dots +80^\circ$
<b>Cryo capability</b>	yes – $\text{LN}_2$ temperature
<b>Detector type</b>	Thinned, backside illuminated CCD, 1340 pixel x 1300 pixel
<b>X-ray magnification</b>	adjustable, value depends on the used objective
<b>Light microscope</b>	BF, DIC, fluorescence
<b>Filter sets for light fluorescence microscope</b>	Filter set 38 from Zeiss, GFP, ex 470, emm 525; Filter set 43 from Zeiss, Cy 3 (rhodamin), ex 545, emm 605
<b>Raw data format</b>	*.spe (WinView); 3.407 MB per image
<b>Data analysis software capabilities available</b>	ImageJ, Imod (eTomo)
<b>Preparation Lab</b>	BioLab (restricted access): safety level S1 (genetically modified organisms (GMOs) belonging to safety level S1 (GenTG) allowed); incubator; plasma cleaner; plunge freezer (liquid ethane); fluorescence light microscope LEICA DMI6000B; cryo light microscopy stage