JAMP Manual

Constituents/standard stand-by settings		
	Keep this place clean and leave it as you found it!	
	Ion gauge main chamber: pressure should be in low 10 ⁻⁷ Pa scale or lower (down to 5x10 ⁻⁸ Pa) Water flow meter for electron gun: Should show 0.2 or more	
	Control panel for turbo pump: Pump should be in "normal operation" Control panel for ion pump(s) and titanium sublimation pump: SIP1+SIP2 should be highlighted, left display should show around 6 kV Ti Pump used for improving vacuum, used when needed	
	lon gun control panel: See details in "Ion gun" chapter Check, that the main "On/Off" is turned to "On"	
	X-ray source: See details in "X-ray source" chapter	
	Control panel for ion gun operation	
	Control panels for SEM operation Upper: wobbler Left: movement Right: focus, magnification, alignment, scansize	
	Exchange chamber and manipulator	

Main chamber, SEM column, motors, ion gun, detector
View from the right side, detector and X-ray source
Water cooling pump for SEM column
Water cooling pump for X-ray source (not turned on at standard standby setting)
Water for SEM column (right, always on) and X-ray source (left)
Control panel and LED schema of the system (all highlighted LED's should be green, no button highlighted, "L" green at Penning gauge)
On/Off of main unit

Insert/remove sample		
	Inserting	
and the second sec	Mount sample on sample holder	
	For cross sectional holder: film towards spring + massive part in direction of transfer!	
	Press "exchange vent"	
	Open door exchange chamber	
	Wait until door opens and N2 stops to flow	
	Insert sample into manipulator in upper ring of sample holder	
	Take care that the manipulator is at the very left end (otherwise MGL1 LED will highlight orange)	
	Close door exchange chamber	
	Press "exchange vent"	
	Wait till pressure in low green region + green light "L" is on	
	Check stage position (at "sample change position"?)	
	Press "open V2"	

	Incort companies onto otages	
	Insert sample onto stage	
	Press "open V2"	
With Theorem 1	Choose right sample holder!	
	-Small sample holder (12mm)	-Big sample holder (20mm) -Cross sectional sample holder
	Tilt max = 90°	Tilt max = 55°
	Be aware that if "Auger Mo holder will be set to "free"! You have to go back to "s choose right holder again!	ister" software crashes, the !! ample change position" and !!
	Removing	
	Check stage position (at "samp	e change position"?)
	SEI detector OFF	
	PCD in (green)	
	Turn on ion gauge and check p	ressure in main and exchange
	Press "open V2"	
	Pick sample with manipulator a	nd withdraw manipulator
	Press "open V2"	
	Press "exchange vent"	
	Open door exchange chamber	
	Wait until door opens and N ₂ sto	ps to flow
	Remove sample	
	Close door as soon as possible ((see "Inserting"!)	and start pumping again

Modes SEM (for SEI detector)
	Mode 3 -> 300V (more e detected)
	Mode 2 -> 200V
	Mode 1 -> 100V
	Mode 0 -> 0V
	Mode -1 -> -300V (to detect BSE, but no BSE detector present!)
Aperture	
	4 -> 30µm (for high-res. SEM) (can be too less for AES)
	3 -> 50µm
	2 -> 70µm
	1 -> 110µm
	After changing aperture align everything again!

Acquiring pictures	
	Ion gauge off?
	Window flanges shut?
	PCD out (white) SEI detector ON
	Focus has to be set to/at 25!
	Accelerating voltage max 30kV
	(after changing accelerating voltage-> alignment!)
	Z max = ± 6mm around eucentric position
	Focus/eucentric position ~ 24mm
	Magnification from 35 to 100.000
	High magnification and big tilt? -> dynamic focus (DFC)!
Search for eucentric positio	n (when using for SEM + AES/Ion gun/XPS)
	Activate guiding lines
	Bring a line to a characteristic point
	Tilt some degrees
	Adjust z (bring point back on line)
	Tilt to 30°
	adjust
	Till back to 0°
	Mark characteristic point again with a guiding line
	Start procedure again
	No movement upon tilting? -> OK!

Using lon gun	
	Make sure, that the main "On/Off" is at "On" at the control unit
6 6	To turn on gas:
	AVC on
	Open the gas till 10x10 ⁻² Pa <mark>(slowly!)</mark>
	Switch the pin upwards
The Confidence (VV) Topology (V)	"Set" in software to 9x10 ⁻² Pa
	Check with "get"
	Example for cleaning/etching: Cu if big peaks appear: oxide layer < 6nm if small peak within the layer is appearing+O+N+C: oxide layer ≈ 3 nm -> ion gun, 60 seconds, channel 3
	lift sample to 30° for AES and 60° for XPS!
	Button on panel "Spot" non-highlighted (scanmode)?
	Flahing grogs 200-200-m
	Etching area: 300x300µm
	Accelerating voltage: IUKV

Channels: Choose a channel corresponding to the tilt and the sample you are using. There are channels for 0°, 30° and 60° + from 500 eV to 3000 eV. Than choose the time you want to etch in seconds.
To turn age off:
Press "close" in software and wait till 2x10-2 Pa
 Switch pin downwards
 Wait until it rises back to 10x10 ⁻²
Close the gas carefully and do not close it brutally!
AVC off
lon gage off
Neutralizing mode:
Works best if tilt ≥ 60° <mark>(check sample holder if it is possible! Not for 20mm holder!)</mark> Best: 75°
Dwell time 50->100
Probe current medium (6)
On controlling panel press -> Spot mode instead of scan mode (highlighted)
sweeps = min. 2 to see a possible peak shift due to charging

AES measurement	
Prink Charlenge Holy 4 Frank Charlenge Holy 7 Frank Charlenge Holy 6 Frank Charlenge Holy 6 Frank Charlenge Holy 6 Frank Charlenge Holy 7 Frank Charlenge Holy 6 Frank Charlenge Holy 7 Frank Charlenge Holy	Best to set probe current to 1x10-8: Sample observ> Probe cond> Auto probe current
	Focus
International International	Observation -> start digital scan If sample might charge: Observation -> transfer image from SEM
The contrast of the contrast o	Before acquisition (after ion gun+focus): Probe cond> OL -> lens clear -> OL (otherwise the magnetic field is not stable and peaks at lower energy are to small/worse)
Wide scan spectrum:	
	"scan": spectrum may shift
	You can defocus to enlarge the spot and the analyzed area
	Start spectrum at 30 eV (0-30 eV is SEM signal)
	Stop spectrum at 2000 eV (higher energy gives BSE-information)
	Choose acceleration voltage 10 kV to 30 kV
Analyzer conditions	
	M1-> const. E pass (for XPS)
	M2 to M5 -> M5 higher count rate, less resolution
	M4: standard
	M1: choose pass energy = for mappings (choose $a = -50 e^{1/3}$
	M2: 0.05% energy = 101 mappings (Choose e.g. $p=50 \text{ eV}$) M2: 0.05% energy resolution = XPS res
	M3: 0.1% energy resolution = XPS res
	= for chem. analysis
	M4: 0.35% energy resolution
	M5 0.5% energy resolution = for mappings (high int. but low res.)
	40000 counts is optimum for detector (with M5)
specifications	
	>0.1% detection limit
	I I-2% quantitative
	Spatial: 10-20 nm Ø

Z: 0-6 nm
Elements: Li-U
SEM: 3 nm, AES: 8 nm
Energy resolution: 0.05% to 0.6%
Chemical analysis: 0.1%
Standard spectra as reference:
aes/std_AES -> M2-M5

rs measurement	
u have to complete the whole p	procedure: start/stop/back to SEM!
	Tilt sample to 60°! (then it is 90° to analyzer)
	Make sure that you are at the eucentric position
	For XPS always use analyzer mode M1 (const. E pass)!
	Start spectrum at 100eV never at 0eV!
	Irradiated area is big (1cm²), analyzed area is defined by magnification (SEM picture)
	To start a measurement:
	Turn on water
	Turn on water pump
	Turn on x-ray source (power switch)
	Press water button till red square disappears
Land Long Barry British Date (1997)	Choose source: Al or Mg?
	Press "Standby"
Add The Constant Hap Add The Constant Hap Product tamps, cut(r) The dial Hap Product tamps, cut(r) The dial Hap Add The dial Hap	In Auger Master software: ??? -> "XPS Aquisition mode"-> On
and the second second second	Press "Hy On"
	Wait until value gets stable
	Press "Operate"
	To stop measurement:
	Press "Standby"
	Proce "Hy On"
	Turn off x-ray source (nower switch)

	PCD out
	e beam irradiate
Analyzer conditions	
	M1-> const. E pass (for XPS)
	M2 to M5 -> M5 higher count rate, less resolution
	M4: standard

Finishing SEM		
	SEI detector OFF	
	PCD in (green)	
	Accelerating voltage back to 10 kV	
See: Insert/remove sample		
Data transfer/transformation		
Finishing working		
	Please record your time of use in the red book	
	Turn off all three screens	
	Turn off gauges if on so far	

XPS measurement – a complete sequence

Pump the sample		
Turn on the ion gun		
	Make sure, that the main "On/Off" is at "On" at the control unit (orange)	
	To turn on the gas (best, when sample is already transferred):	
	Press "FIL ON" (highlights green)	
	Press "AVC" (Auto Value Control) on (highlights green)	
	Open the age till 10x10-2 Rg (slowly)	
	(menulo a little biokente 10 2 m10 2 Da)	
	(maybe a lime higher to 10.3 x 10 ⁻² Pa)	
	Switch the pin upwards	
	If sample is already transferred:	
	Press "Get"	
	If sample is still in exchange chamber:	
	Press "Close"	
Open the X-ray source		
	Turn on water	

	Turn on water pump
	Turn on x-ray source (power switch)
	Press "water" button till red square disappears
	Press "Standby"
	Press "HV On"
	Wait until value gets stable (10 kV)
AES Lon Cun Condition	To set the condition in software:
	Make sure the ion gun valve is closed before sample transfer
Sample Alignment	
	Insert the sample (always the small sample holder Φ 12 m)
	Search for eucentric position $ ightarrow$ tilting to 60° (right side)
	Focus the image (choose the area signal by choosing mag)
	Close the SEI and PCD
Etching the sample	1
	In software set to 9x10 ⁻² Pa by clicking "Set" Check with "Get"
Marcing Contraction Marcing Contraction	Channels:
	Choose a channel corresponding to the tilt and the sample you are using. There are channels for 0° , 30° and 60° from 500 eV to 3000 eV.
	Then choose the time you want to etch in seconds.
	Press "ON" in Etching
	Time is recorded on the panel Press "Reset" to make it back to 0

Perform a wide scan		
$\begin{array}{l} \text{AES} \rightarrow \text{XPS Aquisition} \\ \text{mode} \end{array}$	Check that XPS Aquisition mode is on!! ■ XPS Acquisition mode	
$\begin{array}{l} \text{AES} \rightarrow \text{spectrum} \rightarrow \text{wide} \\ \text{scan spectrum} \rightarrow \\ \text{condition} \rightarrow \text{analysze} \end{array}$	M1; <u>xx</u> eV; Start 100 to 1500 eV; step <u>xx</u> eV; Dwell time 100ms; sweeps 1	
condition \rightarrow Analysis position	Put the spot in the center of image and set Prob. Diam. to 10 μm	
In wide scan spectrum window	Change the saving condition	
	Press the "Operate" button on the X-ray source (250 power)	
Acquisition \rightarrow Start	The result is automatically saved	
	Press "Standby" on the X-ray source	
Perform a detailed scan		
	Press the "Operate" button on the X-ray source (250 power)	
AES \rightarrow condition \rightarrow analyszer condition	M1; <u>xx</u> eV; Start <u></u> to <u></u> eV; step <u>0.05 or 0.01</u> eV; Dwell time 100ms; sweeps <u>3 to 5</u> .	
Acquisition \rightarrow Start	The result is automatically saved	
	Press "Standby" on the X-ray source	
To turn the ion gun off:		
	Press "close" in software and wait till <mark>2x10⁻² Pa</mark> Check with "Get"	
	Switch pin downwards	
	Wait until it rises back to 10x10 ⁻²	
	Close the gas carefully and do not close it brutally!	

	"AVC" off	
	"FIL On" off	
To turn the X-ray source off:		
	check if it is "Standby" on the X-ray source	
	Press "HV On"	
	Turn off X-ray source (power switch)	
	Turn off water pump	
	Close water	
To get back to SEM:		
	In Auger Master software: AES -> "XPS Aquisition mode"-> Off	
	XPS Acquisition mode	
Data Conversion		
Processing \rightarrow Data Conversion \rightarrow "VAMAS" \rightarrow "Yes" \rightarrow "OK" \rightarrow Choose Al Ka \rightarrow "Apply and close" \rightarrow "Cancel" \rightarrow "Exit"		
Transfer the .NPL file from ftp to mobile disc		