# Basics of daily corrector-tuning

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# **Principle remarks**



# **Daily corrector alignment**

- Is fast and fine-tuning against hysteresis of magnetic elements and thermal drift.
  - => If possible, prepare the microscope the evening before (HT, mode, sample)
- Is never a complete realignment of the corrector.
- Timeconsuming alignments indicate:
  - Bad alignment files
  - Broken hardware
  - Tuning beyond requirements or beyond measuring accuracy
  - Omitted check points during the daily alignment procedure

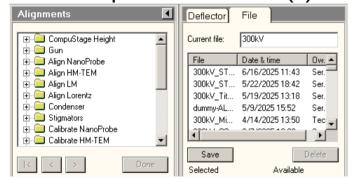
# **Preparations**



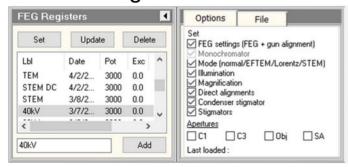
## 1. Load trustful alignment files and choose mode (HT, TEM/STEM)

## **ThermoFisher**

Microscope and corrector(s)

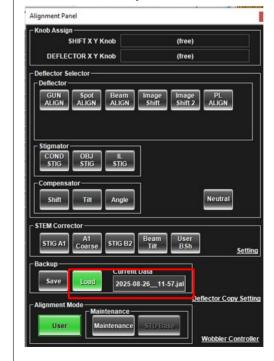


## User settings

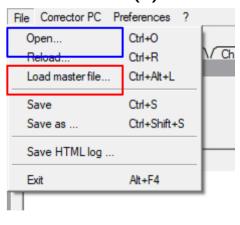


## **JEOL**





## Corrector(s)



# **Check points STEM**



# 2. Verify fix points and do basic pre-alignments

ThermoFisher	JEOL
- Eucentric focus	- Standard focus
- Normalise all	- Lens relaxation
	- Normalise correctors
	- Center beam in imaging mode:
- Leave diffraction, use AT L	JserBeamShift - Switch to ALIGN, use AT ShiftOL
	- Center the right aperture on last condenser lens
	(or on flat area on Ronchigram)
	- Adjust with AT BeamTilt:
- Rotation cent	er of objective - High tension center
	- Focus (in Ronchigram) with z-height on gold cluster sample
	- Select proper magnification (FoV around 500nm)

# **Check points TEM**



# 2. Verify fix points and do basic pre-alignments

ThermoFisher	JEOL
- Eucentric focus	- Standard focus
- Normalise all	- Lens relaxation
	- Normalise correctors
	- Center beam with microscope Beam Shift
	- Find nice amorphous area
	- Check beam tilt pivot points (tiny changes if at all due to hysteresis of objective lens)
	- Adjust microscope BeamTilt w.r.t. specimen detail:
- Rotation center objective	- High tension center
	- Set slight underfocus with z-height
	- Select proper magnification (C1A1 range 12um)

# Standard manual tuning loop for TEM and STEM



## 3. Use corrector UI's measurement procedures and auto-alignment tools

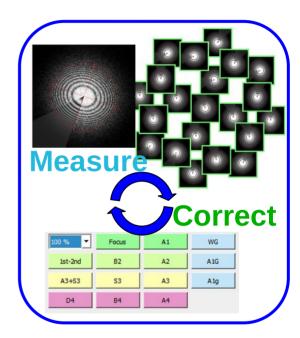
#### **Iterate:**

- Continuous C1A1 measurements (up to second order in STEM, up to first order in TEM)
  - Use auto-alignment tools while measurement is running
- Tableau measurements:fast -> standard (-> enhanced if needed)
  - Use auto-alignment tools after accepting tableau in state of correction

Cross-check with the instruction manual for details.

Concentrate on your experiment, not onto corrector tuning.

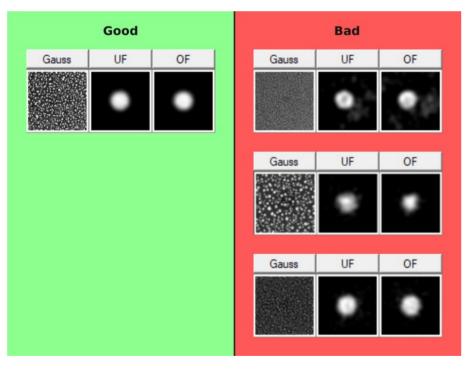
Stop iteration if experimental needs are fulfilled or measurement accuracy is limiting!



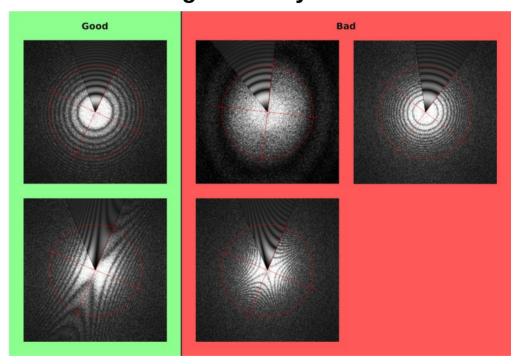
# **Good and bad continuous measurements**



## **STEM:** deconvolution



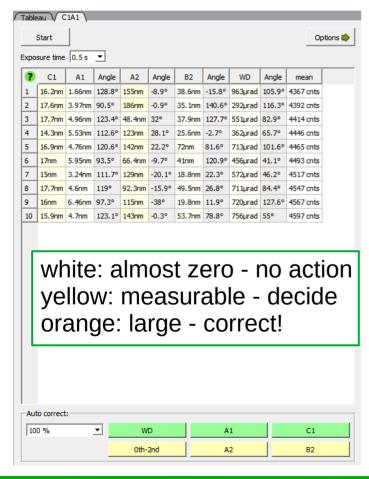
## **TEM:** diffractogram analysis



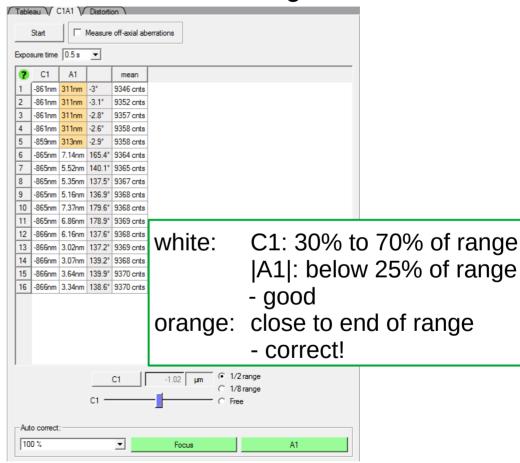
## **Hints at continuous measurements**



## **STEM:** color code for size



## **TEM:** color code for range "violation"

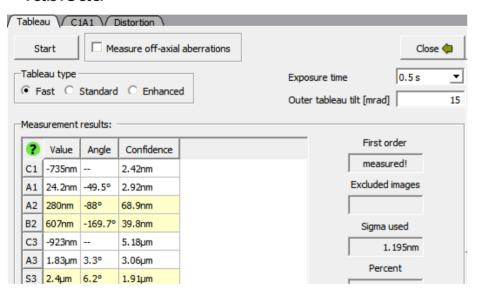


## Hints at tableau measurements and in state of correction

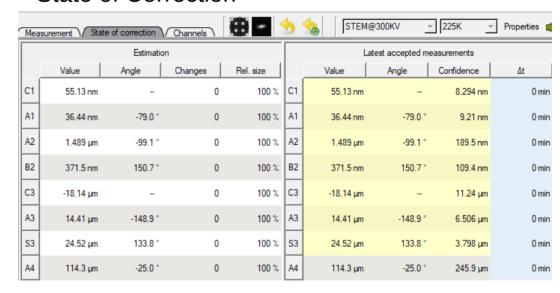


#### TEM and STEM: color code for well measurable aberrations

#### Tableau



## State of Correction



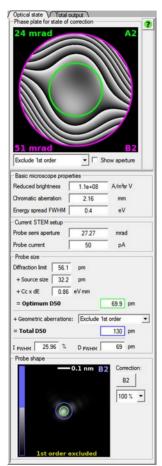
white: almost zero - no action

yellow: above measuring accuracy - correctable

# **STEM:** Tuning towards experimental requirements



#### **STEM: Probe state tool**



## **Desired optical state = "zero" aberrations**

- Reduce all aberrations sufficiently
- ← Phase plate with pi/2 circle and limiting aberration, considered aberrations selectable
- Microscope properties
- ← STEM setup
- ← Ideal and attainable probe size
- → Probe shape and most limiting aberration

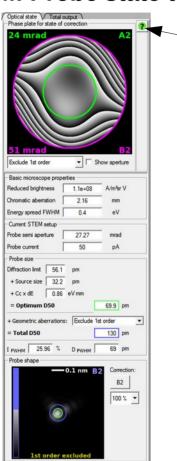
## Most important decision:

Good enough for desired experiment?

# **STEM:** Tuning towards experimental requirements



#### **STEM: Probe state tool**

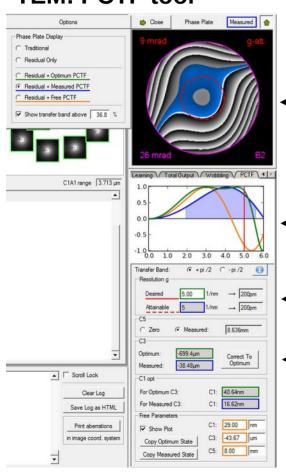


See help dialog behind "Info" button for detailed instruction and complete workflow description.

# **TEM:** Tuning towards experimental requirements



#### **TEM: PCTF tool**



## **Desired optical state** ≠ **zero aberrations**

- Shape phase contrast transfer function with round aberrations
- Reduce other aberrations sufficiently
- ← Phase plate, considered aberrations selectable

- ← Phase contrast transfer function
- Desired and attainable resolution
- Optimisation of spherical aberration C3

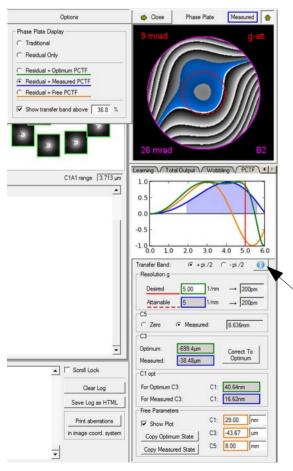
## Most important decision:

Good enough for desired experiment?

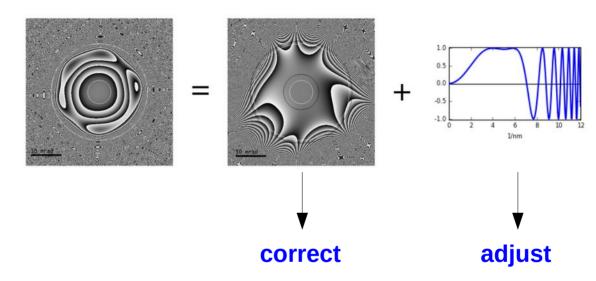
# **TEM: Tuning towards experimental requirements**



#### **TEM: PCTF tool**



#### total aberration = residual aberration + PCTF



See help dialog behind "Info" button for detailed instruction and complete workflow description.

# Finetuning during experiment: Exported elements



## **Corrector GUI**

- **→** Channels
  - → Exported elements

