

# **OBF** System

#### Live Low Dose, Light Element Imaging



JEOL Ltd.

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In the new imaging method '**OBF STEM (Optimum Bright Field STEM)**', raw images acquired by a segmented STEM detector are used as the source for a phase image reconstruction, with dedicated Fourier filters to maximize the signal to noise ratio of retrieved image.

This promising method realizes higher contrast for both heavy and light elements even while operating under extremely low electron dose conditions. Beam sensitive materials difficult to observe with standard ADF and ABF STEM methods can be easily analyzed with higher contrast at a wide range of magnifications.

K. Ooe, T. Seki, et al., Ultramicroscopy 220, 113133 (2021)

### Ultra High Dose-Efficient STEM Imaging

The beam sensitive materials including Metal Organic Frameworks (MOFs) and Zeolites require a reduced electron dose (typically, probe current < 1.0 pA) while maintaining clear atomic contrasts for the framework of light elements. The OBF STEM has an advantage for such low dose experiments, realizing ultra high dose-efficient STEM imaging in an atomic resolution. The OBF STEM image of MOF MIL-101 (**figure 1**) and MFI Zeolite (**figure 2**) can be acquired in a single shot with higher spatial resolution 1.8 Å and 1.0 Å, respectively. The enhanced contrast and resolution can be also confirmed in each stack averaged image (insets).





ABF (inverted)

OBF



Figure 3: SrTiO<sub>3</sub> [001] JEM-ARM300F2 Eta-corr. FHP2, Acc.300 kV, alpha = 24 mrad, Probe current = 0.8, 0.5, 0.3, and 0.1 pA

**-** 0.5 nm

#### High Contrast Imaging for Light Elements

In addition to being highly dose efficient, OBF STEM is also advantageous for light element imaging. Even in a lower acceleration voltage, both higher contrast and spatial resolution can be achieved for light elements.



Figure 4: GaN [110] JEM-ARM200F ASCOR UHR, Acc.60 kV, alpha = 35 mrad



Figure 5: Graphene JEM-ARM200F ASCOR UHR, Acc.60 kV, alpha = 35 mrad

The resolution for light elements becomes much better with a higher acceleration voltage. Each atomic column is now clearly separated with a deep sub-angstrom resolution inside complex structures (**figure 5**) or along higher-index crystallographic axes (**figure 6**).

The quality of OBF STEM is excellent in low dose conditions, and further enhanced under the standard probe conditions of a Cs-corrected electron microscope.



Figure 6:  $\beta$ -Si<sub>3</sub>N<sub>4</sub> [0001] JEM-ARM200F ASCOR UHR, Acc.200 kV, alpha = 24 mrad, Inset) 10 frames averaged



Figure 7: GaN [211] JEM-ARM300F2 Eta-corr. FHP2, Acc.300 kV, alpha = 32 mrad, Inset) 20 frames averaged

0.3 nm

When the OBF method is combined with smaller illumination angles, as in the case of non Cs-corrected instruments or when operating with the objective lens turned off (STEM Lorentz mode), the micro- and mesoscopic structures can be analyzed in a large field of view.

In the fields of biological, polymer, or semiconductor materials, it is often necessary to observe fine features inside a complex structure. OBF STEM is very effective toward this goal due to its enhanced contrast for both lower and higher frequency information.



**Figure 8**: Mitochondria in mouse kidney, unstained JEM-ARM300F2, Acc.300 kV, alpha = 0.13 mrad

Figure 9: Semiconductor device JEM-ARM200F, Acc.200 kV, alpha = 2 mrad

#### Live OBF Imaging

In an actual experiment, live OBF imaging is fundamental for beam sensitive materials as all operations should be performed in a dose limited condition. The live function is included in the OBF system, implemented within the TEM control software, with simple GUI control and real time display updates alongside conventional STEM images.



\*Specifications are subject to change without notice.

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