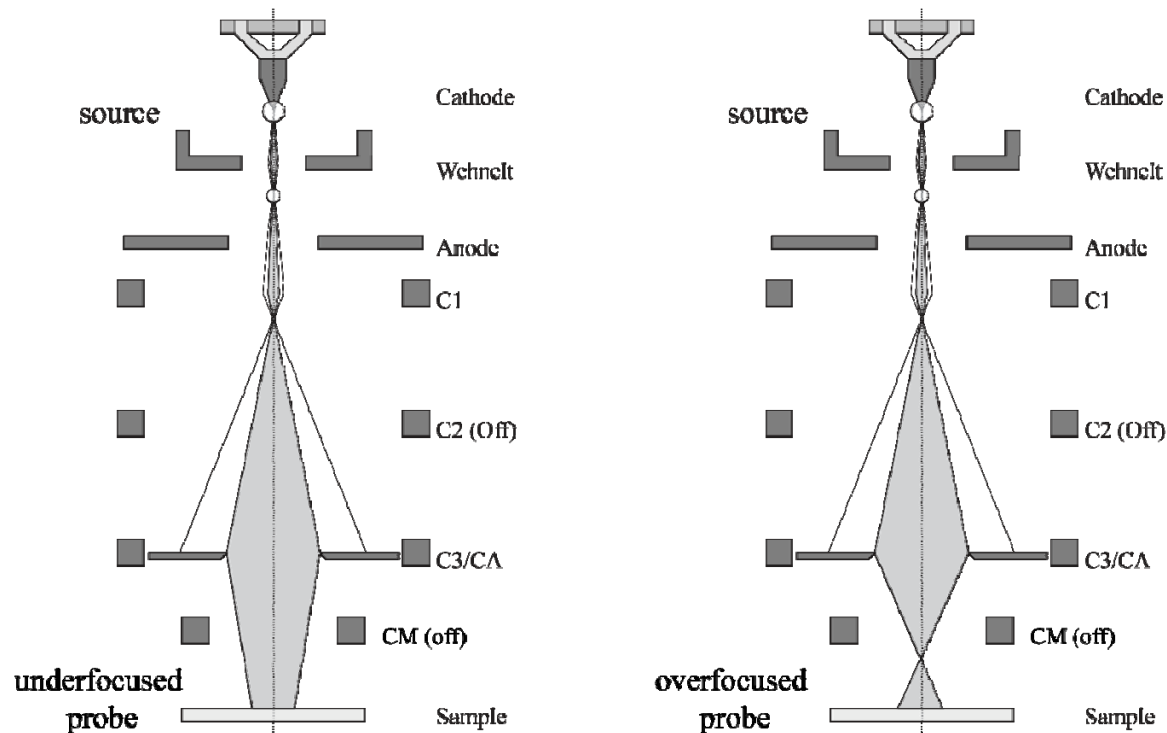


Illumination (Condenser) System

Functions:

- 1) Demagnify source (C1 - Spot Size)
- 2) Focus beam on sample with sufficient intensity
- 3) Vary probe size with magnification
- 4) Allow control of illuminated area (C3)



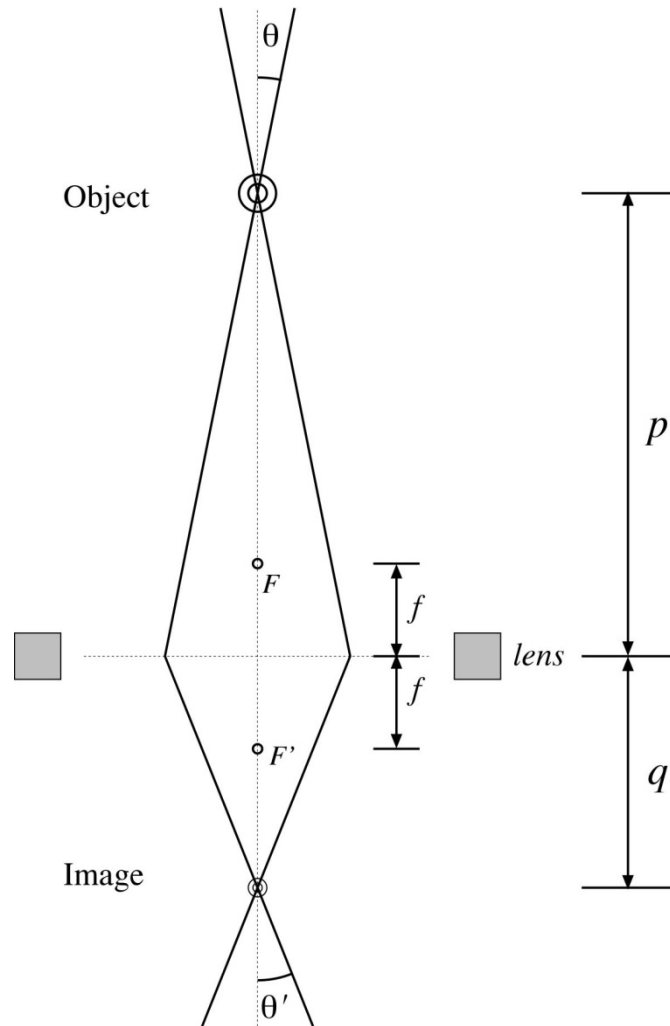
Demagnification of Source

Lateral Magnification

$$M = \frac{q}{p} = \frac{1}{M_\theta} = \frac{\theta}{\theta'}$$

Forming a small probe:

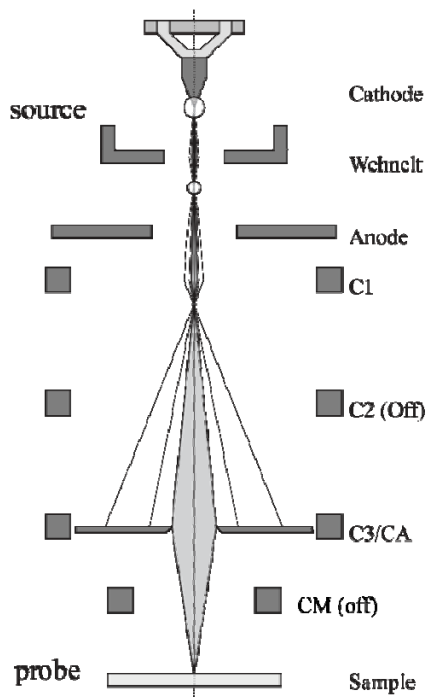
$$\theta' > \theta \Rightarrow q < p \Rightarrow M < 1$$



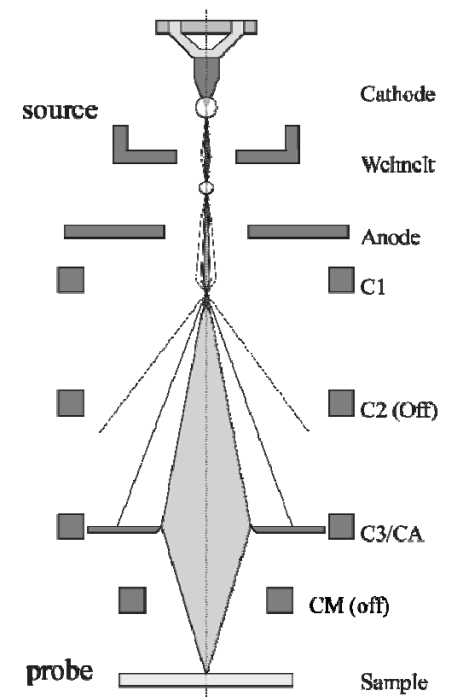
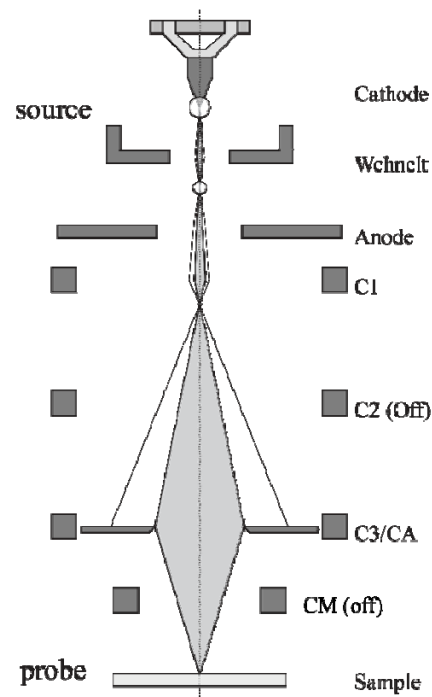
Demagnification requires increasing angular magnification

Convergence angle and probe size

- Convergence angle decreases with decreasing CA size
- Probe size decreases with increasing C1 excitation



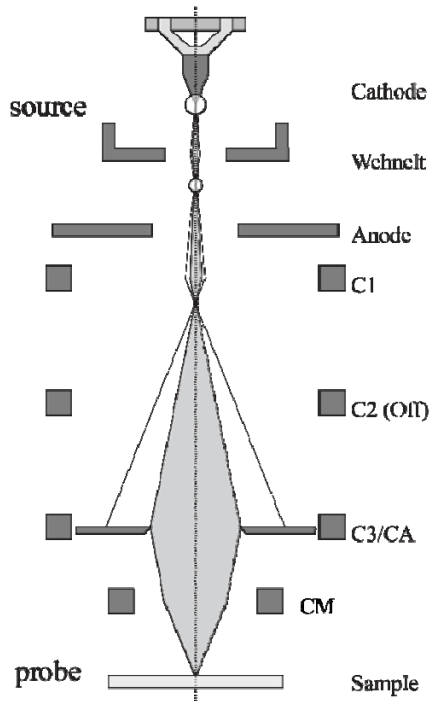
smaller α



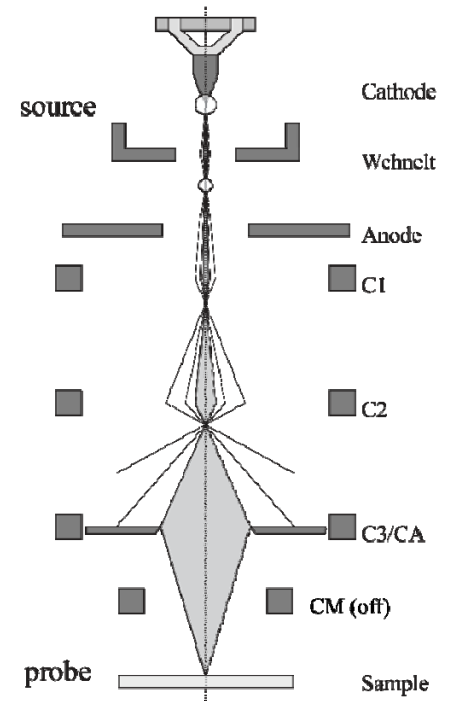
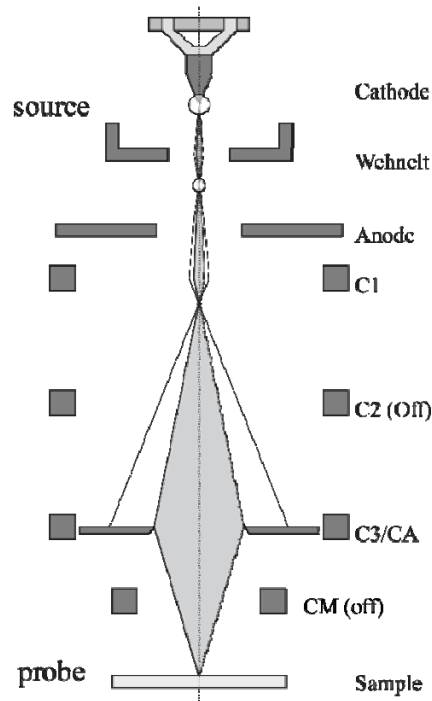
smaller probe

Uses of additional condenser lenses

- Condenser mini-lens (CM) allows larger convergence angles
- C2 lens allows smaller probe sizes



larger α



smaller probe

Comparison: C2 off vs. C2 on

C2 On

$$q_2 + p_3 = L$$

$$p_2 = P$$

$$q_3 = Q$$

$$\frac{1}{f_2} = \frac{1}{p_2} + \frac{1}{q_2} = \frac{1}{P} + \frac{1}{q_2}$$

$$M_2 = \frac{q_2}{p_2} = \frac{q_2}{P} = \frac{1}{\frac{P}{q_2} - 1}$$

$$M_3 = \frac{q_3}{p_3} = \frac{Q}{L - q_2} = \frac{Q}{L - \left(\frac{1}{\frac{1}{f_2} - \frac{1}{P}} \right)}$$

$$d' = M_3 \cdot M_2 \cdot d = \frac{Q}{L \cdot P} \cdot \frac{1}{\frac{1}{f_2} - \frac{1}{L} - \frac{1}{P}} \cdot d$$

$$\Rightarrow M_{C2 \text{ on}} = \frac{Q}{L \cdot P} \cdot \frac{1}{\frac{1}{f_2} - \frac{1}{L} - \frac{1}{P}}$$

C2 Off

$$p_3 = P + L$$

$$q_3 = Q$$

$$d' = M_3 \cdot d = \frac{q_3}{p_3} \cdot d = \frac{Q}{P + L} \cdot d$$

$$\Rightarrow M_{C2 \text{ off}} = \frac{Q}{P + L}$$

Compare: $M_{C2 \text{ on}} < M_{C2 \text{ off}}$

$$\cancel{\phi} \cdot \frac{1}{L \cdot P} \cdot \frac{1}{\frac{1}{f_2} - \frac{1}{L} - \frac{1}{P}} < \frac{\cancel{\phi}}{P + L}$$

$$\frac{1}{P} + \frac{1}{L} < \frac{1}{f_2} - \frac{1}{L} - \frac{1}{P}$$

$$f_2 < \frac{1}{2} \cdot \left(\frac{1}{\frac{1}{P} + \frac{1}{L}} \right)$$

(best to have C2 on and strongly excited.)

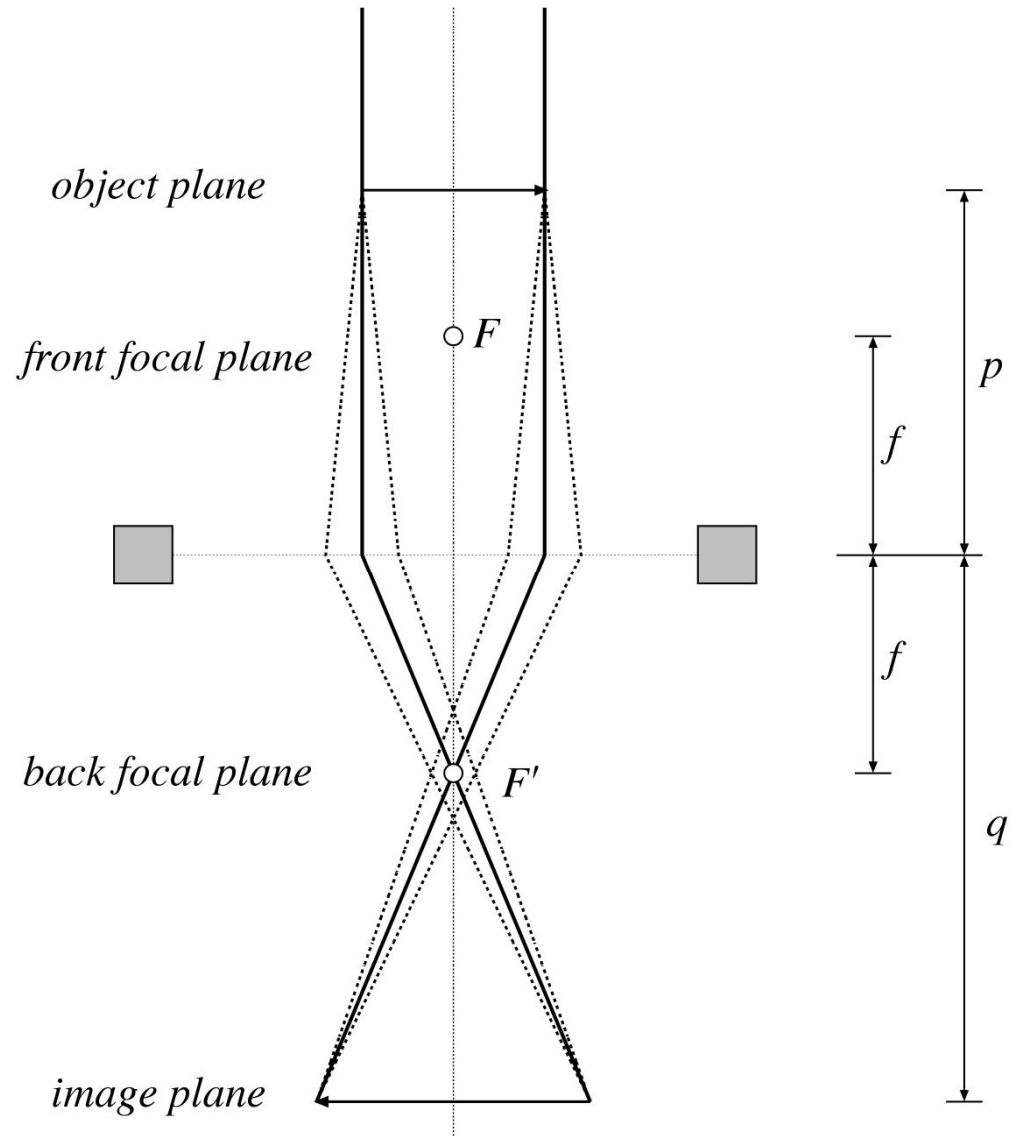
Lens Planes

Parallel Illumination

Diffracted beams deviate from direct beam

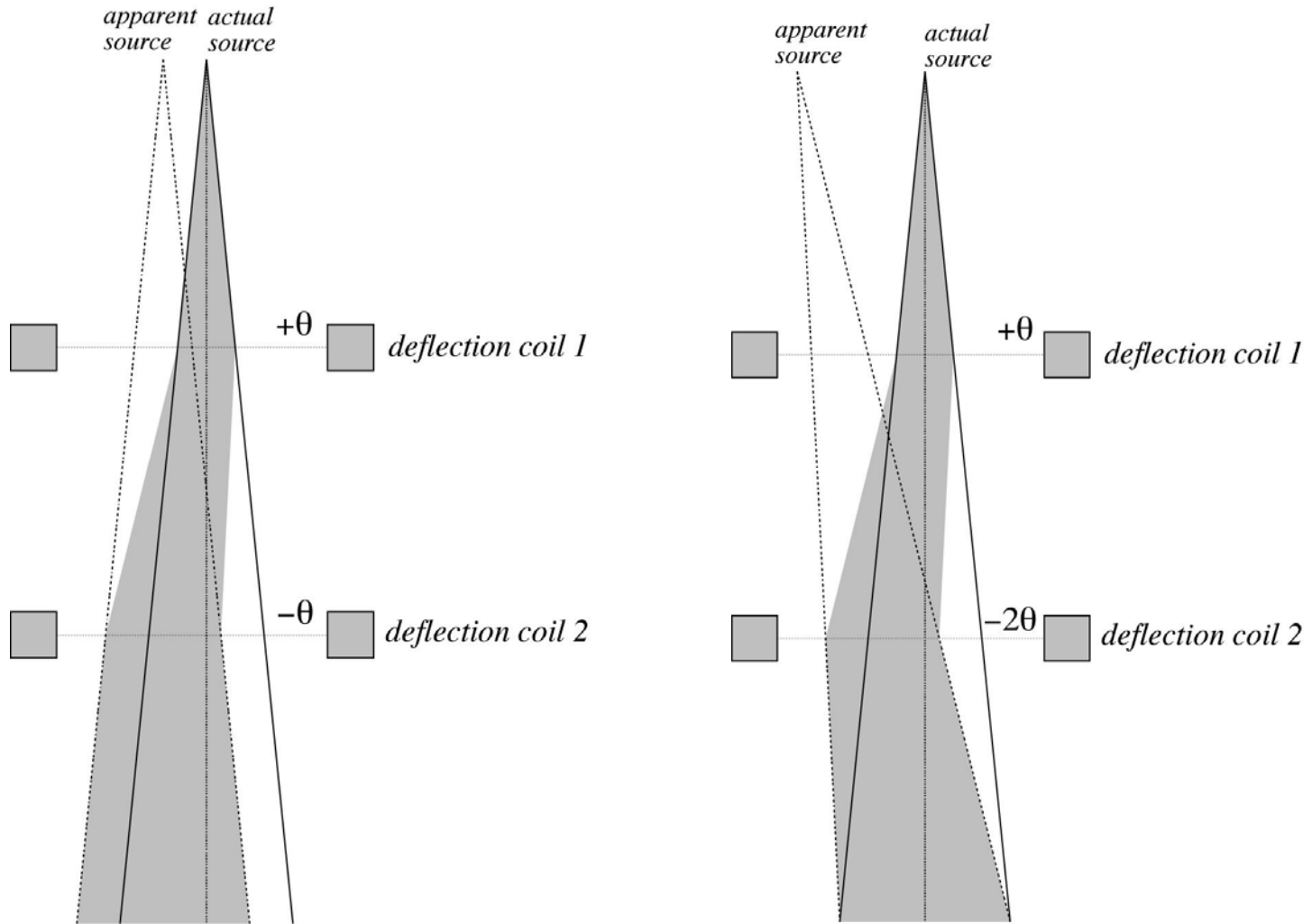
Parallel beams focused to a point in back focal plane

→ diffraction pattern



Beam deflection/tilt

Two deflection coils provide deflection/tilt



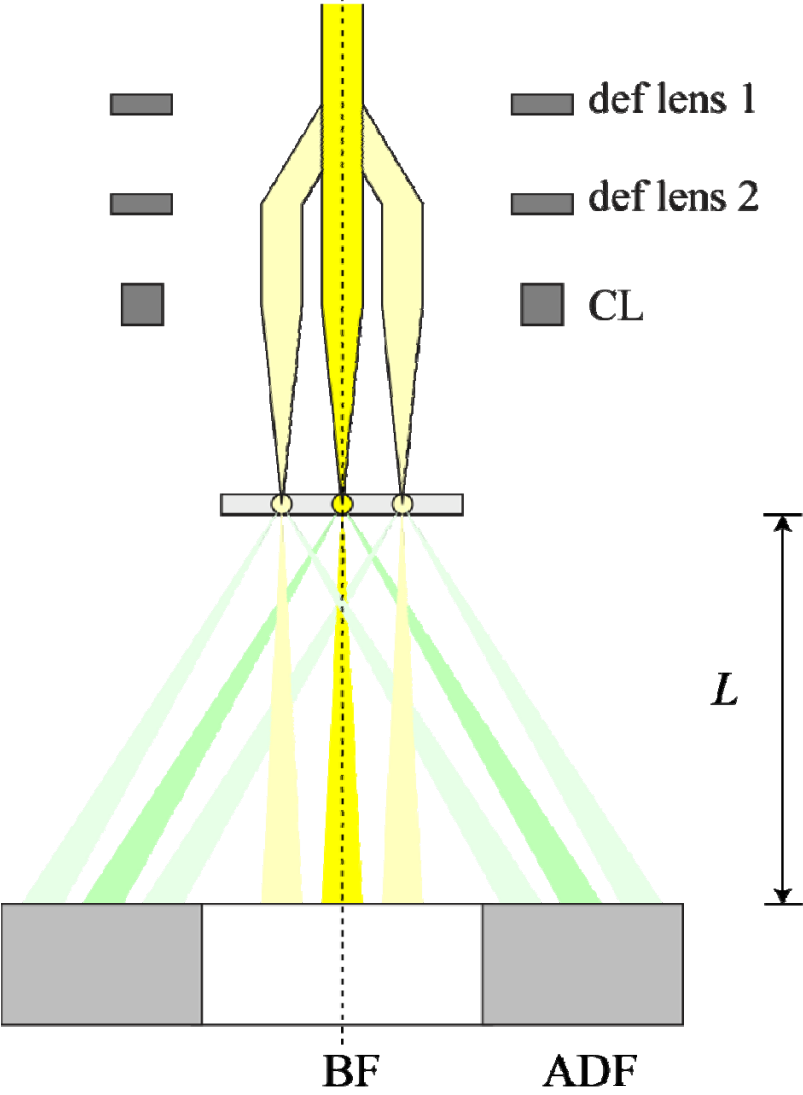
Scanning TEM

Scan beam across sample

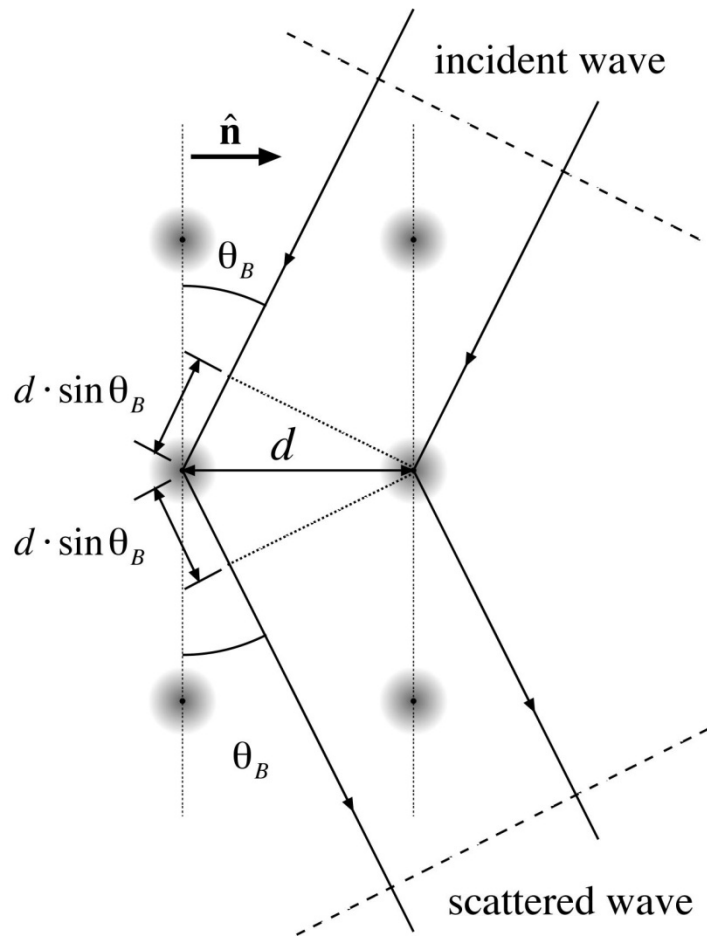
Illumination deflected, not tilted

Axial bright-field detector

Annular dark-field detector



Bragg's law and lattice vectors



Bragg's Law

$$2d \sin \theta_B = n\lambda$$

Reciprocal Lattice Vector

$$\vec{\mathbf{g}} = \left(\frac{1}{d} \right) \hat{\mathbf{n}}$$

Objective aperture placement

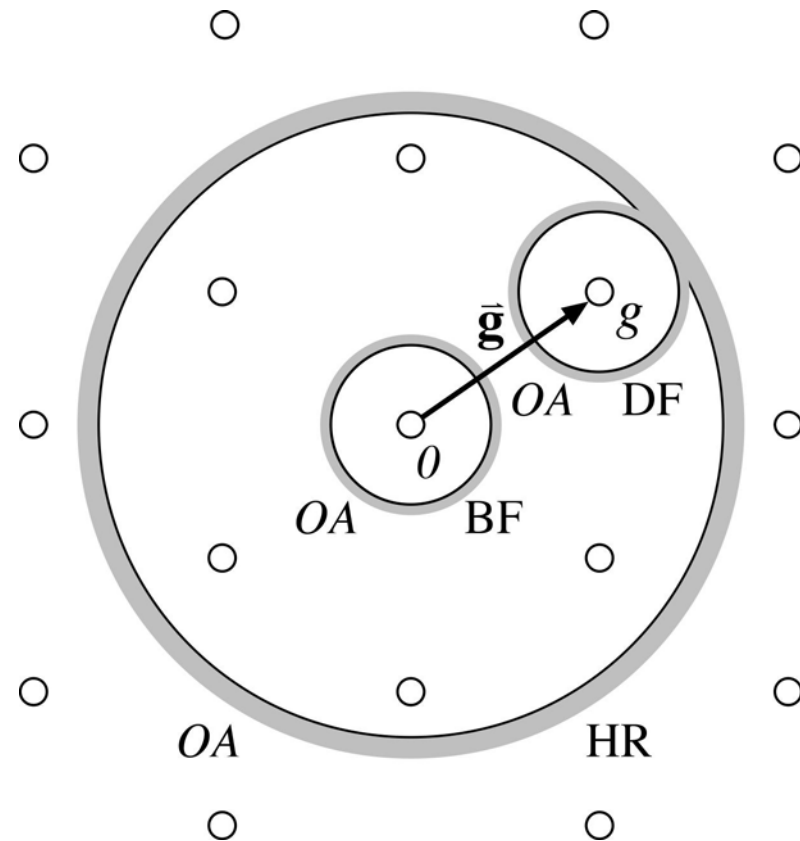
Position of objective aperture in back focal plane

Imaging Modes:

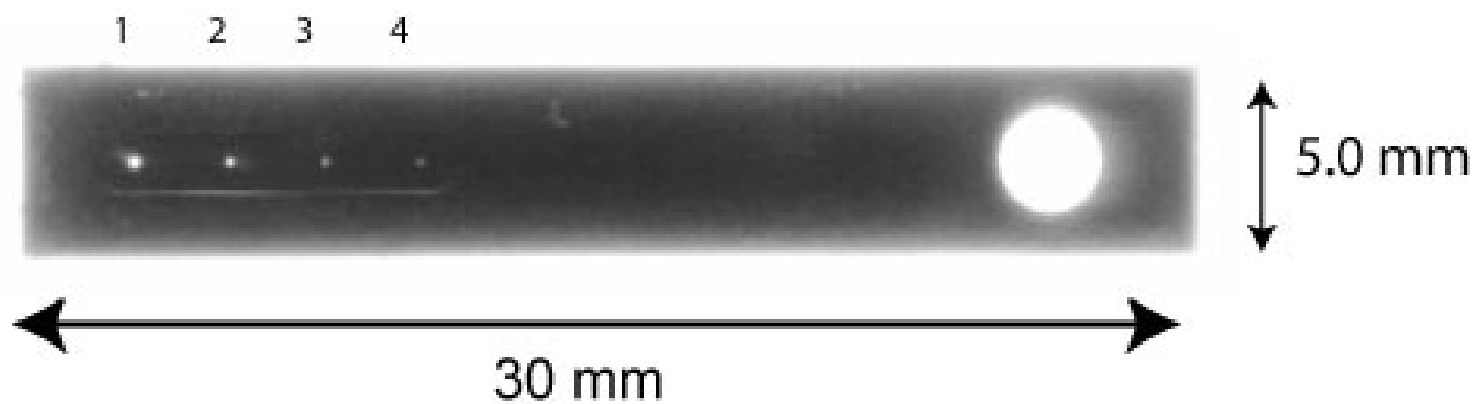
Bright Field: OA includes 0

Dark Field: OA excludes 0

High-Resolution Lattice Image:
 OA includes 0 and several beams



Objective aperture strip for Hitachi TEM



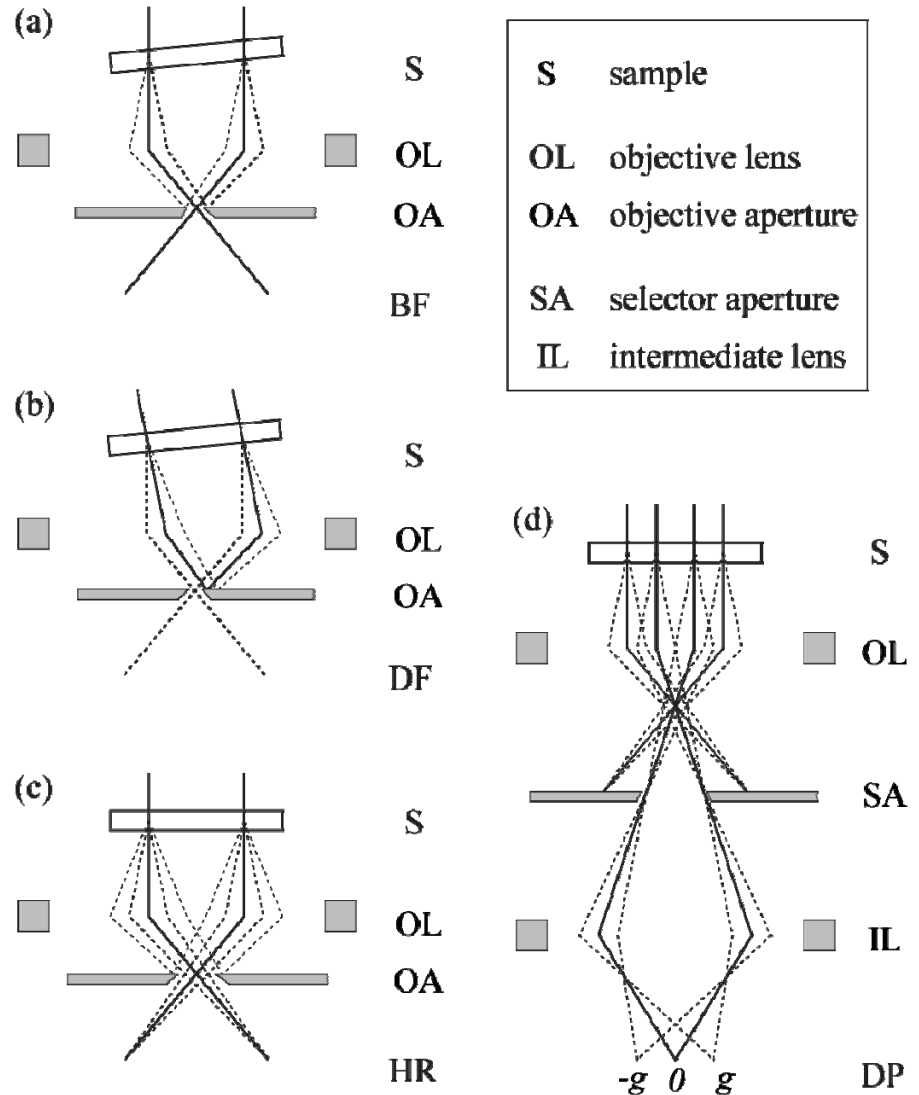
TEM Operational Modes

Bright Field (BF):
OA includes 0
 Two-beam condition

Dark Field (DF):
OA excludes 0
 Two-beam condition

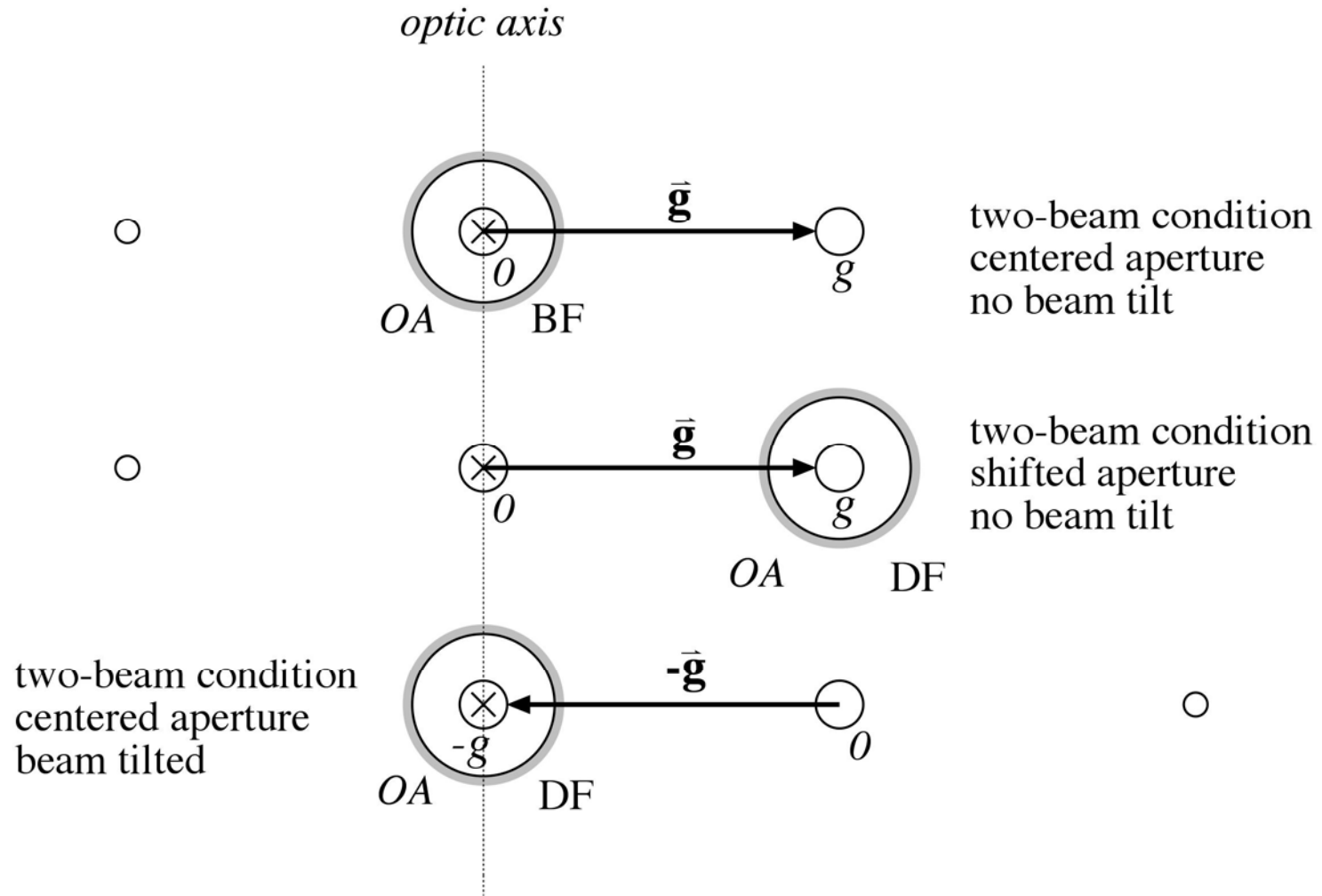
High-Resolution Lattice Image (HR):
OA includes 0
 Orient on low-index zone axis

Selected-Area Diffraction Pattern (DP):
SA in image plane



Bright-/dark-field methods

Objective aperture placement in back focal plane

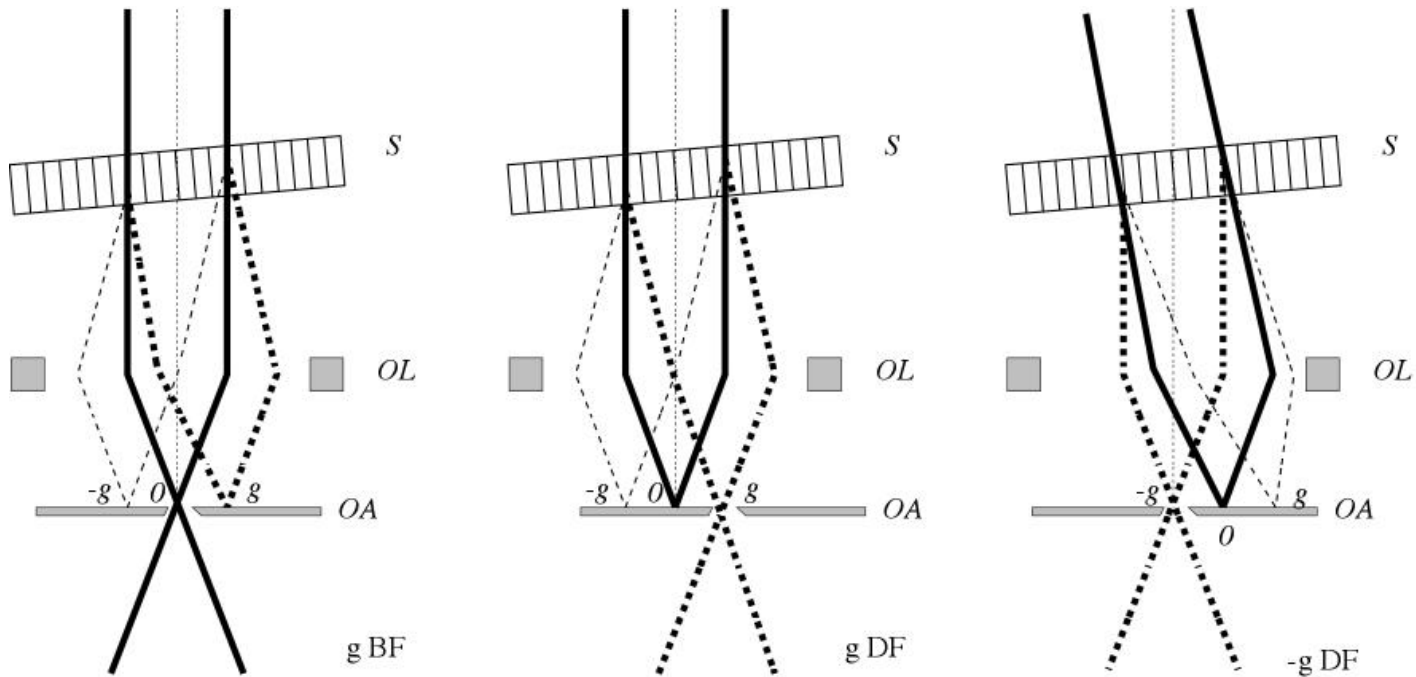


Centered dark-field imaging

Two-beam condition

Tilt the beam to generate dark-field with the scattered beam on-axis

Set up diffraction condition for g , then bring $-g$ parallel to the optic axis.



bright-field

off-axis dark field

centered dark field