Chapter 9 The introduction of EELS *EELS principle*

(Chap. 37, 38, 39, 40)

Textbook: R.F. Egerton, Electron Energy-Loss Spectroscopy in the Electron Microscope, 2nd edition

Particle picture of scattering



elastic

inner-shell inelastic



outer-shell inelastic

TEM beam-specimen interactions and signals



Incident high-energy beam

What is EELS

- Electron Energy Loss Spectroscopy

EEL spectrum is collected series energy loss electrons which generated with the inelastic scattering collision with specimen



Atomic-scale view of electron energy loss in TEM



energy-band diagram



Core ionization edges and the core level diagram



Nomenclature of EELS ionization edges





Ray Tracing for a 90 degree magnetic Sector Spectrometer



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EELS instrumentation spectrum/Imaging



- Below the TEM:
- Serial EELS (e.g. Gatan 607)
- Parallel EELS (e.g. Gatan 666)
- Gatan Enfina
- Gatan Imaging Filter
- In-column:
- Prism-mirror (Leo)
- Omega Filter (Leo, JEOL)

A magnetic prism bends, disperses and focuses an electron beam



Gatan Image Filter (GIF)



Leo-922 energy-filtering TEM



Omega-filter in-column spectrometer (Four magnetic prisms)



The EELS looks like





Intensity

EELS spectral information



Jellium Model

The resonant motion of electron gas would be self-sustaining if there were no damping from the atomic lattice.

The displacement x of a "quasi-free" electron (effective mass m) due to a local electric field E must satisfy the equation of motion.

$$m\ddot{x} + m\Gamma\dot{x} = -eE$$

for a oscillatory field

$$E = E \exp(-i\omega t)$$

The displacement has a solution given by

$$x = (eE/m)(\omega^2 + i\Gamma\omega)^{-1}$$

The displacement x give rise to a polarization P

$$P = -enx = \varepsilon_o \chi E \qquad \qquad \varepsilon = 1 + \frac{P}{\varepsilon_0 \mathscr{E}}$$

 χ is the electronic susceptibility and n is the number of electrons per unit volume

The relative permittivity or dielectric function $\varepsilon(\omega)=1+\chi$ is then given by

$$\varepsilon(\omega) = \varepsilon_1 + i\varepsilon_2 = 1 - \frac{\omega_p^2}{\omega^2 + \Gamma^2} + \frac{i\Gamma\omega_p^2}{\omega(\gamma^2 + \Gamma^2)}$$

 ω_p is the plasmon frequency (the frequency ε_1 passes through 0)

$$\omega_p = \left(ne^2 / (\varepsilon_o m)\right)^{1/2}$$

The energy loss function is defined as

$$\operatorname{Im}\left[\frac{-1}{\varepsilon(\omega)}\right] = \frac{\varepsilon_2}{\varepsilon_1^2 + \varepsilon_2^2} = \frac{\omega\Gamma\omega_p^2}{(\omega^2 - \omega_p^2)^2 + (\omega\Gamma)^2}$$

Drude Model for Volume Plasmon

For RuO₂ a=b=0.449 nm, c=0.31 nm (one unit cell has 2Ru and 4O)

Ru : [Kr] $4d^{7}5s = 8$ O : [He] $2s^{2}2p^{4}=6$

of free electrons = 40=2x(8+2x6)

$$\mathbf{A} = \frac{40}{(4.49)^2 \times (3.1) \times 10^{-30}} \left[\frac{\#}{cm^3}\right] = 6.4 \times 10^{29} [m^{-3}]$$

$$\mathbf{\epsilon}_0 = 10^7 / 4\pi c^2 = 8.842 \times 10^{-12}$$

$$\omega_p = \left(\frac{ne^2}{m\epsilon_0}\right)^{\frac{1}{2}} = \left(\frac{(6.4 \times 10^{29}) \times (1.60219 \times 10^{-19})^2}{(9.1 \times 10^{-31}) \times (8.842 \times 10^{-12})}\right)^{\frac{1}{2}} = 4.5158 \times 10^{16}$$

$$E_p = \hbar \omega_p = \frac{4.5127 \times 10^{16} \times 1.05459 \times 10^{-34}}{1.60219 \times 10^{-19}} = 29.7271 eV$$

While m=9.10956x10⁻³¹ kg

e=1.60219x10⁻¹⁹ C $\epsilon_0 = 10^7/4\pi c^2 = 8.842x10^{-12}$ 1 eV=1.60219x10⁻¹⁹ J

Plasmon loss from RuO2 nanowires



Correlating electron energy levels with EELS edges



Energy-loss spectrum (log-intensity) of YBCO





(a) Energy dispersion curve and (b) DOS $(\rho(E))$ for an electron in a square potential well with infinite sides.



Simplistically speaking, this means that flat regions in a band structure diagram will correspond to peaks in the DOS and therefore peaks in the ELNES.















What EELS can do

Typically applied to: * measurement of specimen thickness * analysis of elemental composition * phase identification via signature in EELS fine structure

Also applicable to studies of:

* electronic band structure and chemical bonding
* atom-specific near-neighbor distributions (RDF)
* Band gap analysis for optoelectronic material
* dielectric response, ε(ω,q)

Measurement of specimen thickness



 $\lambda \sim 100 \text{ nm}$ but depends on Z, E₀ and β Value obtained from calibration specimen or from tables (for common materials) or from parameterized formula

EELS fine structure

(Egerton and Whelan, 1974)



K-edge spectra of diamond and grain from the Allende meteorite

(Blake et al., Nature **332**, 1988, 611)





Oxidation state of Cr in a bacterium (Daulton et al,

M&M 7, 479, 2001)



Compare the EELS and EDX technique

- Prior to the 1980, most EDX detector were protected (from the water vapor and hydrocarbon in the microscope column) by a 10µm thickness beryllium window, which strongly absorbs photons of energy less than 1000eV and percludes analysis of elements of atomic number less than 11.
- With development of ultrathin (UTW) or atmosphericpressure (ATW), elements down to boron can be routinely detected, making EDX competitive with EELS for microanalysis of light elements in a TEM specimen.

EELS and x-Ray Signal Generation



X-ray fluorescence yield (log scale) as a function of atomic number



- The *EELS* is one step signal, while *EDX* is a two step signal (low x-ray fluorescence yield for low Z). In general, the yield rate of the EELS is higher than EDX.
- the signal of EELS concentrates in a small angle range of the transmitted beam, but the EDX signal spans around larger angle range.

(a) These two cause EELS has higher core loss (higher Signal to noise ratio, EELS has less recording time)

X-ray and EELS spectra

(b) EDX has better Signal/ background ratio



Background of EELS:

arises from the inelastic scattering from the atomic electron whose binding energy less than the edge energy

Background of EDX: arises from bremsstrahlung



EDX bulk beam broaden size for SEM system



EDX thin film (TEM) beam broaden size



(c)x-ray has larger interaction volume than that of electron. The ultimate spatial resolution is higher for EELS than for EDX , but the thin crystal is required for EELS





density

The EDX spatial resolution limit

Beam

Energy of Indicate electron beam The EDX spatial resolution (R) equation: $R = \frac{d + R_{max}}{2}$

Specimen 2a

For example:

Specimen thickness :100nm, b is about 10nm Nano EDX Beam size (d) :0.5 nm The resolution limit: *R*=5.25 nm

atomic weight

Follow previous slice, the beta angle for EDX is about 0.82 rad=820mrad

For EELS image mode, the beta angle is about 13.06 mrad

For example: The beam size is about 0.5nm; specimen thickness is about 100nm The EDX beam R_{max} is 5.256nm EELS R_{max} is about 0.6nm

de-localization of energy loss electron



- (d) EELS is an absolute, standardless quantitation technique, but quantification error may exist in the case of crystal.
- (e) Structural information is available, but more operator intensive is required.

(f) Comparison of EELS and XEDS sensitivity (depends on strongly on SNR, but not SBR)(Leapman et al.)

EELS is capable of detecting smaller concentrations of elements of low atomic number



(g) EDX resolution is 50 - 100 eV so there is peak overlap below 1000eV

EELS resolution is ~ 1 eV so edge overlap can be less

> (stainless steel, Zaluzec 1984)



EELS vs. EDXS

EDXS

- X-rays provide elemental information only
- Inefficient signal generation, collection & detection inefficient x-ray mapping
- Slow technique (hours)
- X-ray spectra can contain information from column and other parts of sample
- High detection efficiency for high Z elements
- Energy resolution > 100eV causes frequent overlaps
- Only simple processing required

<u>EELS</u>

- Elemental, Chemical, & Dielectric information
- Very efficient in every respect => higher sensitivity to most elements very efficient mapping technique
- Fast technique (seconds to minutes)
- EELS spectra have no such artifacts
- High detection efficiency for low Z elements
- Energy resolution 0.3-2eV gives far fewer overlaps
- More complex processing required => Needs more hardware & software automation



What is EFTEM

- Energy Filtering Transmission Electron Microscopy



The features of EFTEM

A contrast-enhancement technique:

- it improves contrast in images and diffraction patterns by removing inelastically scattered electrons that produce background "fog".
 A mapping technique:
- it creates elemental maps by forming images with inelastically scattered electrons.

An analytical technique:

 it records and quantifies electron energy-loss spectra to provide precise chemical analysis of TEM samples.

The standard EFTEM elemental mapping images





•銅製程與低介電常數材料已是半導體元件發展趨勢 0.13μm 乃至於 90nm 的製程技術

NTHU

如何在具有圖案結構上之試片決定介電材質之介電常數。





傳統之介電常數量測





影像能譜(ESI)之擷取 Dielectric Function Imaging



Extract image spectrum

Ε.

energy loss





Dielectric Function image







介電常數分佈影像圖(dielectric constant map)



Materials	$\boldsymbol{\epsilon}_{\mathrm{ref}}$	E _{exp}
SiO_2	3.8	4.20 ± 0.31
Si ₃ N ₄	3.6	3.72 ± 0.30
Black Diamond TM	2.5~2.8	2.69±0.27

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Band gap energy Imaging

N a

7.5

5.0

Energy loss (eV)

(a)

(direct band gap)

10.0



2.5

0.0

 $J^{1}(E)$: a = 0.5

(in-direct band gap)

5.0

Energy loss (eV)

(b)

2.5

a=1.5

7.5

10.0

0.0

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AIN/GaN Quantum Well





Band Gap Map

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The map of band energy of AIN/GaN layers is obtained using electron spectroscopy imaging (ESI) technique. The average band-gap energy of AIN and GaN is determined to be about 5.62 \pm 0.35 eV and 3.47 \pm 0.36 eV, respectively.



Tsai, Kai, Chen, L. Chang, Journal of Electron Microscopy (2003)



Yan, Kai, Chen, L. Chang, JEM (2002)



Structure of the specimen



Znoe axis(10-10)



Image to Spectrum





Valence State Mapping





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