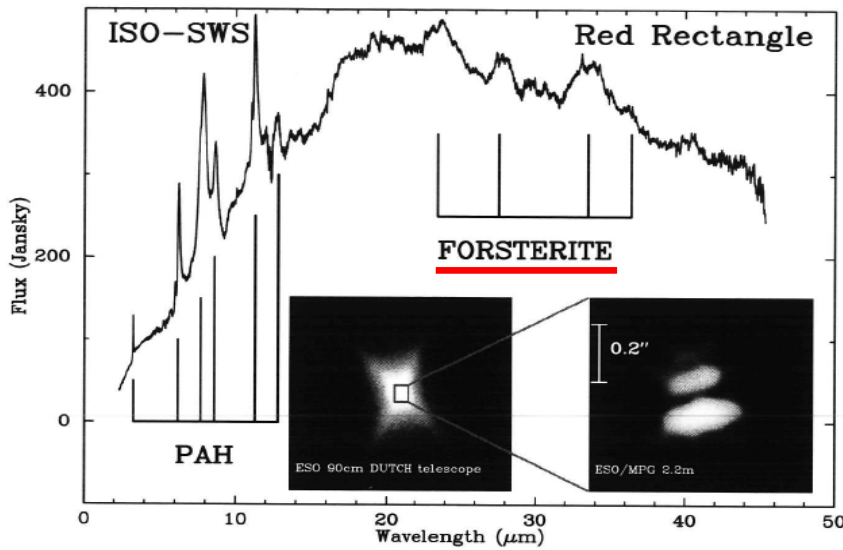


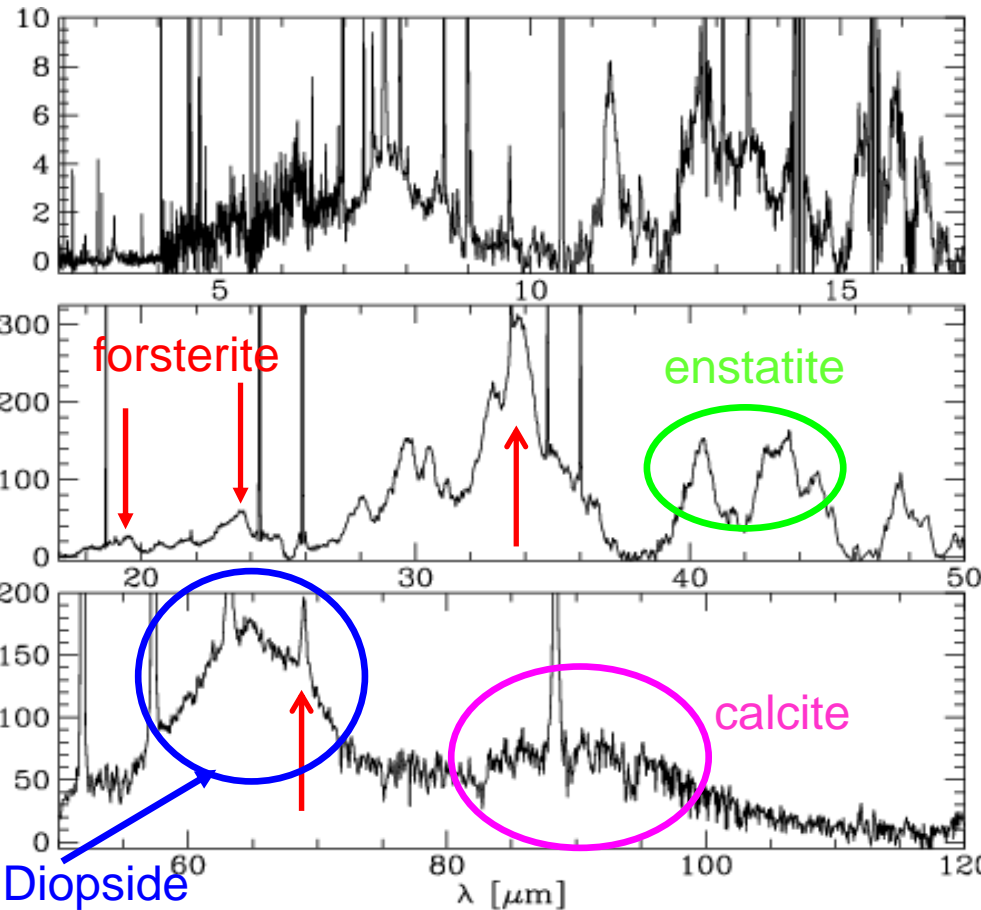
Shape effects of forsterite particles on infrared spectra

C. Koike, Y. Imai, R. Noguchi, H. Chihara,
A. Kumamoto*, C. Kaito*, H. Suto**, A. Tsuchiyama
Osaka University, *Ritsumeikan University,
**Subaru Telescope, NAOJ



Waters et al.
(1998, Nature 391)

ISO observation
In circumstellar
NGC 6302



Molster et al. 2001

Many sharp peaks were
detected



Compare with laboratory data
(absorption of particles,
reflectance)



Identified as minerals
forsterite, enstatite, diopside,
melilite, calcite, ,,,,,,

Infrared Absorption of dust (fine particles) depend on

size

composition (olivine, pyroxene,,,,)

shape (Fo)

coagulation (aggregate)

temperature (cooling at RT, 200K, 100K, 50K, 20K, 8K)

crystallinity (degree of crystallinity)

medium (KBr, PE)

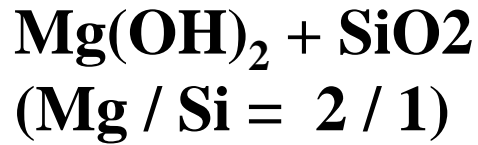
we investigate about forsterite particles

on the shape and coagulation

(Koike et al. 2010, Koike et al. in preparation)

forsterite : Mg_2SiO_4

Sample Preparation for Forsterite Mg_2SiO_4 particles



↓ **evaporation & condensation**

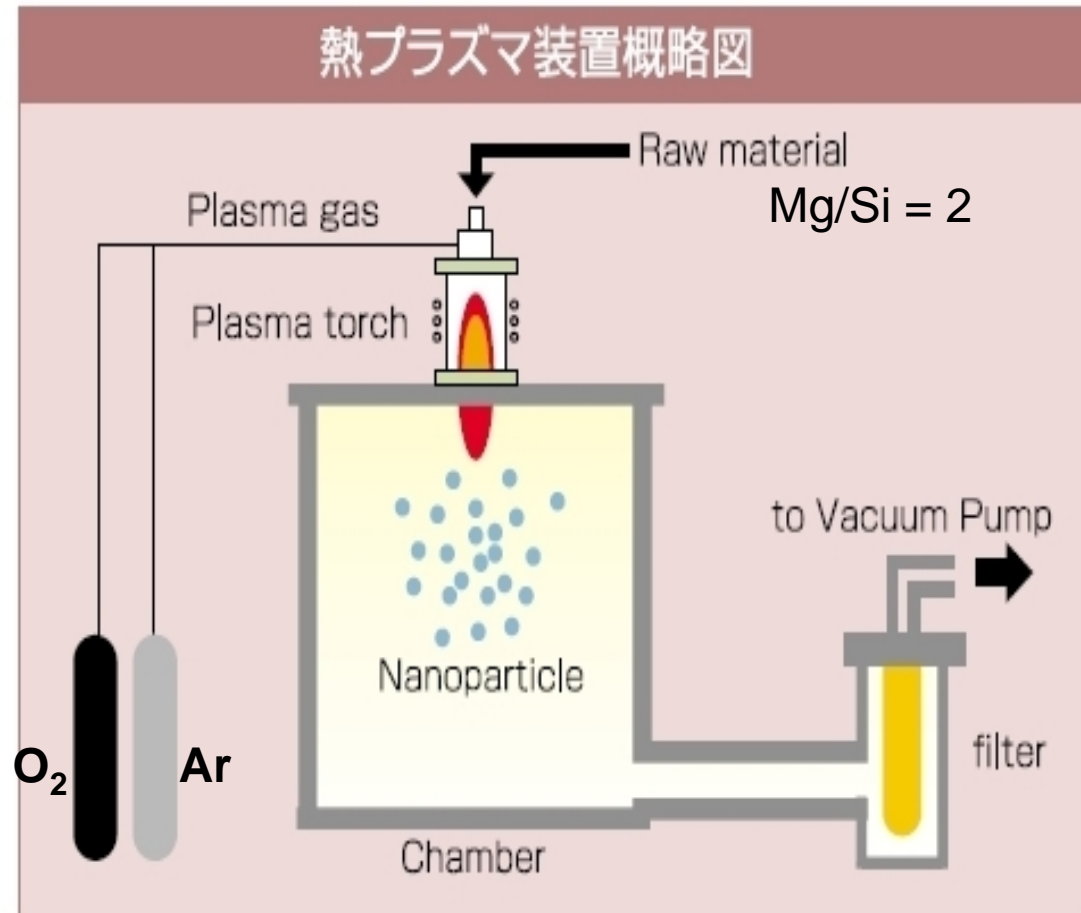
spherical amorphous particles

↓ **annealing and crystallization**

spherical, and irregular forsterite particles

commercial products

in plasma method



Infrared spectra of annealed forsterite particles

Starting products

original amorphous of Nissin products (by XRD)

NiA average size 11 nm : coagulated

NiB average size 80 nm : isolated

↓ annealing at various temperature

shape — depend on annealed temperature

NiA spherical and coagulated → irregular shape

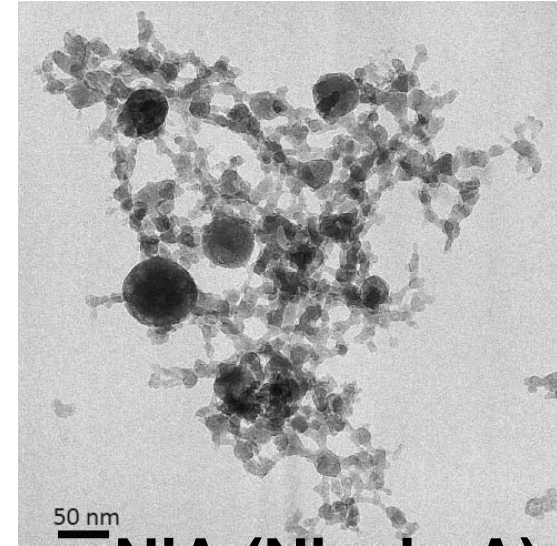
NiB spherical → irregular shape

how to change infrared spectra

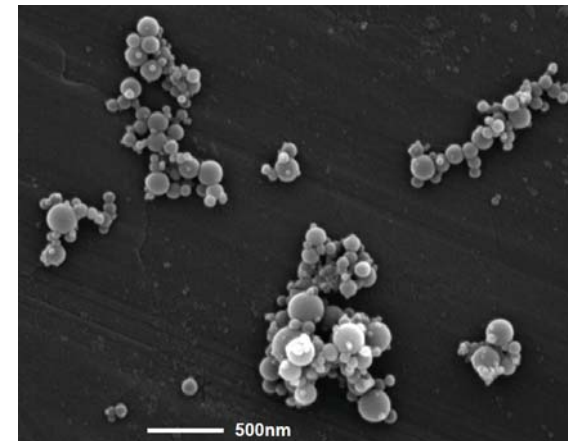
(KBr & PE) (in wide wavelength region)

○ 11 μ m, 33 μ m band

○ 69 μ m band



NiA (Nissin A)



NiB (Nissin B)

Condition for annealing of amorphous products (NiA & NiB)

Low temperature
≤ 1200 °C

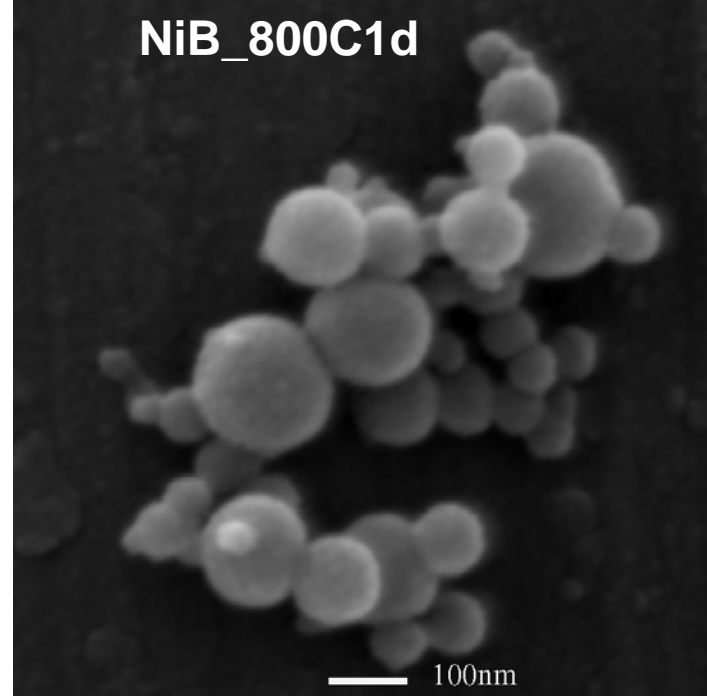
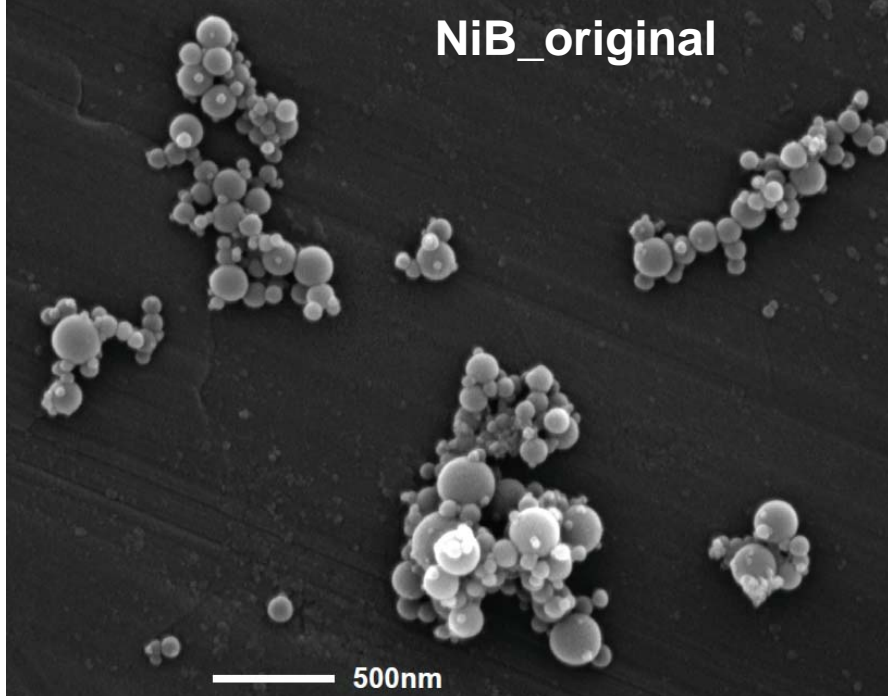
NiA	
	550°C 600°C 650°C 700°C 800°C 1000°C 1100°C 1150°C 1200°C
3h	
24h	○ ○ ○ ○ ○ ○ ○ ○
48h	○

NiB	
	600°C 650°C 700°C 800°C 1100°C 1130°C 1150°C 1200°C
6h	○ ○
24h	○ ○ ○ ○ ○ ○ ○

High temperature
≥ 1200 °C

Annealing temperature		Annealing time		protoenstatite (by XRD)
NiA	1200 °C	3	h	
	1300	3, 16, 18, 60, 96		16 – 96 h
	1400	3		3 h
	1500	2, 3, 4, 10, 14.3, 45.5		2 – 45.5 h
	1600	3		
NiB	1500	3		
	1600	3		

Red: IR measured



**Single
crystal**

NiB_SEM

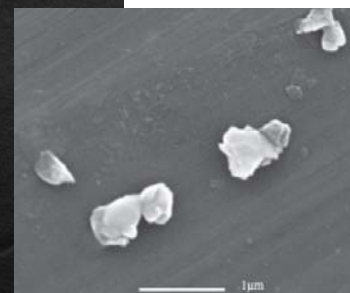
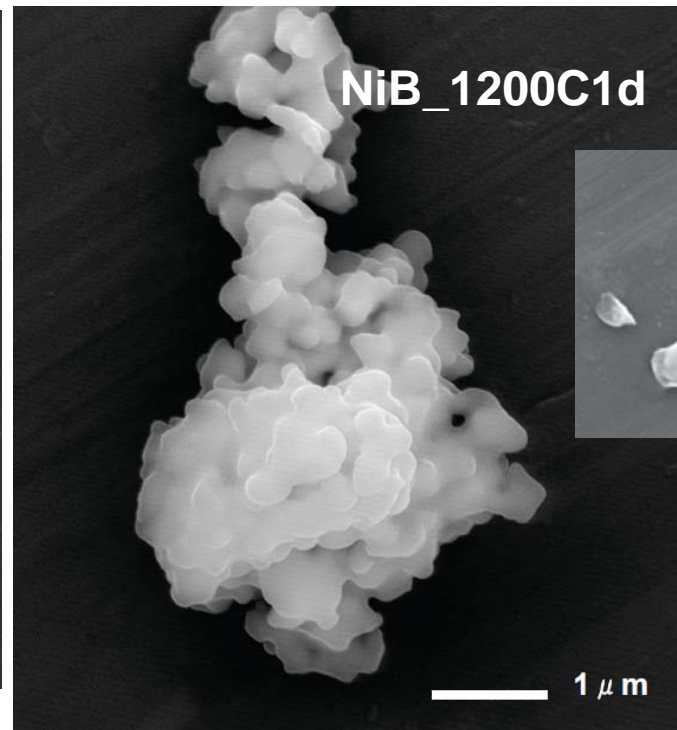
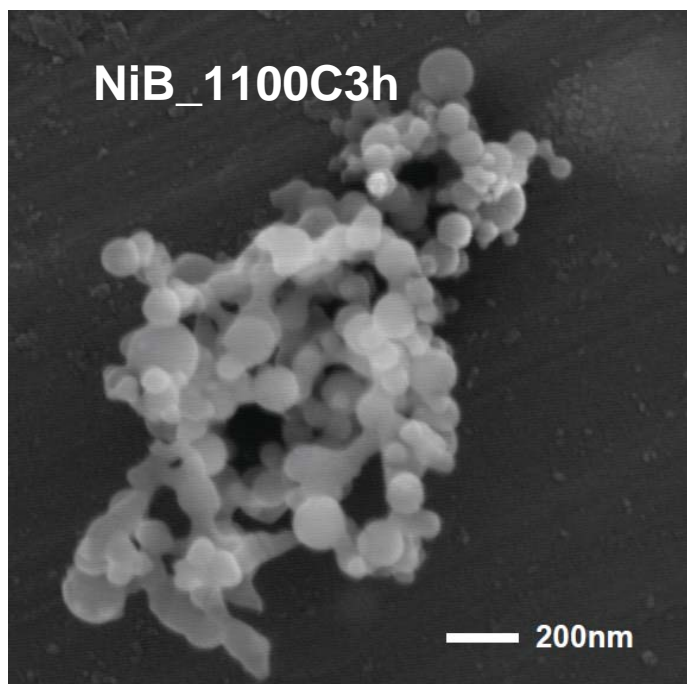
**Spherical
& isolate**



Irregular



**Plate-like
including**



NiA_TEM

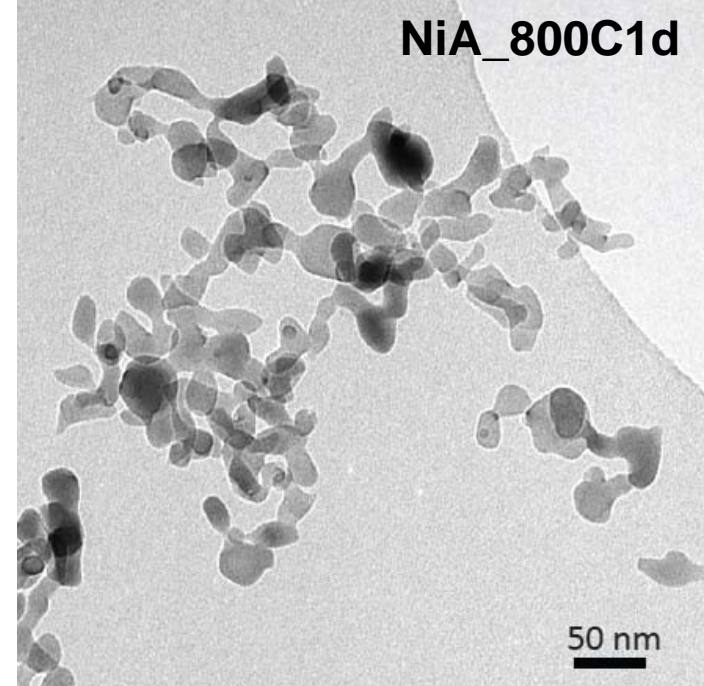
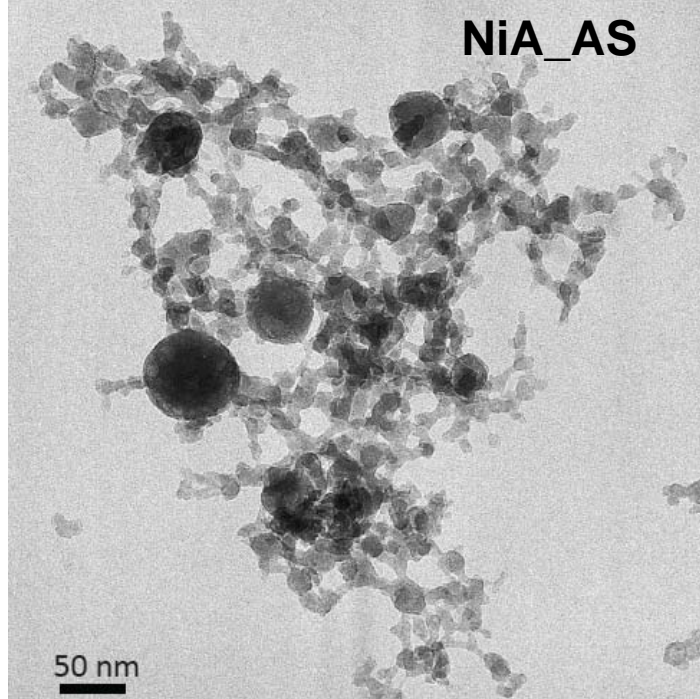
**Spherical &
Coagulation**



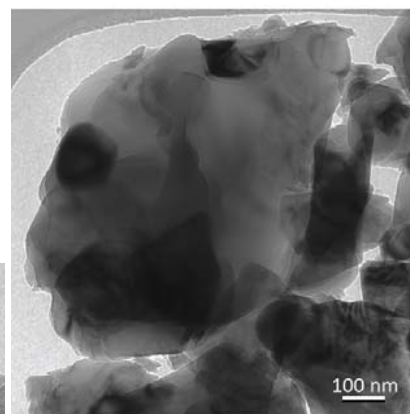
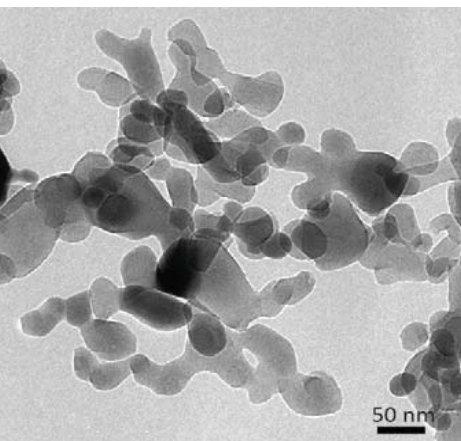
Irregular



**Plate-like
including**

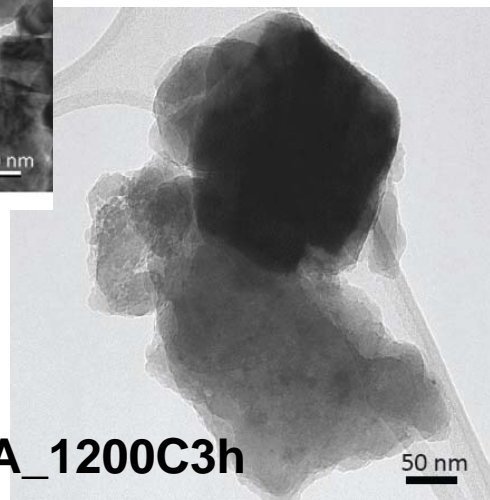


NiA_1000C1d



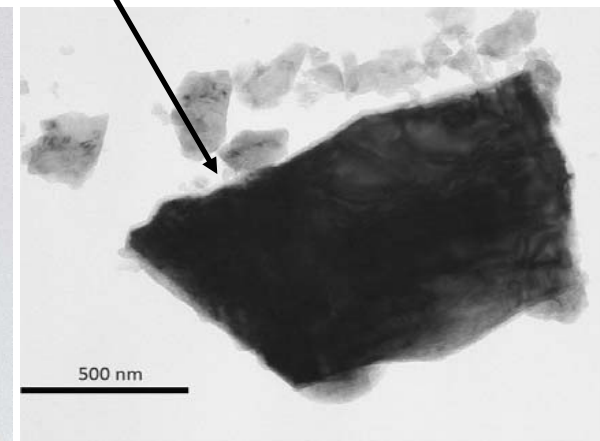
NiA_1150C1d

NiA_1200C3h

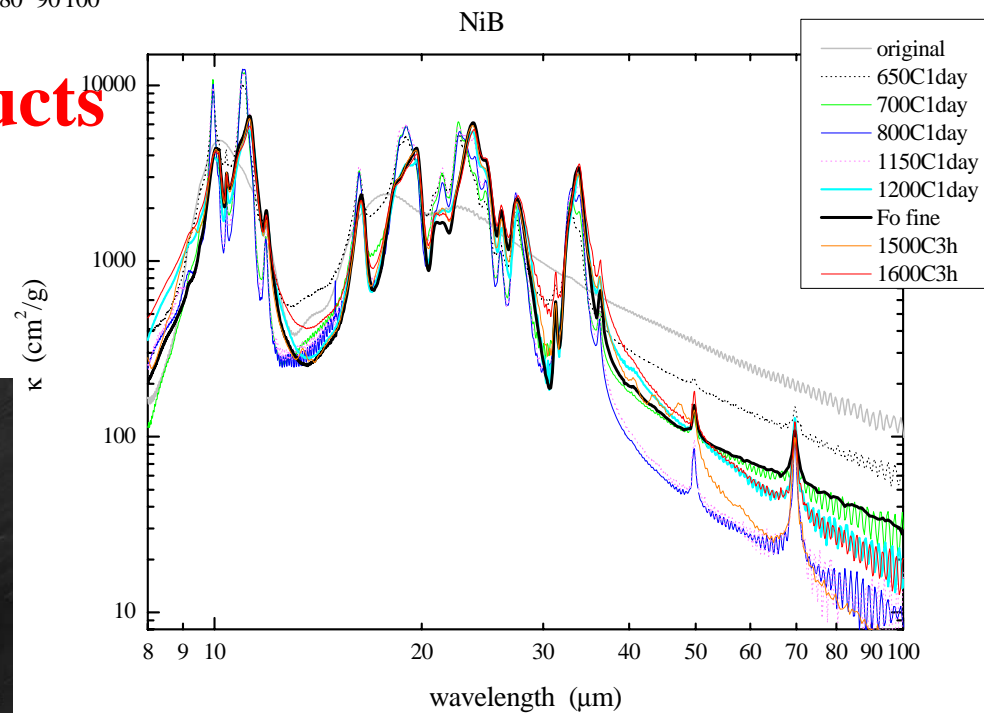
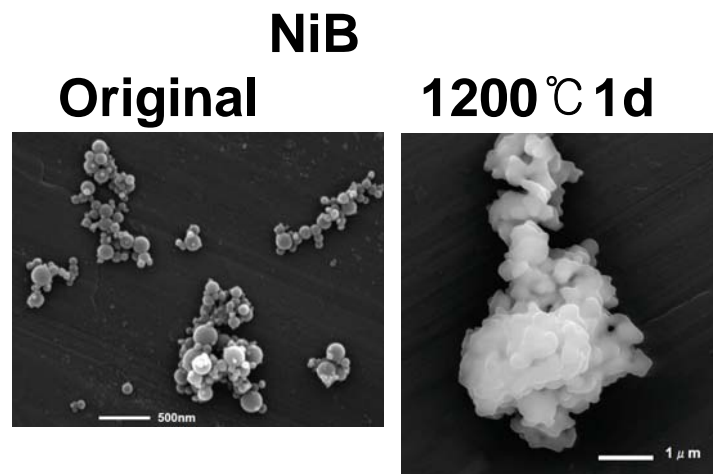
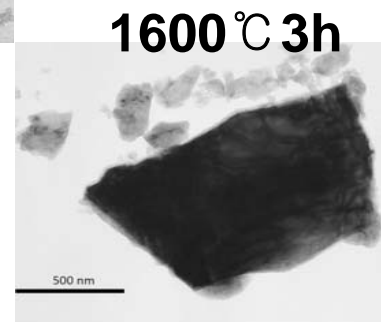
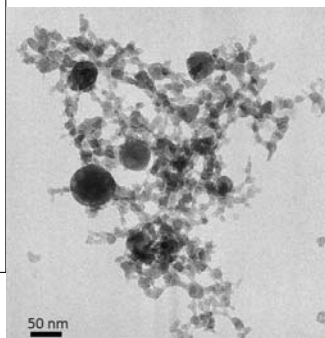
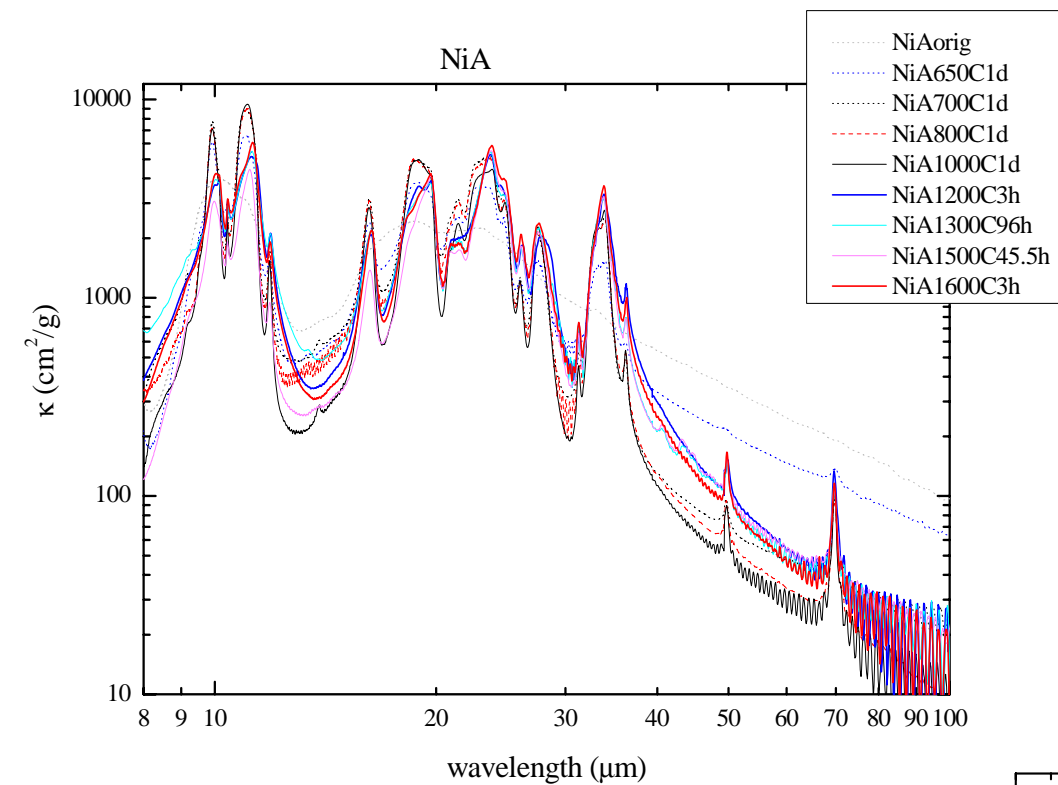


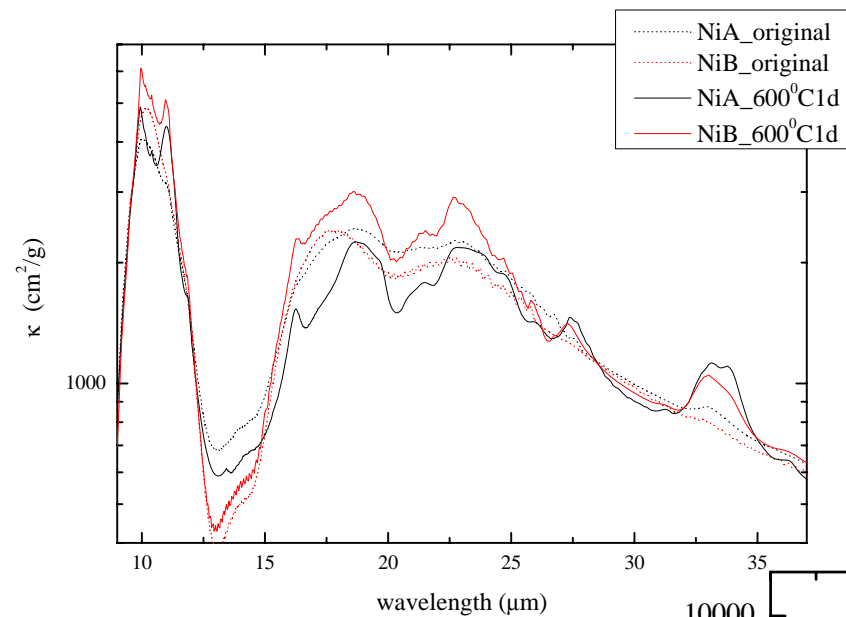
Single crystal

NiA_1600C3h



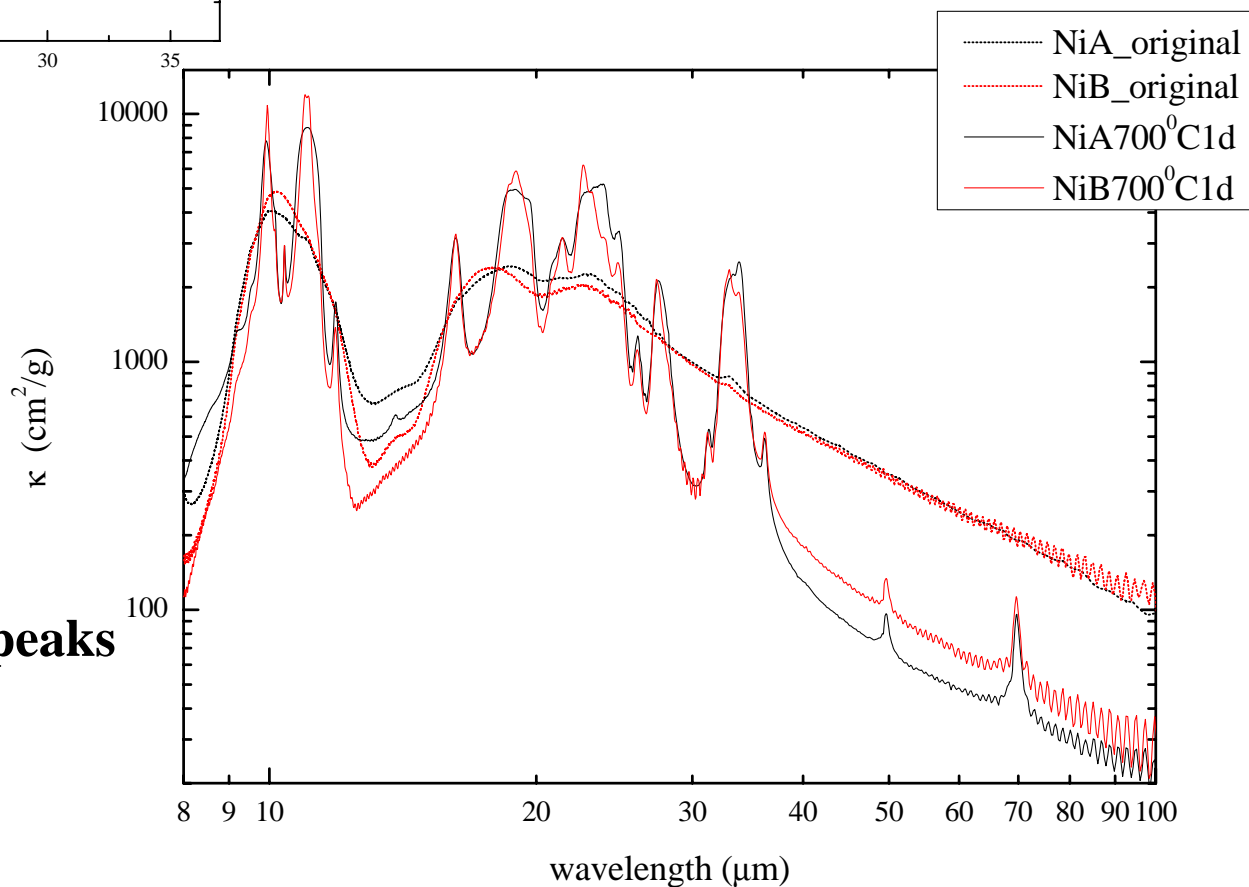
**All spectra of two Ni products
For annealed samples**

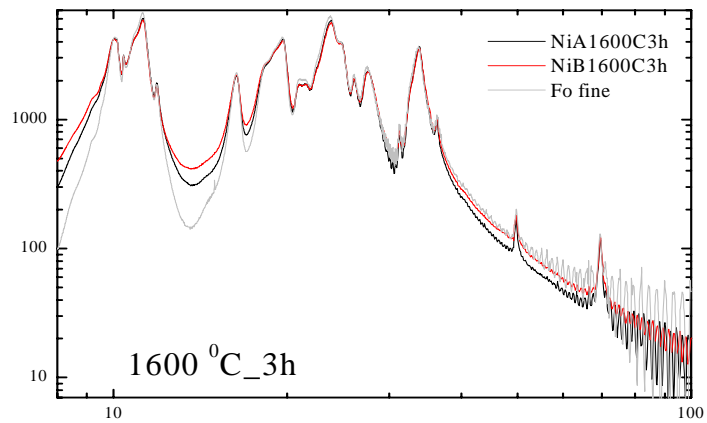
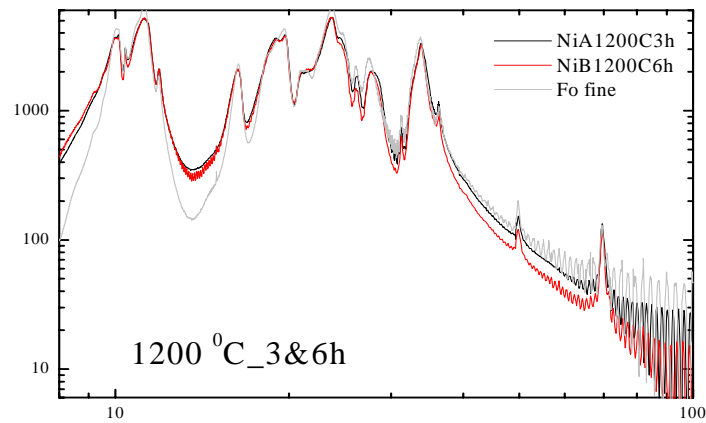
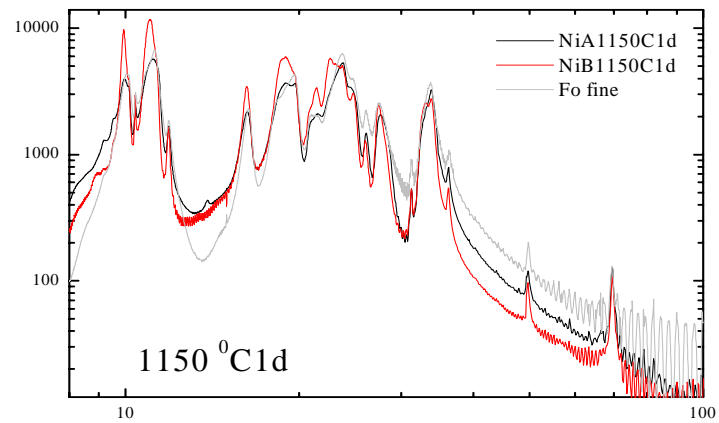
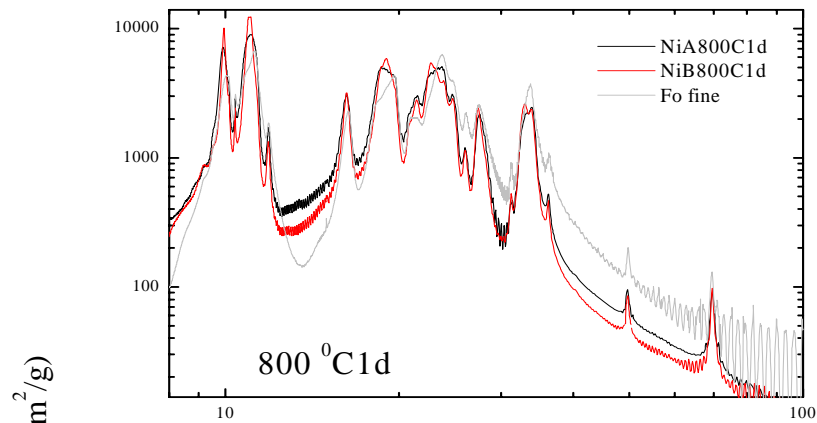




Annealing at 600 °C
Some broad peaks appear at each band
and depend on original products

Annealing at 700 °C
All spectra show sharp peaks
Spectra are depend on
original products





wavelength (μm)

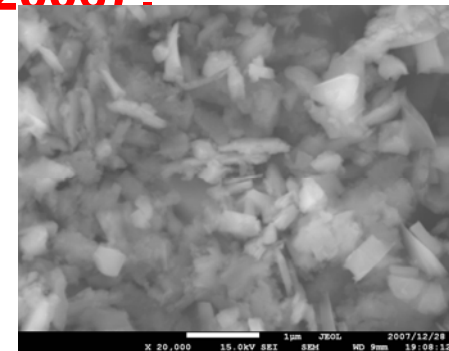
Annealing temperature ≥ 1200 °C

All spectra became similar to that of Fo fine (Koike et al.2006).

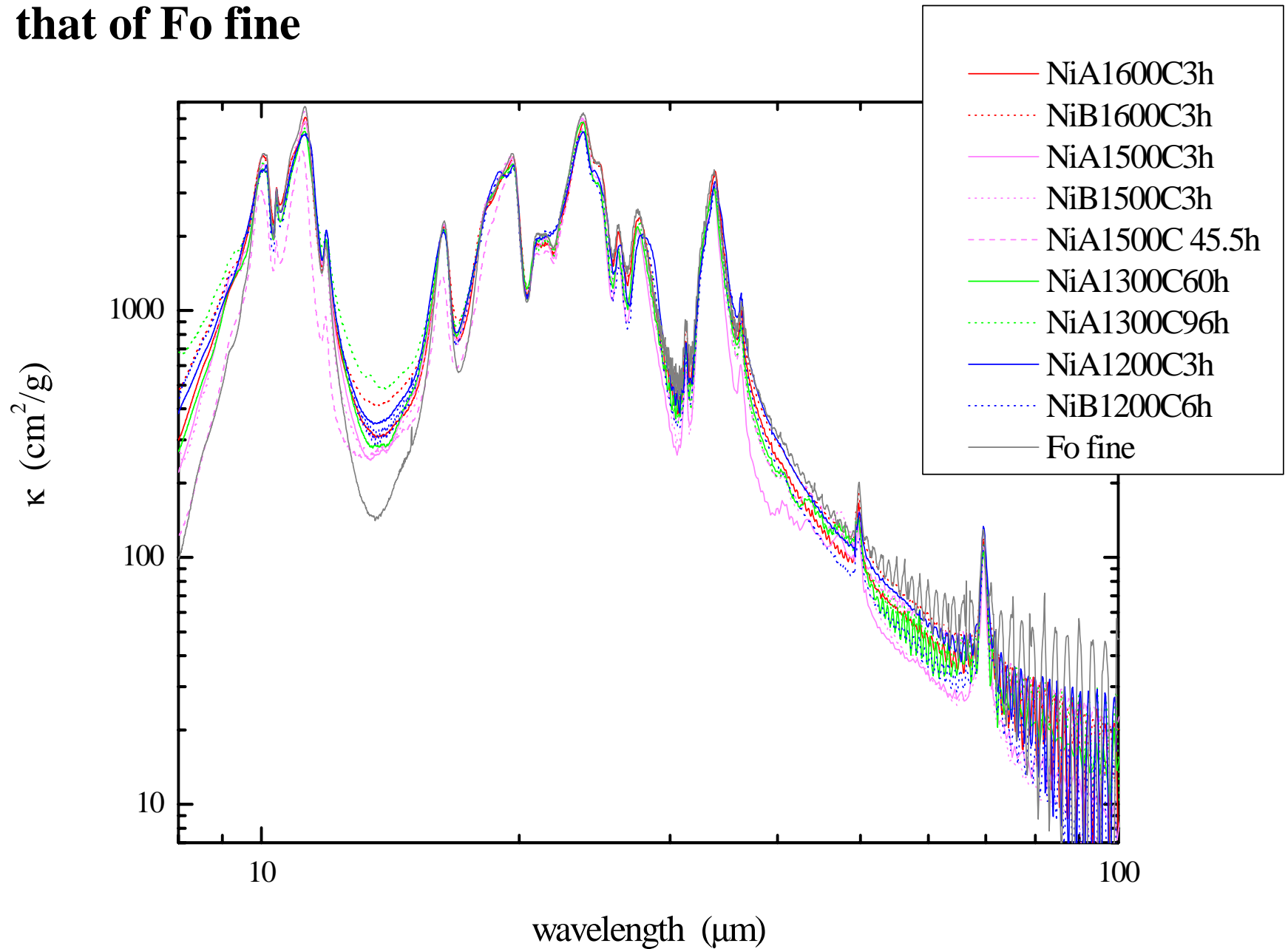
Fo fine

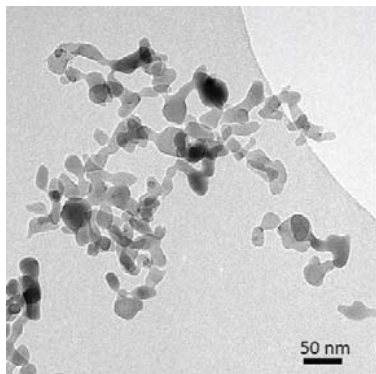
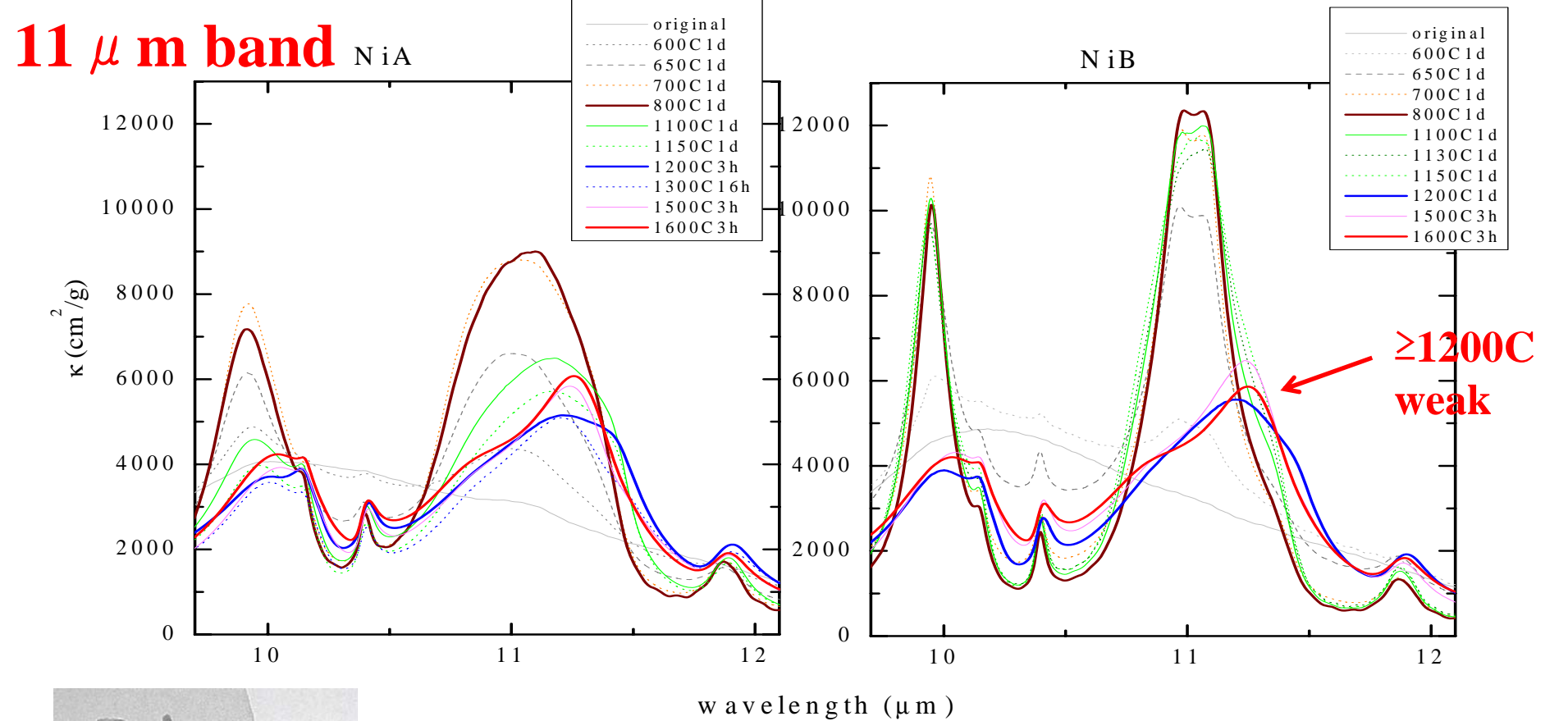
Fo fine

(ground from bulk Fo synthesized by
Czochralski method)
(irregular shape)

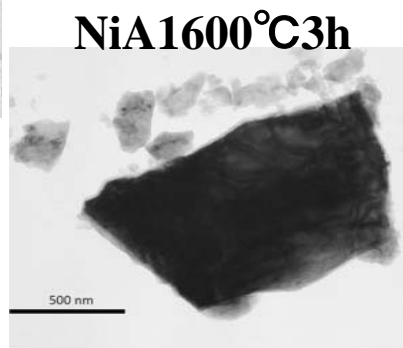


**Both of products have similar spectra for annealing at above 1200 °C
as that of Fo fine**

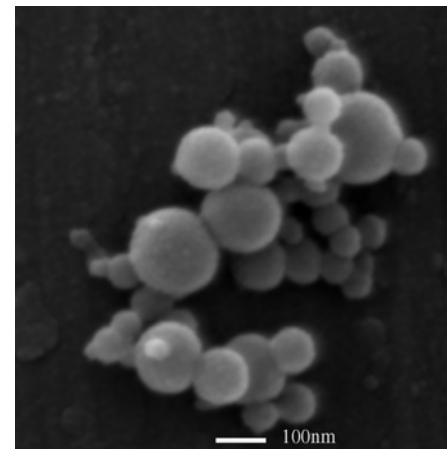




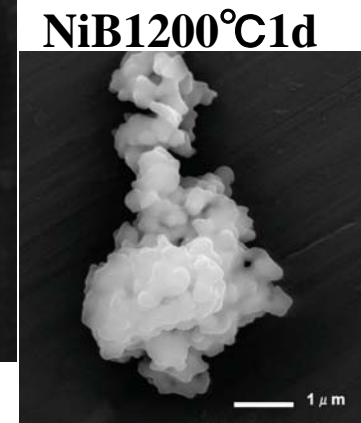
NiA800°C1d



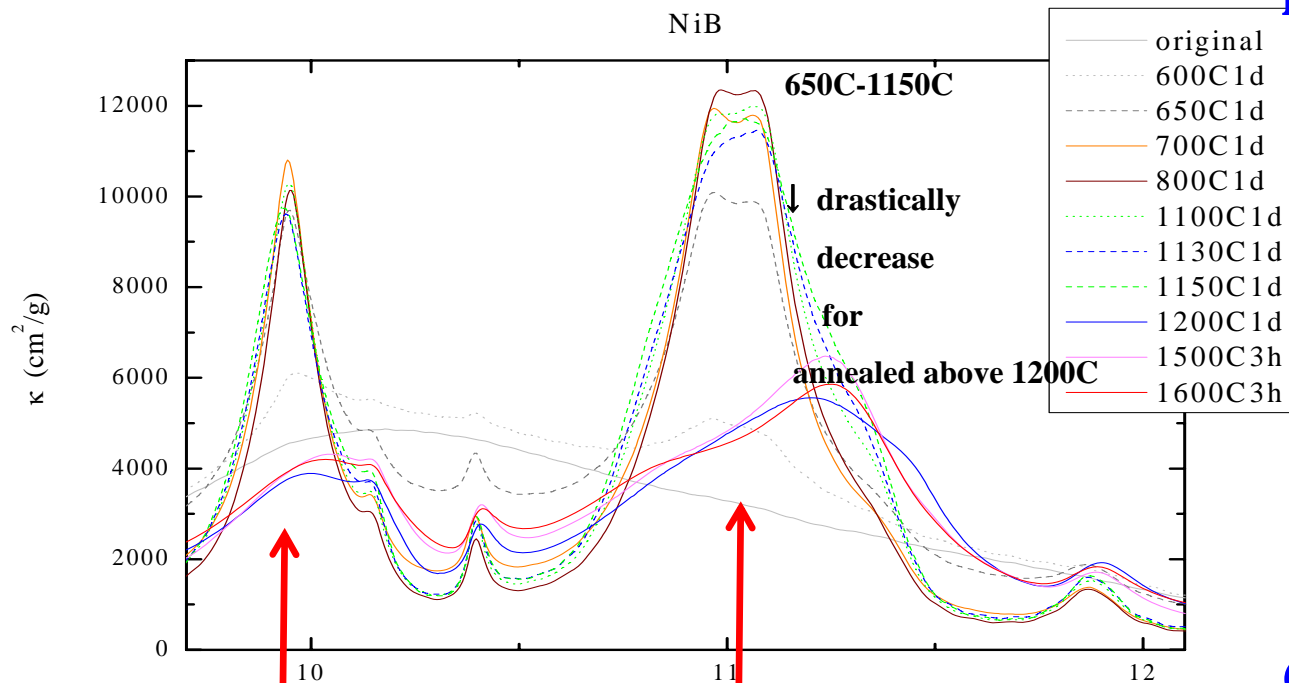
NiA1600°C3h



NiB800°C1d



NiB1200°C1d

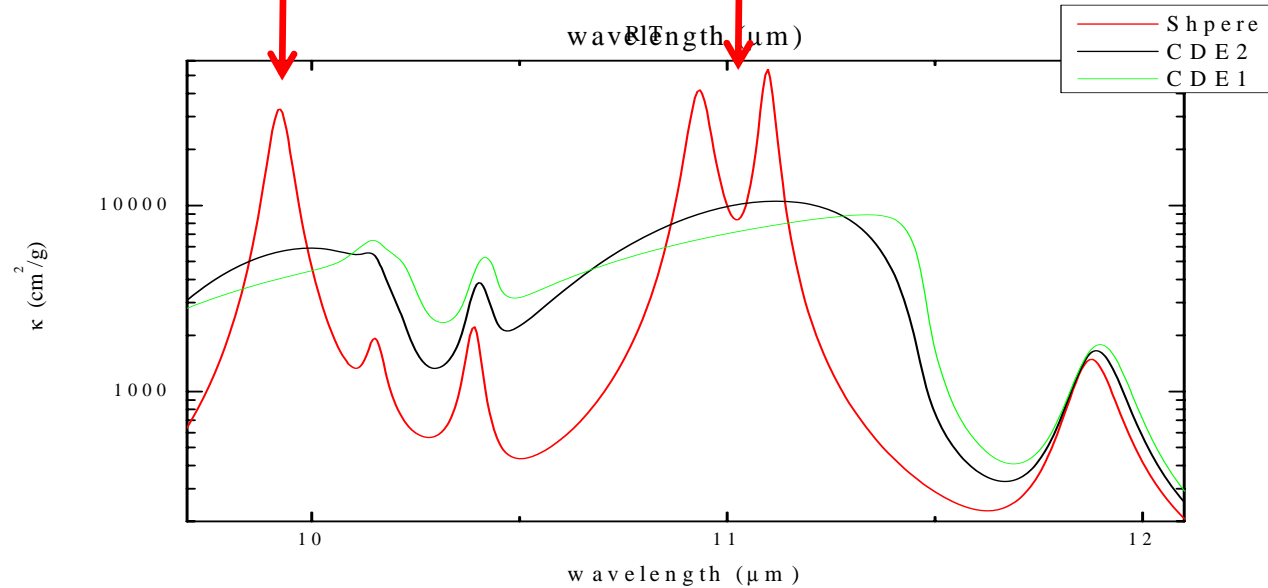


labo data

**9.95 μ m and 11 μ m
peak intensity
strong \rightarrow weak**

**spherical &
coagulate**

irregular (≥ 1200 C)



Calculation (in KBr)

spherical

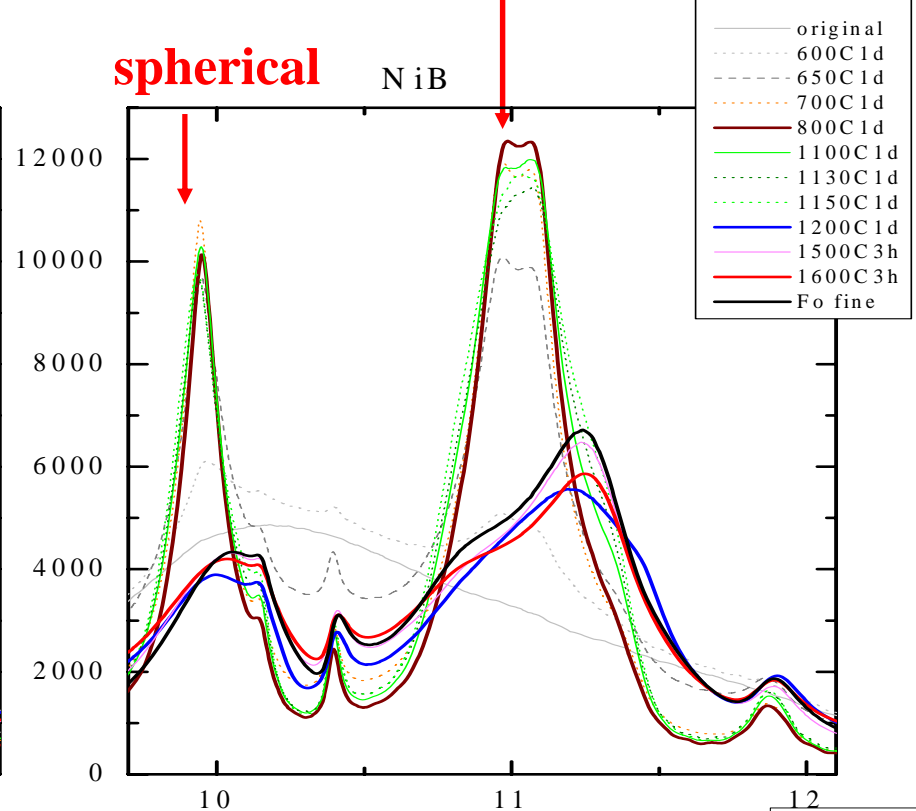
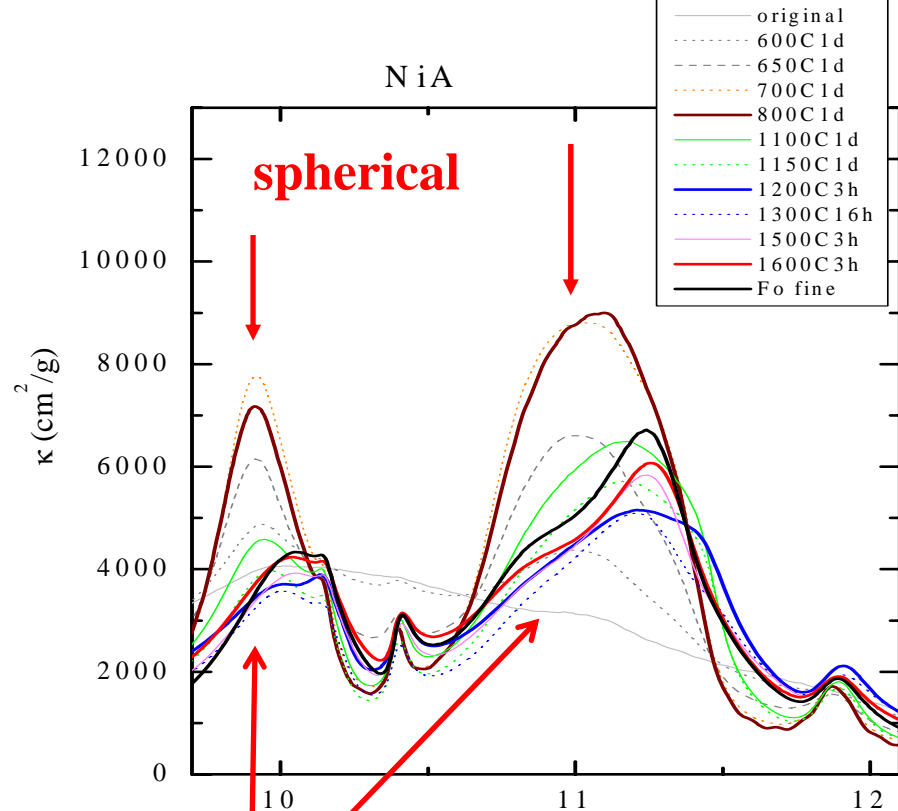
**9.95 μ m
sharp peak**

**11 μ m peak
double peak**

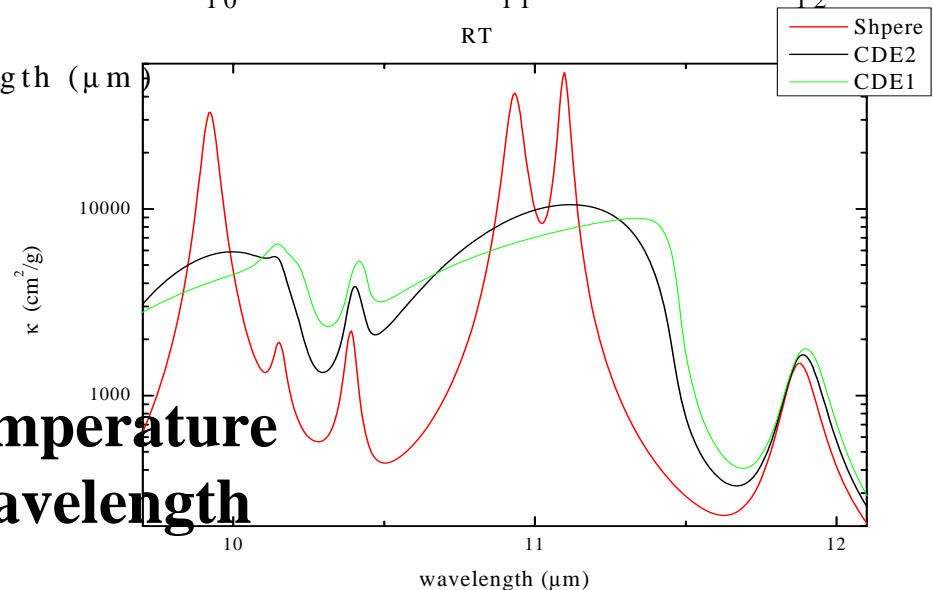
**irregular
broad,
peak position shift**

CDE1: Continuous Distributions of Ellipsoids

(CDE2: near-spherical particle shapes)



wavelength (μm)



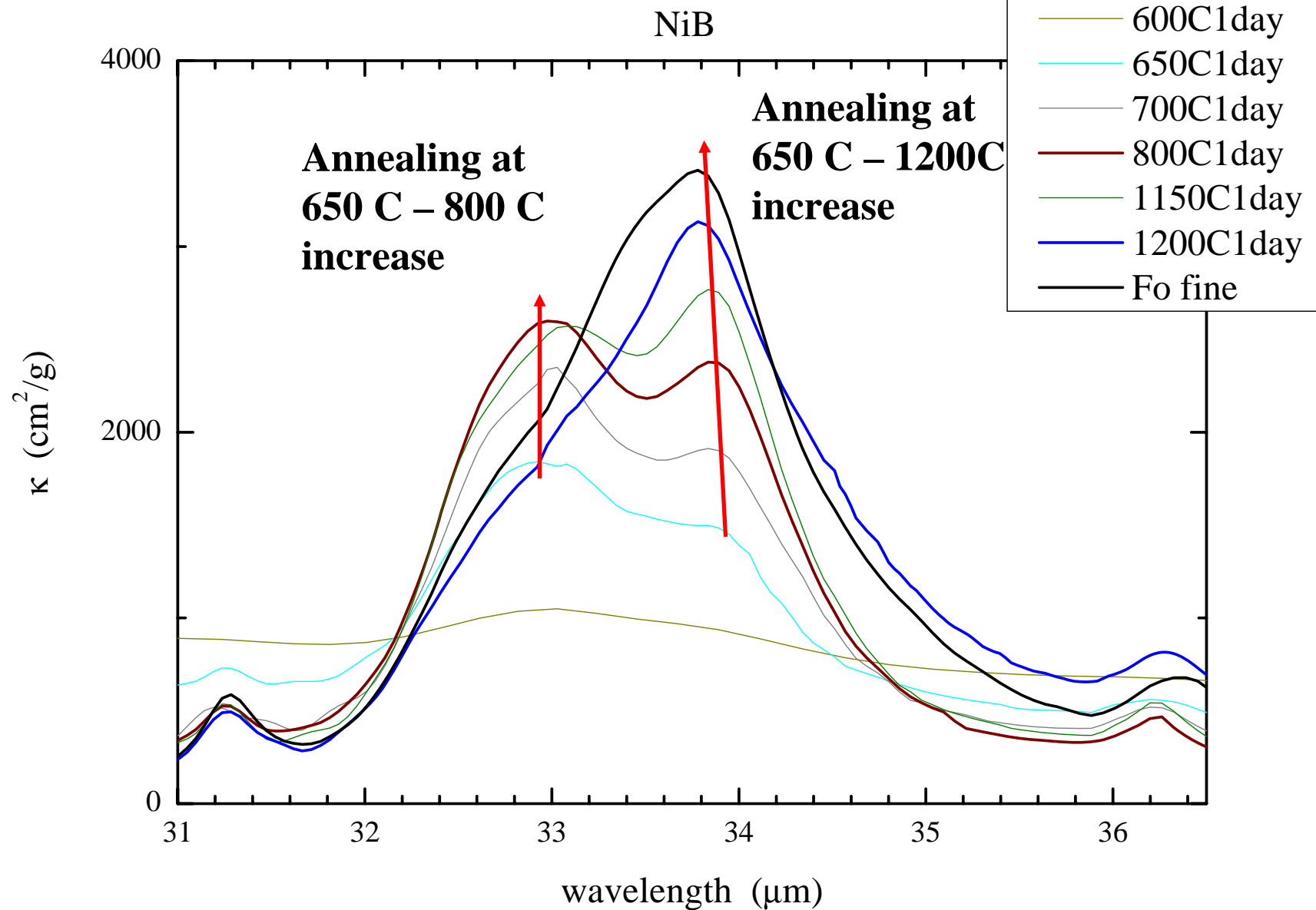
NiA compare to NiB

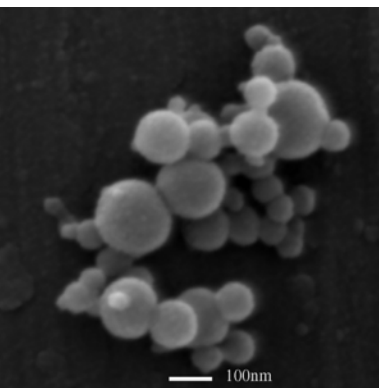
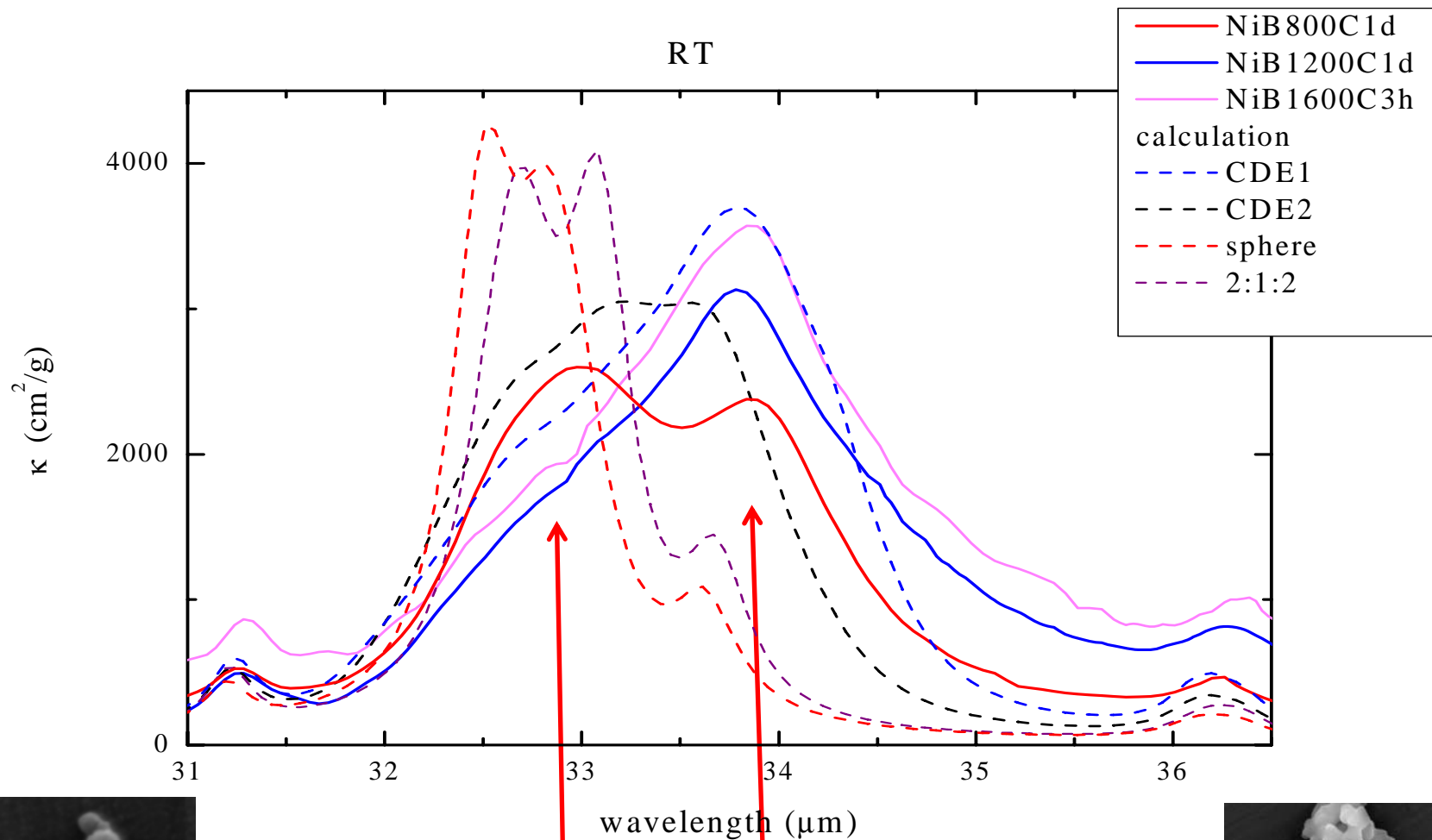
9.95 μm & 11 μm

broad due to coagulation

**NiA & NiB for high annealing temperature
peak position shifts to long wavelength
due to irregular shape**

33 μ m band

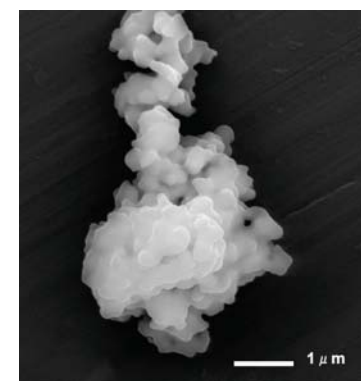




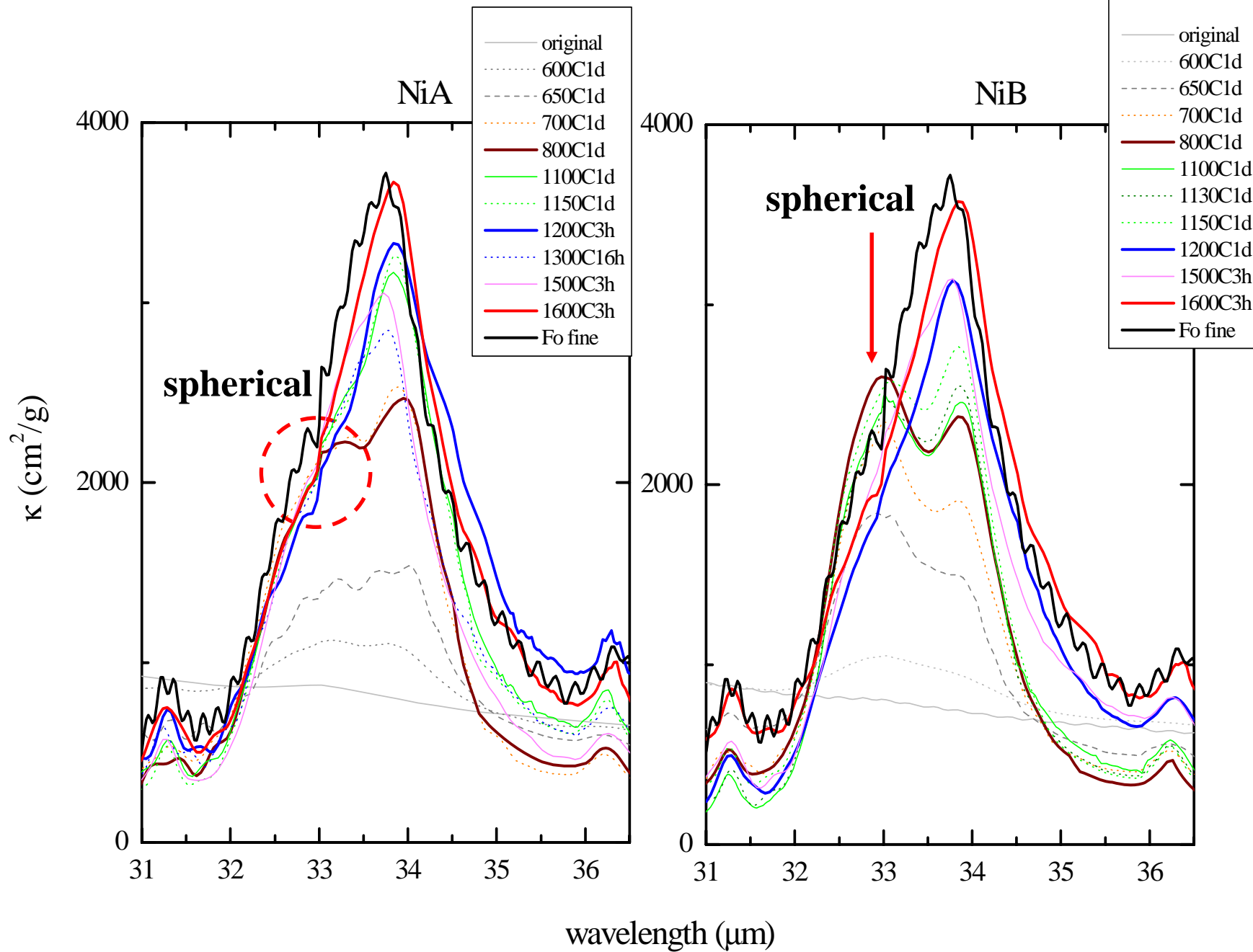
NiB800°C1d

**spherical
& coagulation**

CDE (irregular shape)

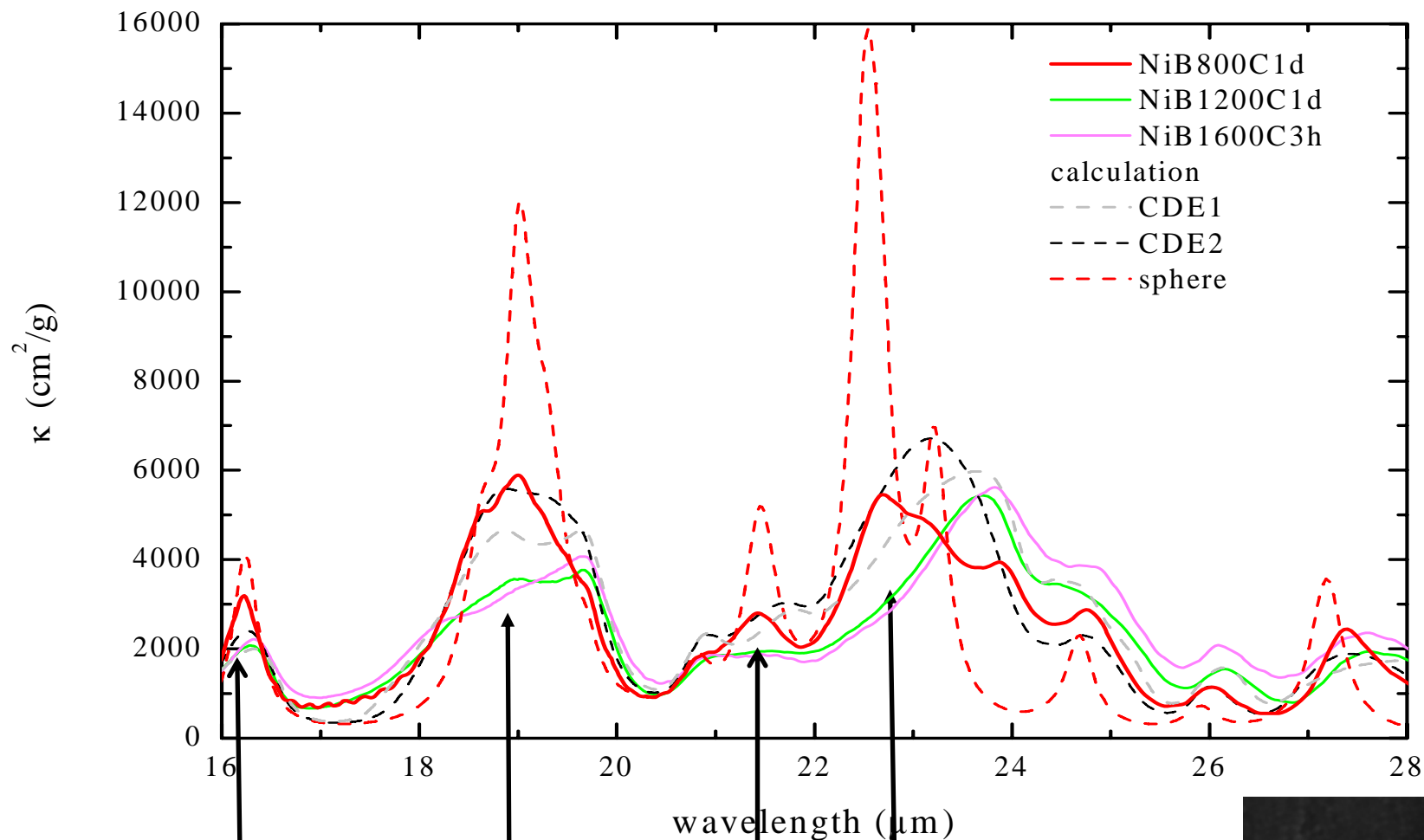


NiB1200°C1d



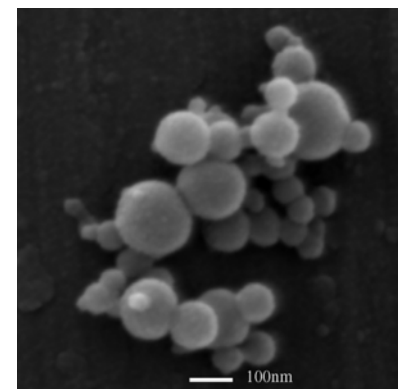
16 – 28 μ m region

RT

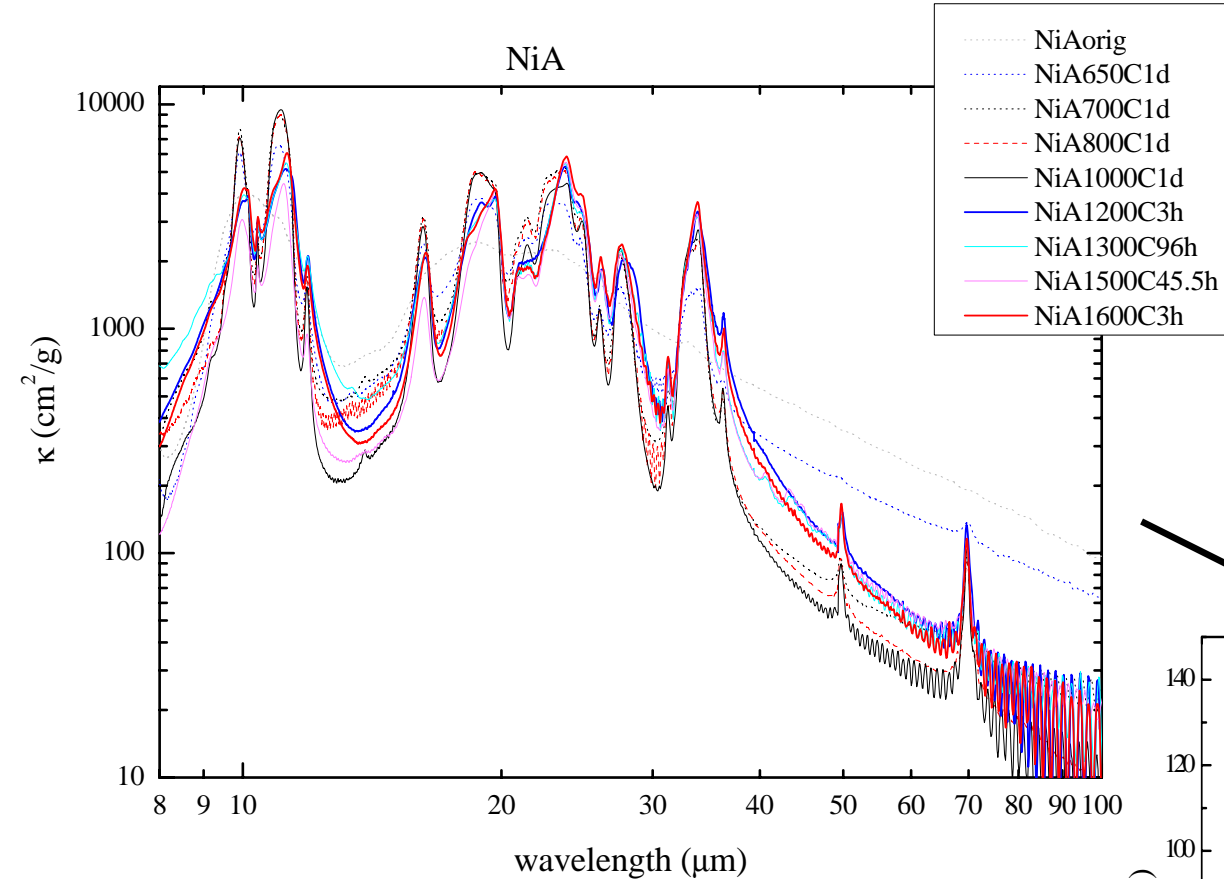


Spherical & coagulation

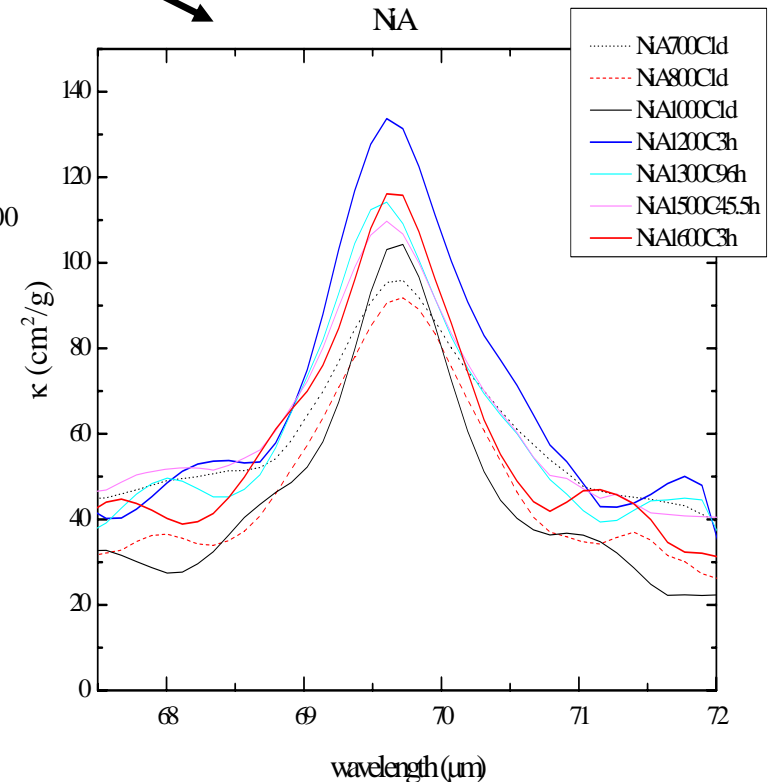
NiB800°C1d



69um band



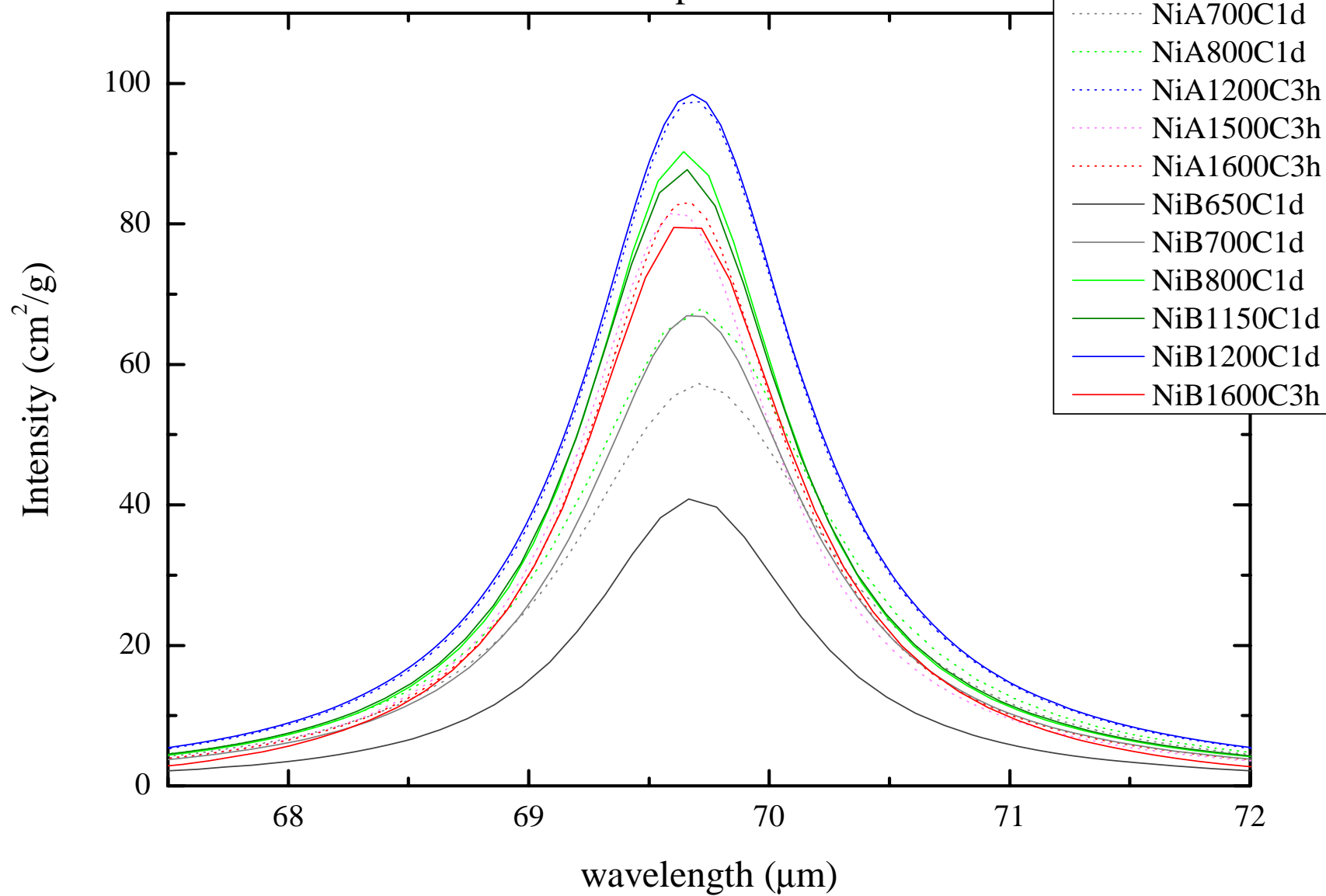
69 μm band



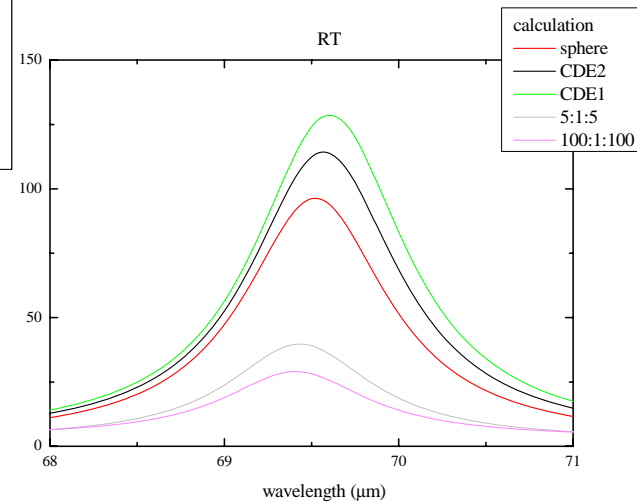
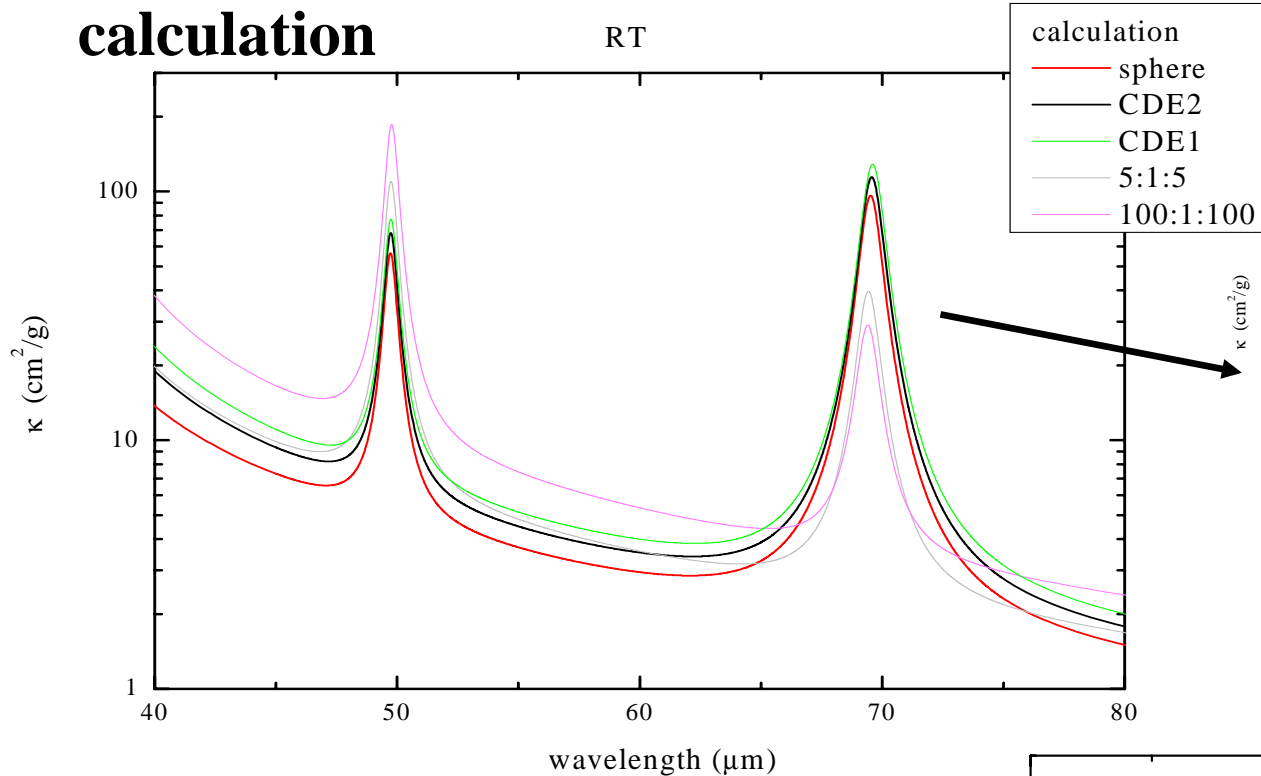
How to change intensity, peak position, and FWHM of 69 um band

These may depend on shape of particles ?

NiA&B peakfit RT



calculation



Calculation (in PE)

Peak intensity

sphere

strong

CDE1, CDE2

elliptical

weak

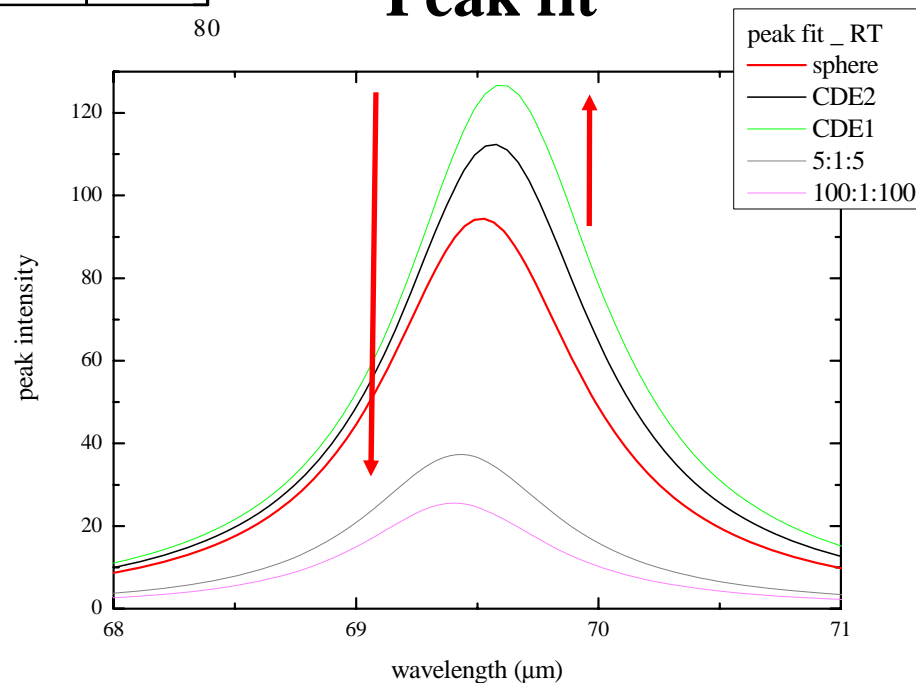
Annealing

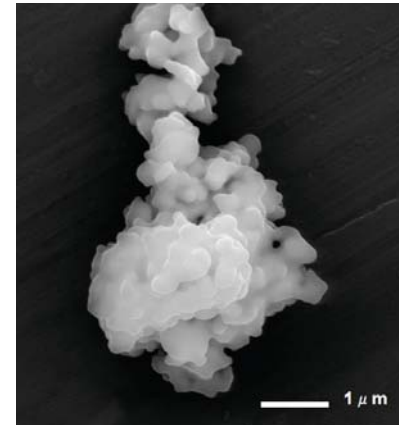
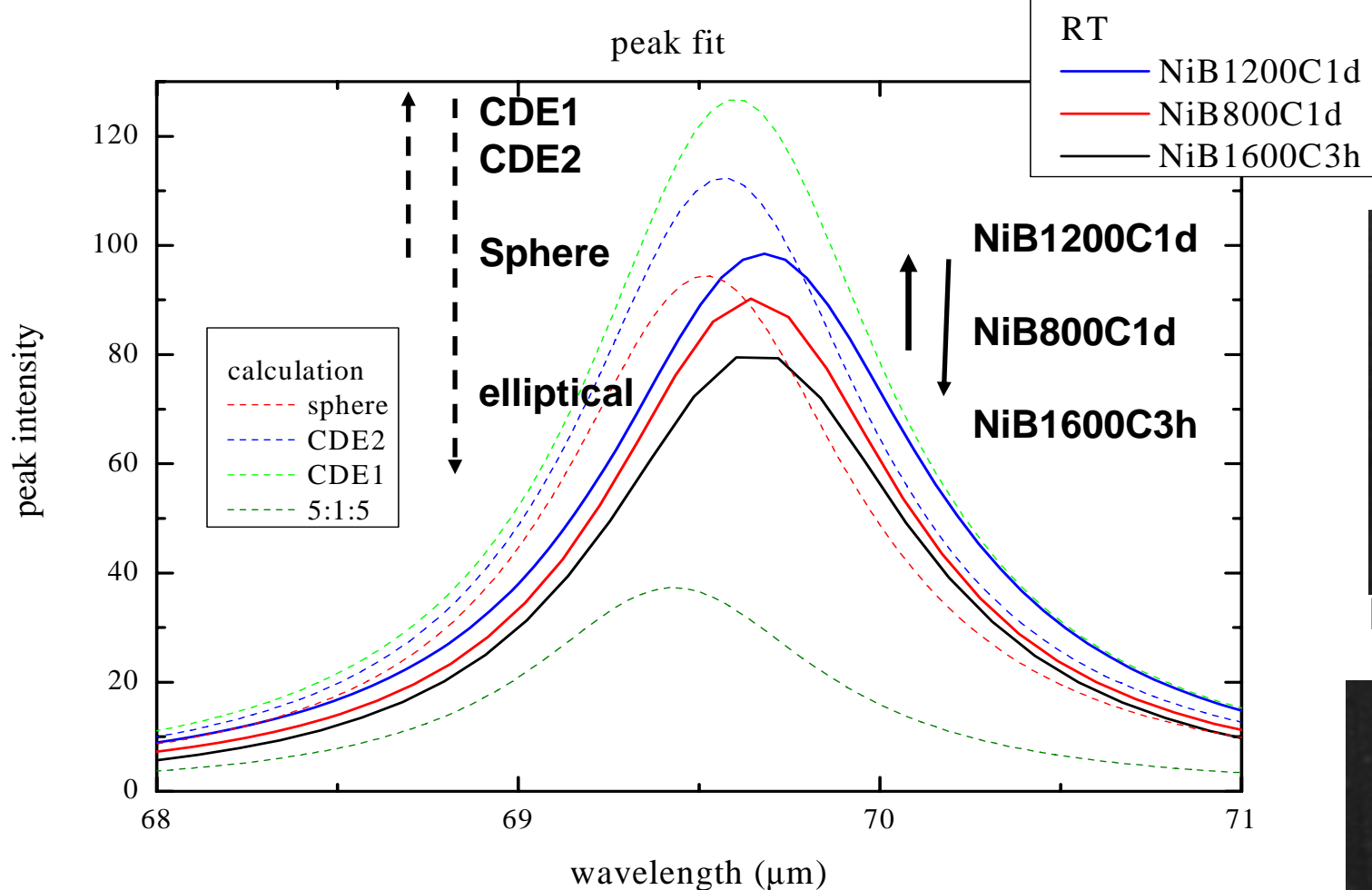
sphere

irregular

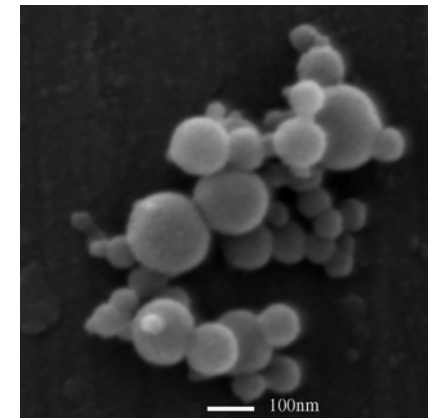
Plate-like
including

Peak fit





NiB1200C1d



NiB800C1d

Difference of Peak intensity
due to shape effects & coagulation
 (irregular & only small part of plate-like)

Difference of peak position among labo data and calculation
 may be due to refractive index of PE ?

LDPE $n = 1.46$ (here assume 1.50) , HDPE $n = ?$

Peak intensity of 69 μm band

peak intensity

strong

weak

annealing temperature

650 $^{\circ}\text{C}$

→

1200 $^{\circ}\text{C}$

→

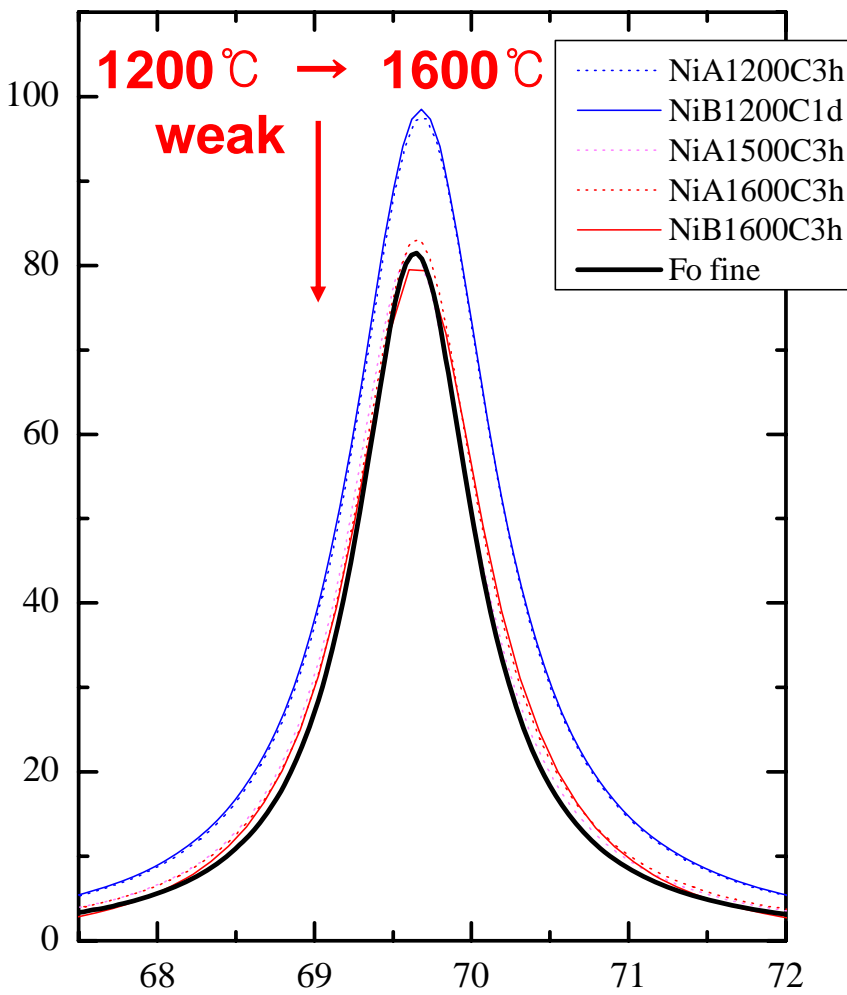
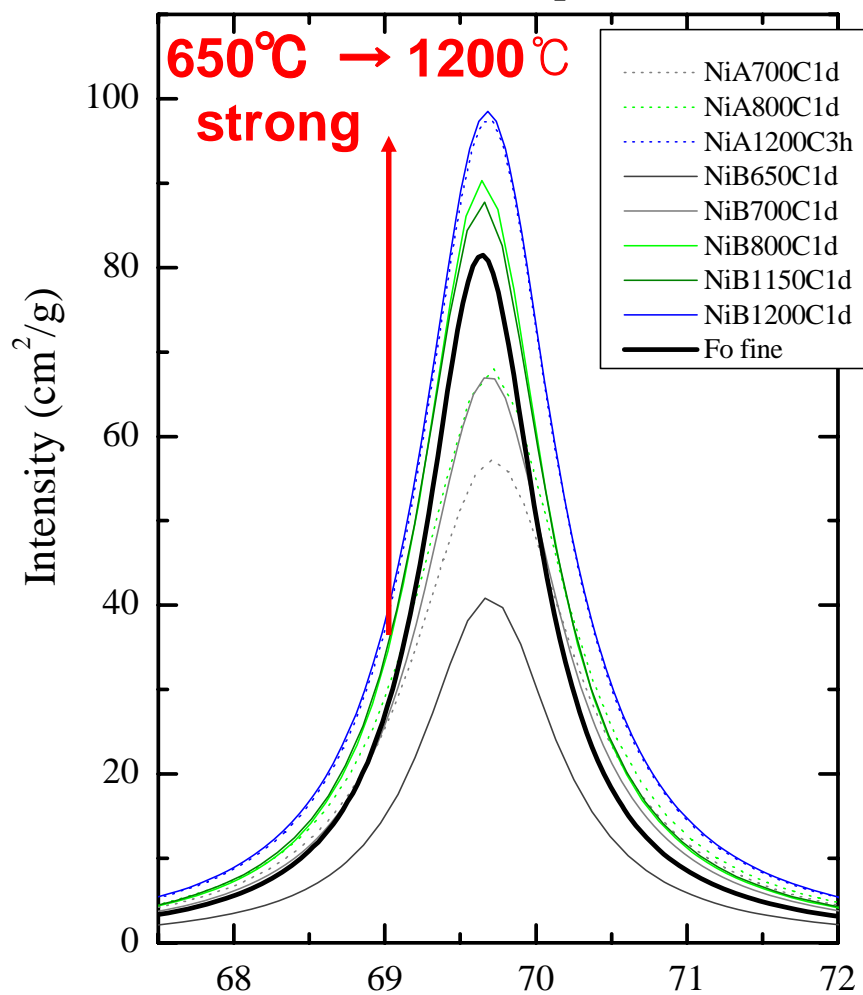
1600 $^{\circ}\text{C}$

crystal growth & shape effects

shape effects

NiA&B_peakfit RT

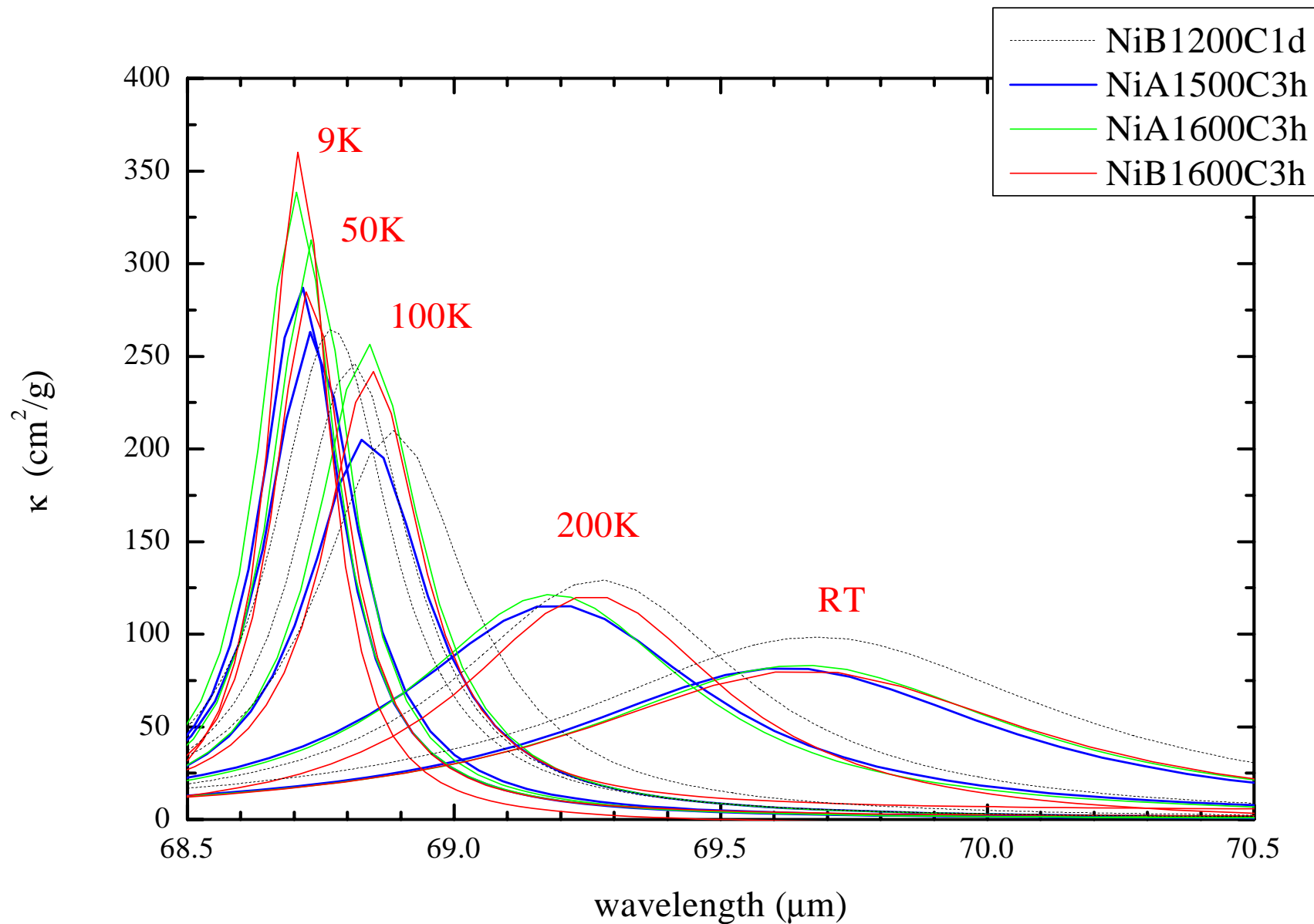
NiA&B_peakfit RT

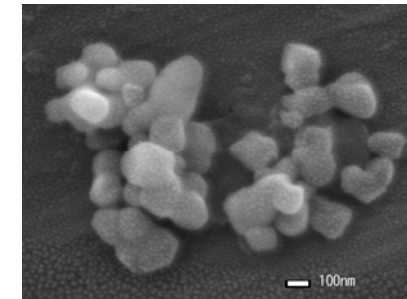
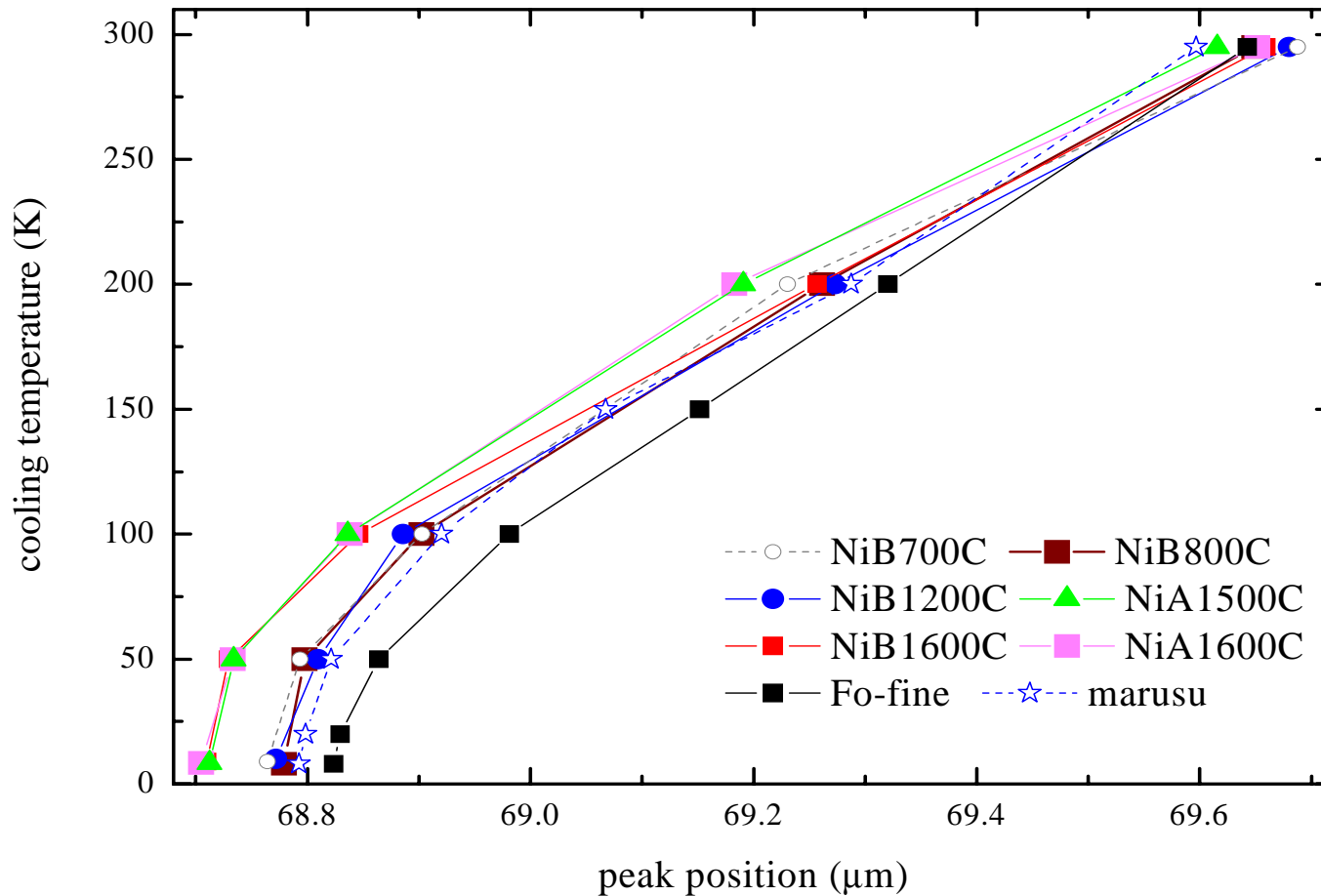


Peak positions are nearly same

wavelength (μm)

69 μ m band at each cooling temperature



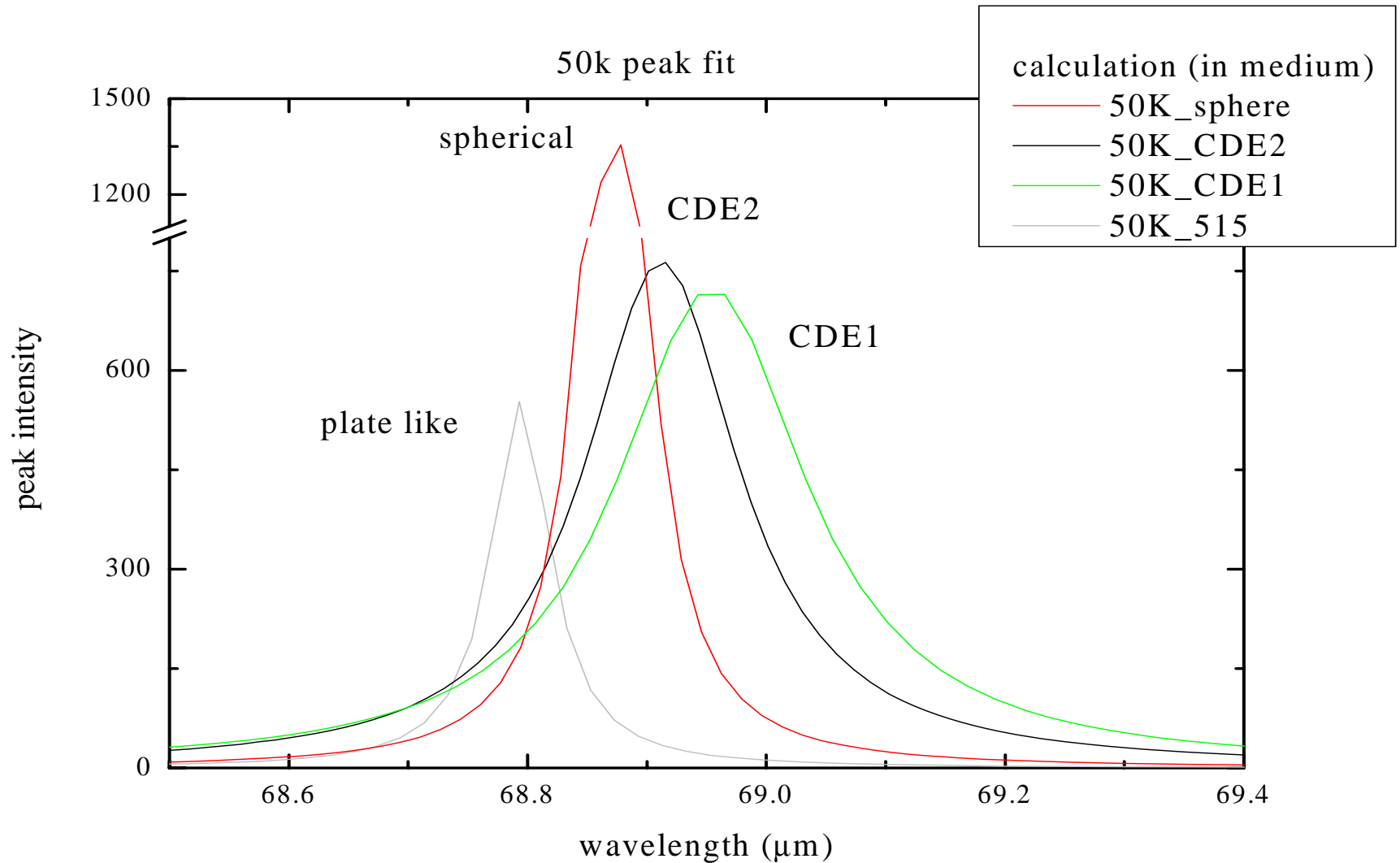


Marusu Fo
(commercial)
(about 0.2 μ m)
(elliptical)

All peak positions shift shorter than that of Fo fine lower 200 K

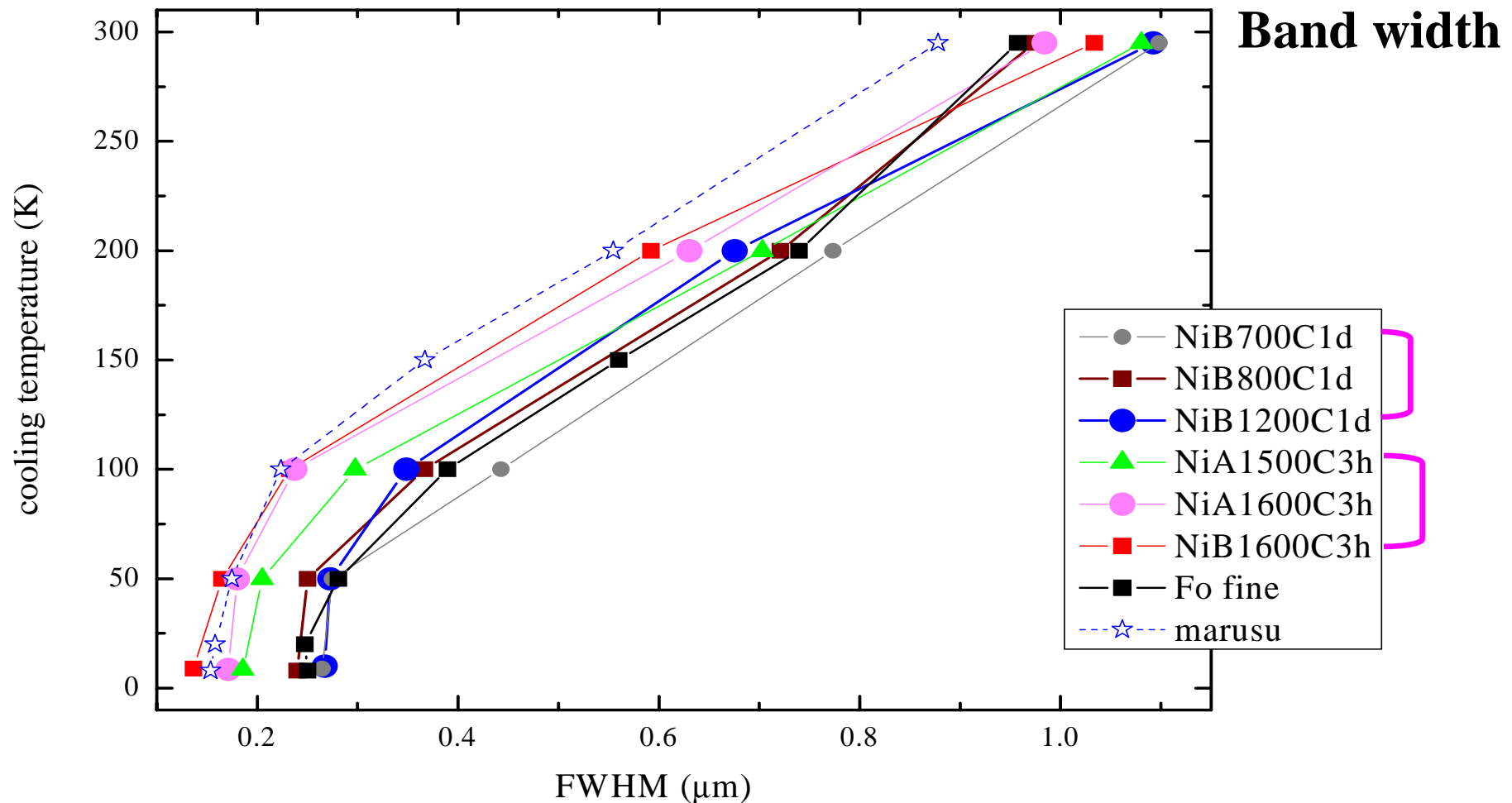
Samples of annealing at 1500C and 1600C

**Peak positions shift to a little shorter wavelength than spherical
this may be due to increase plate-like shape? It is not clear.**



Peak position

plate like : shift to shorter wavelength than spherical



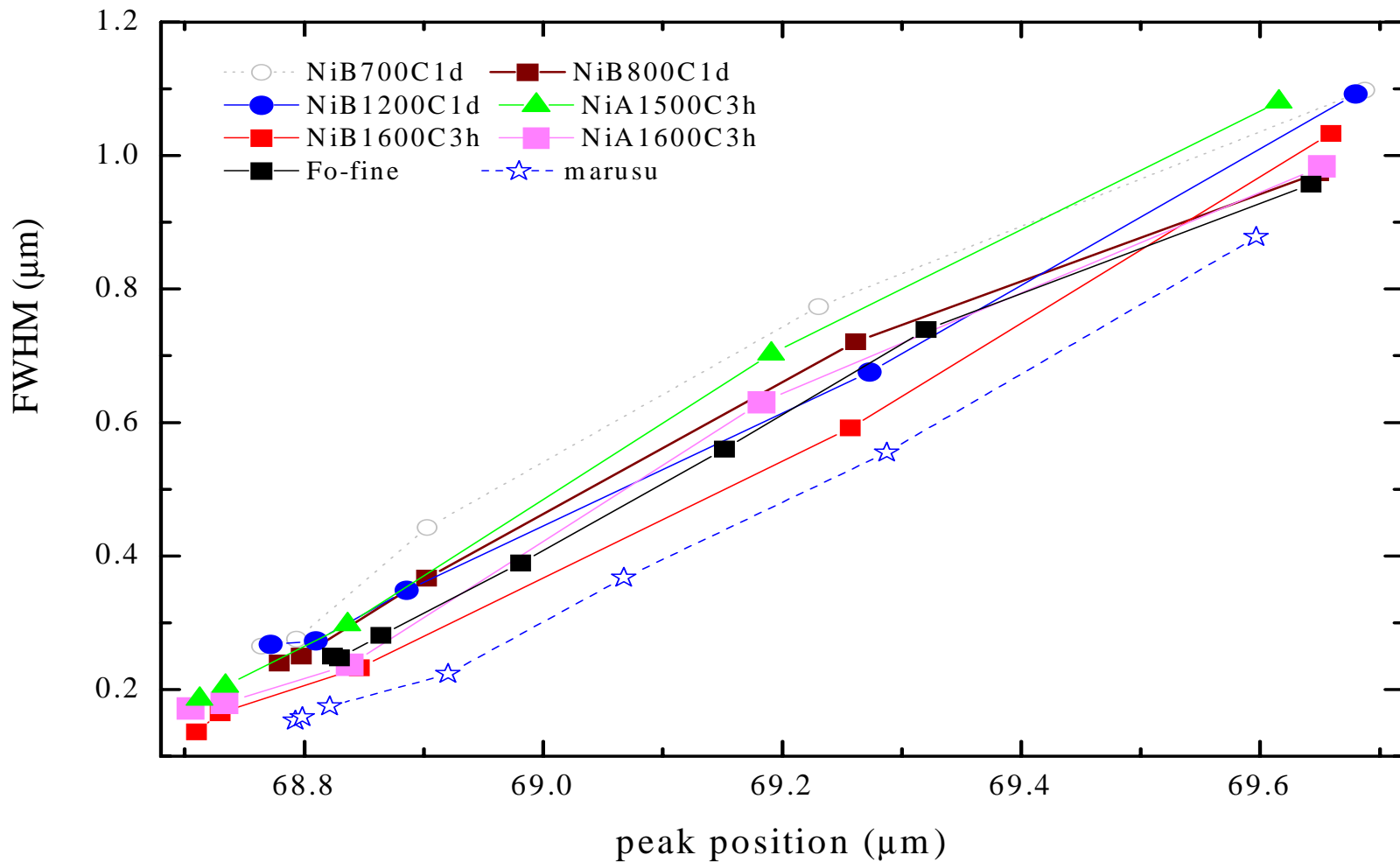
Cooling temperature ↓ : FWHM (Full Width of Half Maximum) ↓

At low temperature : high annealing temperature samples

FWHM became a little sharp

: may be due to similar size ?

Marusu Fo: band is very sharp due to similar size ?



Cooling temperature became low

Peak position & FWHM : correlation

The tendency is same as for annealed sample and Fo fine except for marusu

Summary

Annealing temperature

Below 1200 °C for both products

spectra depend on shape (spherical & coagulation)

11 μ m, 19 μ m, 23 μ m, 33 μ , 69 μ m band

for spherical ---- each peak became sharp and strong

above 1200 °C for both products (irregular)

all spectra became similar to that of Fo fine

69 μ m band for cooling

peak strength

depend on shape (annealing temperature) & coagulation