

Cryo-EM, basic image processing course, ReNaFoBis, Oléron, 22.5.2016, B. Klaholz

To get a screenshot from your displayed images: Maj cmd 4 → selection; Maj cmd 3 → full screen
Capture screen shots under Mac: **Control-Shift-4** → select region of interest with mouse

Control-Shift-3 → full screen shot

Display in imagic: use “ * ” or “quit” to leave, do not use CTRL C;

Use **CTRL Z** and **bg / fg** to create batch job or bring it back

I. Illustration of the Fourier transformation

cp ./FT-effects/* .

[copy over the images to work on]

Files names are:

description:

checker_8	checkerboard array
checker_32	checkerboard array
disc	sharp disc
disc-smooth	smooth disc
square	square

Calculate Fourier transformation of these images:

i

[shortcut for starting IMAGIC program]

IMAGIC-COMMAND : **fft**

** INCFFT2D (vs. Aug. 2005) welcomes you **

Input file, image loc#s [] :

checker_8

Output file, image loc#s [] :

checker_8-fft

[give output file name]

Mode of operation:

FORWARD_FFT REVERSE_FFT AUTO_CORRELATION

SELF_CORRELATION POWER_SPECTRUM AMPLITUDE_SPECTRUM

Please specify option [] : **FORWARD**

[hit return]

[do the same for the other images: produce files:

checker_8-fft, checker_32-fft, disc-fft, disc-smooth-fft, square-fft

quit [or * or Ctrl C] when finished

display these files in IMAGIC:

disp [under linux or within IMAGIC]

Input image file, loc#s [checker_8] : **checker_8**

Size of the display window [600,600] :

[hit return for default]

Type of cursor:

CROSS SQUARE CIRCLE

Please specify option [CROSS] : **[hit return for default]**
Parameters to be changed:
NO_CHANGES(=DISPLAY), SETTINGS, OPTIONS [NO] : **[hit return for default]**

To adjust scale of display:

scale

4

To adjust grey values:

gre

sur

2d

Useful options:

grey [to adjust the dynamic range of the image]

interactive

0,0 [full range] or for example -10,10 [limited range]

file [read in another file]

filename

dev [device, size of display window]

600,1200

erase [removes displayed image, to display freshly another one]

profile [to make profile]

Use cursor to position profile: NO

Starting point (IMAGE coordinates X,Y): 1,1

End point (IMAGE coordinates X,Y): 65,65 [center of a 128,128 image, i.e. center of powerspectrum]

To start a second display:

Ctrl Z

bg [background, batch job]

To quit the display:

*** [or] quit very import!!! (otherwise display problems)**

reactivate a background job: fg [foreground], then stop it with * or quit

disp

Input image file, loc#s [checker_8] : **checker_32** [next file name]
etc.

Switch between display windows to compare the images (do not move displays around such that they remain aligned with respect to each other)

Then display the corresponding FT's, files:

checker_8-fft

checker_32-fft

Switch between these display windows to compare the images

With the same procedure, compare sharp and smooth discs and the square:

disc disc-fft

disc-smooth disc-smooth-fft

square square-fft

When displaying the files **disc-fft** and **disc-smooth-fft** you can draw a profile of the spectrum:

In the display command window:

Parameters to be changed:

NO_CHANGES(=DISPLAY), SETTINGS, OPTIONS [NO] : **profile**

Use cursor to position profile [NO] : **[NO; hit return for default]**

Starting point (IMAGE coordinates X,Y) [1,1] : **70,70** [centre would be 65,65]

End point (IMAGE coordinates X,Y) [128,128] : **128,128**

Parameters to be changed:

NO_CHANGES(=DISPLAY), SETTINGS, OPTIONS [NO] : **[hit return for default]**

Output device (X_WINDOWS, PS, FILE) [X_WINDOWS] : **[hit return for default]**

Display settings:

device

600, 1200

scale

4

file

filename

grey

-10,10

profile

1,1

65,65

**Capture screen shots under Mac: Control-Shift-4 → select region of interest with mouse
Control-Shift-3 → full screen shot**

When interpreting the results, consider that the absolute scales on the y-axis can be different!

II. Pre-processing:

a) Display a digitized micrograph / negative or CMOS camera image of single particles imaged by cryo-EM

In your team directory:

```
cp ./micrograph/* .  
boxer &
```

read in the file called **10719c3** or one of the files called **1.mrc**:

File → read Micrograph

Process → Median Filter 5x5 (makes a block convolution)

**do you see anything?
now better?**

adjust grey values/contrast: **middle mouse button**

change scale to **0.4**

change box size to 96 or 128 [adjust box size to the particle size: should be ~2/3 of the image size; will be smaller than 128 if you use the **1.mrc** image from the CMOS camera; ideally, values of the type of 64, 96, 128, 256, 512, 1024, 2048, 4096 etc. should be used for fast Fourier transform calculations)

select ~6-10 particles

Boxes → Autobox, adjust parameters for a reasonable selection and let it select automatically;
afterwards, deselect some bad images manually

[*if you want to process the next steps with your own data:*

Boxes → Save Box

read in again the file called 10719c3, change box size to 128:

Boxes → Resize Boxes: 128

Boxes → Save Boxed Particles to file name 10719c3_128]

Comment: for CTF correction write out into much larger boxes, e.g. 512

b) Calculate a power-spectrum:

ctfkit

→ Open particle set

File name: 9_ptcl

Adjust grey values to see the power-spectrum better (middle mouse button), adjust parameters to make the predicted spectrum fit with experimental spectrum, adjust defocus value to make the high-resolution peaks fit (not the first peak and first zero which contain information from the particle itself, e.g. secondary structure elements)

Compare with power-spectra from other defocus values: file names: 7_ptcl 1_ptcl 10_ptcl

[*alternative program, not installed at the moment:*

findctf2d &

File → Open Micrograph file name: 10719c3

Moving the mouse over the image indicates the resolution: edge=Nyquist frequency!
(needs to have put 200kV and 3Å for the pixel size into CTF → Edit Microscope Settings)

Increase image size: Tools → Zoom → 200%

An outer mask can be put with the left mouse, and an inner mask with the right mouse button

Find out the defocus value of the micrograph:

CTF → Find CTF]

III. Processing of real experimental data

The basic steps of a structure determination of single particles:

- a) pre-process the data: bandpass-filter
- b) centering / alignment
- c) multivariate statistical analysis (MSA) and classification
- d) angle assignment
- e) 3D reconstruction

In your team directory:

cp ../* .

1) display particle set:

disp [under linux or within IMAGIC]

Input image file, loc#s [] : **CMOS_ctf-append_500**

Size of the display window [600,600] : **[hit return for default]**

Type of cursor:

CROSS SQUARE CIRCLE

Please specify option [CROSS] :

[hit return for default]

Parameters to be changed:

NO_CHANGES(=DISPLAY), SETTINGS, OPTIONS [NO] :

[hit return for default]

To start a second display:

Ctrl Z

2) bandpass-filter

i [shortcut for starting IMAGIC program]

IMAGIC-COMMAND : inc-pre

** INCPREP (vs. 21-May-2007) welcomes you **

Use MPI parallelisation [NO] : **NO [hit return for default]**

Input file, image loc#s [ctf-append_1000] : **CMOS_ctf-append_500**

Output file, image loc#s [ctf-append_1000-bp] : **CMOS_ctf-append_500-bp**

The image will be band-pass filtered.

Please specify:

Low frequency cut off [] : **0.025 [roughly particle size, pixel size: 3Å, Nyquist 6Å]**

Remaining low-freq. transmission [] : **0.1 [leave 10% of low frequencies]**

High frequency cut off [] : **0.5 [high frequency cut off]**

The image will be masked by a circle. Please specify

the mask radius (pixels or fraction of inner radius)

If you specify a drop-off it will be a soft mask.

Mask radius, drop-off [] : **0.999 [keep maximum to the edge of a circular area]**

Desired new sigma [] : **3 [normalise the variance to 3 sigma]**

Invert the image densities [NO] : **[hit return for default]**

Display the filtered version of the particles for comparison

3) calculate the total sum of the particle images which will serve as a reference for particle centering

IMAGIC-COMMAND : **inc-sum**

** SUMMER (vs. 14-June-2007) welcomes you **

Mode of summing:

CONDITIONAL_SUM SOME_SUM TOTAL_SUM

Please specify option [TOTAL_SUM] : **[hit return for default]**

Input file, NO loc#s [] : **CMOS_ctf-append_500-bp**

Output file, image loc#s [] : **CMOS_ctf-append_500-bp_sum**

Display the file **CMOS_ctf-append_500-bp_sum**

4) particle centering:

IMAGIC-COMMAND : **ali-dir**

** ALIDIR (vs. 19-July-2007) welcomes you **

Alignment modes available:

TRANSLATIONAL ROTATIONAL HORIZONTAL VERTICAL

ALL

Please specify option [] : **TRANSLATIONAL**

Correlation functions available:

CCF MCF

Please specify option [] : **CCF**

Input file, image loc#s [] : **CMOS_ctf-append_500-bp**

Output file, image loc#s [] : **CMOS_ctf-append_500-bp_cent1**

Reference file, image loc [] : **CMOS_ctf-append_500-bp_sum**

Give this reference a number (1,2,...) [0] : **[hit return for default]**

Options to filter the reference(s):

NO_FILTER LOWPASS

Please specify option []: **LOWPASS** **[filtering the reference; try also NO_FILTER]**

Halfwidth value for low-pass filter [] : **0.1** **[e.g. 10% of the Nyquist frequency]**

Max shift (pixels/fraction of radius) [] : **0.3** **[e.g. 30% of the image size]**

Full output? [] **NO**

Maximum allowable (radial) shift is ... pixels.

...

IMAGE #-ITER ANGLE XSHIFT YSHIFT CCC

1	1	0.00	-1.14	3.81	0.1768
2	1	0.00	-2.32	0.08	0.1810 etc.

[[**optional:** 5) repeat steps 3 and 4 with the pre-centered images in order to center them even better]]
 final file containing centered images is **CMOS_ctf-append_500-bp_cent2 (or _cent1)**

display the files:

CMOS_ctf-append_500-bp CMOS_ctf-append_500-bp_cent1 (CMOS_ctf-append_500-bp_cent2) to check the success of the centering

6) create a mask for the area to be considered during multivariate statistical analysis (MSA)
 IMAGIC-COMMAND : **test-im**

** TESTIM (vs. 11-July-2007) welcomes you **

Output filename, image loc#s []: **msamask**
 Image dimensions X,Y [96, 96] : **84,84** [hit return for default]
 IMAGIC data formats you can choose:
 PACK INTG REAL COMP RECO
 Please specify option [REAL] : [hit return for default]
 Currently, you can choose:
 ...
 Please specify option [] : **DISC**
 Disc radius (pixel or fraction of inner radius) [] : **0.75** [to be adjusted to particle size]

7) multivariate sta tistical analysis (MSA):

IMAGIC-COMMAND : **msa-run**
 Use MPI parallelisation [NO] : **NO** [hit return for defaults]

** MSA (vs. 3-Sep-2011) welcomes you **

Choose mode of operation:
 FRESH_MSA_REFINE
 Please specify option [FRESH_MSA] : [hit return for default]
 MSA distances:
 EUCLIDIAN CHISQUARE MODULATION
 Please specify option [MODULATION] : [hit return for default]
 Input (= output) file (aligned "images") [] : **CMOS_ctf-append_500-bp_cent1**
 Input MSA mask file [msamask] : [hit return for default]
 Eigenimages output file [: **eigenim**
 Pixel coordinates output file [] : **pixcoos**
 Eigenpixel vectors output file [] : **eigenpix**
 Number of iterations (<65) [] : **25**
 Number of eigenimages (< 70) [] : **40**
 Overcorrection factor (0 < ocf < 0.9) [0.8] : **0.8** [hit return for default]
 Rootname for results file, NO ext. [msa] : [hit return for default]

Display the file **eigenim**

8) hierarchical ascendant classification:

IMAGIC-COMMAND : **msa-class**

** CLASSIFY (vs. Sept. 2006) welcomes you **

Input to be classified:

IMAGES PIXEL-VECTORS SEQUENCES

Please specify option [IMAGES] : **[hit return for default]**
 Input (=output) file (treated by MSA)[] : **CMOS_ctf-append_500-bp_cent1**
 Percentage of images to be ignored [0] : **[hit return for default]**
 Active eigenimages for classification [] : **30**
 Use default classification options [YES] : **[hit return for default]**
 What number of classes do you wish [] : **50** **[total particle number divided by number of members per class (usually 10-20, or 3-5 with high-contrast images)]**

Name of output results files []: **classes0_50**

9) form class averages:

IMAGIC-COMMAND : msa-sum
 ** CLASSUM (vs. 26-Feb-2007) welcomes you **

Input images to be summed [] : **CMOS_ctf-append_500-bp_cent1**
 Rootname of input classification files [] : **classes0_50**
 Output class averages [] : **classsums0_50**
 Downweight small classes [NO] : **[hit return for default]**

Mode of summing statistics:
 NONE VARIANCE S-IMAGE I-IMAGE FT

Please specify option [NONE] : **[hit return for default]**
 Fraction of worst class members to ignore [0] : **[hit return for default]**

Display the file classsums0_50 (and keep it displayed, use Ctrl Z)

10) band-pass filter the class averages:

IMAGIC-COMMAND : band-pass

** INCBAND (vs. Feb. 2007) welcomes you **

Input file, loc#s []: **classsums0_50**
 Output file, image loc#s [] : **classsums0_50-bp**

Filter options available:

BAND-PASS HIGH-PASS LOW-PASS
 INVERSE_BAND-PASS

Please specify option [BAND-PASS] : **BAND-PASS**

The image will be band-pass filtered.

Please specify

Low frequency cut off [] : **0.05** **[remember about Nyquist frequency...]**
 Remaining low freq. transmission [0.005]: **0.005** **[hit return for default]**
 High frequency cut off [] : **0.7**
 ASQ filter the images too [NO]: **[hit return for default]**

Display the file classsums0_50-bp (and keep it displayed, use Ctrl Z, bg to put the job into background)

11) Assigning angles without a reference, based on common lines

IMAGIC-COMMAND :
 ** EULER (vs.) welcomes you **

ang-rec

Pointgroup symmetry:

C1	1	C2	2
C3	3	C4	4
C5	5	C6	6
C7	7	C8	8
C9	9	C10	10
C11	11	C12	12
C13	13	C14	14
C15	15	C16	16
C17	17	C18	18
C19	19	C20	20
C21	21	C22	22
CN	N	D2	222
D3	32	D4	422
D5	52	D6	622
D7	72	D8	822
D9	92	TETRAGONAL 23	
O (CUBIC)	432	ICOSAHEDRAL 532	
NONE			

Please specify option [] : **C1** [C1 Point-group symmetry for an asymmetric object]

Option for angular reconstitution:

NEW	ANCHOR_SET
C1_STARTUP	SELF_SEARCH
SINOGRAM	SINE_CORRELATION
PREDICT_SINECORR_PEAKS	

Please specify option [] :

C1_STARTUP

Input (classum) images, NO loc#s [] :

classums0_50-bp

Loc# of THREE (classum) images to be used [] : **3; 4; 14** [choose 3 different views; separate location numbers by “;”]

Output (ordered) image file [my_ordered] :

[hit return for default; selected

class averages will be put into a new file called my_ordered]

Output sinograms, NO loc#s [my_sino] :

[YES, hit return for default; sinogram file]

ASQ filter the sinogram lines [YES] :

[hit return for default; amplitude square-root filtering]

Linear mask radius for sinograms [] :

0.7 [depends on particle size]

Output sinecorr file, NO loc#s [my_sine] :

[hit return for default; sinogram correlation file]

Wanted angular increment in search [5.0] :

[hit return for default]

Minimum inter-euler stay away angle [30.] :

[hit return for default]

Full output of the results [NO] : **YES**

Are the relative angles clearly bigger than ~40°? If not, select another set of 3 views and start again

Also look at sinograms: display file **my_sino** and sinogram correlation: **my_sine**

12) 3D reconstruction

 IMAGIC-COMMAND : **true**

** TRUE3D (vs. Jan. 2007) welcomes you **

MPI parallelisation:

ONLY_3D ALL NO

 Please specify option [] : **NO**

 Please specify option [] : **ALL in one**

Pointgroup symmetry to be used:

C1 1 C2 2

...

 Please specify option [] : **C1**

 Use default 3D reconstruction options [YES] : **[hit return for default]**

 Input 2D (classum) images, loc#s [] : **my_ordered**

Source of Euler angles:

ANGREC_HEADER_VALUES PLT_FILE

MRA_HEADER_VALUES

 Please specify option [ANGREC_HEADER_VALUES] : **[hit return for default]**

 Output 3D rec. filename, loc#s [] : **3d_0-1** [file which will contain the

3D reconstruction, sections by sections after weighted back-projection]

 Output file for reprojections, NO loc#s [] : **3d_0-1-reproj** [reprojections

according to the same Euler angles as the input images]

 Output file for error projections, NO loc# [] : **3d_0-1-err** [difference between
 reprojection and input image, i.e. reflects amount of error]

 Mask the reconstruction [] : **YES**

 Radius of the mask [] : **0.8**

 Hamming window factor [] : **0.6**

 Object size as fraction of image size []: **0.7**

Now display the files my_ordered and 3d_0-1-reproj for comparison, do they look similar, i.e. is the angle assignment correct?

13) Add more views to the angular reconstitution to improve the structure:

 IMAGIC-COMMAND : **ang-rec**

** EULER (vs. 27-Sep-2006) welcomes you **

Pointgroup symmetry:

C1 1 C2 2

...

 Please specify option []: **C1**

Option for angular reconstitution:

NEW ANCHOR_SET

C1_STARTUP	SELF_SEARCH
SINOGRAM	SINE_CORRELATION
PREDICT_SINECORR_PEAKS	
Please specify option [C1_STARTUP] : NEW	
Option of NEW:	
FRESH ADD REMOVE_PROJ	
Please specify option [ADD] : ADD	
Input (classum) images, NO loc#s []: classums0_50-bp	
Location number(s) wanted [] : 1;23;31;45 [select one image or a series of images]	
Output (ordered) image file [my_ordered] : [hit return for default]	
Output sinograms, NO loc#s [my_sino] : [hit return for default]	
ASQ filter the sinogram lines [YES] : [hit return for default]	
Linear mask radius for sinograms [] : 0.7 [as before]	
Output sinecorr file, NO loc#s [my_sine] : [hit return for default]	
Wanted angular increment in search [5.0] : [hit return for default]	
Full output of the results [YES] : NO	

14) 3D reconstruction with more views

IMAGIC-COMMAND : true

** TRUE3D (vs. Jan. 2011) welcomes you **

MPI parallelisation:

ONLY_3D ALL NO

Please specify option [NO] :	[as before]
Pointgroup symmetry to be used: ...	
Please specify option [C1] :	[as before]
Use default 3D reconstruction options [YES] :	[as before]
Input 2D (classum) images, loc#s [my_ordered] :	[as before]

Source of Euler angles:

ANGREC_HEADER_VALUES PLT_FILE	
MRA_HEADER_VALUES	
Please specify option [ANGREC_HEADER_VALUES] :	[as before]
Output 3D rec. filename, loc#s [3d_0-1] :	3d_0-2
Output file for reprojections, NO loc#s [3d_0-1-reproj] :	3d_0-2-reproj
Output file for error projections, NO loc# [3d_0-1-err]:	3d_0-2-err
Mask the reconstruction [YES] :	[as before]
Radius of the mask [0.85] :	[as before]
Hamming window factor [0.6] :	[as before]
Object size as fraction of image size [0.8] :	[as before]

Compare again new files my_ordered and 3d_0-2-reproj

15) Make forward projections

(could be used as references for a multiple-reference alignment, here only for comparing forward projections of 3d_0-1 and 3d_0-2)

IMAGIC-COMMAND : **threed-for**

** FORWARD (vs. Jan. 2007) welcomes you **

Input 3D image file [] : **3d_0-1**

Output file for forward projections []: **3d_0-1-24**

Threshold 3D density value [-99999] :

[hit return for default]

Option used for current IMAGIC command: FORWARD

Mode of interpolation for projecting:

NEAREST_NEIGHBOUR BILINEAR SPLINE

SINC NARROWING WIDENING

OBLIQUE_SAMPLING HEADERS_ONLY

Please specify option [WIDENING] :

[hit return for default]

Choose projection option:

FILE ANOTHER INTERACTIVE ORTHOGONAL

SPIRAL TETRAHEDRON TOMOGRAPHY STEREO

UNIFORM ICOSAHEDRON ASYM_TRIANGLE RANDOM

Please specify option [ASYM_TRIANGLE] :

[hit return for default]

Pointgroup symmetry to be used:

C1 1 C2 2

Please specify option [C1] :

[hit return for default]

Option to chose Euler angles:

EQUIDIST RANDOM

Please specify option [EQUIDIST] :

[hit return for default]

Option for Euler angle alpha:

ZERO ROTATE

Please specify option [ZERO] :

[hit return for default]

Wanted angular increment in search [] :

45

Please specify option [] :

NO

Do the same for 3d_0-2;

Compare files **3d_0-1-24** and **3d_0-2-24** , did the quality of the reconstruction improve?

If time allows: refine the structure by using forward projections as references: run m-r-a (multi-reference-alignment), then msa and classification and 3D reconstruction (beginning of the iterative procedure)

Optional: 16) display a 3D reconstruction

cp/3D/* .

a) **disp** [as consecutive sections]

b) **threed-se** and **movie** [as a 3D surface]

Once you have saved all screenshots for your report, please shutdown the system before leaving.