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Wadsworth Center

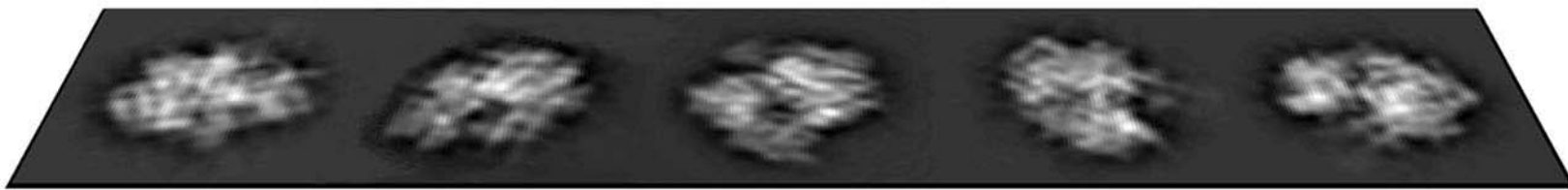
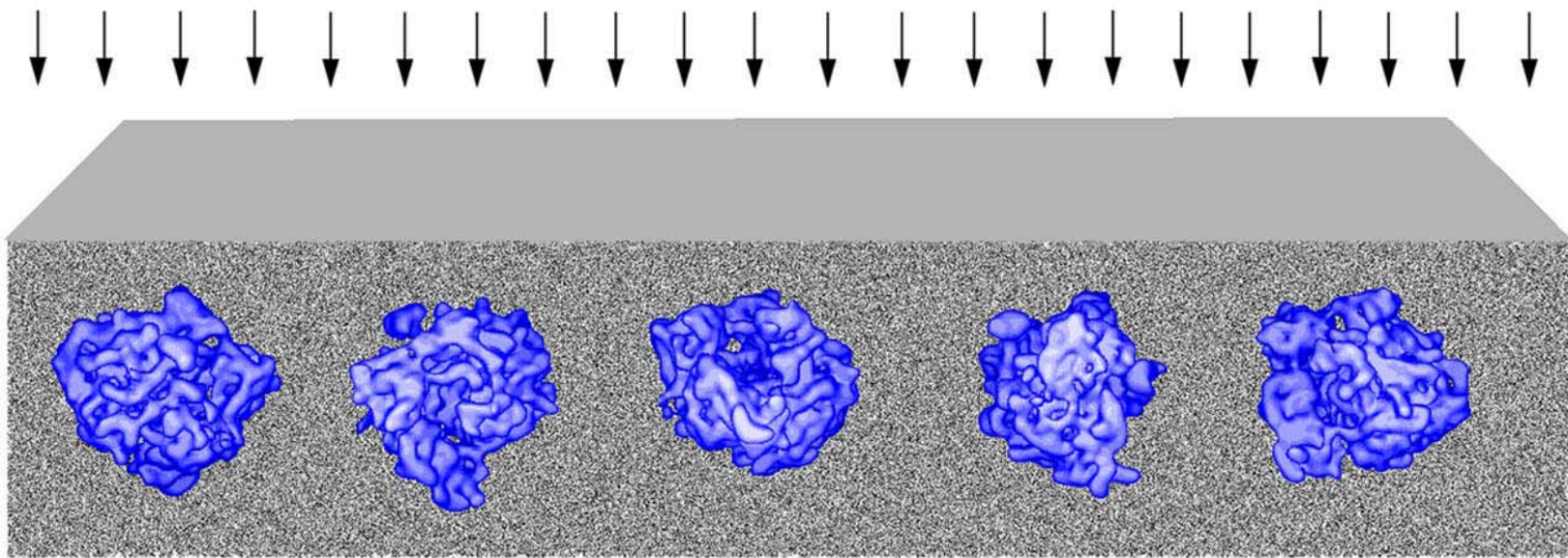
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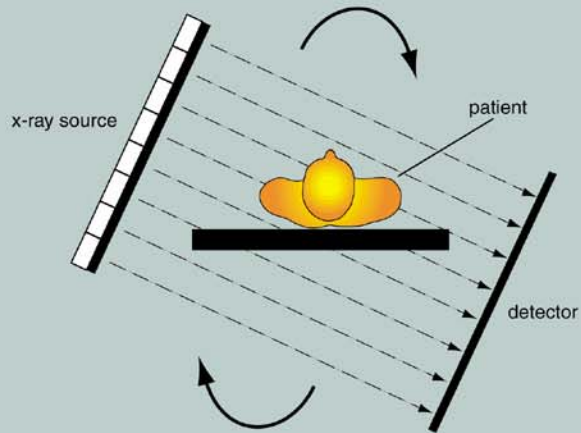
Single-particle reconstruction

Main assumptions:

- 1) All particles in the specimen have identical structure
- 2) All are linked by 3D rigid body transformations (rotations, translations)
- 3) Particle images are interpreted as a “signal” part (= the projection of the common structure) plus “noise”

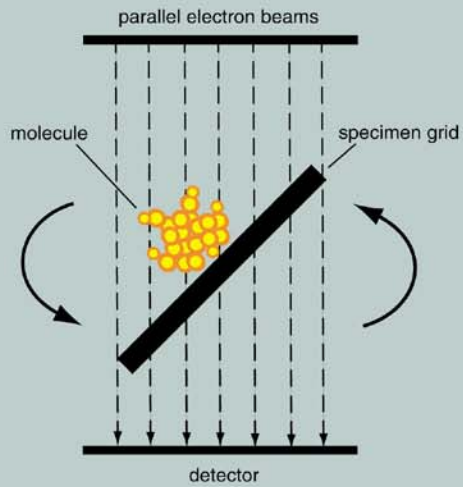
Important requirement:

even angular coverage, without major gaps.



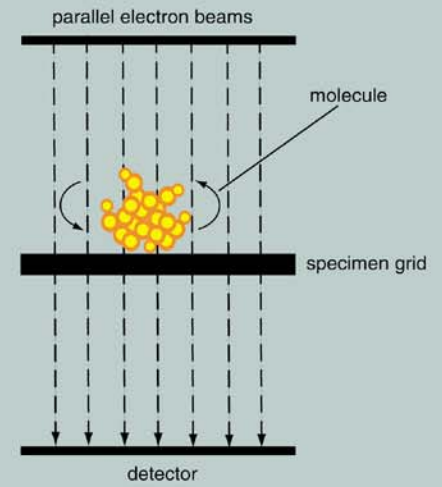
CAT - scan

- beam rotating
- patient stationary



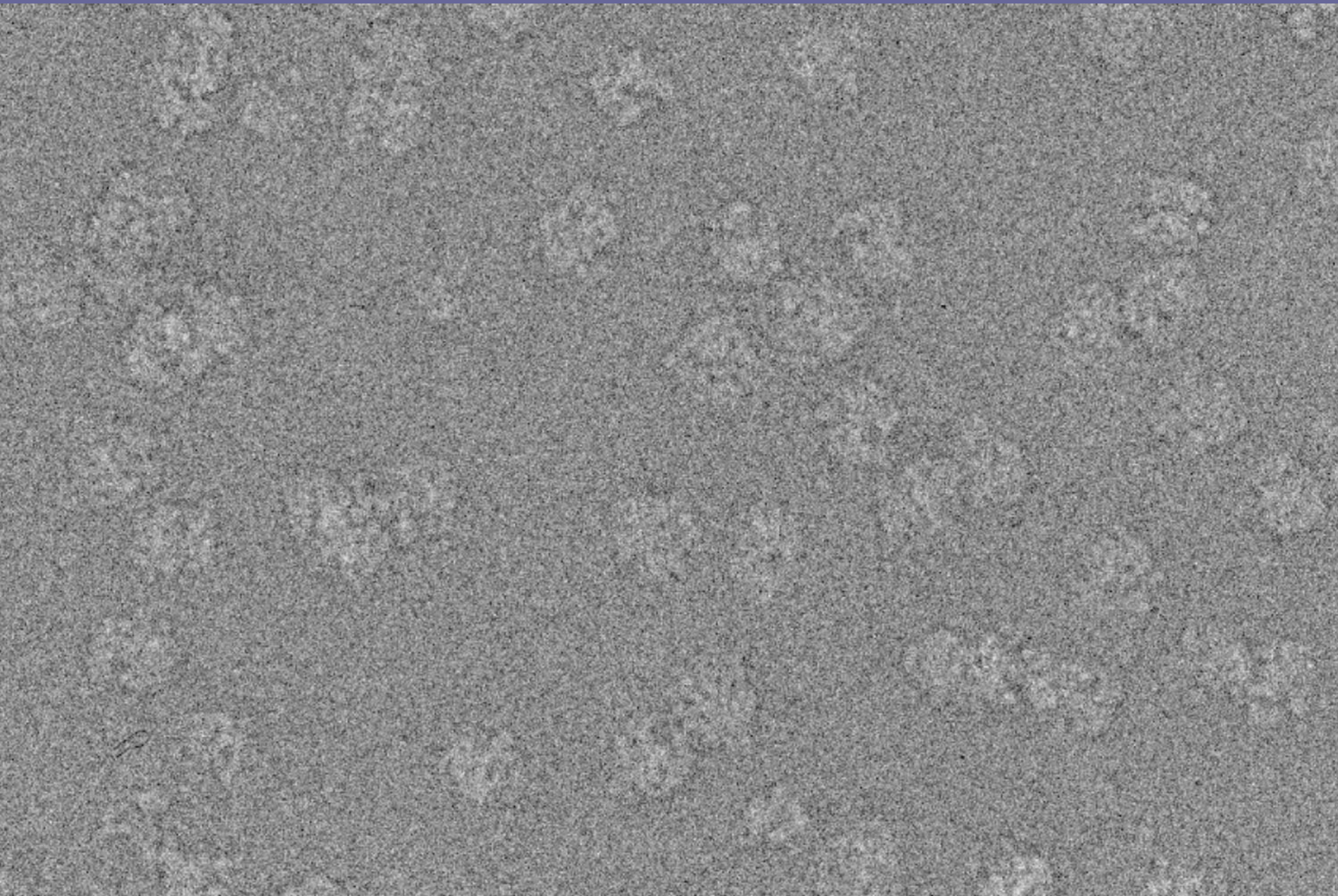
Electron Tomography

- molecule rotating
- beam stationary

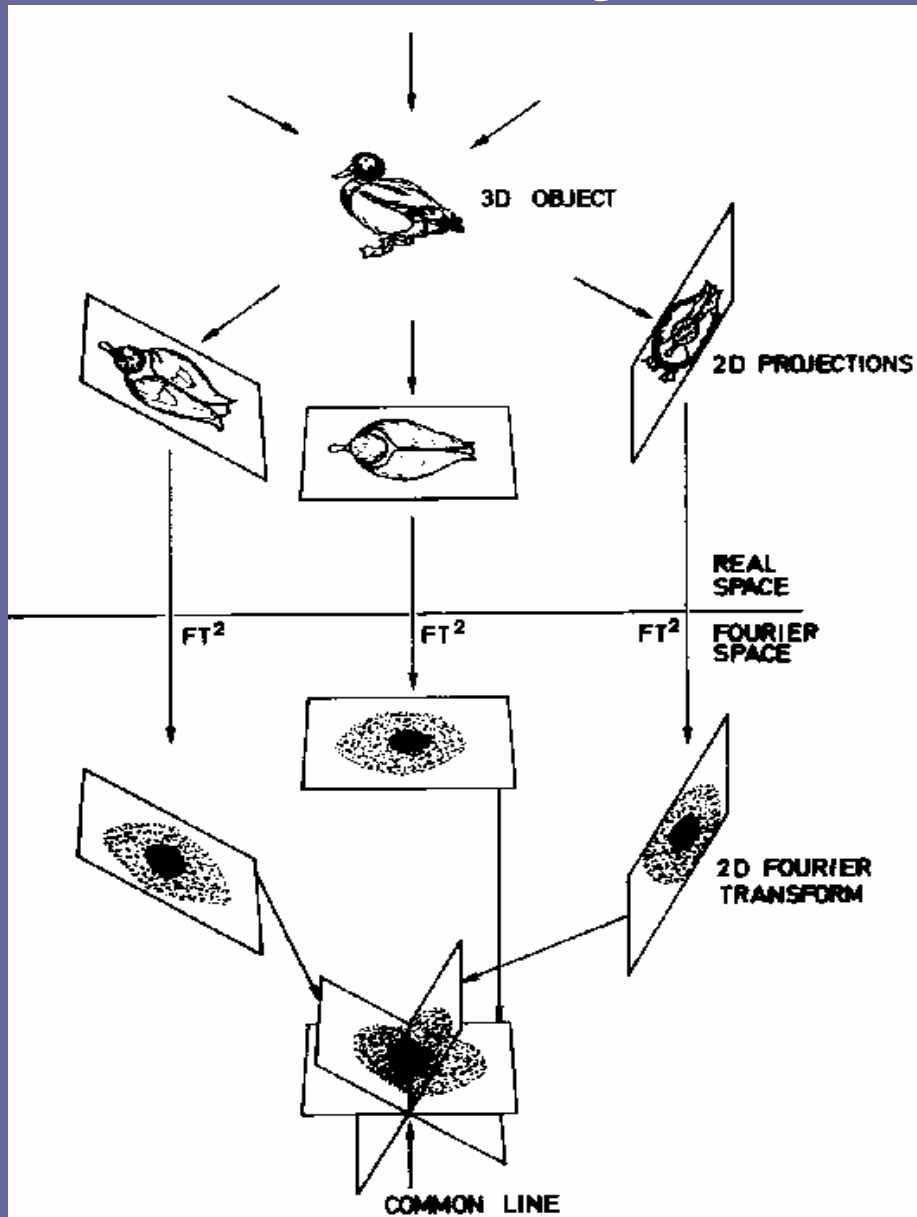


Single particle reconstruction

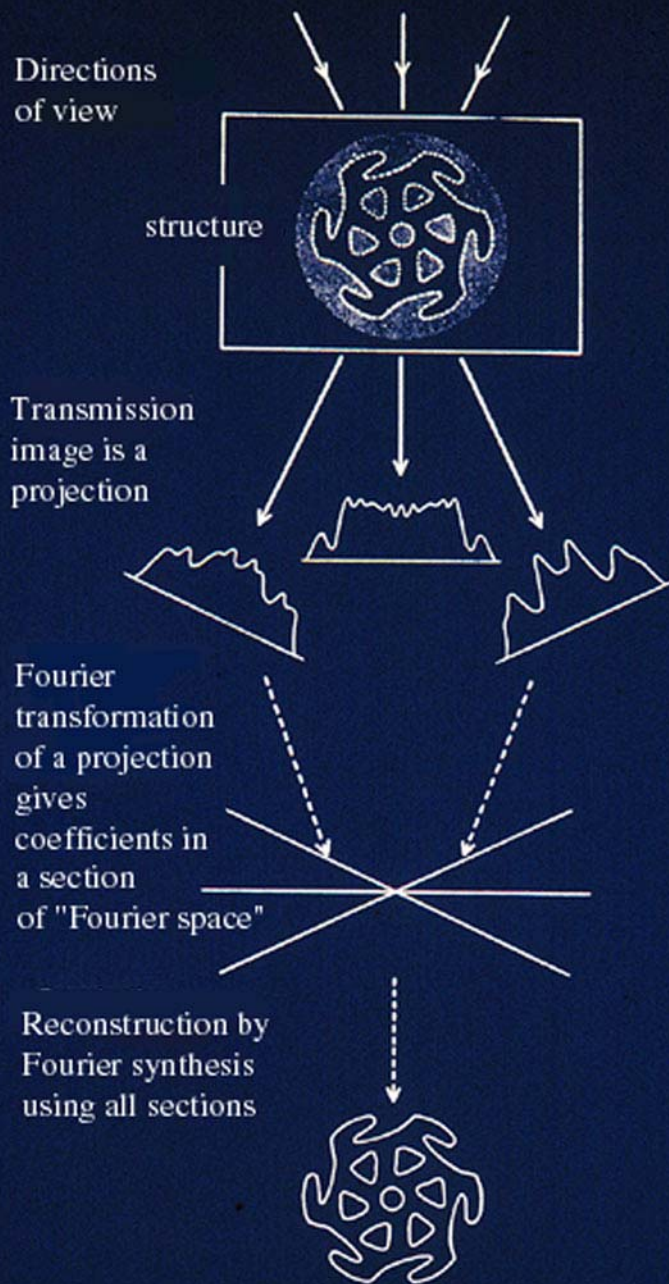
- molecule "rotating"
- beam stationary



Projection Theorem



The two-dimensional Fourier transform of the projection of a three-dimensional density is a central section of three-dimensional Fourier transform of the density perpendicular to the direction of projection.

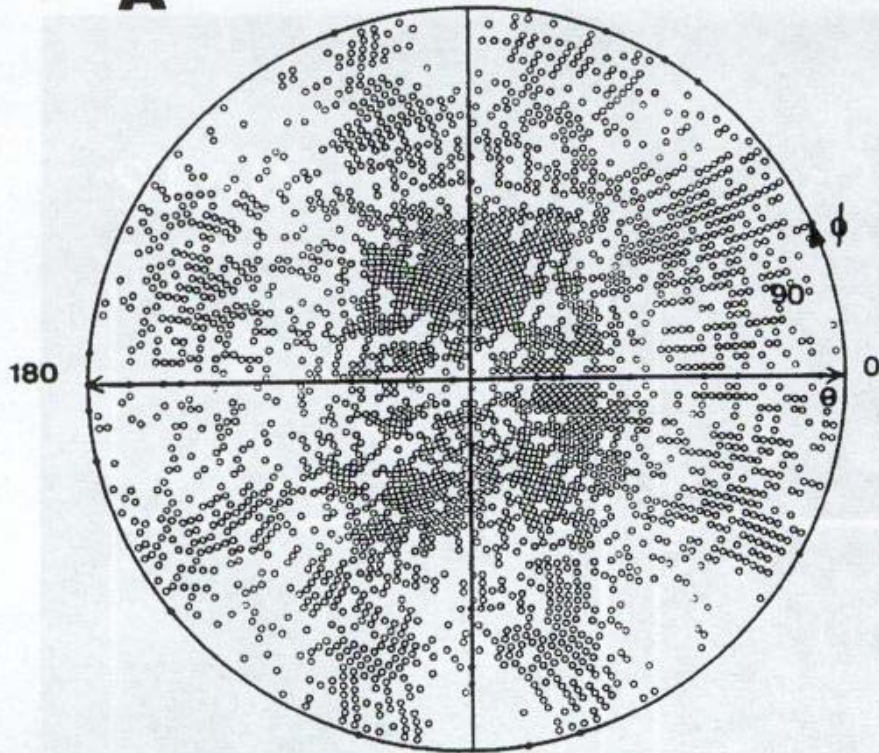


Angular coverage

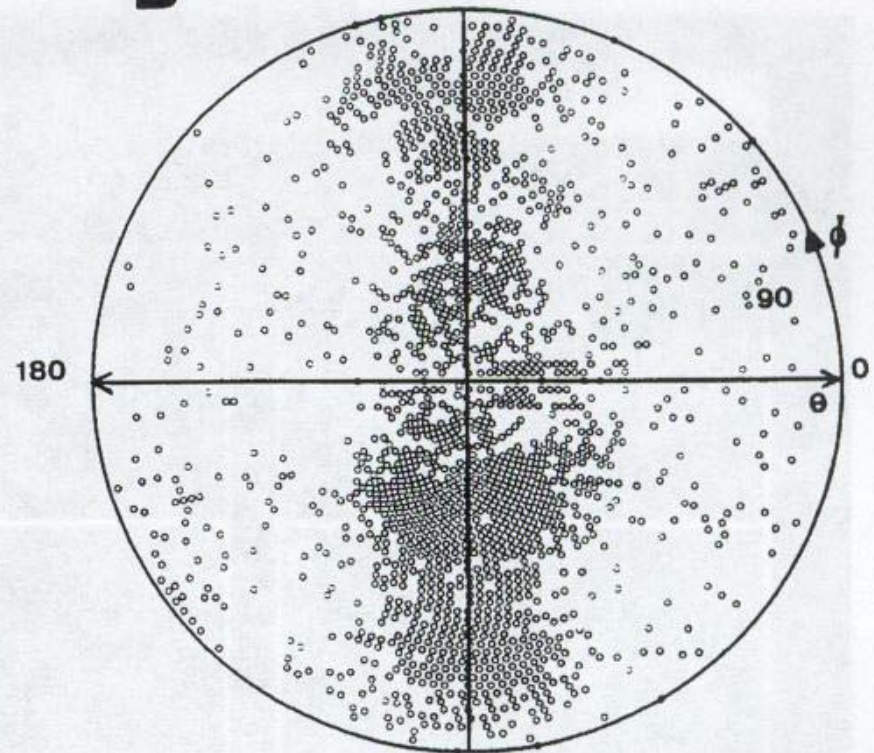
good

bad

A



B



Overview: the necessary steps of a single-particle reconstruction

- 1) *Optical diffraction: quality control, defocus inventory of micrograph batch*
- 2) *Scanning of micrograph batch*
- 3) *Determine defoci, and define defocus groups*
- 4) *Pick particles*
- 5) *Determine particle orientation*
- 6) *3D reconstruction by defocus groups*

---Steve---Steve---Steve---Steve---Steve-----

- 7) *Refinement*
- 8) *CTF correction*
- 9) *Validation*
- 10) *Interpretation: segmentation, docking, etc.*

Overview: tools

- 1) 2D alignment
 - usually by cross-correlation (translational, rotational)
 - (a) reference-based
 - (b) reference-free
- 2) Classification
 - (a) supervised (multi-reference, 3D projection matching)
 - (b) unsupervised
 - (i) K-means
 - (ii) Hierarchical ascendant
 - (c) self-organized maps (SOM)
- 3) Determine resolution
 - (a) phase residual
 - (b) Fourier shell correlation
 - (c) Spectral signal-to-noise ratio (SSNR)
- 3) Low-pass filtration
- 4) Amplitude correction (filter tailored acc. to experimental data)

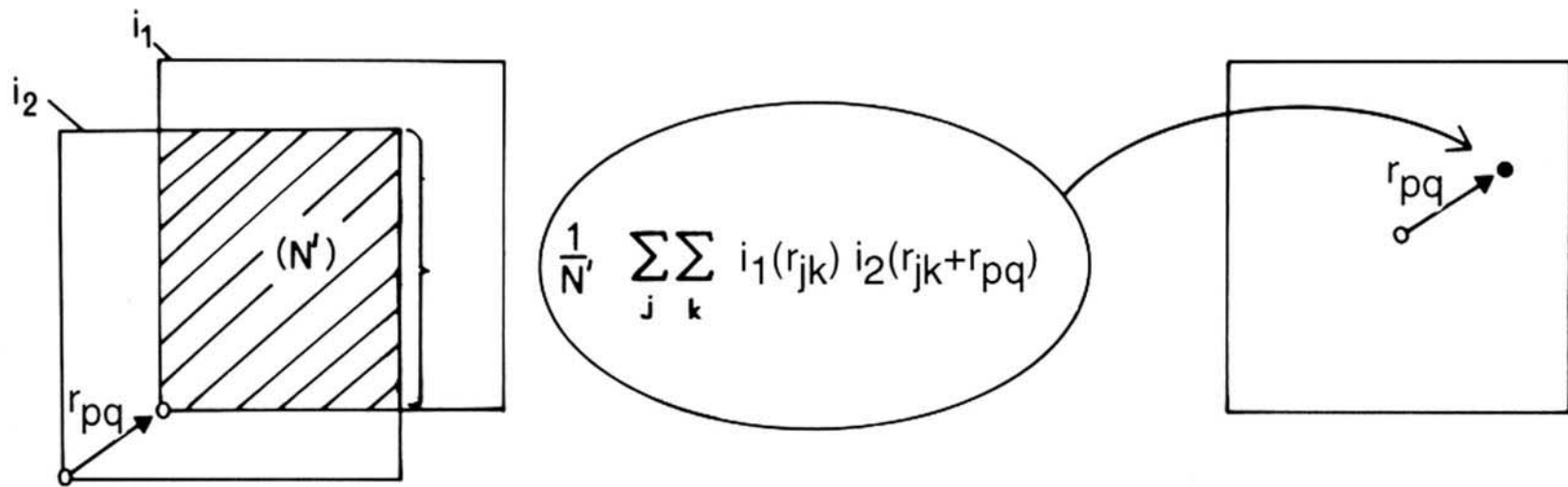


Fig. 3.8. Definition of the cross-correlation function. Image 1 is shifted with respect to image 2 by vector \mathbf{r}_{pq} . In this shifted position, the scalar product of the two images arrays is formed and put into the CCF matrix at position (p, q) . The vector \mathbf{r}_{pq} is now allowed to assume all positions on the sampling grid. In the end, the CCF matrix has an entry in each position. From Frank (1980). Reproduced with permission of Springer-Verlag, New York.

Alignment methods designed to minimize the influence of the reference

"Reference free" iterative alignment (Penczek *et al.*, 1992) :

Two images are randomly picked, aligned, and added.

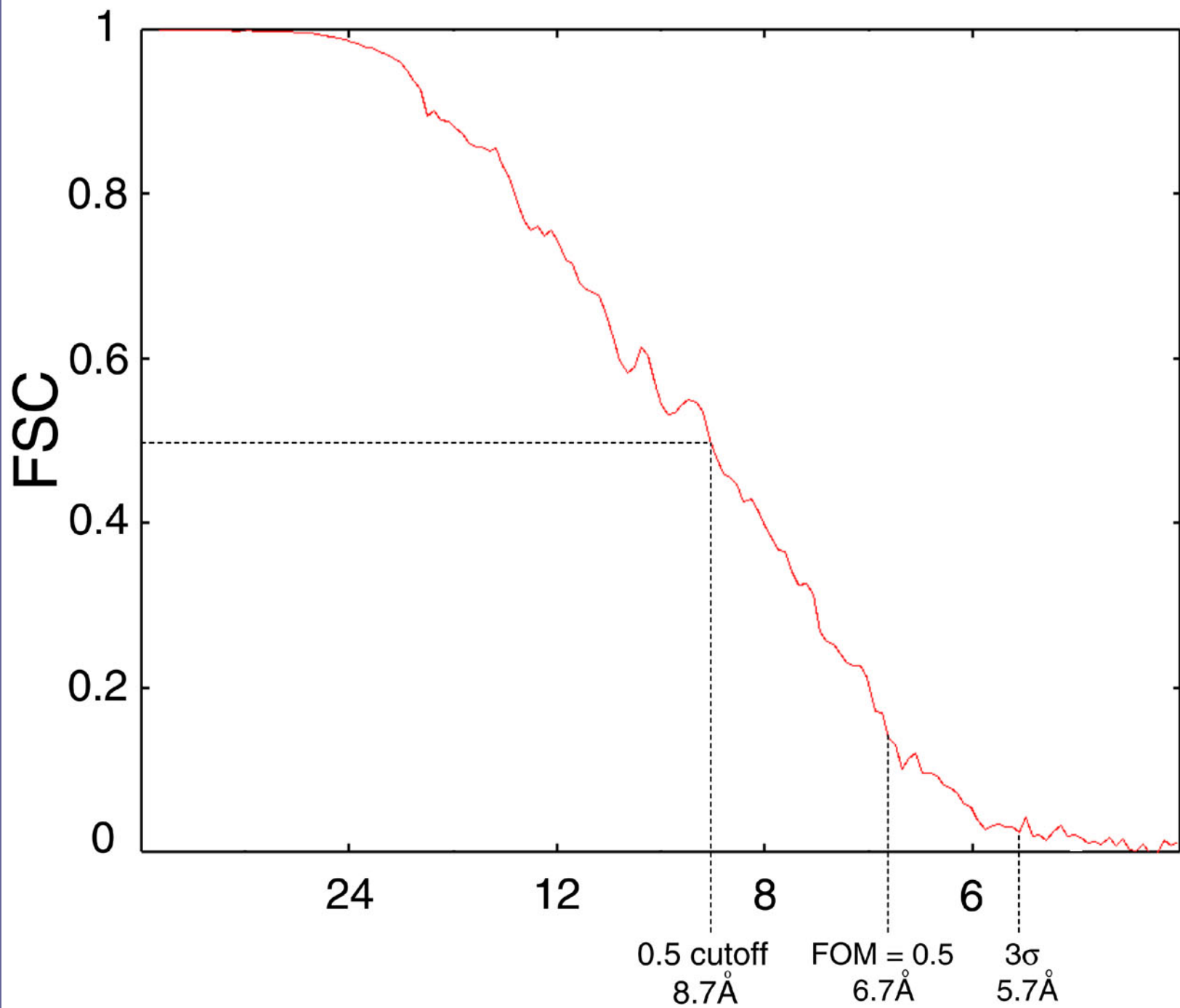
Then, a third image is aligned and added to the previous two. The process is repeated until all images are aligned.

To minimize the influence of the order in which images are picked, the first image is realigned to the [total average - image 1]. Then the second image is realigned to the [total average - image 2], etc ...

The whole process is started again until no improvement is found between one alignment cycle and the next.

Resolution measures & criteria: Fourier shell correlation

$$FSC(k, \Delta k) = \frac{\text{Re} \left| \sum_{[k, \Delta k]} F_1(\mathbf{k}) F_2^*(\mathbf{k}) \right|}{\left[\sum_{[k, \Delta k]} |F_1(\mathbf{k})|^2 |F_2(\mathbf{k})|^2 \right]^{1/2}}$$



Classification

Classification methods are divided into those that are “supervised” and those that are “unsupervised”:

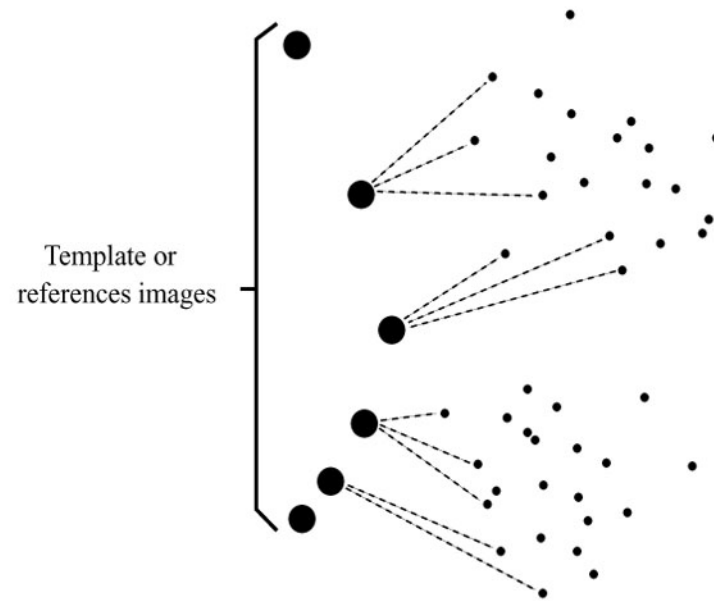
- Supervised: divide or categorize according to similarity with “template” or “reference”.

Example for application: projection matching

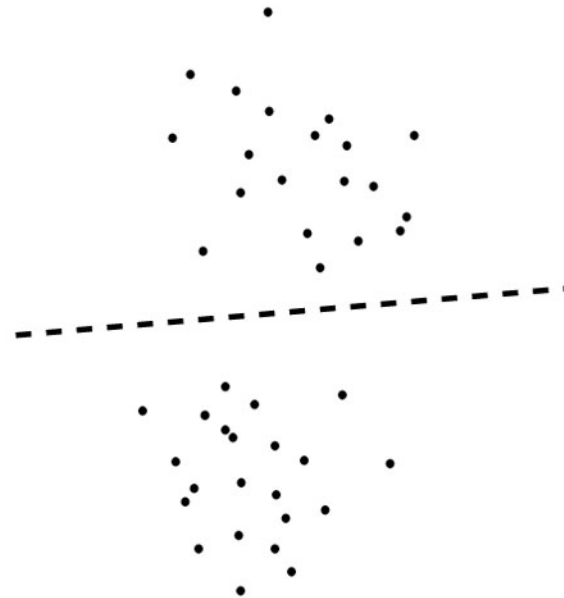
- Unsupervised: divide according to intrinsic properties

Example for application: find classes of projections presenting the same view

Supervised Classification



Unsupervised Classification

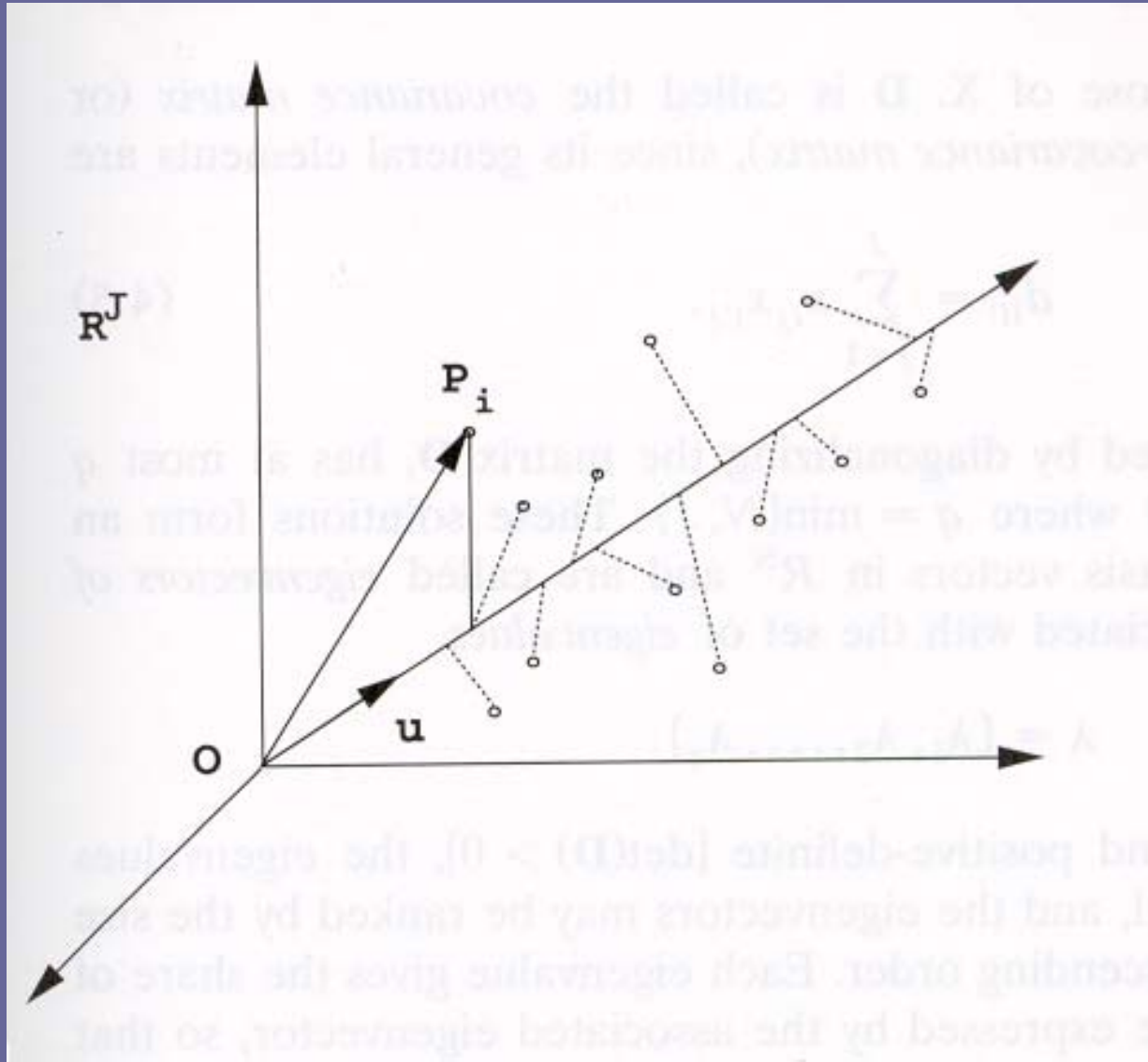


Classification, and the Role of MSA

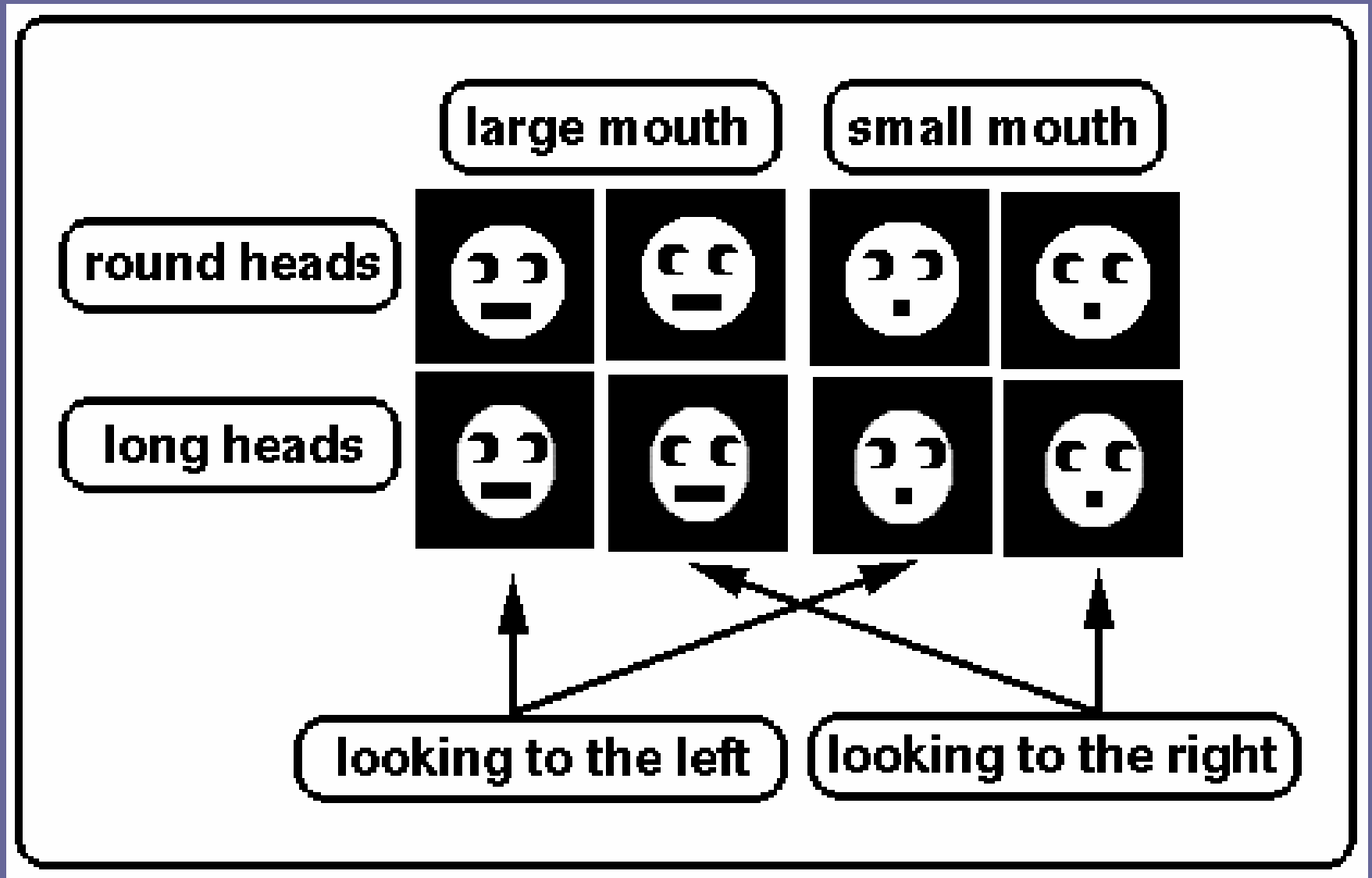
- Classification deals with “objects” in the space in which they are represented.
- For instance, a 64x64 image is an “object” in a 4096-dimensional space since in principle each of its pixels can vary independently.
Let’s say we have 8000 such images. They would form a cloud with 8000 points in this space.
- Unsupervised classification is a method that is designed to find clusters (regions of cohesiveness) in such a point cloud.
- Role of Multivariate Statistical Analysis (MSA): find a space (“factor space”) with reduced dimensionality for the representation of the “objects”. This greatly simplifies classification.
- Reason for the fact that the space of representation can be much smaller than the original space: resolution limitation (neighborhoods behave the same), and correlations due to the physical origin of the variations (e.g., movement of a structural component is represented by correlated additions and subtractions at the leading and trailing boundaries).

Principle of MSA:

Find new coordinate system, tailored to the data

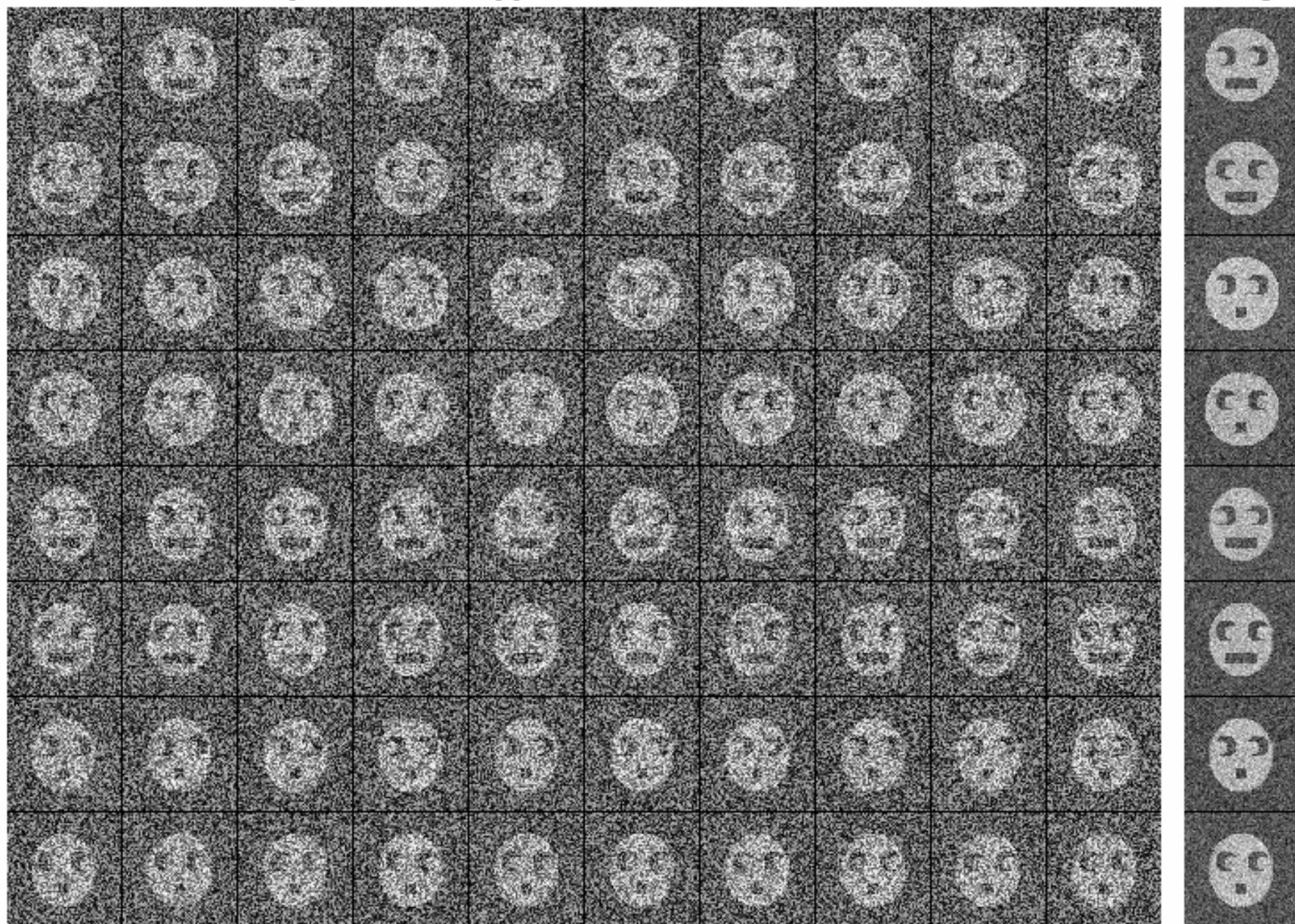


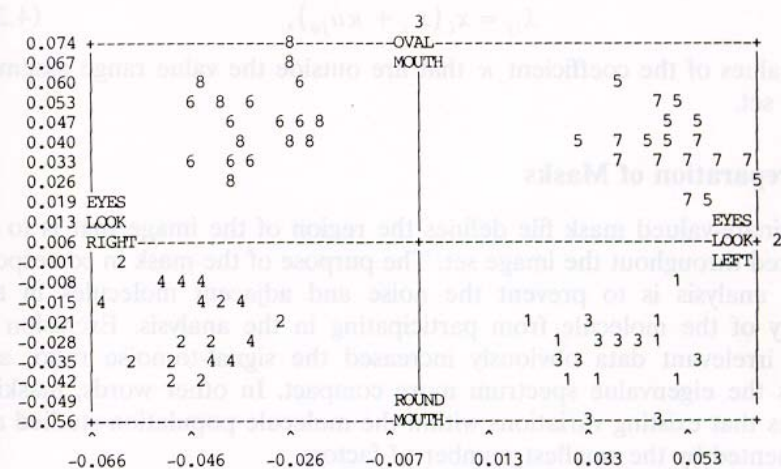
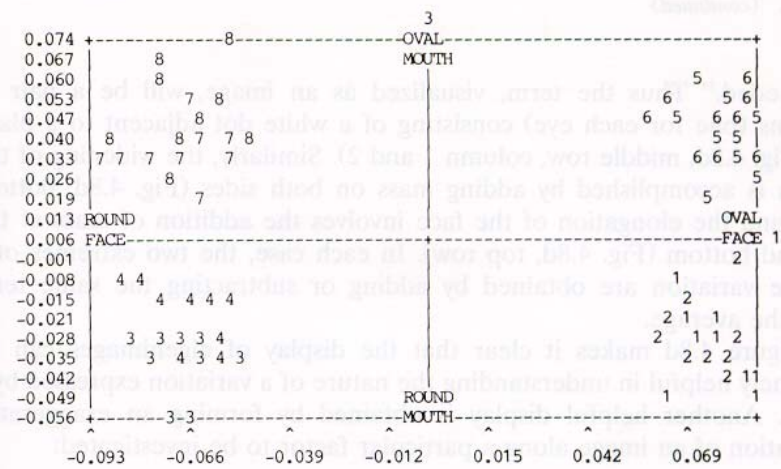
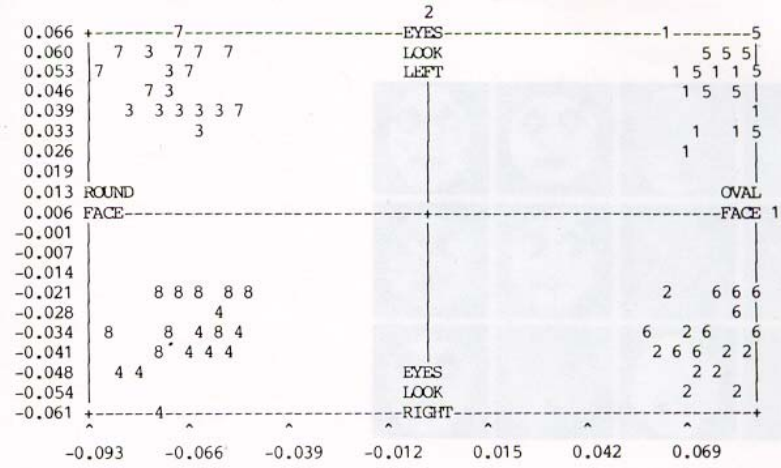
Brétaudière JP and Frank J (1986) Reconstitution of molecule images analyzed by correspondence analysis: A tool for structural interpretation. *J. Microsc.* 144, 1-14.



10 copies of the 8 types of heads + random noise

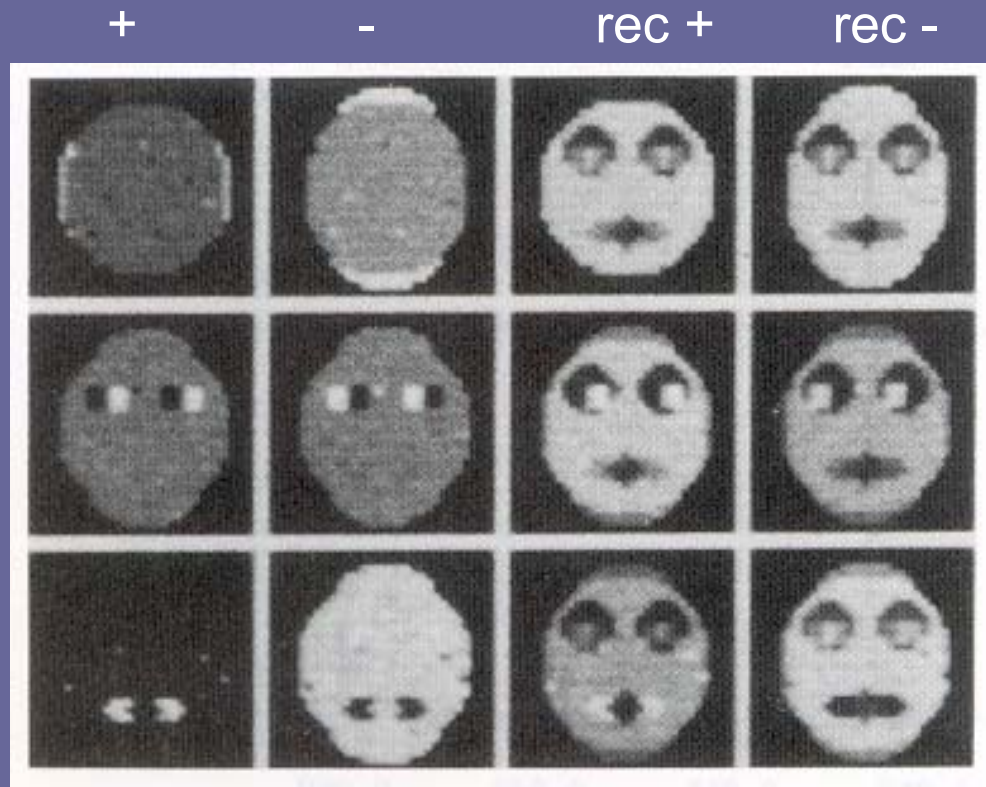
Averages





MSA: eigenimages

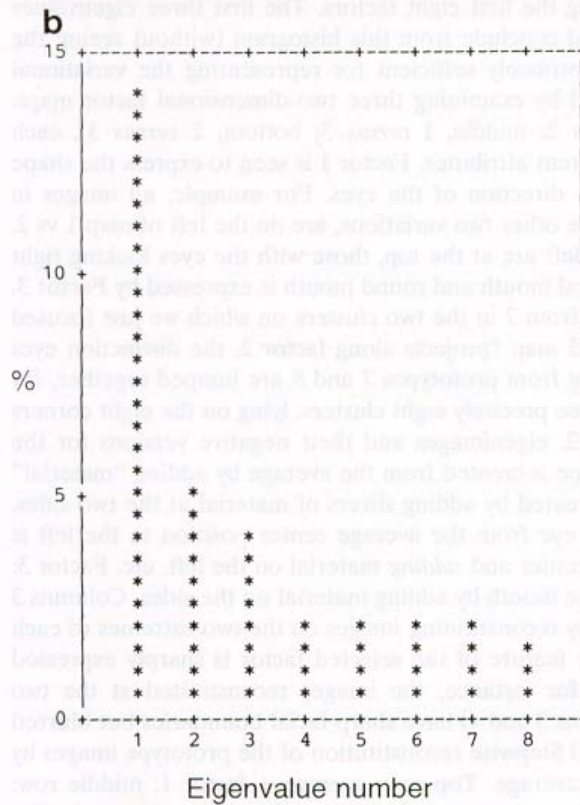
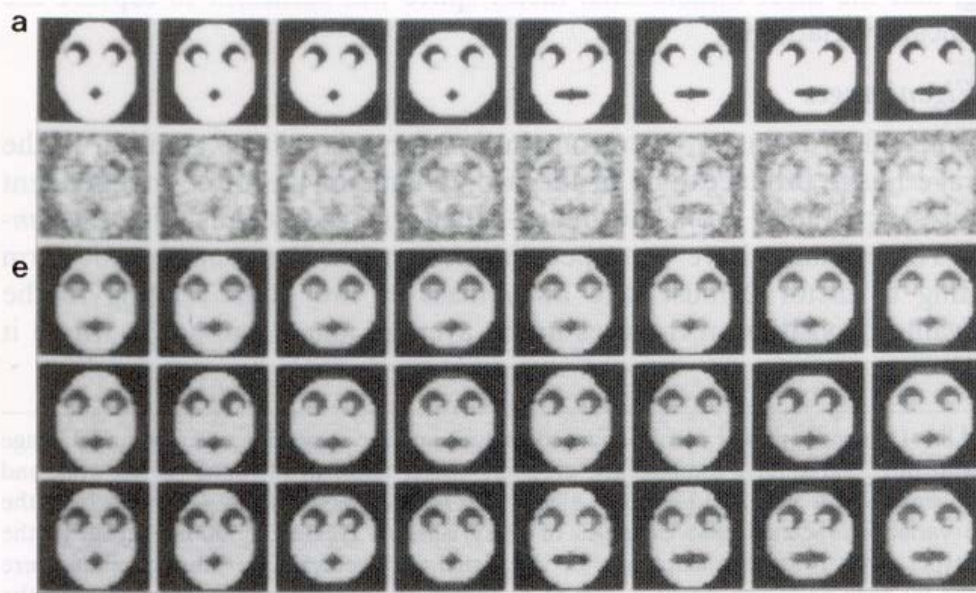
-
- Factor 1
- Factor 2
- Factor 3



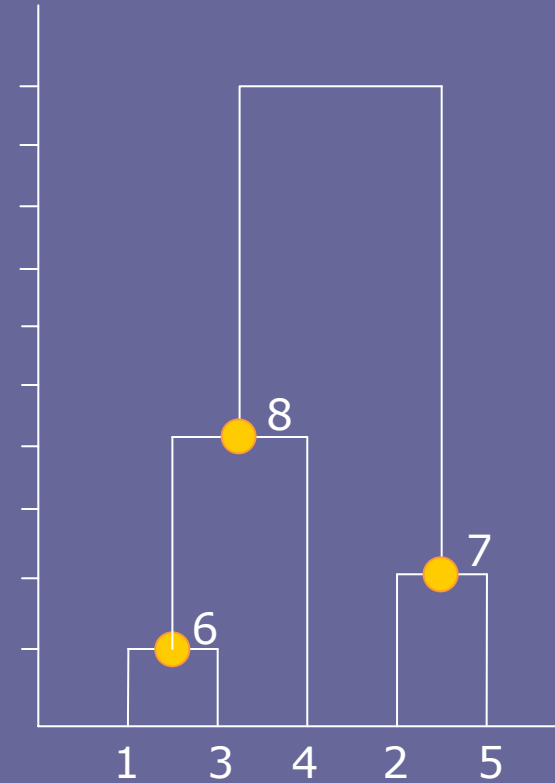
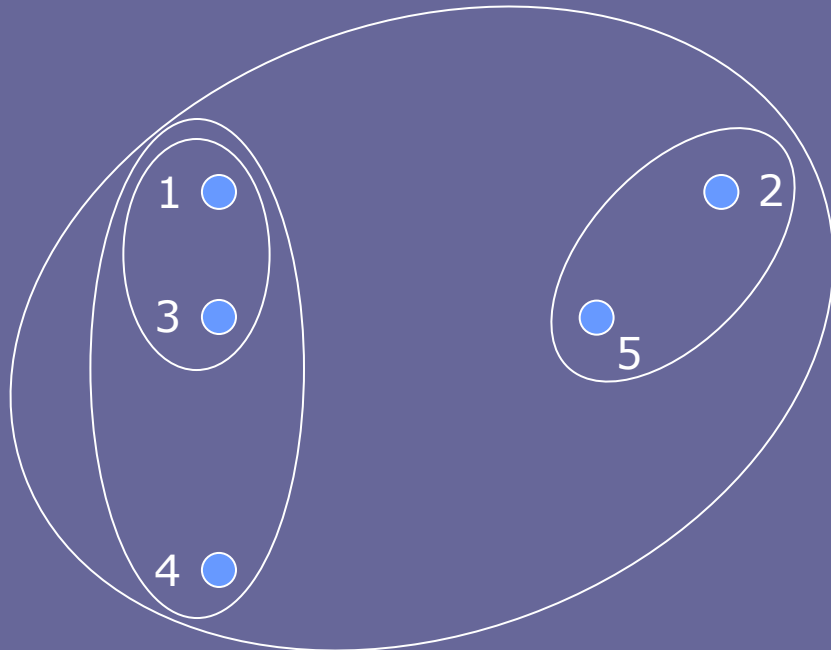
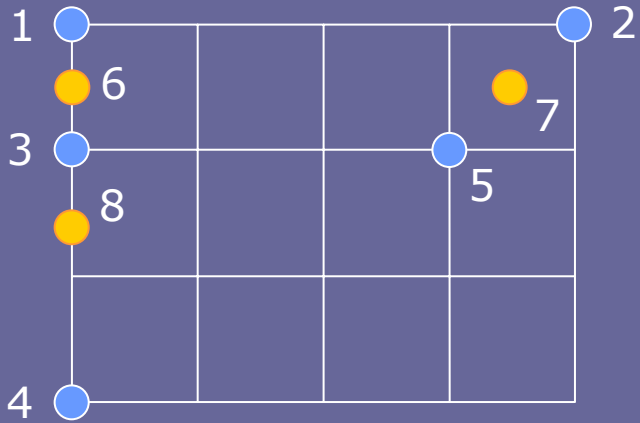
Avrg + F1

Avrg + F1+F2

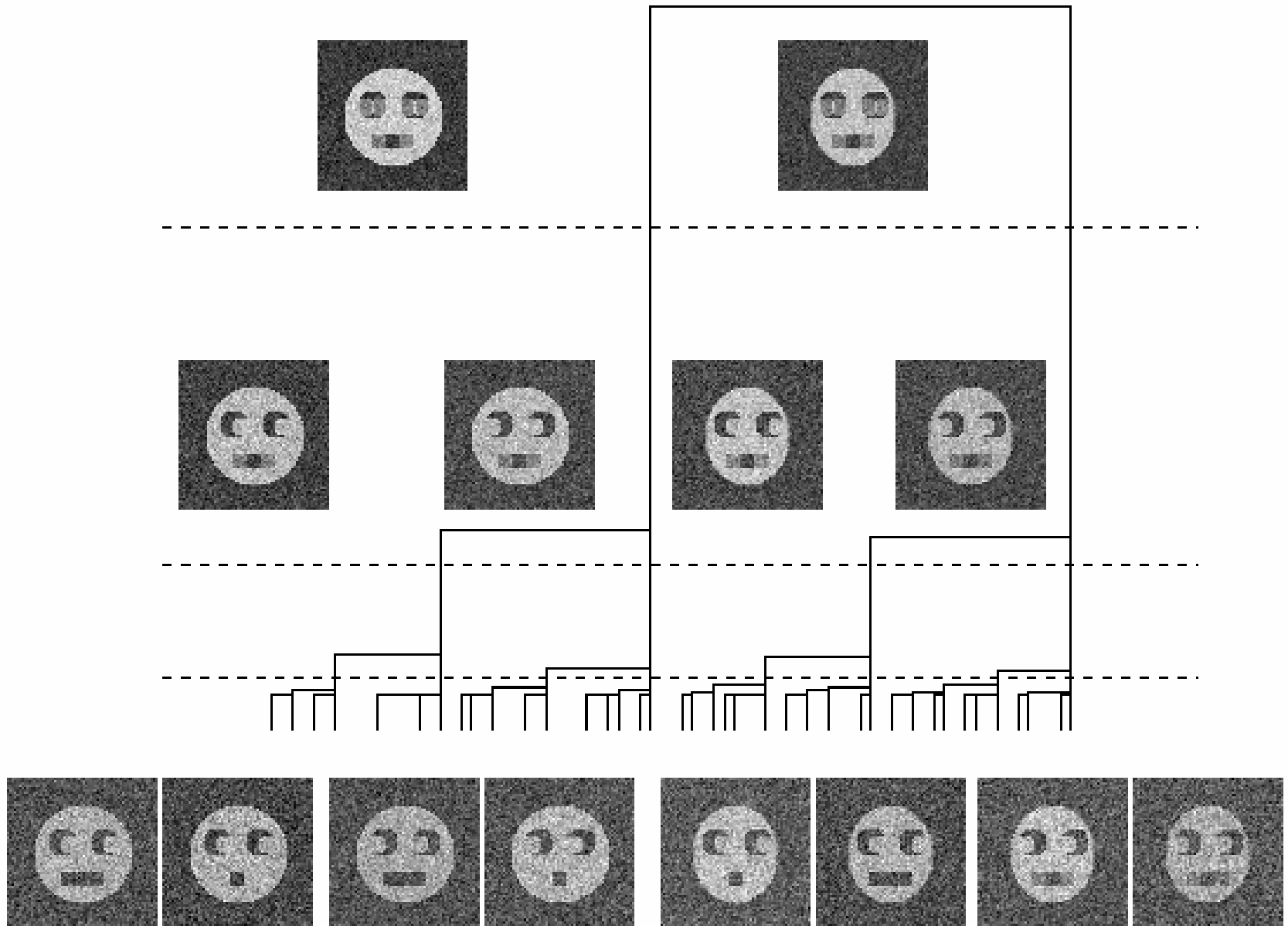
Avrg + F1+F2+F3



Hierarchical Ascendant Classification

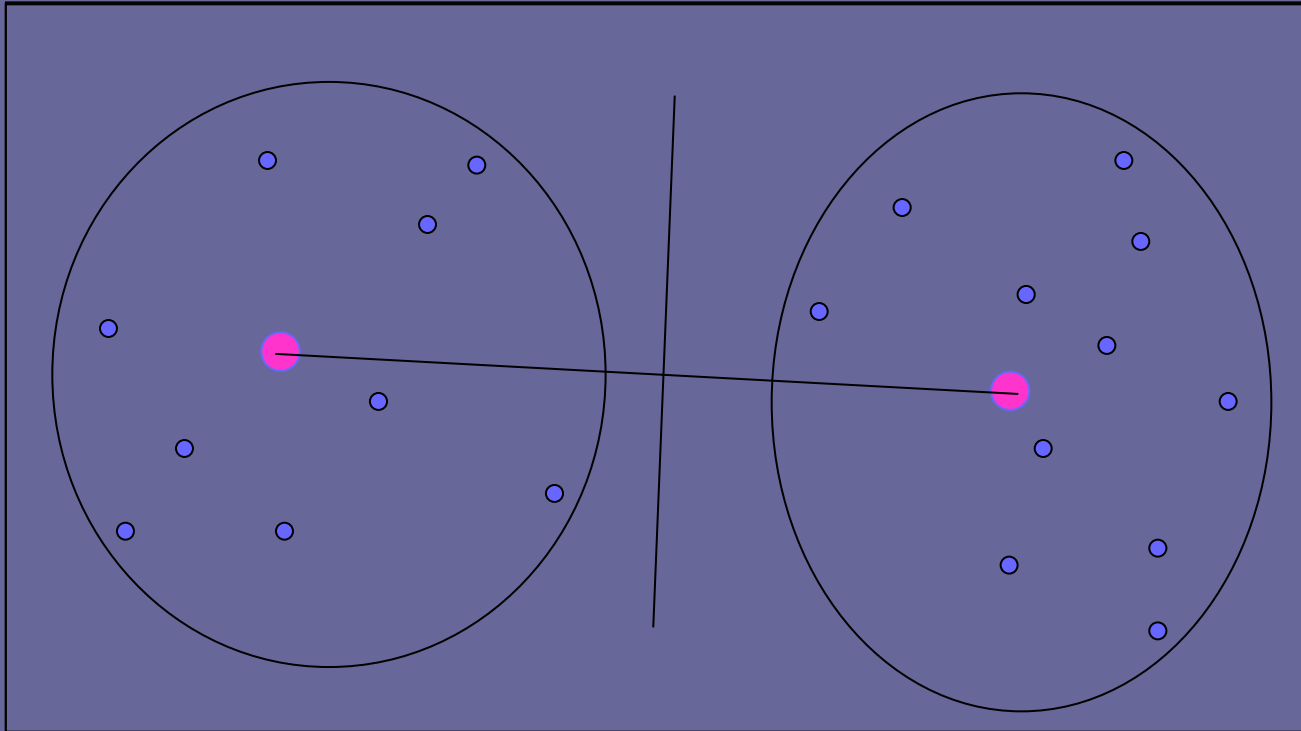


HIERARCHICAL ASCENDENT CLASSIFICATION



Partition methods : *e.g.* "Moving seeds" method

Diday E (1971) La methode des nuées dynamiques. *Rev. Stat. Appl.* 19, 19-34.



Stops when centers don't move from one step to the next
or after a given a selected number of iterations.

Overview: the necessary steps of a single-particle reconstruction

- 1) *Optical diffraction: quality control, defocus inventory of micrograph batch*
- 2) *Scanning of micrograph batch*
- 3) *Determine defoci, and define defocus groups*
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---Steve---Steve---Steve---Steve---Steve-----

- 7) *Refinement*
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Overview: the necessary steps of a single-particle reconstruction -- I

1) *Optical diffraction: quality control, defocus inventory of micrograph batch*

2) *Scanning of micrograph batch*

3) *Determine defoci, and define defocus groups*

4) *Pick particles*

(a) manual

(b) automated

5) *Determine particle orientation*

(a) unknown structure -- bootstrap

(i) random-conical (uses unsupervised classification)

(ii) common lines/ angular reconstitution (uses unsupervised classification)

(b) known structure

(i) reference-based (3D projection matching = supervised classification)

(ii) common lines/ angular reconstitution

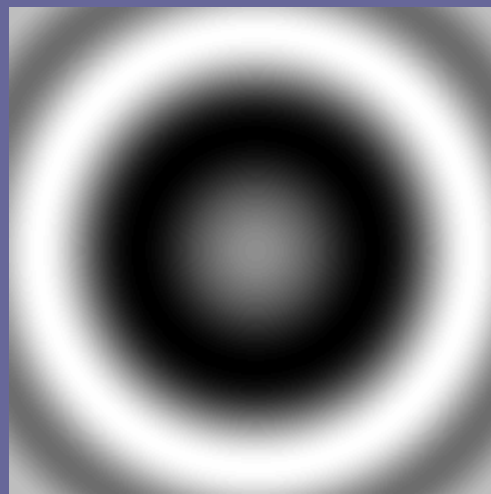
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original object

x

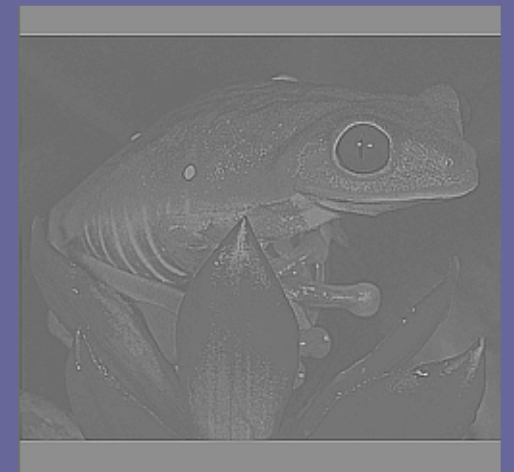
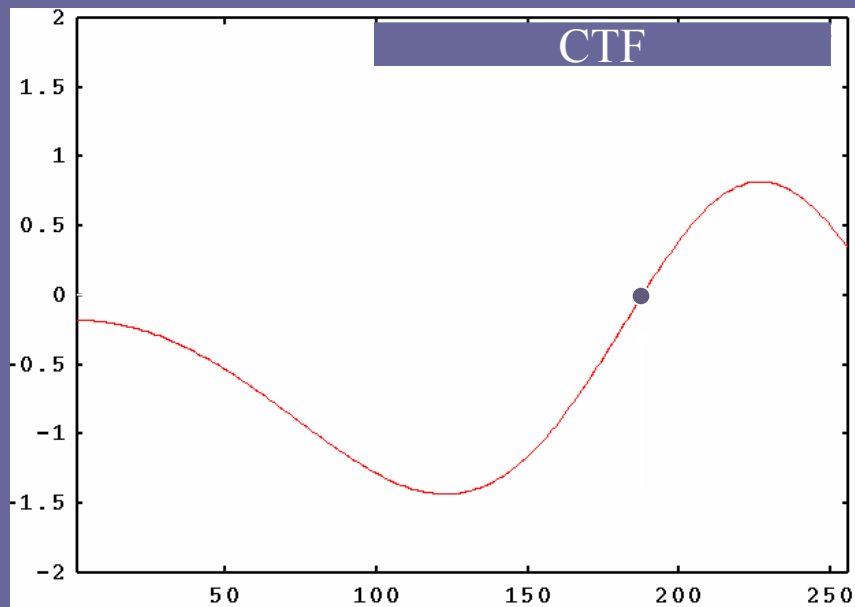


CTF for $\Delta z = 0.400 \mu\text{m}$

=



cryo-EM image

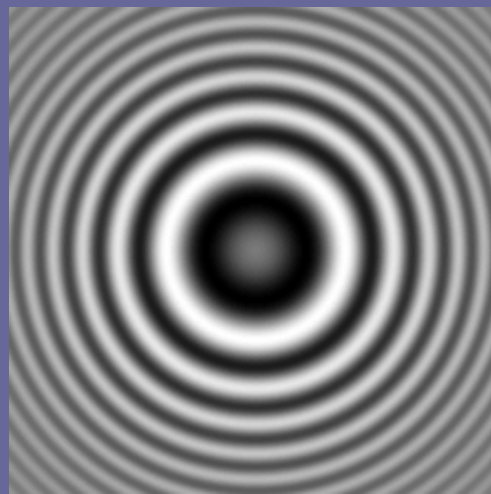


cryo-EM image,
contrast-inverted



original object

\times

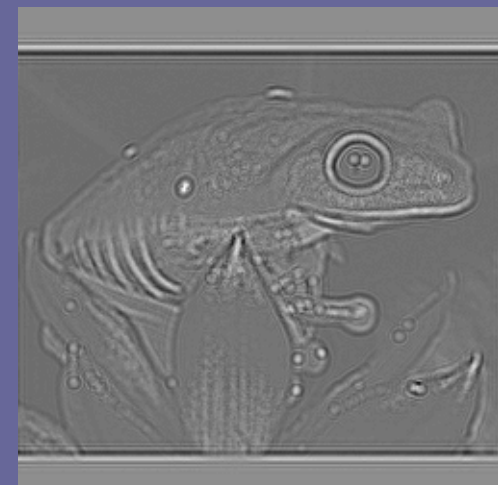
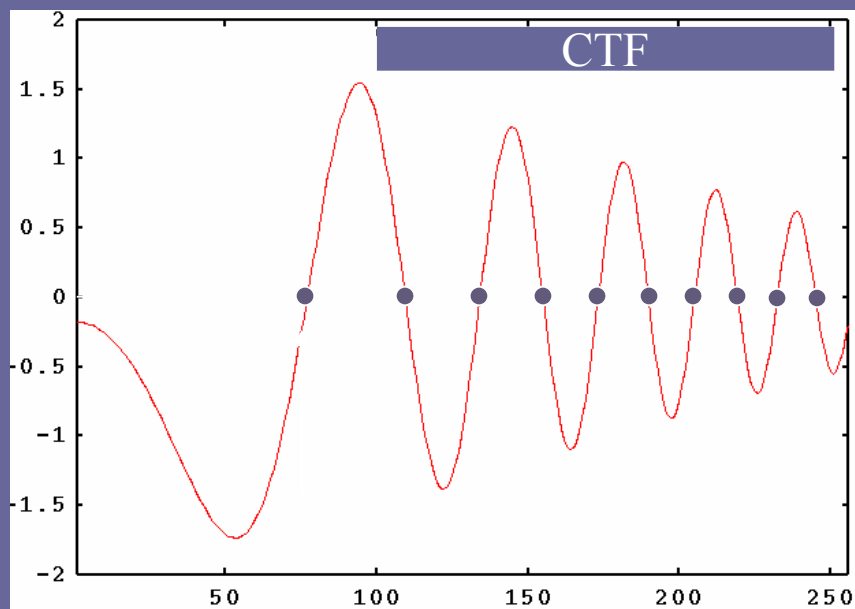


CTF for $\Delta z = 2.500 \mu\text{m}$

$=$



cryo-EM image



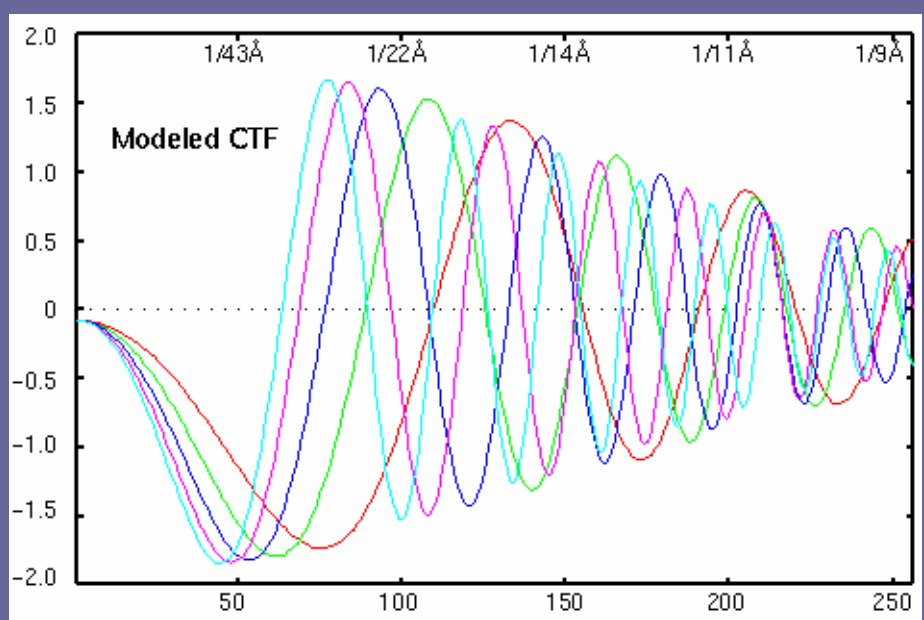
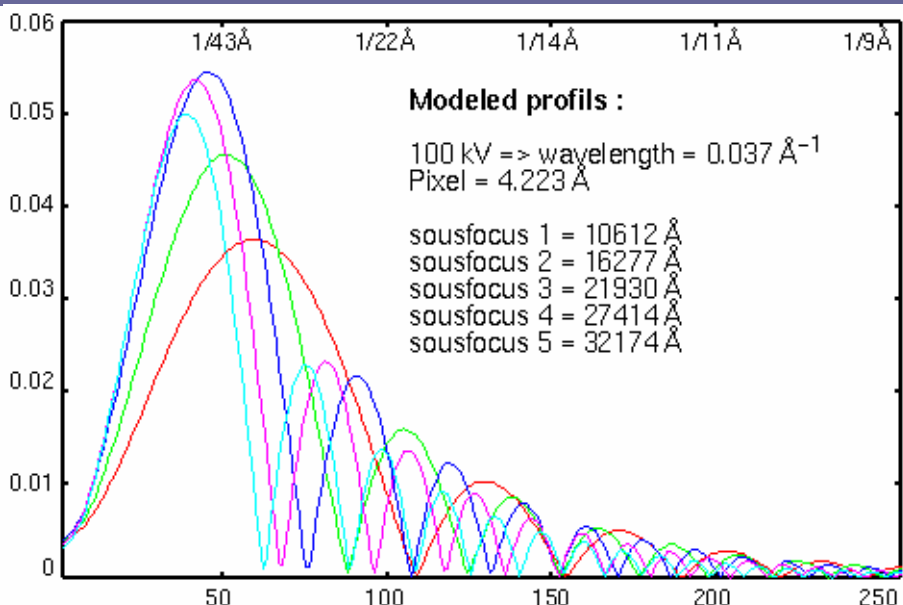
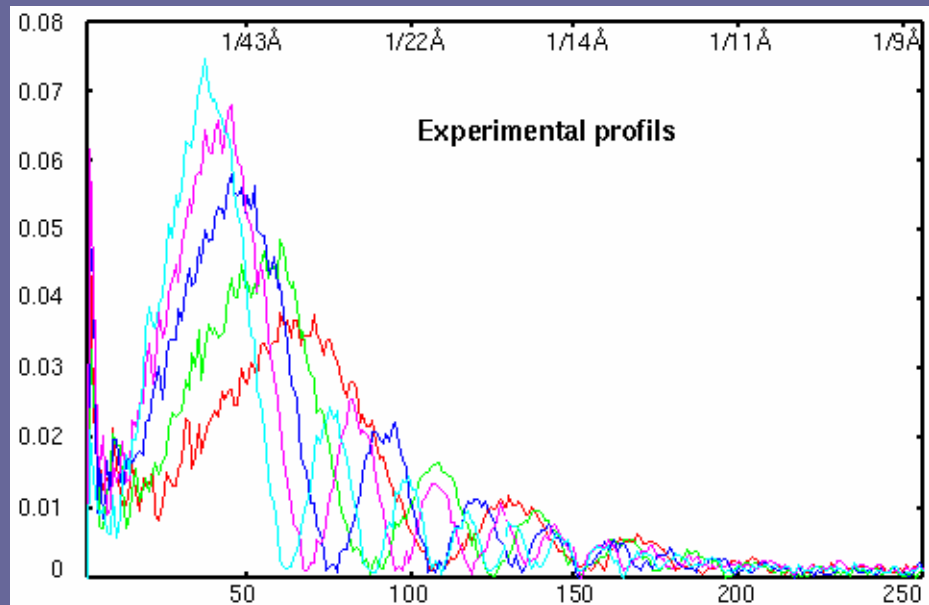
cryo-EM image,
contrast-inverted

Strategy for reconstruction from multiple defocus groups

- Coverage of large defocus range required
- Data collection must be geared toward covering range without major gap
- Characterizing all particles from the same micrograph by the same defocus is OK up to a resolution of $\sim 1/8 \text{ \AA}^{-1}$.

Sequence of steps:

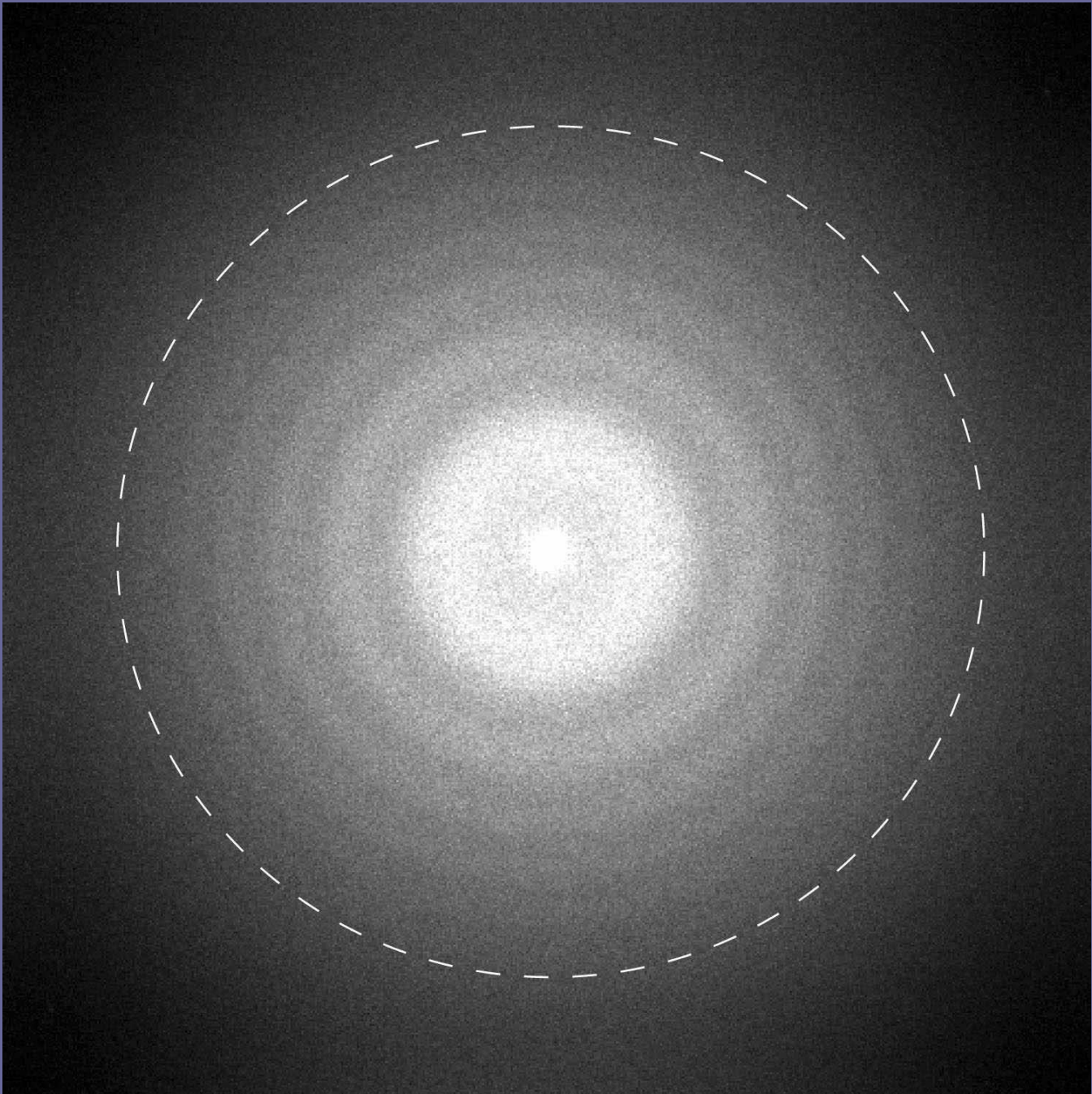
- 1) *Determine defocus for each micrograph*
- 2) *Define defocus groups, by creating supersets of particles from micrographs in a narrow range of defoci*
- 3) *Process particles separately, by defocus group, till the very end (3D reconstruction by defocus groups)*
- 4) *Compute merged, CTF-corrected reconstruction. E.g., Wiener filtering.*



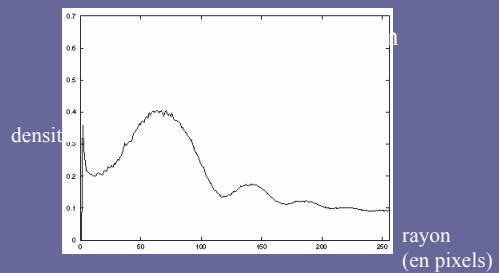
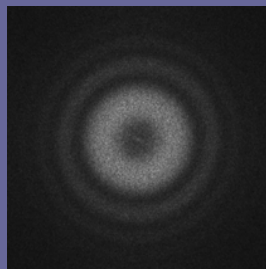
Computation of averaged power spectrum

For each micrograph ...

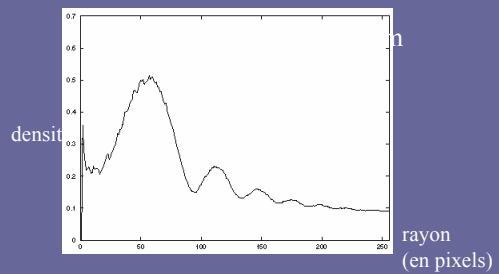
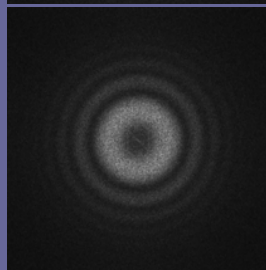
- 1) Divide field into overlapping subfields of $\sim 512 \times 512$
- 2) Compute FFT for each subfield
- 3) Compute $|F(\mathbf{k})|^2$ for each subfield
- 4) Form average over $|F(\mathbf{k})|^2$ of all subfields \Rightarrow averaged, smoothed power spectrum
- 5) Take square root of result \Rightarrow “power spectrum” with reduced dynamic range
- 6) Form azimuthal average \Rightarrow 1D profile, characteristic for the micrograph, ready to be compared with CTF



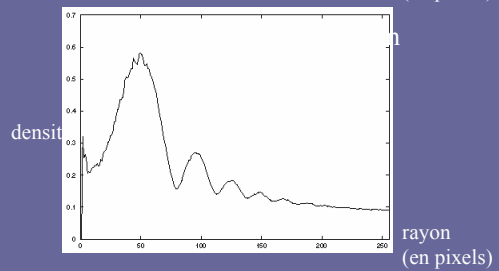
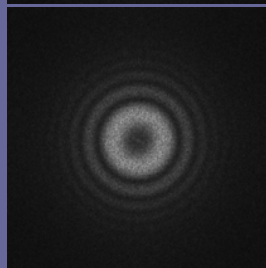
A



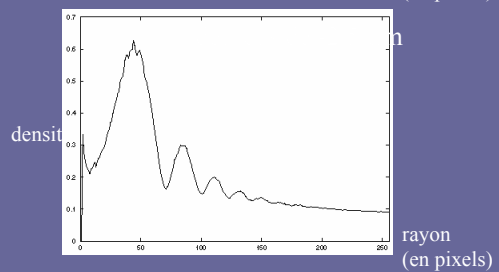
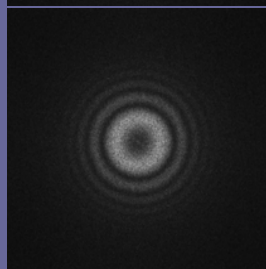
B



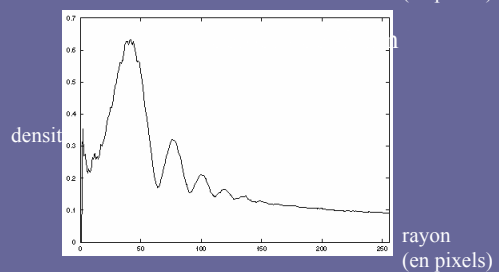
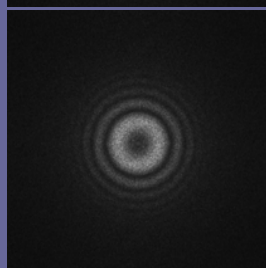
C



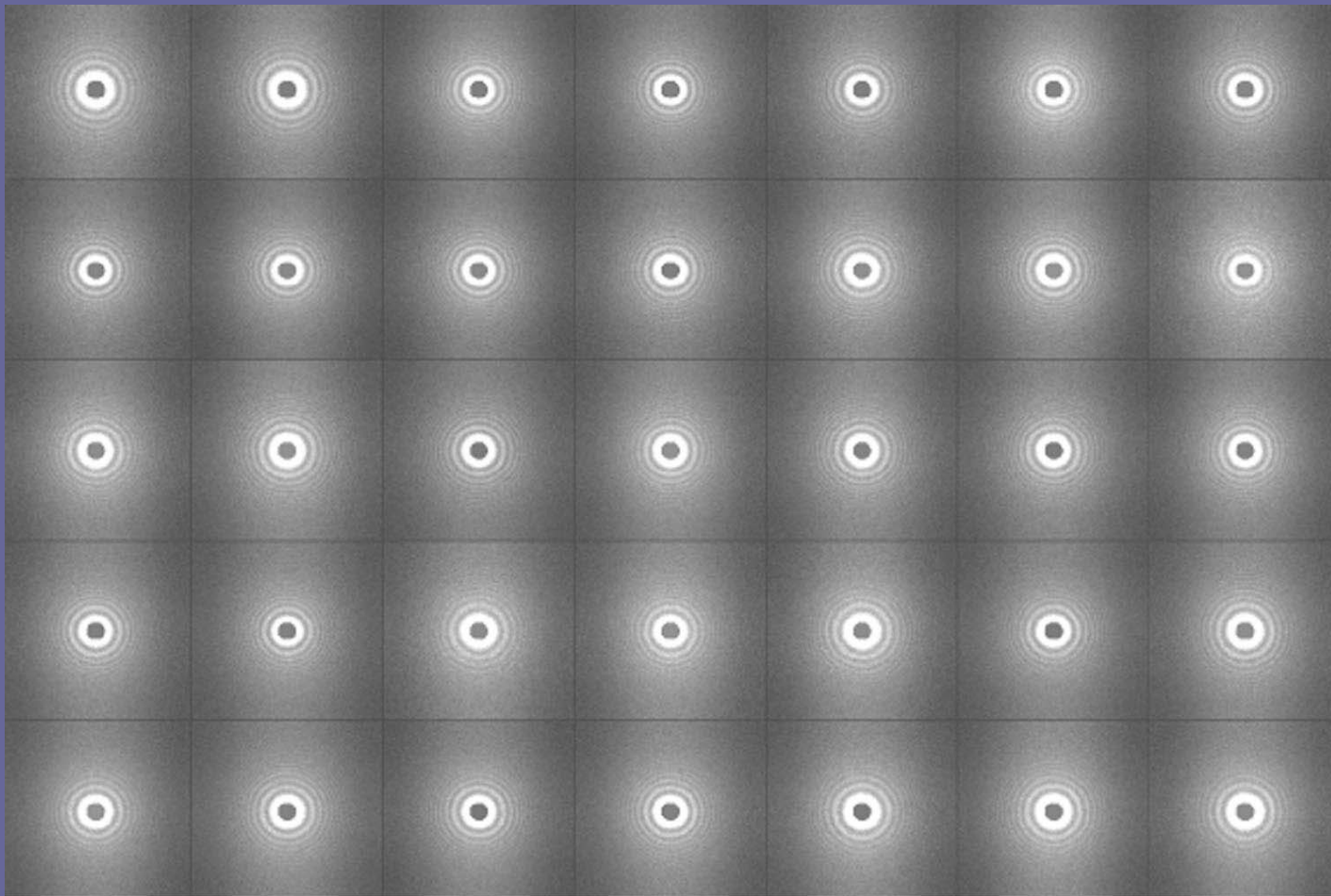
D

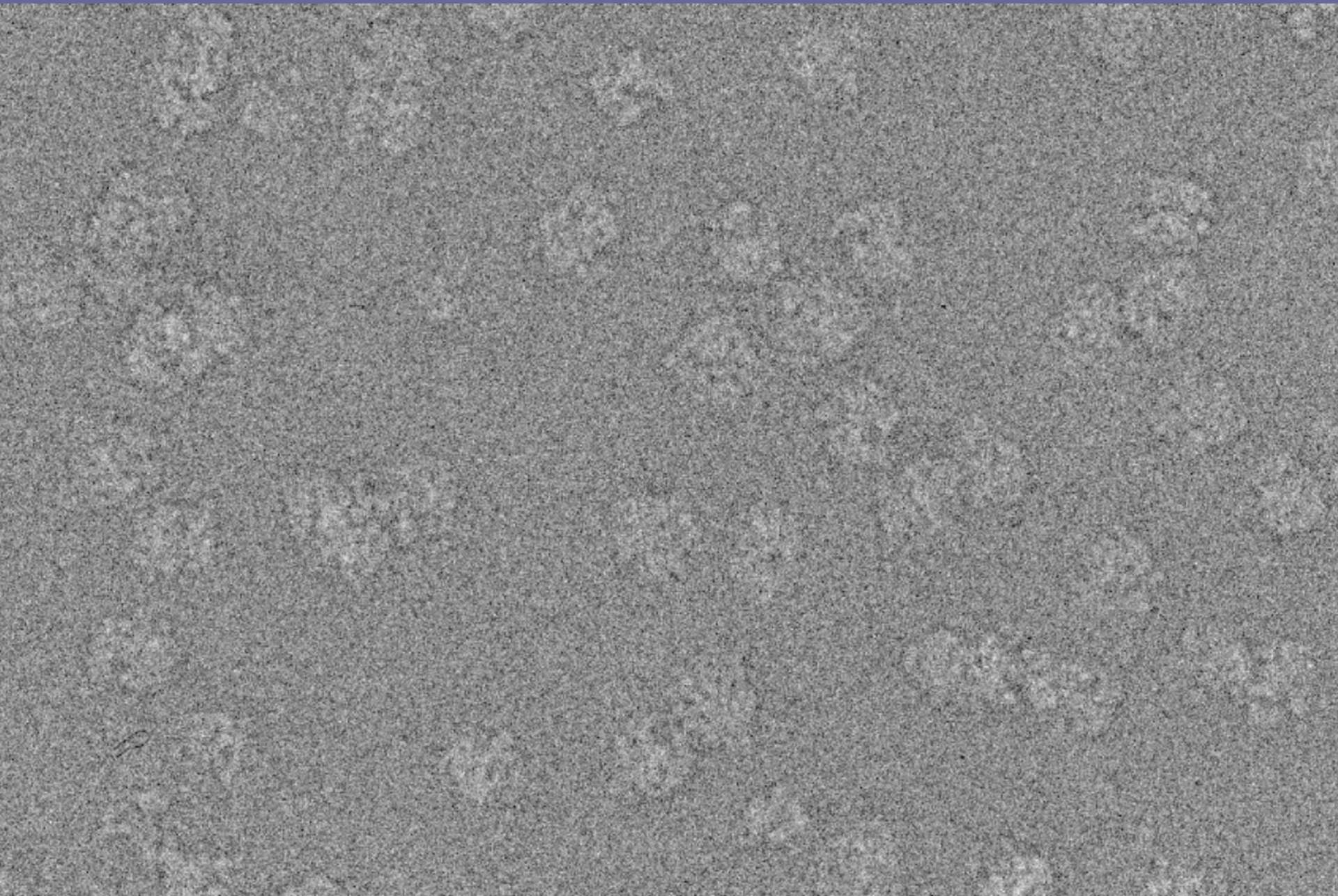


E



Gallery of power spectra from different micrographs





Overview: the necessary steps of a single-particle reconstruction -- I

1) Optical diffraction: quality control, defocus inventory of micrograph batch

2) Scanning of micrograph batch

3) Determine defoci, and define defocus groups

4) Pick particles

(a) manual

(b) automated

5) Determine particle orientation

(a) unknown structure -- bootstrap

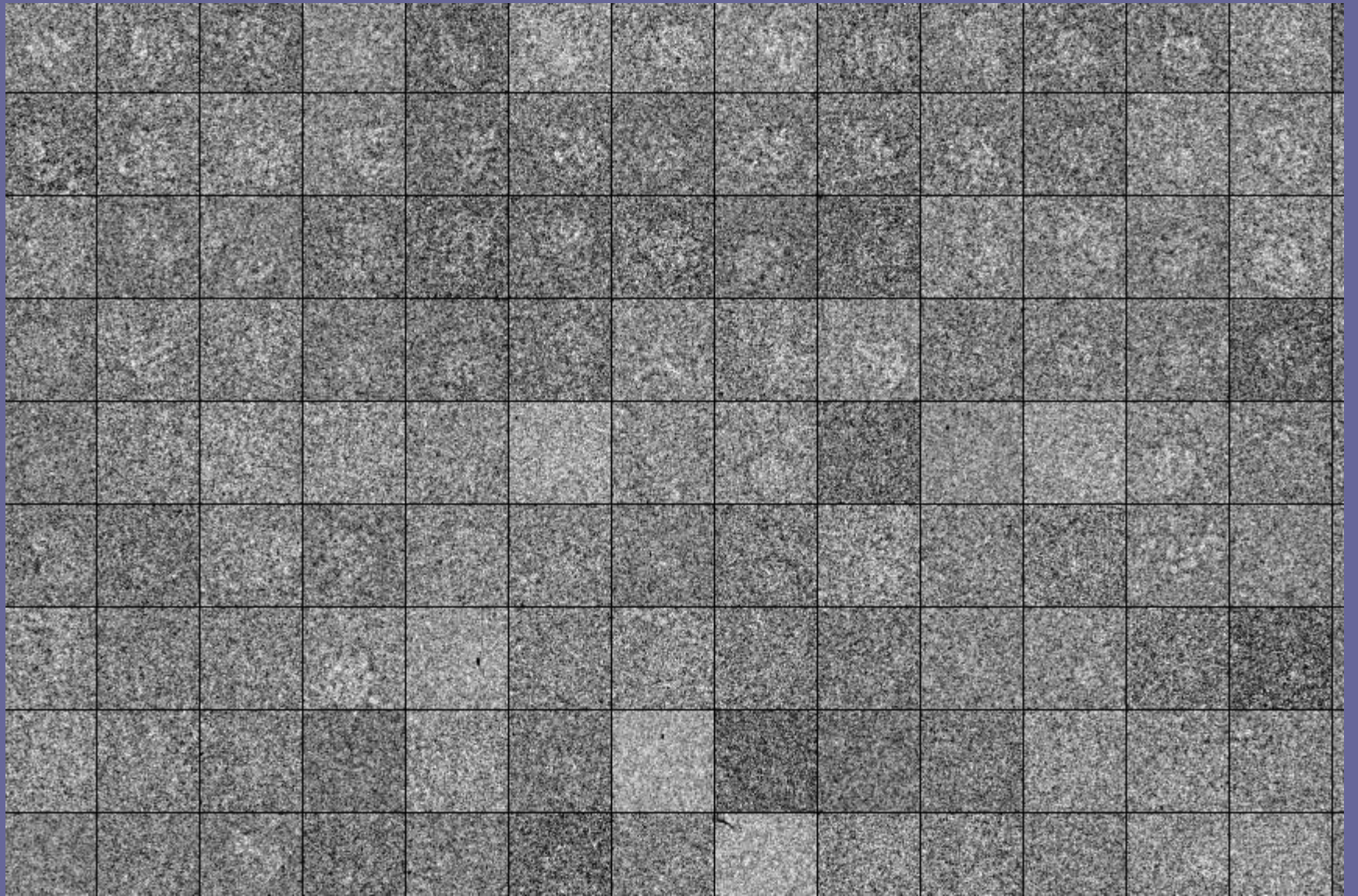
(i) random-conical (uses unsupervised classification)

(ii) common lines/ angular reconstitution (uses unsupervised classification)

(b) known structure

(i) reference-based (3D projection matching = supervised classification)

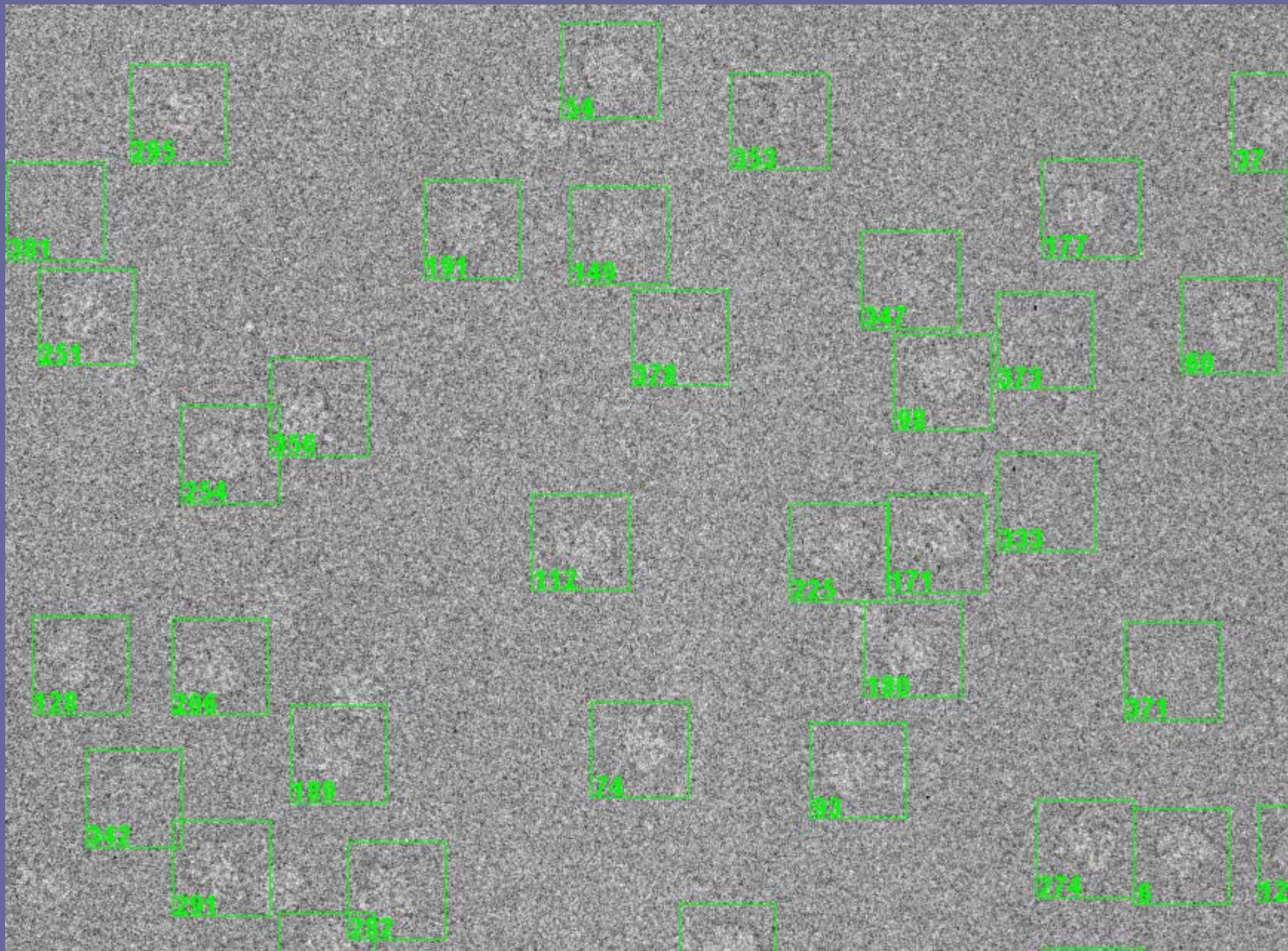
(ii) common lines/ angular reconstitution



Automated particle picking, CCF-based, with local normalization

- (i) Define a reference (e.g., by averaging projections over full Eulerian range);
- (ii) Paste reference into array with size matching the size of the micrograph;
- (iii) Compute CCF via FFT;
- (iv) Compute locally varying variance of the micrograph via FFT (Roseman, 2003);
- (v) ***“Local CCF” = CCF/local variance***
- (vi) Peak search;
- (vii) Window particles ranked by peak size;
- (viii) Fast visual screening.

Advantage of local CCF: avoid problems from background variability



295

24

253

37

231

191

199

171

251

247

69

255

279

95

373

254

111

223

171

333

175

209

136

271

247

199

74

63

291

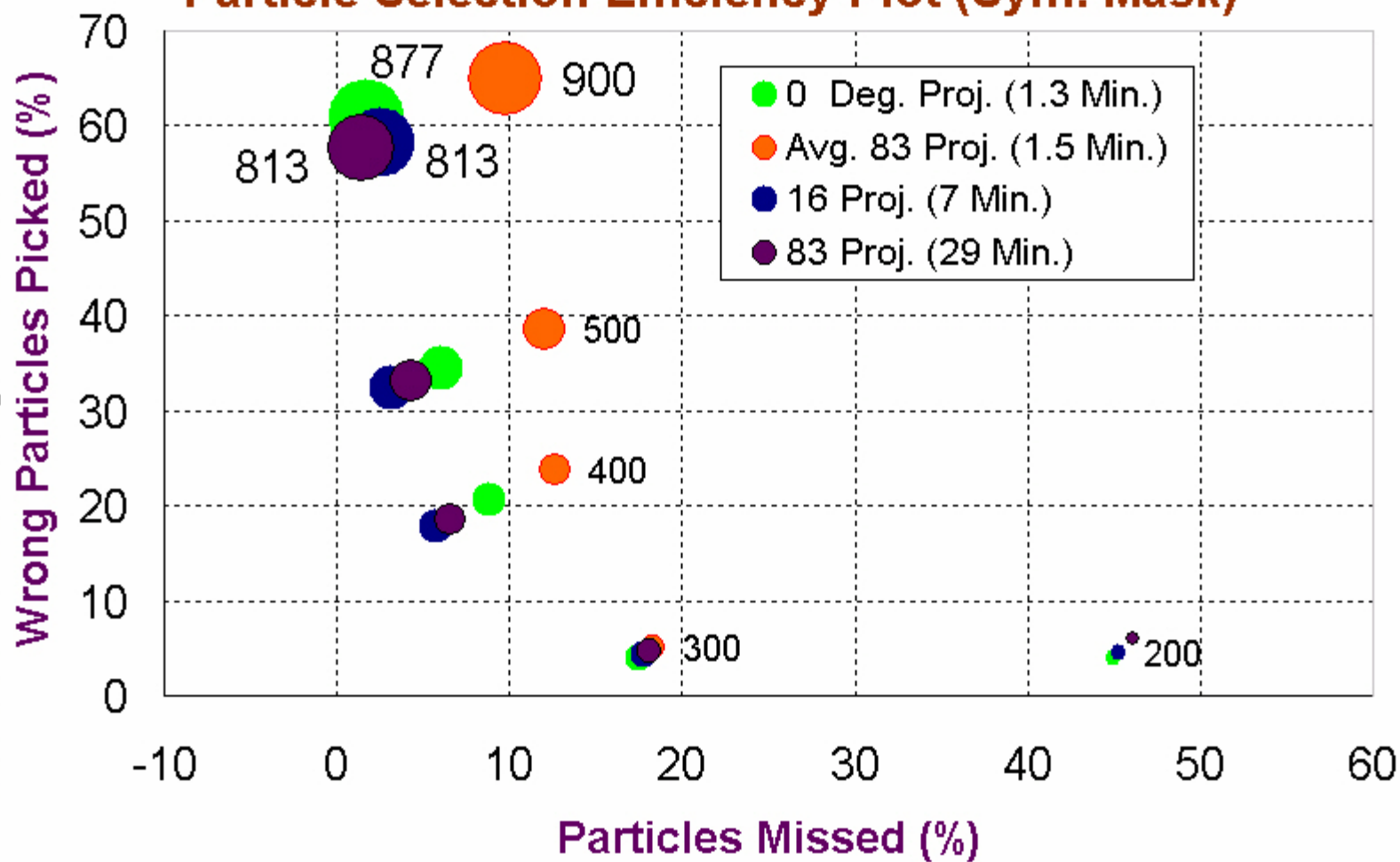
282

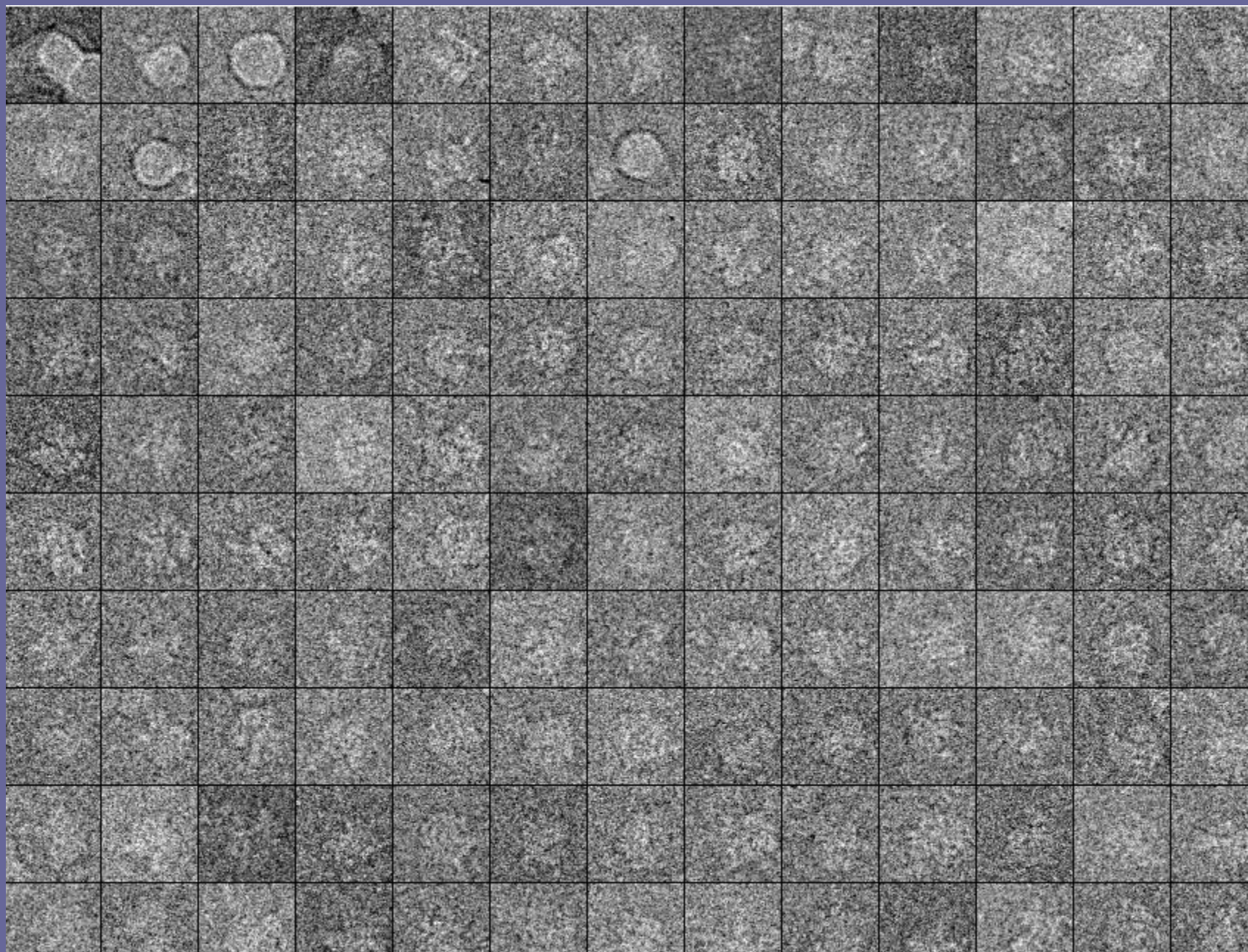
274

8

32

Particle Selection Efficiency Plot (Sym. Mask)





Overview: the necessary steps of a single-particle reconstruction -- I

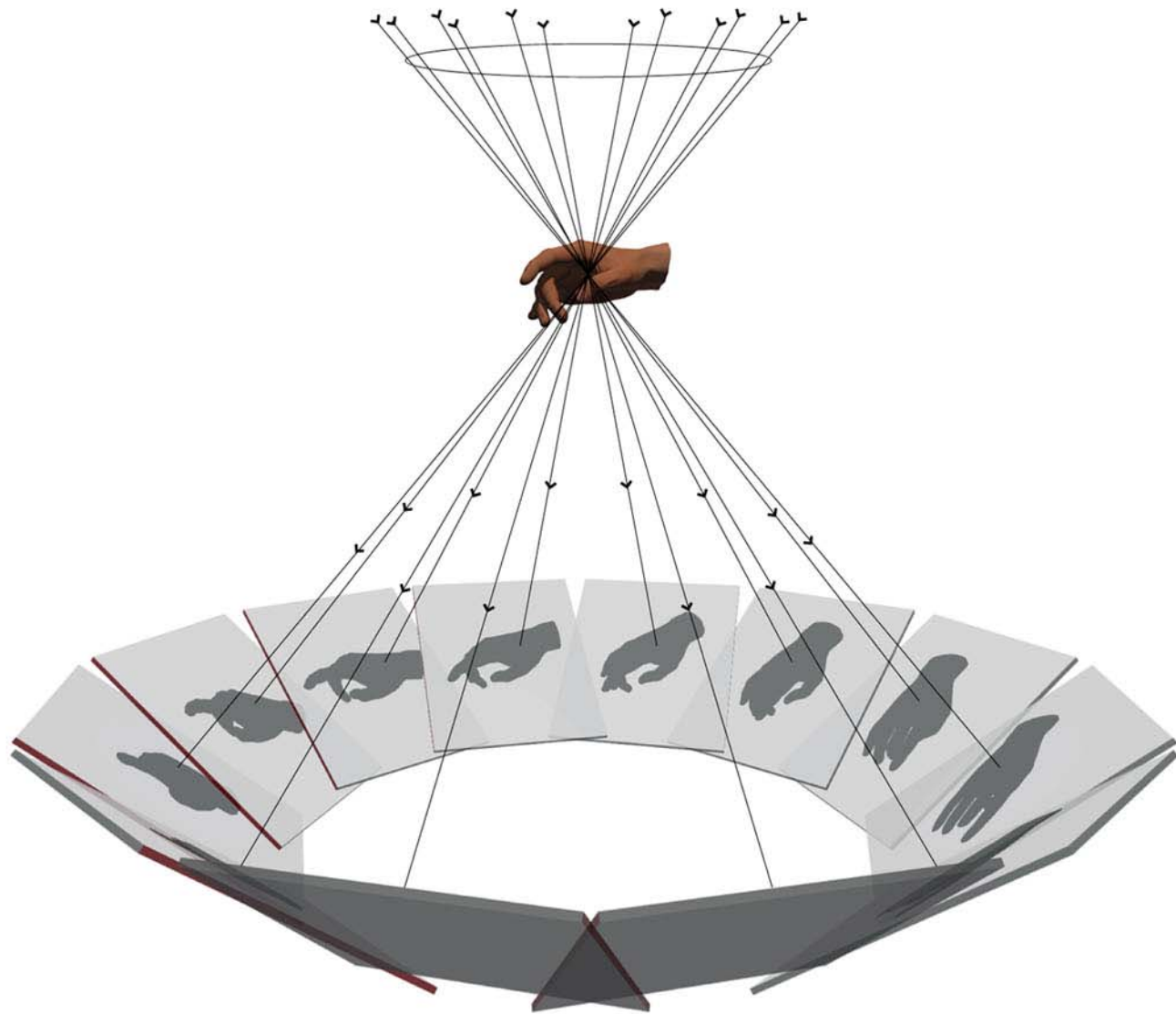
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Random-conical reconstruction

- Premise: all particles exhibit the same view
- Take same field first at theta ~ 50 degrees, then at 0 degrees [in this order, to minimize dose]
- Display both fields side by side
- Pick each particle in both fields
- Align particles from 0-degree field
- This yields azimuths, so that data can be put into the conical geometry*
- Assign azimuths and theta to the tilted particles
- Proceed with 3D reconstruction







Random-conical reconstruction

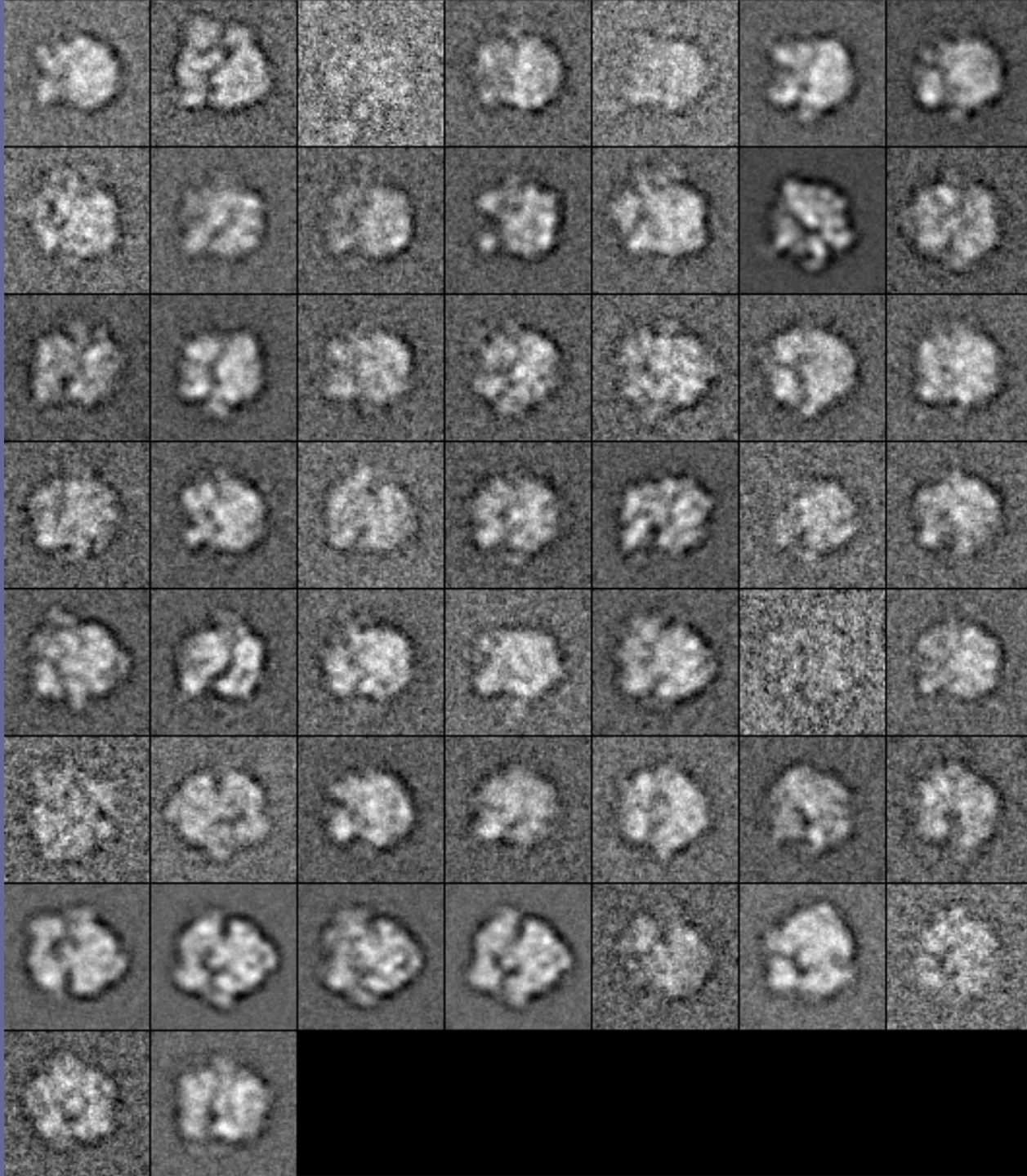
Problems to be solved:

1) Find a subset (view class) of particles that lie in the same orientation on the grid

answer: unsupervised classification of 0-degree particles

2) Missing cone problem

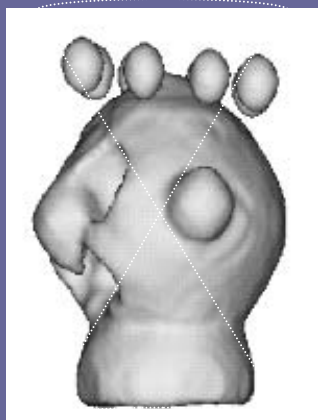
answer: do several random conical reconstructions, each from a different subset (view class), find relative orientations, then make reconstruction from merged projections set.



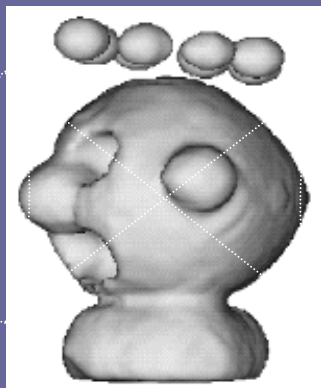
Missing-cone artifacts

Reconstruction
Using top view

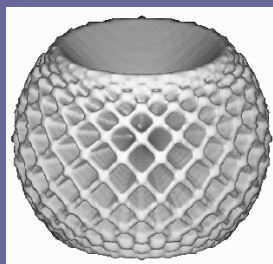
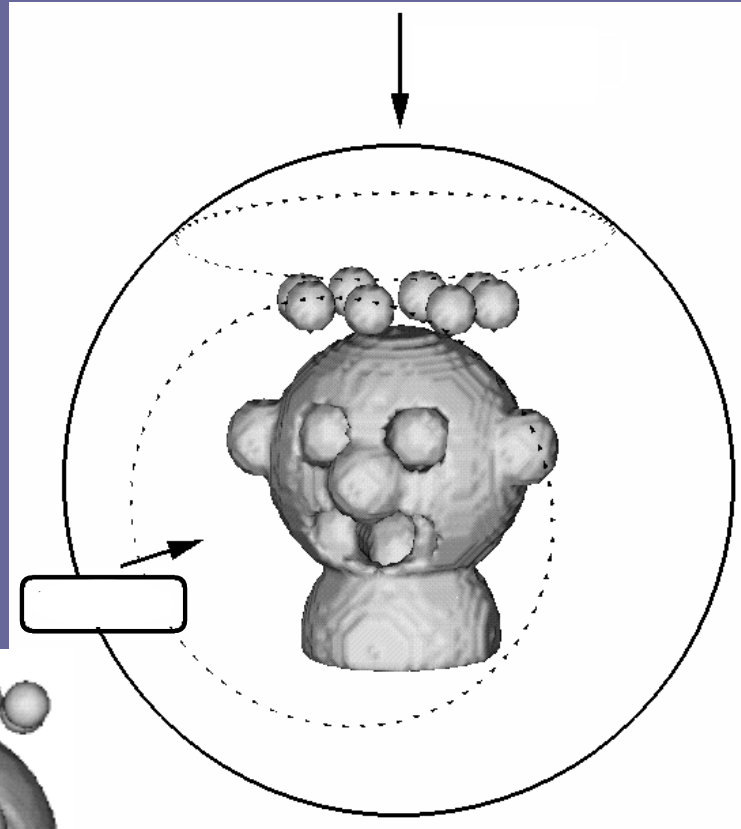
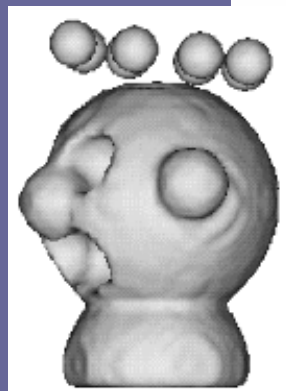
Reconstruction
Using side view



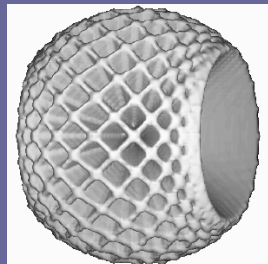
+



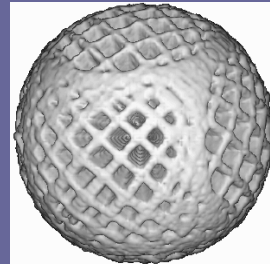
=



+



=



Overview: the necessary steps of a single-particle reconstruction -- I

1) Optical diffraction: quality control, defocus inventory of micrograph batch

2) Scanning of micrograph batch

3) Determine defoci, and define defocus groups

4) Pick particles

(a) manual

(b) automated

5) Determine particle orientation

(a) unknown structure -- bootstrap

(i) random-conical (uses unsupervised classification)

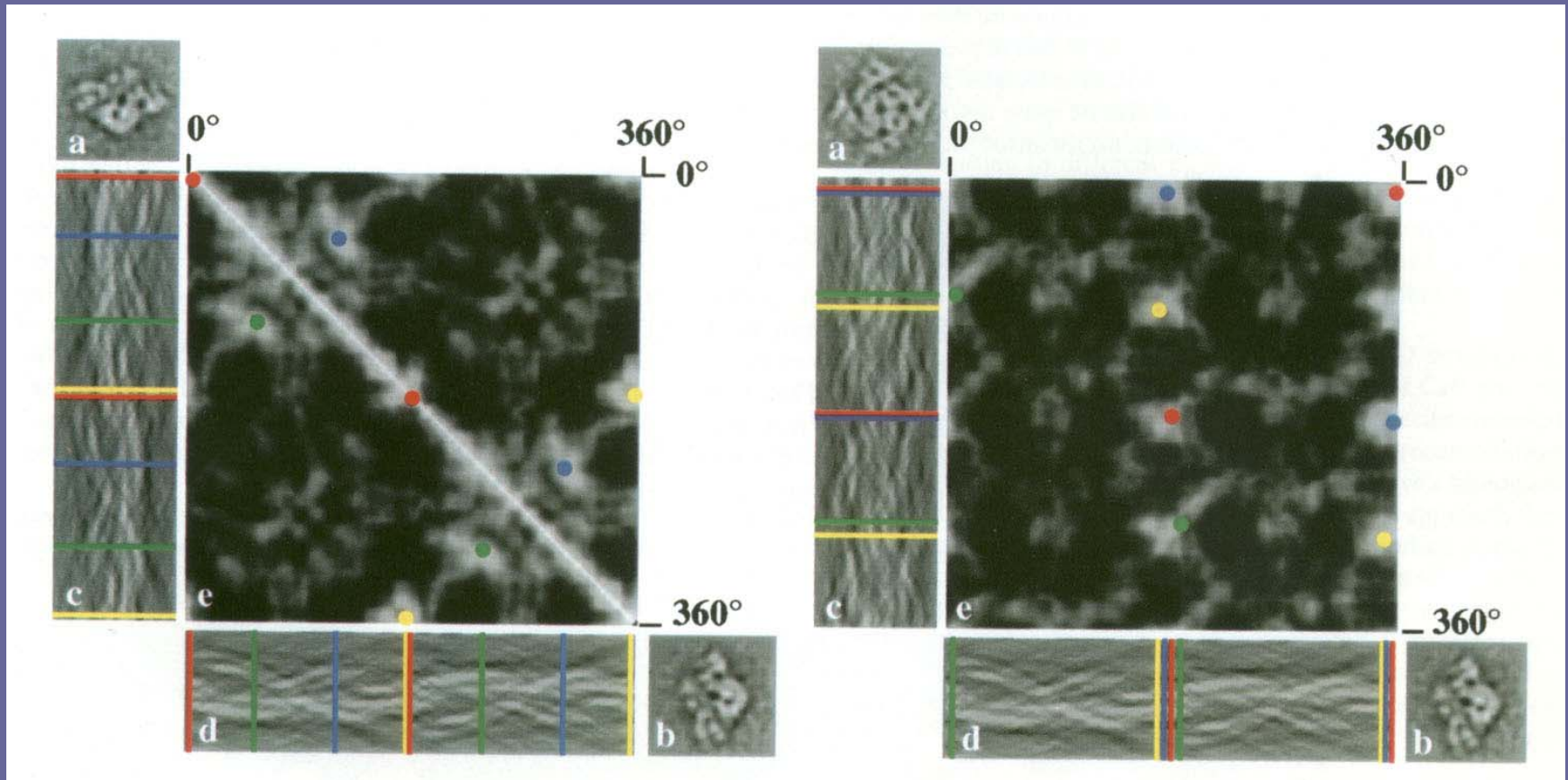
(ii) common lines/ angular reconstitution (uses unsupervised classification)

(b) known structure

(i) reference-based (3D projection matching = supervised classification)

(ii) common lines/ angular reconstitution

Determination of relative orientations by common lines



Serysheva et al. (1995) Nature Struct. Biol. 2: 18-24.

Common lines/ angular reconstitution

- 1) Unsupervised classification, to determine classes of particles exhibiting the same view
- 2) Average images in each class → class averages
- 3) Determine common lines between class averages
stepwise (van Heel, 1967)
simultaneously (Penczek et al., 1996)

Issues:

- unaveraged images are too noisy
- resolution loss due to implicit use of view range
- handedness not defined – tilt or prior knowledge needed

Overview: the necessary steps of a single-particle reconstruction -- I

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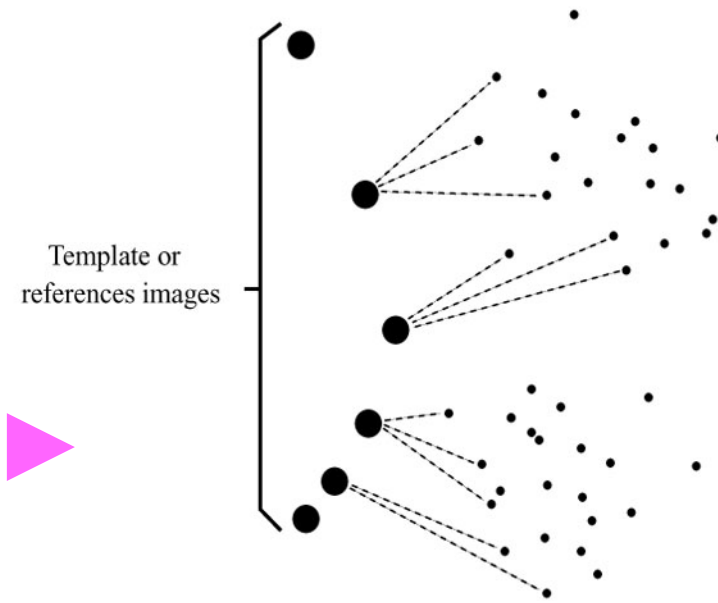
(ii) common lines/ angular reconstitution (uses unsupervised classification)

(b) known structure

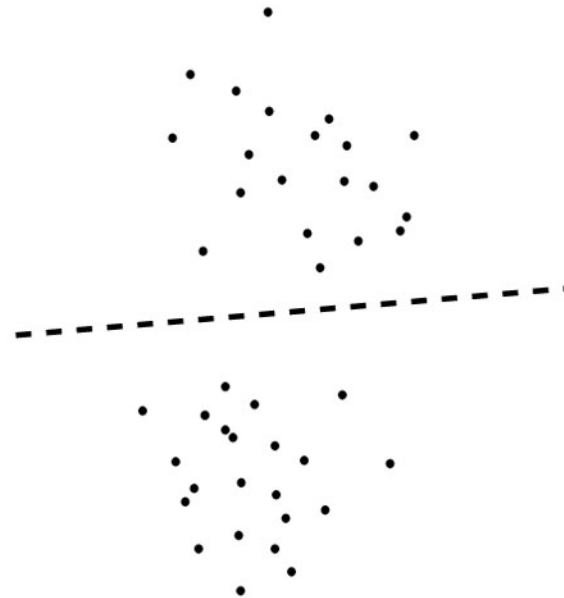
(i) reference-based (3D projection matching = supervised classification)

(ii) common lines/ angular reconstitution

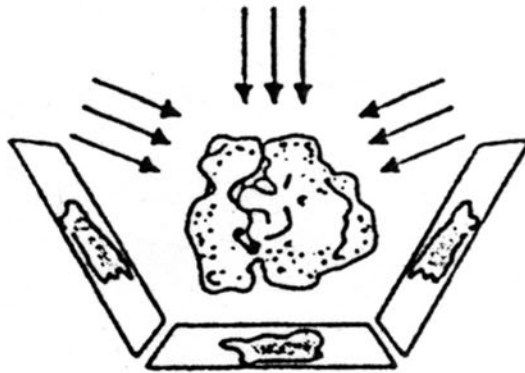
Supervised Classification



Unsupervised Classification



Systematically generated projections
of existing reconstruction



Stack of projections



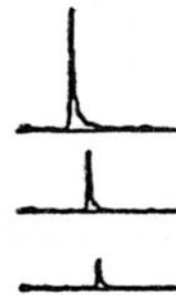
X

Experimental
projection



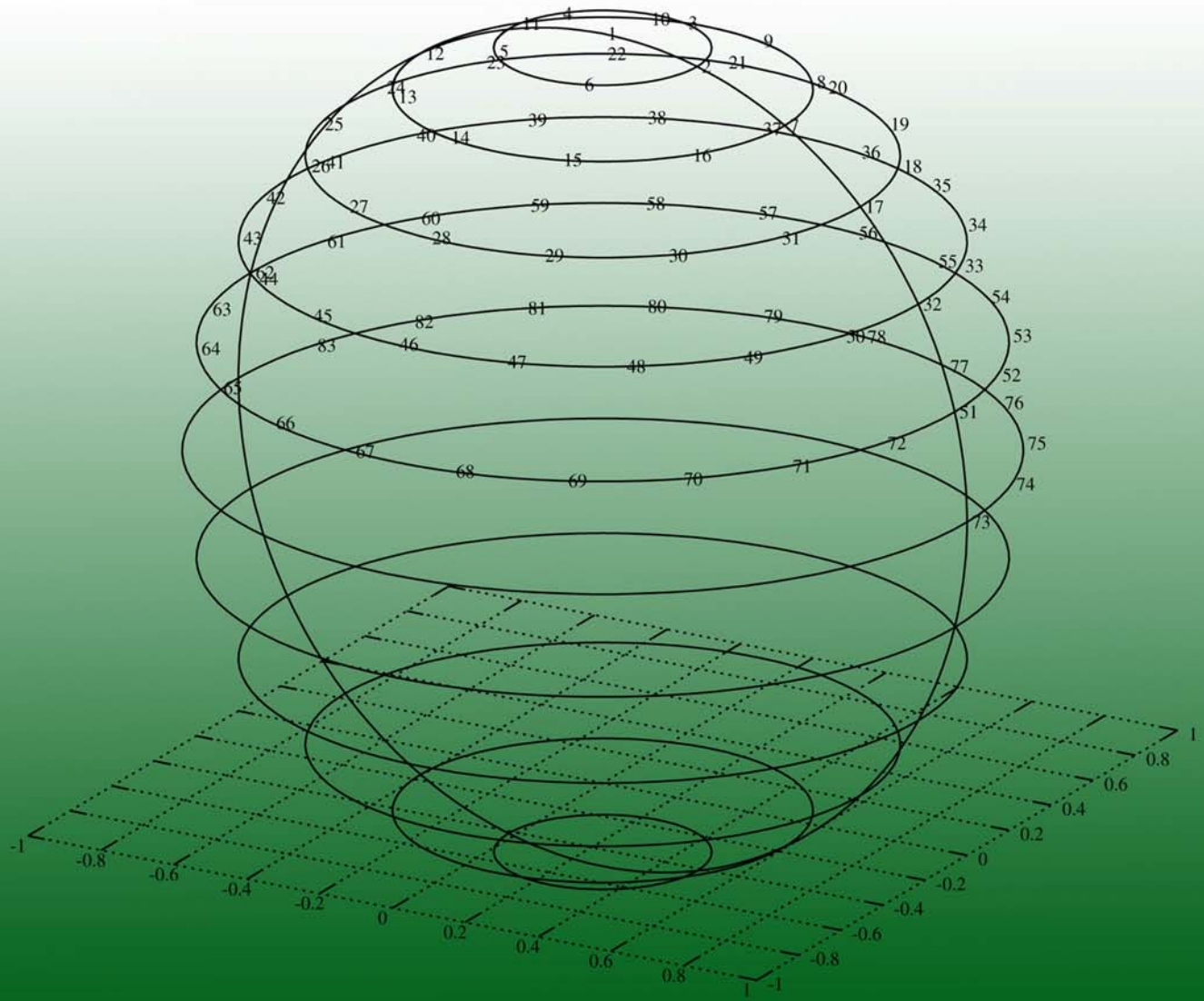
=

Stack of rotational
CCF's

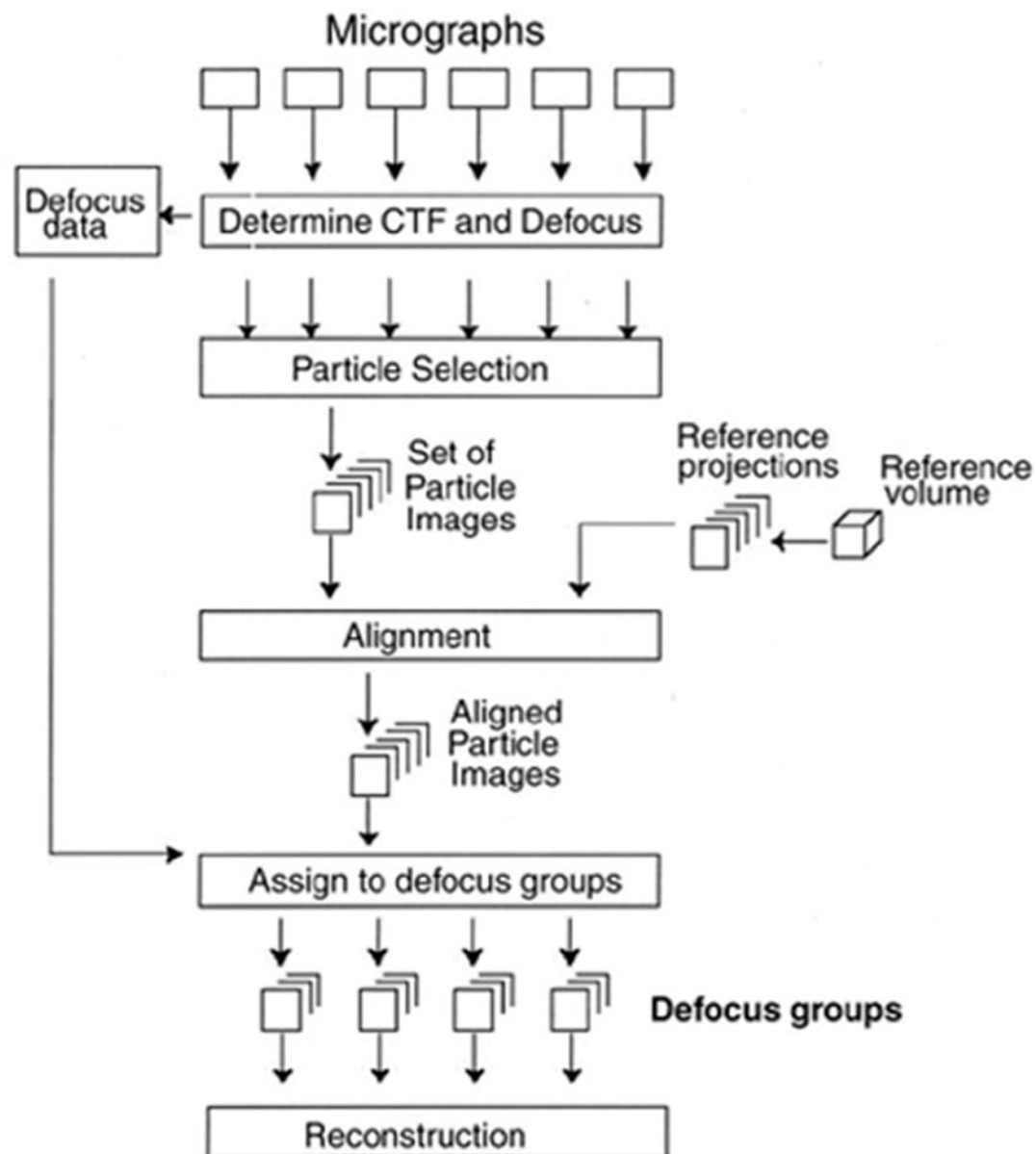


max
→
CCF
coeff's

3 Eulerian
angles



Reference-based Reconstruction



Overview: the necessary steps of a single-particle reconstruction -- II

- 6) *3D reconstruction by defocus group*
 - (a) Fourier interpolation
 - (b) Weighted back-projection
 - (c) Iterative algebraic reconstruction
 - (d) Conjugate gradient

---Steve---Steve---Steve---Steve---Steve-----

- 7) *Refinement*
 - given an initial 3D reference,
iterate the steps {3D projection matching + reconstruction}
 - beware of problem of reference-dependence
- 8) *CTF correction*
- 9) *Validation*
- 10) *Interpretation*: segmentation, docking, etc.

3D reconstruction by defocus group

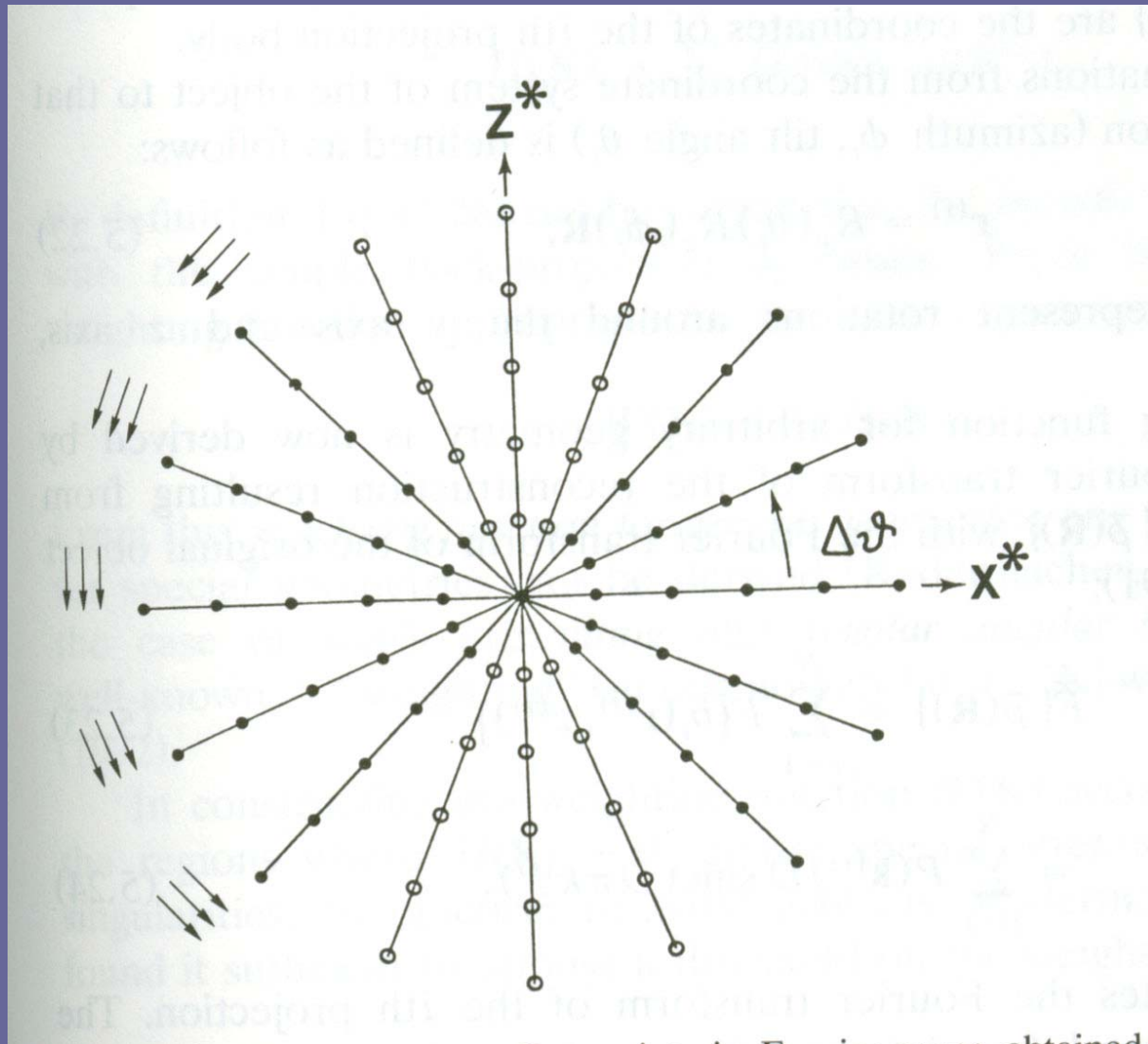
(a) Fourier interpolation

- (b) Weighted back-projection
- (c) Iterative algebraic reconstruction
- (d) Conjugate gradient

Obtain samples on a regular Cartesian grid in 3D Fourier space by interpolation between Fourier values on oblique 2D grids (central sections) running through the origin, each grid corresponding to a projection.

Speed (high) versus accuracy (low). Can be used in the beginning phases of a reconstruction project. However, see new development by Pawel Penczek!

Sample points of adjacent projections are increasingly sparse as we go to higher resolution



3D reconstruction by defocus group

(a) Fourier interpolation

(b) Weighted back-projection

(c) Iterative algebraic reconstruction

(d) Conjugate gradient

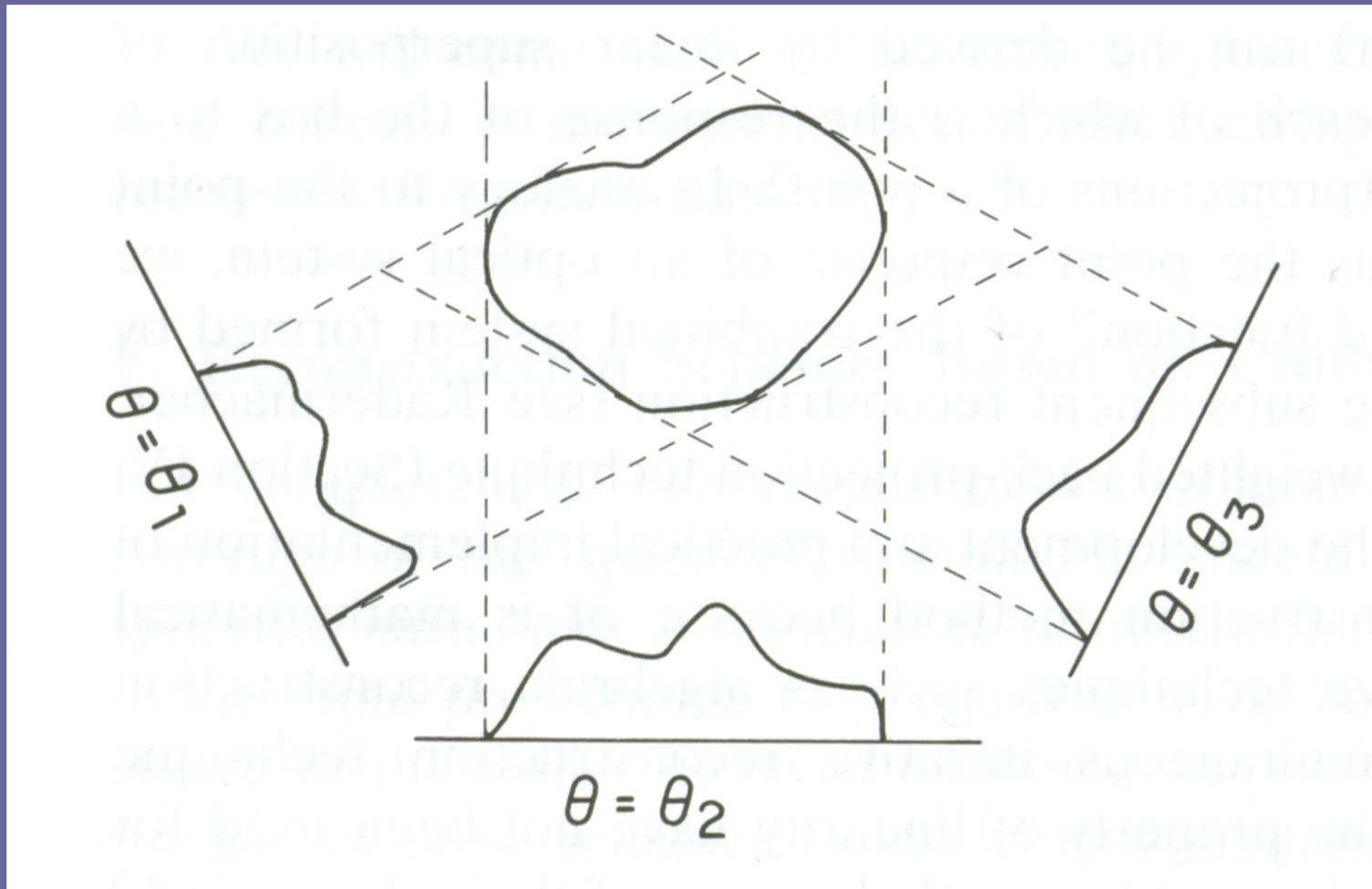
(1) Simple back-projection: Sum over “back-projection bodies”, each obtained by “smearing out” a projection in the viewing direction.

(2) Weighted back-projection: as (1), but “weight” the projections first by multiplying their Fourier transforms with $|K|$ (R^* weighting, in X-ray terminology), then inverting the Fourier transform.

(3) For general geometries, the weighting function is more complicated, and has to be computed every time.

• Weighted back-projection is fast, but does not yield the “smoothest” results. It may show strong artifacts from angular gaps.

Principle of back-projection



3D reconstruction by defocus group

(a) Fourier interpolation

(b) Weighted back-projection

(c) Iterative algebraic reconstruction

(d) Conjugate gradient

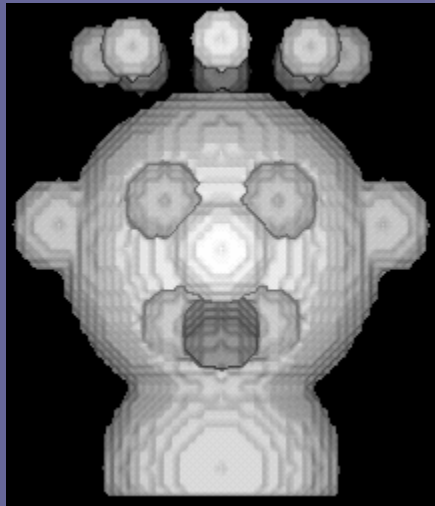
The discrete algebraic projection equation is satisfied, one angle at a time, by adjusting the densities of a starting volume. As iterations proceed, each round produces a better approximation of the object.

The algorithm comes in many variants. It allows constraints to be easily implemented.

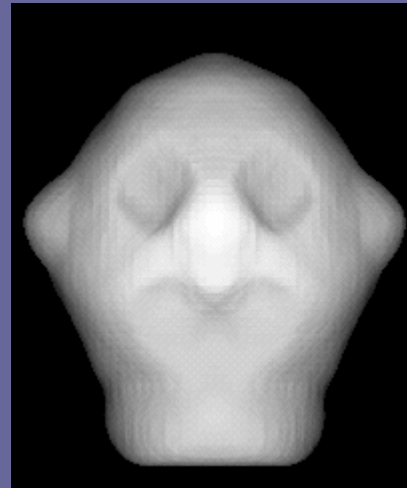
It produces a very smooth reconstruction, and is less affected by angular gaps.

Comparison of some reconstruction algorithms

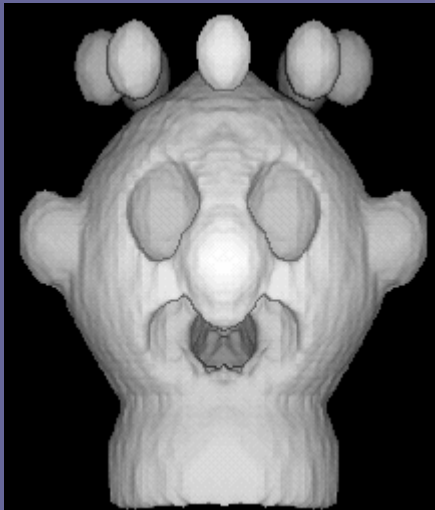
Original object



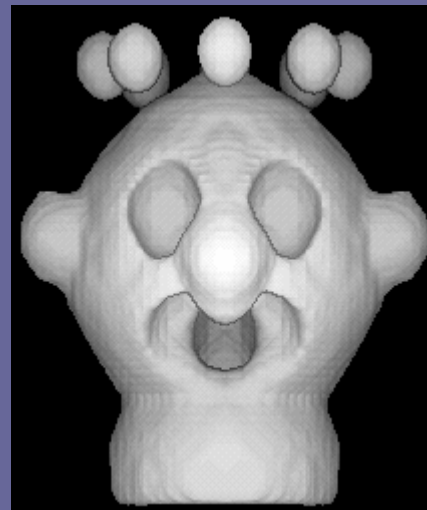
Simple back-projection



Weighted back-projection



Iterative algebraic reconstruction

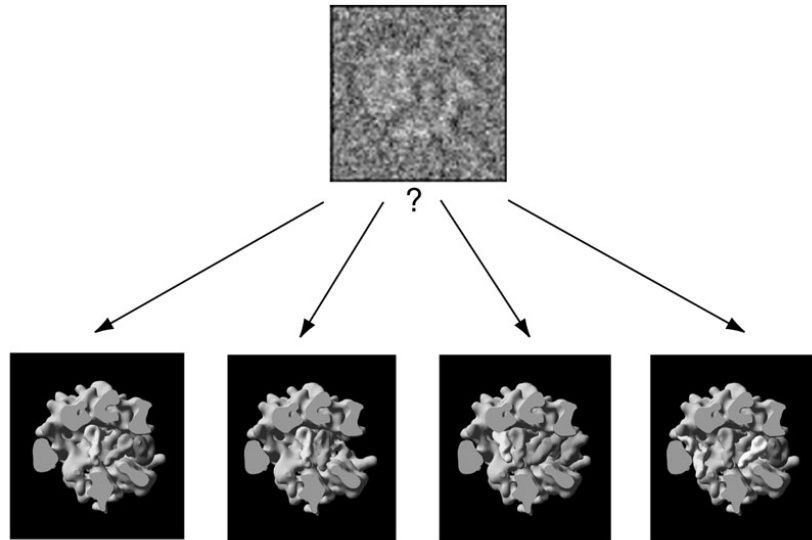


Sources for limited resolution

- 1) Instrumental: partial coherence (envelope function)
- 2) Particles with different height all considered having same defocus (envelope function)
- 3) Numerical: interpolations, inaccuracies
- 4) Failure to exhaust existing information
- 5) Conformational diversity

Conformational diversity: heterogeneous particle population

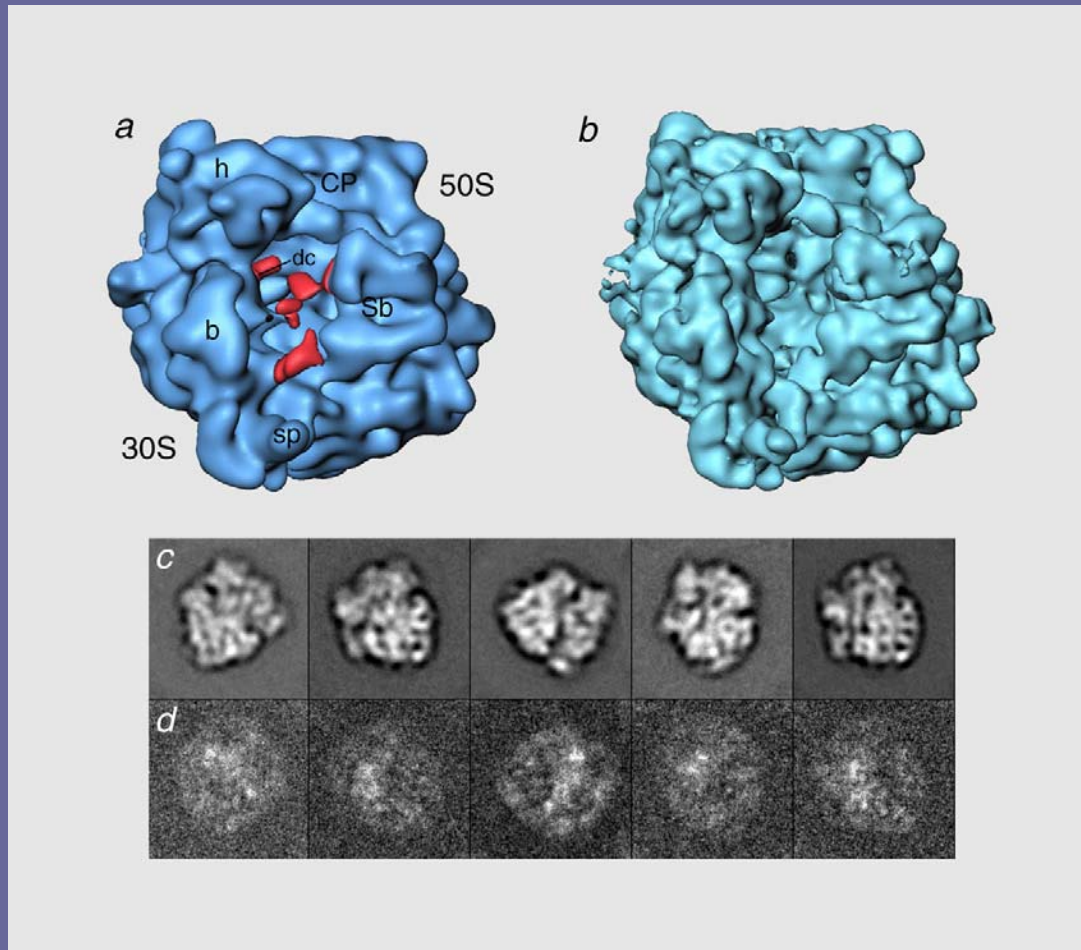
Current approach: assume all conformers are "similar". Treat problem in first approximation as a problem with a single conformer. Then try different models as references to see if population segregates.



Example: low occupancy of ternary complex

reconstruction using all data

empty ribosome (control)



Problem solved by supervised classification

