

SIMONS
ELECTRON
MICROSCOPY
CENTER

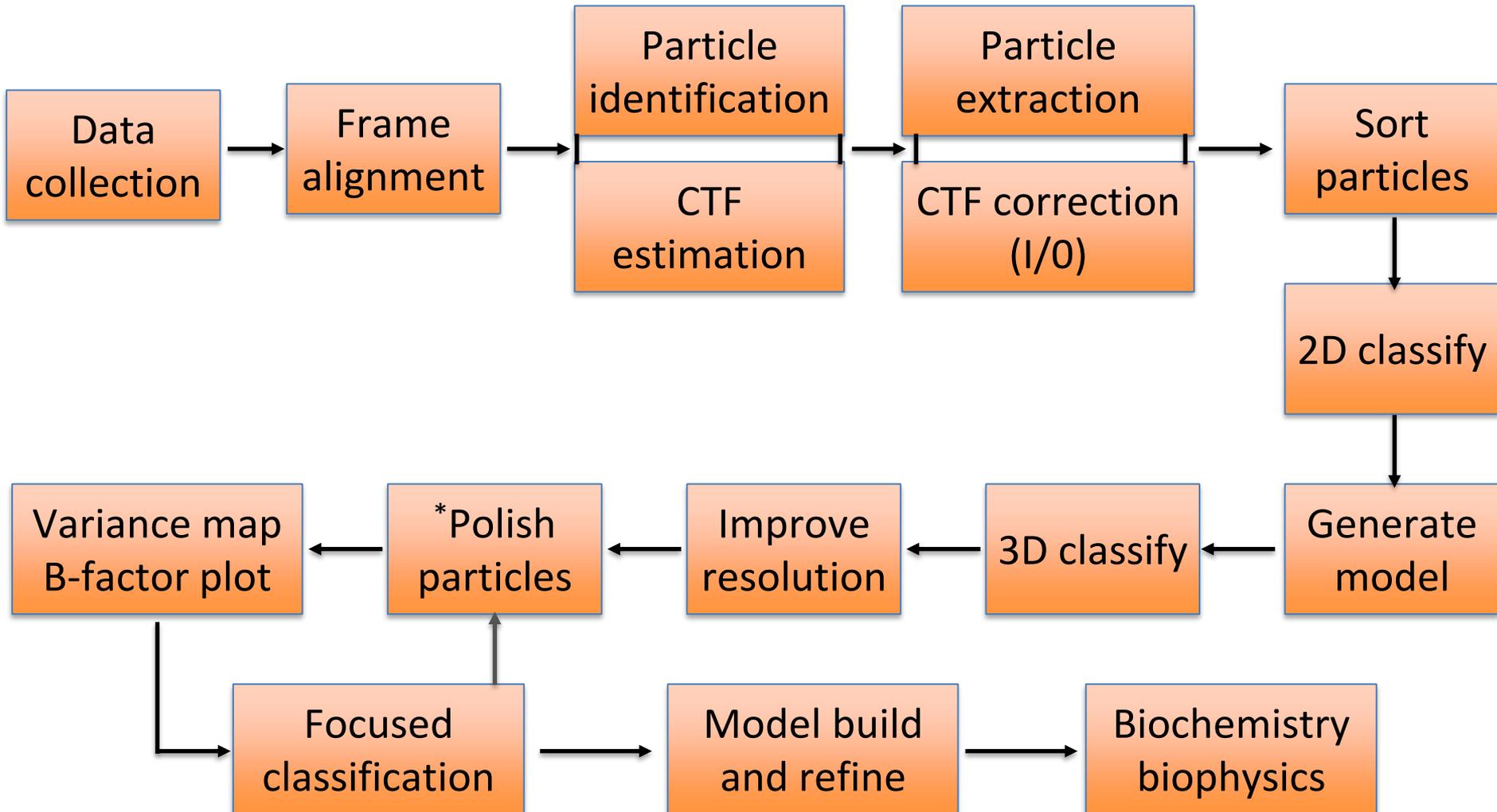
NYSBC 

Winter-Spring 2024 EM Course

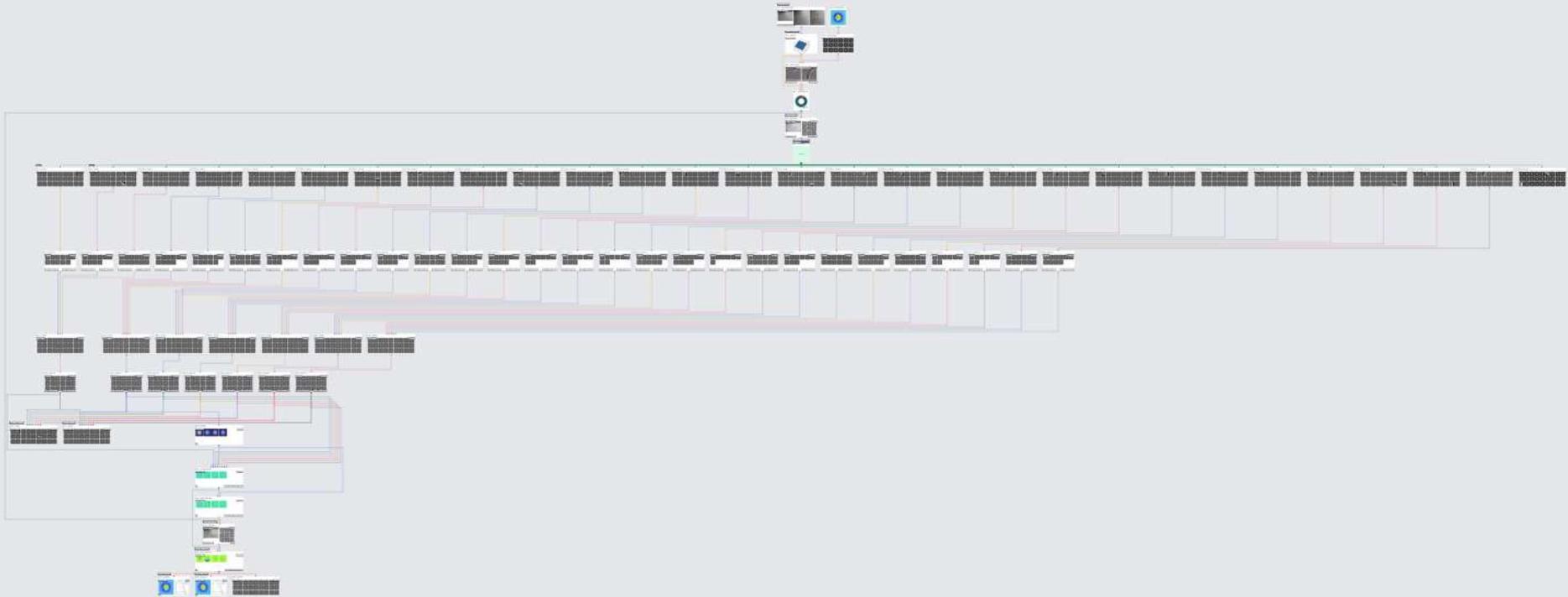
Single-particle analysis (part II)

Reza Khayat & Amedee des Georges

SPA Image analysis



Workflow in cryoSPARC

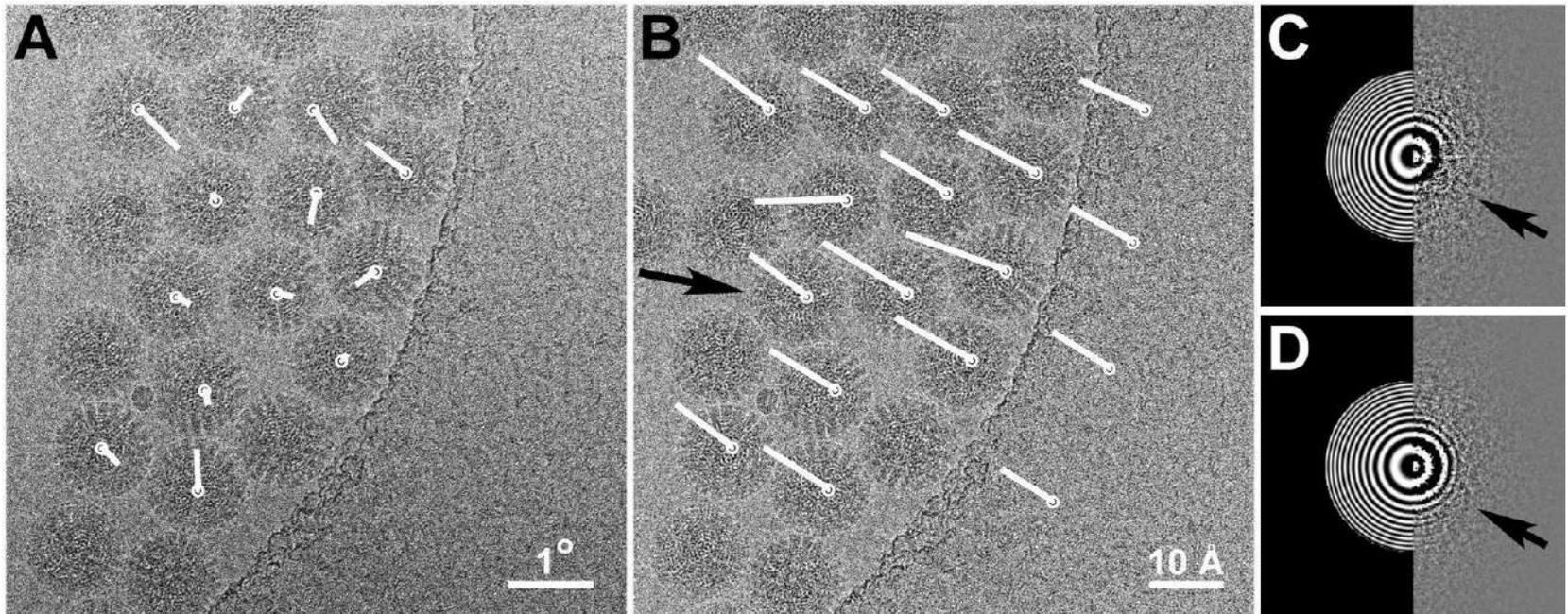


0.10 0.25x 0.5x 1x



Movie frame alignment

- UCSF MotionCor2
- Unblurr
- Warp
- cryoSPARC

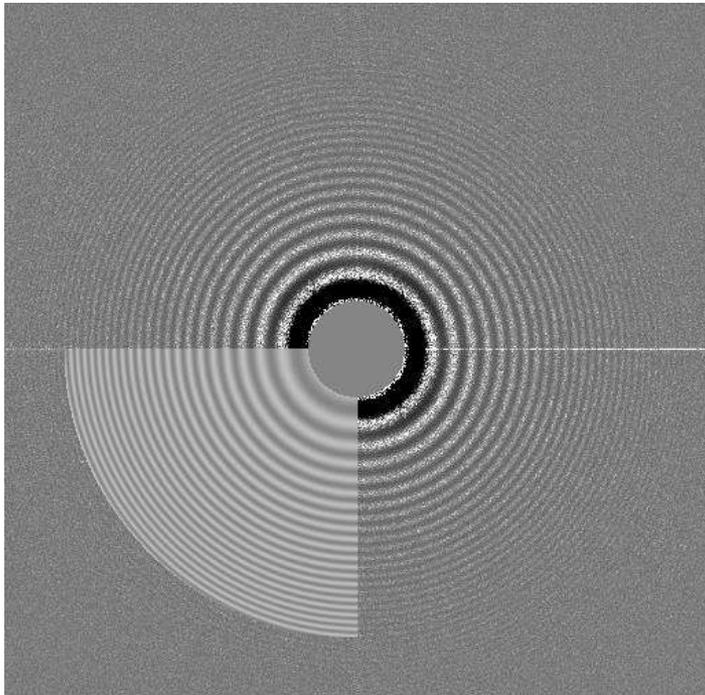


A/C: Unaligned micrographs and power spectrum
B/D: Aligned micrographs and power spectrum

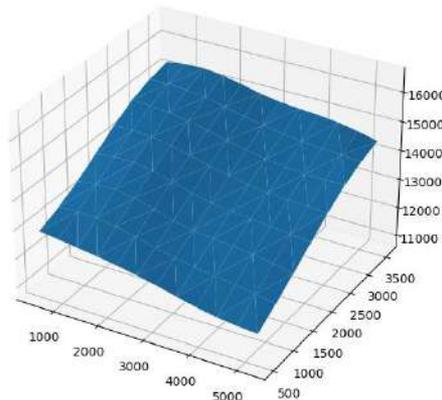
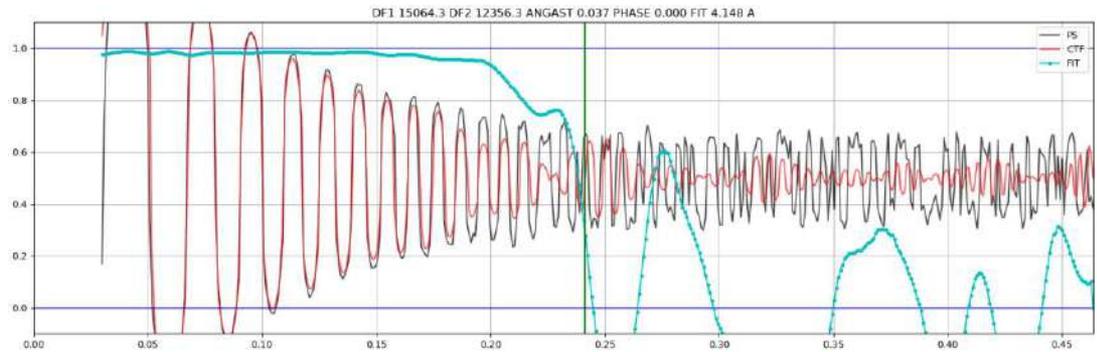
Campbell et al., 2012

CTF Estimation

- ACE
- CTFFIND4
- Sparx/EMAN
- GCTF
- Warp
- CryoSPARC (patch method)



CTFFIND4

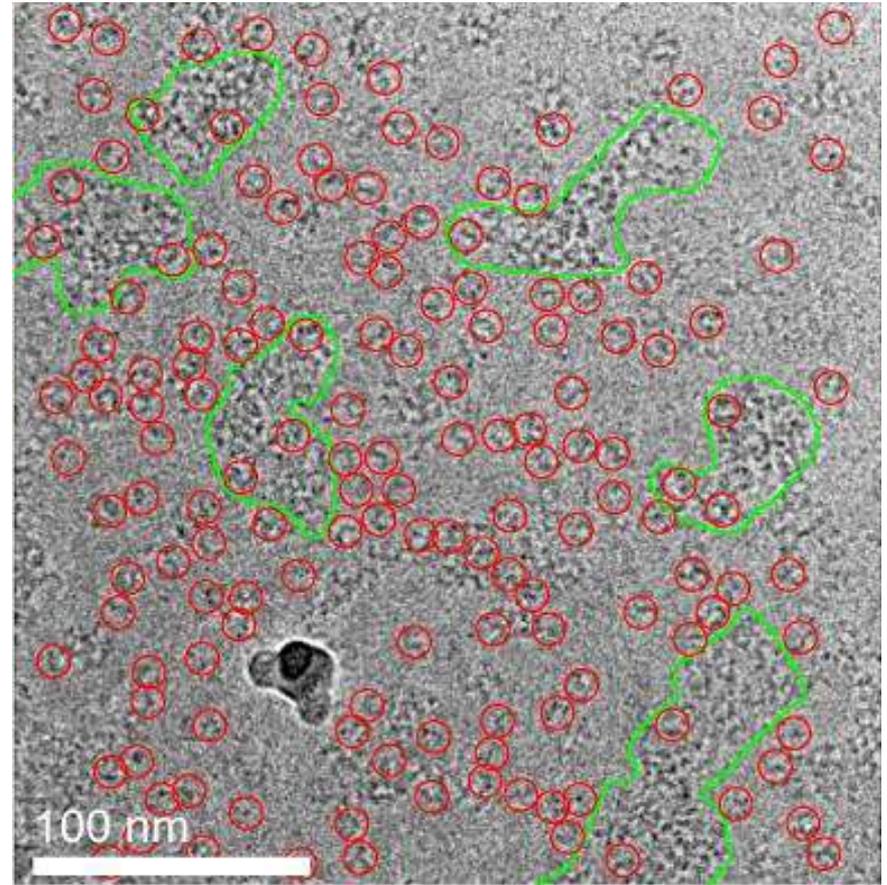


My stuff (CryoSPARC)

Particle Picking

Manual – Blob – Template – Neural network

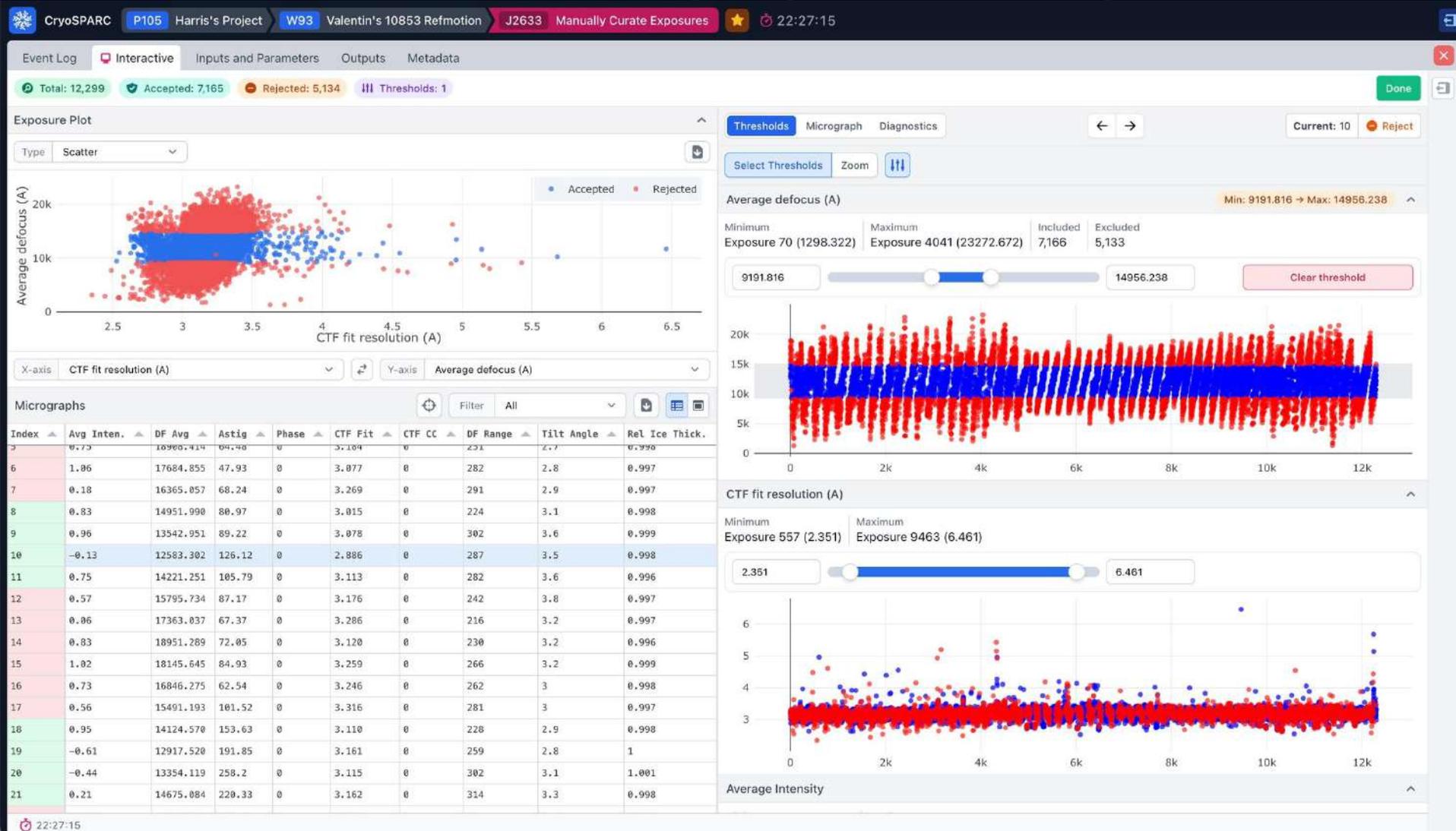
- FindEM
- DoG Picker
- Gautomatch
- Topaz
- Warp
- DeepPicker
- DeepEM
- crYOLO
- PIXER
- DRPnet
- DeepCryoPicker
- AutoCryoPicker



Bepler et al., 2019

- *Start with provided model, get 2D classes, and retrain*

Curation (cryoSPARC)



- Defocus range
- CTF fit resolution
- Number of particles

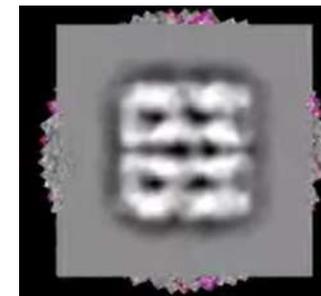
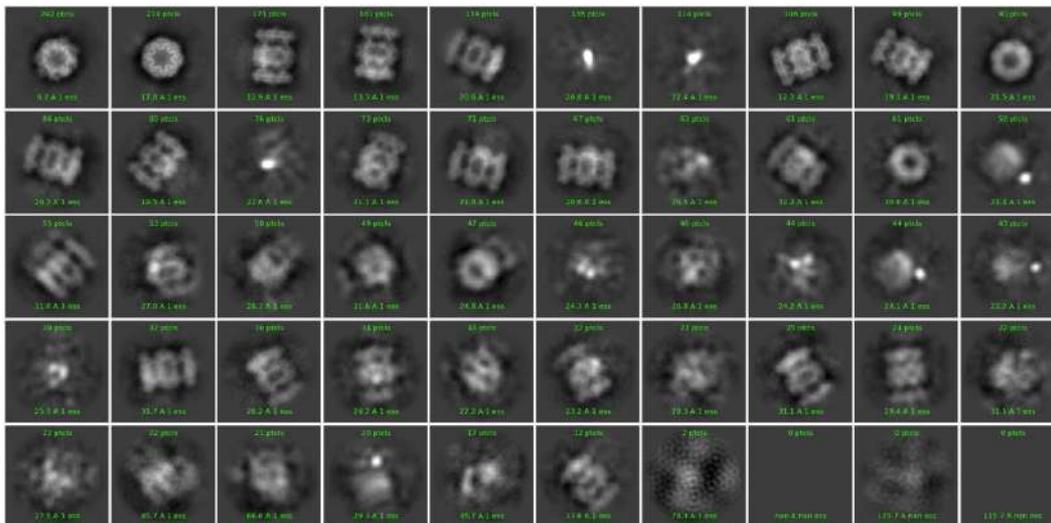
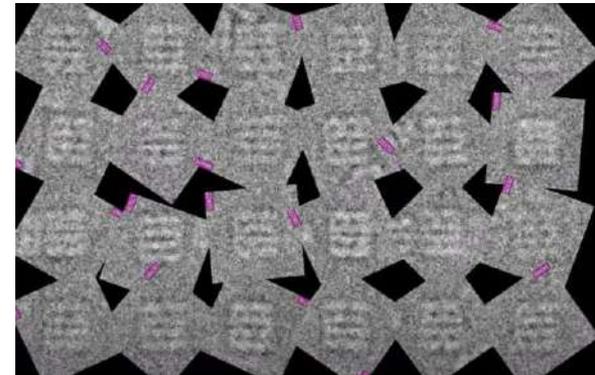
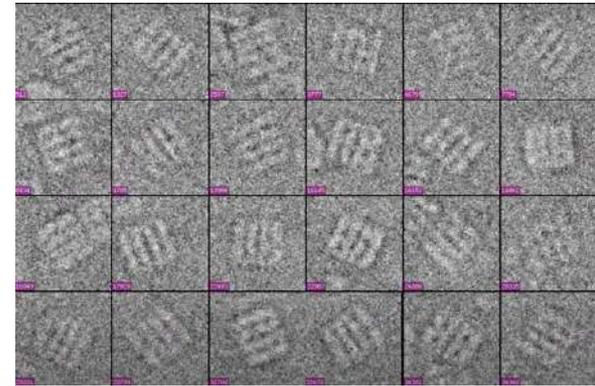
- Tilt angle
- Astigmatism
- Ice thickness

Particle Extraction

- Signal is delocalized according to energy of electron and defocus value of image:
 - Fred Sigworth 2022 lecture: $r = \delta * \lambda * f$
 - r is the radius that surrounds the particle. It describes how far the signal is delocalized.
 - $\delta = \text{defocus in } \text{\AA}$
 - $\lambda = 0.02 \text{ \AA}$ (wavelength of e^- at 300kV)
 - Frequency = desired resolution (e.g. 0.33\AA^{-1} for 3 \AA)
- Even number with low prime factors (2, 3, 5, and 7)
 - I like 32, 64, 128, 256, 320, 384
- You may want to downsize (fourier bin) the particles to expedite initial data processing, and save on drive space.

2D Classification

- Spider
- Sparx/EMAN2
- ISAC
- Relion
- Simple
- cryoSPARC
- Warp
- *Remove "bad" particles*



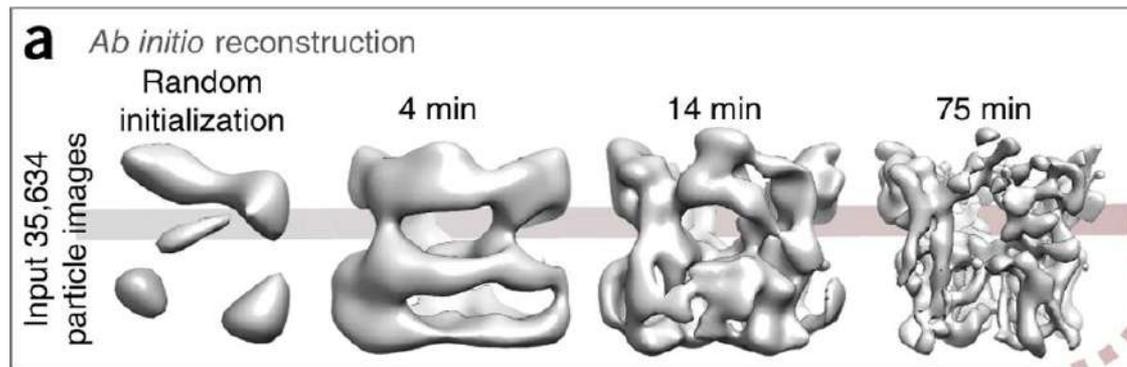
Lander 2009

2D Classification

- Anticipate 100 to 400 particles per class
- Don't ask for too many
- I split my particle stack into stacks of 100K particles and process each separately to get clean-vs-dirty particles
 - Radius of search
 - Relion
 - Tau fudge
 - CTF
 - cryoSPARC
 - Turn off Force Max over poses/shifts
 - Initial classification uncertainty factory (2 and above)
 - Number of iteration to anneal sigma as high as 25
 - Set online-EM iterations to 40
 - Set Batchsize per class to 400
 - Change Re-center mask threshold (possibly as high as 0.75) for centering particles and smearing neighbors
 - set White noise model to off

Initial Model

- Random conical tilt
- Orthogonal conical tilt
- Common-lines
- Tomography with STA
- Random initial parameters, optimize with stochastic gradient descent (SIMPLE, cryoSPARC, and Relion).
- SAXS/SANS
- Hybrid approach
- Structure prediction

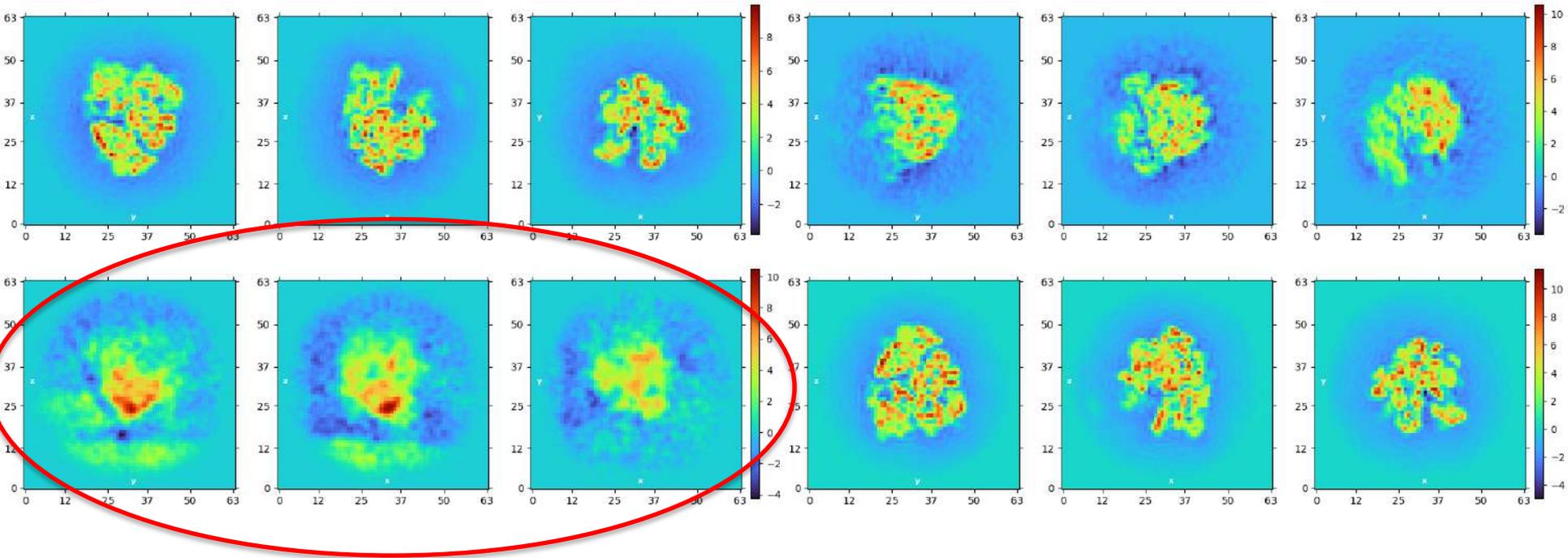


Initial Model

- Generate multiple initial models if uncertain in model
 - Look for continuity in density
 - Look for sausages to indicate α -helices
 - Are projections comparable to class averages?
- Ask for multiple models to be generated
- Starting frequency should have more information than `particle_size / 5`
- Use C_1 symmetry

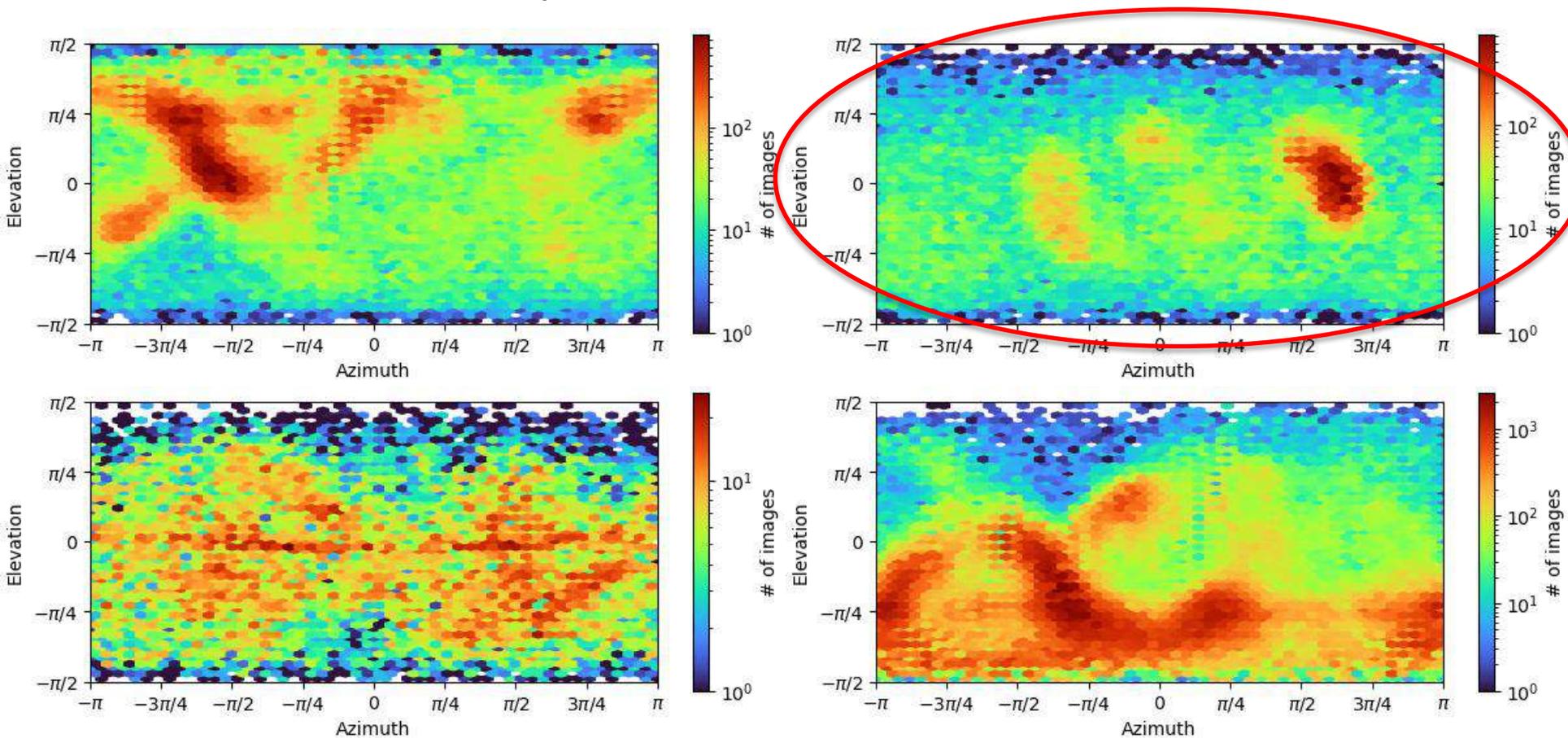
3D Classification

- Can be used to clean data further
 - Discard “bad” particles



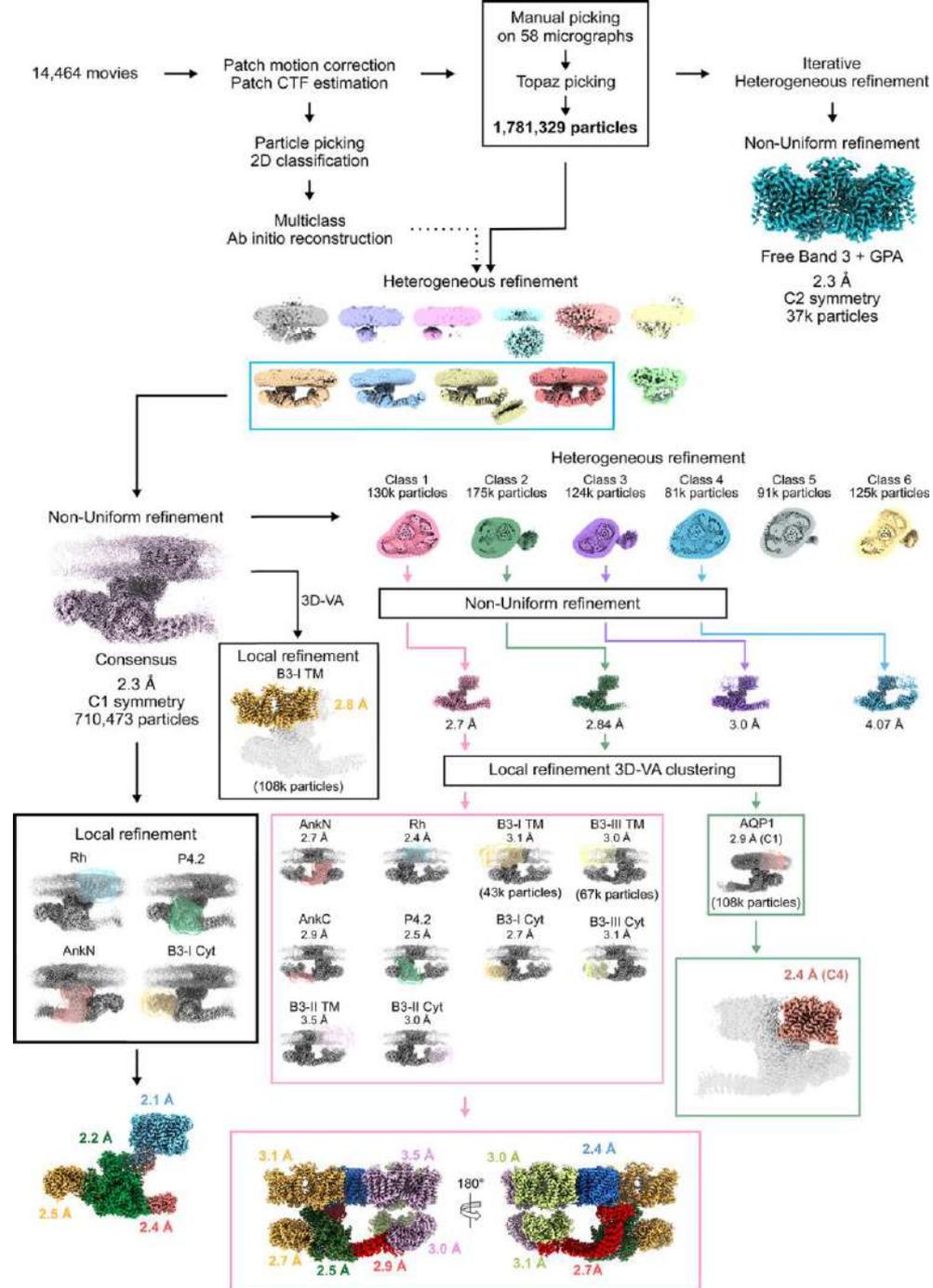
3D Classification

- Can be used to clean data further
 - Discard “bad” particles
 - Discard some preferred orientations



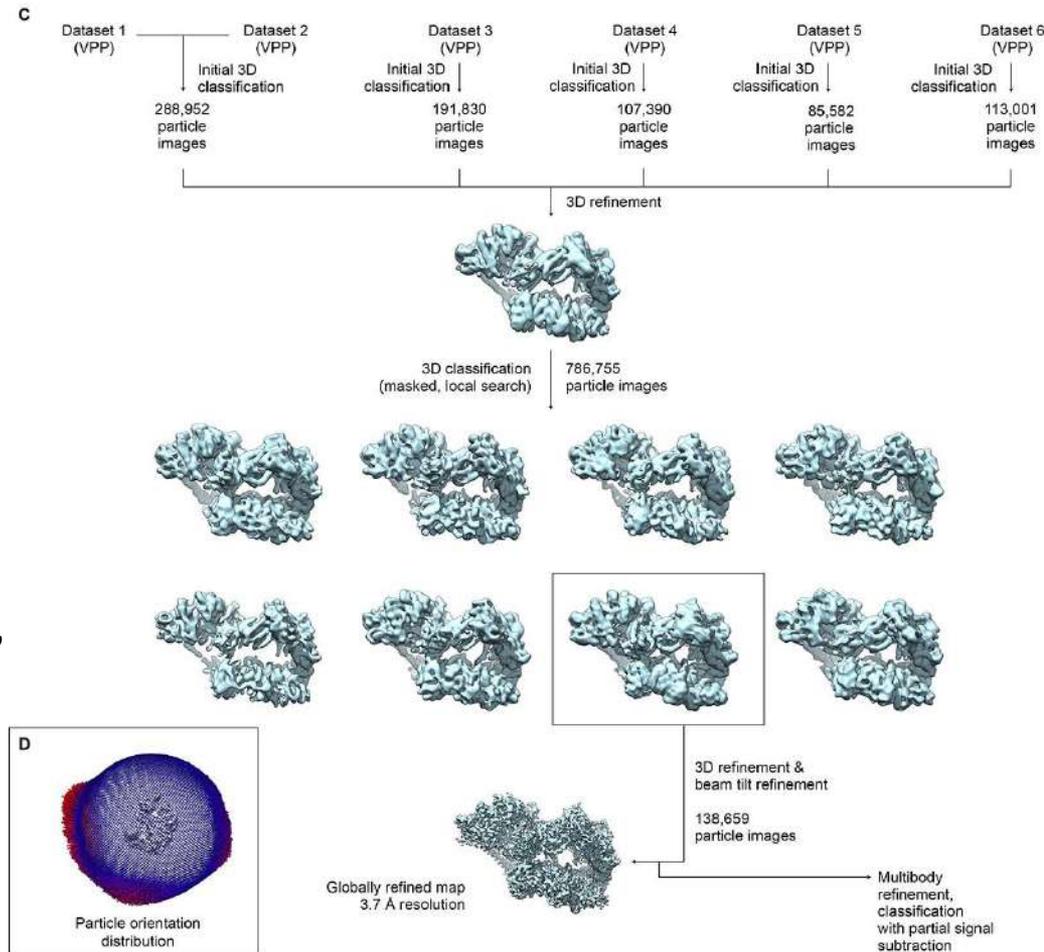
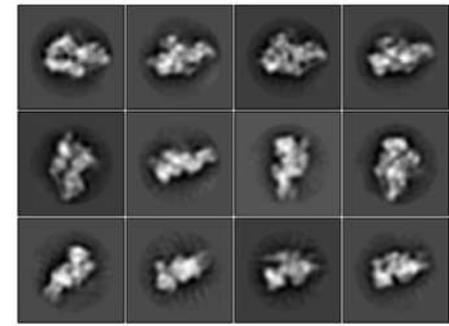
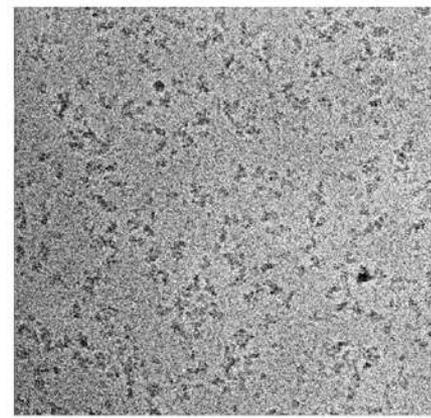
3D Classification

- Configurational and conformational differences
- Multiple ab initio models generate
- 3D classification use to identify configurational differences.
- Local refinement used to identify conformational differences



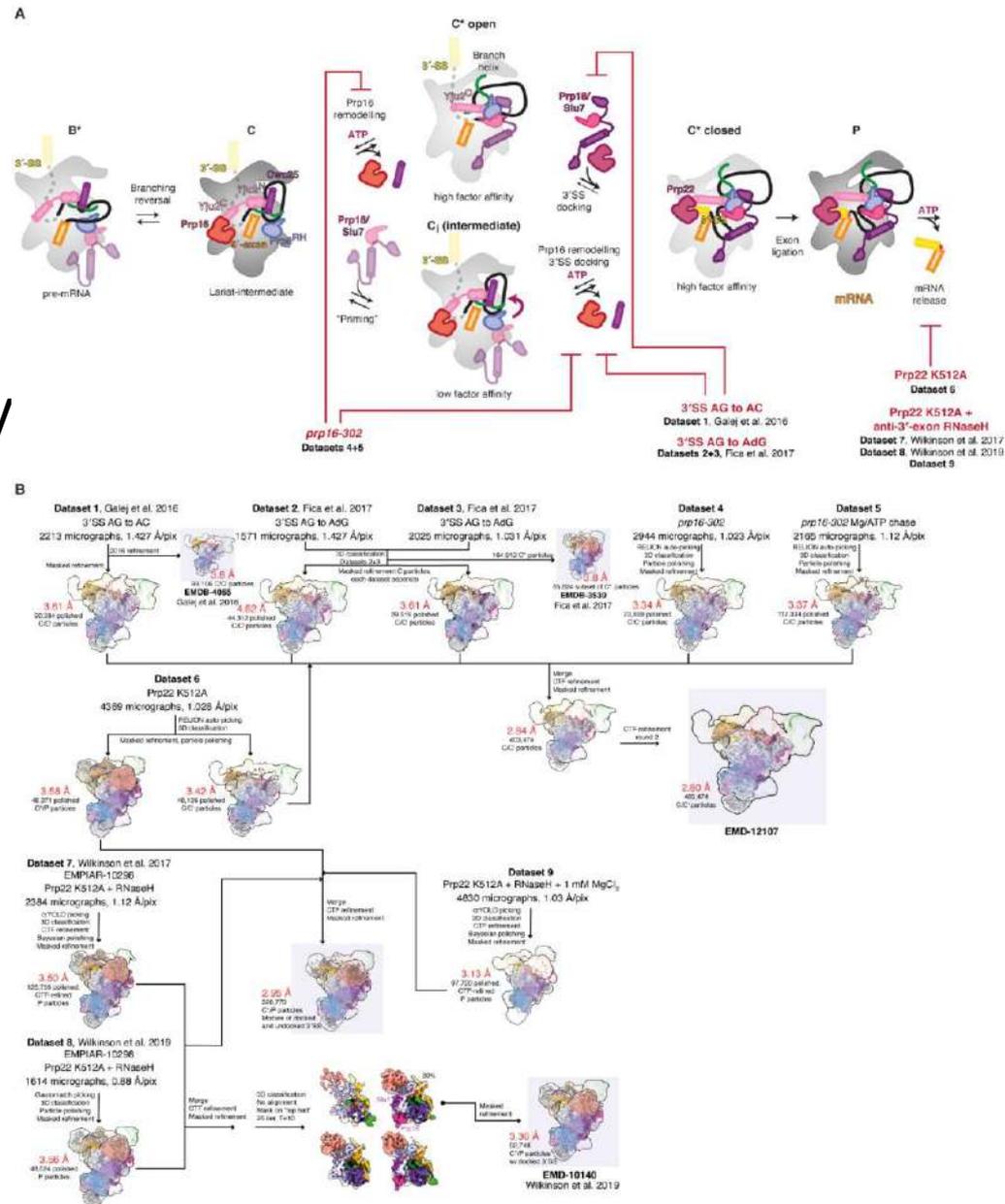
3D Classification

- TFIIH, transcription initiation by RNAPII and NER
- Enrich rare views that 2D classification would discard
- Do 2D classification (B) for sanity check
- Extract particles and perform 3D classification with high tau2_fudge value to enrich for rare views (D).
 - Value empirically determined. Try 1, 5, 10, 20, 50, 100, 200, 500, 1000



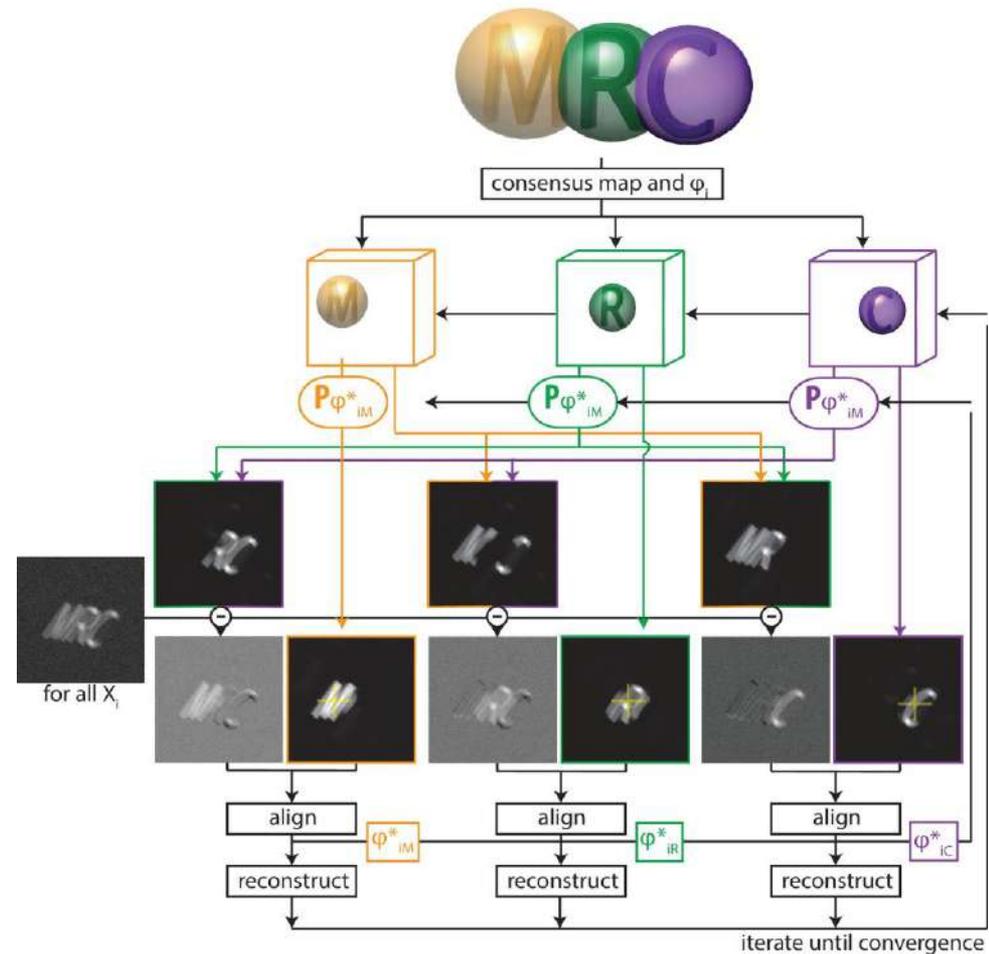
3D Classification

- Spliceosome
- Initial reconstruction is at 2.8 Å; however, lots of domains/proteins at periphery have poor density
- Signal subtraction coupled with focused classification and empirically determined tau2_fudge values (Relion) improve their resolutions
 - Try 1, 5, 10, 20, 50, 100, 200, 500, 1000 in parallel



Signal subtraction

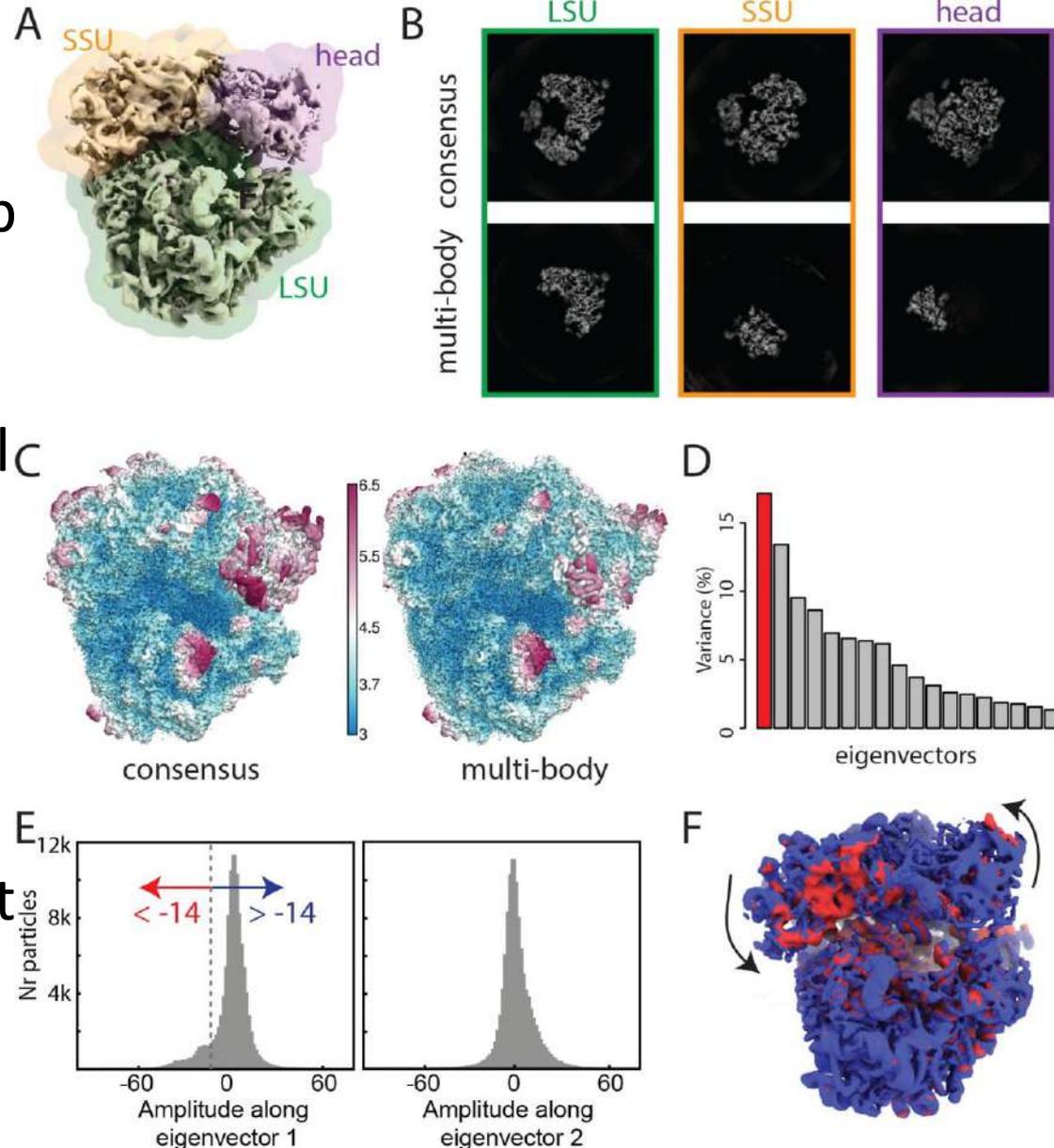
- Improve resolution of desired region(s)
- Generate consensus map
- Mask region of interest
- Use consensus map to subtract everything outside/inside of mask from each particle
- Refine map of remaining signal
- Subtraction does not always work completely. May need to iterate through this process.



Nakane et al., 2018

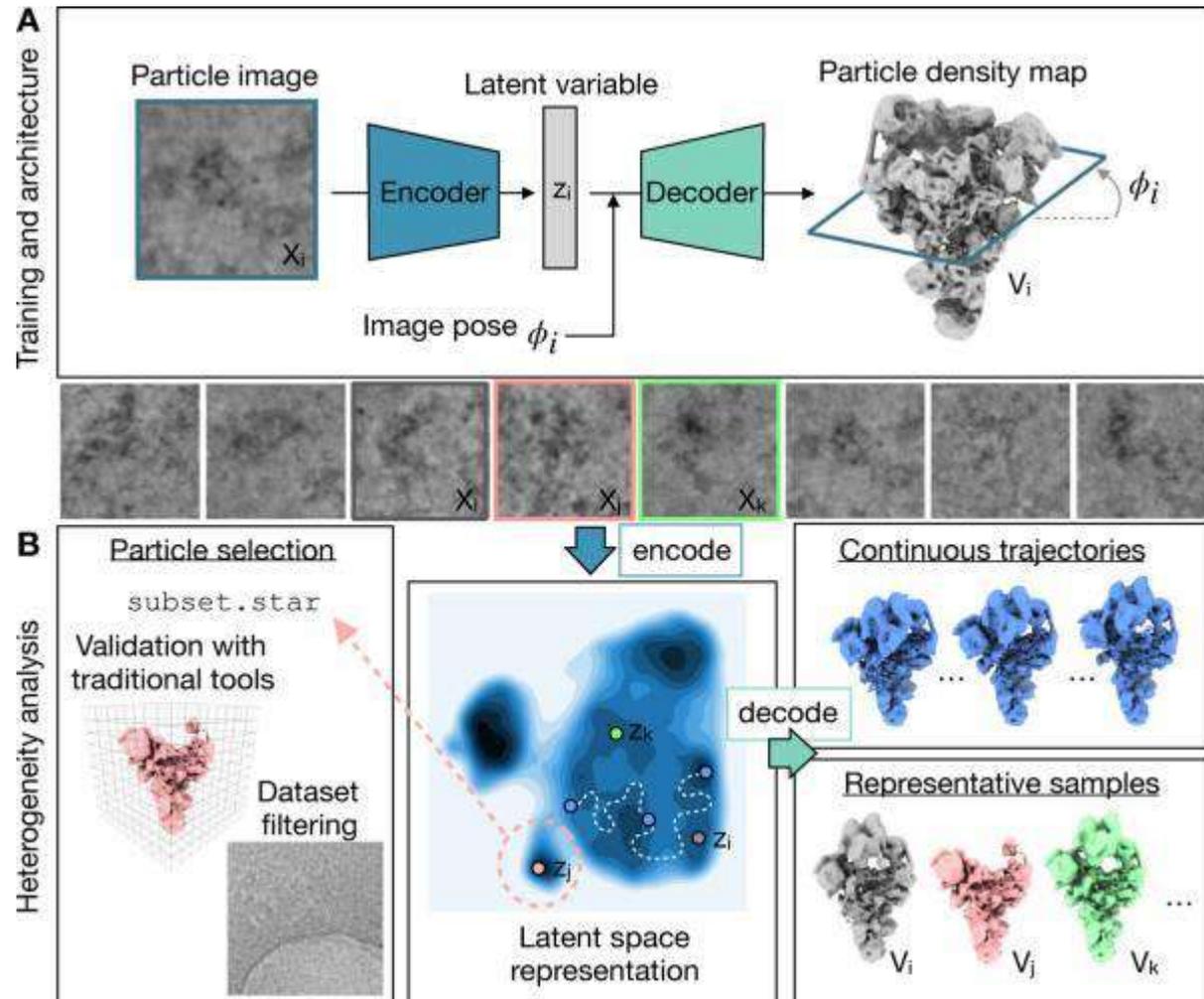
Multi-body refinement

- Improve resolution of desired region(s)
- Generate consensus map
- Mask regions of interest
- Relion will do signal subtraction and principal component analysis to identify rigid body motions.
- There is a lower size limitation
- Mask boundaries will not be trivial to interpret

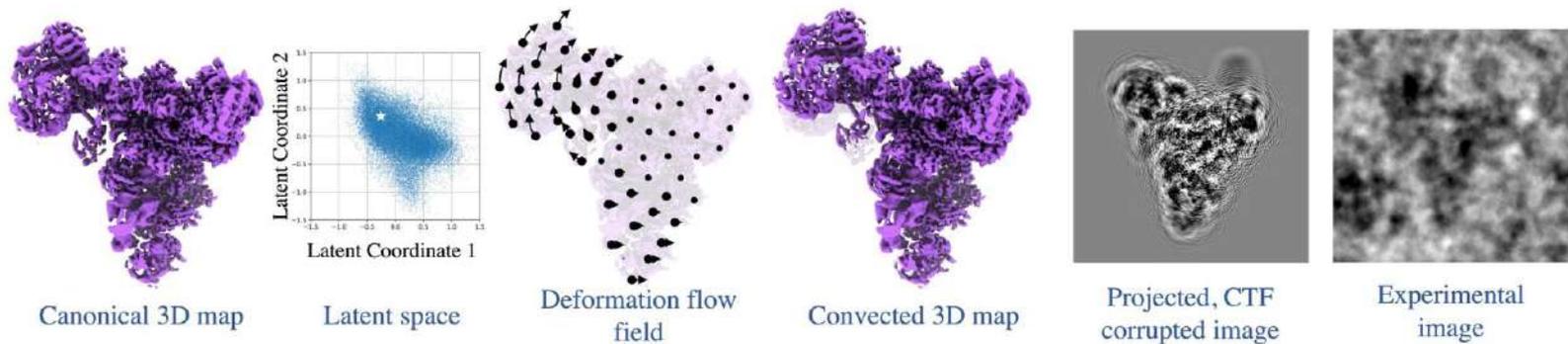
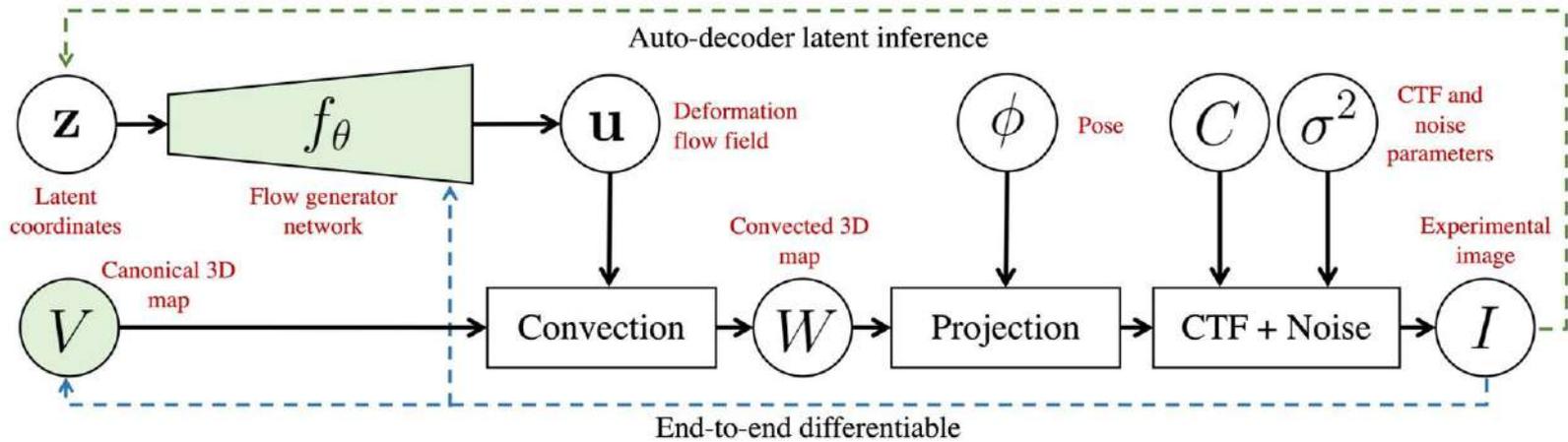


CryoDRGN

- Uses deep generative model to identify data heterogeneity
- The latent space representation (contour map in bottom center) can be used to generate density maps
- Continuous trajectories can be generated for studying motion



3D Flexible refinement



- Uses deep generative model for continuous heterogeneity
- The user defined nonrigid deformation flow field can be used to improve resolution of flexible regions
- Implemented in cryoSPARC