

CryoSPHERE: bridging the gap between AlphaFold and cryoEM experiments



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Introduction

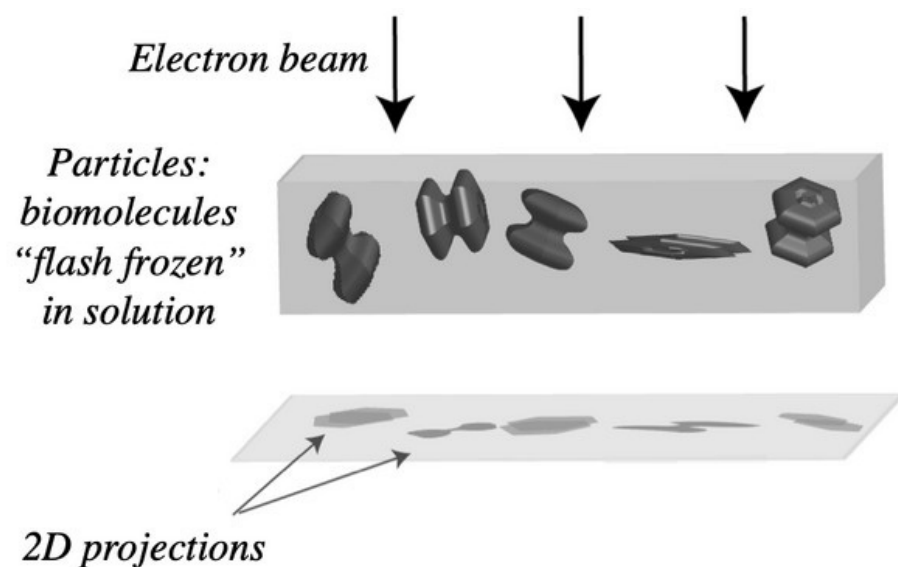


Figure 1: cryoEM acquisition process.

CryoEM reconstruction challenges:

- 100k to 1M images of copies of the same protein
- Orientation of the protein on each image is unknown
- Signal-to-noise ratio very low (0.1 to 0.001)
- On each image, the protein has different shape

We want to recover, for each image, the corresponding shape of the protein.

The current SOTA:

- Too low resolution for very noisy datasets
- Too low resolution for very mobile datasets
- Do not use AlphaFold prior knowledge

Method

CryoSPHERE uses the prior knowledge on the protein structure brought by AlphaFold and learns concurrently:

- How to decompose the protein structure in N segments
- How to move these segments approximately rigidly to recover the conformations

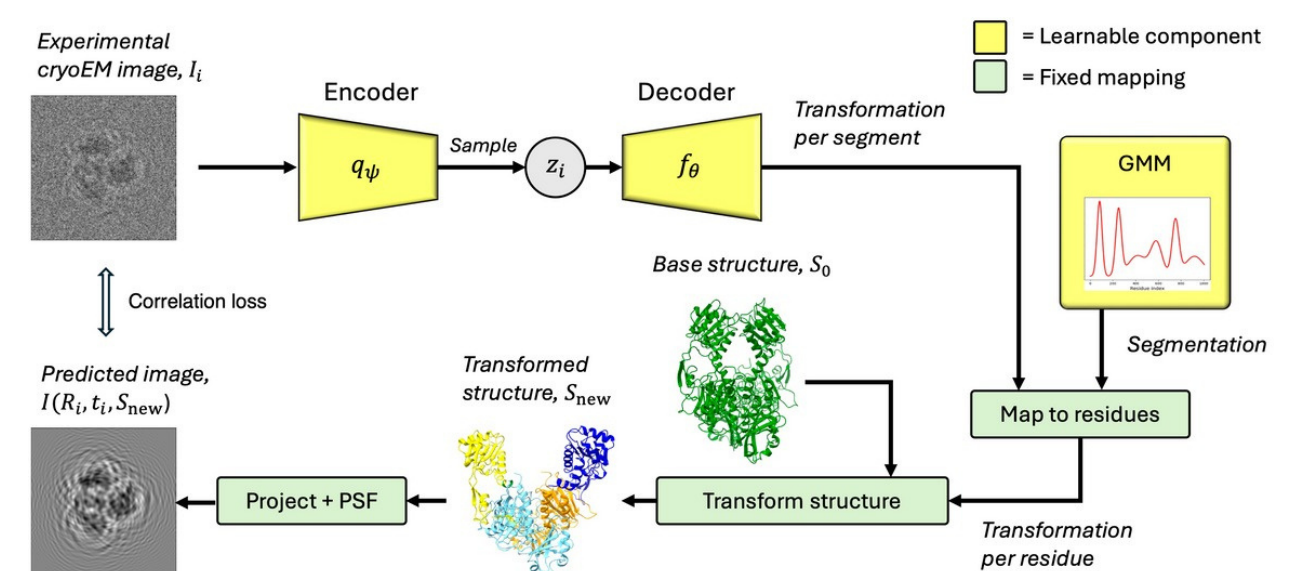


Figure 2: flow chart of cryoSPHERE

Experiments

Toy dataset:

- 10k structures obtained by MD simulation
- 150k images in total with SNR 0.001

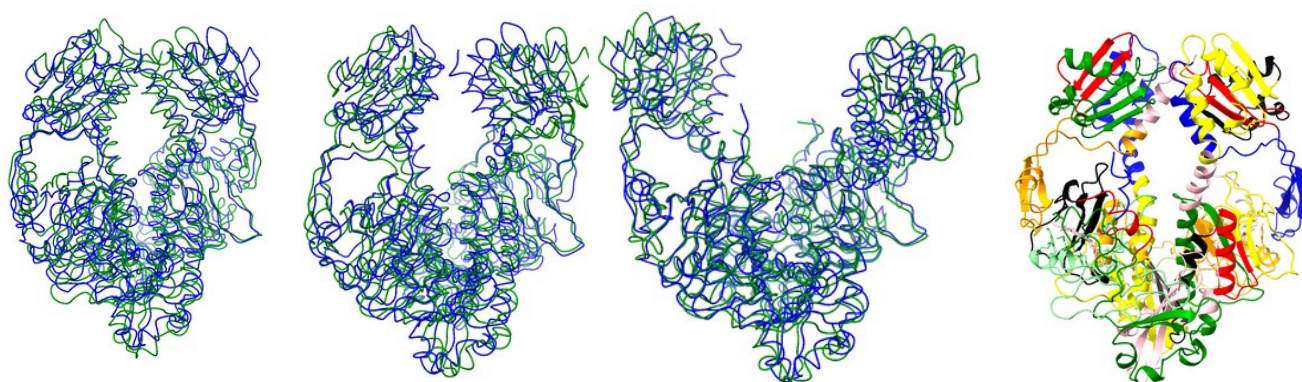


Figure 3: in blue, selection of predicted structures by cryoSPHERE. In green, corresponding ground truth. Rightmost: base structure with colored segments learned by cryoSPHERE.

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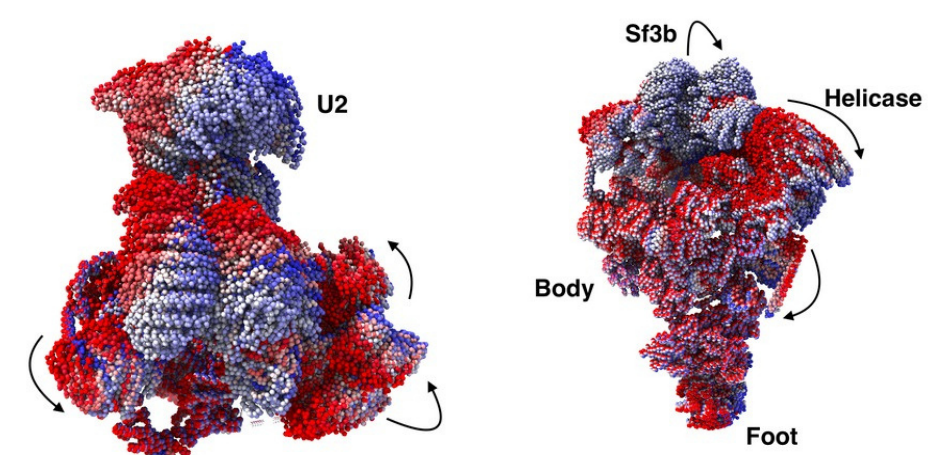


Figure 4: traversal of the latent space, from red to blue. The Sf3b domain gets incurvated down while the helicase move closer to the foot of the protein.

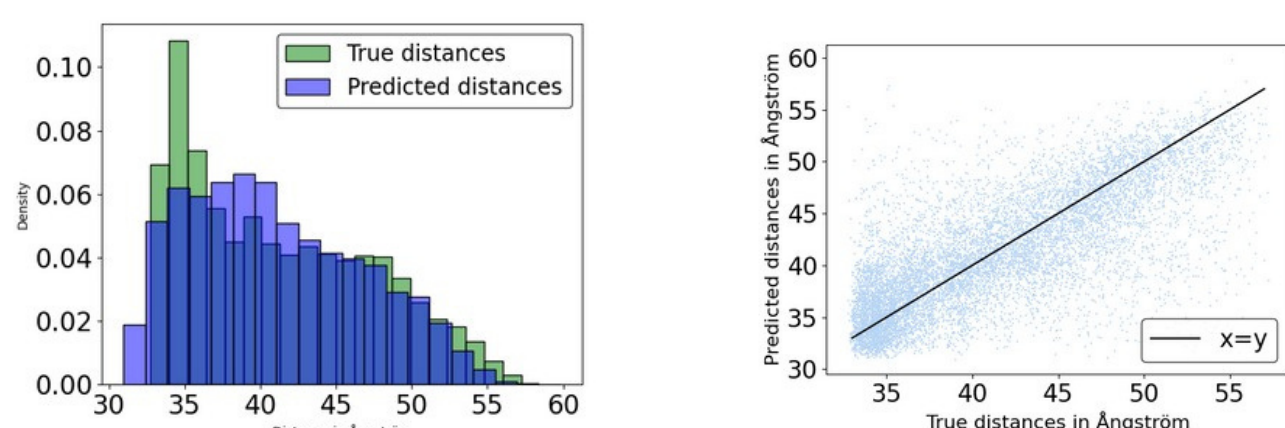


Figure 4: left: histograms of the ground truth and predicted distances between the two upper domains of the protein. Right: predicted vs true distances.

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