



Public Data-based Report



# Validation Report Service

## Cryo-EM Map Validation Report

Report to assess Cryo-EM Volume Map at Level(s) 0, 1

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This report has been generated based on data publicly available at [EMDB](#).

### Basic Entry Information:

**EMDB ID:** [EMD-51709](#)

**Title:** High resolution structure of B. oleracea mitoribosome - Small subunit body protuberance focus

**Authors:** [See EMDB entry link](#)

**Deposited on:** 2024-10-02T00:00:00

**Reported Resolution:** 2.58 Å

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Last update: **April 4, 2026, 3:27am**

## Context

Cryo-electron microscopy is currently one of the most active techniques in Structural Biology. The number of maps deposited at the [Electron Microscopy Data Bank](#) is rapidly growing every year and keeping the quality of the submitted maps is essential to maintain the scientific quality of the field. The ultimate quality measure is the consistency of the map and an atomic model. However, this is only possible for high resolution maps. Over the years there have been many suggestions about validation measures of 3DEM maps. Unfortunately, most of these measures are not currently in use for their spread in multiple software tools and the associated difficulty to access them. To alleviate this problem, we made available a validation grading system that evaluate the information provided to assess the map.

This system grades a map from 0 to 5 depending on the amount of information available. In this way, a map could be validated at Level 0 (the deposited map), 1 (two half maps), 2 (2D classes), 3 (particles), 4 (... + angular assignment), 5 (... + micrographs and coordinates). In addition, we can have three optional qualifiers: A (... + atomic model), W (... + image processing workflow), and O (... + other techniques). To know more about this service read this [paper](#)

This Validation Report Service uses Scipion (see this [link](#) for more detail) as workflow engine and ChimeraX (see this [link](#) for more detail) to generate the 3D views. For more information about the different methods and softwares used for this report, see the references [here](#).



### Summarized overall quality

The map seems to have some problem in its centering or extra space (see Sec. 2.1). There seems to be a problem with the suggested threshold (see Sec. 2.2). There seems to be a problem with the map's background (see Sec. 2.3). The resolution does not seem to be uniform in all directions (see Sec. 4.6).

The average resolution of the map estimated by various methods goes from 2.7Å to 5.3Å with an average of 3.8Å. The resolution reported by the user was 2.6Å. The resolution reported may be overestimated.

**The overall score (passing tests) of this report is 7 out of 12 evaluable items.**

0.a Mass analysis	Sec. 2.1	4 warnings
0.b Mask analysis	Sec. 2.2	2 warnings
0.c Background analysis	Sec. 2.3	2 warnings
0.d B-factor analysis	Sec. 2.4	OK
0.e DeepRes	Sec. 2.5	Could not be measured
0.f LocBfactor	Sec. 2.6	OK
0.g LocOccupancy	Sec. 2.7	Could not be measured
0.h DeepHand	Sec. 2.8	OK
1.a Global resolution	Sec. 4.1	3 warnings
1.b FSC permutation	Sec. 4.2	OK
1.c Blocres	Sec. 4.3	OK
1.d Resmap	Sec. 4.4	Could not be measured
1.e MonoRes	Sec. 4.5	OK
1.f MonoDir	Sec. 4.6	2 warnings
1.g FSO	Sec. 4.7	OK
1.h FSC3D	Sec. 4.8	Could not be measured

## Summary of the warnings across sections.

### Section 2.1 (0.a Mass analysis)

1. **The volume might be significantly decentered in Z.**
2. **The center of mass in X may be significantly shifted. This is common when the refinement is applied exclusively to one protein region.**
3. **The center of mass in Y may be significantly shifted. This is common when the refinement is applied exclusively to one protein region.**
4. **The center of mass in Z may be significantly shifted. This is common when the refinement is applied exclusively to one protein region.**

### Section 2.2 (0.b Mask analysis)

1. **There might be a problem of connectivity at this threshold because more than 5 connected components are needed to reach 95% of the total mask. Probably a smaller threshold will not cause this issue.**
2. **There might be a problem in the construction of the mask, because the overlap is smaller than 0.75. A common reason is that the suggested threshold causes too many disconnected components.**

### Section 2.3 (0.c Background analysis)

1. **The null hypothesis that the background mean is 0 has been rejected because the p-value of the comparison is smaller than 0.001**
2. **There is a significant proportion of outlier values in the background (cdf5 ratio=5932.57)**

### Section 4.1 (1.a Global resolution)

1. **The reported resolution, 2.58 Å, is particularly high with respect to the resolution calculated by the FSC, 4.54 Å**
2. **The reported resolution, 2.58 Å, is particularly high with respect to the resolution calculated by the DPR, 3.32Å.**
3. **The reported resolution, 2.58 Å, is particularly high with respect to the resolution calculated by the SSNR, 4.47Å.**

### Section 4.6 (1.f MonoDir)

1. The distribution of best resolution is not uniform in all directions. The associated p-value is 0.000000.
2. The resolution reported by the user, 2.58Å, is at least 80% smaller than the average directional resolution, 3.91 Å.

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# 1 Input data

Input map: emd\_51709.map

SHA256 hash: 6edd56c8b2ef86bf1a2b682532371155764f1e7a140bf8e029acdcfbdbd50988

Voxel size: 0.937286 (Å)

Visualization threshold: 0.115000

Resolution estimated by user: 2.58

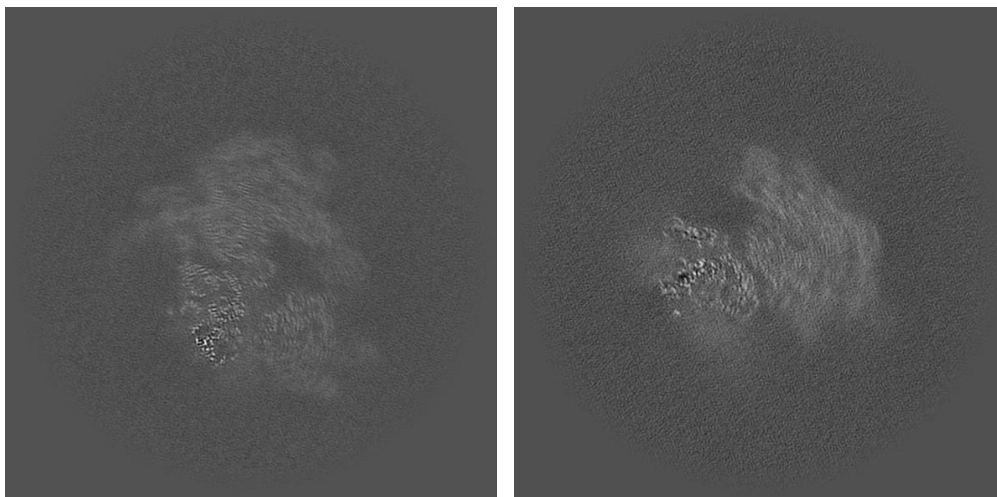
## Orthogonal slices of the input map

### **Explanation:**

In the orthogonal slices of the map, the noise outside the protein should not have any structure (stripes going out, small blobs, particularly high or low densities, ...)

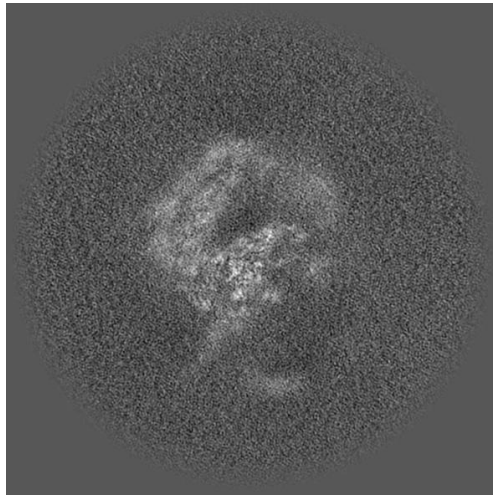
### **Results:**

See Fig. 1.



(a) X Slice 294

(b) Y Slice 294



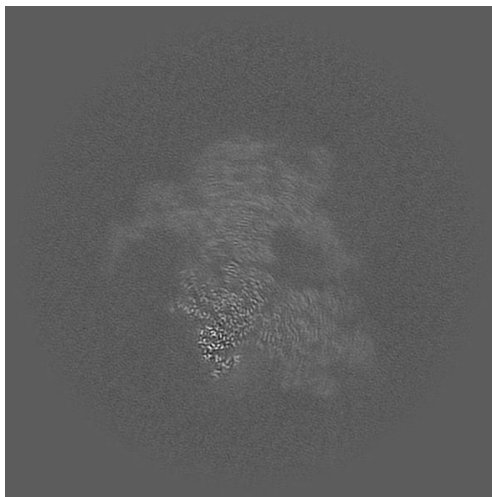
(c) Z Slice 294

Figure 1: Central slices of the input map in the three dimensions

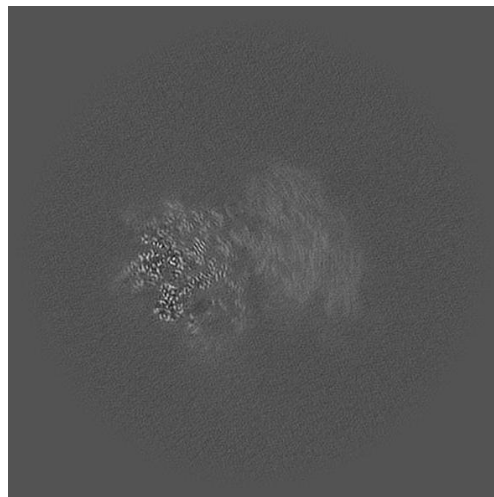
Orthogonal slices of maximum variance of the input map

**Results:**

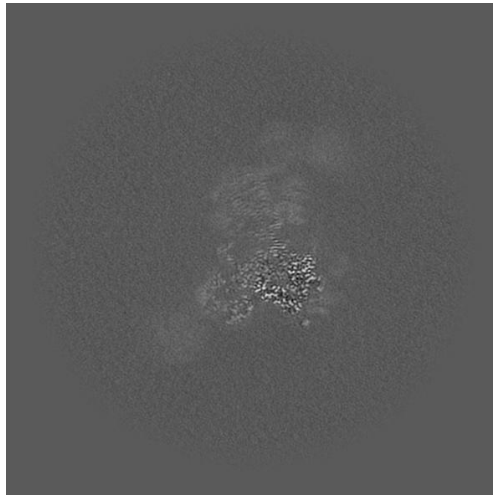
See Fig. 2.



(a) X Slice 307



(b) Y Slice 250



(c) Z Slice 198

Figure 2: Slices of maximum variation in the three dimensions

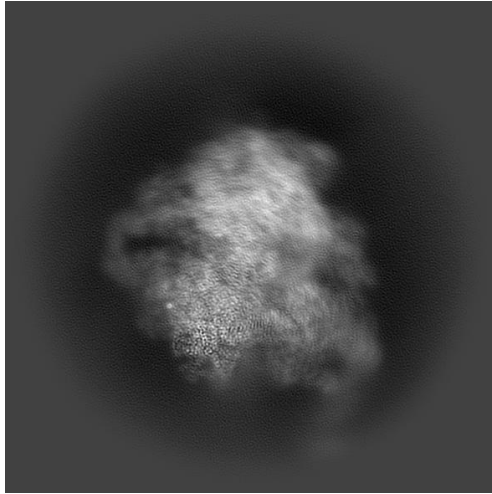
### Orthogonal projections of the input map

#### **Explanation:**

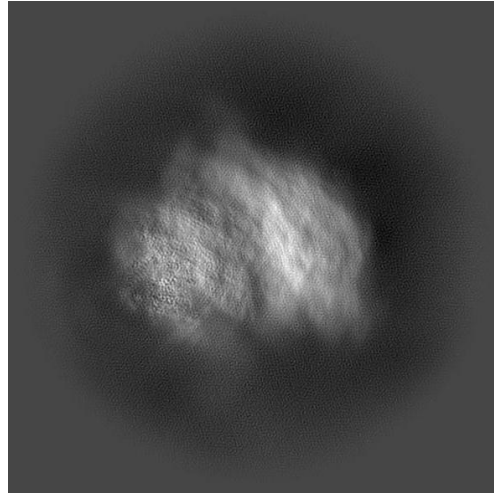
In the projections there should not be stripes (this is an indication of directional overweighting, or angular attraction), and there should not be a dark halo around or inside the structure (this is an indication of incorrect CTF correction or the reconstruction of a biased map).

#### **Results:**

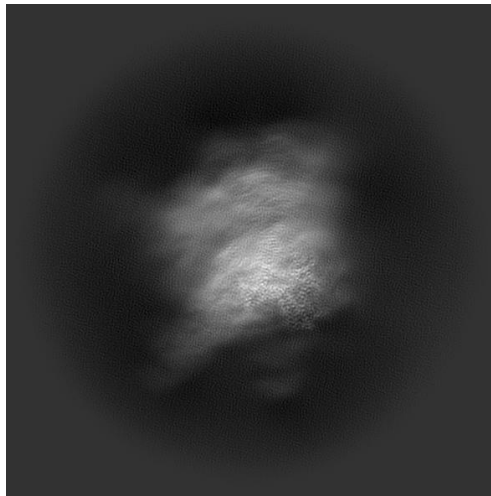
See Fig. 3.



(a) X Projection



(b) Y Projection



(c) Z Projection

Figure 3: Projections in the three dimensions

### Isosurface views of the input map

#### **Explanation:**

An isosurface is the surface of all points that have the same gray value. In these views there should not be many artifacts or noise blobs around the map.

**Results:**  
See Fig. 4.

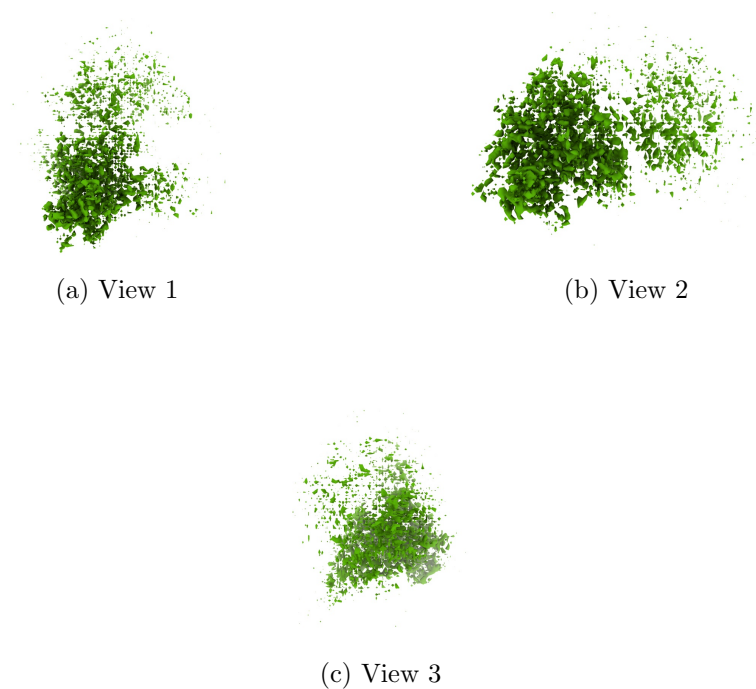


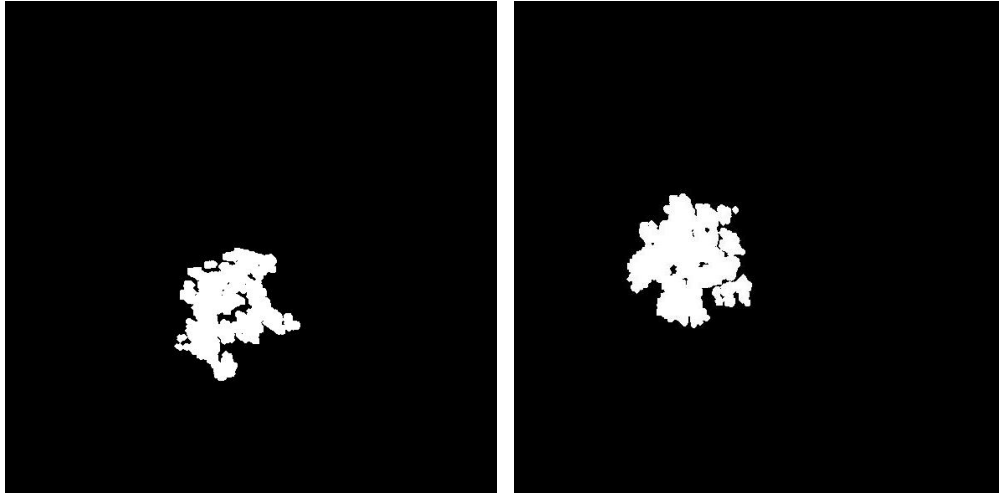
Figure 4: Isosurface at threshold=0.115000. Views generated by ChimeraX at a the following X, Y, Z angles: View 1 (0, -90, -90), View 2 (-90, 0, -90), View 3 (0, 0, 0).

**Orthogonal slices of maximum variance of the mask with hard borders**

**Explanation:**

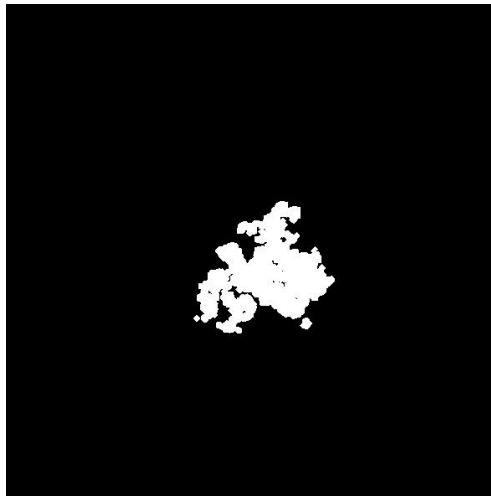
The mask with hard borders has been calculated at the suggested threshold 0.115000, the largest connected component was selected, and then dilated by 2Å.

**Results:**  
See Fig. 5.



(a) X Slice 328

(b) Y Slice 252



(c) Z Slice 201

Figure 5: Slices of maximum variation in the three dimensions of the mask with hard borders

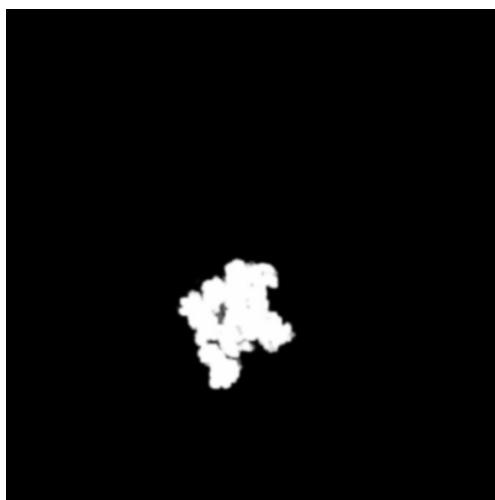
**Orthogonal slices of maximum variance of the mask with soft borders**

**Explanation:**

The mask with soft borders has been calculated at the suggested threshold 0.115000, the largest connected component was selected, and then dilated by

2Å.

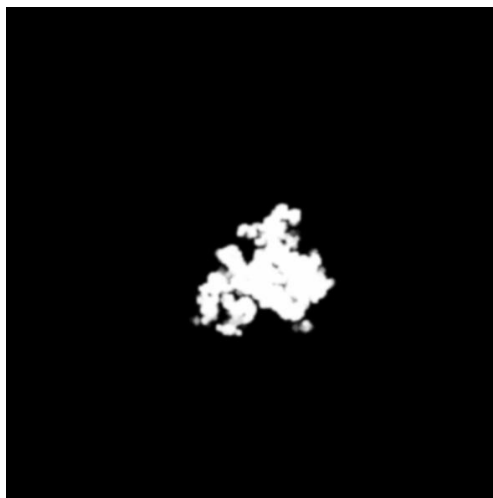
**Results:**  
See Fig. 6.



(a) X Slice 317



(b) Y Slice 252



(c) Z Slice 201

Figure 6: Slices of maximum variation in the three dimensions of the mask with soft borders

## 2 Level 0 analysis

### 2.1 Level 0.a Mass analysis

#### **Explanation:**

The reconstructed map must be relatively well centered in the box, and there should be at least  $30\text{\AA}$  (the exact size depends on the CTF) on each side to make sure that the CTF can be appropriately corrected.

#### **Results:**

The space from the left and right in X are 205.27 and 183.71  $\text{\AA}$ , respectively. There is a decentering ratio  $(\text{abs}(\text{Right-Left})/\text{Size})\%$  of 3.91%

The space from the left and right in Y are 180.90 and 221.20  $\text{\AA}$ , respectively. There is a decentering ratio  $(\text{abs}(\text{Right-Left})/\text{Size})\%$  of 7.31%

The space from the left and right in Z are 125.60 and 275.56  $\text{\AA}$ , respectively. There is a decentering ratio  $(\text{abs}(\text{Right-Left})/\text{Size})\%$  of 27.21%

The center of mass is at  $(x,y,z)=(-633.67,164.72,1723.71)$ . The decentering of the center of mass  $(\text{abs}(\text{Center})/\text{Size})\%$  is 157.77, 21.99, and 243.15, respectively.

**Automatic criteria:** The validation is OK if 1) the decentering and center of mass less than 20% of the map dimensions in all directions, and 2) the extra space on each direction is more than 20% of the map dimensions. For local reconstruction, focused refinement, or similar, warnings are expected.

**WARNINGS:** 4 warnings

1. **The volume might be significantly decentered in Z.**
2. **The center of mass in X may be significantly shifted. This is common when the refinement is applied exclusively to one protein region.**
3. **The center of mass in Y may be significantly shifted. This is common when the refinement is applied exclusively to one protein region.**
4. **The center of mass in Z may be significantly shifted. This is common when the refinement is applied exclusively to one protein region.**

## 2.2 Level 0.b Mask analysis

### **Explanation:**

The map at the suggested threshold should have most of its mass concentrated in a single connected component. It is normal that after thresholding there are a few thousands of very small, disconnected noise blobs. However, their total mass should not exceed 10%. The raw mask (just thresholding) and the mask constructed for the analysis (thresholding + largest connected component + dilation) should significantly overlap. Overlap is defined by the overlapping coefficient  $(\text{size}(\text{Raw AND Constructed})/\text{size}(\text{Raw}))$  that is a number between 0 and 1, the closer to 1, the more they agree.

### **Results:**

Raw mask: At threshold 0.115000, there are 14644 connected components with a total number of voxels of 320153 and a volume of 263617.05 Å<sup>3</sup> (see Fig. 7). The size and percentage of the total number of voxels for the raw mask are listed below (up to 95% of the mass or the first 100 clusters, whatever happens first), the list contains (No. voxels (volume in Å<sup>3</sup>), percentage, cumulated percentage):

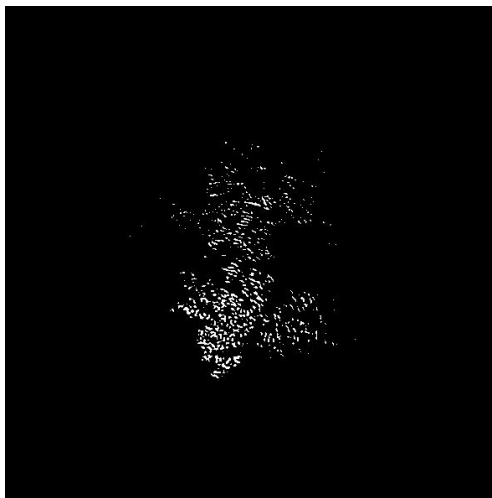
(173365 (142750.40), 54.15, 54.15)(4996 (4113.75), 1.56, 55.71)(3860 (3178.36), 1.21, 56.92)(3629 (2988.15), 1.13, 58.05)(3321 (2734.54), 1.04, 59.09)(2994 (2465.29), 0.94, 60.02)(2489 (2049.47), 0.78, 60.80)(2011 (1655.88), 0.63, 61.43)(1791 (1474.73), 0.56, 61.99)(1172 (965.04), 0.37, 62.35)(1142 (940.33), 0.36, 62.71)(1012 (833.29), 0.32, 63.03)(969 (797.88), 0.30, 63.33)(900 (741.07), 0.28, 63.61)(769 (633.20), 0.24, 63.85)(766 (630.73), 0.24, 64.09)(745 (613.44), 0.23, 64.32)(729 (600.27), 0.23, 64.55)(680 (559.92), 0.21, 64.76)(658 (541.80), 0.21, 64.97)(648 (533.57), 0.20, 65.17)(646 (531.92), 0.20, 65.37)(643 (529.45), 0.20, 65.57)(555 (456.99), 0.17, 65.75)(548 (451.23), 0.17, 65.92)(535 (440.52), 0.17, 66.08)(532 (438.05), 0.17, 66.25)(512 (421.59), 0.16, 66.41)(510 (419.94), 0.16, 66.57)(500 (411.70), 0.16, 66.73)(485 (399.35), 0.15, 66.88)(479 (394.41), 0.15, 67.03)(479 (394.41), 0.15, 67.18)(474 (390.30), 0.15, 67.33)(471 (387.83), 0.15, 67.47)(440 (362.30), 0.14, 67.61)(416 (342.54), 0.13, 67.74)(411 (338.42), 0.13, 67.87)(382 (314.54), 0.12, 67.99)(382 (314.54), 0.12, 68.11)(369 (303.84), 0.12, 68.22)(353 (290.66), 0.11, 68.33)(347 (285.72), 0.11, 68.44)(335 (275.84), 0.10, 68.55)(330 (271.73), 0.10, 68.65)(314 (258.55), 0.10, 68.75)(312 (256.90), 0.10, 68.84)(308 (253.61), 0.10, 68.94)(306 (251.96), 0.10, 69.04)(300 (247.02), 0.09, 69.13)(299 (246.20), 0.09, 69.22)(297 (244.55), 0.09, 69.32)(292 (240.44), 0.09, 69.41)(280 (230.55), 0.09, 69.49)(278 (228.91), 0.09, 69.58)(278 (228.91), 0.09, 69.67)(276 (227.26), 0.09, 69.75)(276 (227.26), 0.09, 69.84)(272 (223.97), 0.08, 69.93)(271 (223.14), 0.08, 70.01)(267 (219.85), 0.08, 70.09)(267 (219.85), 0.08, 70.18)(265 (218.20), 0.08, 70.26)(264 (217.38), 0.08, 70.34)(264 (217.38), 0.08, 70.42)(264 (217.38), 0.08, 70.51)(260 (214.09), 0.08, 70.59)(256 (210.79), 0.08, 70.67)(255 (209.97), 0.08, 70.75)(252 (207.50), 0.08, 70.83)(249 (205.03), 0.08, 70.90)(246 (202.56), 0.08, 70.98)(245 (201.74), 0.08, 71.06)(238 (195.97), 0.07, 71.13)(236 (194.32), 0.07, 71.21)(236 (194.32), 0.07, 71.28)(234 (192.68), 0.07, 71.35)(233 (191.85), 0.07, 71.43)(232 (191.03), 0.07, 71.50)(228 (187.74), 0.07, 71.57)(228 (187.74), 0.07, 71.64)(228 (187.74), 0.07, 71.71)(226 (186.09), 0.07, 71.78)(224 (184.44), 0.07, 71.85)(222 (182.80), 0.07, 71.92)(222 (182.80), 0.07, 71.99)(219 (180.33), 0.07, 72.06)(219 (180.33), 0.07, 72.13)(213 (175.39), 0.07, 72.19)(210 (172.92), 0.07, 72.26)(208 (171.27), 0.06, 72.32)(204 (167.98),

0.06, 72.39)(203 (167.15), 0.06, 72.45)(202 (166.33), 0.06, 72.51)(202 (166.33),  
0.06, 72.58)(199 (163.86), 0.06, 72.64)(198 (163.04), 0.06, 72.70)(196 (161.39),  
0.06, 72.76)(188 (154.80), 0.06, 72.82)

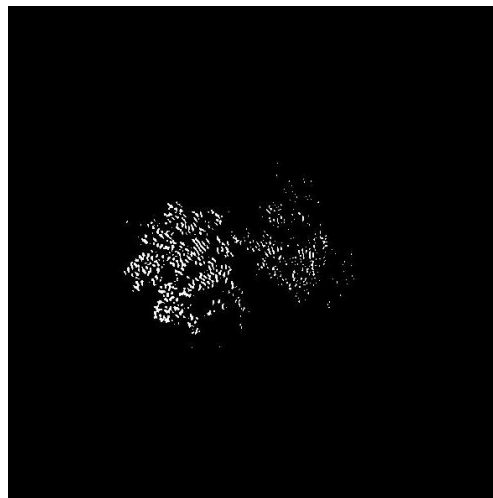
Number of components to reach 95% of the mass: 4782

The average size of the remaining 9862 components is 1.62 voxels ( 0.82  $\text{\AA}^3$ ). Their size go from 4 voxels ( 3.29  $\text{\AA}^3$ ) to 1 voxels ( 0.82  $\text{\AA}^3$ ).

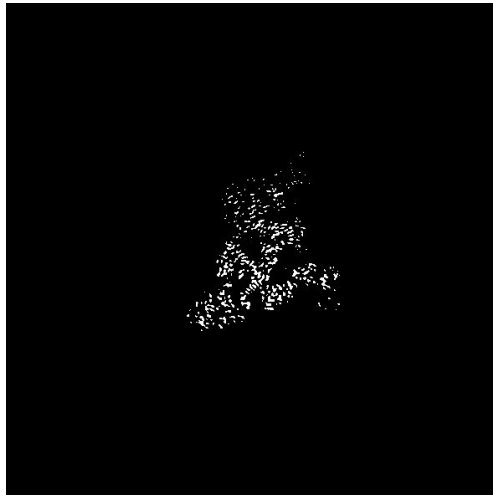
The slices of the raw mask can be seen in Fig. 7.



(a) X Slice 307



(b) Y Slice 251



(c) Z Slice 215

Figure 7: Maximum variance slices in the three dimensions of the raw mask

The following table shows the variation of the mass enclosed at different thresholds (see Fig. 8):

Threshold	Voxel mass	Molecular mass(kDa)	# Aminoacids
0.0340	7511758.00	5124.49	46586.23
0.0680	1749332.00	1193.39	10848.96
0.1020	490698.00	334.75	3043.20
0.1360	180902.00	123.41	1121.91
0.1700	93582.00	63.84	580.37
0.2040	57053.00	38.92	353.83
0.2380	37461.00	25.56	232.32
0.2721	25452.00	17.36	157.85
0.3061	17622.00	12.02	109.29
0.3401	12334.00	8.41	76.49
0.3741	8614.00	5.88	53.42
0.4081	6087.00	4.15	37.75
0.4421	4139.00	2.82	25.67
0.4761	2843.00	1.94	17.63
0.5101	1888.00	1.29	11.71
0.5441	1239.00	0.85	7.68
0.5781	753.00	0.51	4.67
0.6121	433.00	0.30	2.69
0.6461	236.00	0.16	1.46
0.6801	125.00	0.09	0.78
0.7141	59.00	0.04	0.37
0.7482	28.00	0.02	0.17
0.7822	8.00	0.01	0.05
0.8162	4.00	0.00	0.02

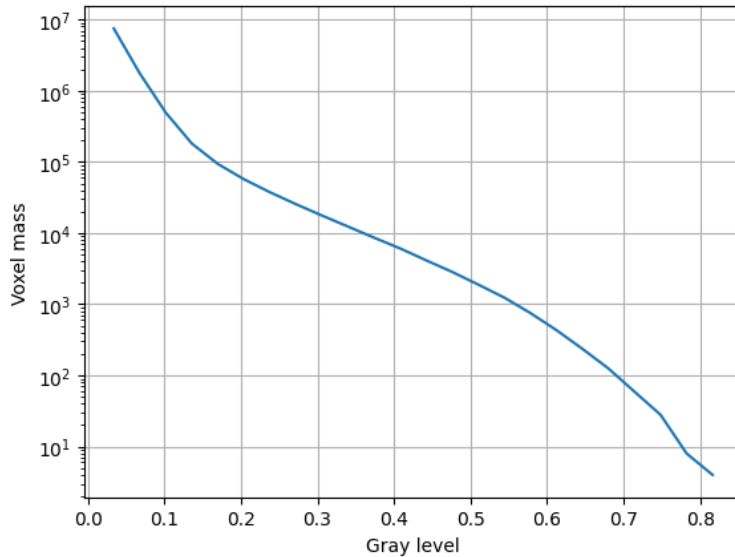


Figure 8: Voxel mass as a function of the gray level.

Constructed mask: After keeping the largest component of the previous mask and dilating it by  $2\text{\AA}$ , there is a total number of voxels of 1112718 and a volume of  $916222.67 \text{\AA}^3$ . The overlap between the raw and constructed mask is 0.57.

**Automatic criteria:** The validation is OK if 1) to keep 95% of the mass we need to keep at most 5 connected components; and 2) the average volume of the blobs outside the given threshold has a size smaller than  $5\text{\AA}^3$ ; and 3) the overlap between the raw mask and the mask constructed for the analysis is larger than 75%.

**WARNINGS:** 2 warnings

1. **There might be a problem of connectivity at this threshold because more than 5 connected components are needed to reach 95% of the total mask. Probably a smaller threshold will not cause this issue.**
2. **There might be a problem in the construction of the mask, because the overlap is smaller than 0.75. A common reason is that the suggested threshold causes too many disconnected**

**components.**

## **2.3 Level 0.c Background analysis**

### **Explanation:**

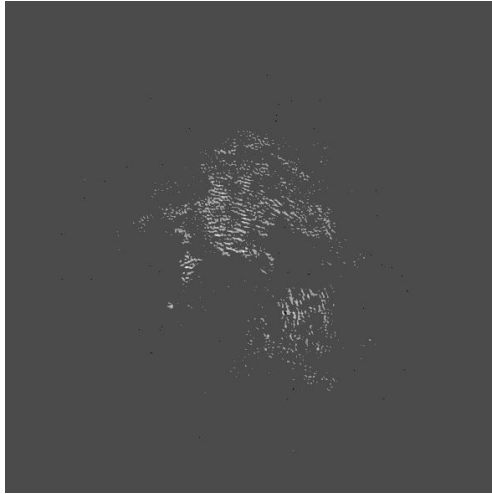
Background is defined as the region outside the macromolecule mask. The background mean should be zero, and the number of voxels with a very low or very high value (below 5 standard deviations of the noise) should be very small and they should be randomly distributed without any specific structure. Sometimes, you can see some structure due to the symmetry of the structure.

### **Results:**

The null hypothesis that the background mean is 0 was tested with a one-sample Student's t-test. The resulting t-statistic and p-value were -256.21 and 0.000000, respectively.

The mean and standard deviation (sigma) of the background were -0.000302 and 0.016754. The percentage of background voxels whose absolute value is larger than 5 times the standard deviation is 0.34 % (see Fig. 9). The same percentage from a Gaussian would be 0.000057% (ratio between the two percentages: 5932.570807).

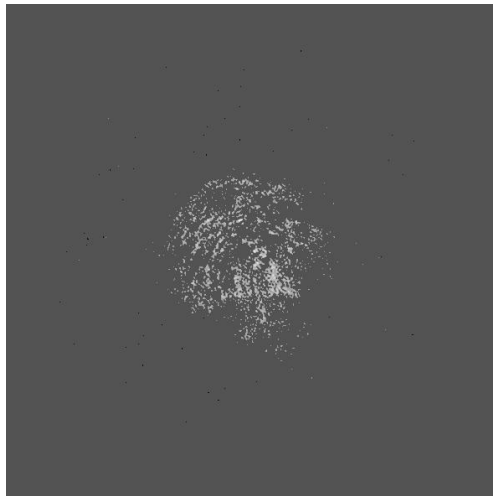
Slices of the background beyond 5\*sigma can be seen in Fig. 9.



(a) X Slice 295



(b) Y Slice 288



(c) Z Slice 341

Figure 9: Maximum variance slices in the three dimensions of the parts of the background beyond  $5 \cdot \sigma$

**Automatic criteria:** The validation is OK if 1) the p-value of the null hypothesis that the background has 0 mean is larger than 0.001; and 2) the number of voxels above or below 5 sigma is smaller than 20 times the amount

expected for a Gaussian with the same standard deviation whose mean is 0.

**WARNINGS:** 2 warnings

1. **The null hypothesis that the background mean is 0 has been rejected because the p-value of the comparison is smaller than 0.001**
2. **There is a significant proportion of outlier values in the background (cdf5 ratio=5932.57)**

## 2.4 Level 0.d B-factor analysis

### **Explanation:**

The B-factor line (see this [link](#) for more details) fitted between 15Å and the resolution reported should have a slope that is between 0 and 300 Å<sup>2</sup>.

### **Results:**

Fig. 10 shows the logarithm (in natural units) of the structure factor (the module squared of the Fourier transform) of the experimental map, its fitted line, and the corrected map. The estimated B-factor was -3.0. The fitted line was  $\log(|F|^2) = -0.8/R^2 + (-13.1)$ .

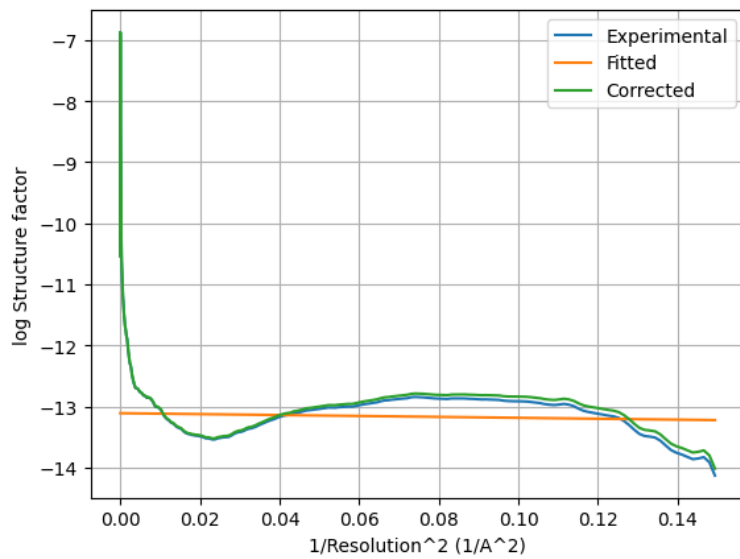
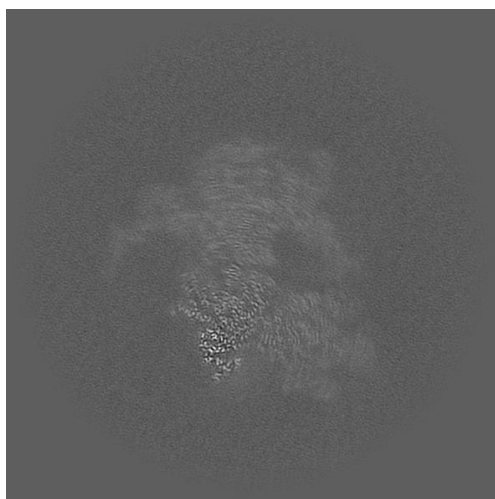
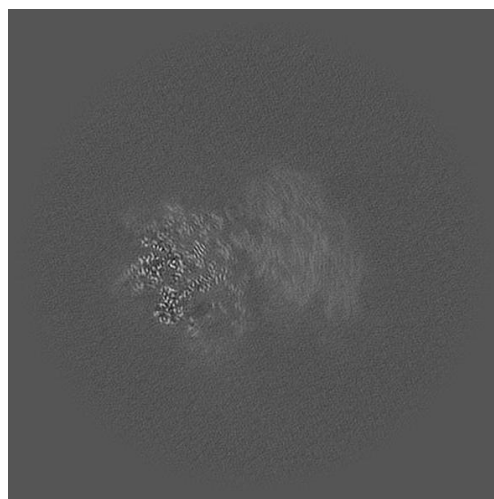


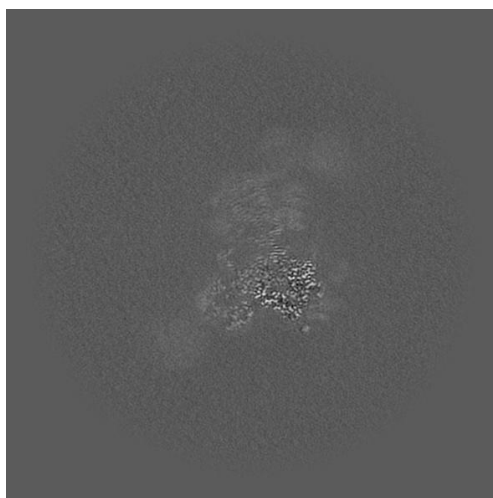
Figure 10: Guinier plot. The X-axis is the square of the inverse of the resolution in Å.



(a) X Slice 307



(b) Y Slice 250



(c) Z Slice 198

Figure 11: Slices of maximum variation in the three dimensions of the B-factor corrected map

**Automatic criteria:** The validation is OK if the B-factor is in the range  $[-300,0]$ .

**STATUS:** OK

## 2.5 Level 0.e Local resolution with DeepRes

### Explanation:

DeepRes (see this [link](#) for more details) measures the local resolution using a neural network that has been trained on the appearance of atomic structures at different resolutions. Then, by comparing the local appearance of the input map to the appearance of the atomic structures a local resolution label can be assigned.

### Results:

**ERROR: The protocol failed.**

**STATUS:** Could not be measured

## 2.6 Level 0.f Local B-factor

### Explanation:

LocBfactor (see this [link](#) for more details) estimates a local resolution B-factor by decomposing the input map into a local magnitude and phase term using the spiral transform.

### Results:

Fig. 12 shows the histogram of the local B-factor according to LocBfactor. Some representative percentiles are:

Percentile	Local B-factor ( $\text{\AA}^{-2}$ )
2.5%	-158.16
25%	-133.46
50%	-121.66
75%	-110.27
97.5%	-88.43

Fig. 13 shows some representative views of the local B-factor.

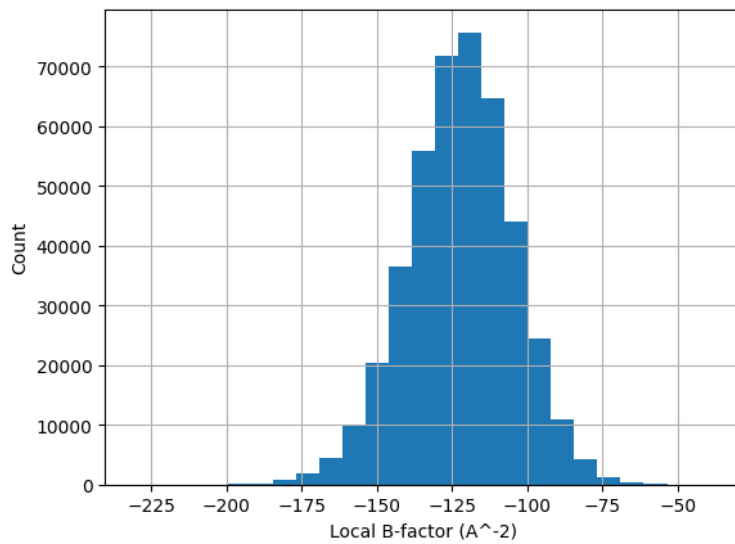
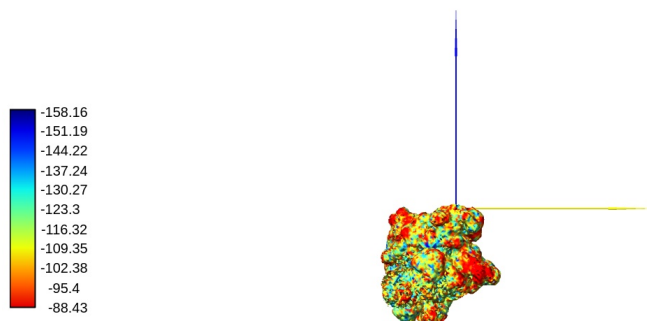
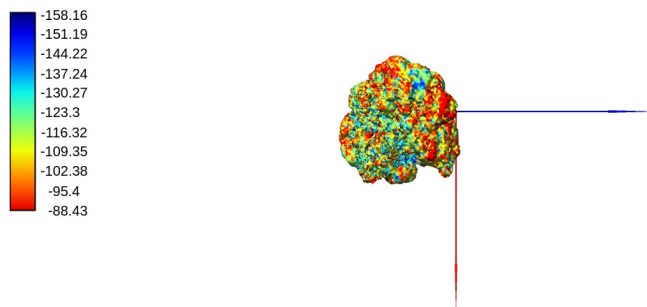


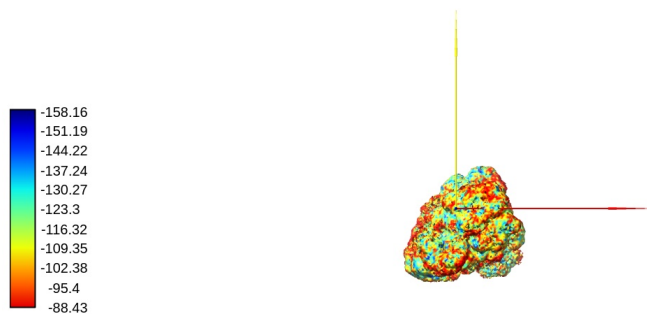
Figure 12: Histogram of the local B-factor according to LocBfactor.



(a) View 1



(b) View 2



(c) View 3

Figure 13: Local B-factor according to LocBfactor. Views generated by ChimeraX at a the following X, Y, Z angles: View 1 (0, -90, -90), View 2 (-90, 0, -90), View 3 (0, 0, 0).

**Automatic criteria:** The validation is OK if the median B-factor is in the range [-300,0].

STATUS: [OK](#)

## 2.7 Level 0.g Local Occupancy

### Explanation:

LocOccupancy (see this [link](#) for more details) estimates the occupancy of a voxel by the macromolecule.

### Results:

**ERROR: The protocol failed.**

STATUS: **Could not be measured**

## 2.8 Level 0.h Hand correction

### Explanation:

Deep Hand (see this [link](#) for more details) determines the correction of the hand for those maps with a resolution smaller than 5Å. The method calculates a value between 0 (correct hand) and 1 (incorrect hand) using a neural network to assign its hand.

### Results:

Deep hand assigns a score of 0.280 to the input volume.

**Automatic criteria:** The validation is OK if the deep hand score is smaller than 0.5.

STATUS: [OK](#)

## 3 Half maps

Half map 1: emd\_51709\_half\_map\_1.map

SHA256 hash: 3122dfcc8d628cf9dd0eb0998e9c126bf60bafba51d33c02433553ebd233522a

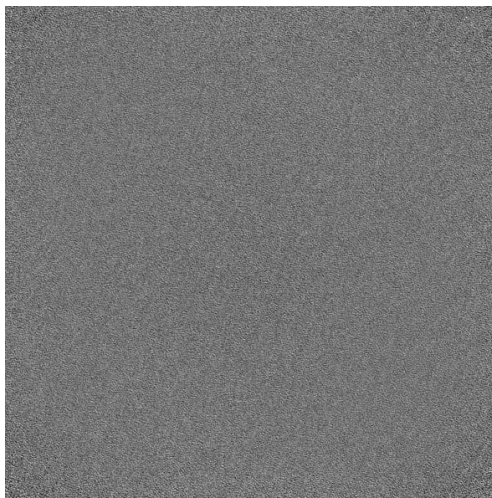
Half map 2: emd\_51709\_half\_map\_2.map

SHA256 hash: 460fdf791de51000bf13bee260b174a724dde85d7b812d547dc9854ad60cc51f

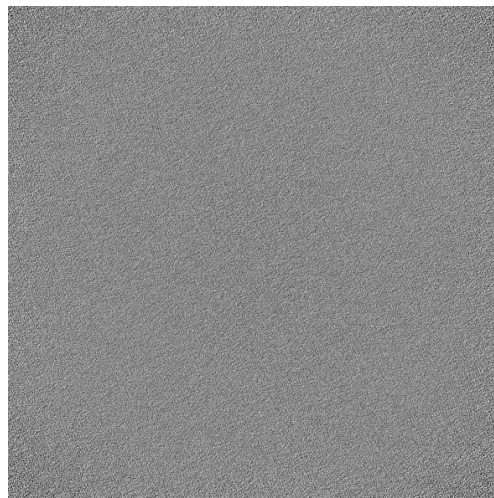
Slices of the first half map can be seen in Fig. 14.

Slices of the second half map can be seen in Fig. 15.

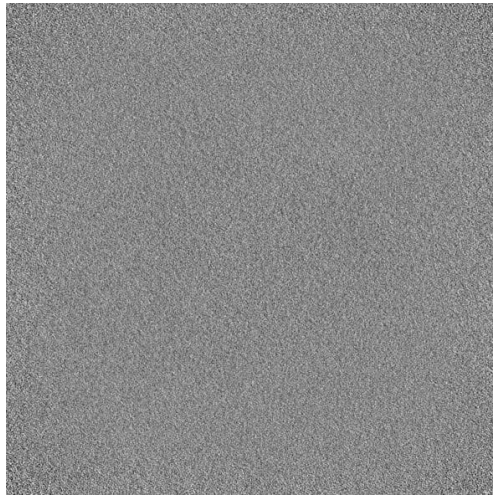
Slices of the difference between both maps can be seen in Fig. 16. There should not be any structure in this difference. Sometimes some patterns are seen if the map is symmetric.



(a) X Slice 0

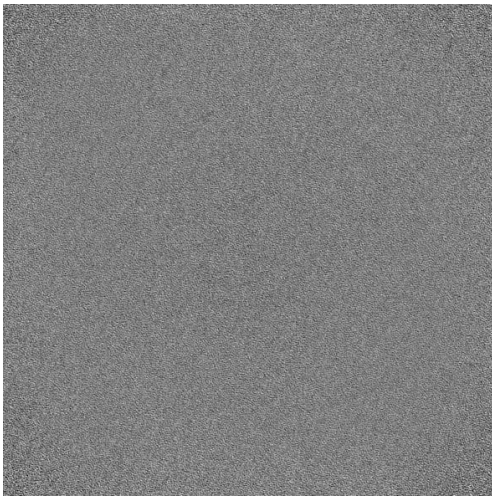


(b) Y Slice 0

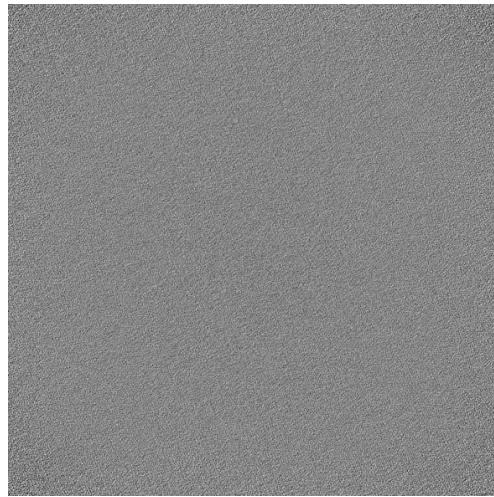


(c) Z Slice 0

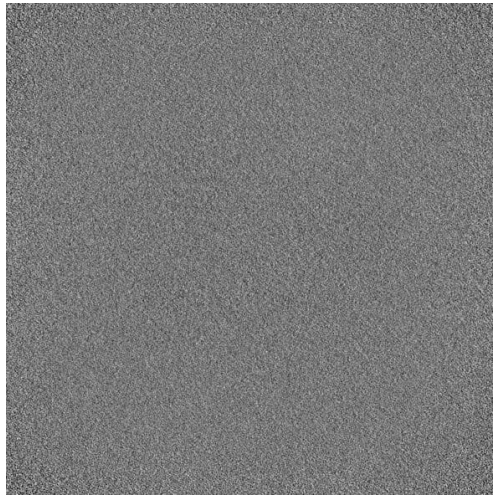
Figure 14: Slices of maximum variation in the three dimensions of Half 1



(a) X Slice 0

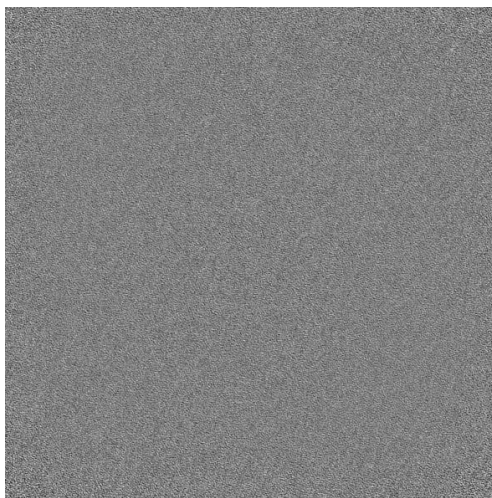


(b) Y Slice 0

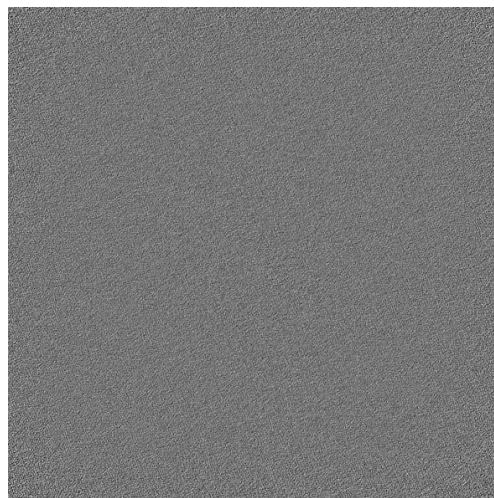


(c) Z Slice 0

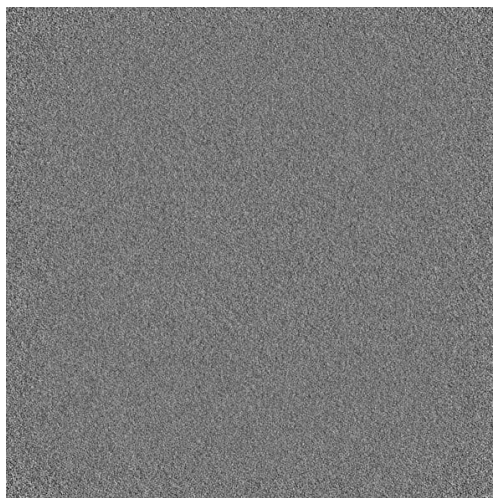
Figure 15: Slices of maximum variation in the three dimensions of Half 2



(a) X Slice 0



(b) Y Slice 0



(c) Z Slice 0

Figure 16: Slices of maximum variation in the three dimensions of the difference Half1-Half2.

## 4 Level 1 analysis

### 4.1 Level 1.a Global resolution

**Explanation:** The Fourier Shell Correlation (FSC) between the two half maps is the most standard method to determine the global resolution of a map. However, other measures exist such as the Spectral Signal-to-Noise Ratio and the Differential Phase Residual. There is a long debate about the right thresholds for these measures. Probably, the most clear threshold is the one of the SSNR (SSNR=1). For the DPR we have chosen  $103.9^\circ$  and for the FSC, the standard 0.143. For a deep discussion of all these thresholds, see this [link](#). Note that these thresholds typically result in resolution values that are at the lower extreme of the local resolution range, meaning that this resolution is normally in the first quarter. It should not be understood as the average resolution of the map.

Except for the noise, the FSC and DPR should be approximately monotonic. They should not have any “coming back” behavior. If they have, this is typically due to the presence of a mask in real space or non-linear process-

ing.

**Results:**

Fig. 17 shows the FSC and the 0.143 threshold. The resolution according to the FSC is 4.54Å. The map information is well preserved (FSC>0.9) up to 21.22Å.

Fig. 18 shows the DPR and the 103.9° threshold. The resolution according to the DPR is 3.32Å.

Fig. 19 shows the SSNR and the SSNR=1 threshold. The resolution according to the SSNR is 4.47Å.

The mean resolution between the three methods is 4.11Å and its range is within the interval [ 3.32, 4.54]Å.

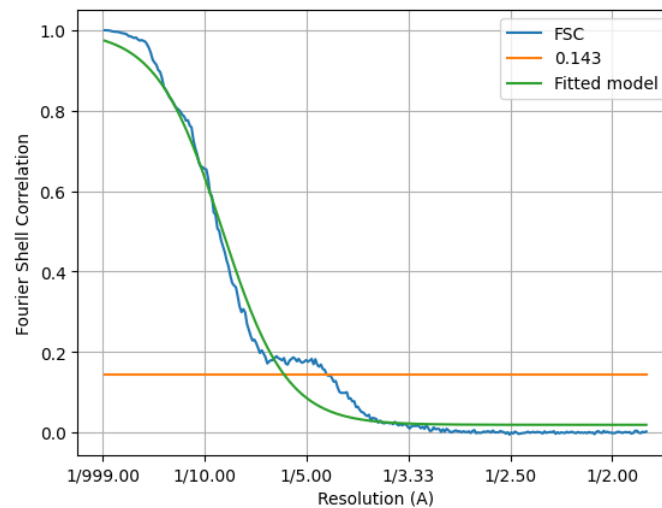


Figure 17: Fourier Shell correlation between the two halves.

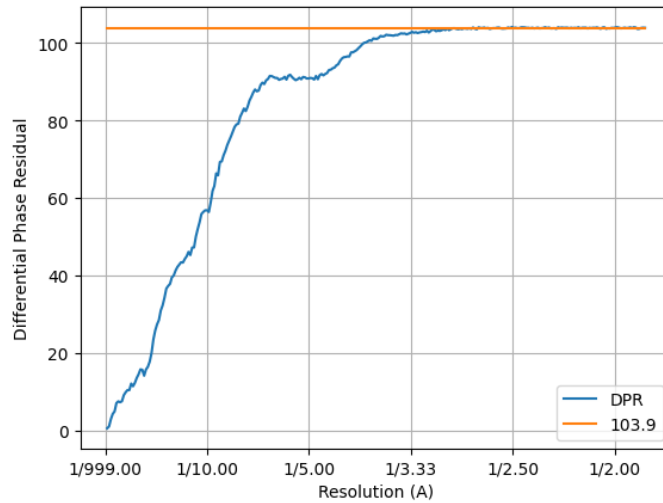


Figure 18: Differential Phase Residual between the two halves.

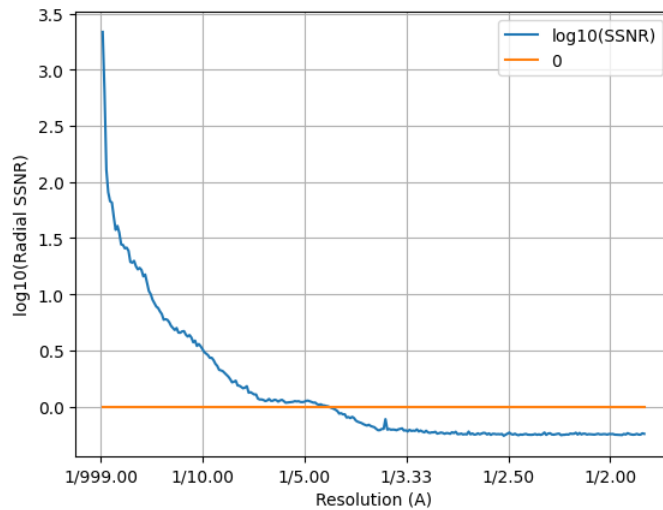


Figure 19: Spectral Signal-to-Noise Ratio estimated from the two halves.

**Automatic criteria:** The validation is OK if the user provided resolution is larger than 0.8 times the resolution estimated by 1) FSC, 2) DPR, and 3) SSNR.

**WARNINGS:** 3 warnings

1. **The reported resolution, 2.58 Å, is particularly high with respect to the resolution calculated by the FSC, 4.54 Å**
2. **The reported resolution, 2.58 Å, is particularly high with respect to the resolution calculated by the DPR, 3.32Å.**
3. **The reported resolution, 2.58 Å, is particularly high with respect to the resolution calculated by the SSNR, 4.47Å.**

## 4.2 Level 1.b FSC permutation

### Explanation:

This method (see this [link](#) for more details) calculates a global resolution by formulating a hypothesis test in which the distribution of the FSC of noise is calculated from the two maps.

### Results:

The resolution at 1% of FDR was 2.8. The estimated B-factor was -30.42  
Fig. 20 shows the estimated FSC and resolution.

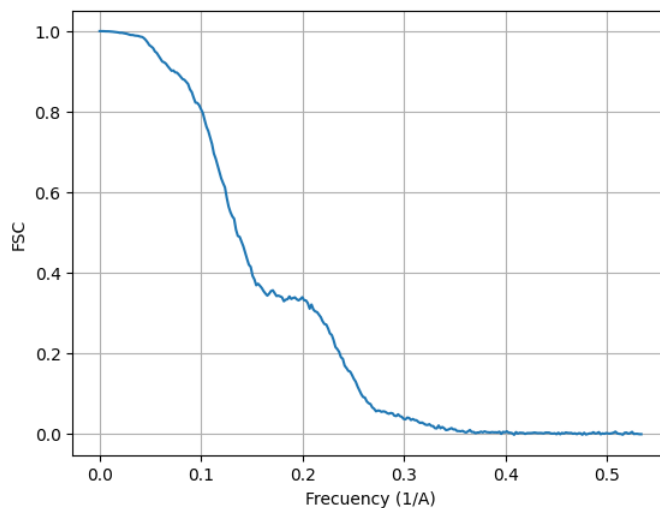


Figure 20: FSC and resolution estimated by a permutation test.

**Automatic criteria:** The validation is OK if the user provided resolu-

tion is larger than 0.8 times the resolution estimated by FSC permutation.

**STATUS:** [OK](#)

### 4.3 Level 1.c Local resolution with Blocres

**Explanation:**

This method (see this [link](#) for more details) computes a local Fourier Shell Correlation (FSC) between the two half maps.

**Results:**

Fig. 21 shows the histogram of the local resolution according to Blocres. Some representative percentiles are:

Percentile	Resolution(Å)
2.5%	2.60
25%	2.93
50%	3.23
75%	3.60
97.5%	4.68

The reported resolution, 2.58 Å, is at the percentile 1.7. Fig. 22 shows some representative views of the local resolution.

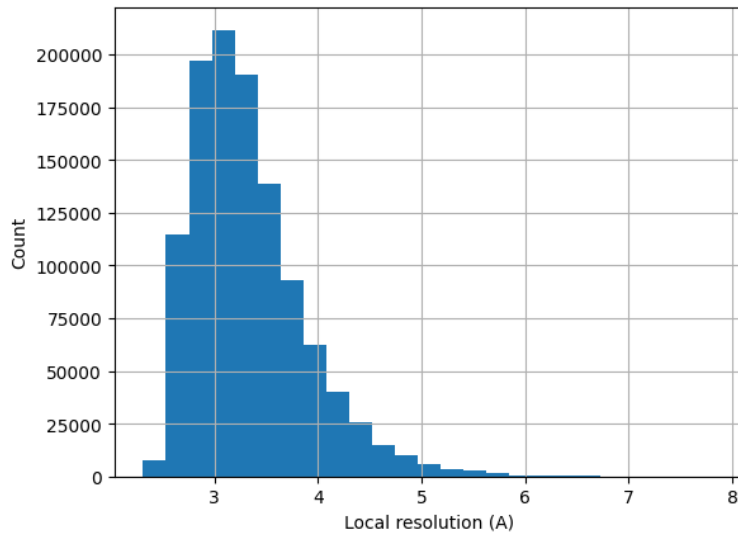
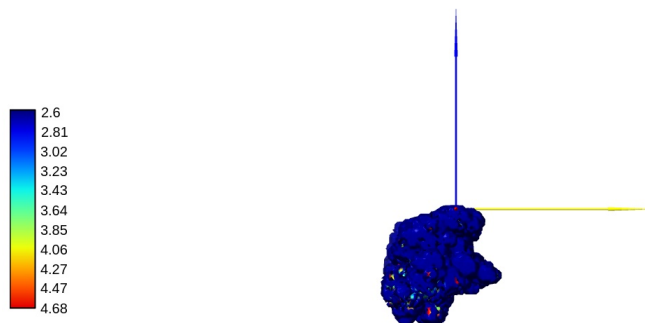
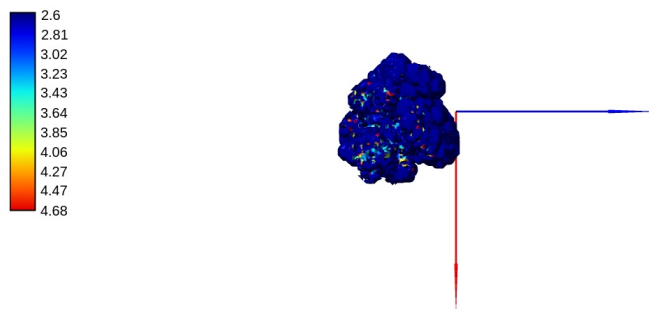


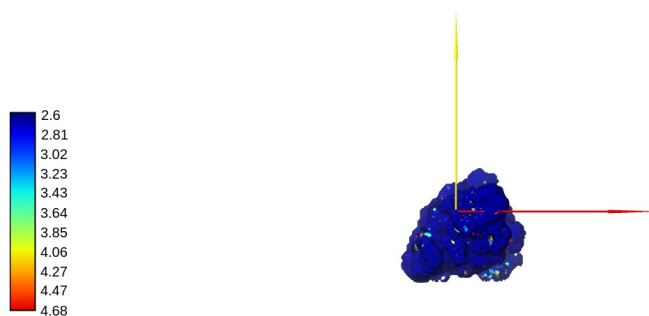
Figure 21: Histogram of the local resolution according to blocres.



(a) View 1



(b) View 2



(c) View 3

Figure 22: Local resolution according to Blocres. Views generated by ChimeraX at the following X, Y, Z angles: View 1 (0, -90, -90), View 2 (-90, 0, -90), View 3 (0, 0, 0).

**Automatic criteria:** The validation is OK if the percentile of the user provided resolution is larger than 0.1% of the percentile of the local resolution as estimated by BlocRes.

STATUS: [OK](#)

#### 4.4 Level 1.d Local resolution with Resmap

**Explanation:**

This method (see this [link](#) for more details) is based on a test hypothesis testing of the superiority of signal over noise at different frequencies.

**Results:**

**ERROR: The protocol failed.**

STATUS: [Could not be measured](#)

#### 4.5 Level 1.e Local resolution with MonoRes

**Explanation:**

MonoRes (see this [link](#) for more details) evaluates the local energy of a point with respect to the distribution of energy in the noise. This comparison is performed at multiple frequencies and for each one, the monogenic transformation separates the amplitude and phase of the input map. Then the energy of the amplitude within the map is compared to the amplitude distribution observed in the noise, and a hypothesis test is run for every voxel to check if its energy is significantly above the level of noise.

**Results:**

Fig. [23](#) shows the histogram of the local resolution according to MonoRes. Some representative percentiles are:

Percentile	Resolution( $\text{\AA}$ )
2.5%	2.53
25%	3.97
50%	5.33
75%	7.60
97.5%	9.80

The reported resolution, 2.58  $\text{\AA}$ , is at the percentile 3.2. Fig. 24 shows some representative views of the local resolution

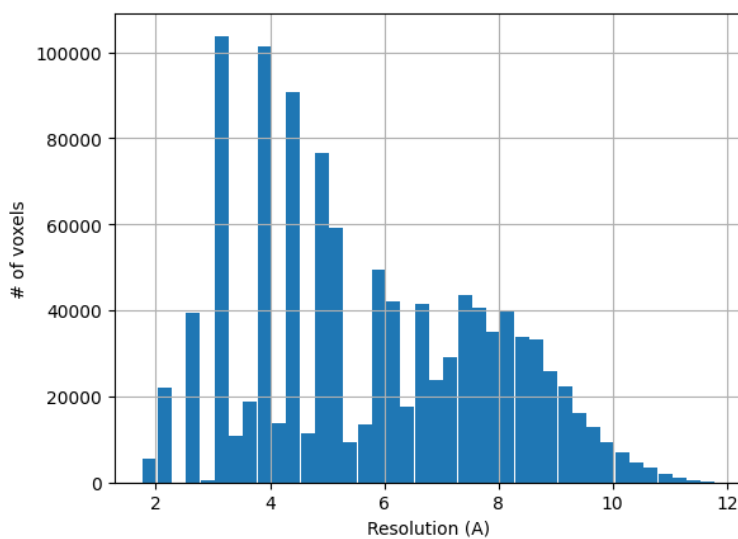
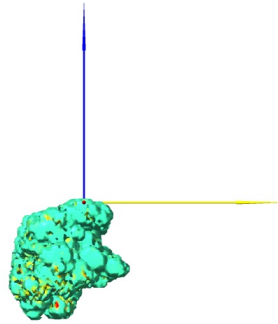
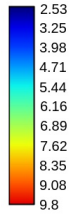
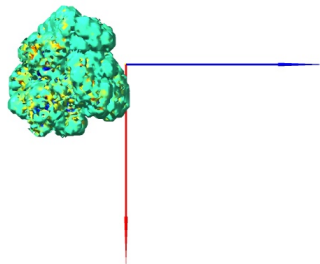
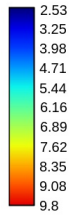


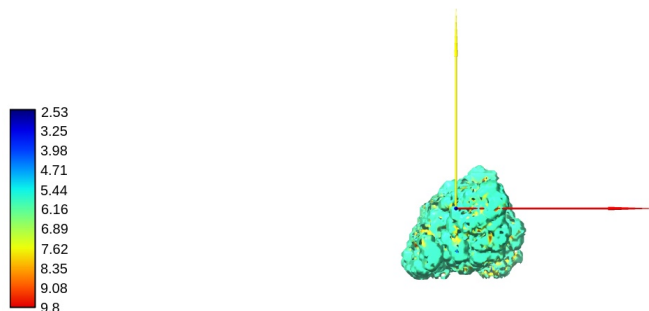
Figure 23: Histogram of the local resolution according to MonoRes.



(a) View 1



(b) View 2



(c) View 3

Figure 24: Local resolution according to MonoRes. Views generated by ChimeraX at a the following X, Y, Z angles: View 1 (0, -90, -90), View 2 (-90, 0, -90), View 3 (0, 0, 0).

**Automatic criteria:** The validation is OK if the percentile of the user provided resolution is larger than 0.1% of the percentile of the local resolution as estimated by MonoRes.

**STATUS:** [OK](#)

## 4.6 Level 1.f Local and directional resolution with MonoDir

### Explanation:

MonoDir (see this [link](#) for more details) extends the concept of local resolution to local and directional resolution by changing the shape of the filter applied to the input map. The directional analysis can reveal image alignment problems.

The histogram of best resolution voxels per direction (Directional Histogram 1D) shows how many voxels in the volume have their maximum resolution in that direction. Directions are arbitrarily numbered from 1 to N. This histogram should be relatively flat. We perform a Kolmogorov-Smirnov test to check its uniformity. If the null hypothesis is rejected, then the di-

rectional resolution is not uniform. It does not mean that it is wrong, and it could be caused by several reasons: 1) the angular distribution is not uniform, 2) there are missing directions, 3) there is some anisotropy in the data (including some preferential directional movement).

Ideally, the radial average of the minimum, maximum, and average resolution at each voxel (note that these are spatial radial averages) should be flat and as low as possible. If they show some slope, this is associated with inaccuracies in the angular assignment. These averages make sense when the shells are fully contained within the protein. As the shells approach the outside of the protein, these radial averages make less sense.

### Results:

Fig. 25 shows the 1D directional histogram and Fig. 26 the 2D directional histogram. We compared the 1D directional histogram to a uniform distribution using a Kolmogorov-Smirnov test. The D statistic was 0.086963, and the p-value of the null hypothesis 0.000000.

The radial average of the minimum, maximum and average resolution at each voxel is shown in Fig. 27. The overall mean of the directional resolution is 3.91

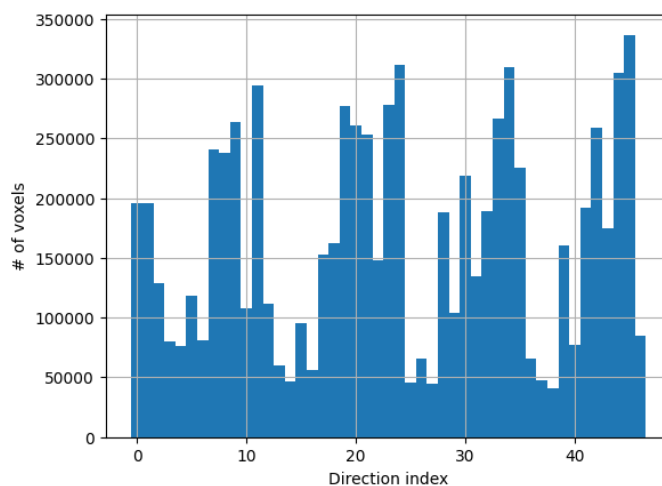


Figure 25: Histogram 1D of the best direction at each voxel.

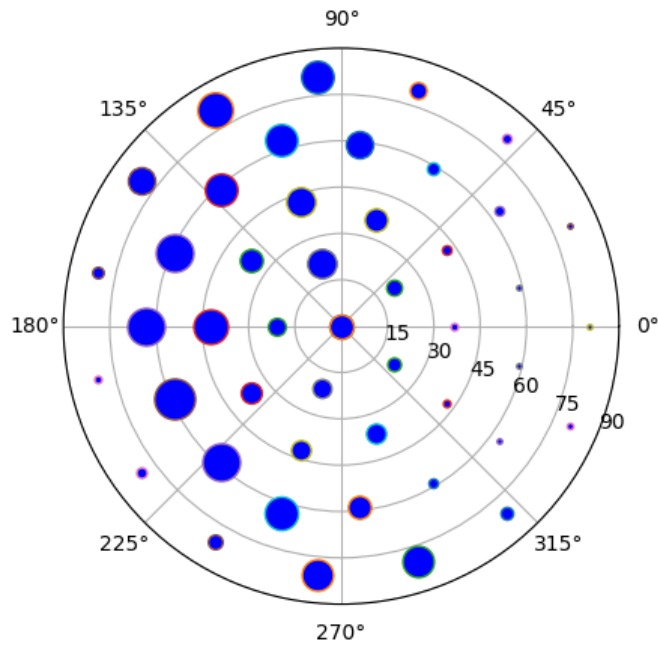


Figure 26: Histogram 2D of the best direction at each voxel. The azimuthal rotation is circular, while the tilt angle is the radius. The size of the point is proportional to the number of voxels whose maximum resolution is in that direction (this count can be seen in Fig. 25).

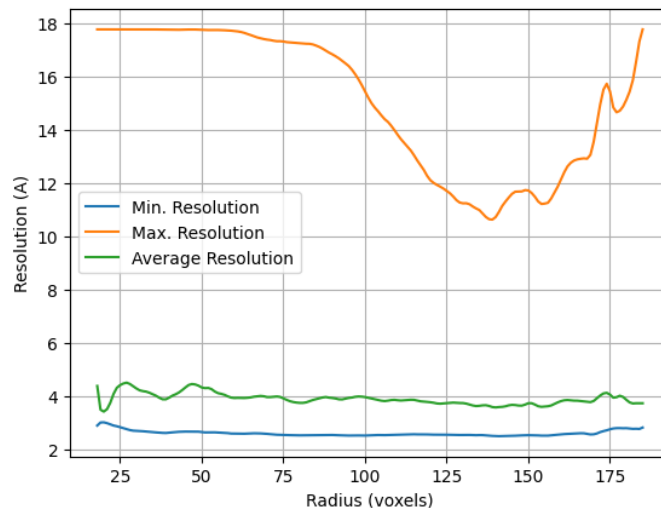


Figure 27: Radial averages (in space) of the minimum, maximum and average resolution at each voxel.

**Automatic criteria:** The validation is OK if 1) the null hypothesis that the directional resolution is not uniform is not rejected with a threshold of 0.001 for the p-value, and 2) the resolution provided by the user is not smaller than 0.8 times the average directional resolution.

**WARNINGS:** 2 warnings

1. **The distribution of best resolution is not uniform in all directions. The associated p-value is 0.000000.**
2. **The resolution reported by the user, 2.58Å, is at least 80% smaller than the average directional resolution, 3.91 Å.**

## 4.7 Level 1.g Fourier Shell Occupancy

### Explanation:

This method (see this [link](#) for more details) calculates the anisotropy of the energy distribution in Fourier shells. This is an indirect measure of anisotropy of the angular distribution or the presence of heterogeneity. A natural threshold for this measure is 0.5. However, 0.9 and 0.1 are also interesting values that define the frequency at which the occupancy is 90% and 10%, respectively. This region is shaded in the plot.

## Results:

Fig. 28 shows the Fourier Shell Occupancy and its anisotropy. The directional resolution is shown in Fig. 29. The resolution according to the FSO is 2.66Å. Fourier shells are occupied at between 90 and than 10% in the range [ 3.52, 2.38]Å.

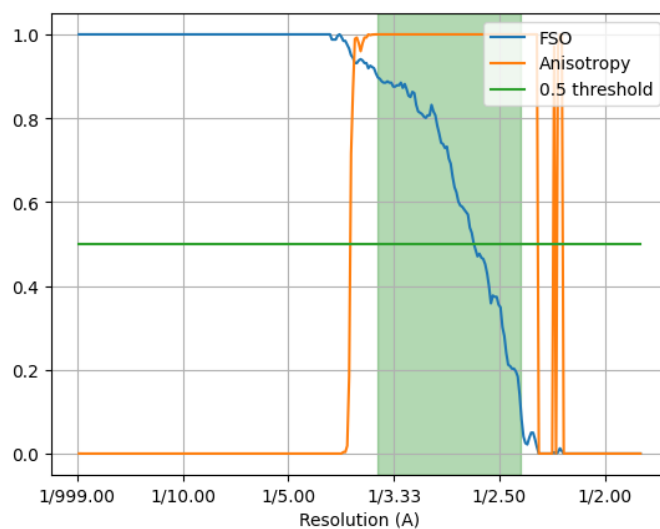


Figure 28: FSO and anisotropy.

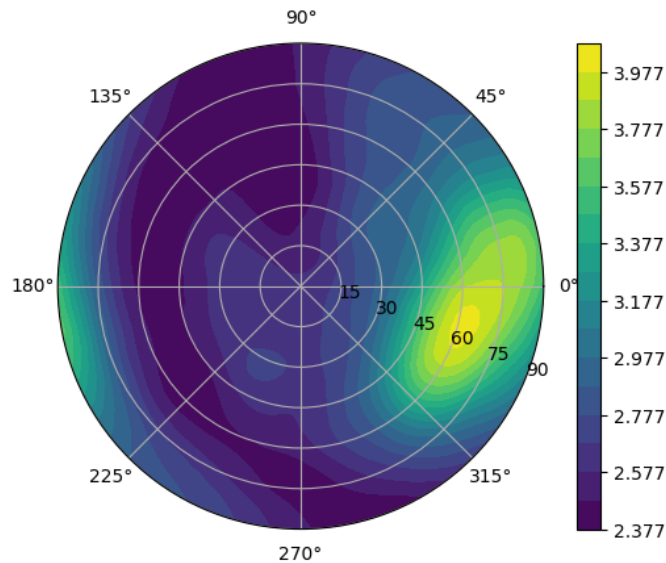


Figure 29: Directional resolution in the projection sphere.

**Automatic criteria:** The validation is OK if the resolution provided by the user is not smaller than 0.8 times the resolution estimated by the first cross of FSO below 0.5.

**STATUS:** OK

## 4.8 Level 1.h Fourier Shell Correlation 3D

### Explanation:

This method (see this [link](#) for more details) analyzes the FSC in different directions and evaluates its homogeneity.

### Results:

**ERROR:** The protocol failed.

**STATUS:** Could not be measured