



Public Data-based Report



Validation Report Service

Cryo-EM Map Validation Report

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

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

Basic Entry Information:

EMDB ID: [EMD-68835](#)

Title: Cryo-EM structure of a triangular-prism protein-DNA complex (D17-11-2t-10a/10b-T)

Authors: [See EMDB entry link](#)

Deposited on: 2026-01-30T00:00:00

Reported Resolution: 18.61 Å

Contact Us:

Instruct Image Processing Center ([I²PC](#))
Biocomputing Unit ([BCU](#))
i2pc@cnb.csic.es
[VRS Website](#)

National Center for Biotechnology (CNB)
St/ Darwin, 3 (Autonomous University of Madrid)
28049 Cantoblanco, Madrid (Spain)

Last update: **September 5, 2026, 5:18am**

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 is no problem with the suggested threshold. There seems to be a problem with the map's background (see Sec. [2.3](#)).

The overall score (passing tests) of this report is 1 out of 3 evaluable items.

0.a Mass analysis	Sec. 2.1	1 warnings
0.b Mask analysis	Sec. 2.2	OK
0.c Background analysis	Sec. 2.3	2 warnings
0.d B-factor analysis	Sec. 2.4	Does not apply

Summary of the warnings across sections.

Section 2.1 (0.a Mass analysis)

1. **The center of mass in Z may be significantly shifted. This is common when the refinement is applied exclusively to one protein region.**

Section 2.3 (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=10235.18)**

Contents

1	Input data	6
2	Level 0 analysis	13
2.1	Level 0.a Mass analysis	13
2.2	Level 0.b Mask analysis	14
2.3	Level 0.c Background analysis	18
2.4	Level 0.d B-factor analysis	20
3	Half maps	21

1 Input data

Input map: emd_68835.map

SHA256 hash: 94817718b3559879fbb2274fd5dfbec908afc28208f08aab5d1e86a4c11306a0

Voxel size: 0.970000 (Å)

Visualization threshold: 0.069200

Resolution estimated by user: 18.61

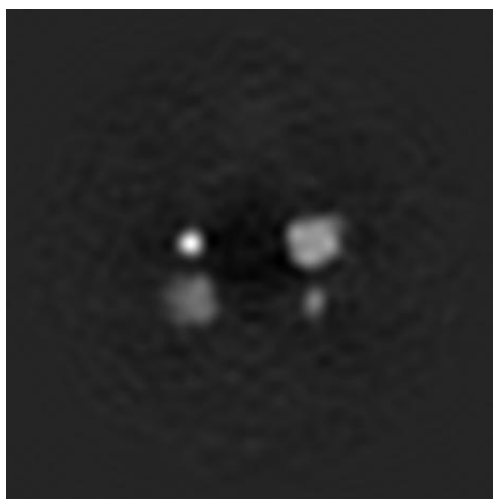
Orthogonal slices of the input map

Explanation:

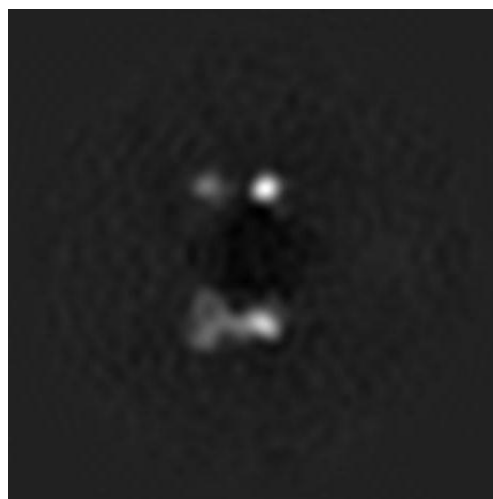
In the orthogonal slices of the map, the noise outside the protein should not have any structure (stripes going out, small blobs, particularly high or low densities, ...)

Results:

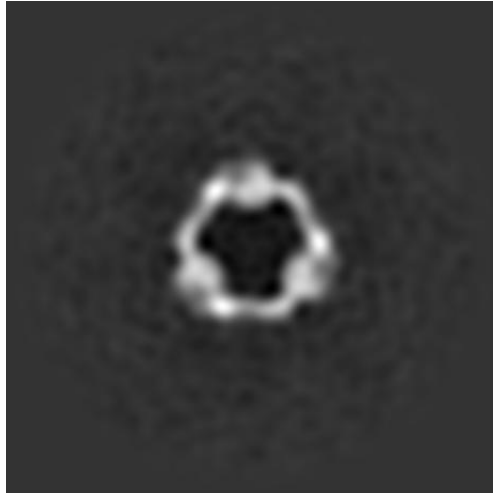
See Fig. 1.



(a) X Slice 230



(b) Y Slice 230



(c) Z Slice 230

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

Orthogonal slices of maximum variance of the input map

Results:

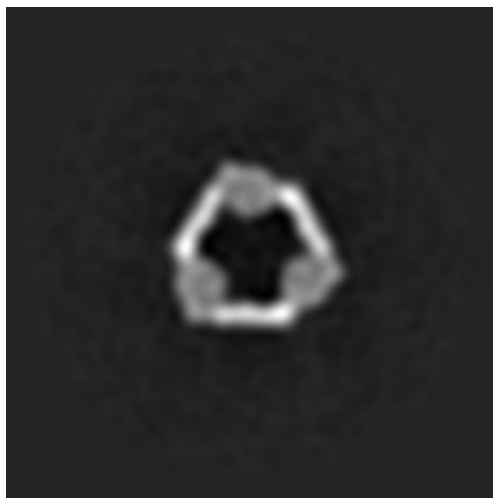
See Fig. 2.



(a) X Slice 188



(b) Y Slice 174



(c) Z Slice 241

Figure 2: Slices of maximum variation in the three dimensions

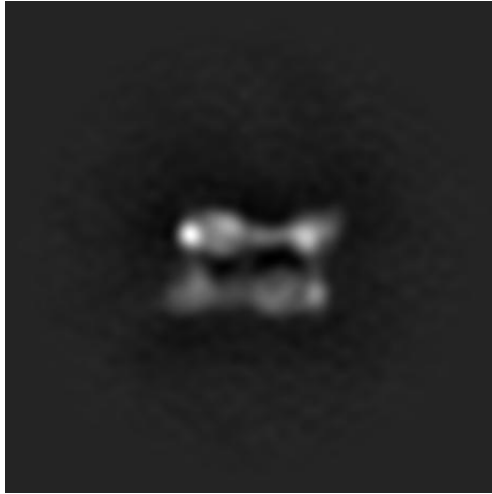
Orthogonal projections of the input map

Explanation:

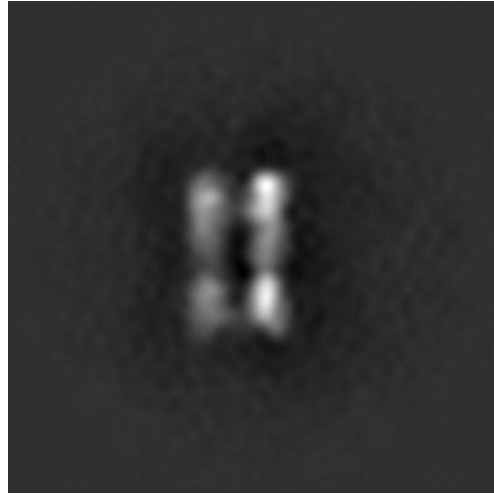
In the projections there should not be stripes (this is an indication of directional overweighting, or angular attraction), and there should not be a dark halo around or inside the structure (this is an indication of incorrect CTF correction or the reconstruction of a biased map).

Results:

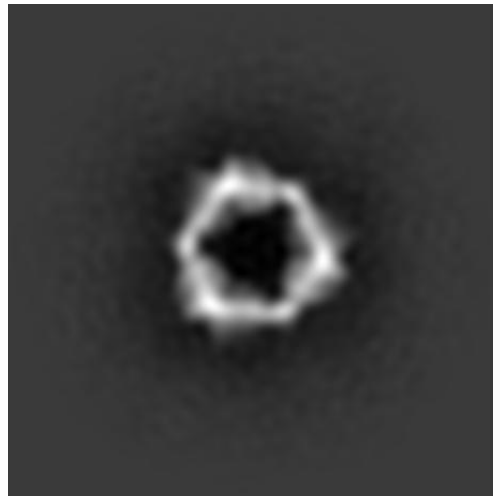
See Fig. [3](#).



(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.

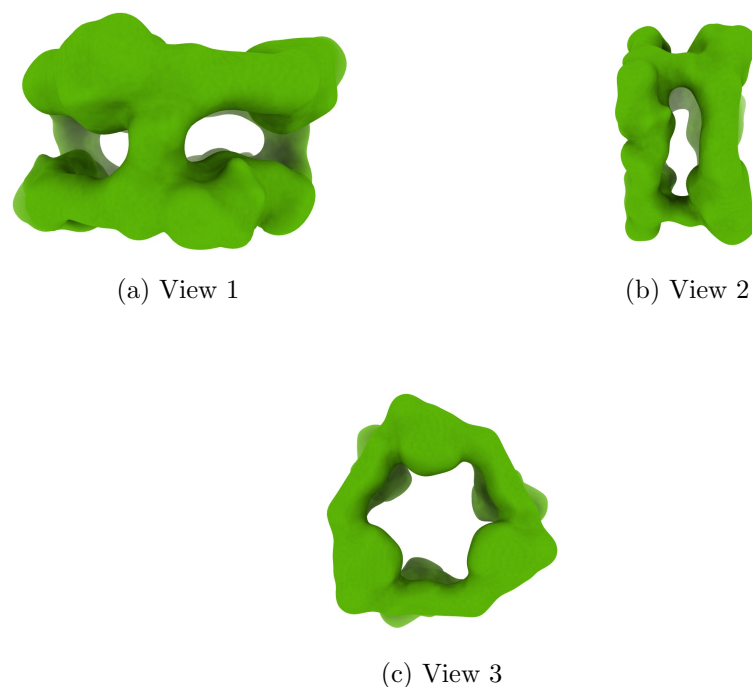


Figure 4: Isosurface at threshold=0.069200. Views generated by ChimeraX at a the following X, Y, Z angles: View 1 (0, -90, -90), View 2 (-90, 0, -90), View 3 (0, 0, 0).

Orthogonal slices of maximum variance of the mask with hard borders

Explanation:

The mask with hard borders has been calculated at the suggested threshold 0.069200, the largest connected component was selected, and then dilated by 2Å.

Results:
See Fig. 5.

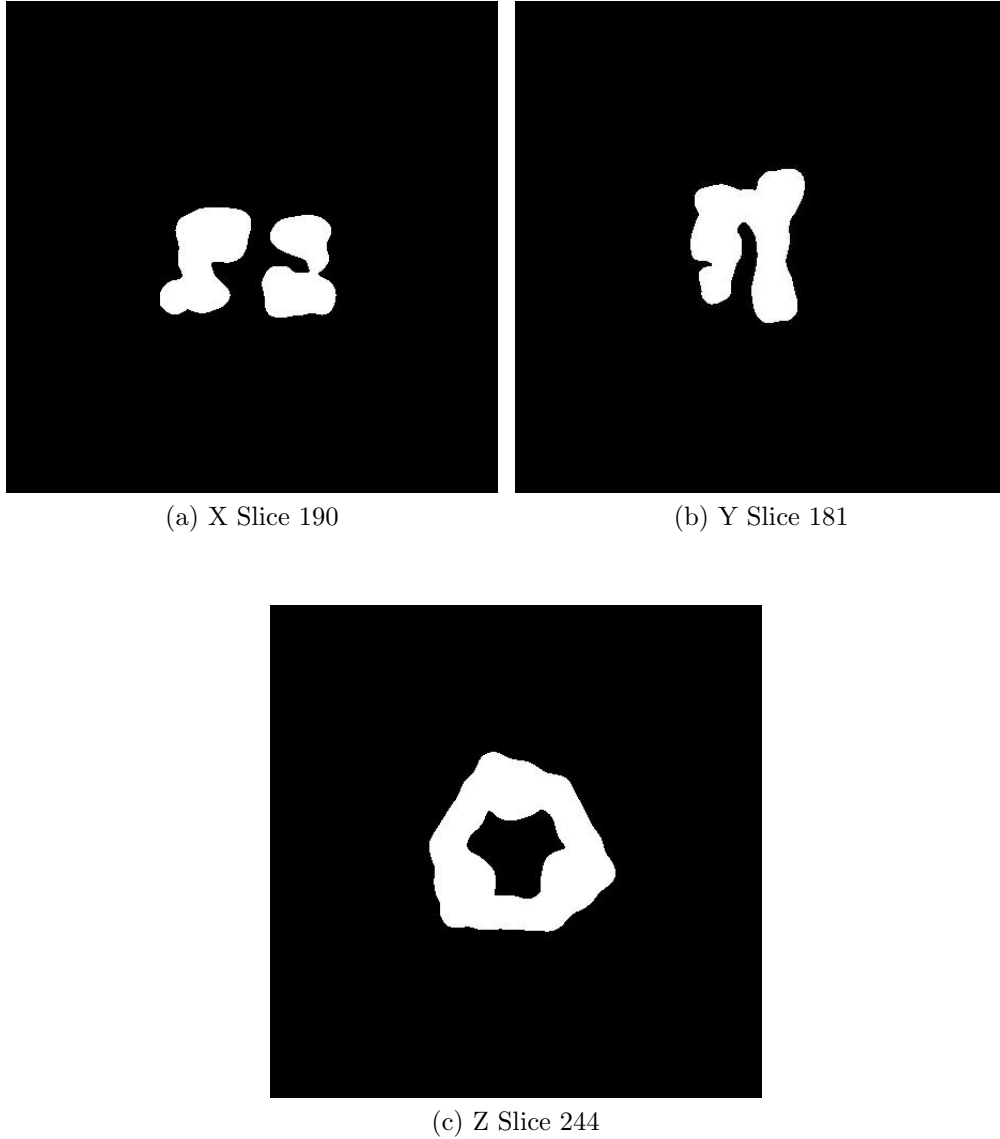


Figure 5: Slices of maximum variation in the three dimensions of the mask with hard borders

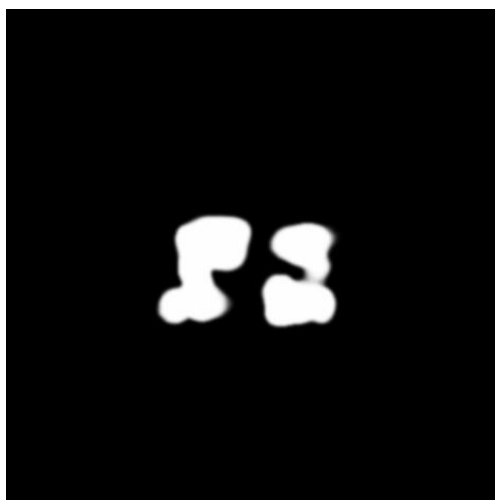
Orthogonal slices of maximum variance of the mask with soft borders

Explanation:

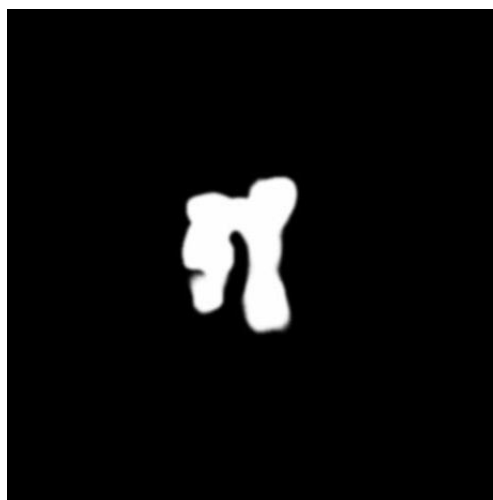
The mask with soft borders has been calculated at the suggested threshold 0.069200, the largest connected component was selected, and then dilated by

2Å.

Results:
See Fig. 6.



(a) X Slice 191



(b) Y Slice 181



(c) Z Slice 243

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 144.53 and 131.92 $\text{\AA}$, respectively. There is a decentering ratio $(\text{abs}(\text{Right-Left})/\text{Size})\%$ of 2.83%

The space from the left and right in Y are 135.80 and 132.89 $\text{\AA}$, respectively. There is a decentering ratio $(\text{abs}(\text{Right-Left})/\text{Size})\%$ of 0.65%

The space from the left and right in Z are 159.08 and 184.30 $\text{\AA}$, respectively. There is a decentering ratio $(\text{abs}(\text{Right-Left})/\text{Size})\%$ of 5.65%

The center of mass is at (x,y,z)=(219.21,223.41,503.77). The decentering of the center of mass (abs(Center)/Size)% is 2.34, 1.43, and 59.52, 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: 1 warnings

1. **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, there 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 (size(Raw AND Constructed)/size(Raw)) that is a number between 0 and 1, the closer to 1, the more they agree.

Results:

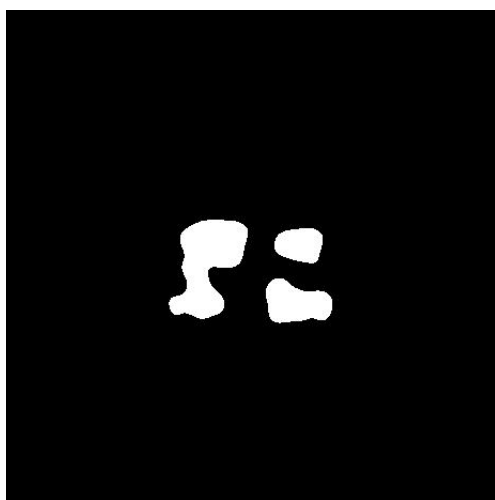
Raw mask: At threshold 0.069200, there are 1 connected components with a total number of voxels of 720047 and a volume of 657167.46 Å³ (see Fig. 7). The size and percentage of the total number of voxels for the raw mask are listed below (up to 95% of the mass or the first 100 clusters, whatever happens first), the list contains (No. voxels (volume in Å³), percentage, cumulated percentage):

(720047 (657167.46), 100.00, 100.00)

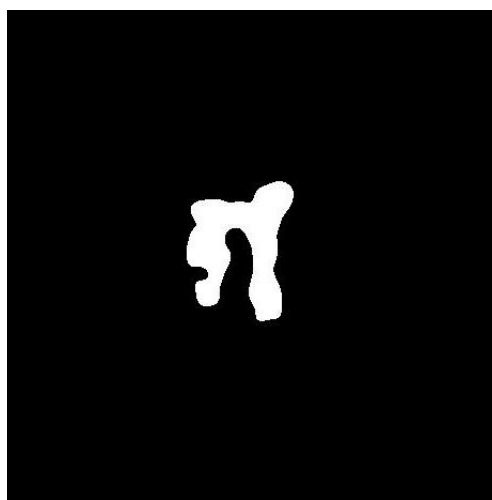
Number of components to reach 95% of the mass: 1

The average size of the remaining 0 components is nan voxels (nan $\text{\AA}^3$).
Their size go from 720047 voxels (657167.46 $\text{\AA}^3$) to 720047 voxels (657167.46 $\text{\AA}^3$).

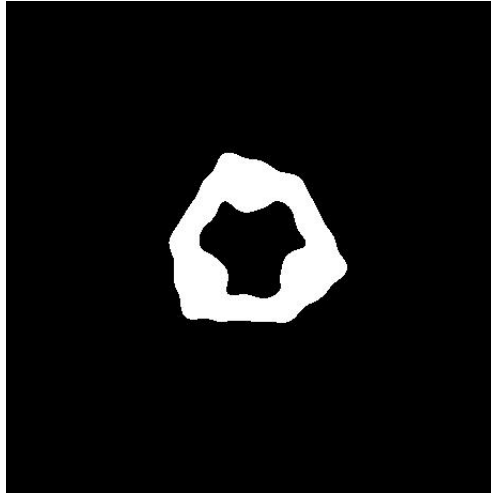
The slices of the raw mask can be seen in Fig. 7.



(a) X Slice 188



(b) Y Slice 179



(c) Z Slice 244

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
0.0198	1152177.00	871.22	7920.17
0.0396	945039.00	714.59	6496.28
0.0593	788475.00	596.21	5420.05
0.0791	654655.00	495.02	4500.16
0.0989	534951.00	404.50	3677.30
0.1187	427660.00	323.38	2939.77
0.1385	331168.00	250.41	2276.48
0.1583	263088.00	198.93	1808.49
0.1780	215760.00	163.15	1483.15
0.1978	169709.00	128.33	1166.60
0.2176	119334.00	90.23	820.31
0.2374	79171.00	59.87	544.23
0.2572	55284.00	41.80	380.03
0.2770	43172.00	32.64	296.77
0.2967	33696.00	25.48	231.63
0.3165	25725.00	19.45	176.84
0.3363	18893.00	14.29	129.87
0.3561	13086.00	9.89	89.95
0.3759	8328.00	6.30	57.25
0.3957	4599.00	3.48	31.61
0.4154	2624.00	1.98	18.04
0.4352	1624.00	1.23	11.16
0.4550	847.00	0.64	5.82
0.4748	295.00	0.22	2.03

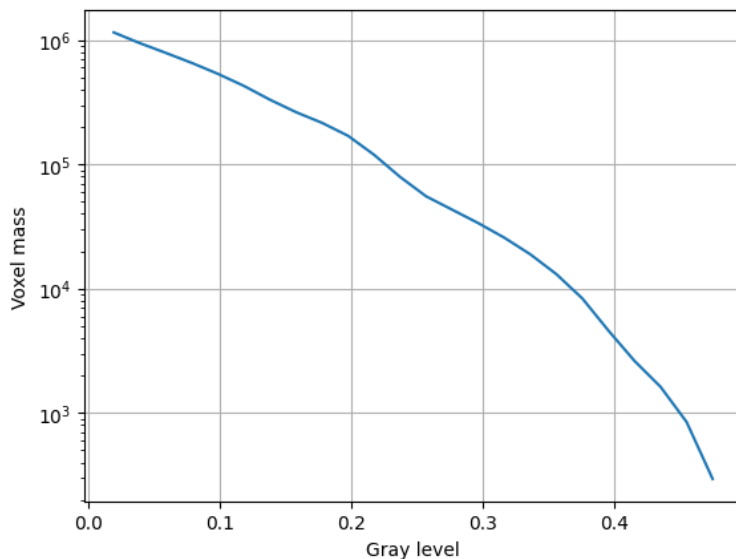


Figure 8: Voxel mass as a function of the gray level.

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 1102319 and a volume of $1006056.79\text{ \AA}^3$. The overlap between the raw and constructed mask is 1.00.

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%.

STATUS: OK

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

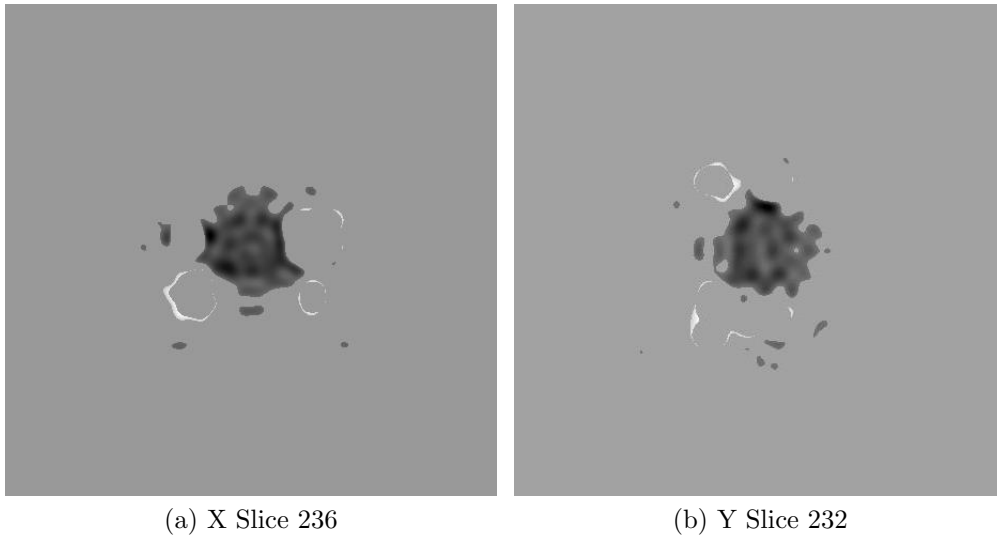
small and they should be randomly distributed without any specific structure. Sometimes, you can see some structure due to the symmetry of the structure.

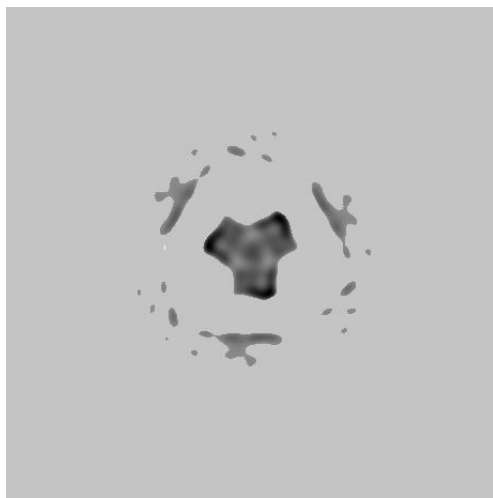
Results:

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 -2936.91 and 0.000000, respectively.

The mean and standard deviation (σ) of the background were -0.001363 and 0.004554. The percentage of background voxels whose absolute value is larger than 5 times the standard deviation is 0.59 % (see Fig. 9). The same percentage from a Gaussian would be 0.000057% (ratio between the two percentages: 10235.182697).

Slices of the background beyond 5σ can be seen in Fig. 9.





(c) Z Slice 241

Figure 9: Maximum variance slices in the three dimensions of the parts of the background beyond 5σ

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=10235.18)**

2.4 Level 0.d B-factor analysis

Explanation:

The B-factor line (see this [link](#) for more details) fitted between $15\text{\AA}$ and the resolution reported should have a slope that is between 0 and $300\text{ \AA}^2$.

Results:

This method cannot be applied to maps with a resolution worse than 8Å.

STATUS: Does not apply

3 Half maps

Half map 1: emd_68835_half_map_1.map

SHA256 hash: ea3d91d44569a94e962f17036f88eb94fdbab01a2f11cd8d4d3d5fb99f742c65

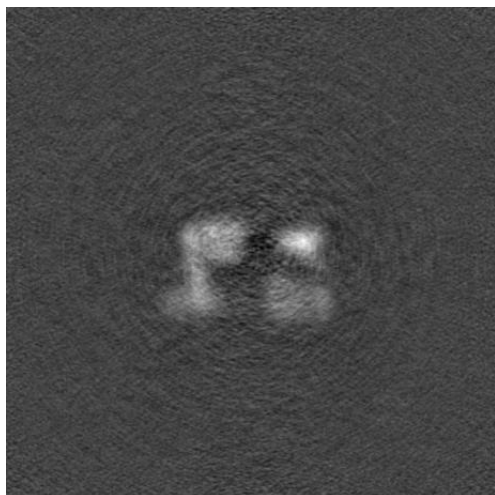
Half map 2: emd_68835_half_map_2.map

SHA256 hash: 0e525e2f446662f92f319056ac2a9a48f9663d40fa836383cc6f6e927a13d261

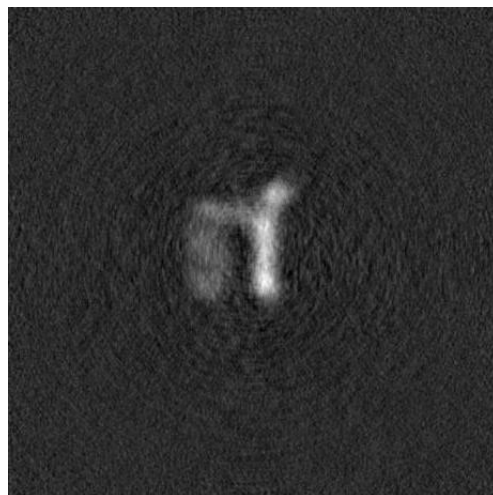
Slices of the first half map can be seen in Fig. 10.

Slices of the second half map can be seen in Fig. 11.

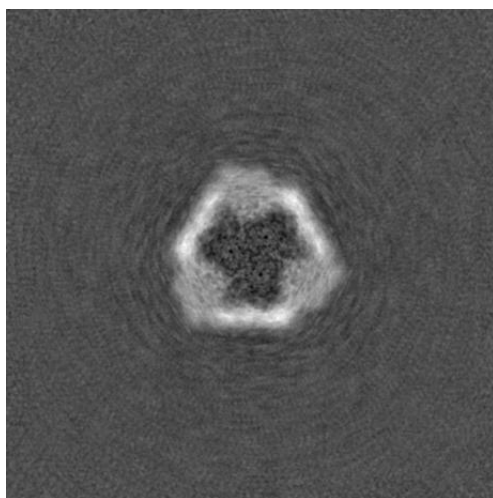
Slices of the difference between both maps can be seen in Fig. 12. There should not be any structure in this difference. Sometimes some patterns are seen if the map is symmetric.



(a) X Slice 190

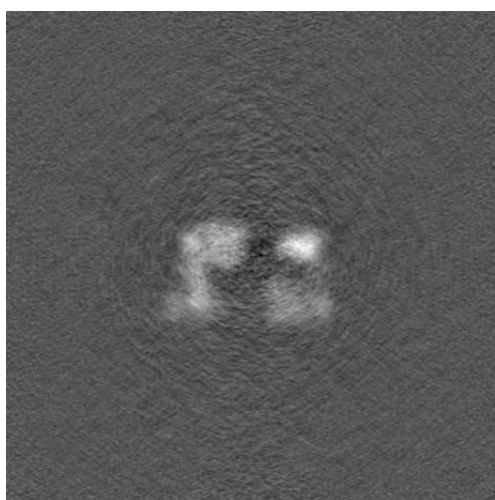


(b) Y Slice 173

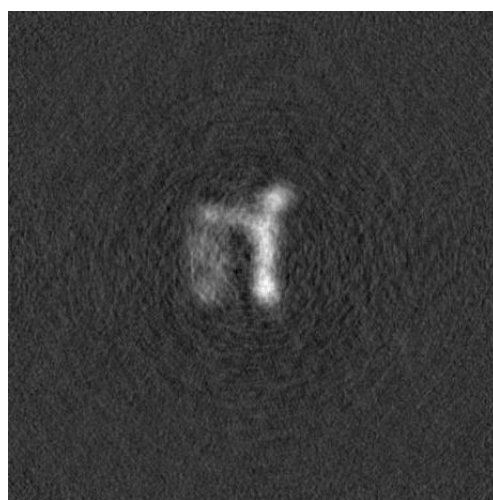


(c) Z Slice 241

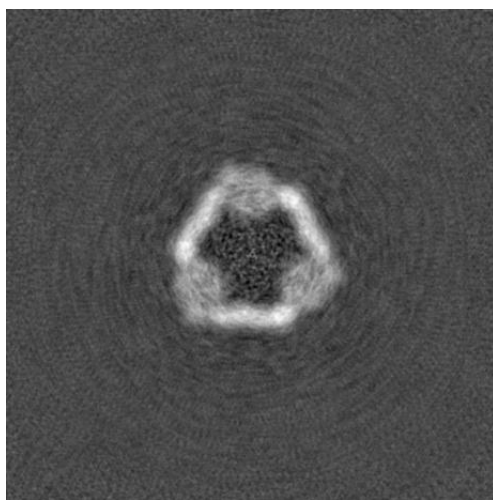
Figure 10: Slices of maximum variation in the three dimensions of Half 1



(a) X Slice 190

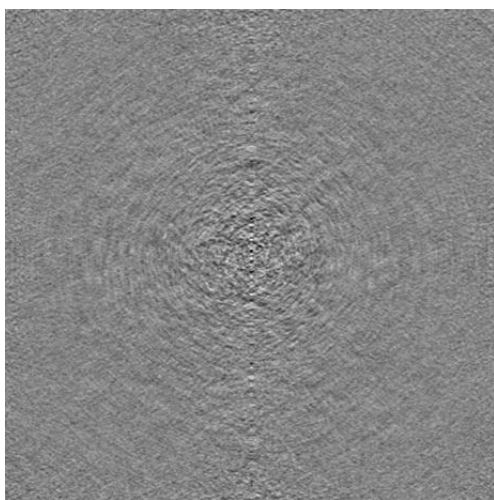


(b) Y Slice 174

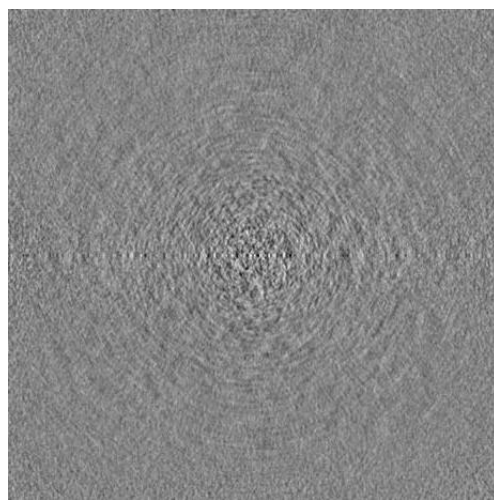


(c) Z Slice 241

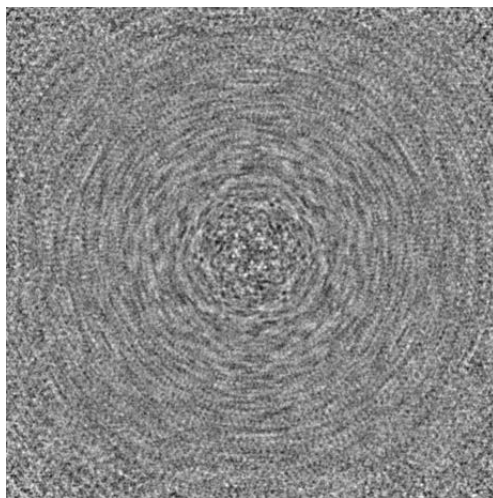
Figure 11: Slices of maximum variation in the three dimensions of Half 2



(a) X Slice 230



(b) Y Slice 230



(c) Z Slice 0

Figure 12: Slices of maximum variation in the three dimensions of the difference Half1-Half2.