



## wwPDB EM Validation Summary Report ⓘ

Feb 10, 2025 – 12:25 PM JST

PDB ID : 8YBY  
EMDB ID : EMD-34563  
Title : State - I: Spike 2-up RBD with THSC20.HVTR26 (Fab26) - single Fab masked  
Authors : Rencilin, C.F.; Bhattacharya, J.; Dutta, S.  
Deposited on : 2024-02-16  
Resolution : 4.40 Å(reported)

This is a wwPDB EM Validation Summary Report for a publicly released PDB entry.

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<https://www.wwpdb.org/validation/2017/EMValidationReportHelp>

with specific help available everywhere you see the ⓘ symbol.

The types of validation reports are described at

<http://www.wwpdb.org/validation/2017/FAQs#types>.

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The following versions of software and data (see [references ⓘ](#)) were used in the production of this report:

EMDB validation analysis : 0.0.1.dev113  
MolProbity : 4.02b-467  
Percentile statistics : 20231227.v01 (using entries in the PDB archive December 27th 2023)  
MapQ : 1.9.13  
Ideal geometry (proteins) : Engh & Huber (2001)  
Ideal geometry (DNA, RNA) : Parkinson et al. (1996)  
Validation Pipeline (wwPDB-VP) : 2.40

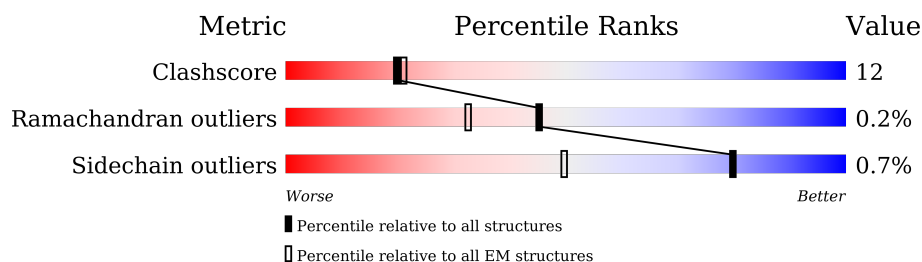
# 1 Overall quality at a glance

The following experimental techniques were used to determine the structure:

*ELECTRON MICROSCOPY*

The reported resolution of this entry is 4.40 Å.

Percentile scores (ranging between 0-100) for global validation metrics of the entry are shown in the following graphic. The table shows the number of entries on which the scores are based.



Metric	Whole archive (#Entries)	EM structures (#Entries)
Clashscore	210492	15764
Ramachandran outliers	207382	16835
Sidechain outliers	206894	16415

The table below summarises the geometric issues observed across the polymeric chains and their fit to the map. The red, orange, yellow and green segments of the bar indicate the fraction of residues that contain outliers for  $\geq 3$ , 2, 1 and 0 types of geometric quality criteria respectively. A grey segment represents the fraction of residues that are not modelled. The numeric value for each fraction is indicated below the corresponding segment, with a dot representing fractions  $\leq 5\%$ . The upper red bar (where present) indicates the fraction of residues that have poor fit to the EM map (all-atom inclusion  $< 40\%$ ). The numeric value is given above the bar.

Mol	Chain	Length	Quality of chain
1	B	1273	
1	D	1273	
1	E	1273	
2	A	231	
3	C	216	

## 2 Entry composition

There are 3 unique types of molecules in this entry. The entry contains 28024 atoms, of which 0 are hydrogens and 0 are deuteriums.

In the tables below, the AltConf column contains the number of residues with at least one atom in alternate conformation and the Trace column contains the number of residues modelled with at most 2 atoms.

- Molecule 1 is a protein called Spike glycoprotein.

Mol	Chain	Residues	Atoms					AltConf	Trace
1	B	1033	Total	C	N	O	S	0	0
			8031	5134	1334	1526	37		
1	D	1067	Total	C	N	O	S	0	0
			8350	5327	1395	1589	39		
1	E	1067	Total	C	N	O	S	0	0
			8350	5327	1395	1589	39		

There are 6 discrepancies between the modelled and reference sequences:

Chain	Residue	Modelled	Actual	Comment	Reference
B	986	PRO	LYS	conflict	UNP P0DTC2
B	987	PRO	VAL	conflict	UNP P0DTC2
D	986	PRO	LYS	conflict	UNP P0DTC2
D	987	PRO	VAL	conflict	UNP P0DTC2
E	986	PRO	LYS	conflict	UNP P0DTC2
E	987	PRO	VAL	conflict	UNP P0DTC2

- Molecule 2 is a protein called THSC20.HVTR26 (Fab26) - Heavy Chain.

Mol	Chain	Residues	Atoms					AltConf	Trace
2	A	230	Total	C	N	O	S	0	0
			1693	1063	287	335	8		

- Molecule 3 is a protein called THSC20.HVTR26 (Fab26) - Light Chain.

Mol	Chain	Residues	Atoms					AltConf	Trace
3	C	215	Total	C	N	O	S	0	0
			1600	999	263	331	7		



LYS	TRP	PRO	TRP	TRP	TYR	THR	ILE	TRP	LEU	GLY	ILE	ALA	GLY	LEU	ILE	ALA	VAL	MET	VAL	THR	THR	ILE	MET	LEU	CYS	CYS	MET	THR	ASN	CYS	CYS	SER	CYS	LYS	GLY	CYS	CYS	LYS	PHE	ASP	GLU	ASP	SER	GLU	PRO	VAL	LEU	LYS	GLY	VAL	LYS	LEU
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● Molecule 1: Spike glycoprotein

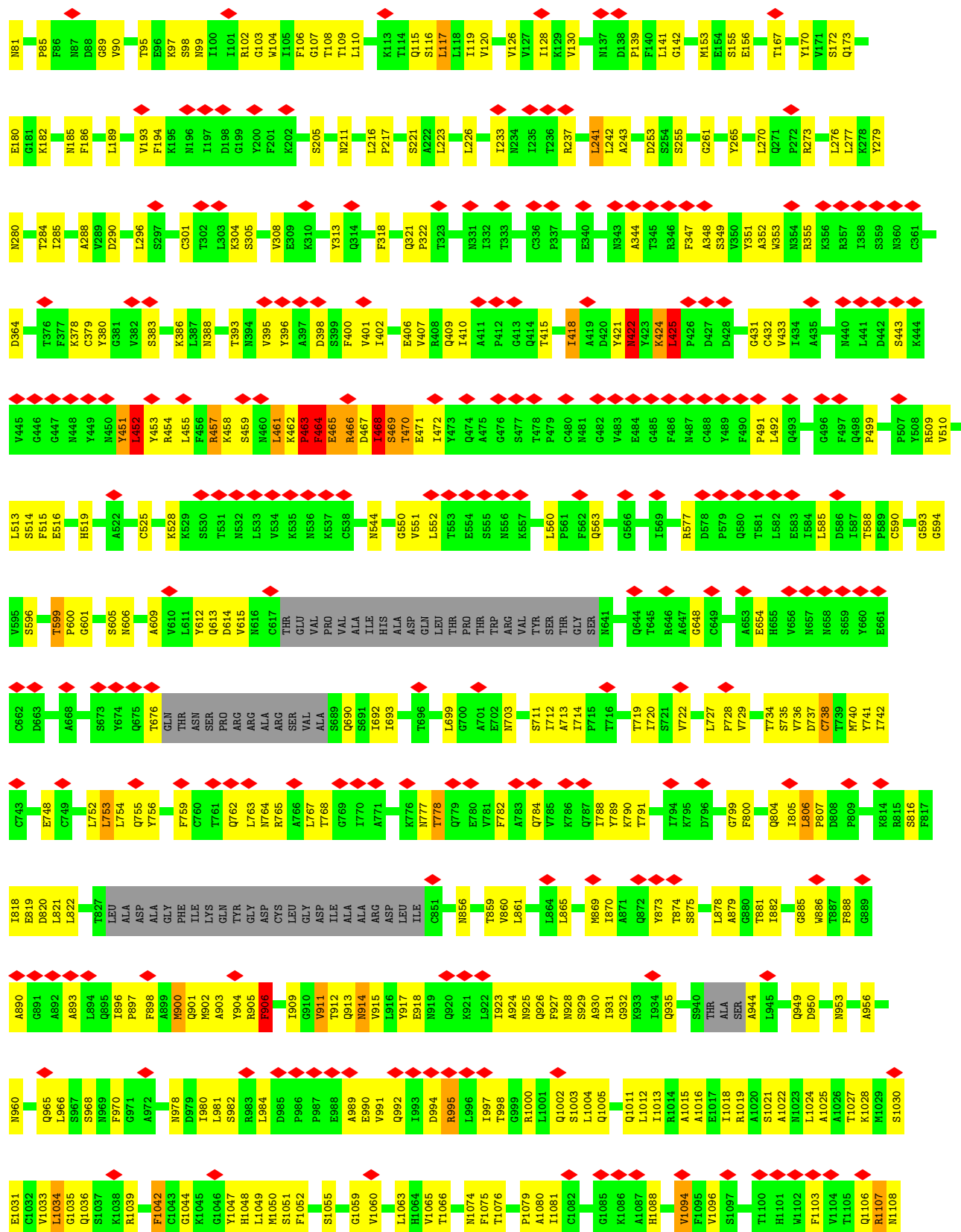
Chain D: 

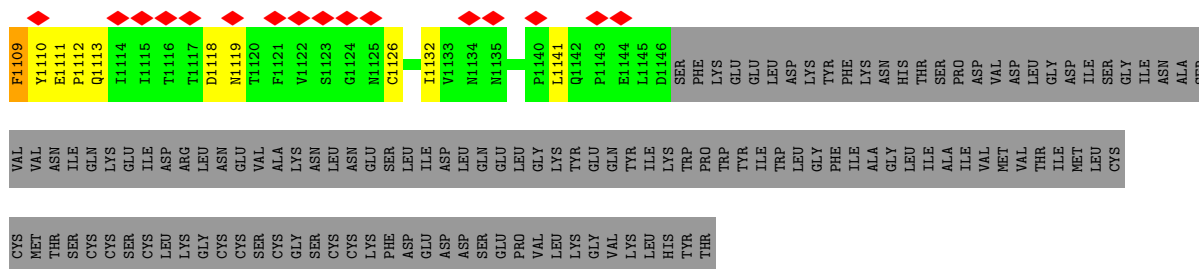
ASP	VAL	ASP	LEU	GLY	ASP	ILE	SER	VAL	VAL	ASN	ILE	GLN	GLY	ILE	ASP	ARG	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	GLU	ILE	ASP	ARG	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS	ASN	LEU	ASN	GLY	VAL	ALA	LYS
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● Molecule 1: Spike glycoprotein

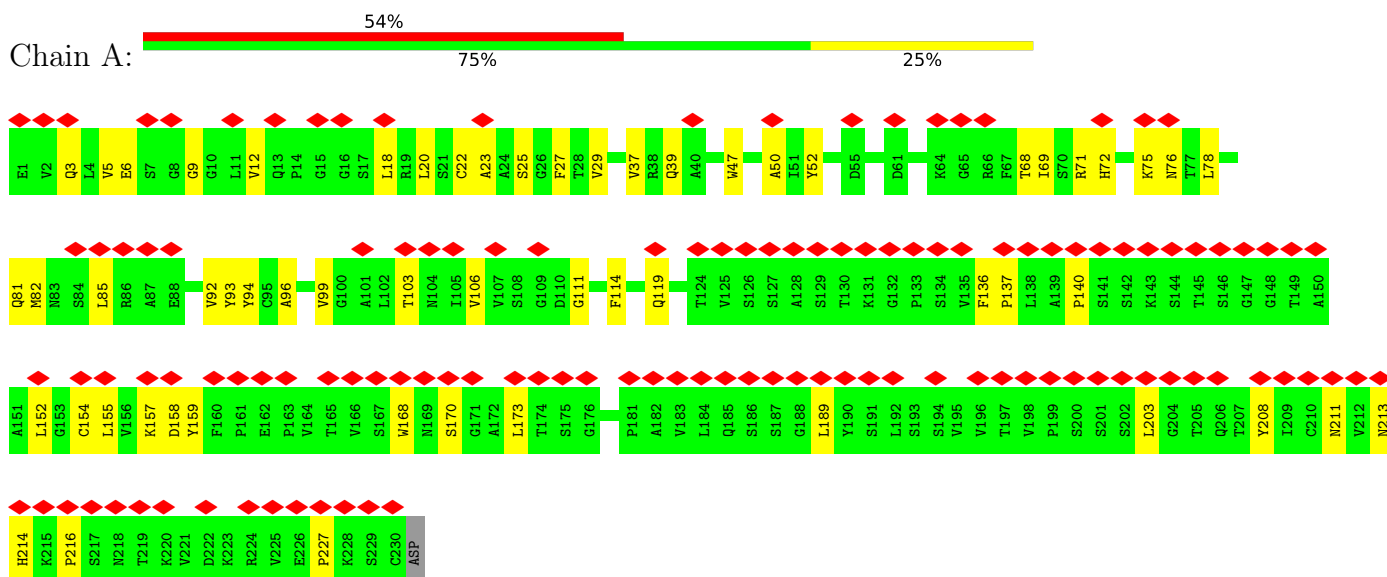
Chain E: 

MET	PHE	VAL	PHE	LEU	VAL	VAL	LEU	LEU	PRO	VAL	SER	Q14	M17	T22	T29	N30	R34	G35	V36	Y37	D40	R44	S45	S46	V47	L48	H49	Q52	D53	L54	F55	L56	P57	F58	V62	H66	A67	I68	H69	V70	SER	GLY	THR	THR	ASN	ASN	GLY	T76	K77	D80
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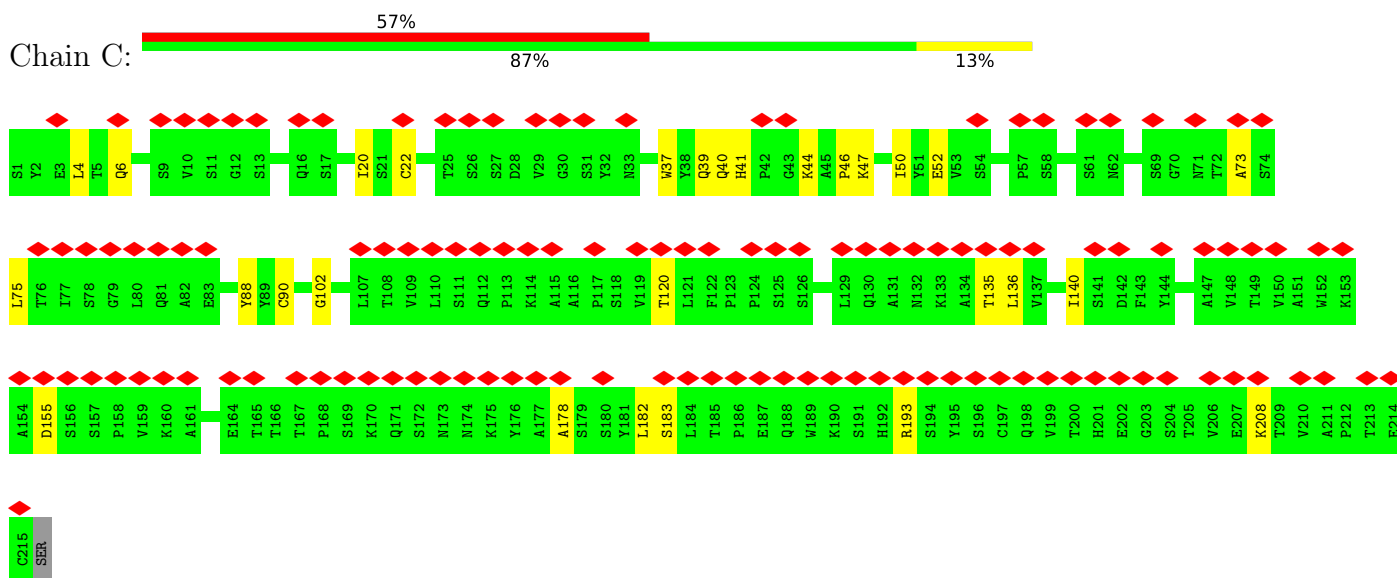




• Molecule 2: THSC20.HVTR26 (Fab26) - Heavy Chain



• Molecule 3: THSC20.HVTR26 (Fab26) - Light Chain



## 4 Experimental information

Property	Value	Source
EM reconstruction method	SINGLE PARTICLE	Depositor
Imposed symmetry	POINT, Not provided	
Number of particles used	109596	Depositor
Resolution determination method	FSC 0.143 CUT-OFF	Depositor
CTF correction method	PHASE FLIPPING AND AMPLITUDE CORRECTION	Depositor
Microscope	FEI TALOS ARCTICA	Depositor
Voltage (kV)	200	Depositor
Electron dose ( $e^-/\text{\AA}^2$ )	50	Depositor
Minimum defocus (nm)	1500	Depositor
Maximum defocus (nm)	2750	Depositor
Magnification	Not provided	
Image detector	GATAN K2 SUMMIT (4k x 4k)	Depositor
Maximum map value	27.086	Depositor
Minimum map value	-12.980	Depositor
Average map value	-0.000	Depositor
Map value standard deviation	1.000	Depositor
Recommended contour level	2.82	Depositor
Map size (Å)	331.2, 331.2, 331.2	wwPDB
Map dimensions	360, 360, 360	wwPDB
Map angles (°)	90.0, 90.0, 90.0	wwPDB
Pixel spacing (Å)	0.92, 0.92, 0.92	Depositor



## 5 Model quality [i](#)

### 5.1 Standard geometry [i](#)

The Z score for a bond length (or angle) is the number of standard deviations the observed value is removed from the expected value. A bond length (or angle) with  $|Z| > 5$  is considered an outlier worth inspection. RMSZ is the root-mean-square of all Z scores of the bond lengths (or angles).

Mol	Chain	Bond lengths		Bond angles	
		RMSZ	# $ Z  > 5$	RMSZ	# $ Z  > 5$
1	B	0.47	2/8214 (0.0%)	0.70	13/11178 (0.1%)
1	D	0.51	1/8544 (0.0%)	0.65	5/11626 (0.0%)
1	E	0.46	0/8544	0.84	23/11626 (0.2%)
2	A	0.28	0/1731	0.54	0/2359
3	C	0.27	0/1639	0.51	0/2236
All	All	0.46	3/28672 (0.0%)	0.71	41/39025 (0.1%)

Chiral center outliers are detected by calculating the chiral volume of a chiral center and verifying if the center is modelled as a planar moiety or with the opposite hand. A planarity outlier is detected by checking planarity of atoms in a peptide group, atoms in a mainchain group or atoms of a sidechain that are expected to be planar.

Mol	Chain	#Chirality outliers	#Planarity outliers
1	B	0	10
1	D	0	6
1	E	0	23
3	C	0	1
All	All	0	40

All (3) bond length outliers are listed below:

Mol	Chain	Res	Type	Atoms	Z	Observed(Å)	Ideal(Å)
1	D	1000	ARG	CA-C	20.94	2.07	1.52
1	B	234	ASN	CG-ND2	10.10	1.58	1.32
1	B	234	ASN	CB-CG	5.36	1.63	1.51

The worst 5 of 41 bond angle outliers are listed below:

Mol	Chain	Res	Type	Atoms	Z	Observed(°)	Ideal(°)
1	E	425	LEU	CA-CB-CG	16.27	152.72	115.30
1	D	1000	ARG	O-C-N	-15.28	98.26	122.70
1	D	1000	ARG	CB-CA-C	12.79	135.97	110.40
1	D	1000	ARG	N-CA-CB	-11.59	89.73	110.60

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Mol	Chain	Res	Type	Atoms	Z	Observed(°)	Ideal(°)
1	E	1034	LEU	CA-CB-CG	10.39	139.19	115.30

There are no chirality outliers.

5 of 40 planarity outliers are listed below:

Mol	Chain	Res	Type	Group
1	B	114	THR	Peptide
1	B	115	GLN	Peptide
1	B	117	LEU	Peptide
1	B	195	LYS	Mainchain
1	B	528	LYS	Peptide

## 5.2 Too-close contacts

In the following table, the Non-H and H(model) columns list the number of non-hydrogen atoms and hydrogen atoms in the chain respectively. The H(added) column lists the number of hydrogen atoms added and optimized by MolProbity. The Clashes column lists the number of clashes within the asymmetric unit, whereas Symm-Clashes lists symmetry-related clashes.

Mol	Chain	Non-H	H(model)	H(added)	Clashes	Symm-Clashes
1	B	8031	0	7820	211	0
1	D	8350	0	8140	195	0
1	E	8350	0	8141	351	0
2	A	1693	0	1664	34	0
3	C	1600	0	1544	19	0
All	All	28024	0	27309	677	0

The all-atom clashscore is defined as the number of clashes found per 1000 atoms (including hydrogen atoms). The all-atom clashscore for this structure is 12.

The worst 5 of 677 close contacts within the same asymmetric unit are listed below, sorted by their clash magnitude.

Atom-1	Atom-2	Interatomic distance (Å)	Clash overlap (Å)
1:D:1002:GLN:HG2	1:E:756:TYR:CE1	1.40	1.51
1:D:1000:ARG:CA	1:D:1000:ARG:C	2.07	1.22
1:D:1002:GLN:CG	1:E:756:TYR:CE1	2.31	1.12
1:E:753:LEU:HD12	1:E:756:TYR:CZ	1.85	1.11
1:D:1002:GLN:HG2	1:E:756:TYR:CD1	1.88	1.08

There are no symmetry-related clashes.

## 5.3 Torsion angles

### 5.3.1 Protein backbone

In the following table, the Percentiles column shows the percent Ramachandran outliers of the chain as a percentile score with respect to all PDB entries followed by that with respect to all EM entries.

The Analysed column shows the number of residues for which the backbone conformation was analysed, and the total number of residues.

Mol	Chain	Analysed	Favoured	Allowed	Outliers	Percentiles	
1	B	1017/1273 (80%)	882 (87%)	134 (13%)	1 (0%)	48	83
1	D	1055/1273 (83%)	956 (91%)	98 (9%)	1 (0%)	48	83
1	E	1055/1273 (83%)	873 (83%)	178 (17%)	4 (0%)	30	67
2	A	228/231 (99%)	209 (92%)	19 (8%)	0	100	100
3	C	213/216 (99%)	199 (93%)	14 (7%)	0	100	100
All	All	3568/4266 (84%)	3119 (87%)	443 (12%)	6 (0%)	45	78

5 of 6 Ramachandran outliers are listed below:

Mol	Chain	Res	Type
1	E	464	PHE
1	B	196	ASN
1	D	1079	PRO
1	E	452	LEU
1	E	463	PRO

### 5.3.2 Protein sidechains

In the following table, the Percentiles column shows the percent sidechain outliers of the chain as a percentile score with respect to all PDB entries followed by that with respect to all EM entries.

The Analysed column shows the number of residues for which the sidechain conformation was analysed, and the total number of residues.

Mol	Chain	Analysed	Rotameric	Outliers	Percentiles	
1	B	891/1112 (80%)	884 (99%)	7 (1%)	79	85
1	D	933/1112 (84%)	932 (100%)	1 (0%)	92	95
1	E	933/1112 (84%)	919 (98%)	14 (2%)	60	75
2	A	190/191 (100%)	189 (100%)	1 (0%)	86	90

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Mol	Chain	Analysed	Rotameric	Outliers	Percentiles	
3	C	183/184 (100%)	183 (100%)	0	100	100
All	All	3130/3711 (84%)	3107 (99%)	23 (1%)	80	87

5 of 23 residues with a non-rotameric sidechain are listed below:

Mol	Chain	Res	Type
1	E	778	THR
1	E	914	ASN
1	E	911	VAL
1	E	995	ARG
1	B	995	ARG

Sometimes sidechains can be flipped to improve hydrogen bonding and reduce clashes. 5 of 46 such sidechains are listed below:

Mol	Chain	Res	Type
1	D	953	ASN
1	E	314	GLN
1	D	1002	GLN
1	E	99	ASN
1	E	690	GLN

### 5.3.3 RNA [i](#)

There are no RNA molecules in this entry.

## 5.4 Non-standard residues in protein, DNA, RNA chains [i](#)

There are no non-standard protein/DNA/RNA residues in this entry.

## 5.5 Carbohydrates [i](#)

There are no oligosaccharides in this entry.

## 5.6 Ligand geometry [i](#)

There are no ligands in this entry.

## 5.7 Other polymers [i](#)

There are no such residues in this entry.

## 5.8 Polymer linkage issues [i](#)

There are no chain breaks in this entry.

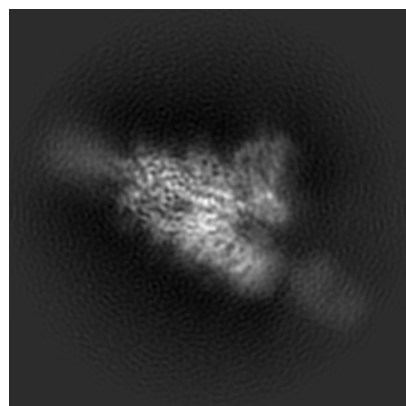
## 6 Map visualisation [i](#)

This section contains visualisations of the EMDB entry EMD-34563. These allow visual inspection of the internal detail of the map and identification of artifacts.

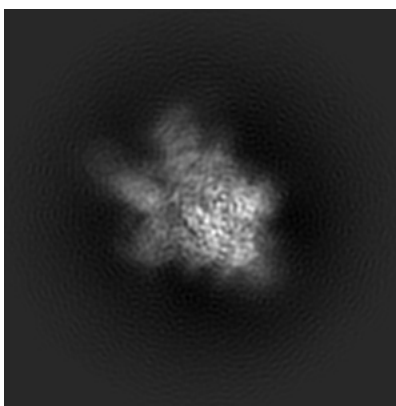
Images derived from a raw map, generated by summing the deposited half-maps, are presented below the corresponding image components of the primary map to allow further visual inspection and comparison with those of the primary map.

### 6.1 Orthogonal projections [i](#)

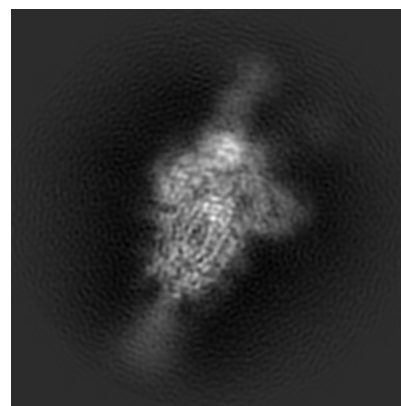
#### 6.1.1 Primary map



X

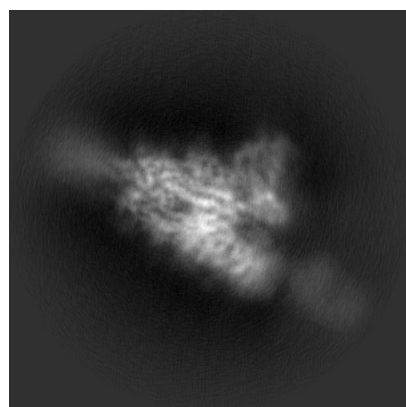


Y

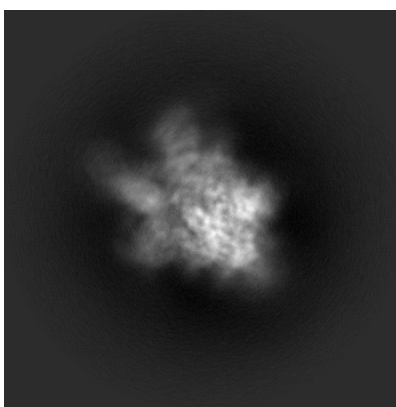


Z

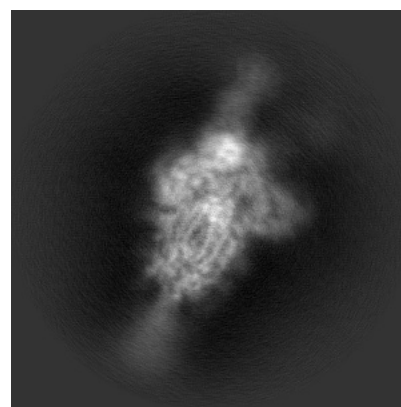
#### 6.1.2 Raw map



X



Y

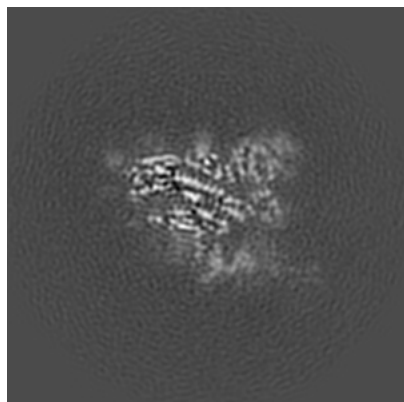


Z

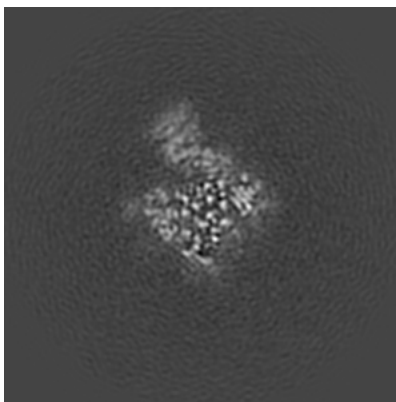
The images above show the map projected in three orthogonal directions.

## 6.2 Central slices [i](#)

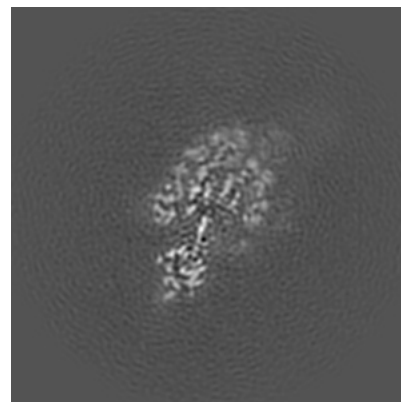
### 6.2.1 Primary map



X Index: 180

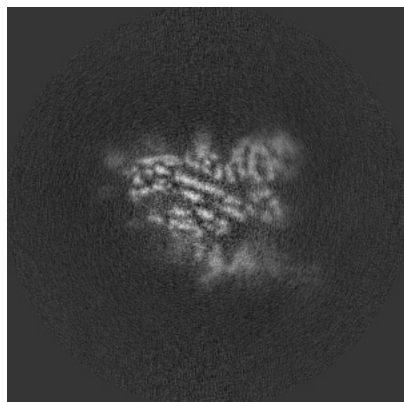


Y Index: 180

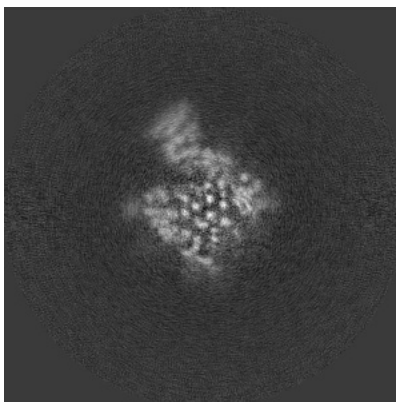


Z Index: 180

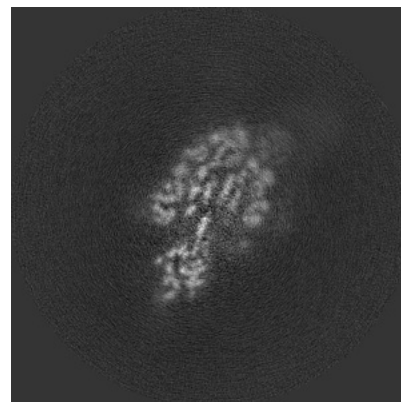
### 6.2.2 Raw map



X Index: 180



Y Index: 180

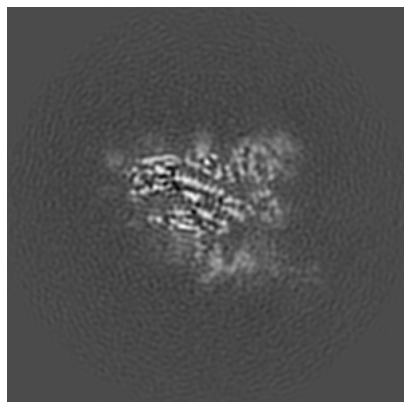


Z Index: 180

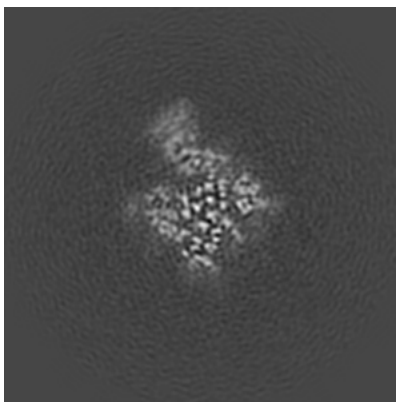
The images above show central slices of the map in three orthogonal directions.

## 6.3 Largest variance slices [i](#)

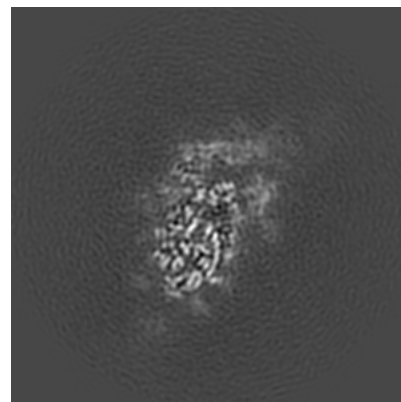
### 6.3.1 Primary map



X Index: 180

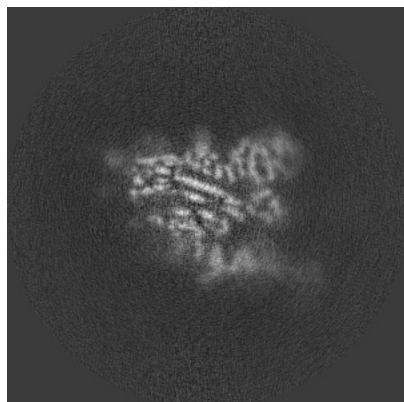


Y Index: 178

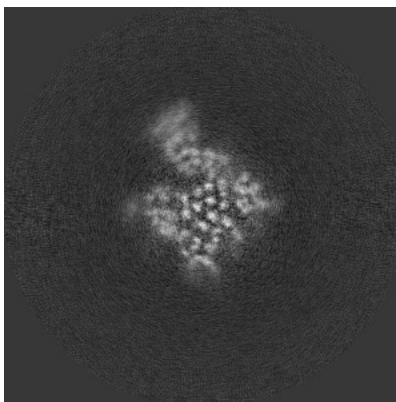


Z Index: 193

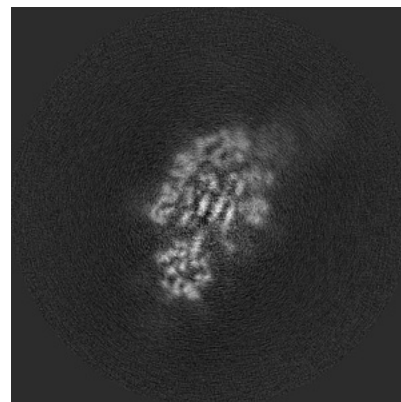
### 6.3.2 Raw map



X Index: 181



Y Index: 178



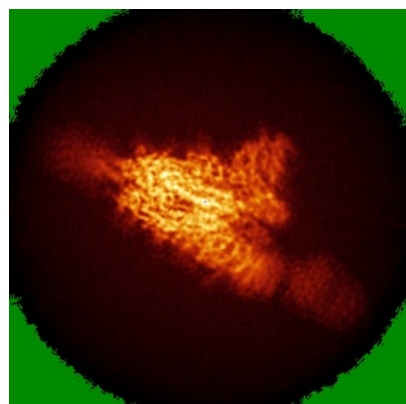
Z Index: 185

The images above show the largest variance slices of the map in three orthogonal directions.

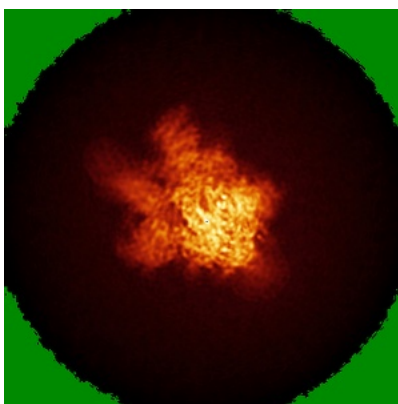


## 6.4 Orthogonal standard-deviation projections (False-color) [i](#)

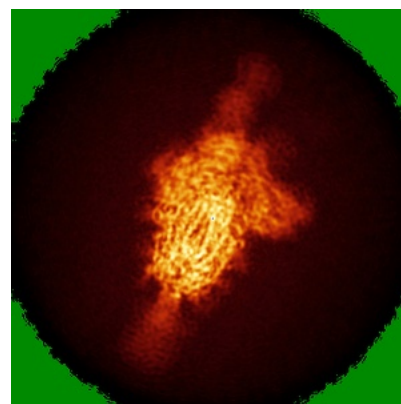
### 6.4.1 Primary map



X

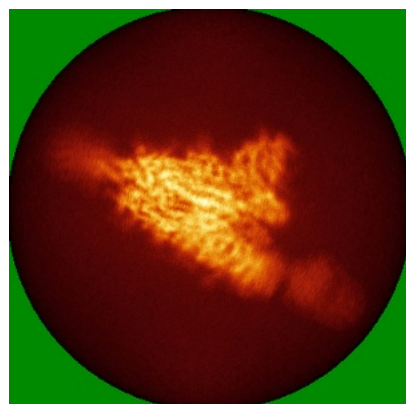


Y

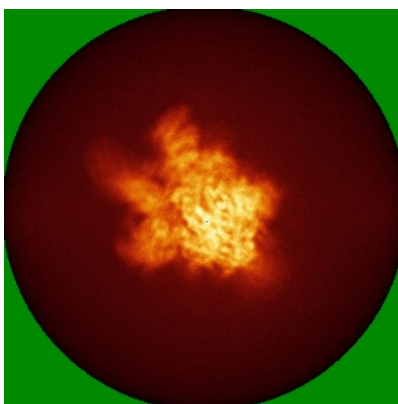


Z

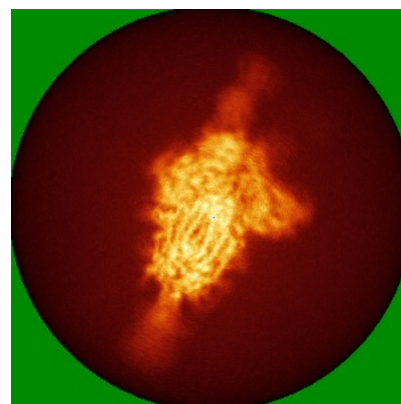
### 6.4.2 Raw map



X



Y

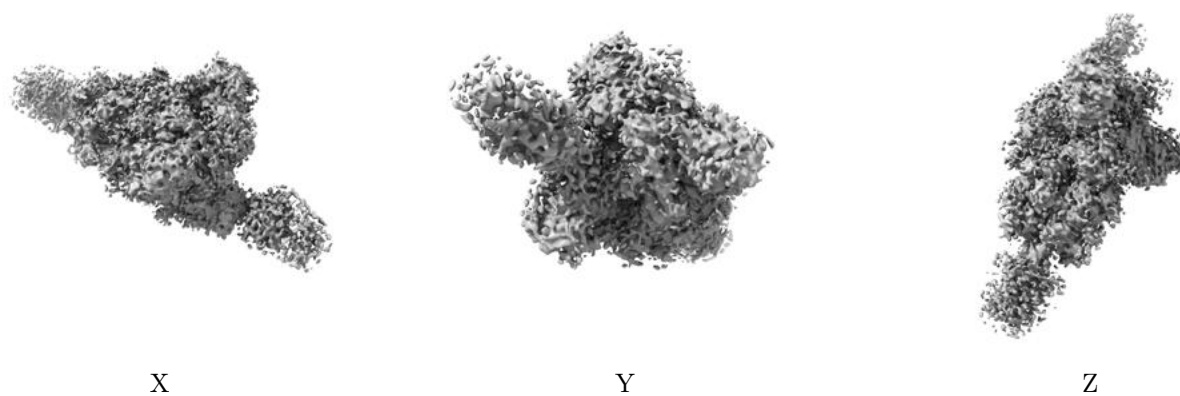


Z

The images above show the map standard deviation projections with false color in three orthogonal directions. Minimum values are shown in green, max in blue, and dark to light orange shades represent small to large values respectively.

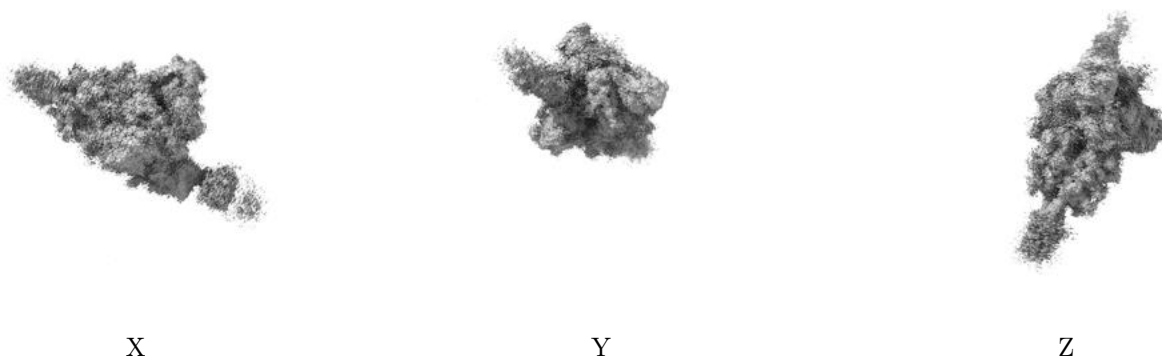
## 6.5 Orthogonal surface views [i](#)

### 6.5.1 Primary map



The images above show the 3D surface view of the map at the recommended contour level 2.82. These images, in conjunction with the slice images, may facilitate assessment of whether an appropriate contour level has been provided.

### 6.5.2 Raw map



These images show the 3D surface of the raw map. The raw map's contour level was selected so that its surface encloses the same volume as the primary map does at its recommended contour level.

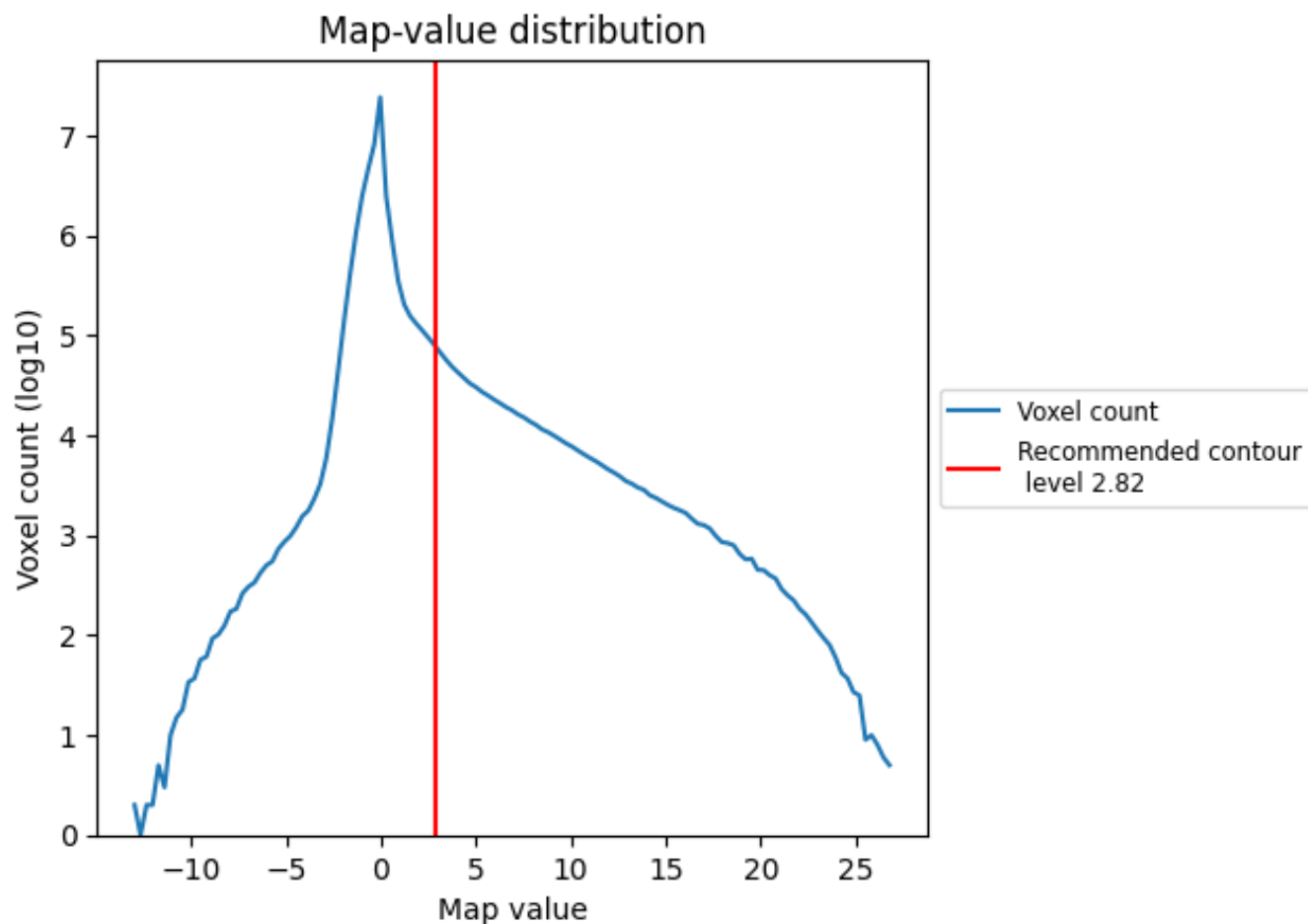
## 6.6 Mask visualisation [i](#)

This section was not generated. No masks/segmentation were deposited.

## 7 Map analysis [i](#)

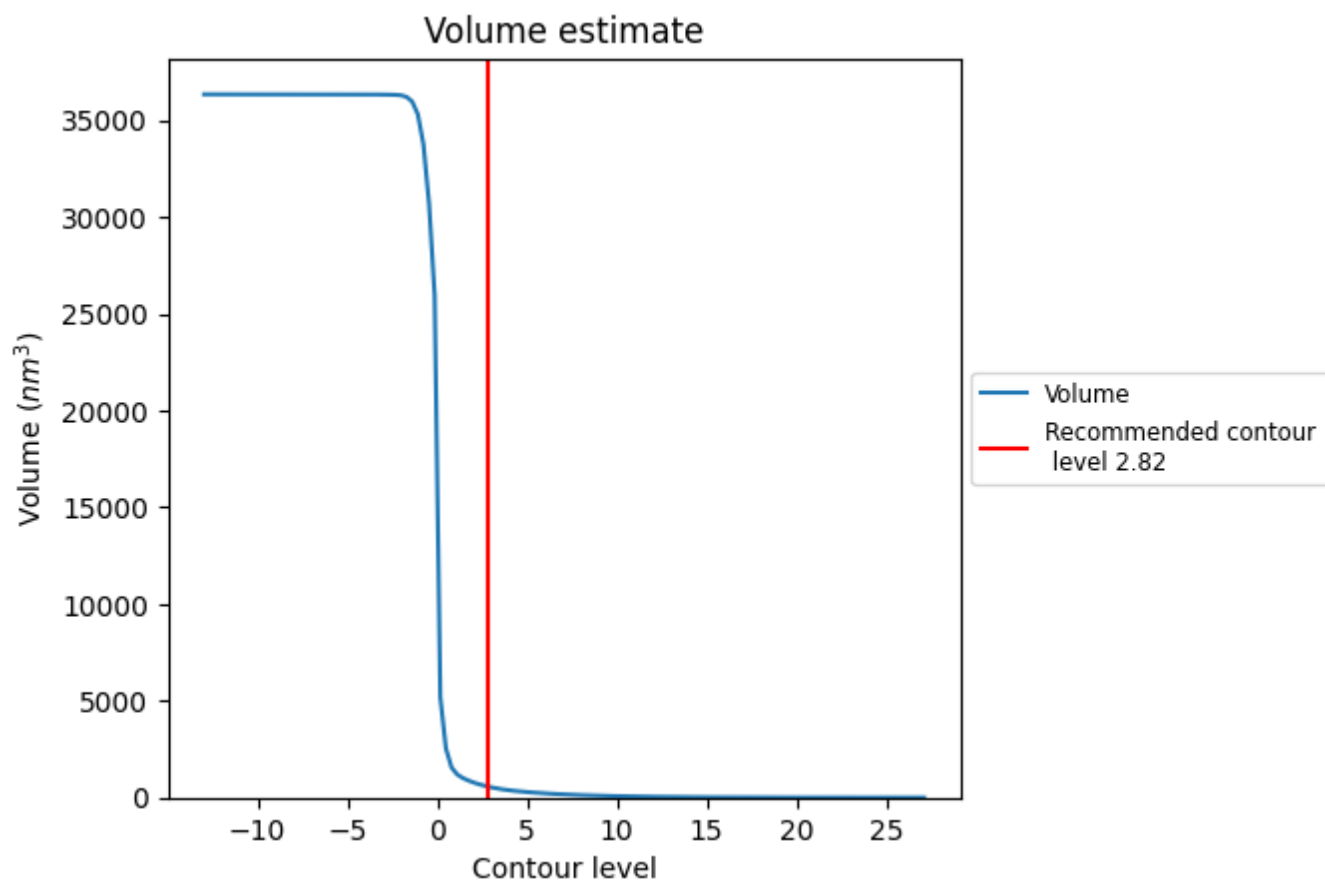
This section contains the results of statistical analysis of the map.

### 7.1 Map-value distribution [i](#)



The map-value distribution is plotted in 128 intervals along the x-axis. The y-axis is logarithmic. A spike in this graph at zero usually indicates that the volume has been masked.

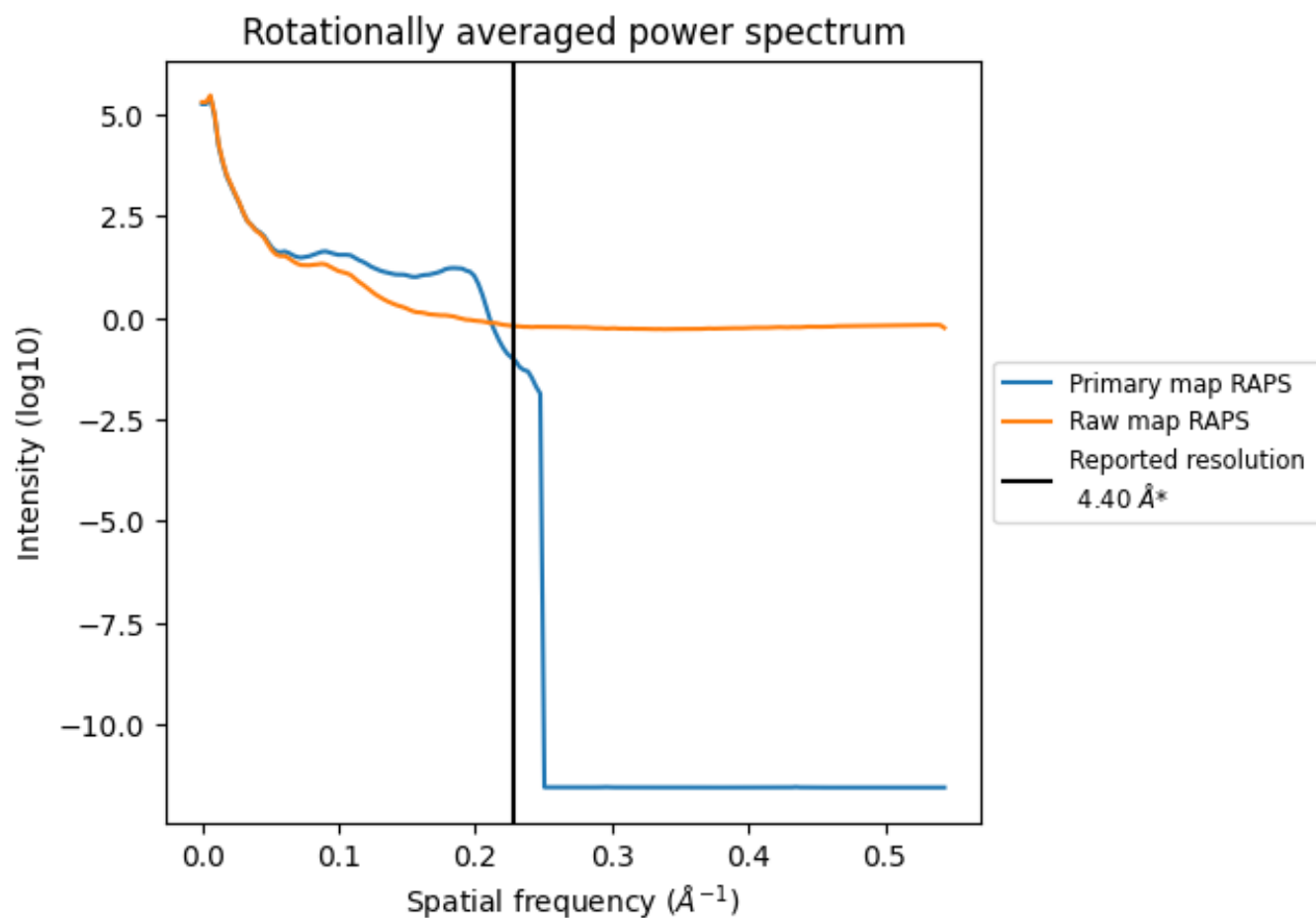
## 7.2 Volume estimate [i](#)



The volume at the recommended contour level is 566 nm<sup>3</sup>; this corresponds to an approximate mass of 511 kDa.

The volume estimate graph shows how the enclosed volume varies with the contour level. The recommended contour level is shown as a vertical line and the intersection between the line and the curve gives the volume of the enclosed surface at the given level.

### 7.3 Rotationally averaged power spectrum ⓘ

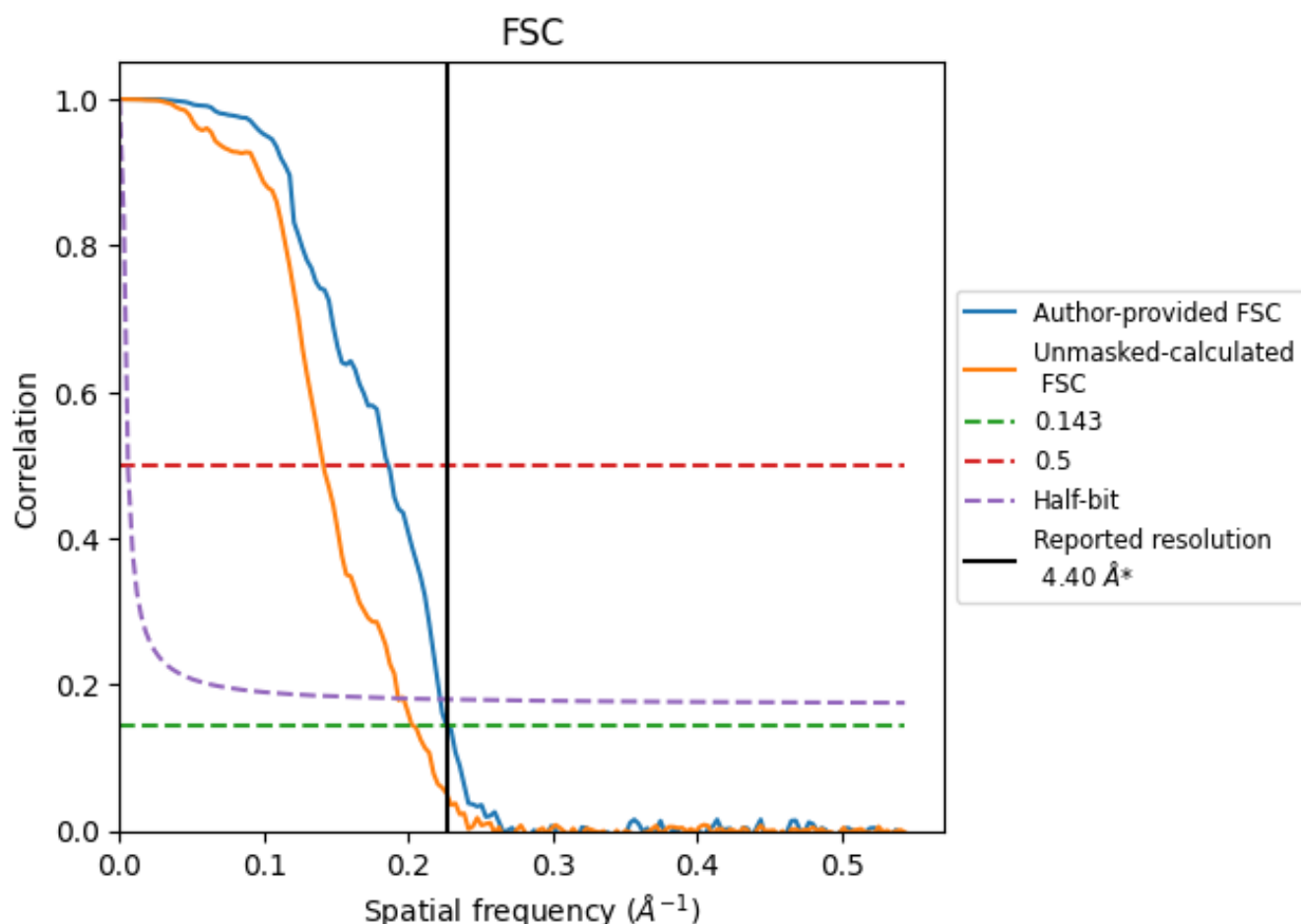


\*Reported resolution corresponds to spatial frequency of 0.227  $\text{\AA}^{-1}$

## 8 Fourier-Shell correlation [i](#)

Fourier-Shell Correlation (FSC) is the most commonly used method to estimate the resolution of single-particle and subtomogram-averaged maps. The shape of the curve depends on the imposed symmetry, mask and whether or not the two 3D reconstructions used were processed from a common reference. The reported resolution is shown as a black line. A curve is displayed for the half-bit criterion in addition to lines showing the 0.143 gold standard cut-off and 0.5 cut-off.

### 8.1 FSC [i](#)



\*Reported resolution corresponds to spatial frequency of 0.227 Å<sup>-1</sup>

## 8.2 Resolution estimates [i](#)

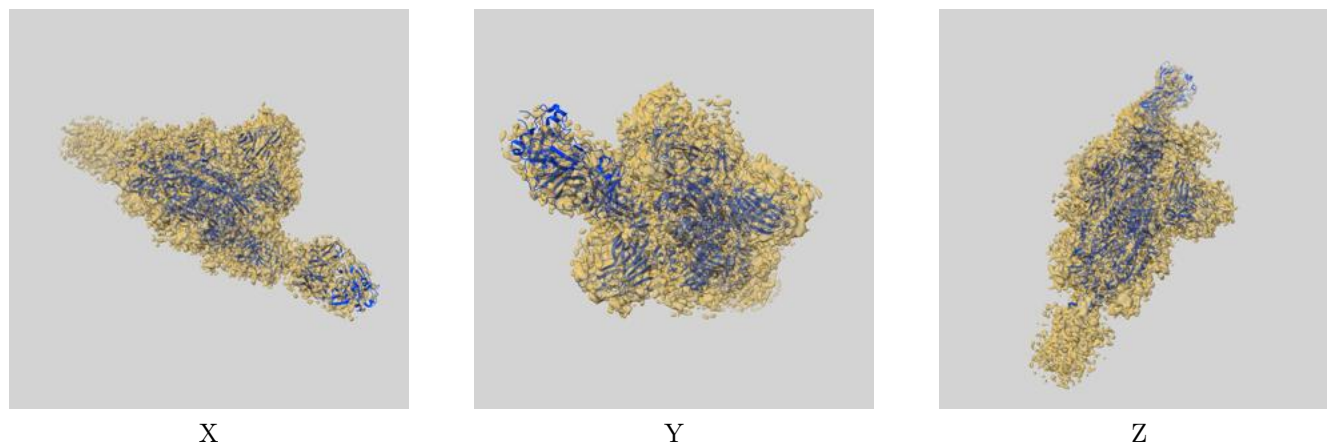
Resolution estimate (Å)	Estimation criterion (FSC cut-off)		
	0.143	0.5	Half-bit
Reported by author	4.40	-	-
Author-provided FSC curve	4.41	5.38	4.50
Unmasked-calculated*	4.91	7.09	5.18

\*Resolution estimate based on FSC curve calculated by comparison of deposited half-maps. The value from deposited half-maps intersecting FSC 0.143 CUT-OFF 4.91 differs from the reported value 4.4 by more than 10 %

## 9 Map-model fit [i](#)

This section contains information regarding the fit between EMDB map EMD-34563 and PDB model 8YBY. Per-residue inclusion information can be found in [section 3](#) on [page 4](#).

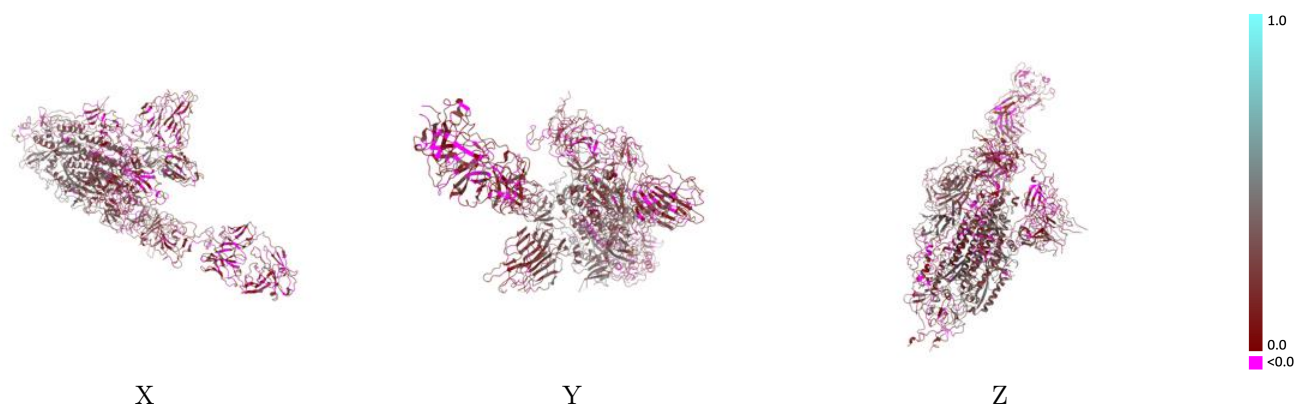
### 9.1 Map-model overlay [i](#)



The images above show the 3D surface view of the map at the recommended contour level 2.82 at 50% transparency in yellow overlaid with a ribbon representation of the model coloured in blue. These images allow for the visual assessment of the quality of fit between the atomic model and the map.

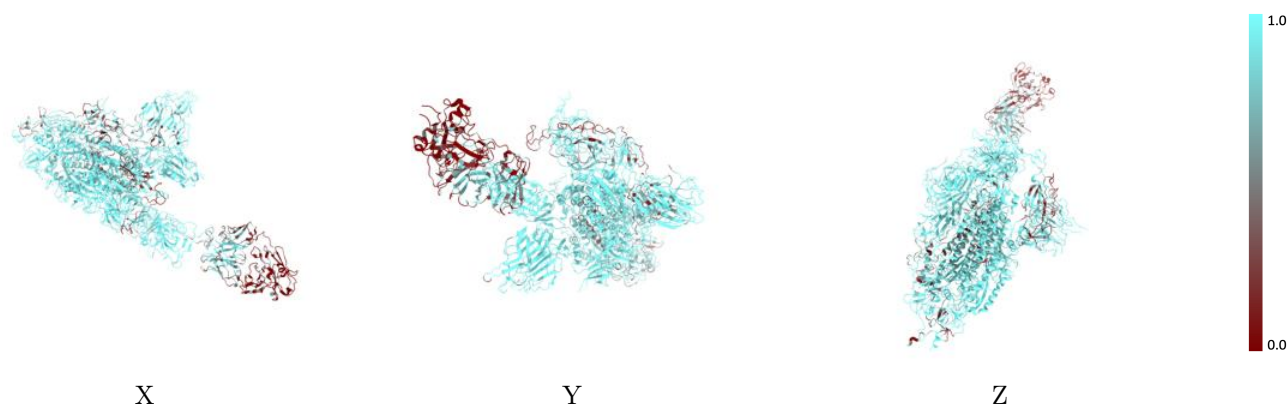


## 9.2 Q-score mapped to coordinate model [i](#)



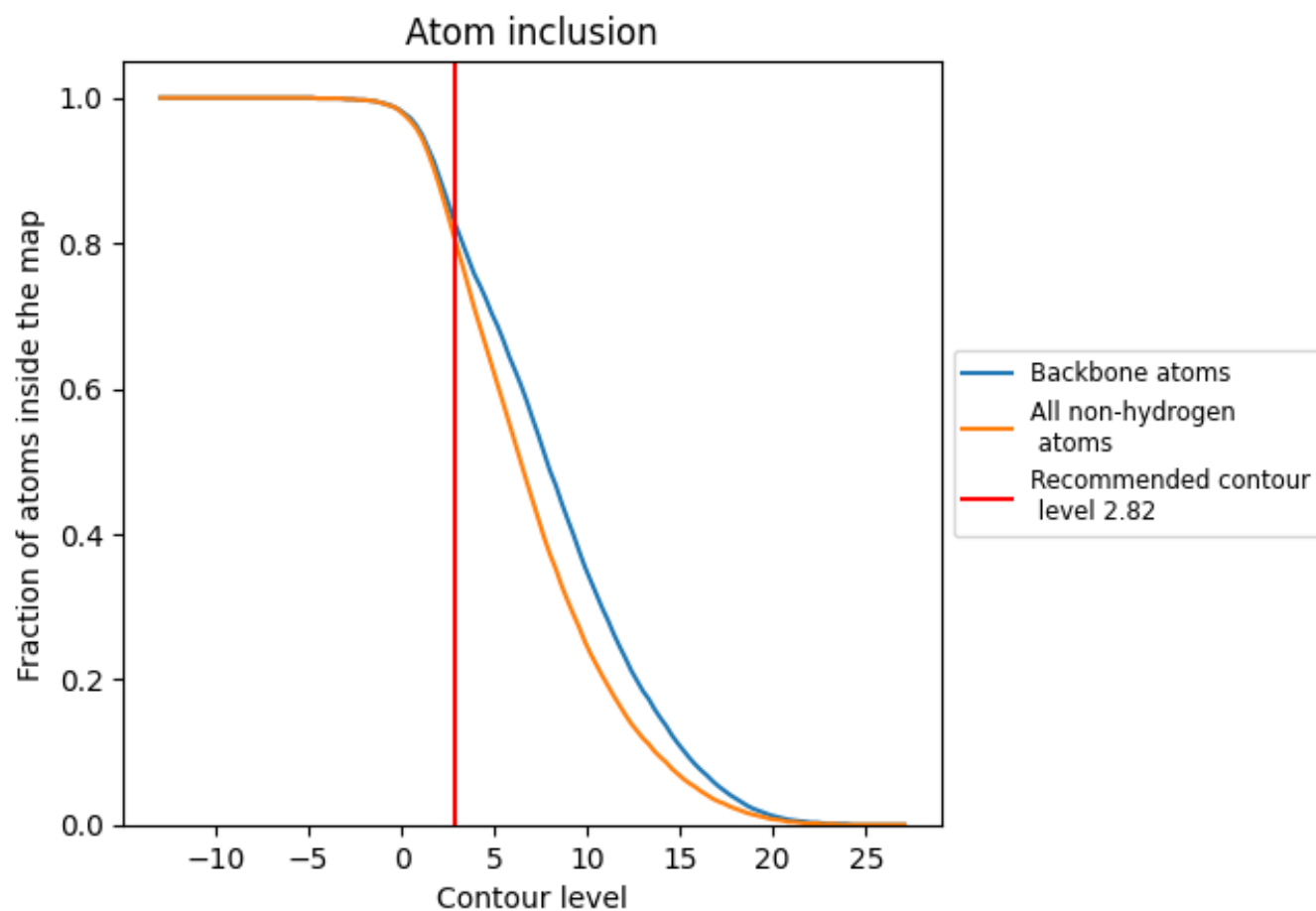
The images above show the model with each residue coloured according to its Q-score. This shows their resolvability in the map with higher Q-score values reflecting better resolvability. Please note: Q-score is calculating the resolvability of atoms, and thus high values are only expected at resolutions at which atoms can be resolved. Low Q-score values may therefore be expected for many entries.

## 9.3 Atom inclusion mapped to coordinate model [i](#)



The images above show the model with each residue coloured according to its atom inclusion. This shows to what extent they are inside the map at the recommended contour level (2.82).

## 9.4 Atom inclusion [i](#)



At the recommended contour level, 83% of all backbone atoms, 81% of all non-hydrogen atoms, are inside the map.

9.5 Map-model fit summary ⓘ

The table lists the average atom inclusion at the recommended contour level (2.82) and Q-score for the entire model and for each chain.

Chain	Atom inclusion	Q-score
All	<div></div> 0.8090	<div></div> 0.2290
A	<div></div> 0.4090	<div></div> 0.1310
B	<div></div> 0.9600	<div></div> 0.3040
C	<div></div> 0.3620	<div></div> 0.1290
D	<div></div> 0.9460	<div></div> 0.3030
E	<div></div> 0.6930	<div></div> 0.1210

