

Full wwPDB X-ray Structure Validation Report (i)

Aug 15, 2023 – 10:23 PM EDT

PDB ID : 1SJ5

Title : Crystal structure of a duf151 family protein (tm0160) from thermotoga mar-

itima at 2.8 A resolution

Authors: Spraggon, G.; Panatazatos, D.; Klock, H.E.; Wilson, I.A.; Woods Jr., V.L.;

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Deposited on : 2004-03-02

Resolution : 2.80 Å(reported)

This is a Full wwPDB X-ray Structure Validation Report for a publicly released PDB entry.

We welcome your comments at validation@mail.wwpdb.org
A user guide is available at
https://www.wwpdb.org/validation/2017/XrayValidationReportHelp
with specific help available everywhere you see the (i) symbol.

The types of validation reports are described at http://www.wwpdb.org/validation/2017/FAQs#types.

The following versions of software and data (see references (1)) were used in the production of this report:

 $\begin{array}{ccc} & Mol Probity & : & 4.02b\text{-}467 \\ & Xtriage \text{ (Phenix)} & : & 1.13 \end{array}$

EDS: 2.35

Percentile statistics : 20191225.v01 (using entries in the PDB archive December 25th 2019)

 $Refmac \quad : \quad 5.8.0158$

CCP4 : 7.0.044 (Gargrove)

Ideal geometry (proteins) : Engh & Huber (2001) Ideal geometry (DNA, RNA) : Parkinson et al. (1996)

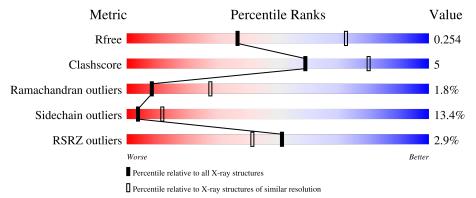
Validation Pipeline (wwPDB-VP) : 2.35

1 Overall quality at a glance (i)

The following experimental techniques were used to determine the structure: $X\text{-}RAY\ DIFFRACTION$

The reported resolution of this entry is 2.80 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	Similar resolution
Metric	$(\# \mathrm{Entries})$	$(\# ext{Entries}, ext{ resolution range}(\mathring{A}))$
R_{free}	130704	3140 (2.80-2.80)
Clashscore	141614	3569 (2.80-2.80)
Ramachandran outliers	138981	3498 (2.80-2.80)
Sidechain outliers	138945	3500 (2.80-2.80)
RSRZ outliers	127900	3078 (2.80-2.80)

The table below summarises the geometric issues observed across the polymeric chains and their fit to the electron density. The red, orange, yellow and green segments of the lower bar indicate the fraction of residues that contain outliers for >=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 <=5% The upper red bar (where present) indicates the fraction of residues that have poor fit to the electron density. The numeric value is given above the bar.

Mol	Chain	Length	Quality of chain					
1	A	164	66%	17%	• • 13%			
1	В	164	66%	16%	• 16%			



2 Entry composition (i)

There is only 1 type of molecule in this entry. The entry contains 2173 atoms, of which 0 are hydrogens and 0 are deuteriums.

In the tables below, the ZeroOcc column contains the number of atoms modelled with zero occupancy, 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 conserved hypothetical protein TM0160.

	\mathbf{Mol}	Chain	Residues	Atoms			ZeroOcc	AltConf	Trace		
Ī	1	Λ	143	Total	С	N	N O S		0	0	0
	1	Λ	140	1111	712	192	204	3	0	U	U
	1	B	137	Total	С	N	О	S	0	0	0
	1	Ъ	131	1062	683	179	197	3	U	U	0

There are 40 discrepancies between the modelled and reference sequences:

A -11 MET - cloning artifact UNP Q9W A -10 GLY - cloning artifact UNP Q9W A -9 SER - cloning artifact UNP Q9W A -8 ASP - cloning artifact UNP Q9W A -7 LYS - cloning artifact UNP Q9W A -6 ILE - cloning artifact UNP Q9W A -5 HIS - cloning artifact UNP Q9W A -4 HIS - cloning artifact UNP Q9W A -2 HIS - cloning artifact UNP Q9W A -1 HIS - cloning artifact UNP Q9W A 0 HIS - cloning artifact UNP Q9W A 146 ARG - SEE REMARK 999 UNP Q9W A 147 ASP - SEE REMARK 999 UNP Q9W						
A -10 GLY - cloning artifact UNP Q9W A -9 SER - cloning artifact UNP Q9W A -8 ASP - cloning artifact UNP Q9W A -7 LYS - cloning artifact UNP Q9W A -6 ILE - cloning artifact UNP Q9W A -5 HIS - cloning artifact UNP Q9W A -4 HIS - cloning artifact UNP Q9W A -3 HIS - cloning artifact UNP Q9W A -1 HIS - cloning artifact UNP Q9W A 0 HIS - cloning artifact UNP Q9W A 38 ALA CYS engineered mutation UNP Q9W A 146 ARG - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W	Chain	Residue	Modelled	Actual	Comment	Reference
A -9 SER - cloning artifact UNP Q9W A -8 ASP - cloning artifact UNP Q9W A -7 LYS - cloning artifact UNP Q9W A -6 ILE - cloning artifact UNP Q9W A -5 HIS - cloning artifact UNP Q9W A -4 HIS - cloning artifact UNP Q9W A -3 HIS - cloning artifact UNP Q9W A -1 HIS - cloning artifact UNP Q9W A 0 HIS - cloning artifact UNP Q9W A 38 ALA CYS engineered mutation UNP Q9W A 146 ARG - SEE REMARK 999 UNP Q9W A 147 ASP - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W	A	-11	MET	-	cloning artifact	UNP Q9WY07
A -8 ASP - cloning artifact UNP Q9W A -7 LYS - cloning artifact UNP Q9W A -6 ILE - cloning artifact UNP Q9W A -5 HIS - cloning artifact UNP Q9W A -4 HIS - cloning artifact UNP Q9W A -3 HIS - cloning artifact UNP Q9W A -2 HIS - cloning artifact UNP Q9W A -1 HIS - cloning artifact UNP Q9W A 0 HIS - cloning artifact UNP Q9W A 146 ARG - SEE REMARK 999 UNP Q9W A 147 ASP - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W	A	-10	GLY	-		UNP Q9WY07
A -7 LYS - cloning artifact UNP Q9W A -6 ILE - cloning artifact UNP Q9W A -5 HIS - cloning artifact UNP Q9W A -4 HIS - cloning artifact UNP Q9W A -3 HIS - cloning artifact UNP Q9W A -1 HIS - cloning artifact UNP Q9W A 0 HIS - cloning artifact UNP Q9W A 38 ALA CYS engineered mutation UNP Q9W A 146 ARG - SEE REMARK 999 UNP Q9W A 147 ASP - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W	A	-9	SER	-	cloning artifact	UNP Q9WY07
A -6 ILE - cloning artifact UNP Q9W A -5 HIS - cloning artifact UNP Q9W A -4 HIS - cloning artifact UNP Q9W A -3 HIS - cloning artifact UNP Q9W A -1 HIS - cloning artifact UNP Q9W A 0 HIS - cloning artifact UNP Q9W A 38 ALA CYS engineered mutation UNP Q9W A 146 ARG - SEE REMARK 999 UNP Q9W A 147 ASP - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W A 149 ILE - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W <	A	-8	ASP	-	cloning artifact	UNP Q9WY07
A -5 HIS - cloning artifact UNP Q9W A -4 HIS - cloning artifact UNP Q9W A -3 HIS - cloning artifact UNP Q9W A -2 HIS - cloning artifact UNP Q9W A -1 HIS - cloning artifact UNP Q9W A 0 HIS - cloning artifact UNP Q9W A 38 ALA CYS engineered mutation UNP Q9W A 146 ARG - SEE REMARK 999 UNP Q9W A 147 ASP - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W A 152 ARG - SEE REMARK 999 UNP Q9W <	A	-7	LYS	-	cloning artifact	UNP Q9WY07
A -4 HIS - cloning artifact UNP Q9W A -3 HIS - cloning artifact UNP Q9W A -2 HIS - cloning artifact UNP Q9W A -1 HIS - cloning artifact UNP Q9W A 0 HIS - cloning artifact UNP Q9W A 38 ALA CYS engineered mutation UNP Q9W A 146 ARG - SEE REMARK 999 UNP Q9W A 147 ASP - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W A 149 ILE - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W A 152 ARG - SEE REMARK 999 UNP Q9W A 153 SER - SEE REMARK 999 UNP Q9W A 154 SER - SEE REMARK 999 UNP Q9W A 155 SER - SEE REMARK 999 UNP Q9W A 156 CONING ARTIFACT UNP Q9W C CONING ARTIFAC	A	-6	ILE	-	cloning artifact	UNP Q9WY07
A -3 HIS - cloning artifact UNP Q9W A -2 HIS - cloning artifact UNP Q9W A -1 HIS - cloning artifact UNP Q9W A 0 HIS - cloning artifact UNP Q9W A 38 ALA CYS engineered mutation UNP Q9W A 146 ARG - SEE REMARK 999 UNP Q9W A 147 ASP - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W A 149 ILE - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W A 152 ARG - SEE REMARK 999 UNP Q9W B -11 MET - cloning artifact UNP Q9W <	A	-5	HIS	-	cloning artifact	UNP Q9WY07
A -2 HIS - cloning artifact UNP Q9W A -1 HIS - cloning artifact UNP Q9W A 0 HIS - cloning artifact UNP Q9W A 38 ALA CYS engineered mutation UNP Q9W A 146 ARG - SEE REMARK 999 UNP Q9W A 147 ASP - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W A 149 ILE - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W A 152 ARG - SEE REMARK 999 UNP Q9W B -11 MET - cloning artifact UNP Q9W B -10 GLY - cloning artifact UNP Q9W <td>A</td> <td>-4</td> <td>HIS</td> <td>-</td> <td>cloning artifact</td> <td>UNP Q9WY07</td>	A	-4	HIS	-	cloning artifact	UNP Q9WY07
A -1 HIS - cloning artifact UNP Q9W A 0 HIS - cloning artifact UNP Q9W A 38 ALA CYS engineered mutation UNP Q9W A 146 ARG - SEE REMARK 999 UNP Q9W A 147 ASP - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W A 149 ILE - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W A 152 ARG - SEE REMARK 999 UNP Q9W B -11 MET - cloning artifact UNP Q9W B -10 GLY - cloning artifact UNP Q9W	A	-3	HIS	-	cloning artifact	UNP Q9WY07
A 0 HIS - cloning artifact UNP Q9W A 38 ALA CYS engineered mutation UNP Q9W A 146 ARG - SEE REMARK 999 UNP Q9W A 147 ASP - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W A 149 ILE - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W A 152 ARG - SEE REMARK 999 UNP Q9W B -11 MET - cloning artifact UNP Q9W B -10 GLY - cloning artifact UNP Q9W	A	-2	HIS	-		UNP Q9WY07
A 38 ALA CYS engineered mutation UNP Q9W A 146 ARG - SEE REMARK 999 UNP Q9W A 147 ASP - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W A 149 ILE - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W A 152 ARG - SEE REMARK 999 UNP Q9W B -11 MET - cloning artifact UNP Q9W B -10 GLY - cloning artifact UNP Q9W	A	-1	HIS	-	cloning artifact	UNP Q9WY07
A 146 ARG - SEE REMARK 999 UNP Q9W A 147 ASP - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W A 149 ILE - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W A 152 ARG - SEE REMARK 999 UNP Q9W B -11 MET - cloning artifact UNP Q9W B -10 GLY - cloning artifact UNP Q9W	A	0	HIS	_	cloning artifact	UNP Q9WY07
A 147 ASP - SEE REMARK 999 UNP Q9W A 148 LEU - SEE REMARK 999 UNP Q9W A 149 ILE - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W A 152 ARG - SEE REMARK 999 UNP Q9W B -11 MET - cloning artifact UNP Q9W B -10 GLY - cloning artifact UNP Q9W	A	38	ALA	CYS	engineered mutation	UNP Q9WY07
A 148 LEU - SEE REMARK 999 UNP Q9W A 149 ILE - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W A 152 ARG - SEE REMARK 999 UNP Q9W B -11 MET - cloning artifact UNP Q9W B -10 GLY - cloning artifact UNP Q9W	A	146	ARG	_	SEE REMARK 999	UNP Q9WY07
A 149 ILE - SEE REMARK 999 UNP Q9W A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W A 152 ARG - SEE REMARK 999 UNP Q9W B -11 MET - cloning artifact UNP Q9W B -10 GLY - cloning artifact UNP Q9W	A	147	ASP	-	SEE REMARK 999	UNP Q9WY07
A 150 ASN - SEE REMARK 999 UNP Q9W A 151 SER - SEE REMARK 999 UNP Q9W A 152 ARG - SEE REMARK 999 UNP Q9W B -11 MET - cloning artifact UNP Q9W B -10 GLY - cloning artifact UNP Q9W	A	148	LEU	_	SEE REMARK 999	UNP Q9WY07
A 151 SER - SEE REMARK 999 UNP Q9W A 152 ARG - SEE REMARK 999 UNP Q9W B -11 MET - cloning artifact UNP Q9W B -10 GLY - cloning artifact UNP Q9W	A	149	ILE	_	SEE REMARK 999	UNP Q9WY07
A 152 ARG - SEE REMARK 999 UNP Q9W B -11 MET - cloning artifact UNP Q9W B -10 GLY - cloning artifact UNP Q9W	A	150	ASN	-	SEE REMARK 999	UNP Q9WY07
B -11 MET - cloning artifact UNP Q9W B -10 GLY - cloning artifact UNP Q9W	A	151	SER	-	SEE REMARK 999	UNP Q9WY07
B -10 GLY - cloning artifact UNP Q9W	A	152	ARG	_	SEE REMARK 999	UNP Q9WY07
	В	-11	MET			UNP Q9WY07
R 0 SFR cloning artifact UND OOM	В	-10	GLY	<u> </u>		UNP Q9WY07
-5 SER - Coming at mace ONI Q9W	В	-9	SER	- cloning artifact		UNP Q9WY07
B -8 ASP - cloning artifact UNP Q9W	В	-8	ASP	9		UNP Q9WY07
B -7 LYS - cloning artifact UNP Q9W	В	-7	LYS	-	cloning artifact	UNP Q9WY07

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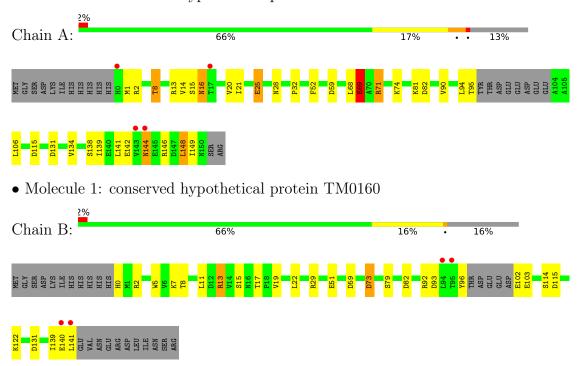
Chain	Residue	Modelled	Actual	Comment	Reference
В	-6	ILE	-	cloning artifact	UNP Q9WY07
В	-5	HIS	-	cloning artifact	UNP Q9WY07
В	-4	HIS	-	cloning artifact	UNP Q9WY07
В	-3	HIS	-	cloning artifact	UNP Q9WY07
В	-2	HIS	-	cloning artifact	UNP Q9WY07
В	-1	HIS	-	cloning artifact	UNP Q9WY07
В	0	HIS	-	cloning artifact	UNP Q9WY07
В	38	ALA	CYS	engineered mutation	UNP Q9WY07
В	146	ARG	-	SEE REMARK 999	UNP Q9WY07
В	147	ASP	-	SEE REMARK 999	UNP Q9WY07
В	148	LEU	-	SEE REMARK 999	UNP Q9WY07
В	149	ILE	-	SEE REMARK 999	UNP Q9WY07
В	150	ASN	-	SEE REMARK 999	UNP Q9WY07
В	151	SER	-	SEE REMARK 999	UNP Q9WY07
В	152	ARG	-	SEE REMARK 999	UNP Q9WY07



3 Residue-property plots (i)

These plots are drawn for all protein, RNA, DNA and oligosaccharide chains in the entry. The first graphic for a chain summarises the proportions of the various outlier classes displayed in the second graphic. The second graphic shows the sequence view annotated by issues in geometry and electron density. Residues are color-coded according to the number of geometric quality criteria for which they contain at least one outlier: green = 0, yellow = 1, orange = 2 and red = 3 or more. A red dot above a residue indicates a poor fit to the electron density (RSRZ > 2). Stretches of 2 or more consecutive residues without any outlier are shown as a green connector. Residues present in the sample, but not in the model, are shown in grey.

• Molecule 1: conserved hypothetical protein TM0160





4 Data and refinement statistics (i)

Property	Value	Source
Space group	P 1 21 1	Depositor
Cell constants	44.00Å 52.14Å 73.33Å	Depositor
a, b, c, α , β , γ	90.00° 97.64° 90.00°	Depositor
Resolution (Å)	72.55 - 2.80	Depositor
resolution (A)	31.64 - 2.80	EDS
% Data completeness	98.2 (72.55-2.80)	Depositor
(in resolution range)	98.3 (31.64-2.80)	EDS
R_{merge}	(Not available)	Depositor
R_{sym}	0.08	Depositor
$< I/\sigma(I) > 1$	2.44 (at 2.81Å)	Xtriage
Refinement program	REFMAC 5.1.24	Depositor
R, R_{free}	0.216 , 0.276	Depositor
It, It free	0.227 , 0.254	DCC
R_{free} test set	379 reflections (4.65%)	wwPDB-VP
Wilson B-factor (\mathring{A}^2)	58.5	Xtriage
Anisotropy	0.476	Xtriage
Bulk solvent $k_{sol}(e/Å^3)$, $B_{sol}(Å^2)$	0.33, 39.3	EDS
L-test for twinning ²	$ < L >=0.49, < L^2>=0.33$	Xtriage
Estimated twinning fraction	No twinning to report.	Xtriage
F_o, F_c correlation	0.92	EDS
Total number of atoms	2173	wwPDB-VP
Average B, all atoms (Å ²)	23.0	wwPDB-VP

Xtriage's analysis on translational NCS is as follows: The largest off-origin peak in the Patterson function is 7.86% of the height of the origin peak. No significant pseudotranslation is detected.

²Theoretical values of <|L|>, $<L^2>$ for acentric reflections are 0.5, 0.333 respectively for untwinned datasets, and 0.375, 0.2 for perfectly twinned datasets.



¹Intensities estimated from amplitudes.

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		
WIOI		RMSZ	# Z >5	RMSZ	# Z > 5	
1	A	0.78	0/1128	0.98	3/1534 (0.2%)	
1	В	0.74	0/1080	0.95	$4/1471 \ (0.3\%)$	
All	All	0.76	0/2208	0.96	7/3005 (0.2%)	

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	В	0	1

There are no bond length outliers.

All (7) bond angle outliers are listed below:

Mol	Chain	Res	Type	Atoms	\mathbf{Z}	$\mathbf{Observed}(^{o})$	$\operatorname{Ideal}({}^{o})$
1	A	115	ASP	CB-CG-OD2	8.18	125.66	118.30
1	В	131	ASP	CB-CG-OD2	7.49	125.04	118.30
1	A	59	ASP	CB-CG-OD2	6.81	124.43	118.30
1	A	82	ASP	CB-CG-OD2	6.43	124.09	118.30
1	В	73	ASP	CB-CG-OD2	6.39	124.05	118.30
1	В	115	ASP	CB-CG-OD2	5.64	123.38	118.30
1	В	93	ASP	CB-CG-OD2	5.21	122.98	118.30

There are no chirality outliers.

All (1) planarity outliers are listed below:

Mol	Chain	Res	Type	Group
1	В	51	GLU	Peptide



5.2 Too-close contacts (i)

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	A	1111	0	1160	16	0
1	В	1062	0	1093	5	0
All	All	2173	0	2253	21	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 5.

All (21) 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:A:28:ASN:HB3	1:A:146:ARG:NH2	1.94	0.81
1:A:28:ASN:CG	1:A:146:ARG:HH22	1.96	0.69
1:A:28:ASN:CB	1:A:146:ARG:HH22	2.09	0.65
1:A:28:ASN:HB3	1:A:146:ARG:HH22	1.62	0.59
1:A:144:ASN:O	1:A:148:LEU:HG	2.04	0.57
1:A:134:VAL:O	1:A:138:SER:OG	2.18	0.56
1:A:20:VAL:O	1:A:32:PRO:HA	2.06	0.55
1:A:149:ILE:O	1:A:149:ILE:HG22	2.09	0.51
1:A:28:ASN:CB	1:A:146:ARG:NH2	2.66	0.47
1:A:8:THR:HG22	1:A:21:ILE:CG1	2.45	0.46
1:A:141:LEU:HB2	1:A:146:ARG:HE	1.81	0.45
1:B:73:ASP:HB2	1:B:92:ARG:HB2	1.99	0.45
1:A:71:ARG:HD2	1:A:94:LEU:HD21	1.98	0.44
1:B:13:ARG:HB2	1:B:13:ARG:HH11	1.82	0.44
1:B:59:ASP:OD1	1:B:122:LYS:NZ	2.44	0.44
1:A:141:LEU:CD1	1:A:146:ARG:HG2	2.48	0.43
1:A:90:VAL:CG1	1:A:106:LEU:HD22	2.49	0.42
1:A:14:VAL:O	1:A:16:ASN:N	2.54	0.41
1:B:5:TRP:O	1:B:22:LEU:HA	2.20	0.41
1:B:7:LYS:O	1:B:8:THR:HG23	2.21	0.41
1:A:68:LEU:O	1:A:69:GLU:C	2.59	0.40

There are no symmetry-related clashes.



5.3 Torsion angles (i)

5.3.1 Protein backbone (i)

In the following table, the Percentiles column shows the percent Ramachandran outliers of the chain as a percentile score with respect to all X-ray entries followed by that with respect to entries of similar resolution.

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	Perc	entiles
1	A	139/164 (85%)	126 (91%)	9 (6%)	4 (3%)	4	15
1	В	133/164 (81%)	121 (91%)	11 (8%)	1 (1%)	19	49
All	All	272/328 (83%)	247 (91%)	20 (7%)	5 (2%)	8	28

All (5) Ramachandran outliers are listed below:

Mol	Chain	Res	Type
1	A	15	SER
1	A	148	LEU
1	A	25	GLU
1	A	69	GLU
1	В	82	ASP

5.3.2 Protein sidechains (i)

In the following table, the Percentiles column shows the percent sidechain outliers of the chain as a percentile score with respect to all X-ray entries followed by that with respect to entries of similar resolution.

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	3
1	A	123/144 (85%)	107 (87%)	16 (13%)	4 13	
1	В	116/144 (81%)	100 (86%)	16 (14%)	3 11	
All	All	239/288 (83%)	207 (87%)	32 (13%)	4 12	

All (32) residues with a non-rotameric sidechain are listed below:

-		
1 A	1	MET

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Mol	Chain	Res	Type
1	A	2	ARG
1	A	8	THR
1	A	13	ARG
1	A	16	ASN
1	A	25	GLU
1	A	52	PHE
1	A	69	GLU
1	A	71	ARG
1	A	74	LYS
1	A	81	LYS
1	A	95	THR
1	A	131	ASP
1	A	139	ILE
1	A	142	GLU
1	A	144	ASN
1	В	0	HIS
1	В	2	ARG
1	В	11	LEU
1	В	13	ARG
1	В	15	SER
1	В	17	THR
1	В	19	VAL
1	В	29	ARG
1	В	79	SER
1	В	96	TYR
1	В	102	GLU
1	В	103	GLU
1	В	114	SER
1	В	139	ILE
1	В	140	GLU
1	В	141	LEU

Sometimes side chains can be flipped to improve hydrogen bonding and reduce clashes. All (1) such side chains are listed below:

Mol	Chain	Res	Type
1	В	83	ASN

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 monosaccharides 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 Fit of model and data (i)

6.1 Protein, DNA and RNA chains (i)

In the following table, the column labelled '#RSRZ>2' contains the number (and percentage) of RSRZ outliers, followed by percent RSRZ outliers for the chain as percentile scores relative to all X-ray entries and entries of similar resolution. The OWAB column contains the minimum, median, 95^{th} percentile and maximum values of the occupancy-weighted average B-factor per residue. The column labelled 'Q< 0.9' lists the number of (and percentage) of residues with an average occupancy less than 0.9.

Mol	Chain	Analysed	<RSRZ $>$	# RSRZ > 2	$OWAB(A^2)$	Q < 0.9
1	A	143/164 (87%)	-0.16	4 (2%) 53 43	17, 22, 30, 35	0
1	В	137/164 (83%)	-0.10	4 (2%) 51 41	16, 23, 28, 33	0
All	All	280/328 (85%)	-0.13	8 (2%) 51 41	16, 23, 29, 35	0

All (8) RSRZ outliers are listed below:

Mol	Chain	Res	Type	RSRZ
1	A	143	VAL	5.5
1	A	144	ASN	3.3
1	В	94	LEU	2.5
1	A	17	THR	2.5
1	В	95	THR	2.4
1	A	0	HIS	2.3
1	В	140	GLU	2.3
1	В	141	LEU	2.0

6.2 Non-standard residues in protein, DNA, RNA chains (i)

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

6.3 Carbohydrates (i)

There are no monosaccharides in this entry.

6.4 Ligands (i)

There are no ligands in this entry.



6.5 Other polymers (i)

There are no such residues in this entry.

