



# wwPDB X-ray Structure Validation Summary Report ⓘ

Mar 14, 2018 – 04:04 am GMT

PDB ID : 1VQR  
Title : Crystal structure of a virulence factor (cj0248) from campylobacter jejuni subsp. jejuni at 2.25 Å resolution  
Authors : Joint Center for Structural Genomics (JCSG)  
Deposited on : 2004-12-17  
Resolution : 2.25 Å(reported)

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

We welcome your comments at [validation@mail.wwpdb.org](mailto: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 ⓘ symbol.

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

MolProbity	:	4.02b-467
Xtriage (Phenix)	:	1.13
EDS	:	trunk31020
Percentile statistics	:	20171227.v01 (using entries in the PDB archive December 27th 2017)
Refmac	:	5.8.0158
CCP4	:	7.0 (Gargrove)
Ideal geometry (proteins)	:	Engh & Huber (2001)
Ideal geometry (DNA, RNA)	:	Parkinson et al. (1996)
Validation Pipeline (wwPDB-VP)	:	trunk31020

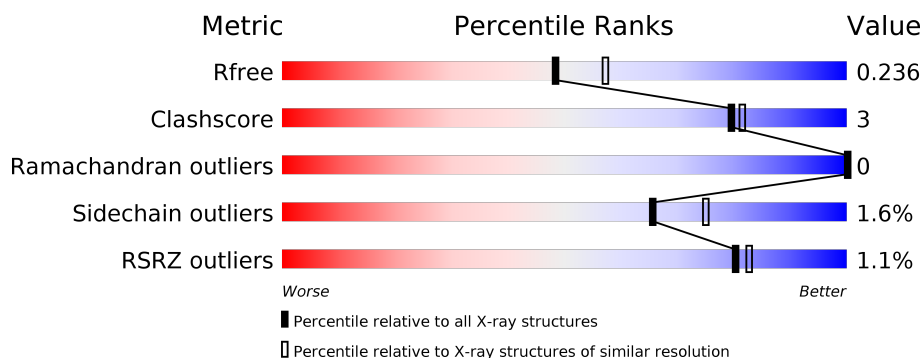
# 1 Overall quality at a glance

The following experimental techniques were used to determine the structure:

*X-RAY DIFFRACTION*

The reported resolution of this entry is 2.25 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)	Similar resolution (#Entries, resolution range(Å))
$R_{free}$	111664	1178 (2.26-2.26)
Clashscore	122126	1286 (2.26-2.26)
Ramachandran outliers	120053	1253 (2.26-2.26)
Sidechain outliers	120020	1254 (2.26-2.26)
RSRZ outliers	108989	1158 (2.26-2.26)

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 on the lower bar indicate the fraction of residues that contain outliers for  $\geq 3$ , 2, 1 and 0 types of geometric quality criteria. 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 electron density. The numeric value is given above the bar.

Mol	Chain	Length	Quality of chain
1	A	297	
1	B	297	
1	C	297	
1	D	297	

## 2 Entry composition

There are 2 unique types of molecules in this entry. The entry contains 8688 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 hypothetical protein Cj0248.

Mol	Chain	Residues	Atoms						ZeroOcc	AltConf	Trace
1	A	284	Total	C	N	O	S	Se	0	1	0
			2210	1424	362	415	3	6			
1	B	269	Total	C	N	O	S	Se	0	1	0
			2076	1351	338	379	3	5			
1	C	281	Total	C	N	O	S	Se	0	0	0
			2149	1392	350	398	3	6			
1	D	267	Total	C	N	O	S	Se	0	2	0
			2060	1339	337	377	3	4			

There are 72 discrepancies between the modelled and reference sequences:

Chain	Residue	Modelled	Actual	Comment	Reference
A	-11	MSE	-	LEADER SEQUENCE	UNP Q9PIP7
A	-10	GLY	-	LEADER SEQUENCE	UNP Q9PIP7
A	-9	SER	-	LEADER SEQUENCE	UNP Q9PIP7
A	-8	ASP	-	LEADER SEQUENCE	UNP Q9PIP7
A	-7	LYS	-	LEADER SEQUENCE	UNP Q9PIP7
A	-6	ILE	-	LEADER SEQUENCE	UNP Q9PIP7
A	-5	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
A	-4	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
A	-3	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
A	-2	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
A	-1	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
A	0	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
A	1	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
A	5	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
A	40	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
A	52	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
A	91	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
A	144	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
B	-11	MSE	-	LEADER SEQUENCE	UNP Q9PIP7
B	-10	GLY	-	LEADER SEQUENCE	UNP Q9PIP7
B	-9	SER	-	LEADER SEQUENCE	UNP Q9PIP7

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Chain	Residue	Modelled	Actual	Comment	Reference
B	-8	ASP	-	LEADER SEQUENCE	UNP Q9PIP7
B	-7	LYS	-	LEADER SEQUENCE	UNP Q9PIP7
B	-6	ILE	-	LEADER SEQUENCE	UNP Q9PIP7
B	-5	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
B	-4	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
B	-3	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
B	-2	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
B	-1	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
B	0	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
B	1	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
B	5	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
B	40	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
B	52	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
B	91	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
B	144	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
C	-11	MSE	-	LEADER SEQUENCE	UNP Q9PIP7
C	-10	GLY	-	LEADER SEQUENCE	UNP Q9PIP7
C	-9	SER	-	LEADER SEQUENCE	UNP Q9PIP7
C	-8	ASP	-	LEADER SEQUENCE	UNP Q9PIP7
C	-7	LYS	-	LEADER SEQUENCE	UNP Q9PIP7
C	-6	ILE	-	LEADER SEQUENCE	UNP Q9PIP7
C	-5	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
C	-4	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
C	-3	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
C	-2	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
C	-1	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
C	0	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
C	1	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
C	5	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
C	40	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
C	52	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
C	91	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
C	144	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
D	-11	MSE	-	LEADER SEQUENCE	UNP Q9PIP7
D	-10	GLY	-	LEADER SEQUENCE	UNP Q9PIP7
D	-9	SER	-	LEADER SEQUENCE	UNP Q9PIP7
D	-8	ASP	-	LEADER SEQUENCE	UNP Q9PIP7
D	-7	LYS	-	LEADER SEQUENCE	UNP Q9PIP7
D	-6	ILE	-	LEADER SEQUENCE	UNP Q9PIP7
D	-5	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
D	-4	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
D	-3	HIS	-	LEADER SEQUENCE	UNP Q9PIP7

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Chain	Residue	Modelled	Actual	Comment	Reference
D	-2	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
D	-1	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
D	0	HIS	-	LEADER SEQUENCE	UNP Q9PIP7
D	1	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
D	5	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
D	40	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
D	52	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
D	91	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7
D	144	MSE	MET	MODIFIED RESIDUE	UNP Q9PIP7

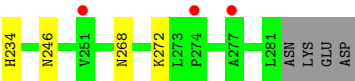
- Molecule 2 is water.

Mol	Chain	Residues	Atoms	ZeroOcc	AltConf
2	A	68	Total O 68 68	0	0
2	B	47	Total O 47 47	0	0
2	C	36	Total O 36 36	0	0
2	D	42	Total O 42 42	0	0



- Molecule 1: hypothetical protein Cj0248





## 4 Data and refinement statistics

Property	Value	Source
Space group	C 1 2 1	Depositor
Cell constants a, b, c, $\alpha$ , $\beta$ , $\gamma$	123.14Å 122.09Å 96.90Å 90.00° 120.58° 90.00°	Depositor
Resolution (Å)	61.04 – 2.25 61.05 – 2.25	Depositor EDS
% Data completeness (in resolution range)	97.3 (61.04-2.25) 97.3 (61.05-2.25)	Depositor EDS
$R_{merge}$	(Not available)	Depositor
$R_{sym}$	0.05	Depositor
$\langle I/\sigma(I) \rangle$ <sup>1</sup>	1.91 (at 2.25Å)	Xtriage
Refinement program	REFMAC	Depositor
R, $R_{free}$	0.193 , 0.233 0.199 , 0.236	Depositor DCC
$R_{free}$ test set	2850 reflections (5.01%)	wwPDB-VP
Wilson B-factor (Å <sup>2</sup> )	44.5	Xtriage
Anisotropy	0.160	Xtriage
Bulk solvent $k_{sol}$ (e/Å <sup>3</sup> ), $B_{sol}$ (Å <sup>2</sup> )	0.32 , 50.3	EDS
L-test for twinning <sup>2</sup>	$\langle  L  \rangle = 0.49$ , $\langle L^2 \rangle = 0.32$	Xtriage
Estimated twinning fraction	No twinning to report.	Xtriage
$F_o, F_c$ correlation	0.95	EDS
Total number of atoms	8688	wwPDB-VP
Average B, all atoms (Å <sup>2</sup> )	48.0	wwPDB-VP

Xtriage's analysis on translational NCS is as follows: *The largest off-origin peak in the Patterson function is 4.75% of the height of the origin peak. No significant pseudotranslation is detected.*

<sup>1</sup>Intensities estimated from amplitudes.

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



## 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	A	0.71	0/2249	0.73	2/3044 (0.1%)
1	B	0.69	0/2117	0.68	0/2873
1	C	0.70	2/2186 (0.1%)	0.65	0/2970
1	D	0.64	0/2105	0.65	0/2859
All	All	0.69	2/8657 (0.0%)	0.68	2/11746 (0.0%)

All (2) bond length outliers are listed below:

Mol	Chain	Res	Type	Atoms	Z	Observed(Å)	Ideal(Å)
1	C	220	GLU	CD-OE2	9.32	1.35	1.25
1	C	220	GLU	CD-OE1	6.15	1.32	1.25

All (2) bond angle outliers are listed below:

Mol	Chain	Res	Type	Atoms	Z	Observed(°)	Ideal(°)
1	A	147	ARG	NE-CZ-NH2	-5.35	117.62	120.30
1	A	198	ARG	NE-CZ-NH2	-5.27	117.66	120.30

There are no chirality outliers.

There are no planarity outliers.

### 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	2210	0	2187	14	0
1	B	2076	0	2048	14	0

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Mol	Chain	Non-H	H(model)	H(added)	Clashes	Symm-Clashes
1	C	2149	0	2110	12	0
1	D	2060	0	2021	12	0
2	A	68	0	0	1	0
2	B	47	0	0	1	0
2	C	36	0	0	1	0
2	D	42	0	0	2	0
All	All	8688	0	8366	51	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 3.

The worst 5 of 51 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:C:5:MSE:SE	1:C:5:MSE:CE	2.16	1.44
1:D:129[B]:ASN:ND2	2:D:325:HOH:O	2.25	0.70
1:A:216:HIS:ND1	2:A:343:HOH:O	2.28	0.65
1:C:5:MSE:HG2	1:C:184:LEU:O	2.03	0.59
1:D:155:ASN:O	1:D:159:GLN:HG2	2.04	0.58

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	Percentiles	
1	A	281/297 (95%)	279 (99%)	2 (1%)	0	100	100
1	B	264/297 (89%)	258 (98%)	6 (2%)	0	100	100
1	C	279/297 (94%)	273 (98%)	6 (2%)	0	100	100
1	D	263/297 (89%)	259 (98%)	4 (2%)	0	100	100
All	All	1087/1188 (92%)	1069 (98%)	18 (2%)	0	100	100

There are no Ramachandran outliers to report.

### 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 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	
1	A	237/256 (93%)	232 (98%)	5 (2%)	56	65
1	B	218/256 (85%)	215 (99%)	3 (1%)	69	79
1	C	224/256 (88%)	220 (98%)	4 (2%)	62	71
1	D	216/256 (84%)	213 (99%)	3 (1%)	69	79
All	All	895/1024 (87%)	880 (98%)	15 (2%)	65	73

5 of 15 residues with a non-rotameric sidechain are listed below:

Mol	Chain	Res	Type
1	B	184	LEU
1	B	279	GLU
1	D	27	ARG
1	B	129	ASN
1	C	266	LEU

Some sidechains can be flipped to improve hydrogen bonding and reduce clashes. All (5) such sidechains are listed below:

Mol	Chain	Res	Type
1	A	36	ASN
1	C	118	ASN
1	C	173	ASN
1	C	216	HIS
1	C	246	ASN

### 5.3.3 RNA ⓘ

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 carbohydrates 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<sup>th</sup> 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(Å <sup>2</sup> )	Q<0.9
1	A	278/297 (93%)	-0.23	1 (0%) 92 93	38, 47, 60, 72	0
1	B	264/297 (88%)	-0.20	1 (0%) 92 93	37, 47, 60, 69	0
1	C	275/297 (92%)	0.01	5 (1%) 68 72	37, 47, 60, 70	0
1	D	262/297 (88%)	-0.02	5 (1%) 66 70	37, 47, 58, 66	0
All	All	1079/1188 (90%)	-0.11	12 (1%) 80 83	37, 47, 60, 72	0

The worst 5 of 12 RSRZ outliers are listed below:

Mol	Chain	Res	Type	RSRZ
1	C	254	LEU	4.5
1	C	251	VAL	3.2
1	D	98	ASN	3.1
1	C	247	ALA	2.9
1	D	277	ALA	2.5

### 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 carbohydrates 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.