



Full wwPDB NMR Structure Validation Report ⓘ

May 9, 2022 – 12:50 PM JST

PDB ID : 7BZH
Title : Solution structure of a DNA binding protein from *Sulfolobus islandicus*
Authors : Zhang, Z.; Liu, X.
Deposited on : 2020-04-28

This is a Full wwPDB NMR 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/NMRValidationReportHelp>

with specific help available everywhere you see the ⓘ symbol.

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

MolProbity : 4.02b-467
Percentile statistics : 20191225.v01 (using entries in the PDB archive December 25th 2019)
RCI : v_1n_11_5_13_A (Berjanski et al., 2005)
PANAV : Wang et al. (2010)
ShiftChecker : 2.28.1
Ideal geometry (proteins) : Engh & Huber (2001)
Ideal geometry (DNA, RNA) : Parkinson et al. (1996)
Validation Pipeline (wwPDB-VP) : 2.28.1

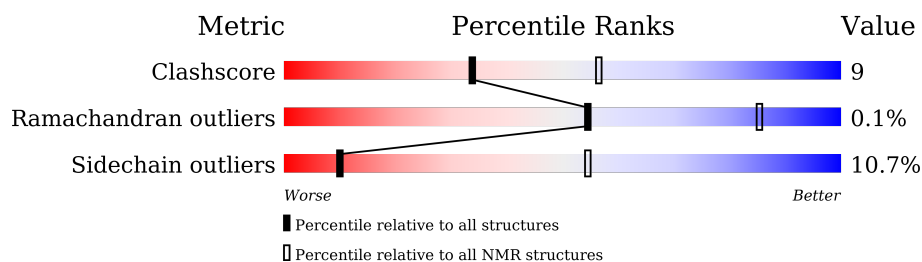
1 Overall quality at a glance

The following experimental techniques were used to determine the structure:

SOLUTION NMR

The overall completeness of chemical shifts assignment is 79%.

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)	NMR archive (#Entries)
Clashscore	158937	12864
Ramachandran outliers	154571	11451
Sidechain outliers	154315	11428

The table below summarises the geometric issues observed across the polymeric chains and their fit to the experimental data. The red, orange, yellow and green segments indicate the fraction of residues that contain outliers for ≥ 3 , 2, 1 and 0 types of geometric quality criteria. A cyan segment indicates the fraction of residues that are not part of the well-defined cores, and 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 $\leq 5\%$

Mol	Chain	Length	Quality of chain
1	A	59	 66% 22% • 8%

2 Ensemble composition and analysis

This entry contains 20 models. Model 1 is the overall representative, medoid model (most similar to other models).

The following residues are included in the computation of the global validation metrics.

Well-defined (core) protein residues			
Well-defined core	Residue range (total)	Backbone RMSD (Å)	Medoid model
1	A:4-A:57 (54)	0.17	1

Ill-defined regions of proteins are excluded from the global statistics.

Ligands and non-protein polymers are included in the analysis.

The models can be grouped into 3 clusters. No single-model clusters were found.

Cluster number	Models
1	1, 2, 4, 5, 8, 11, 13, 15, 16, 17, 19
2	3, 7, 9, 10, 14, 20
3	6, 12, 18

3 Entry composition [i](#)

There is only 1 type of molecule in this entry. The entry contains 1032 atoms, of which 521 are hydrogens and 0 are deuteriums.

- Molecule 1 is a protein called Sul7s.

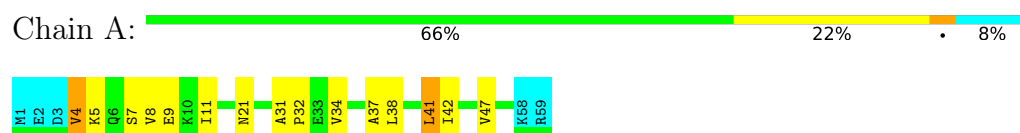
Mol	Chain	Residues	Atoms						Trace
1	A	59	Total	C	H	N	O	S	0
			1032	333	521	86	91	1	

4 Residue-property plots

4.1 Average score per residue in the NMR ensemble

These plots are provided for all protein, RNA, DNA and oligosaccharide chains in the entry. The first graphic is the same as shown in the summary in section 1 of this report. The second graphic shows the sequence where residues are colour-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. Stretches of 2 or more consecutive residues without any outliers are shown as green connectors. Residues which are classified as ill-defined in the NMR ensemble, are shown in cyan with an underline colour-coded according to the previous scheme. Residues which were present in the experimental sample, but not modelled in the final structure are shown in grey.

- Molecule 1: Sul7s

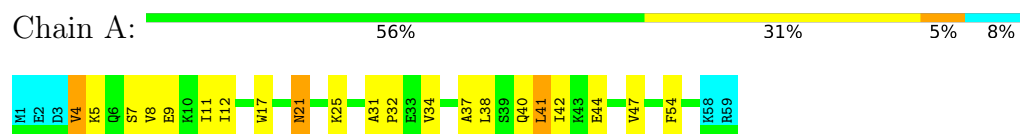


4.2 Scores per residue for each member of the ensemble

Colouring as in section 4.1 above.

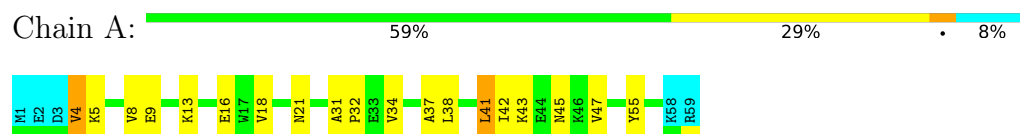
4.2.1 Score per residue for model 1 (medoid)

- Molecule 1: Sul7s



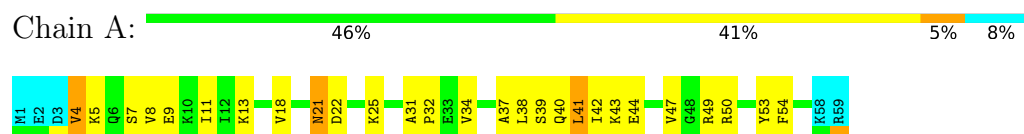
4.2.2 Score per residue for model 2

- Molecule 1: Sul7s



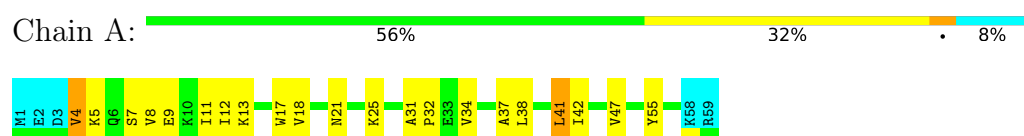
4.2.3 Score per residue for model 3

- Molecule 1: Sul7s



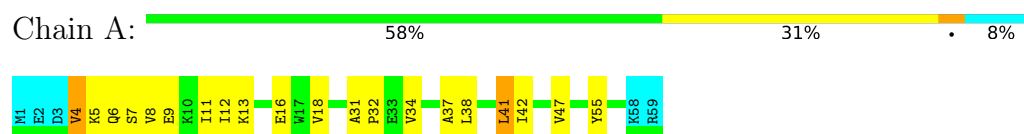
4.2.4 Score per residue for model 4

- Molecule 1: Sul7s



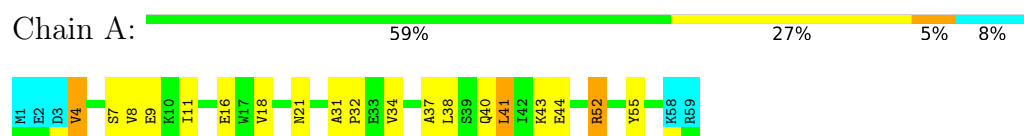
4.2.5 Score per residue for model 5

- Molecule 1: Sul7s



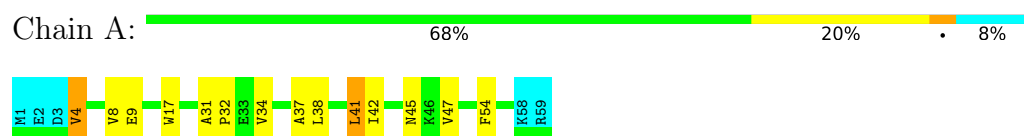
4.2.6 Score per residue for model 6

- Molecule 1: Sul7s



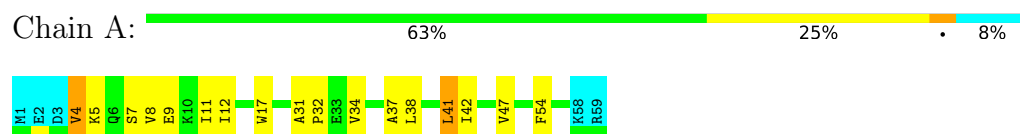
4.2.7 Score per residue for model 7

- Molecule 1: Sul7s



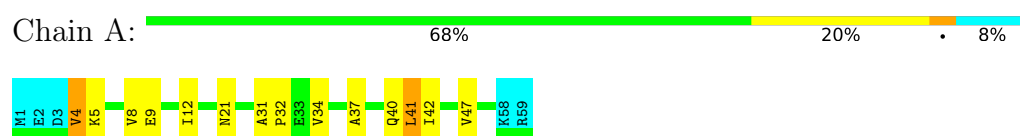
4.2.8 Score per residue for model 8

- Molecule 1: Sul7s



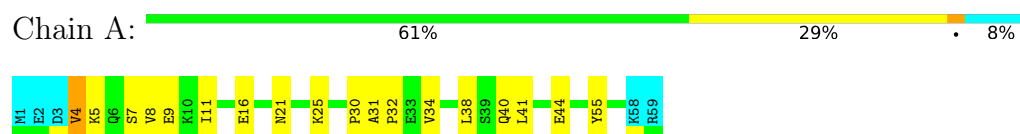
4.2.9 Score per residue for model 9

- Molecule 1: Sul7s



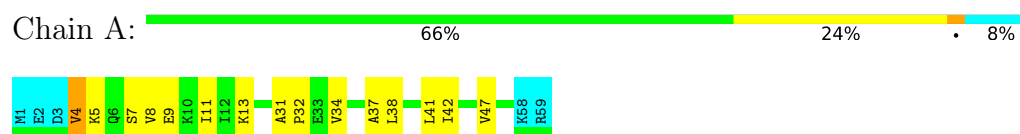
4.2.10 Score per residue for model 10

- Molecule 1: Sul7s



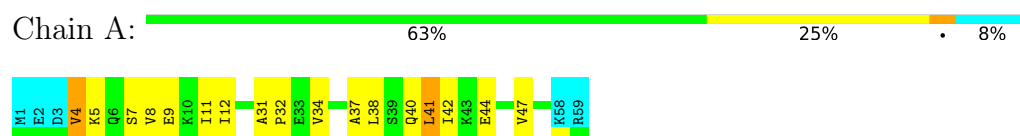
4.2.11 Score per residue for model 11

- Molecule 1: Sul7s



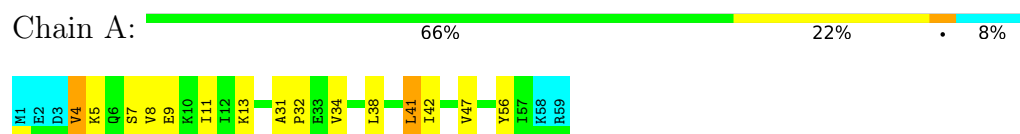
4.2.12 Score per residue for model 12

- Molecule 1: Sul7s



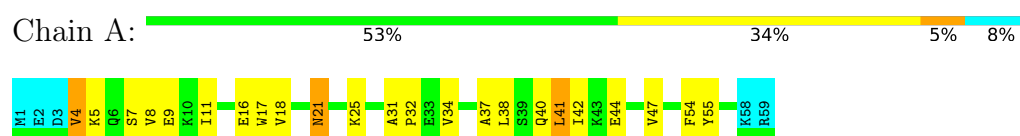
4.2.13 Score per residue for model 13

- Molecule 1: Sul7s



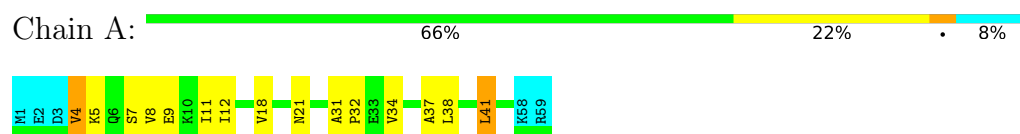
4.2.14 Score per residue for model 14

- Molecule 1: Sul7s



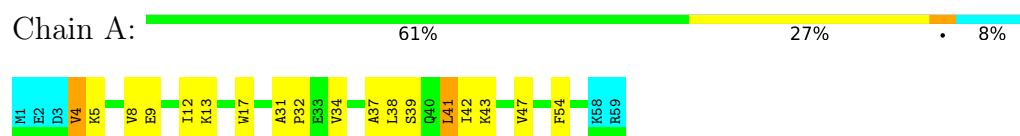
4.2.15 Score per residue for model 15

- Molecule 1: Sul7s



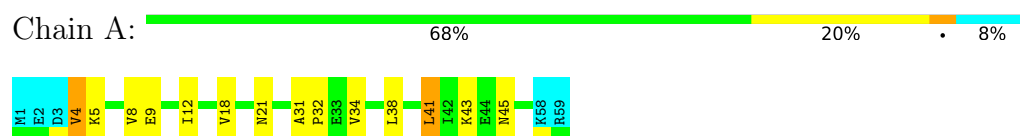
4.2.16 Score per residue for model 16

- Molecule 1: Sul7s



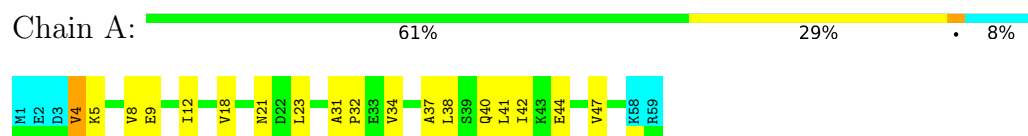
4.2.17 Score per residue for model 17

- Molecule 1: Sul7s



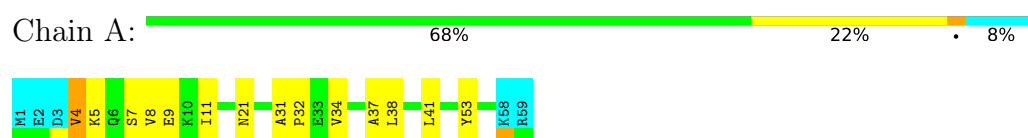
4.2.18 Score per residue for model 18

- Molecule 1: Sul7s



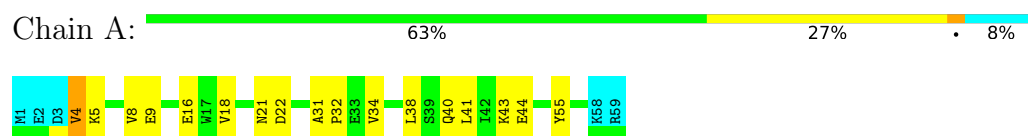
4.2.19 Score per residue for model 19

- Molecule 1: Sul7s



4.2.20 Score per residue for model 20

- Molecule 1: Sul7s



5 Refinement protocol and experimental data overview

The models were refined using the following method: *simulated annealing*.

Of the 100 calculated structures, 20 were deposited, based on the following criterion: *structures with the lowest energy*.

The following table shows the software used for structure solution, optimisation and refinement.

Software name	Classification	Version
CNS	refinement	
ARIA	structure calculation	

The following table shows chemical shift validation statistics as aggregates over all chemical shift files. Detailed validation can be found in section 7 of this report.

Chemical shift file(s)	working_cs.cif
Number of chemical shift lists	1
Total number of shifts	765
Number of shifts mapped to atoms	704
Number of unparsed shifts	0
Number of shifts with mapping errors	0
Number of shifts with mapping warnings	61
Assignment completeness (well-defined parts)	79%

6 Model quality [i](#)

6.1 Standard geometry [i](#)

There are no covalent bond-length or bond-angle outliers.

There are no bond-length outliers.

There are no bond-angle outliers.

There are no chirality outliers.

There are no planarity outliers.

6.2 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 each 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 averaged over the ensemble.

Mol	Chain	Non-H	H(model)	H(added)	Clashes
1	A	465	474	474	8±2
All	All	9300	9480	9480	169

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 9.

All unique clashes are listed below, sorted by their clash magnitude.

Atom-1	Atom-2	Clash(Å)	Distance(Å)	Models	
				Worst	Total
1:A:16:GLU:O	1:A:55:TYR:HA	0.59	1.97	20	6
1:A:42:ILE:HA	1:A:47:VAL:O	0.58	1.98	11	14
1:A:5:LYS:O	1:A:9:GLU:HG2	0.56	2.01	19	17
1:A:7:SER:O	1:A:11:ILE:HG12	0.56	2.00	8	6
1:A:9:GLU:O	1:A:13:LYS:HG3	0.54	2.03	16	5
1:A:31:ALA:N	1:A:32:PRO:HD2	0.53	2.19	18	20
1:A:39:SER:HA	1:A:42:ILE:HD12	0.52	1.81	16	2
1:A:8:VAL:O	1:A:12:ILE:HG12	0.51	2.05	8	9
1:A:5:LYS:HG3	1:A:41:LEU:HD21	0.51	1.83	5	7
1:A:4:VAL:O	1:A:8:VAL:HG23	0.51	2.06	18	20
1:A:37:ALA:O	1:A:41:LEU:HD22	0.50	2.07	4	15
1:A:9:GLU:HB3	1:A:41:LEU:HG	0.49	1.84	7	5
1:A:49:ARG:HG2	1:A:54:PHE:CE2	0.49	2.42	3	1
1:A:21:ASN:O	1:A:25:LYS:HD3	0.49	2.08	10	5

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Atom-1	Atom-2	Clash(Å)	Distance(Å)	Models	
				Worst	Total
1:A:40:GLN:O	1:A:44:GLU:HG3	0.48	2.09	1	7
1:A:17:TRP:HA	1:A:54:PHE:O	0.47	2.09	7	5
1:A:7:SER:O	1:A:11:ILE:HG13	0.47	2.09	19	7
1:A:49:ARG:HG2	1:A:54:PHE:CD2	0.45	2.45	3	1
1:A:18:VAL:CG2	1:A:23:LEU:HG	0.45	2.42	18	1
1:A:37:ALA:O	1:A:41:LEU:HG	0.44	2.12	18	1
1:A:9:GLU:O	1:A:13:LYS:HB2	0.43	2.13	5	1
1:A:5:LYS:HG3	1:A:41:LEU:CD2	0.43	2.43	5	6
1:A:18:VAL:HB	1:A:22:ASP:OD1	0.42	2.15	3	1
1:A:40:GLN:O	1:A:44:GLU:HG2	0.41	2.15	14	1
1:A:50:ARG:HG3	1:A:53:TYR:CE1	0.41	2.51	3	1
1:A:17:TRP:CZ3	1:A:55:TYR:HB3	0.41	2.50	4	1
1:A:37:ALA:O	1:A:40:GLN:HB2	0.41	2.16	9	1
1:A:13:LYS:HG2	1:A:56:TYR:CD2	0.41	2.51	13	1
1:A:12:ILE:HD13	1:A:47:VAL:HG11	0.41	1.92	5	1
1:A:52:ARG:HD3	1:A:52:ARG:H	0.41	1.75	6	1

6.3 Torsion angles [i](#)

6.3.1 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 PDB entries followed by that with respect to all NMR 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	A	54/59 (92%)	50±1 (93±2%)	4±1 (7±1%)	0±0 (0±0%)	54	85
All	All	1080/1180 (92%)	1007 (93%)	72 (7%)	1 (0%)	54	85

All 1 unique Ramachandran outliers are listed below.

Mol	Chain	Res	Type	Models (Total)
1	A	30	PRO	1

6.3.2 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 PDB entries followed by that with respect to all NMR 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	A	50/55 (91%)	45±2 (89±3%)	5±2 (11±3%)	10	54
All	All	1000/1100 (91%)	893 (89%)	107 (11%)	10	54

All 12 unique residues with a non-rotameric sidechain are listed below. They are sorted by the frequency of occurrence in the ensemble.

Mol	Chain	Res	Type	Models (Total)
1	A	4	VAL	20
1	A	34	VAL	20
1	A	38	LEU	19
1	A	41	LEU	16
1	A	21	ASN	11
1	A	18	VAL	8
1	A	43	LYS	6
1	A	45	ASN	3
1	A	6	GLN	1
1	A	52	ARG	1
1	A	53	TYR	1
1	A	22	ASP	1

6.3.3 RNA [i](#)

There are no RNA molecules in this entry.

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

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

6.5 Carbohydrates [i](#)

There are no monosaccharides in this entry.

6.6 Ligand geometry [i](#)

There are no ligands in this entry.

6.7 Other polymers [i](#)

There are no such molecules in this entry.

6.8 Polymer linkage issues [i](#)

There are no chain breaks in this entry.

7 Chemical shift validation

The completeness of assignment taking into account all chemical shift lists is 79% for the well-defined parts and 78% for the entire structure.

7.1 Chemical shift list 1

File name: working_cs.cif

Chemical shift list name: *assigned_chem_shift_list*

7.1.1 Bookkeeping

The following table shows the results of parsing the chemical shift list and reports the number of nuclei with statistically unusual chemical shifts.

Total number of shifts	765
Number of shifts mapped to atoms	704
Number of unparsed shifts	0
Number of shifts with mapping errors	0
Number of shifts with mapping warnings	61
Number of shift outliers (ShiftChecker)	0

The following assigned chemical shifts were not mapped to the molecules present in the coordinate file.

- No matching atoms found in structure. All 61 occurrences are reported below.

Chain	Res	Type	Atom	Shift Data		
				Value	Uncertainty	Ambiguity
A	35	TYR	HN	9.085	0.001	1
A	13	LYS	HN	8.255	0.001	1
A	11	ILE	HN	7.541	0.001	1
A	12	ILE	HN	7.88	0.001	1
A	5	LYS	HN	8.041	0.001	1
A	46	LYS	HN	8.031	0.002	1
A	20	PHE	HN	9.847	0.001	1
A	34	VAL	HN	7.872	0.002	1
A	31	ALA	HN	9.002	0.002	1
A	47	VAL	HN	6.785	0.001	1
A	36	ASP	HN	8.178	0.001	1
A	21	ASN	HN	8.423	0.001	1
A	56	TYR	HN	8.804	0.001	1
A	38	LEU	HN	8.348	0.002	1

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Chain	Res	Type	Atom	Shift Data		
				Value	Uncertainty	Ambiguity
A	8	VAL	HN	8.384	0.002	1
A	7	SER	HN	8.065	0.001	1
A	50	ARG	HN	9.075	0.002	1
A	10	LYS	HN	7.734	0.001	1
A	58	LYS	HN	8.111	0.001	1
A	41	LEU	HN	8.0	0.001	1
A	54	PHE	HN	9.423	0.003	1
A	6	GLN	HN	8.267	0.001	1
A	44	GLU	HN	7.931	0.001	1
A	59	ARG	HN	8.085	0.001	1
A	45	ASN	HN	8.215	0.001	1
A	18	VAL	HN	9.183	0.003	1
A	49	ARG	HN	8.394	0.002	1
A	48	GLY	HN	8.431	0.002	1
A	37	ALA	HN	7.695	0.001	1
A	55	TYR	HN	8.488	0.001	1
A	25	LYS	HN	7.274	0.001	1
A	4	VAL	HN	8.191	0.001	1
A	9	GLU	HN	8.26	0.0	1
A	24	LEU	HN	7.315	0.0	1
A	57	ILE	HN	7.416	0.003	1
A	53	TYR	HN	7.943	0.003	1
A	42	ILE	HN	8.243	0.002	1
A	43	LYS	HN	7.991	0.001	1
A	16	GLU	HN	8.194	0.002	1
A	33	GLU	HN	7.421	0.002	1
A	23	LEU	HN	7.68	0.001	1
A	27	ILE	HN	7.811	0.001	1
A	19	THR	HN	8.921	0.001	1
A	29	TYR	HN	7.728	0.001	1
A	17	TRP	HN	8.257	0.002	1
A	15	ARG	HN	8.165	0.002	1
A	14	ASP	HN	7.878	0.001	1
A	22	ASP	HN	7.785	0.001	1
A	26	TYR	HN	7.46	0.001	1
A	40	GLN	HN	7.891	0.001	1
A	39	SER	HN	8.335	0.001	1
A	3	ASP	HN	8.489	0.001	1
A	52	ARG	HN	8.728	0.004	1
A	15	ARG	HD1	2.928	0.0	1
A	15	ARG	HG1	1.578	0.003	1

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Chain	Res	Type	Atom	Shift Data		
				Value	Uncertainty	Ambiguity
A	49	ARG	HD1	3.097	0.002	1
A	59	ARG	HD1	3.213	0.001	1
A	5	LYS	HD1	1.648	0.0	1
A	43	LYS	HD1	1.689	0.004	1
A	46	LYS	HD1	1.679	0.004	1
A	58	LYS	HE1	3.103	-1.0	1

7.1.2 Chemical shift referencing

The following table shows the suggested chemical shift referencing corrections.

Nucleus	# values	Correction \pm precision, ppm	Suggested action
$^{13}\text{C}_\alpha$	59	-0.41 ± 0.15	None needed (< 0.5 ppm)
$^{13}\text{C}_\beta$	57	0.01 ± 0.09	None needed (< 0.5 ppm)
$^{13}\text{C}'$	53	0.09 ± 0.13	None needed (< 0.5 ppm)
^{15}N	53	0.53 ± 0.51	None needed (imprecise)

7.1.3 Completeness of resonance assignments

The following table shows the completeness of the chemical shift assignments for the well-defined regions of the structure. The overall completeness is 79%, i.e. 585 atoms were assigned a chemical shift out of a possible 745. 0 out of 9 assigned methyl groups (LEU and VAL) were assigned stereospecifically.

	Total	^1H	^{13}C	^{15}N
Backbone	208/264 (79%)	54/105 (51%)	104/108 (96%)	50/51 (98%)
Sidechain	342/403 (85%)	211/238 (89%)	127/143 (89%)	4/22 (18%)
Aromatic	35/78 (45%)	34/40 (85%)	0/37 (0%)	1/1 (100%)
Overall	585/745 (79%)	299/383 (78%)	231/288 (80%)	55/74 (74%)

The following table shows the completeness of the chemical shift assignments for the full structure. The overall completeness is 78%, i.e. 636 atoms were assigned a chemical shift out of a possible 820. 0 out of 9 assigned methyl groups (LEU and VAL) were assigned stereospecifically.

	Total	^1H	^{13}C	^{15}N
Backbone	224/289 (78%)	59/115 (51%)	112/118 (95%)	53/56 (95%)
Sidechain	377/453 (83%)	233/268 (87%)	140/159 (88%)	4/26 (15%)
Aromatic	35/78 (45%)	34/40 (85%)	0/37 (0%)	1/1 (100%)
Overall	636/820 (78%)	326/423 (77%)	252/314 (80%)	58/83 (70%)

7.1.4 Statistically unusual chemical shifts [i](#)

There are no statistically unusual chemical shifts.

7.1.5 Random Coil Index (RCI) plots [i](#)

The image below reports *random coil index* values for the protein chains in the structure. The height of each bar gives a probability of a given residue to be disordered, as predicted from the available chemical shifts and the amino acid sequence. A value above 0.2 is an indication of significant predicted disorder. The colour of the bar shows whether the residue is in the well-defined core (black) or in the ill-defined residue ranges (cyan), as described in section 2 on ensemble composition.

Random coil index (RCI) for chain A:

