



Full wwPDB NMR Structure Validation Report ⓘ

Feb 13, 2017 – 02:10 am GMT

PDB ID : 2RQH
Title : Structure of GSPT1/ERF3A-PABC
Authors : Osawa, M.; Nakanishi, T.; Hosoda, N.; Uchida, S.; Hoshino, T.; Katada, I.; Shimada, I.
Deposited on : 2009-05-08

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

<http://wwpdb.org/validation/2016/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:

Cyrange : Kirchner and Güntert (2011)
NmrClust : Kelley et al. (1996)
MolProbity : 4.02b-467
Percentile statistics : 20161228.v01 (using entries in the PDB archive December 28th 2016)
RCI : v_1n_11_5_13_A (Berjanski et al., 2005)
PANAV : Wang et al. (2010)
ShiftChecker : trunk28760
Ideal geometry (proteins) : Engh & Huber (2001)
Ideal geometry (DNA, RNA) : Parkinson et al. (1996)
Validation Pipeline (wwPDB-VP) : recalc28949

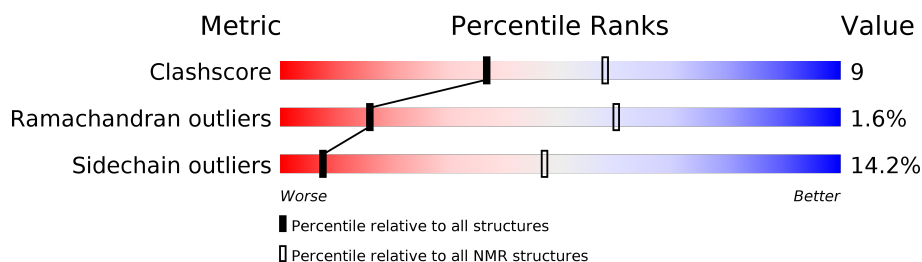
1 Overall quality at a glance

The following experimental techniques were used to determine the structure:

SOLUTION NMR

The overall completeness of chemical shifts assignment was not calculated.

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	125131	11601
Ramachandran outliers	121729	10391
Sidechain outliers	121581	10367

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	22	 32% 9% 59%
2	B	83	 57% 16% • 27%

2 Ensemble composition and analysis

This entry contains 20 models. Model 15 is the overall representative, medoid model (most similar to other models). The authors have identified model 1 as representative, based on the following criterion: *lowest energy*.

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:77-A:85, B:556-B:616 (70)	0.28	15

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 2 clusters. No single-model clusters were found.

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

3 Entry composition [i](#)

There are 2 unique types of molecules in this entry. The entry contains 1599 atoms, of which 806 are hydrogens and 0 are deuteriums.

- Molecule 1 is a protein called G1 to S phase transition 1.

Mol	Chain	Residues	Atoms					Trace
1	A	22	Total	C	H	N	O	0
			333	112	166	29	26	

- Molecule 2 is a protein called Polyadenylate-binding protein 1.

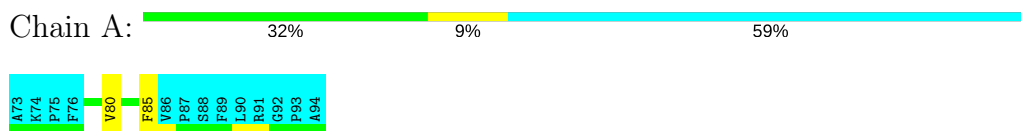
Mol	Chain	Residues	Atoms						Trace
2	B	83	Total	C	H	N	O	S	0
			1266	392	640	107	122	5	

4 Residue-property plots

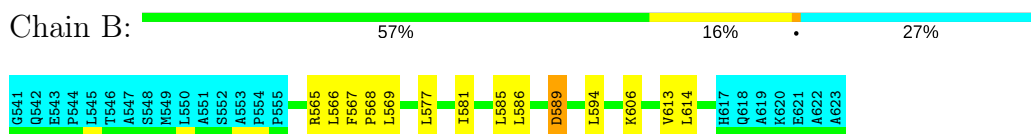
4.1 Average score per residue in the NMR ensemble

These plots are provided for all protein, RNA and DNA 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: G1 to S phase transition 1



- Molecule 2: Polyadenylate-binding protein 1

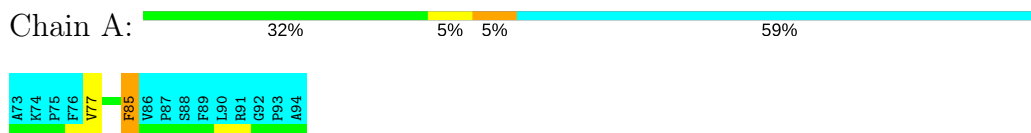


4.2 Scores per residue for each member of the ensemble

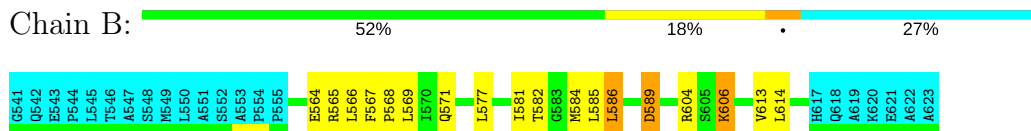
Colouring as in section 4.1 above.

4.2.1 Score per residue for model 1

- Molecule 1: G1 to S phase transition 1

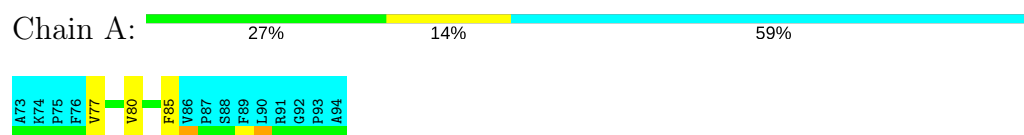


- Molecule 2: Polyadenylate-binding protein 1

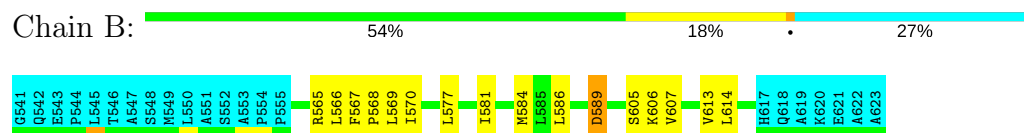


4.2.2 Score per residue for model 2

- Molecule 1: G1 to S phase transition 1

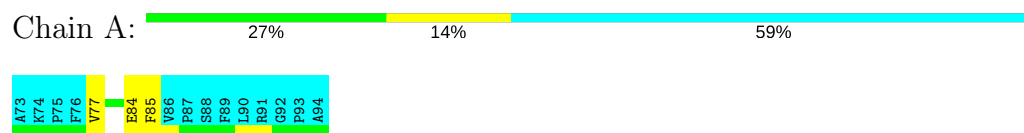


- Molecule 2: Polyadenylate-binding protein 1

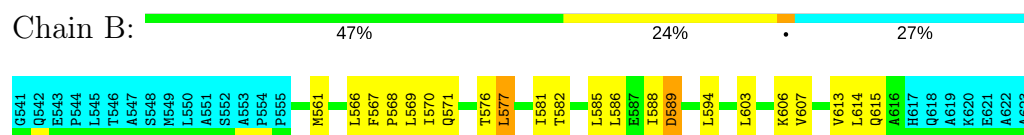


4.2.3 Score per residue for model 3

- Molecule 1: G1 to S phase transition 1

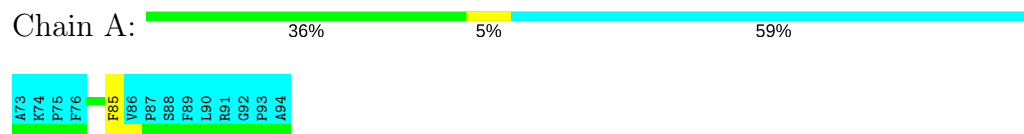


- Molecule 2: Polyadenylate-binding protein 1

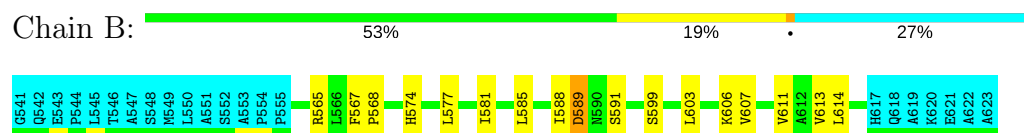


4.2.4 Score per residue for model 4

- Molecule 1: G1 to S phase transition 1

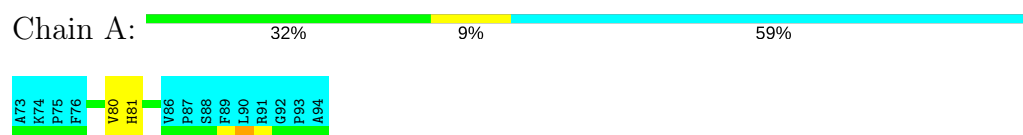


- Molecule 2: Polyadenylate-binding protein 1

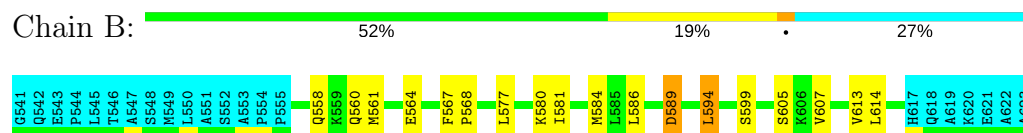


4.2.5 Score per residue for model 5

- Molecule 1: G1 to S phase transition 1

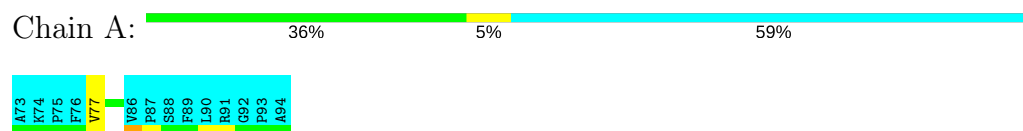


- Molecule 2: Polyadenylate-binding protein 1

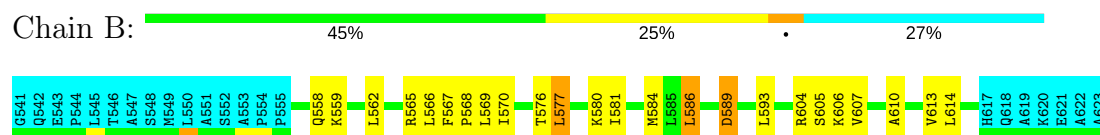


4.2.6 Score per residue for model 6

- Molecule 1: G1 to S phase transition 1

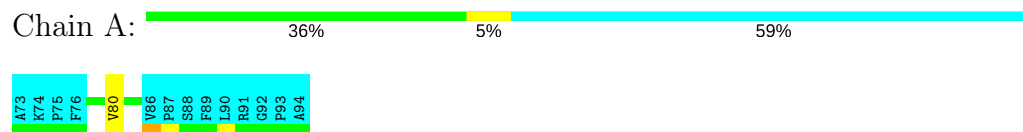


- Molecule 2: Polyadenylate-binding protein 1

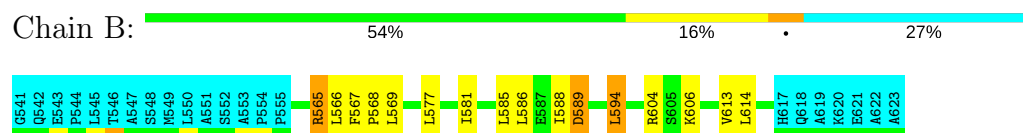


4.2.7 Score per residue for model 7

- Molecule 1: G1 to S phase transition 1

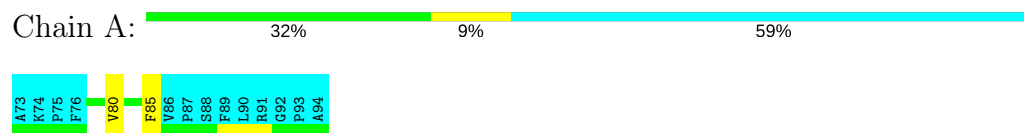


- Molecule 2: Polyadenylate-binding protein 1

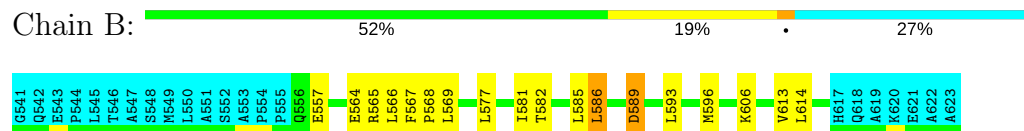


4.2.8 Score per residue for model 8

- Molecule 1: G1 to S phase transition 1

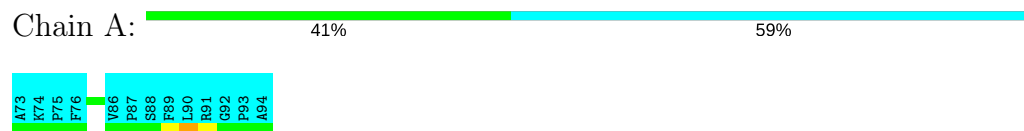


- Molecule 2: Polyadenylate-binding protein 1

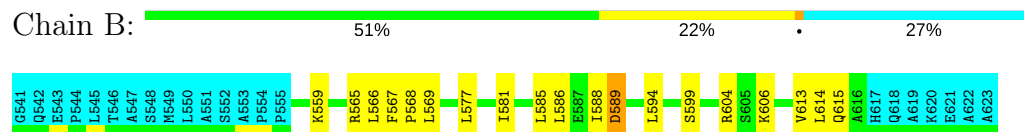


4.2.9 Score per residue for model 9

- Molecule 1: G1 to S phase transition 1

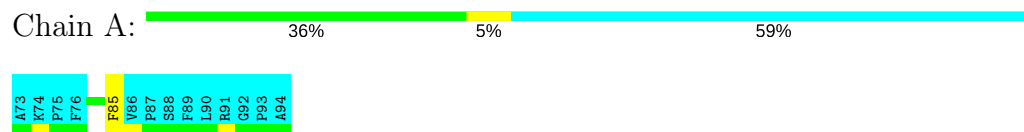


- Molecule 2: Polyadenylate-binding protein 1

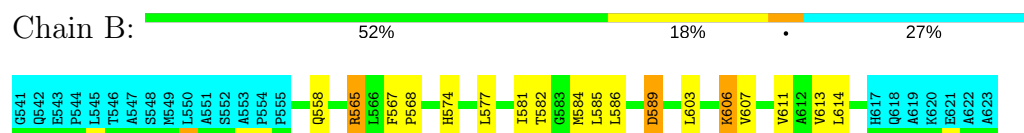


4.2.10 Score per residue for model 10

- Molecule 1: G1 to S phase transition 1

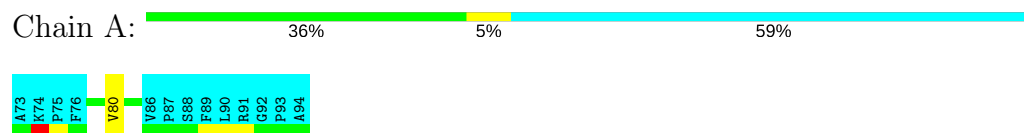


- Molecule 2: Polyadenylate-binding protein 1

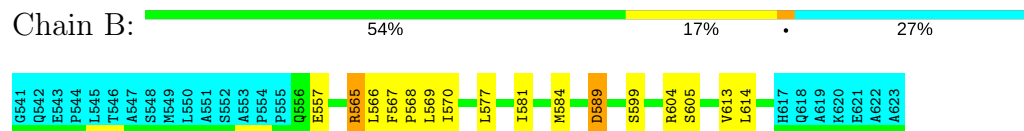


4.2.11 Score per residue for model 11

- Molecule 1: G1 to S phase transition 1

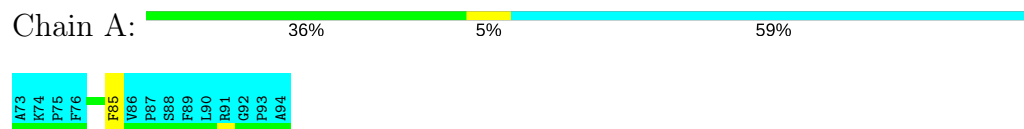


- Molecule 2: Polyadenylate-binding protein 1

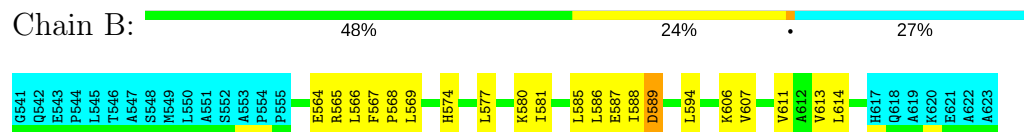


4.2.12 Score per residue for model 12

- Molecule 1: G1 to S phase transition 1

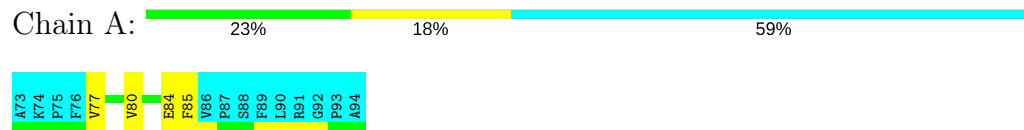


- Molecule 2: Polyadenylate-binding protein 1

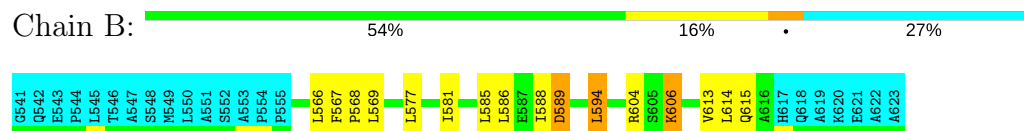


4.2.13 Score per residue for model 13

- Molecule 1: G1 to S phase transition 1

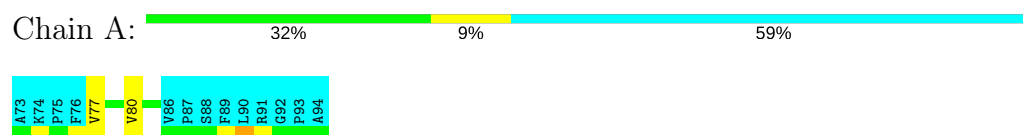


- Molecule 2: Polyadenylate-binding protein 1

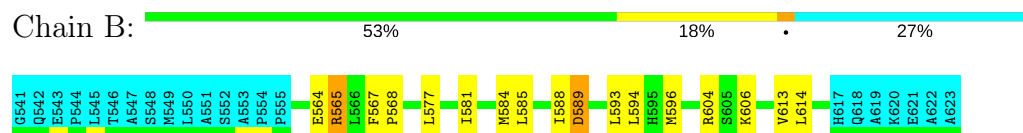


4.2.14 Score per residue for model 14

- Molecule 1: G1 to S phase transition 1

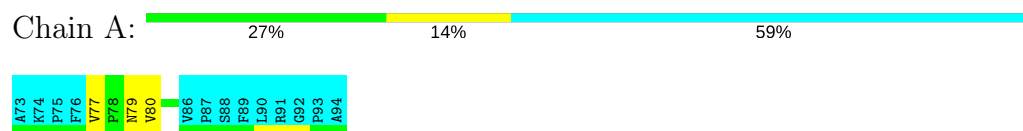


- Molecule 2: Polyadenylate-binding protein 1

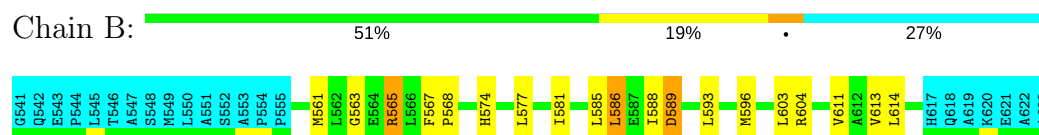


4.2.15 Score per residue for model 15 (medoid)

- Molecule 1: G1 to S phase transition 1

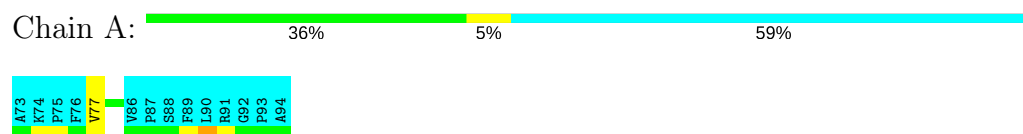


- Molecule 2: Polyadenylate-binding protein 1

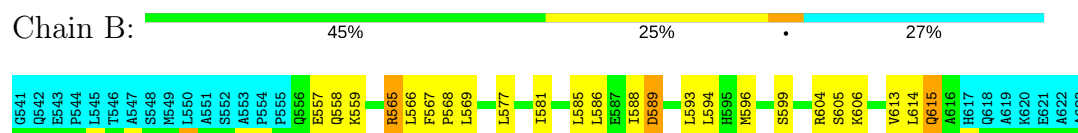


4.2.16 Score per residue for model 16

- Molecule 1: G1 to S phase transition 1

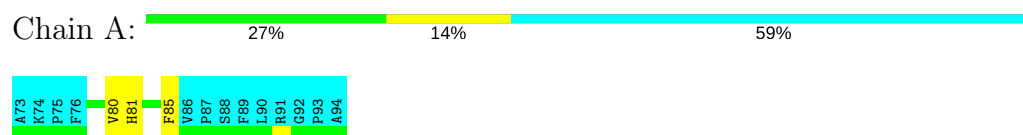


- Molecule 2: Polyadenylate-binding protein 1

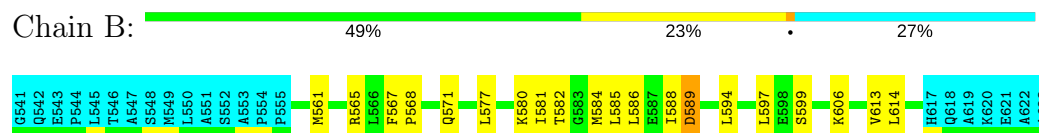


4.2.17 Score per residue for model 17

- Molecule 1: G1 to S phase transition 1

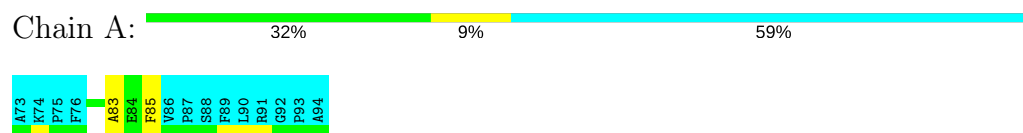


- Molecule 2: Polyadenylate-binding protein 1

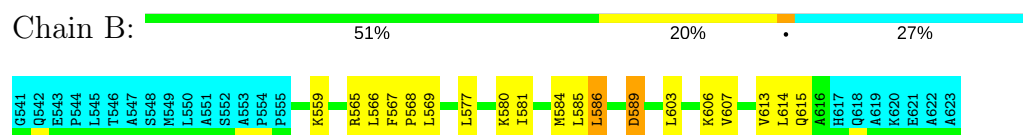


4.2.18 Score per residue for model 18

- Molecule 1: G1 to S phase transition 1

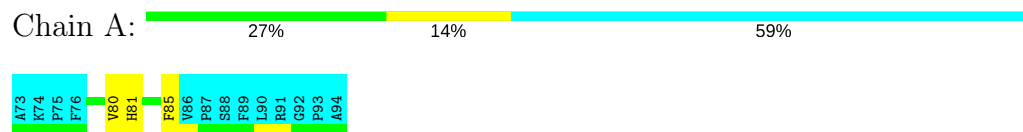


- Molecule 2: Polyadenylate-binding protein 1

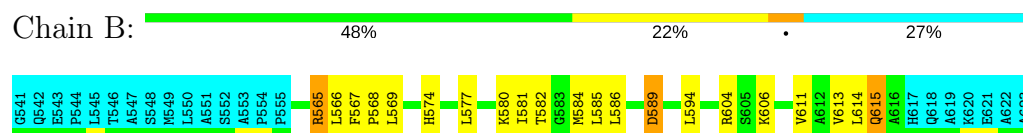


4.2.19 Score per residue for model 19

- Molecule 1: G1 to S phase transition 1



- Molecule 2: Polyadenylate-binding protein 1



4.2.20 Score per residue for model 20

- Molecule 1: G1 to S phase transition 1

Chain A:  36% 5% 59%



- Molecule 2: Polyadenylate-binding protein 1

Chain B:  51% 22% 27%



5 Refinement protocol and experimental data overview

The models were refined using the following method: *TORSION ANGLE DYNAMICS*.

Of the 100 calculated structures, 20 were deposited, based on the following criterion: *TARGET FUNCTION*.

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

Software name	Classification	Version
CYANA	structure solution	2.1
CYANA	refinement	2.1

No chemical shift data was provided. No validations of the models with respect to experimental NMR restraints is performed at this time.

6 Model quality

6.1 Standard geometry

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

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	69	63	63	2±1
2	B	473	490	490	10±2
All	All	10840	11060	11060	207

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:80:VAL:HG21	2:B:614:LEU:HD23	0.87	1.44	14	10
2:B:582:THR:HG22	2:B:586:LEU:HD23	0.68	1.63	10	3
1:A:85:PHE:HB2	2:B:586:LEU:HD21	0.64	1.70	19	7
2:B:603:LEU:O	2:B:607:VAL:HG23	0.60	1.96	20	5
2:B:581:ILE:HD12	2:B:607:VAL:HG13	0.59	1.73	6	4
2:B:574:HIS:CD2	2:B:611:VAL:HG21	0.58	2.32	15	2
2:B:593:LEU:HD23	2:B:596:MET:CE	0.57	2.29	8	2
2:B:574:HIS:CE1	2:B:611:VAL:HG21	0.57	2.34	19	3
1:A:80:VAL:HG13	1:A:81:HIS:CD2	0.57	2.34	5	1
2:B:581:ILE:HG22	2:B:610:ALA:CB	0.55	2.31	20	2
2:B:585:LEU:HD13	2:B:588:ILE:CD1	0.55	2.32	14	8
2:B:593:LEU:HD23	2:B:596:MET:HE2	0.55	1.78	20	2
2:B:594:LEU:O	2:B:594:LEU:HD23	0.55	2.01	16	1

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Atom-1	Atom-2	Clash(Å)	Distance(Å)	Models	
				Worst	Total
2:B:581:ILE:O	2:B:585:LEU:HD23	0.54	2.02	18	15
2:B:596:MET:CE	2:B:603:LEU:HD12	0.54	2.33	15	1
2:B:593:LEU:HD23	2:B:596:MET:SD	0.53	2.43	15	3
2:B:566:LEU:HA	2:B:569:LEU:HD12	0.53	1.81	13	13
2:B:585:LEU:HD13	2:B:588:ILE:HD11	0.51	1.80	15	8
1:A:80:VAL:HG13	1:A:81:HIS:ND1	0.51	2.20	17	2
2:B:614:LEU:HD22	2:B:614:LEU:O	0.51	2.05	15	7
1:A:85:PHE:CB	2:B:586:LEU:HD21	0.50	2.35	19	3
2:B:614:LEU:HD13	2:B:614:LEU:C	0.50	2.27	4	7
2:B:594:LEU:O	2:B:594:LEU:HD13	0.50	2.06	7	1
1:A:80:VAL:HG21	2:B:614:LEU:CD2	0.50	2.29	14	1
2:B:585:LEU:HD11	2:B:606:LYS:HB3	0.49	1.83	10	8
2:B:582:THR:O	2:B:586:LEU:HD23	0.49	2.06	1	3
1:A:77:VAL:HG22	1:A:77:VAL:O	0.49	2.08	1	3
2:B:567:PHE:N	2:B:568:PRO:HD2	0.48	2.24	20	20
2:B:614:LEU:HD13	2:B:615:GLN:N	0.48	2.23	16	5
2:B:614:LEU:C	2:B:614:LEU:HD13	0.46	2.31	15	7
2:B:570:ILE:HG21	2:B:581:ILE:HD11	0.46	1.88	20	5
2:B:574:HIS:NE2	2:B:611:VAL:HG21	0.46	2.25	12	1
2:B:585:LEU:O	2:B:593:LEU:HD21	0.45	2.11	15	2
2:B:594:LEU:HD13	2:B:594:LEU:O	0.45	2.11	13	1
2:B:614:LEU:O	2:B:614:LEU:HD22	0.45	2.12	9	5
1:A:80:VAL:HG22	1:A:81:HIS:N	0.43	2.29	19	2
2:B:581:ILE:HG22	2:B:610:ALA:HB1	0.43	1.90	20	1
2:B:594:LEU:HD23	2:B:594:LEU:O	0.43	2.14	12	1
2:B:588:ILE:HD13	2:B:606:LYS:NZ	0.43	2.29	17	1
1:A:77:VAL:HG13	1:A:77:VAL:O	0.42	2.15	13	3
1:A:77:VAL:O	1:A:77:VAL:HG13	0.42	2.15	2	3
2:B:576:THR:HB	2:B:577:LEU:HD12	0.42	1.89	6	2
2:B:562:LEU:HD13	2:B:593:LEU:HD13	0.42	1.91	6	1
2:B:567:PHE:CB	2:B:568:PRO:CD	0.41	2.98	17	17
2:B:607:VAL:O	2:B:611:VAL:HG23	0.41	2.14	12	2
2:B:559:LYS:O	2:B:586:LEU:HD13	0.41	2.16	6	1
1:A:77:VAL:O	1:A:77:VAL:HG22	0.41	2.16	15	1
2:B:563:GLY:HA3	2:B:586:LEU:HD22	0.41	1.92	15	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	9/22 (41%)	6±1 (69±9%)	3±1 (29±9%)	0±0 (1±3%)	21	68
2	B	61/83 (73%)	57±1 (93±2%)	3±1 (5±2%)	1±0 (2±0%)	16	60
All	All	1400/2100 (67%)	1259 (90%)	119 (8%)	22 (2%)	16	60

All 3 unique Ramachandran outliers are listed below. They are sorted by the frequency of occurrence in the ensemble.

Mol	Chain	Res	Type	Models (Total)
2	B	589	ASP	20
1	A	83	ALA	1
1	A	85	PHE	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	7/17 (41%)	7±0 (97±6%)	0±0 (3±6%)	51	90
2	B	53/68 (78%)	45±2 (84±4%)	8±2 (16±4%)	6	44
All	All	1200/1700 (71%)	1029 (86%)	171 (14%)	7	47

All 26 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)
2	B	589	ASP	20
2	B	577	LEU	20
2	B	613	VAL	20
2	B	586	LEU	13

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Mol	Chain	Res	Type	Models (Total)
2	B	565	ARG	11
2	B	604	ARG	10
2	B	606	LYS	10
2	B	584	MET	10
2	B	594	LEU	8
2	B	580	LYS	6
2	B	599	SER	6
2	B	564	GLU	5
2	B	605	SER	5
2	B	561	MET	5
2	B	559	LYS	3
2	B	615	GLN	3
2	B	557	GLU	3
2	B	571	GLN	3
1	A	85	PHE	2
2	B	614	LEU	2
2	B	587	GLU	1
2	B	558	GLN	1
2	B	591	SER	1
1	A	84	GLU	1
1	A	79	ASN	1
2	B	560	GLN	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 carbohydrates 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

No chemical shift data were provided