

# Full wwPDB X-ray Structure Validation Report (i)

Dec 14, 2023 – 05:12 am GMT

PDB ID		
Title	:	Structure of the Cys65Asp mutant of phenylacetone monooxygenase: oxidised state
Authors	:	Brondani, P.B.; Dudek, H.M.; Martinoli, C.; Mattevi, A.; Fraaije, M.W.
Deposited on	:	2014-04-24
Resolution	:	1.81  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:

MolProbity	:	FAILED
Xtriage (Phenix)	:	1.13
$\mathrm{EDS}$	:	FAILED
Percentile statistics	:	20191225.v01 (using entries in the PDB archive December 25th 2019)
Ideal geometry (proteins)	:	Engh & Huber $(2001)$
Ideal geometry (DNA, RNA)	:	Parkinson et al. (1996)
Validation Pipeline (wwPDB-VP)	:	2.36

## 1 Overall quality at a glance (i)

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

The reported resolution of this entry is 1.81 Å.

There are no overall percentile quality scores available for this entry.

ENTRY-COMPOSITION INFOmissingINFO

SEQUENCE-PLOTS INFOmissingINFO



## 2 Data and refinement statistics (i)

Property	Value	Source	
Space group	P 32 2 1	Depositor	
Cell constants	107.74Å 107.74Å 106.80Å	Depositor	
a, b, c, $\alpha$ , $\beta$ , $\gamma$	$90.00^{\circ}$ $90.00^{\circ}$ $120.00^{\circ}$	Depositor	
Resolution (Å)	93.30 - 1.81	Depositor	
% Data completeness	99.8 (93.30-1.81)	Depositor	
(in resolution range)		-	
R <sub>merge</sub>	0.06	Depositor	
R <sub>sym</sub>	(Not available)	Depositor	
$< I/\sigma(I) > 1$	$2.37 (at 1.82 \text{\AA})$	Xtriage	
Refinement program	REFMAC 5.8.0049	Depositor	
$R, R_{free}$	0.193 , $0.230$	Depositor	
Wilson B-factor ( $Å^2$ )	32.3	Xtriage	
Anisotropy	0.037	Xtriage	
L-test for twinning <sup>2</sup>	$< L >=0.49, < L^2>=0.32$	Xtriage	
Estimated twinning fraction	0.025 for -h,-k,l	Xtriage	
Total number of atoms	4590	wwPDB-VP	
Average B, all atoms $(Å^2)$	36.0	wwPDB-VP	

EDS failed to run properly - this section is therefore incomplete.

Xtriage's analysis on translational NCS is as follows: The largest off-origin peak in the Patterson function is 3.51% of the height of the origin peak. No significant pseudotranslation is detected.

<sup>&</sup>lt;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.



<sup>&</sup>lt;sup>1</sup>Intensities estimated from amplitudes.

## 3 Model quality (i)

### 3.1 Standard geometry (i)

MolProbity failed to run properly - this section is therefore empty.

#### 3.2 Too-close contacts (i)

MolProbity failed to run properly - this section is therefore empty.

#### 3.3 Torsion angles (i)

#### 3.3.1 Protein backbone (i)

MolProbity failed to run properly - this section is therefore empty.

#### 3.3.2 Protein sidechains (i)

MolProbity failed to run properly - this section is therefore empty.

#### 3.3.3 RNA (i)

MolProbity failed to run properly - this section is therefore empty.

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

validation-pack failed to run properly - this section is therefore empty.

#### 3.5 Carbohydrates (i)

validation-pack failed to run properly - this section is therefore empty.

### 3.6 Ligand geometry (i)

validation-pack failed to run properly - this section is therefore empty.

#### 3.7 Other polymers (i)

validation-pack failed to run properly - this section is therefore empty.



## 3.8 Polymer linkage issues (i)

There are no chain breaks in this entry.



## 4 Fit of model and data (i)

## 4.1 Protein, DNA and RNA chains (i)

EDS failed to run properly - this section is therefore empty.

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

EDS failed to run properly - this section is therefore empty.

## 4.3 Carbohydrates (i)

EDS failed to run properly - this section is therefore empty.

## 4.4 Ligands (i)

EDS failed to run properly - this section is therefore empty.

### 4.5 Other polymers (i)

EDS failed to run properly - this section is therefore empty.

