

Biotransformation of benzo[a]pyrene by *Pannonibacter* sp. JPA3 and the degradation mechanism research through the initially oxidized benzo[a]pyrene-4,5-dihydrodiol to downstream metabolites

Jingnan Jin^{a,*}, Yahui Shi^a, Baozhong Zhang^a, Dongjin Wan^a, Qingye Zhang^b, Ying Li^a

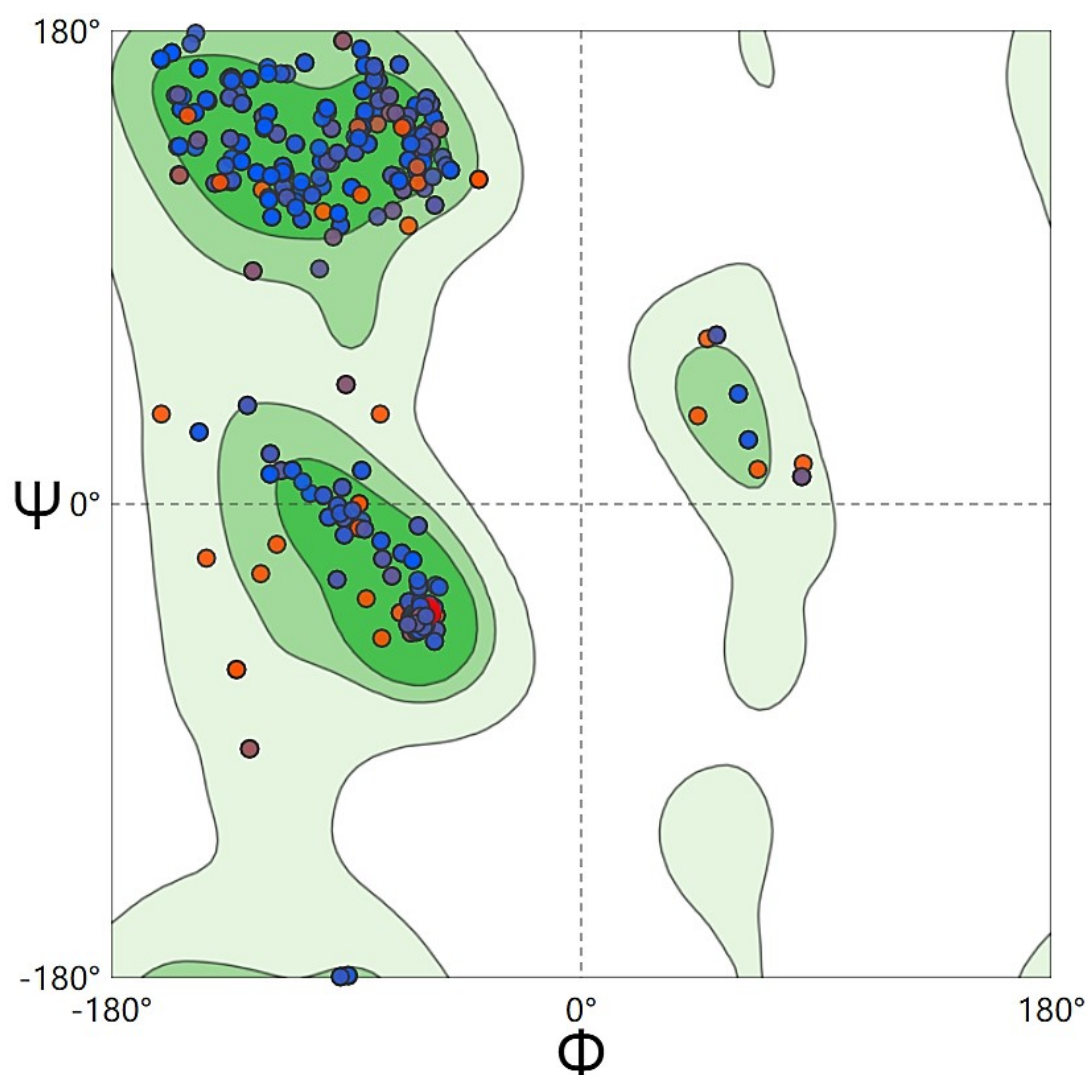


Fig. S1. The Ramachandran Plots of modeling structure of target enzyme in the strain.

Table S1

The constitution of the mineral salt medium.

Elements	Concentration
KH_2PO_4	0.50 g/L
$\text{Na}_2\text{HPO}_4 \cdot 12\text{H}_2\text{O}$	1.26 g/L
$(\text{NH}_4)_2\text{SO}_4$	3.00 g/L
$\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$	0.54 g/L
$\text{MnSO}_4 \cdot \text{H}_2\text{O}$	0.15 mg/L
$\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$	0.50 mg/L
$\text{ZnSO}_4 \cdot 7\text{H}_2\text{O}$	0.24 mg/L
$\text{CoCl}_2 \cdot 6\text{H}_2\text{O}$	3.66 mg/L
$\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$	0.30 mg/L

Table S2

Amplification system of of 16S rDNA

Elements	Concentration
2×Taq MasterMix	12.50 μL
each primer	1.00 μL
genomic DNA	0.50 μL
ddH ₂ O	10.00 μL

Table S3

The amplification program of 16S rDNA.

Temperature	Time
94 °C	1 min
94 °C	1 min
30 times 54 °C	1 min
72 °C	1 min
72 °C	10 min

Table S4

Parameters of HP-5 chromatogram column.

Elements	Concentration
Carrier gas	40 cm s^{-1}
Column temperature	60 °C for 1 min 320 °C for 10 °C min^{-1}
Electron ionization energy	70 eV
Ion source temperature	230 °C

Table S5

Procedure of PCR amplification for dioxygenase gene.

Temperature	Time
95 °C	1 min
30 times	95 °C 55 °C 72 °C
72 °C	10 min