Deciphering the Role of V88L Substitution in NDM-24 metallo-β-lactamase

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Running Title: V88L substitution in NDM-24

Table S1. Oligonucleotides used in this study.

Primers	Sequence (5′- 3′)[a]
NDM-F	CGGGATCCATGGAATTGCCCAATATTATG
NDM-R	AACTGCAGTCAGCGCAGCTTGTCGGCCAT
BamHI-**TEV**-NDM-F	ATGGATCC**GAAAACCTGTATTTCCAAGGC**CAGCAAATGGAAACTGGCGAC
XhoI-NDM-R	ATCTCGAGTCAGCGCAGCTTGTCGGCCATG

[a]The sequences of restriction site were shown with underline and TEV protease recognition sequence was in bold.

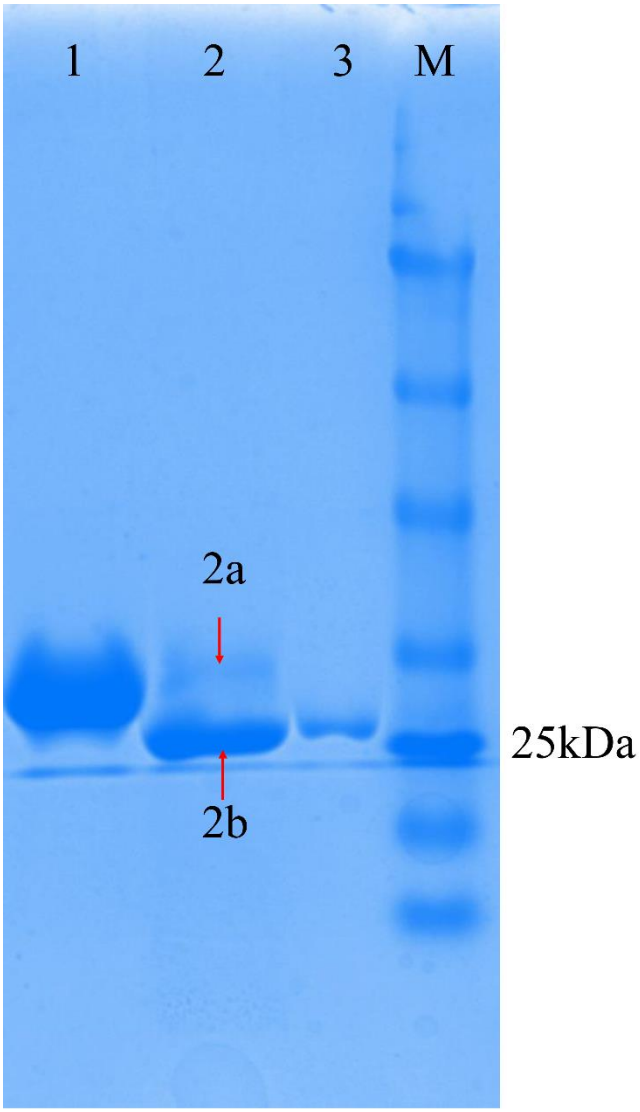

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Figure S1. SDS-PAGE of NDM-24. Lane 1: NDM-24 containing His Tags;

Lane 2: NDM was cleaved by using Turbo tobacco etch virus (TEV) protease to remove His Tags (Accelagen, San Diego, CA, USA): tagged protein (2a) and untagged protein (2b); Lane 3: untagged protein; Lane M: Marker

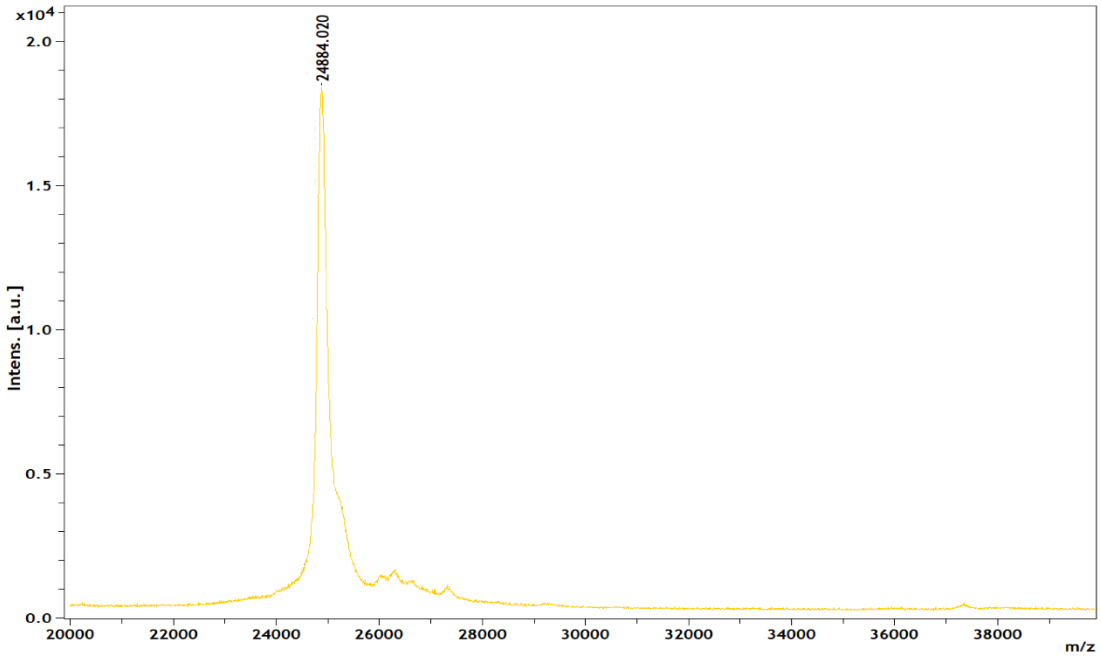

Figure S2. Molecular mass spectrometry of NDM-24 estimated by MALDI-TOF.

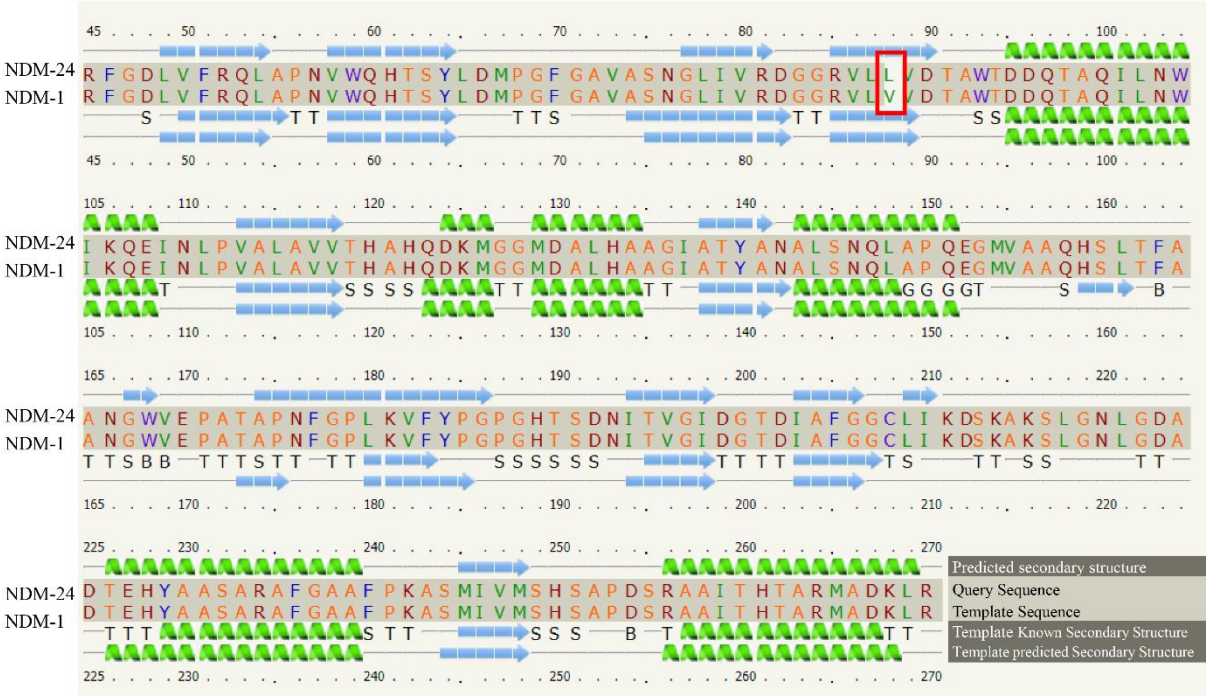

Figure S3. Predicted secondary structure of NDM-24.