Supplementary Information

STRUCTURE AND FUNCTIONAL ANALYSIS OF LPTC, A CONSERVED MEMBRANE PROTEIN INVOLVED IN THE LIPOPOLYSACCHARIDE EXPORT PATHWAY IN ESCHERICHIA COLI

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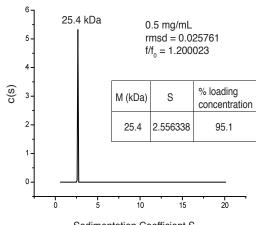
Legend for Supplemental Figures

<u>Fig. S1.</u> Determination of the molecular weight and composition of purified His₆-LptC₍₂₄₋₁₉₁₎. The molecular weight and subunit composition of purified His₆-LptC₍₂₄₋₁₉₁₎ was determined by analytical ultracentrifugation. Purified His₆-LptC₍₂₄₋₁₉₁₎ at 0.5 mg/mL in buffer C was centrifuged in a Beckman Coulter Optima XL-1 analytical ultracentrifuge at 45,000 rpm for 16 hr at 16 °C. A_{280 nm} measurements were carried out 5 min intervals during the ultracentrifugation.

Fig. S2. ClustalW alignments of LptA and LptC. *Panels A* and *B* provide primary and secondary structure alignments of LptA and LptC, respectively. Residue numbering corresponds to LptA from *E. coli* K-12 (without gaps). *Panels C* and *D* provide the primary structure alignments of LBP-14, a synthetic fragment derived from the LPS binding protein, to LptA and LptC, respectively. Alignments were performed with ClustalW (http://www.ebi.ac.uk/Tools/clustalw2/index.html) (1). Residues with high sequence identity are colored, and non-conserved residues are shown as black letters. The residues within the loops regions are indicated.

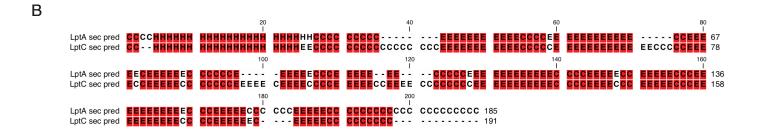
Reference:

1. Thompson, J., Higgins, D., and Gibson, T. (1994) Nucleic Acids Res 22, 4673-4680

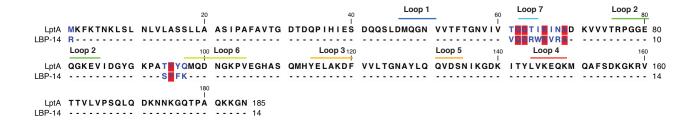


Sedimentation Coefficient S





С



D

			20 I	40 I		60 I	L	.oop 1 80
LptC	MSKARRWVII	VLSLAVLVN	I GINMAEKDDT	AQVVVNNNDP	TYKSEHTDTL	VYNPEGALSY	RLIAQHVEYY	SDQAVSWFTQ 80
LBP-14								
	Lo	op 7	100 Loop 2	120	Loop 6	140	Loop 3	Loop 5 160
			D KAKLTNDRML S					TLYGTTFNSS 159
LptC		KNAELIEKV	R TSYEIQNKQT					