Replacement of Lipopolysaccharide with Free Lipid A Molecules in Escherichia coli Mutants Lacking All Core Sugars

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Supporting Information

Supporting Figure 1. ESI/MS demonstrating the accumulation of lipid IV_A in *kdtA* mutant CMR100. Exponentially-growing cells in 50 mL LB broth supplemented with 1 mM IPTG and 100 μg/mL ampicillin at 30 °C were harvested in late log phase. The total lipids were extracted with a two-phase neutral Bligh-Dyer system (46), re-dissolved in chloroform/methanol/piperidine (2:1:0.03, v/v/v), and immediately analyzed in the negative ion mode by direct infusion ESI/MS, using an ABI QSTAR XL quadrupole time-of-flight mass spectrometer. **Panel A.** Major glycerophospholipid ions of the control strain DY330(pWMsbA) between *m/z* 698 and 722 consist mainly of molecular species of PE and PG, as indicated. **Panel B.** The *kdtA* deletion mutant CMR100 contains similar glycerophospholipids, but accumulates additional peaks (*red*), which are interpreted as the [M-2H]²⁻ and [M-3H+Na]²⁻ ions of lipid IV_A. The other regions of these spectra did not show significant differences.

