

TABLE S1. Primers used in this study.

Primer	Sequence (5' to 3')	Modification, for use
T7kan-2-SacI-F	<u>GAGCTCCCTATA</u> GTGAGTCGTATTACCATCA TCGATGAATTGTG	<u>SacI</u> , Km ^r gene amplification
T7kan-2-SacI-R	<u>GAGCTCTGAGGCCGGTCTCC</u> TATAG	<u>SacI</u> , Km ^r gene amplification
KAN-2 FP-1	ACCTACAACAAAGCTCTCATCAACC	Inverse PCR
KAN-2 RP-1	GCAATGTAACATCAGAGATTTGAG	Inverse PCR
3177_F-O	CATTGTCAGCGAATTACC	SO3177 disruption
3177_5-O_SpeI	ATG <u>ACTAGTCGGACATTA</u> ATCTATCTCCC	<u>SpeI</u> , SO3177 disruption
3177_5-I	CTGATCGGTGCAAAAGTTCTGACTGTTACC ACAAAG	Linker sequence, SO3177 disruption
3177_3-I	AACTTTGCACCGATCAGAACCGGATGATT AAAGGT	Linker sequence, SO3177 disruption
3177_3-O-SpeI	ACG <u>ACTAGTT</u> CATTATTGCAC	<u>SpeI</u> , SO3177 disruption
3177_R-O	TTCGCCAAGGAGGATAAC	SO3177 disruption
SO3177-F-HindIII	ACGGA <u>AGCTT</u> GGAAAGAATTAAAGTTTTG	<u>HindIII</u> , SO3177 complementation
SO3177-R-XbaI	ACCG <u>TCTAGAGTT</u> ATTAAACACCTTTAAC	<u>XbaI</u> , SO3177 complementation

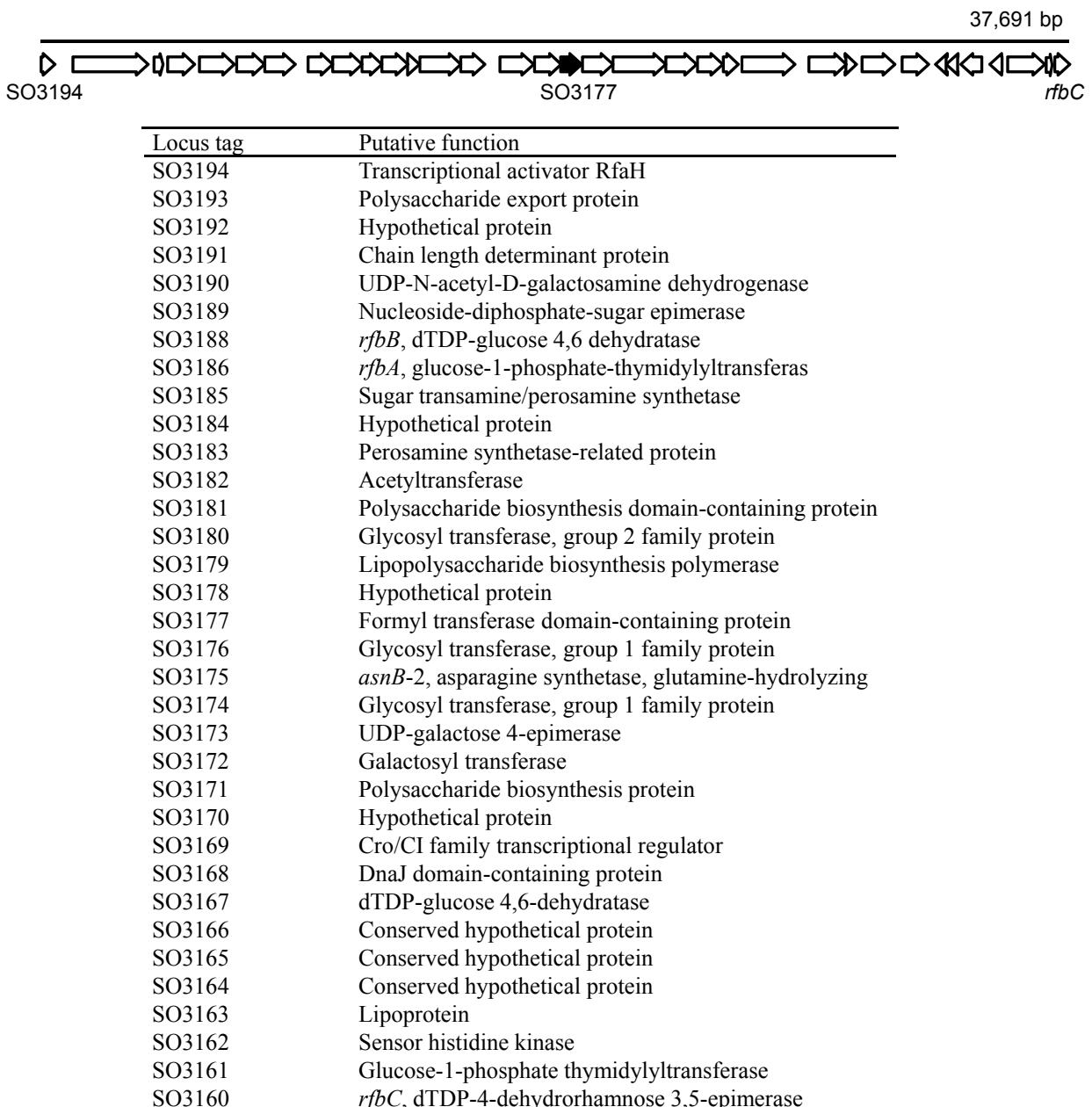


FIG. S1. Putative polysaccharide biosynthesis gene cluster, including SO3177, in the genome of *S. oneidensis* MR-1. ORFs are indicated in order of locus tag from the left.



FIG. S2. Silver-stained SDS-PAGE gel, showing sizes of LPSs extracted from the wild-type MR-1 (lane 1), 4A (lane 2), and Δ SO3177 (lane 3). The arrow indicates the positions of LPSs from *S. oneidensis*.