

Supplementary Material

Transcriptomic Response of *L. Monocytogenes* to Co-Culture with *S. Cerevisiae*

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Table S1. Primer sequences, amplicon sizes, and PCR conditions used for the gene transcription assay.

Genes	Primer Sequence	Concentration (μM)	Amplicon Size (bp)	PCR Efficiency
IGS	IGS_f: GGCCTATAGCTCAGCTGGTTA	1.2	135	2.03
	IGS_r: GCTGAGCTAAGGCCCGTAAA	1.2	-	-
<i>rpob</i>	<i>rpob</i> _f: CCGCGATGCGAAAACAAT	0.9	69	2.04
	<i>rpob</i> _r: CCWACAGAGATACGGTTATCRAATGC	0.9	-	-
16S	16S_f: GATGCATAGCCGACCTGAGA	0.9	114	2.05
	16S_r: CTCCGTCAGACTTTCGTCCA	0.9	-	-
<i>prfA</i>	<i>prfA</i> _f: CTATTTGCGGTCAACTTTTAATCCT	0.9	100	2.09
	<i>prfA</i> _r: CCTAACTCCTGCATTGTAAATTATCC	0.9	-	-
<i>sigB</i>	<i>sigB</i> _f: CCAAGAAAATGGCGATCAAGAC	1.2	166	2.13
	<i>sigB</i> _r: CGTTGCATCATATCTTCTAATAGCT	1.2	-	-
<i>hly</i>	<i>hly</i> _f: TACATTAGTGGAAGATGG	1.2	153	1.98
	<i>hly</i> _r: ACATTCAAGCTATTATTACA	1.2	-	-
<i>plcA</i>	<i>plcA</i> _f: CTAGAAGCAGGAATACGGTACA	1.2	115	1.94
	<i>plcA</i> _r: ATTGAGTAATCGTTTCTAAT	1.2	-	-
<i>plcB</i>	<i>plcB</i> _f: CAGGCTACCACTGTGCATATGAA	0.9	72	2.00
	<i>plcB</i> _r: CCATGTCTTCYGTGCTTGATAATTG	0.9	-	-
<i>inlA</i>	<i>inlA</i> _f: AATGCTCAGGCAGCTACAMTTACA	0.9	114	2.12
	<i>inlA</i> _r: CGTGTCTGTTACRTTCGTTTTTCC	0.9	-	-
<i>inlB</i>	<i>inlB</i> _f: AAGCAMGATTTTCATGGGAGAGT	0.9	78	2.04
	<i>inlB</i> _r: TTACCGTTCCATCAACATCATAACTT	0.9	-	-
<i>inlC</i>	<i>inlC</i> _f: ACTGGTCAGAAATGTGTGAATGA	0.9	80	2.06
	<i>inlC</i> _r: CCATCTGGGTCTTTGACAGT	0.9	-	-
<i>inlJ</i>	<i>inlJ</i> _f: TCGTAAATGCTCACATCCAAG	0.9	81	2.03
	<i>inlJ</i> _r: TTGCCCTTCAGCATCCAAGT	0.9	-	-

Thermocycling conditions: initial denaturation at 95 °C for 20 s and then 40x (95 °C for 10 s, 60 °C for 30 s, 72 °C for 30 s). Melting curve analysis: 95 °C for 15 s and then 60 °C for 1 min and raised to 95 °C at 0.3 °C/s.