

Supplementary Information

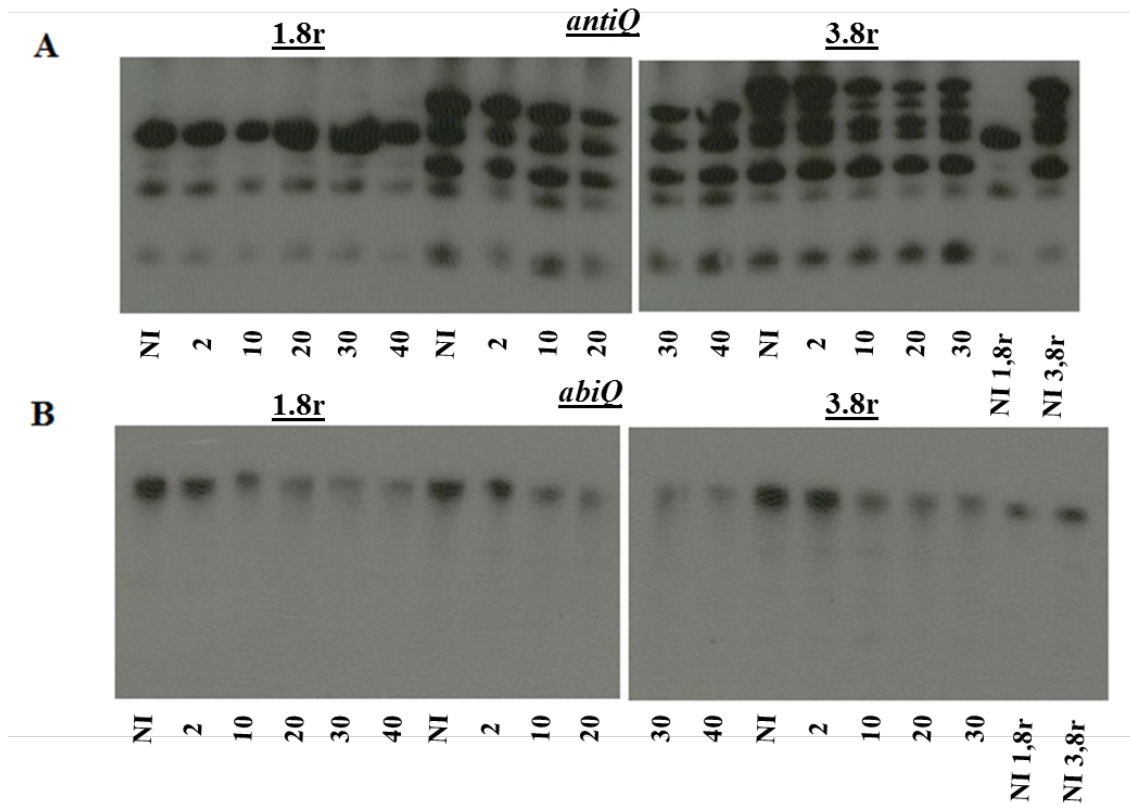
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Table S1. Primers used in this study.

Function	Primers	Sequence (5'-3') ^a	T _m (°C)	Notes
5'RACE PCR				
	JS2	ccatcttttcatgagcagctt	53,9	cDNA of AbiQ _{op} transcripts
	AbiQFwd	ttttacgaatgacgttaagattc	49,3	Control+, cDNA
	AbiQRev	CAGAATTCGAATTCcgaagaggattattatattgcc	50,9	Control+/-, cDNA
	PolyC	GACTCGAGTCGACATCGAccccccccccccccc	73,8	PCR RACE-AbiQ
	M13Fwd	gtaaacgacggccagt	52,6	MCS of pBS-KS
	M13Rev	caggaaacagctatgac	47	MCS of pBS-KS
Revertant (<i>antiQ</i> repeats)				
	Mut1.8Fwd	attactttatacctttgcaagctacaaaaaggggaa	59,6	Revertant 1.8R
	Mut1.8Rev	ttcccctttttagcttgacaaggtataaaagta	59,6	Revertant 1.8R
	Mut3.8Fwd	attactttatacctttgcaagctacaaaaaggggaa	61	Revertant 3.8R
	Mut3.8Rev	ggtcccctttttagcttgacaaggtataaaagta	61	Revertant 3.8R
Point mutant (<i>antiQ</i>)				
	antiQ(A13C)Fwd	gatattataattgtaaggaatccaagccatcgttggaaatgataaaatggagtatcc	63,5	Mutant A13C
	antiQ(A13C)Rev	ggatactccaattttatcaattccaacgatggcttggatattccttacaattataatc	63,5	Mutant A13C
	antiQ(A24C)Fwd	gatattataattgtaaggaatccaagccatagttggaaatgataaaatggagtatcc	63,5	Mutant A24C
	antiQ(A24C)Rev	ggatactccaattttagcaattccaactatggcttggatattccttacaattataatc	63,5	Mutant A24C
	antiQ(T25C)Fwd	gatattataattgtaaggaatccaagccatagttggaaatgataaaatggagtatcc	63,5	Mutant T25C
	antiQ(T25C)Rev	ggatactccaattttgcaattccaactatggcttggatattccttacaattataatc	63,5	Mutant T25C
	antiQ(A26C)Fwd	gatattataattgtaaggaatccaagccatagttggaaatgataaaatggagtatcc	65	Mutant A26C
	antiQ(A26C)Rev	ccaactatggcttggatactccaattgatcaattccaactatggcttggatattccttac	65	Mutant A26C
	antiQ(A28C)Fwd	gatattataattgtaaggaatccaagccatagttggaaatgataaaatggagtatcc	65	Mutant A28C
	antiQ(A28C)Rev	ccaactatggcttggatactccaattgatcaattccaactatggcttggatattccttac	65	Mutant A28C
	antiQ(G32A)Fwd	gatattataattgtaaggaatccaagccatagttggaaatgataaaatggagtatcc	65	Mutant G32A
	antiQ(G32A)Rev	ccaactatggcttggatactccaattgatcaattccaactatggcttggatattccttaca	65	Mutant G32A
Vector (MCS)				
	pNZ-F	aatgtcactaacctgccccg	57,4	MCS of pNZ123
	pNZ-R	cattgaacatgctgaagagc	52,1	MCS of pNZ123

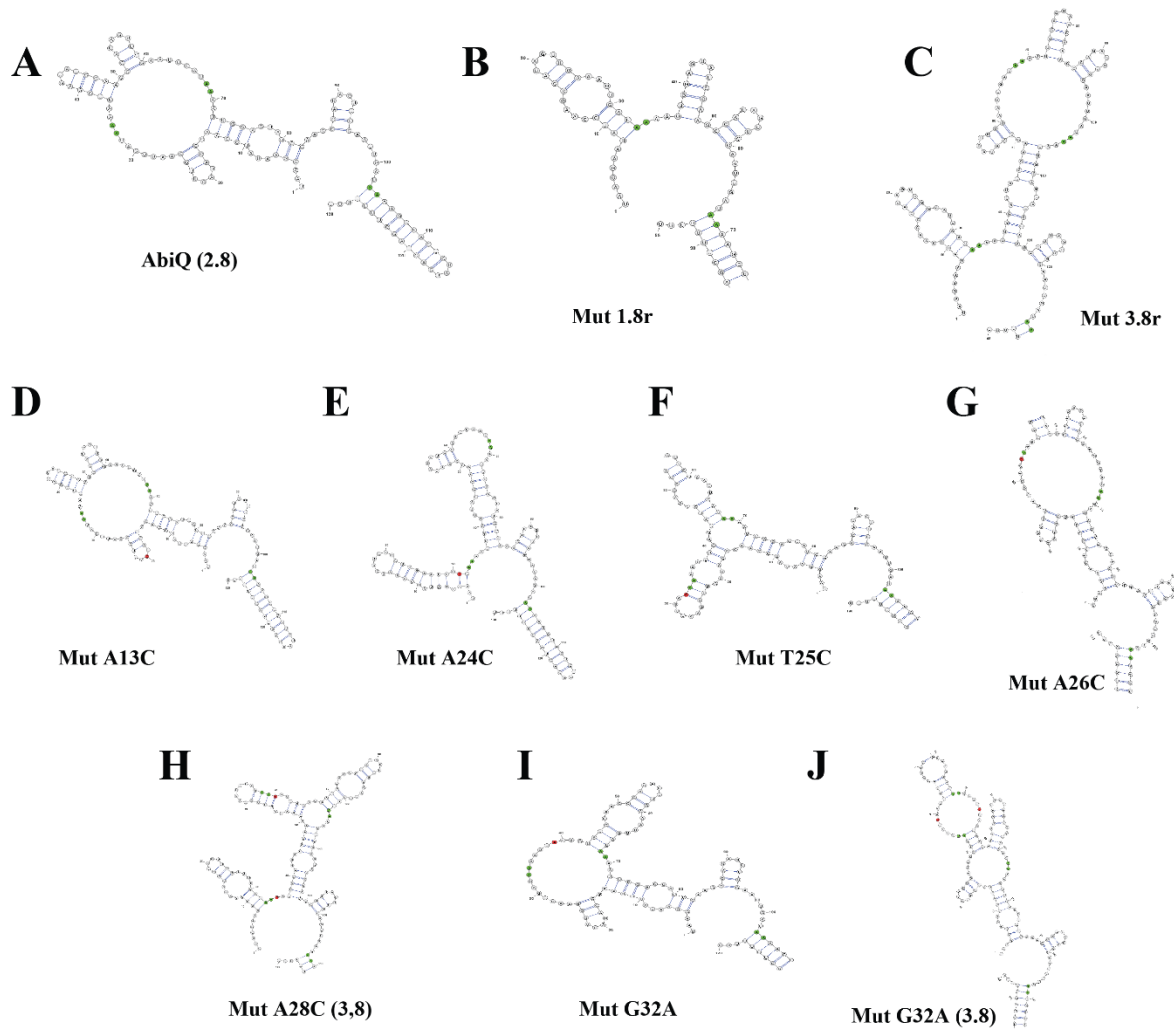
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^a Capital letters, 5' flanking region. Dotted underline, recognition site of restriction enzyme. Solid underline, first repetition of *antiQ* (35nt). Bold letters, mutated nucleotides.



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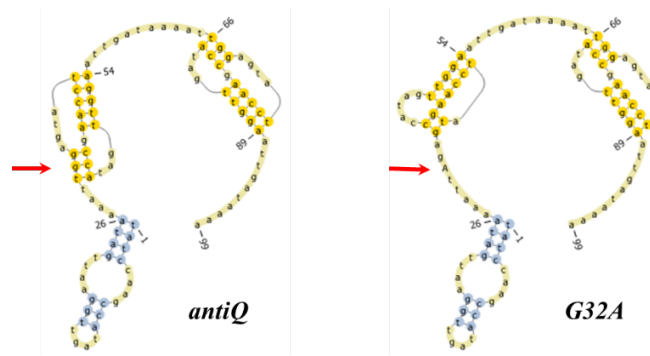
8 **Figure S1.** Northern blot experiment targeting *antiQ* (Panel A) and *abiQ* (Panel B) transcripts
 9 from *L. lactis* IL1403 AbiQ wild-type and mutants (1.8 and 3.8 repetitions) during phage
 10 infection by P008.



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13 **Figure S2.** Comparison of the secondary structure of *antiQ* non-coding RNA (A), mutants with
 14 variable numbers of repeats (B, C) and point mutations mutants (D-J). RNAfold software (mode:
 15 mfe-partition function) has been used in combination with VARNA visualization software to
 16 characterize all the complete *antiQ* RNA structures (1, 2). Green nucleotides surround the ABIQ
 17 cleavage site and the specifically mutated nucleotides are shown in red.



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20 **Figure S3.** Comparison of pseudoknot structure of *antiQ* non-coding RNA (left) and the G32A
21 mutants (right). Pseudoknot structure analysis was performed using pKnotsRG software and the
22 visualization software Pseudoviewer 3.0 (3-5). To simplify analysis, only the nucleotides of the
23 repeats (99 nt) were used as the sequence and uracil was replaced by thymine.

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References

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2. **Darty K, Denise A, Ponty Y.** 2009. VARNA: Interactive drawing and editing of the RNA secondary structure. *Bioinformatics* **25**:1974-1975.
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4. **Byun Y, Han K.** 2006. PseudoViewer: web application and web service for visualizing RNA pseudoknots and secondary structures. *Nucleic Acids Res.* **34**:W416-422.
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