

A. CIPer	/5Phos-----AS-----U2-----R-----U1-----ES-----3'
CSP-CIPer	GTAGTATCTACCACAGTAACAAA
- <i>gyrA</i>	<i>TGTGCTGATAGACGGACAGGGCAA C</i>
+ <i>parC</i>	CCAAATCTGCCGACCACGC <i>CGC</i>
B. PCR	5' -3'
B-F-NP	/5Bio/ACACATTACTCACCTATT
R-NP	TAGAAGTTAGTGTAGGTATG
GP5+	TTTGTACTGTGGTAGATACTAC
GP6+	5/Bio/GAAAAATAAACTGTAAATCATATT
GYRA2-1	TAACGTGAATGCCGCTACAA
GYRA2-2	/5Bio/CCCTGTCCGTCTATCAGCACA
PARC-1	GCGTGGTCGGCGAGATTTGG
PARC-2	/5Bio/GCGAACCGAAGTTGCCGATGC
C. Sequencing	5' -3'
GP5+	TTTGTACTGTGGTAGATACTAC
MSP-16 (1)	ATTATGTGCTGCCATATCTACTT
MSP-31 (1)	GTGCTGCAATTGCAAACAGT
MSP-59 (1)	CTACTACTTCTCTATTCTAA
MSP-39 (1)	CTATAGAGTCTTCATACCT
MSP-18 (2)	ACAGTCTCCTGTACCTGGG
MSP-33 (2)	TGAGAATTTAAAGAATATATAAGACA
MSP-52 (2)	TATGTGCTGAGGTAAAAAG
MSP-56 (2)	TACTAACATGACTATTAGTACT
MSP-45 (3)	CTCTACACAAAATCCTGTGC
MSP-35 (3)	GTTCTGCTGTGTCTTCTAGT
MSP-58 (3)	ACTGAAGTAACTAAGGAAGG
MSP-51 (3)	ACTATTAGCACTGCCACTGC
GYRA2-4	AGCGAAATTTGCCCATACG
PARC-3	ATCCGCACGGCGACAGTTCC

Figure S7. Oligonucleotides used in the present study. Bold italic marked sequences in the anchor sites of CIPers -*gyrA* and +*parC* were added to the originally described primers to obtain higher annealing temperature values for the anchor site. The number in parenthesis following the multiple sequencing primers (MSPs) denotes which primer pool the MSP belongs to.