

Steven J Hancock, Minh-Duy Phan, Kate M Peters, Brian M Forde, Teik Min Chong, Wai-Fong Yin, Kok-Gan Chan, David L Paterson, Timothy R Walsh, Scott A Beatson, Mark A Schembri. 2017. Identification of IncA/C Plasmid Replication and Maintenance Genes and Development of a Plasmid Multilocus Sequence Typing Scheme. [Antimicrobial Agents and Chemotherapy, 61\(2\), PMID: 27872077](#).

The A/C pMLST scheme uses internal fragments of the following 4 genes:

- *repA*: replication gene
- *parA*: putative partitioning gene
- *parB*: putative partitioning gene
- *053*: putative partitioning gene

Primer	Sequence (5'-3')	Primer information	Amplicon size (bp)
<i>repA</i> -F	AAGAGAACCAAAGACAAAGAC	Amplify <i>repA</i>	982
<i>repA</i> -R	GCTGCTTACGCTTGTTGGA		
<i>parA</i> -F	AAAAGTAATCAGCTTCGCCA	Amplify <i>parA</i>	780
<i>parA</i> -R	TAGCCACCTTCTCTAATAG		
<i>parB</i> -F	TGTCCGAACTTGCTAAAGC	Amplify <i>parB</i>	1128
<i>parB</i> -R	CTGACACAGGCACATGAA		
<i>053</i> -F	AGATCTCACAGGACATGAA	Amplify <i>053</i>	250
<i>053</i> -R	TTCAAGAACGAAGACCTGT		

Amplification of the four loci used in PMLST was performed with KAPA HiFi DNA polymerase (KAPA Biosystems) with cycling program:

95°C for 3min; 25 cycles of 98°C for 20sec, 60°C for 15 sec 72°C for 30 sec and a final extension of 72°C for 3min.