

mecC	ATGAAAAAAATTTATATTAGTGTGCTAGTTCCTTTTACTAATTATGATT-----	48
mecA	ATGAAAAAGATAAAAAAT---TGTTCCACTTATTTTAATAGTTGTAGTTGTCGGGTTTGGT	57
	***** * * * * * * * * * * * * * * *	
mecC	ATAATAACTTGGTTATTCAAAGATGACGATATTGAGAAAACAATTAGTTCTATTGAAAAA	108
mecA	ATATATTTTATGCTTCAAAGATAAAGAAATTAATAATACTATTGATGCAATTGAAGAT	117
	*** ** * * * * * * * * * * * * * * *	
mecC	GGAAACTATAACGAAGTATATAAAAAATAGTTCAGAAAAATCTAACTGGCATATGGAGAA	168
mecA	AAAAATTTCAAACAAGTTTATAAAGATAGCAGTTATATTTCTAAAAGCGATAATGGTGAA	177
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mecC	GAAGAAATTGTAGATAGGAATAAAAAATTTACAAGATTTAAGTGTCAATAACTTAAAA	228
mecA	GTAGAAATGACTGAACGTCGATAAAAAATATATAATAGTTTAGGCGTTAAAGATATAAAC	237
	* * * * * * * * * * * * * * * * * * * *	
mecC	ATTACTAATCATGAAATTAAAAAACTGGAAAAGATAAAAAGCAAGTTGATGTTAAATAT	288
mecA	ATTCAGGATCGTAAAAATAAAAAAGTATCTAAAAATAAAAAACGAGTAGATGCTCAATAT	297
	*** ** * * * * * * * * * * * * * * * *	
mecC	AACATATATACAAAATATGGAACATACGACGTAATACACAATTAACTTTATTTATGAA	348
mecA	AAAATTAACAACTACGCTAACATTGATCGCAACGTTCAATTTAATTTTGTAAAGAA	357
	** * * * * * * * * * * * * * * * *	
mecC	GATAAGCATTGGAAATTAGATTGGAGACCAGACGTAATAGTACCTGGTTTGAAAAATGGA	408
mecA	GATGGTATGTGGAAGTTAGATTGGGATCATAGCGTCATTATTCCAGGAATGCAGAAAGAC	417
	*** ** * * * * * * * * * * * * * * * *	
mecC	CAGAAAATTAATATAGAAACATTAAAAATCAGAGCGAGGCAAAAATAAAGATAGAAATGGT	468
mecA	CAAGCATACATATTGAAAATTTAAATCAGAACGTGGTAAAATTTTAGACCGAAACAAT	477
	** * * * * * * * * * * * * * * * *	
mecC	ATAGAATTAGCTAAAACCTGGAATACATATGAAATCGGTATTGTCCCTAACAAAACACCC	528
mecA	GTGGAATTGGCCAATACAGGAACAGCATATGAGATAGGCATCGTTCCAAAGAATGTATCT	537
	* * * * * * * * * * * * * * * * * * * *	
mecC	AAAGAAAAATATGATGATATTGCTCGTGACTTACAAATTGATACAAAAGCTATAACCAAT	588
mecA	AAAAAGATTATAAAGCAATCGCTAAAGAACTAAGTATTTCTGAAGACTATATCAAACAA	597
	*** * * * * * * * * * * * * * * * *	
mecC	AAAGTTAATCAAAAATGGGTTTCAGCCAGATTCAATTTGTACCAATTA AAAAGATAAATAAA	648
mecA	CAATGGATCAAAATTTGGGTACAAGATGATACCTTCGTTCCACTTAAACCGTTAAAAAA	657
	* * * * * * * * * * * * * * * * * * * *	
mecC	CAAGATGAATATATAGACAAATTAATTAAATCATACAATTTACAAATAAACACTATAAAA	708
mecA	ATGGATGAATATTTAAGTGATTTCGCAAAAAAATTTCATCTTACAACCTAATGAAACAGAA	717
	***** * * * * * * * * * * * * * * *	
mecC	AGCCGTGTTTATCCATTGAACGAAGCAACAGTACACCTTTTAGGTTATGTGGGTCCAATT	768
mecA	AGTCGTAACCTATCCTCTAGGAAAAGCGACTTCACATCTATTAGGTTATGTTGGTCCCATT	777
	** * * * * * * * * * * * * * * * *	
mecC	AATTCTGACGAGTTAAAAAGTAAGCAATTTAGAACTATAGCAAAAATACTGTTATTGGA	828
mecA	AACTCTGAAGAATTAACCAAAAAGAATATAAAGGCTATAAAGATGATGCAGTTATTGGT	837
	** * * * * * * * * * * * * * * * *	
mecC	AAAAAAGGCTTAGAACGCCTCTATGATAACAATTGCAAAACACTGATGGTTTTAAGGTA	888
mecA	AAAAAGGACTCGAAAACTTTACGATAAAAAGCTCCAACATGAAGATGGCTATCGTGTC	897
	***** * * * * * * * * * * * * * * *	
mecC	TCCATTGCAAATACTTATGACAATAAACCTTTAGACACATTATTGGAGAAAAAGGCTGAA	948
mecA	ACAATCGTTGACGATAATAGCAATACAATCGCACATACATTAATAGAGAAAAAGAAAAAA	957
	* * * * * * * * * * * * * * * * * * * *	
mecC	AACGGAAAAGATCTTCATTTAACTATAGATGCTAGAGTACAAGAAAGTATTTATAAACAT	1008
mecA	GATGGCAAGATATTCAACTAACTATTGATGCTAAAGTTCAAAAGAGTATTTATAACAAC	1017
	* * * * * * * * * * * * * * * * * * * *	
mecC	ATGAAAAATGACGATGGATCTGGTACAGCATTACAACCAAAAACCTGGAGAAATTTAGCT	1068
mecA	ATGAAAAATGATTATGGCTCAGGTACTGCTATCCACCCTCAAACAGGTGAATTATTAGCA	1077
	***** * * * * * * * * * * * * * * *	
mecC	TTGGTAAGTACCCCATCGTACGATGTTTATCCATTTCATGAATGGATTAAAGCAATAATGAC	1128
mecA	CTTGTAAGCACACCTTCATATGACGCTCATCCATTTATGTATGGCATGAGTAACGAAGAA	1137
	* * * * * * * * * * * * * * * * * * * *	
mecC	TACCGTAAATTAATAACAATAAAAAAGAGCCTTTGCTCAACAAATTTCAATCACTACA	1188

<i>mecA</i>	TATAATAAATTAACCGAAGATAAAAAAGAACCTCTGCTCAACAAGTTCAGATTACAAC	1197
	** ***** *	
<i>mecC</i>	TCACCAGGTTCAACCCAAAAATATTAACATCTATTATAGCCTTAAAGAAAATAAACTA	1248
<i>mecA</i>	TCACCAGGTTCAACTCAAAAAATATTAACAGCAATGATTGGGTAAATAACAAAACATTA	1257
	***** * * * *	
<i>mecC</i>	GACAAAAATACTAATTTTGATATTTATGGTAAGGGTTGGCAAAAAGATGCATCATGGGGG	1308
<i>mecA</i>	GACGATAAAACAAGTTATAAAATCGATGGTAAGGGTTGGCAAAAAGATAAATCTTGGGGT	1317
	** *	
<i>mecC</i>	AATTATAATATCACAAAGATTAAAGTAGTAGACGGCAATATCGATTTAAAGCAAGCAATA	1368
<i>mecA</i>	GGTTACAACGTTACAAGATATGAAGTGGTAAATGGTAATATCGACTTAAACAAGCAATA	1377
	* *	
<i>mecC</i>	GAATCATCAGACAACATATTTTTCGCCGCATTGCATTAGCATTAGGAGCCAAAAATTT	1428
<i>mecA</i>	GAATCATCAGATAACATTTCTTTGCTAGAGTAGCACTCGAATTAGGCAGTAAGAAATTT	1437
	***** *	
<i>mecC</i>	GAGCAAGGTATGCAAGATTGGGAATCGGTGAAATATCCCGAGTGATTATCCCTTTTAT	1488
<i>mecA</i>	GAAAAAGGCATGAAAAAAGTAGGTGTTGGTGAAGATATACCAAGTGATTATCCATTTTAT	1497
	** *	
<i>mecC</i>	AAAGCACAAATCTCAAAATAGTAATTTAAAAAATGAAATATTATTAGCAGATTCAGGATAT	1548
<i>mecA</i>	AATGCTCAAATTTCAAACAAAATTTAGATAATGAAATATTATTAGCTGATTCAGGTTAC	1557
	* *	
<i>mecC</i>	GGCCAAGGCGAGATACTAGTAAACCCATACAAAATTTTATCAATATACAGTGCTTTAGAA	1608
<i>mecA</i>	GGACAAGGTGAAATACTGATTAACCCAGTACAGATCCTTTCAATCTATAGCGCATAGAA	1617
	* *	
<i>mecC</i>	AATAACGGAATATACAAAATCCTCATGTTTACGTAAAACAAAATCTCAAATATGGAAA	1668
<i>mecA</i>	AATAATGGCAATATTAACGCACCTCACTATTAAAGACACGAAAAACAAAGTTTGAAG	1677
	***** *	
<i>mecC</i>	AAAGATATTATACCTAAAAAAGACATAGATATATTAATAATGGTATGGAACGTGTAGTT	1728
<i>mecA</i>	AAAAATATTATTTCCAAAGAAAATATCAATCTATTAAGTATGATGCAACAAGTCGTA	1737
	** *	
<i>mecC</i>	AATAAAACACATAGGGATGATATATACAAAAATTATGCCGAATTATTGGTAAATCTGGC	1788
<i>mecA</i>	AATAAAACACATAAAGAAGATATTTATAGATCTTATGCAAACCTAATTGGCAAAATCCGGT	1797
	***** *	
<i>mecC</i>	ACAGCAGAATTAATAATGAATCAAGGGGAACTGGAAGACAAATAGGTTGGTTTGTTC	1848
<i>mecA</i>	ACTGCAGAACTCAAATGAAACAAGGAGAACTGGCAGACAAATGGGTGGTTTATATCA	1857
	** *	
<i>mecC</i>	TATAATAAAAAATAATCCTAATATGTTAATGGCGATTAATGTTAAAGACGTTCAAAATAAA	1908
<i>mecA</i>	TATGATAAAGATAATCCAACATGATGATGGCTATTAATGTTAAAGATGTACAAGATAAA	1917
	** *	
<i>mecC</i>	GGGATGGCCAGCTATAATGCTACTATATCTGGAAGGTTTATGATGATTTGTATGATAAT	1968
<i>mecA</i>	GGAAATGGCTAGCTACAATGCCAAAATCTCAGGTAAAGTGTATGATGAGCTATATGAGAAC	1977
	* *	
<i>mecC</i>	GGAAAACTCAATTTGATATAGATCAGTAA	1998
<i>mecA</i>	GGTAATAAAAAATACGATATAGATGAATAA	2007
	** *	

Figure S1. Comparison of *mecC* (*mecA*_{LGA251}) and *mecA* sequences. The nucleotide sequence of *mecA* is from *S. aureus* USA300 (accession no. CP000730) and the nucleotide sequence of *mecC* is from *S. aureus* LGA251 (accession no. NC_017349).

Table 1. Concept related to *S. aureus* molecular typing and virulence factors.

Concept or term	Definition or explanation
Multilocus sequence typing (MLST)	Molecular typing technique that allows characterize isolates of microbial species using the DNA sequences of internal fragments of multiple housekeeping genes. This method gives a combination of several alleles that determine the sequence type (ST) of a microorganism
Clonal complex (CC)	Determined thanks to MLST technique is a cluster of sequence types (STs) in an eBURST diagram in which all STs are linked as Single Locus Variant (SLVs) to at least one other sequence type (ST)
Single Locus Variant (SLVs)	Strains that differ in only one loci
<i>spa</i> -typing	Molecular typing technique that implicates the sequencing of the polymorphic X region of the staphylococcal protein A gene (<i>spa</i>).
SCC <i>mec</i>	Genetic structure that contains the genes that confer resistance to methicillin (<i>mecA/mecC</i>), and that is integrated into the chromosome.
<i>hla</i> , <i>hly</i> , <i>hld</i>	Hemolysin genes: these genes encode hemolysins that are virulence factors that allow the rupture of cell membranes
<i>edinB</i>	Epidermal cell differentiation inhibitor gene that encodes a exotoxin that specifically inhibit host protein RhoA
<i>lukED</i>	Leukotoxin gene that encodes a pore-forming toxin
<i>cap8</i>	Operon that encodes the type 8 capsular polysaccharide that is an essential factor for bacteria protection, especially when it infects a host, conferring anti-phagocytic ability
<i>ica</i>	Intercellular adhesion gene cluster that leads to the biosynthesis of polysaccharide intercellular adhesion (PIA) molecules, being related to biofilm formation
Pyrogenic Toxin Superantigen (PTSAg) genes	Group of exocellular toxins that share biochemical characteristics and that includes: the toxic shock syndrome toxin (TSST) and staphylococcal enterotoxins (SEA, SEB, SEC, SED, SEE, SEG, SEH, SEI, SEJ, SEK, SEL, SEM, SEN, SEO, SEP, SEQ, SER, SEU).
<i>tst</i>	Gene that encodes the toxic shock syndrome toxin (TSST) associated with severe clinical symptoms
<i>sea</i> , <i>seb</i> , <i>sec</i> , <i>sed</i> , <i>see</i> , <i>seg</i> , <i>seh</i> , <i>sei</i> , <i>sej</i> , <i>sek</i> , <i>sel</i> , <i>sem</i> , <i>sen</i> , <i>seo</i> , <i>sep</i> , <i>seq</i> , <i>ser</i> , <i>seu</i>	Staphylococcal enterotoxin genes related to food poisoning
PVL (Panton-Valentine Leukocidin)	Leukotoxin related to alterations of cell permeability and leukocyte destruction

Table S2. SCC_{mec} types described in *S. aureus* as combination of the *mec* complex and the *ccr* type, indicating in parentheses the structure of the *mec* complex and the *ccr* genes that constitute the *ccr* type.

SCC _{mec} type	<i>mec</i> complex	<i>ccr</i> type
I	B (IS1272- Δ <i>mecR1-mecA</i> -IS431)	1 (<i>ccrA1ccrB1</i>)
II	A (<i>mecI-mecR1-mecA</i> -IS431)	2 (<i>ccrA2ccrB2</i>)
III	A (<i>mecI-mecR1-mecA</i> -IS431)	3 (<i>ccrA3ccrB3</i>)
IV	B (IS1272- Δ <i>mecR1-mecA</i> -IS431)	2 (<i>ccrA2ccrB2</i>)
V	C2 (IS431- Δ <i>mecR1-mecA</i> -IS431) (IS431s in opposite direction)	5 (<i>ccrC1</i>)
VI	B (IS1272- Δ <i>mecR1-mecA</i> -IS431)	4 (<i>ccrA4ccrB4</i>)
VII	C1 (IS431- Δ <i>mecR1-mecA</i> -IS431)	5 (<i>ccrC1</i>)
VIII	A (<i>mecI-mecR1-mecA</i> -IS431)	4 (<i>ccrA4ccrB4</i>)
IX	C2 (IS431- Δ <i>mecR1-mecA</i> -IS431) (IS431s in opposite direction)	1 (<i>ccrA1ccrB1</i>)
X	C1 (IS431- Δ <i>mecR1-mecA</i> -IS431)	7 (<i>ccrA1ccrB6</i>)
XI	E (<i>blaZ-mecC-mecR1-mecI</i>)	8 (<i>ccrA1ccrB3</i>)
XII	C2-like (Δ IS431- <i>mecA</i> - Δ <i>mecR1</i> -IS431)	9 (<i>ccrC2</i>)
XIII	A (IS431- <i>mecI-mecR1-mecA</i> -IS431)	9 (<i>ccrC2-new</i>)
XIV	A (<i>mecI-mecR1-mecA</i> -IS431)	1 (<i>ccrA1ccrB1</i>)