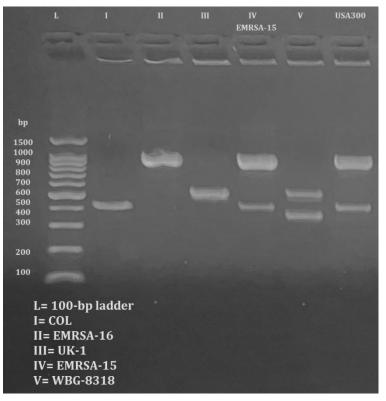
## Molecular Analysis – SCCmec typing results:

Target	Primer	Sequence $(5' \rightarrow 3)$	Product,	SCCmec Types:				
			bp	Ι	II	III	IV	V
ccrA2-	F	ATTGCCTTGATAATAGCCYTCT	937		$\checkmark$		$\checkmark$	
В	R	TAAAGGCATCAATGCACAAACACT						
ccrC	F	CGTCTATTACAAGATGTTAAGGATAAT	518			. /		. /
	R	CCTTTATAGACTGGATTATTCAAAATAT				V		$\mathbf{V}$
IS1272	F	GCCACTCATAACATATGGAA	415				. /	
	R	CATCCGAGTGAAACCCAAA					$\mathbf{V}$	
mecA-	F	TATACCAAACCCGACAACTAC	- 359	]				. /
IS431	R	CGGCTACAGTGATAACATCC						V

## Table 1 Primers used for SCCmec typing with their corresponding types

## Multiplex PCR conditions:

Initial denaturation for 4 min at 94°C, followed by 30 cycles of denaturation (30 s at 94°C), annealing (30 s at 55°C) and extension (60 s at 72°C), and a final extension for 4 min at 72°C. Electrophoresis was performed on a 1.5% agarose gel via ethidium bromide staining, afterwards the gel was visualized under ultraviolet illumination



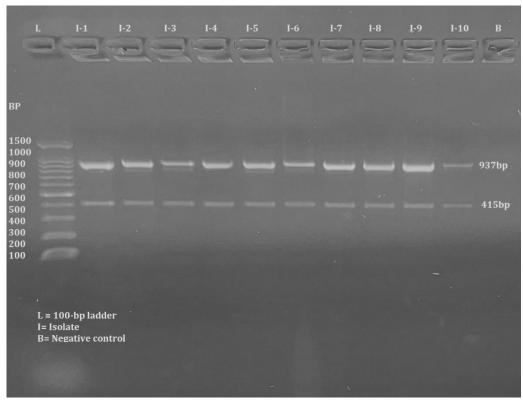


Figure 1 SCCmec typing for the reference strains. Specific genes were targeted in this multiplex PCR assay: IS1272 for type I (415bp), ccrA2-B for type II (937bp), ccrC for type III (518bp), ccrA2-B and IS1272 for type IV (937bp and 415bp, respectively) and ccrC and mecA-IS431 for type V (518bp and 359bp, respectively). 100-bp ladder (Promega, USA) was used in this assay.

Figure 2 SCCmec typing for the 10 clinical isolates. Specific genes were targeted in this multiplex PCR assay: IS1272 for type I (415bp), ccrA2-B for type II (937bp), ccrC for type III (518bp), ccrA2-B and IS1272 for type IV (937bp and 415bp, respectively) and ccrC and mecA-IS431 for type V (518bp and 359bp, respectively). 100-bp ladder (Promega, USA) was used in this assay.