

**Introduction:** *Staphylococcus aureus* colonises the nares of approximately 30% of healthy people but it is also a major cause of infection including skin and soft tissue and bloodstream infections. MRSA arises from the acquisition of either *mecA* or *mecC*, carried on mobile staphylococcal cassette chromosome *mec* (SCC*mec*) elements by methicillin-susceptible *S. aureus* (MSSA) clones. There is convincing evidence that *mec* and SCC*mec* originated in coagulase-negative staphylococci (CoNS), which also colonise the nares, throat and skin of healthy individuals.

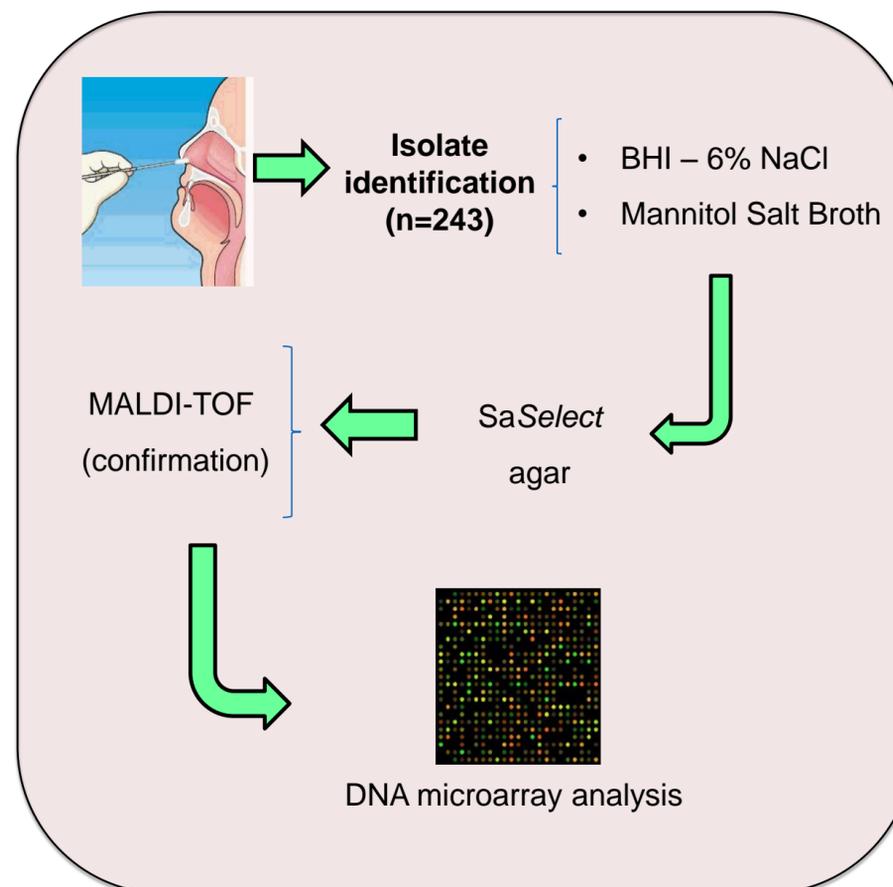
**Methods:** Nasal swabs (eSwab Copan®) were obtained from 243 students without any previous hospital contact (ethical approval number REC949 by RCSI Research Committee). All the samples were screened using two step enrichment procedure to recover *S. aureus* (MRSA or MSSA), MRSE and other CoNS from the swab (Huber et al. 2011).

**References:**

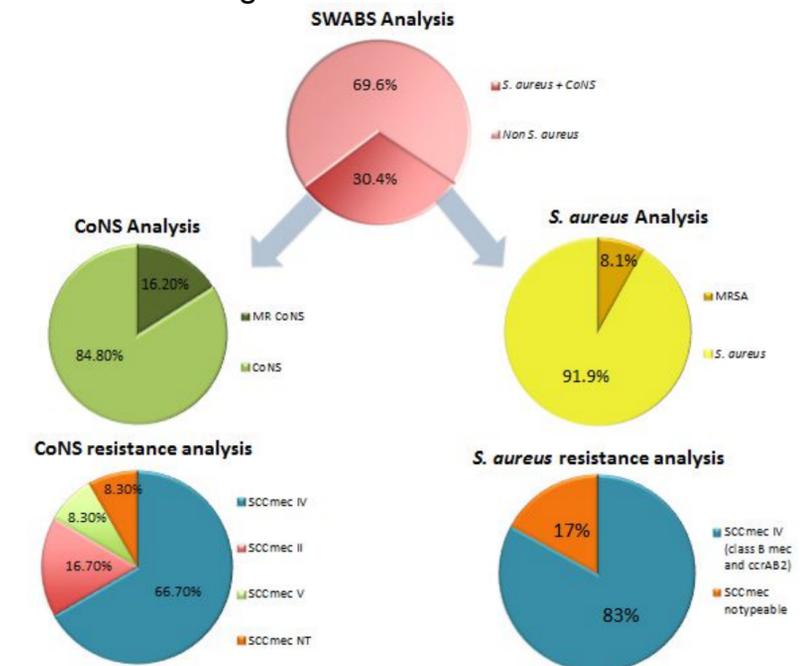
•Lindsay JA. 2010. Genomic variation and evolution of *Staphylococcus aureus*. Int J Med Microbiol. 300:98–103.

•Barbier F., Ruppe E., Hernandez D., Lebeaux D., 2010. Methicillin-resistant coagulase-negative staphylococci in the community: high homology of SCC*mec* IVa between *Staphylococcus epidermidis* and major clones of methicillin-resistant *Staphylococcus aureus*. J. Infect. Dis., 202: 270–81.

The swabs were enriched in Brain Heart Infusion (BHI) supplemented with 6% NaCl for 24 h at 37°C followed by further enrichment in mannitol salt broth for 24 h at 37°C. An aliquot (100 µl) of a 1/1000 dilution from this culture was spread onto SaSelect agar (Bio-Rad®, Hercules, CA, USA). The criteria for presumptive identification of different staphylococci growing on SaSelect were defined following the manufacturer instructions. DNA microarray was performed according to the manufacturer's instruction.



**Results:** Overall, class B *mec* (13/18) and *ccrAB2* (14/18) were common to MRSA and MR-CoNS and SCC*mec* IV predominated among MRSA and MR-CoNS.



**Conclusion:** The occurrence of similar SCC*mec* elements among *S. aureus* and CoNS in healthy individuals, supports the evidence that CoNS are a reservoir of mobile genetic elements including SCC*mec* that may contribute to the *S. aureus* gene pool even in the absence of significant antibiotic selective pressure. Further studies will include analysis of the entire DNA sequence of these SCC*mec* elements from MRSA and MR-CoNS to determine the full extent of their similarity.

**Acknowledgements:**

- Funded by RCSI School of Postgraduate Studies,
- Funded by CAPES/Science Without Border