

## **Poster Overview**

Nr.	Title	Presenter	Authors
P287	Comparison of the in vitro antibacterial activity of the clinically accepted glycopeptide antibiotics with three novel semisynthetic glycopeptides	Vladimir Vimberg	Vladimir Vimberg   Aniko Borbas   Pal Herczeghp   Pauline Cavanagh   Vaclava Adamkova   Gabriela Balíková Novotná
P288	WHOLE GENOME SEQUENCING OF STAPHYLOCOCCUS AUREUS CLONAL COMPLEX 45 ISOLATES DOES NOT DISTINGUISH NASAL CARRIAGE FROM BACTEREMIA	Chandler Roe	Chandler Roe   Marc Stegger   Berit Lilje   Kim Lee Ng   Elizabeth Driebe   David M. Engelthaler   Paal Andersen
P289	Diverse and unexpected staphylococcal responses to skin lipids	Mal Horsburgh	Xizhang Zhao   Josephine Moran   Mal Horsburgh
P290	Community-Associated MRSA Colonization: Look Beyond Your Nose	Maghnus O'Seaghdha	Olivia Huber   Joanna O'Gorman   Bouchra Benghomari   Gillian Weaver   Sara Tavana   Angel Wong   Jyl Richards   Susan Duty   Elizabeth Scott   Maghnus O'Seaghdha
P291	JAK-INHIBITORS IN S AUREUS INFECTIONS	Anders Jarneborn	Anders Jarneborn   Majd Mohammad   Tao Jin
P292	Complete sequences of two Staphylococcus aureus plasmids carrying genes for resistance to antibiotics, heavy metals, biocides and/or virulence	Sofia Santos Costa	Sofia Santos Costa   Kristina Kadlec   Miguel Viveiros   José Melo-Cristino   Stefan Schwarz   Isabel Couto
P293	RELATIONSHIP BETWEEN ANTIBIOTIC RESISTANCE OF STAPHYLOCOCCUS AUREUS AND RADIOGRAPHY IMAGES IN PATIENTS WITH DIABETIC FOOT ULCERS	Maria Stańkowska	Maria Stańkowska   Katarzyna Garbacz   Marek Bronk
P294	EVALUATION OF SCCMECFINDER IN NON- AUREUS STAPHYLOCOCCI	H. Kaya	H. Kaya   M. Stegger   T. B. Johannesen   A. R. Larsen
P296	DIVERSE SCCMEC- FUSC ELEMENTS AMONG DISTINCT COMMUNITY-ASSOCIATED METHICILLIN-RESISTANT STAPHYLOCOCCUS AUREUS STRAINS	Anna Shore	Grainne Brennan   Tanya Fleming   Peter Kinnevey   Brian O'Connell   David C. Coleman   Anna Shore
P297	Fibrinolytic and antibiotic treatment of MRSA intravascular biofilm infection	Mikkel Illemann Johansen	Mikkel Illemann Johansen   Søren Jensen-Fangel   Søren Rahbek   Ole Halfdan Larsen   Rikke Louise Meyer   Nis Jørgensen
P298	Oxazolidinone resistance in coagulase-negative staphylococci from household animals	Stefan Schwarz	Chengtao Sun   Peng Zhang   Xing Ji   Run Fan   Baoli Chen   Yang Wang   Stefan Schwarz   Congming Wu
P299	Acquisition of daptomycin resistance influences virulence traits in MRSA strains	Adriana Rosato	Agustina Taglialegna   Maria Celeste Varela   Roberto Rosato   Adriana Rosato

## **Poster Abstracts**

## [P292] COMPLETE SEQUENCES OF TWO Staphylococcus aureus PLASMIDS CARRYING GENES FOR RESISTANCE TO ANTIBIOTICS, HEAVY METALS, BIOCIDES AND/OR VIRULENCE

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**Aim:** To gain detailed insight into the genetic organization of two multidrug resistance plasmids, pSM31 and pSM39, from clinical *S. aureus* isolates collected in a hospital in Lisbon, Portugal, and to identify factors that may explain their selection and persistence.

**Methods:** Plasmid pSM31 was sequenced by Sanger sequencing using cloning and primer walking whereas pSM39 was sequenced by next generation sequencing. Sequence analysis was carried out using the BlastN and BlastP programs and the ORF Finder software from NCBI.

**Results:** The two plasmids fully characterized in this work were non-conjugative multiresistance plasmids. pSM31 (27,424 bp) is closely related to the family of globally dispersed plasmids from the pGSA23 group harbouring determinants for resistance to  $\beta$ -lactams and cadmium as well several enterotoxin genes. pSM39 (26,036 bp) is a newly described member of the pGSA11 group that carries genes conferring resistance to  $\beta$ -lactams, biocides, cadmium, zinc and mercury and comprises a region that may have been acquired by recombination with a *S. epidermidis* plasmid.

**Conclusions:** The sequence analysis of these two plasmids confirmed their involvement in antimicrobial resistance among *S. aureus* strains isolated from a clinical setting. It also provided genetic evidence for the potential risk of co-selection of antibiotic resistance genes and/or virulence genes by selective pressure of antibiotics, biocides and/or heavy metals.

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