

## **POSTER PRESENTATION**

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# SCCMEC and SPA typing of meticillin resistance Staphylococcus aureus isolated from infections from Southern Poland

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## Introduction

The spa gene encodes protein A and is used for typing of methicillin-resistant *Staphylococcus aureus*, MRSA, such as the mec operon carried by staphylococcal cassette chromosome (SCCmec).

## **Objectives**

The aim of this study was the molecular typing of MRSA isolated from different forms of infections for epidemiological purposes using spa typing method and SCCmec classification.

## **Methods**

A total of 90 MRSA isolates coming from eight hospitals from southern Poland were tested. Isolates originated from: bloodstream and respiratory tract infections (36), surgical site infections (30), chronic wounds (24). S pa typing was performed as described previously [1], using the spa typing website (http://www.spaserver.ridom.de/). Staphylococcal cassette chromosome mec (SCC mec) typing was performed as described previously [2].

#### Results

The majority of MRSA strains were of SCCmec type II (42.2%) or SCCmec IV (21.1%). Eleven strains was marked as SCCmec III (12.2%), 8 as SCCmec V (8.9%), 4 as SCCmec I (4.4%) and one as SCCmec VI.

The spa type t003 was the most frequently observed (37.8% of strains), then t138 (14.4%), t008 and t037 and t041 (4.4%),

SCC *mec* type II and spa-t003 together were characteristic for 25% of the chronic wounds and 27% in SSI, but 39% in invasive infections.

SCCmec III and t138 occurred in 10% of the strains from two hospitals, SCCmecIV and t003 occurred in 6.7% strains from two hospitals.

### **Conclusion**

Epidemiological and molecular studies of MRSA isolates allowed to detail insight into the problem of staphylococcal infections. Those methods are less time-consuming than PFGE and give the opportunity for the observation of the current situation and epidemiological trends for resistant strains on the level of ward/hospital or even whole region (supported by a grant DEC-2011/03/B/NZ7/01911).

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#### References

- 1. Harmsen: J ClinMicrobiol 2003, 41(12):5442-5448.
- 2. Kondo: Antimicrob Agents Chemother 2007, 51(1):264-74.

#### Disclosure of interest

None declared.

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