# The isolation rates of oxacillin-susceptible mecA-positive Staphylococcus aureus at Nagasaki University Hospital in Japan



Yui Shigeishi<sup>1</sup>, Yasuhide Kawamoto<sup>1</sup>, Daisuke Sasaki<sup>1</sup>, Kosuke Kosai<sup>2</sup>, Fujiko Mitsueda-Kaseida<sup>1,2</sup>, Katsunori Yanagihara<sup>1,2</sup>

1. Department of Laboratory Medicine, Nagasaki University Hospital, Nagasaki, Japan

2. Department of Laboratory Medicine, Nagasaki University Graduate School of Biomedical Sciences, Nagasaki, Japan

**RES-144** 

# INTRODUCTION

Methicillin-resistant S. aureus (MRSA) has been defined by antimicrobial susceptibility testing. Recently, PCR amplification of *mecA* or *mecC* genes has also been used for MRSA detection, although it is not yet the standard method. Oxacillin-susceptible mecA-positive S. aureus (OS-MRSA) can be easily misidentified as MSSA by antimicrobial susceptibility testing1.

OS-MRSA has been reported in many countries, but few reports indicate isolation rates 1,2,3.

Therefore, we investigated the isolation rates of OS-MRSA at our hospital.

# MATERIALS AND METHODS

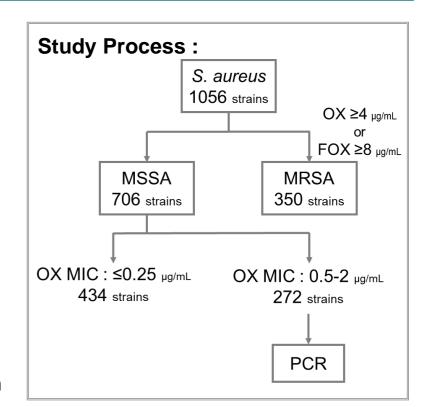
We included 1056 S. aureus strains isolated at Nagasaki University Hospital from 2023 to 2024.

#### **Determination of MIC:**

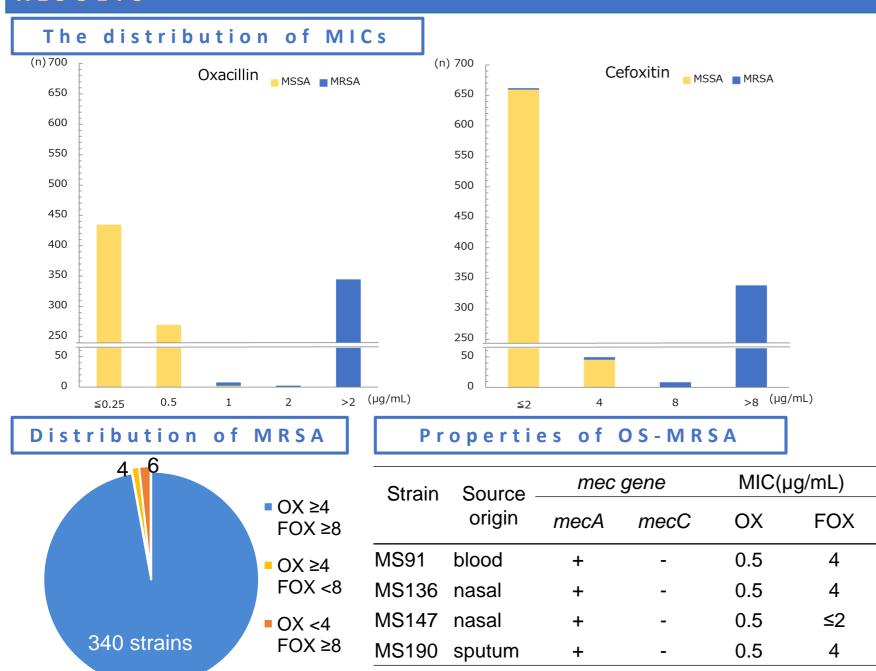
Antimicrobial susceptibility testing for all isolates were determined using the BD Phoenix. The strains with oxacillin (OX) MIC of ≥4 µg/mL or cefoxitin (FOX) MIC of ≥8 µg/mL were defined as MRSA according to CLSI M1004, and their MIC distributions were analyzed.

#### Detection of *mecA* and *mecC* gene:

PCR was performed to detect the presence of mecA and mecC genes 5, particularly in MSSA strains with oxacillin MICs ranging from 0.5 to  $2 \mu g/mL^1$ .



### RESULTS



#### The distribution of MICs:

- The rate of MRSA was 33.1% (350/1056) and that of MSSA was 66.9% (706/1056).
- The MIC distribution for oxacillin was as follows:  $\leq 0.25 \,\mu \text{g/mL}$  in 41.1% (434/1056), 0.5 μg/mL in 25.5% (269/1056), 1 μg/mL in 0.7% (7/1056), 2 μg/mL in 0.2% (2/1056), and  $>2 \mu g/mL$  in 32.6% (344/1056).
- The MIC distribution for cefoxitin was as follows:  $\leq 2 \mu g/mL$  in 62.6% (661/1056), 4 μg/mL in 4.6% (49/1056), 8 μg/mL in 0.8% (8/1056), and >8 μg/mL in 32.0% (338/1056).

#### **Distribution of MRSA:**

· Among MRSA, 6/350 (1.71%) were susceptible to oxacillin.

## **Properties of OS-MRSA:**

• The *mecA* gene was detected in 1.5% (4/272) of MSSA strains with oxacillin MICs ranging from 0.5 to 2  $\mu$ g/mL.

# DISCUSSION

- · Six MRSA strains were susceptible to oxacillin, and discordant susceptibility results between oxacillin and cefoxitin in MRSA was consistent with previously reported 6.
- The isolation rate of OS-MRSA at our hospital was 1.5%, consistent with previously reported results (1.2 $\sim$ 1.8%) <sup>1,2,3</sup>.
- · In one case, OS-MRSA was isolated from a blood culture, suggesting a possible bloodstream infection. Since the antibiotics used for MSSA and MRSA differ, accurate identification of OS-MRSA is critically important.
- · In this study, the oxacillin MIC for OS-MRSA was 0.5 μg/mL, representing the lowest MIC within the range of the examined strains (oxacillin MICs: 0.5–2 μg/mL). Therefore, it was suggested that even strains with lower oxacillin MICs (<0.5 μg/mL) might be identified as OS-MRSA.

# CONCLUSION

The findings of this study revealed the frequency of OS-MRSA.

5) National Institute of Infectious Diseases home page (https://id-info.jihs.go.jp/relevant/manual/010/MRSA20240318.pdf)

However, the mecA gene detection was performed only for MSSA strains with oxacillin MICs between 0.5 and 2 µg/mL. Therefore, comprehensive surveillance including all MSSA strains is necessary to accurately assess the prevalence of OS-MRSA.

# REFERENCE

- 1) Y. Hososaka et al. J Infect Chemother (2007) 13:79–86
- 2) J.Liu et al. Journal of Microbiology, Immunology and Infection (2021) 54, 1070e1077

6) C-M. Ho et al. / Diagnostic Microbiology and Infectious Disease 86 (2016) 405-411

- 4) CLSI M100 ED35:2025 (https://em100.edaptivedocs.net/)
- 3) K. Saeed et al. Infection (2014) 42:843-847

# CONTACT

Yui Shigeishi

Department of Laboratory Medicine, Nagasaki University Hospital,

1-7-1, Sakamoto, Nagasaki 852-8501, Japan

E-mail: y-shigeishi@nagasaki-u.ac.jp

TEL: +81-95-819-7574 FAX: +81-95-819-7422