Molecular Epidemiology of Methicillin Resistant Staphylococcus aureus Isolated from Newborns in a Hospital in Rio de Janeiro, Brazil

MM Loureiro, BA de Moraes, MRR Quadra**, GS Pinheiro**, PN Suffys*, MD Asensi/+ 

Laboratório de Enterobacterias, Departamento de Bacteriologia *Laboratório de Biologia Molecular e Diagnóstico de Doenças Infecciosas, Departamento de Bioquímica e Biologia Molecular, Instituto Oswaldo Cruz, Av. Brasil 4365, 21045-900, Rio de Janeiro, RJ, Brasil **Hospital Maternidade Alexander Fleming II, Sistema Único de Saúde, Rio de Janeiro, RJ, Brasil

Methicillin resistant Staphylococcus aureus (MRSA) is an organism that is frequently transmitted in hospitals and perinatal units. The MRSA is considered a public health problem in neonatology because of its strong potential for dissemination in the wards associated with high rates of morbidity and mortality. In this study we describe the bacteriological, epidemiological and molecular characteristics of MRSA isolated from anterior nares and blood cultures of newborns hospitalized in a public maternity hospital in the city of Rio de Janeiro, Brazil. The frequency of MRSA isolated from nasal swabs of newborns was 47.8% (43/90). The genetic analysis of MRSA strains from anterior nares, showed 8 different pulsed field gel electrophoresis patterns (PFGE). Upon analysis of PFGE patterns of the 12 MRSA strains isolated from blood cultures, 8 different patterns were observed, 9 (75%) strains were genetic related to nasal secretion isolates patterns. In conclusion, our data demonstrate the importance of screening of newborns for the presence of MRSA in Brazilian hospitals and the usefulness of genetic typing of these pathogen during epidemiologic studies. This should lead to a better knowledge on the significancy and spreading of MRSA in the hospitals.

Key words: newborns - nasal carriers - methicillin resistant Staphylococcus aureus - blood culture - pulsed field gel electrophoresis - hospital infection - Rio de Janeiro - Brazil

The newborns in intensive and/or intermediate care units are considerably susceptible to acquire hospital infections of several microorganisms (Pujol et al. 1994, Haley et al. 1995). Methicillin resistant Staphylococcus aureus (MRSA) has emerged over the past 30 years as an important cause of hospital infections and is actually an organism that is frequently transmitted in hospitals and perinatal units (Haley et al. 1982, Boyce et al. 1994, Na’Was et al. 1998). Outbreaks causing several pathological problems to the newborns and mothers have been reported (Pujol et al. 1996, Corbella et al. 1997). The MRSA is considered a public health problem in neonatology because of its strong potential for dissemination in the wards, high levels of antibiotic resistance, associated with high rates of morbidity and mortality (Soares et al. 1997, Tambic et al. 1997, Conterno et al. 1998).

The prevalence of MRSA in Brazil is considered elevated, specially in large hospitals (Branchini et al. 1993, Sader et al. 1994a), and in despite of the wide distribution, there are some hospitals that never had MRSA outbreak (Teixeira et al. 1996).


Epidemiological transmission studies of MRSA have been traditionally performed by phenotypic characterization (Bannerman et al. 1995, Na’Was et al.1998), therefore, the increasing of the molecu-
lar epidemiology studies revolutioned our understanding of diseases transmission and allowed to recognize and investigate outbreaks in ways never before possible (Ostrost 1999). The identification of several new clones and distribution of endemics and epidemics strains became feasible through the use of molecular tools as pulsed field gel electrophoresis (PFGE) and restriction fragment length polymorphism (RFLP) of chromossomal DNA associated with hybridization techniques (Kreiswirth et al. 1993, Teixeira et al. 1995, Tambic et al. 1997).

In this study we describe the bacteriological and epidemiological characteristics of MRSA, isolated from anterior nares and blood of newborns hospitalized in a public maternity hospital in the city of Rio de Janeiro, Brazil, and their genetic analysis by PFGE as an epidemiological marker.

**MATERIALS AND METHODS**

Patients - The Hospital Maternidade Alexander Fleming II is a maternity-childhood hospital in the city of Rio de Janeiro, Brazil, providing assistance and perinatal care including a neonatology intermediate care unity (NIU) with 40 beds and a neonatology intensive care unity (NICU) with 15 beds.

After the detection of a positive blood culture for MRSA from a hospitalized newborn in the NICU in April 1998 (index case), a prospective study was realized for detection of MRSA in anterior nares of hospitalized newborns and health care workers in the NICU and NIU. A total of 332 nasal swabs were taken from anterior nares in following groups: newborns group 1 (NBG 1), samples from 90 newborns collected at the period between 04/13 and 04/16/98; NBG 2, 82 samples from newborns with MRSA colonization belonged to NBG 1 submitted to descolonization treatment by nasal application of 2% mupirocin ointment and bathing in 4% chlorexidine solution; NBG 3, samples from 56 newborns hospitalized, collected at the period between 04/27 and 05/04/98, after the adoption of control procedures and treatment of the colonized newborns group (NBG 1); NBG 4, samples from 104 neonatology health care workers collected in the same period of the newborns included in NBG 1. In addition, with the objective to control the MRSA transmission, all staff adopted contact isolation precautions such as handwashing, gloving and gown use for the management of the infected newborns.

All MRSA (12 strains) isolated from blood cultures of hospitalized newborns with hospital infection, during the period from April 1997 to April 1999 were genetically analyzed and compared to those isolated from anterior nares secretions. As we detected a interval of 8 months (08/14/97-04/01/98) without positive blood culture for MRSA, we considered the strain 5 BC isolated in 04/01/98, as the index case to study the patients and staff carriers.

**Bacterial strains and susceptibility testing -** Culture for *S. aureus* using nasal secretion and blood as well as identification of the organism were performed according to Kloos and Bannerman (1995). The MRSA characterization was made by antimicrobial susceptibility tests performed with disk diffusion method according to NCCLS (National Committee for Clinical Laboratory Standards 1997). The following concentrations of the antimicrobials drugs were used: penicillin (10 U), ampicillin (10 mcg), cephalotin (30 mcg), oxacillin (1 mcg), amikacin (30 mcg), erythromycin (15 mcg) and vancomycin (30 mcg). The reference strain *S. aureus* (ATCC 25923) was used as a quality control.

**PFGE -** The chromossomal DNA restriction analysis of 43 MRSA isolated from anterior nares and 12 from blood were performed by PFGE. Therefore, strains were submitted to chromosomal DNA extraction and processing according to the procedure described by Sader et al. (1994b). Nucleic acids present in the agarose plugs were digested with 10U *SmaI* at 30°C during 20 h and electrophoresis was performed at 6V/cm during 23 h in 0.5X TBE at 13°C in 1% agarose gel using a ramping switch time of 1-50 sec. All agarose gels were made for twice to verify the reproducibility of the tests.

Band patterns were analyzed using Gelcompar 4.1 (Applied Maths, Belgium), without using internal markers. The similarities between fingerprints were determined by construction of a similarity matrix using the Dice coefficient with 1.5% position tolerance, what generated a dendrogram using the UPGMA clustering algorithm.

Definition of clonal structure of MRSA isolates was according to Tenover et al. (1995): PFGE patterns with more than 90% similarity under the stringency conditions defined in this study were considered as the same clone.

**RESULTS**

The frequency of MRSA isolated from nasal swabs in the four groups analyzed was higher in NBG 1 (43 positive cultures; 47.8%). In NBG 2, two positive cultures were obtained from two newborns carriers (NBG 1) submitted to descolonization treatment. No positive cultures were obtained from NBG 3 and NBG 4. Upon 43 newborns with positive nasal swabs for MRSA, 32 (74.4%) were asymptomatic, 6 (14%) showed a clinical picture compatible with *S. aureus* infection (pneumonia, sepsis or meningitis), while 5 more (11.6%) died with generalized infection caused by MRSA.
All MRSA strains were resistant to all antimicrobials utilized in the susceptibility test, with exception to vancomycin.

Upon genetic analysis of MRSA isolates from anterior nares, 8 different PFGE patterns were observed among the 43 strains analyzed (Fig. 1-A) and in the construction of a dendrogram, these patterns were divided into two main genetic clusters (Fig. 2). The first cluster, designed cluster I, had 86% percent of the strains isolated from nasal swabs and consisted of pattern A (26 strains); from four more probably and closely related patterns were observed in 11 strains (A1: 5 strains; A2: 2 strains; A3: 1 strain and A4: 3 strains). The second, designed cluster II, consisted of 3 different patterns observed in 6 more strains (B: 2 strains; C: 3 strains and D: 1 strain).

We detected 8 different profiles in the PFGE analysis of the 12 MRSA strains isolated from blood, along two years (Fig.1-B, C). A dendrogram constructed from all profiles (Fig. 3) revealed that 9 (75%) strains were genetic related to nasal secretion isolates, with the following distribution: 50% of MRSA isolates (2BC, 6BC, 8BC, 9BC, 10BC and 12BC) had 90 to 96% of similarity to clone A1; 25% (3BC, 4BC, and 5BC) had 94 to 100% of similarity to the clone A4 and the patterns 1BC, 7BC and 11BC apparently form a distinct genetic group in this analysis.

The distribution of blood culture MRSA chromosomal profiles according to isolation date and type of anterior nares clone (Table) showed...
that the clone A4 had similar PFGE profiles in the strain that started the epidemiological study (5BC) and the strains 3BC and 4BC, both isolated from blood culture 8 months before. The clone A1, also was observed similar PFGE profile in one strain (12BC) isolated from blood culture one year later the investigation of nasal carriers.

**DISCUSSION**

Infections with MRSA increase significantly the cost of hospitalization, mortality and morbidity rates of hospitalized patients, especially newborns (Jarvis 1987, Wenzel 1994, Smith & Doebbeling 1996). Detection of nasal MRSA carriers in hospitalized patients and in hospital staff by a variety of procedures has been associated with decreasing levels of infections and prevention of the transmission of these microrganisms (Blumberg & Klugman 1994).

Isolation of MRSA in nasal secretion of almost half of the hospitalized newborns from NBG 1 investigated in this study revealed the existence of significant colonization of the organism in that

**TABLE**

<table>
<thead>
<tr>
<th>Blood culture methicillin resistant <em>S. aureus</em> DNA profiles according to isolation date and type of anterior nares clone</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chromosomal DNA profiles</td>
</tr>
<tr>
<td>1BC</td>
</tr>
<tr>
<td>2BC</td>
</tr>
<tr>
<td>3BC</td>
</tr>
<tr>
<td>4BC</td>
</tr>
<tr>
<td>5BCa</td>
</tr>
<tr>
<td>6BC</td>
</tr>
<tr>
<td>7BC</td>
</tr>
<tr>
<td>8BC</td>
</tr>
<tr>
<td>9BC</td>
</tr>
<tr>
<td>10BC</td>
</tr>
<tr>
<td>11BC</td>
</tr>
<tr>
<td>12BC</td>
</tr>
</tbody>
</table>

*a*: index case that motivate the study of colonization; 
DP: different pattern
population. The treatment with 2% mupirocin ointment, the drug of choice for treatment of nasal colonization with MRSA, associated with bathing in a 4% solution of chlorhexidine were efficient to eliminate the MRSA from NBG 1. After this treatment, the control cultures for descolonization were negative (NBG 2). These data are in agreement with other reports on the efficacy of prevention of transmission of the pathogen by these measures (Blumberg & Klugman 1994, Chatfield et al. 1994, Coello et al. 1994, Teixeira et al. 1996, Tambic et al. 1997). Therefore, one month after the drug administration two patients revealed MRSA in anterior nares. Because the control cultures were negative, probably these newborns were reinfected by MRSA. Unfortunately, we did not have these two MRSA strains to include into epidemiological and molecular analysis.

The fact that 74.4% of the individuals with colonized anterior nares studied were asymptomatic demonstrates the need for large scale monitoring of nasal carriers of MRSA by culture. Earlier studies have shown the participation of clinical staff in the spreading of MRSA in hospitals through manipulation of carriers without the adoption of precaution measures, and the importance of nasal carriage for development of disease (Tambic et al. 1997, Villari et al. 1998). In this study however, none of the members of the hospital staff was found to be a carrier of MRSA and probably transmitted the microorganism by hands.

Because of its reproducibility and discriminatory capacity, the separation of cleaved genomic DNA by PFGE has been shown to be an excellent typing technique for MRSA in comparison with more classical or other molecular typing methods (Bannerman et al. 1995, Haley et al. 1995, Na’Was et al. 1998).

In this study, 60% of the isolates from nasal secretion (26 strains) had an identical PFGE profile, suggesting that transmission of a single strain occurred extensively within the hospital unit. Eighty-six percent of the nasal strains belonged to a cluster I of at least 90% similarity meaning that most of the strains isolated during a 4-day period were highly related. Also, 75% of the MRSA strains isolated from blood during a period of two years had a similarity of more than 90% to patterns of strains isolated from nasal secretion; this observation suggests that strains are endemic in this hospital unit and these results are in agreement with Haley et al. (1995), who described the persistence of a MRSA strain in the neonatal intensive care unit in a hospital in Texas during three years, even after implementation of control measures.

Our observations of similar PFGE patterns in strains isolated from the nose and blood, specially in newborns who developed MRSA pathologies agree with earlier suggestions that the respiratory tract is the primary port of entry for colonization and can lead to pneumonia, septicemia and death (Pujol et al. 1994, 1996, Corbella et al. 1997, Cordero et al. 1997).

In conclusion, our data demonstrated the importance of screening the newborns for the presence of MRSA in Brazilian hospitals and the usefulness of genetic typing of these pathogens during epidemiologic studies on a small and large scale basis. This should lead to a better knowledge on the significance and spreading of MRSA in the hospitals.

REFERENCES


