

# Impact of admission screening for methicillin-resistant Staphylococcus aureus on the length of stay in an emergency department.

AUTHOR(S)

Peadar Gilligan, Michael Quirke, Stephen Winder, Hilary Humphreys

**CITATION** 

Gilligan, Peadar; Quirke, Michael; Winder, Stephen; Humphreys, Hilary (2010): Impact of admission screening for methicillin-resistant Staphylococcus aureus on the length of stay in an emergency department.. Royal College of Surgeons in Ireland. Journal contribution. https://hdl.handle.net/10779/rcsi.10768778.v2

HANDLE

10779/rcsi.10768778.v2

LICENCE

#### CC BY-NC-SA 4.0

This work is made available under the above open licence by RCSI and has been printed from <a href="https://repository.rcsi.com">https://repository.rcsi.com</a>. For more information please contact <a href="mailto:repository@rcsi.com">repository@rcsi.com</a>

**URL** 

https://repository.rcsi.com/articles/journal\_contribution/Impact\_of\_admission\_screening\_for\_methicillin-resistant\_Staphylococcus\_aureus\_on\_the\_length\_of\_stay\_in\_an\_emergency\_department\_/10768778/2

2 3 4	_	dmission screening for meticillin-resistant nureus (MRSA) on the length-of-stay in an emergency department
5		
6 7	<sup>1</sup> P Gilligan, <sup>1</sup> M Quirke, <sup>2</sup>	S Winder, <sup>3,4</sup> H Humphreys.
8 9	<sup>1</sup> Emergency Department	and <sup>2</sup> Information Technology Department, Beaumont
10	Hospital, Dublin, <sup>3</sup> Depar	tment of Microbiology, Beaumont Hospital, Dublin,
11	<sup>4</sup> Department of Clinical 1	Microbiology, Royal College of Surgeons in Ireland, Dublin
12		
13	Address for corresp	ondence
14 15 16 17 18 19 20 21	Dr Peadar Gilligan Consultant in Emergency Emergency Department Beaumont Hospital Beaumont Road Dublin 9 Ireland	Medicine
22	Funding source Th	is work was not supported by an external research grant or
23 24	fu	nding from a commercial or other source.
25	<b>Conflict of interest</b>	HH has received research funding from Pfizer, Steris
26		Corporation, 3M, Inov8 Science and Cepheid in the last
27		three years. He has also received lecture or consultancy
28		fees from 3M, Novartis and Astellas.
29		
30		

MRSA and Emergency Department length-of-stay

1

**Running title** 

## Abstract

1

2	Preventing and controlling meticillin-resistant Staphylococcus aureus (MRSA)
3	includes early detection and isolation. In the emergency department (ED), such
4	measures have to be balanced with the requirement to treat patients urgently and
5	transfer quickly to an acute hospital bed. We assessed the contribution that previous
6	MRSA risk group identification and selective <u>rescreening of those patients who were</u>
7	previously documented in the research hospital as being MRSA positive made to a
8	patient's stay in a busy, overcrowded ED. Patients with a previous diagnosis of
9	MRSA colonisation were flagged automatically as a "risk group" (RG) patient on
10	their arrival in the ED and were compared with those non-risk group (NRG), i.e. not
11	previously <u>proven in the research hospital to be infected or colonised with MRSA.</u>
12	Over an 18 month period, there were 16,456 admissions via the ED of which 985
13	(6%) were RG patients. The expected median times to be admitted following a request
14	for a ward bed for NRG and RG patients were 10.4 hours and 12.9 hours,
15	respectively. Female sex, older age (over 65 years old) and RG status all
16	independently predicted a statistically significant longer stay in the ED following a
17	request for a hospital bed. We consider that national and local policies for MRSA
18	need to balance the welfare of patients in the ED with the need to comply with best
19	practice, when there are inadequate ED and in-patient isolation facilities. Patients with

MRSA requiring emergency admission must have a bed to go to.

Emergency department, MRSA, surveillance, isolation, boarders

22

21

Key words:

#### Introduction

1

2 Healthcare-associated infection affects 5-10% of patients in an acute hospital, many of these infections are device-related and meticillin-resistant Staphylococcus aureus 3 (MRSA) accounts for approximately 16% of all infections. 1,2 General infection 4 prevention and control measures, such as the use of standard precautions, as well as 5 6 specific measures for MRSA are justified as the outcome from MRSA infections is 7 less favourable compared with other infections. A meta-analysis comparing 8 bloodstream infection due to MRSA and methicillin-susceptible S. aureus showed a significant increase in mortality associated with MRSA bloodstream infection.<sup>3</sup> 9 10 Preventing and controlling MRSA requires a multi-faceted approach that includes 11 early detection and isolation, a common feature of many national guidelines.<sup>4</sup> In the 12 13 UK and elsewhere, there is a move towards the introduction of universal screening, 14 i.e. the taking of swabs from all patients irrespective of risk, and this has provoked 15 some discussion on its merits.<sup>5</sup> 16 17 In the emergency department (ED), infection prevention and control measures have to 18 be balanced with the requirement to treat patients urgently and to transfer patients 19 requiring admission as quickly as possible to acute hospital beds. Furthermore, in 20 North America and elsewhere, the phenomenon of community-acquired MRSA is 21 being seen, particularly in EDs, in patient's without known risk factors for healthcareassociated MRSA. We have documented that anything that prolongs the patient's ED 22 stay may adversely affect patient welfare and compound overcrowding.<sup>7</sup> 23 24 national and local guidelines on the prevention and control of MRSA largely focus on in-patients but their implementation have implications for the ED. Prolonging a 25

- 1 patient's stay in the ED will further compound overcrowding, which is a recognised
- 2 contributor to spread of hospital acquired infection.<sup>8</sup>

3

- 4 We assessed the contribution that previous MRSA risk group identification and
- 5 selective <u>rescreening of these patients made to their stay in a busy, overcrowded ED.</u>

6

#### **Materials and Methods**

2	Institution	and	patients
_	IIIDILLIULU	uiiu	parient

The study was conducted in the ED of an urban academic teaching hospital with an annual census of approximately 46,000 patient visits, an admission rate of approximately 23% and an average occupancy with patients awaiting admission (boarders) of 105%. The observational study was approved by the chair of the institution's ethics committee. From the first of November 2006 to the 30<sup>th</sup> of April 2008, data were gathered in the ED's Oracle database on all patient attendances that resulted in admission. This Oracle database was interrogated using the Diver Solution

programme which is a data warehousing solution that facilitates the gathering of data

11 from different databases.

Selective screening, involved the taking of swabs from nose, groin and wounds (if present) from patients at the time of their re-attendance to the ED where there was a previous positive swab for MRSA in the research hospital. The prior diagnosis of MRSA colonisation or infection in the research hospital automatically gave rise to the patients being flagged as a "risk group" (RG) patient on the ED information technology system. For the purposes of this study the term "risk group" is applied only to those diagnosed with MRSA in the hospital laboratory on a previous admission. All such risk group patients were screened for evidence of ongoing MRSA colonisation. Previous MRSA carriers were declared clear on the basis of three negative swab results but even those cleared of infection remained as RG on return to the hospital.

Deleted: H

1	
2	Patients with a previous diagnosis of MRSA colonisation and / or infection, whether
3	or not they had negative screens subsequently, were flagged automatically as a "risk
4	group" (RG) patient on their arrival in the ED using the Oracle database. Those
5	patients with a prior diagnosis of MRSA colonisation were compared with those
6	without it with respect to age, sex, triage category, total ED processing time and time
7	from a bed on a ward being requested until the patient was admitted. Triage is the
8	systematic prioritisation of patients. The Manchester Triage System uses a series of
9	criteria to decide what level of priority a patient should have on the basis of their
10	presentation. The National Triage scale from the United Kingdom is used in many
11	EDs and is a five point scale from immediate (red), very urgent (orange), urgent
12	(yellow), standard (green) to non-urgent (blue).9
13	
14	The results from all patients screened for MRSA by the hospital's diagnostic
15	laboratory were entered on the laboratory information system and this informs the
16	hospital's inpatient system at 6 am on the morning following MRSA results being
17	available. The hospital's inpatient system informed in turn the Oracle database, and if
18	the patient subsequently re-attended the ED, that patient was automatically flagged as
19	a RG patient and selectively screened for evidence of ongoing MRSA colonization.
20	Previously MRSA positive or RG patients were screened with swabs from the nose,
21	groin and broken areas of skin, and swabs were cultured on MRSA Select

Chromogenic Agar (Bio-Rad Life Science group, France). Patients positive for

and continued on hospital wards after admission.  $\frac{10}{}$ 

MRSA were decolonised according to guidelines, and this was commenced in the ED

1 The ED is open plan with fourteen cubicles and two side rooms. Clinically stable 2 patients with a prior history of MRSA were isolated, where possible in these rooms, 3 pending the availability of other isolation facilities in the hospital. However, these 4 side rooms do not have separate toilet facilities or an ante room, and frequently the 5 number of patients requiring isolation for MRSA and for other indications exceeds the 6 capacity of the two rooms. Other patients in the department wait on trolleys or chairs, 7 usually in close proximity to other patients in the open plan area. 8 9 Statistical analysis 10 Cox proportional hazards methods were used to evaluate relative probabilities of 11 being admitted for RG patients versus non-RG patients (NRG) who acted as controls. 12 The interval time to event analysis determines whether a patient category has an 13 increased or decreased chance of admittance at a particular time point and the result is 14 defined by a hazard ratio. 15 16 A multi-factorial model was used to examine if risk group identification was 17 independently significant in the presence of confounding variables such as age and 18 sex. In addition the model was stratified into triage categories. Stata (version10, 19 College Station, Texas) was used to analyse the data and a p value less than 0.05 was 20 deemed to be significant.

#### Results

1

2 Over the 18 month period of the study there were 16,456 admissions via the 3 ED. Of these, 985 (6%) had a prior diagnosis of MRSA colonisation, i.e. were RG 4 patients. Amongst the NRG patients, 48.4% were female compared with 45.2% of RG 5 patients. The Manchester triage category for those who were triaged in RG and NRG 6 patients are compared in Table 1. Over the time frame of the study 161 of the 16,456 7 subsequently admitted patients did not undergo triage and are not represented in table 8 1. 9 10 The total time from arrival in the ED to admission to a ward bed was a median of 20.3 11 hours. For NRG and RG patients it was a median of 19.9 hours, [IQR 10.5 – 29.8 12 hours] and 22.6 hours [IQR 12.2 – 33.4 hours], respectively. 13 14 The expected median time to be admitted following a request for a ward bed was 10.5 15 hours; NRG patients waited a median of 10.4 hours [IQR 3.1 – 20.6 hours] compared 16 to 12.9 hours [IQR 4.3 – 26.6 hours] for RG patients. The results of the Cox model 17 revealed that older age (>65 years old) and female sex were statistically significant 18 factors influencing the time spent in the ED from arrival to a bed request but MRSA 19 colonisation was not (Table 2). However, female sex, older age and RG status all 20 independently predicted a longer stay in the ED following a request for a hospital bed, 21 i.e. RG status did not impact on the ED's and the on-call team's processing of patients 22 but did influence the time taken to allocate a ward bed. (Table 2). 23 24

### **Discussion**

1

- 2 Risk factors associated with healthcare-associated MRSA infection include advanced
- 3 age, male gender, previous hospitalisation, nursing home care, length of
- 4 hospitalisation, a stay in intensive care, chronic medical illness, prior antibiotic use,
- 5 presence of indwelling devices, asymptomatic colonisation with MRSA and exposure
- 6 to an infected or colonised patient.  $\frac{11}{1}$  In this study we applied the term risk group (RG)
- 7 only to those with a prior diagnosis of MRSA colonisation or infection in our hospital
- 8 laboratory. All such patients were electronically flagged as RG on their return to the
- 9 ED. The flagging of patients with a prior diagnosis of MRSA and selective
- 10 rescreening of this group in this study was associated with a prolonged ED stay.

11

- 12 In the United States it is estimated that the rate of MRSA colonisation in the
- community is about 2% of the population.  $^{12}$ Gopal *et al* found that 6.7% of screened
- adult emergency admissions to their UK based hospital were colonised with MRSA.<sup>8</sup>
- 15 It is estimated that a non-isolated MRSA carrier will infect 0.14 patients per day in the
- 16 absence of decolonisation. <sup>13</sup> Consequently, the early identification of colonized /
- 17 infected patients and the prompt implementation of contact precautions are important
- 18 | in preventing MRSA transmission in hospitals. 8,14-17

19

- With regards to the limitations of our research the RG status of a patient is not
- 21 validated but is assumed to reflect all patients with a previous positive result for
- 22 MRSA colonisation in our hospital. The study required the analysis of a real time
- computer database which is dependent upon the medical and nursing staff putting in
- data in a time sensitive manner. We have no reason to believe that any failure to do so
- would have been more prevalent in either those with or without MRSA.

Deleted: H

Overcrowding in EDs is a distressing and potentially dangerous phenomenon in many
health systems. In our hospital, the elderly, those with prior colonisation with MRSA
and women wait longest for an acute hospital bed when they require emergency
admission. The research hospital has insufficient acute beds to provide for emergency
admissions and this is further compounded by the fact that many patients in the
hospital experience delayed discharges owing to lack of nursing home beds or step
down facilities in the catchment area. The reason for the delay for females may be due
to the fact that our hospital does not usually house males and females in the same bay
of a ward and as more males are admitted as emergencies, finding a "female bed" can
be problematic. The expected time for admission from time of arrival in the ED was a
median of 22.6 hours for RG patients compared to 19.9 hours for NRG patients. The
additional 2.7 hours for those requiring selective screening and in an individual
patient sick enough to require hospital admission on an already unacceptably long
wait for a hospital bed is of concern and further compounds overcrowding in our ED.
Paradoxically, the implementation of screening to identify patients early who require
isolation or cohorting in hospital impacts negatively on the provision of emergency
care. Overcrowding in EDs has already been shown to increase ambulance diversions
to other units, delay treatments, increase waiting times and walk outs and lead to
longer lengths of hospital stay as well as increasing patient morbidity and mortality. <sup>7</sup>
The early identification and recognition of patients with potentially transmissible
diseases and their early isolation is desirable and appropriate as illustrated by the
SARS epidemic earlier this decade. However, it is neither desirable nor appropriate
that such patients have more prolonged stays in the most overcrowded part of an acute

1	hospital i.e. the ED. Vichard et al argue that "sepsis containment units" where patients
2	with MRSA can be isolated helps to prevent cross contamination, <sup>18</sup> but the ED can
3	not be expected to house patients for prolonged periods in the absence of isolation
4	rooms or cohort facilities on wards.
5	
6	Dantas et al noted that prolonged stay in the ED posed a risk for colonization and the
7	transmission of multidrug-resistant bacteria and for contracting HCAI, all associated
8	with increased mortality. 19 Cunningham et al documented that overcrowding and the
9	rapid turnover of patients in acute hospital settings contributes to cross-infection with
10	MRSA and they argue that adequate acute capacity would help to address this. 20 Borg
11	has described the correlation between workload indices and increased HCAI. <sup>21</sup>
12	
13	The results of the study reported here show that the selective screening of patients
14	with a prior diagnosis of MRSA colonisation prolongs their ED stay and increases the
15	workload of already busy ED nursing staff, potentially increasing MRSA
16	transmission. Reducing overcrowding in neonatal intensive care units has been shown
17	to be effective in controlling endemic MRSA spread <sup>22</sup> and it is plausible that reducing
18	ED overcrowding would have a similar positive effect.
19	
20	The delay in being admitted to a ward bed from the ED in this study has been shown
21	to be partly related to selective screening for MRSA. Being over 65 years old, being
22	female or having MRSA should not mean that the patient will have a longer wait in
23	the ED when they require acute admission. Being able to clarify the patients MRSA
24	status sooner would probably help to facilitate earlier transfer to a bed. PCR testing
25	for MRSA may be of benefit in this regard. Hospital ward staff are reluctant to accept
ļ	

1 patients from our ED with MRSA without there being an isolation room available **Deleted:** ontaminating the ward and other patients. 2 because of the risk of cross-infection. Clearly, leaving patients for prolonged periods 3 in the ED which is the most overcrowded part of any acute hospital is not appropriate. Deleted: cannot behave as though the f 4 Acute hospitals must accept that factors contributing to the spread of infection are **Deleted:** that increase Deleted: only 5 jmportant on in-patient wards and in the ED. Having identified the problem the Deleted: the Deleted: do not equally apply in the 6 research hospital has now allocated areas within each ward that allow for the setting 7 up of cohorts of patients with MRSA, if no single rooms are available. Another 8 alternative suggestion is to not selectively screen patients during the ED component of 9 their hospital stay if in so doing ward placement is made more difficult and to allow 10 patients to be screened on the wards. 11 12 Harbarth et al investigated the use of a universal rapid MRSA admission screening in 13 a surgical department and although they did not demonstrate a reduction in 14 nosocomial MRSA, they acknowledge that others have recommended universal admission screening as a means to control MRSA. 23-25 Robiecsek et al in their study 15 16 of universal admission MRSA surveillance with isolation and decolonization of 17 patients who tested positive for MRSA, found that this was associated with a greater 18 than 50% reduction in healthcare—associated MRSA bloodstream, respiratory, urinary Deleted: wound 19 tract and surgical site infections during admission and for up to 30 days after discharge.<sup>26</sup> 20 21 22 The early isolation of patients with MRSA is the standard in most centres but 23 adequate facilities must be available both on hospital wards and in EDs. We have 24 shown that selective screening of patients with a prior diagnosis of MRSA 25 colonisation in our hospital prolongs the wait for an acute hospital bed and

- 1 compounds ED overcrowding. National and local policies for MRSA control need to
- 2 address this by balancing the welfare of patients in the ED with the need to comply
- 3 with best practice when there are inadequate ED and in-patient isolation facilities.
- 4 Patients with MRSA requiring emergency admission to hospital must have a bed to go

5 to.

6

We are grateful to all the staff in the ED and the infection prevention and control team.

#### References

- 2 1. Smyth ETM, McIlvenny G, Enstone JE, *et al.* Four country healthcare associated infection prevalence survey 2006: overview of the results. *J Hosp Infect* 2008; **69**: 230-248.
- 5 2. Humphreys H, Newcombe RG, Enstone J, *et al.* Four country healthcare associated infection prevalence survey 2006: risk factor analysis. *J Hosp Infect* 2008; **69**: 249-257.
- 8 3. Cosgrove SE, Sakoulas G, Perencevich EN, Schwaber MJ, Karchmer AW, Carmeli Y. Comparison of mortality associated with methicillin-resistant and methicillin-susceptible *Staphylococcus aureus* bacteremia: A meta-analysis. *Clin Infect Dis* 2003; **36**: 53-9.
- 4. Humphreys H. National guidelines for the control and prevention of methicillinresistant *Staphylococcus aureus* –what do they tell us? *Clin Microbiol Infect* 2007; **13**: 846-853.
- 5. Dancer SJ. Considering the introduction of universal MRSA screening. *J Hosp Infect* 2008; 69: 315-320.
- 6. Moran GJ, Krishnadasan A, Gorwitz RJ, *et al.* Methicillin-resistant *S. aureus* infections among patients in the Emergency Department. *N Engl J Med* 2006; **355**; 666-74.
- 7. Gilligan P, Winder S, Singh I, Gupta V, Kelly P, Hegarty D. The boarders in the Emergency Department (BED) Study. *Emerg Med J* 2008; **25**: 265-269.
- 8. Gopal Rao G, Michalczk P, Nayeem N, Walker G, Wigmore L. Prevalence and Risk factors for methicillin-resistant *Staphylococcus aureus* in adult emergency admissions a case for screening all patients? *J Hosp Infect* 2007;**66**:15-21.
- Kevin Mackway-Jones on behalf of the Manchester Triage Group. Emergency
   Triage, Manchester Triage Group. *BMJ Publishing Group* 1997, London. ISBN 0-7279-1126-0.
- 28 10. Coia JE, Duckworth GJ, Edwards DI, *et al.* Guidelines for the control and prevention of meticillin-resistant Staphylococcus aureus (MRSA) in healthcare facilities. *J Hosp Infect* 2006;**635**:S1-S44.
- 11. Haddadin AS. Methicillin-resistant *Staphylococcus aureus* in the Intensive Care
   Unit. *Posatgrad Med J* 2002;**78**:385-92.
- 12. Kuehnert MJ, Kruszon- Moran D, Hill HA, et al. Prevalence of Staphylococcus
   aureus nasal colonisation in the United States, 2001-2002. J Infect Dis
   2006;193:172-179.
- 13. Jernigan JA, Titus MG, Groschel DH, Getchell-White S, Farr BM. Effectiveness
   of contact isolation during a hospital outbreak of methicillin-resistant
   Staphylococcus aureus. Am J Epidemiol. 1996;143:496-504.
- 14. Muto CA, Jernigan JA, Ostrowsky BE, et al. SHEA guideline for preventing
   nosocomial transmission of multidrug resistant strains of *Staphylococcus aureus* and enterococcus. *Infect Control Hosp Epidemiol* 2003;24:3622-386.
- 42 15. Department of Health. Screening for methicillin-resistant *Staphylococcus aureus* 43 (MRSA) colonization A strategy for NHS Trusts: Summary of best practice.
   44 England: Department of Health; November 2006.
- 16. Davis KA, Stewart JJ, Crouch HK, Florez CE, Hospenthal DR. Methicillin resistant *Staphylococcus aureus* (MRSA) nares colonization at hospital admission
- 47 and its effects on subsequent MRSA infection. Clin Infect Dis 2004; **39**:776-82.
- 48 17. Wallin TR, Hern HG, Frazee BW. Community-associated methicillin-resistant 49 Staphylococcus aureus. Emerg Med Clin North Am. 2008;**26**:431-55.

1 2

- 18. Vichard P, Talan D, Jeunet L. The importance of isolating septic patients in an autonomous unit in orthopaedic surgery and traumatology. *Bull Acad Natl Med* 2003;**187**:905-18.
- 19. Dantas SRPE, Moretti-Branchini ML. Impact of antibiotic-resistant pathogens colonizing the respiratory secretions of patients in an extended-care area of the Emergency Department. *Infect Control Hosp Epidemiol* 2003;**24**:351-355.
- 20. Cunningham JB, Kernahan WG, Sowney R. Bed occupancy and turnover interval as determinant factors in MRSA infections in acute settings in Northern Ireland: 1 April 2001 to 31 March 2003. *J Hosp Infect* 2005;**61**:189-193.
- 21. Borg MA. Bed occupancy and overcrowding as determinant factors in the incidence of MRSA infections within general ward settings. *J Hosp Infect* 2003; **54**:316-318.
- 22. Haley RW, Cushion NB, Tenover FC, et al. Eradication of endemic methicillinresistant *Staphylococcus aureus* infection from a neonatal intensive care unit. *J Infect Dis* 1995;**171**:614-624.
- 23. Harbarth S, Fankhauser C, Schrenzel J, et al. Universal screening for methicillinresistant *Staphylococcus aureus* at hospital admission and nosocomial infection in surgical patients. *JAMA*. 2008;**299**:1149-1157.
- 24. Weber SG, Huang SS, Oriola S, *et al.* Legislative mandates for use of active surveillance cultures to screen for methicillin resistant *Staphylococcus aureus* and vancomycin-resistant enterococci: position statement from the joint SHEA and APIC Task Force. *Infect Control Hosp Epidemiol* 2007;**28**:249-260.
  - 25. Diekma DJ, Edmond MB. Look before you leap: active surveillance for multidrug-resistant organisms. *Clin Infect Dis* 2007;44:1101-1107.
- 26. Robiecsek A, Beaumont JL, Paule SM, *et al.* Universal Surveillance for methicillin-resistant *Staphylococcus aureus* in 3 affiliated Hospitals. *Ann Intern Med* 2008;**148**:409-418.

 ED-MRSA Screening(3)

27-8-09

1 **Table 1.** Triage categories of admitted patients from the ED with and without MRSA.

$^{\circ}$	
1.	
_	

Category	Immediate	Very urgent	Urgent	Standard	Non-urgent
	(Red)	(Orange)	(Yellow)	(Green)	(Blue)
NRG	171	5936	8864	322	25
(%)	(1.1)	(38.8)	(57.9)	(2.1)	(.2)
RG	18	459	489	10	1
(%)	(1.8)	(47)	(50.1)	(1)	(.1)

<sup>3</sup> NRG = Non-risk Group (not previously MRSA positive),

<sup>4</sup> RG = Risk Group (previously MRSA positive).

 Table 2. Cox model of variables associated with prolonged ED stay from time of

arrival to time of hospital admission.

5

Variable	Hazard ratio from time of arrival to bed request [95% CI]		Hazard ratio from bed request to admission [95% CI]	
RG	0.946	[0.887 - 1.010]*	0.874	[0.819 - 0.933]**
Age > 65 years	0.891	[0.863 - 0.919]**	0.780	[0.756 - 0.805]**
Female Sex	0.902	[0.875 - 0.930]**	0.883	[0.857 - 0.911]**

RG, Risk group, i.e. previously MRSA positive p = 0.094 \*\* p < 0.001