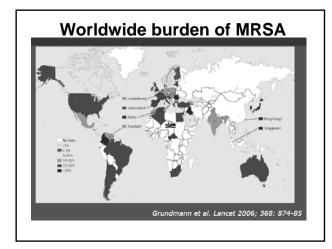
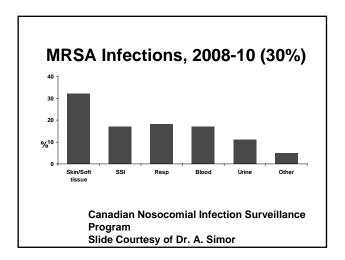
MRSA Pneumonia Manal Tadros PGY5 **Medical Microbiology Department** University of Toronto **Disclosures** None **OBJECTIVES** Describe the epidemiology & pathogenesis of MRSA pneumonia Present preliminary results of The **Canadian MRSA Pneumonia Outcome** Study (CaMPOS)





Epidemiology of MRSA pneumonia

- MRSA causes about 20%-40% of hospital-acquired pneumonia (HAP/HCAP) and ventilator-associated pneumonia (VAP) in US and other countries
- 1%-5% of Community associated pneumonia (CAP)
- Special concern about CA-MRSA strains: US 300/CMRSA-10

Mandell and Wundernik CID 2012:54;1134-36

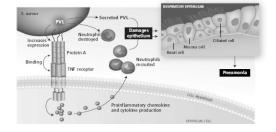
Epidemiology of MRSA pneumonia cont.

- US300/CMRSA-10 reported to cause severe pneumonia: high fever, cavitary lung lesions hypotension, and hemoptysis followed by rapid progression to septic shock and requirement for ventilator support
- High mortality rate (50%) was initially reported¹
- MR 37% in recent reports²

1 Dufour et al CID 2002 2 Haque et al JCM 2012

Dathogonocic of MARSA Decimalistics of Exponential growth phase (exponential growth phase) Secreted proteins (exponential growth phase) A Coagulase Enterotoxin B TSST-1 Colleges-binding protein Colleges-binding protein Exponential growth phase Colleges-binding protein Colleges-binding growth phase Consultationary pha

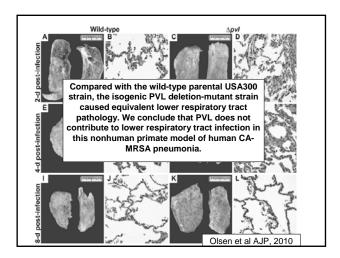
Panton- Valentine Leukocidin



Kahl BC, Peters G. Microbiology: mayhem in the lung. Science 2007; 315:182-1083

Lack of a Major Role of *Staphylococcus aureus* Panton-Valentine Leukocidin in Lower Respiratory Tract Infection in Nonhuman Primates

Olsen et al, *The American Journal of Pathology,* Vol. 176, No. 3, March 2010



Other Virulence Factors

- Alpha-Hemolysin¹
- Arginine Catabolic Mobile element²
- Accessory Gene Regulator³

1 Wardenburg et al Nat Med, 2007 2 Diep et al, JID,2008 3 Schewizer et al AAC, 2010

The Canadian MRSA Pneumonia Study (CaMPOS)

Study Objective:

 To determine the epidemiology, incidence, and outcome of MRSA pneumonia in adult patients admitted to Canadian hospitals

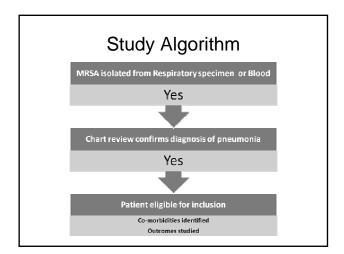
Sites:

- 1 year surveillance in eleven hospital sites from different areas across Canada.
- Funding:

Sponsored by Pfizer

PΙ

• Dr. A. E. Simor



Laboratory Investigations

The initial clinical (respiratory) or blood isolate of MRSA from each patient was analyzed for:

- Antimicrobial susceptibilities,
- molecular type by PFGE, SCC_{mec} type
- PVL gene detection.

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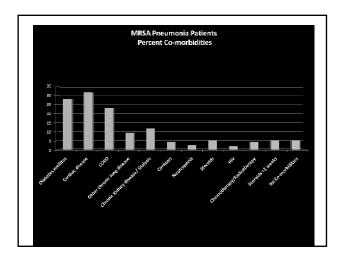
CaMPOS

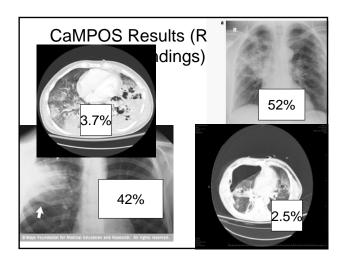
Results

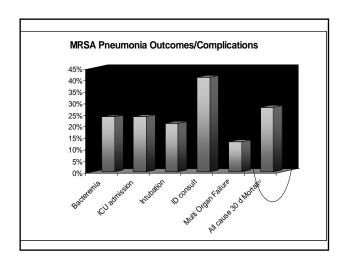
- 161 cases of MRSA pneumonia
- CAP= 45 (28%)
- HCAP=116 (72%)

36 ICU associated, 23 VAP

- Mean age was 64, SD 17.2
- Overall MRSA pneumonia rate was 0.47/10,000 patient-days
- HA- MRSA pneumonia rate was 0.33/10,000 patient days







Co-variates	Died at 30 d	Survived at	U	Univariate anlysis Multivariate		tivariate a	analysis	
	n=45	30 d n=116			-			-
			OR	95%CI	<i>p</i> value	OR	95%CI	pvalue
Age >65 n(%)	27(60)	57(49)	1.5	0.7-3.1	0.2	0.9	0.9-1.0	0.747
Males n (%)	30(66.7)	78(67.2)	0.9	0.4-2.0	1			
MRSA bacteremia n (%)	18(40)	20(17.2)	3.4	1.5-7.4	0.003	1.5	0.4-5.5	0.4
HCAP n (%)	36(80)	80(68.9)	1.8	0.7-4.1	0.17	1.4	0.3-7.0	0.6
CAP n (%)	9(20)	36(31.0)	0.5	0.2-1.2	0.17			
MRSA VAP n(%)	5(11.1)	18(15.5)	0.6	0.2-1.9	0.6			
ID consulted n(%)	17(37.8)	49 (42.2)	0.8	0.4-1.7	0.7			
COPD n(%)	14 (31.1)	23 (19.2)	1.8	0.8-3.9	0.14			
DM n (%)	14 (31.1)	31(26.7)	1.2	0.5-2.6	0.5			
Cirrhosis n(%)	5 (11.1)	2 (1.7)	7.1	1.3-38.1	0.01			
Steroids >2 weeks n(%)	2(4.4)	6(5.1)	0.8	0.1-4.3	1			
Multi-organ failure	16(35.5)	5(4.3)	12.2	4.1-36.2	<0.001	8.2	1.7-38.6	0.008
Appropriate empiric therapy used n(%) (Vanco/Linezolid)	15(33.3)	36 (31)	1.1	0.5-2.3	0.8			
Mean Vanco troughs (SD)	14.5(6.7)	14.3 (10.1)			0.9			
PFGE type CMRSA 10 n(%)	11(27)	29(28)	8.0	0.36-1.9	0.6			
PFGE type CMRSA 2 n(%)	26(63.4)	57(54.8)	1.4	0.6-3.0	0.3			
PVL+	11(27)	30(29)	0.9	0.4-2.0	$\overline{}$	2.4	0.5-10.7	0.2

Conclusions

- MRSA pneumonia rates in Canadian hospitals are relatively low
- This infection was associated with 28% 30-day mortality. Variables associated with increased mortality included the presence of bacteremia or cirrhosis.
- Microbial factors (such as PFGE type, PVL gene,) were not associated with increased mortality

THANK YOU				
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Dr. S. Haider	Infection Control Departments			

