

Evaluation of the Charlson Weighted Comorbidity Index and the MacCabe Classification as Predictors of Mortality in Patients with Nosocomial Bloodstream Infection due to *Pseudomonas aeruginosa*

ABSTRACT

Background: *Pseudomonas aeruginosa* bacteremia is associated with a high mortality rate. Many factors have been shown to increase the risk of mortality. However, the impact of underlying disease as measured by the Charlson weighted comorbidity index (WCI) and the MacCabe classification in patients with nosocomial bloodstream infection (nBSI) due to *P. aeruginosa* has not been evaluated.

Objective: To evaluate the utility of the Charlson index and the MacCabe classification as predictors of mortality in patients with *P. aeruginosa* nBSI.

Methods: Charlson and MacCabe scores were assessed for a cohort of 57 patients with *P. aeruginosa* nBSI. Charlson WCI was dichotomized into scores of <3 and ≥3 points. The MacCabe classification was dichotomized into non-fatal vs. ultimately fatal and rapidly fatal diseases. Univariate analysis was performed to evaluate the association of the following variables with mortality: Charlson index, MacCabe classification, age, gender, BSI acquired in the intensive care unit, appropriate antimicrobial therapy, and cardiovascular, respiratory, renal, hematologic and hepatic failure. Variables significant in univariate analysis were entered into a logistic regression model. Due to the collinearity between these two comorbidity variables, separate stepwise logistic regression (SLR) models were used to assess independent predictors for mortality.

Results: In univariate analysis, the Charlson WCI and MacCabe classification were able to predict overall mortality, OR 4.46 (CI95 1.3-15.2, p=0.014) and OR 3.96 (CI95 1.3-11.9, p=0.013), respectively. In the first model (Charlson WCI) hematologic failure (OR 7.5; CI95 2.04-71.8, p=0.006) and cardiovascular failure (OR 5.5; CI95 1.3-19.5, p=0.017) were found to be independently predictive of mortality in patients with *P. aeruginosa* BSI. Similarly, in the second model (MacCabe classification), variables independently related to death were the development of hematologic failure (OR 6.6; CI95 1.7-53.9, p=0.010) and cardiovascular failure (OR=6.1, CI95 1.4-20.6, p=0.014).

Conclusions: The development of hematologic and cardiovascular failure were independent predictors of death. Neither the Charlson index nor the MacCabe classification predicts mortality in patients with *P. aeruginosa* nBSI, suggesting that underlying disease processes are less important than the development of associated end organ dysfunction in predicting survival.

INTRODUCTION

Pseudomonas aeruginosa remains an important pathogen in community and hospital settings. It ranks among the top ten pathogens causing bloodstream infections (BSI) in the United States and Canada, and the attributable mortality of *P. aeruginosa* bacteremia is as high as 34%.

Many factors have been shown to increase the risk of mortality. However, the impact of underlying disease as measured by the Charlson weighted comorbidity index (WCI) and the MacCabe classification in patients with nosocomial bloodstream infection (nBSI) due to *P. aeruginosa* has not been evaluated. The objective of our study was to evaluate the utility of the Charlson index and the MacCabe classification as predictors of mortality in patients with *P. aeruginosa*

METHODS

Setting: The Virginia Commonwealth University Medical Center is a 820-bed tertiary care facility in Richmond, Virginia. The hospital houses 9 intensive care units and a burn unit; approximately 30,000 patients are admitted annually.

Study design: Charlson and MacCabe scores were assessed for a cohort of 57 adults with *P. aeruginosa* nBSI. Charlson WCI was dichotomized into scores of <3 and ≥3 points. The MacCabe classification was dichotomized into non-fatal vs. ultimately fatal and rapidly fatal diseases.

Statistical methods: Mean values were compared using 2 sample t tests for independent samples. Proportions were compared using a χ^2 test. All tests of significance were 2-tailed, and α was set at 0.05. Independent predictors of mortality were identified by means of stepwise logistic regression analysis, using variables found to be significant in univariate analysis. Due to the collinearity between the Charlson WCI and the MacCabe classification, separate stepwise logistic regression (SLR) models were used to assess independent predictors for mortality. To also estimate the hazard ratio of progression to death for these two comorbidity variables, we used the multivariable-adjusted Cox proportional hazard regression model.

RESULTS

Table 1: The Charlson Weighted Index of Comorbidity

Assigned Weight for Disease	Condition
1	Myocardial infarct
	Congestive heart failure
	Peripheral vascular disease
	Cerebrovascular disease
	Dementia
	Chronic pulmonary disease
	Connective tissue disease
	Ulcer disease
	Mild liver disease
	Diabetes
2	Hemiplegia
	Moderate or severe renal disease
	Diabetes with end organ damage
	Any tumor
	Leukemia
3	Lymphoma
6	Moderate or severe liver disease
	Metastatic solid tumor
	AIDS

Table 2: The MacCabe Classification

Class	Prognosis
1	None or Non-fatal disease
2	Ultimately fatal disease
3	Rapidly fatal disease

Figure 1: Survival in cohorts stratified by the dichotomized Charlson WCI

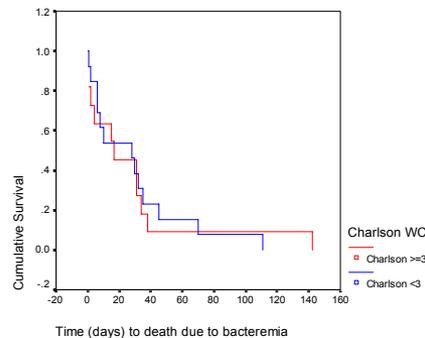
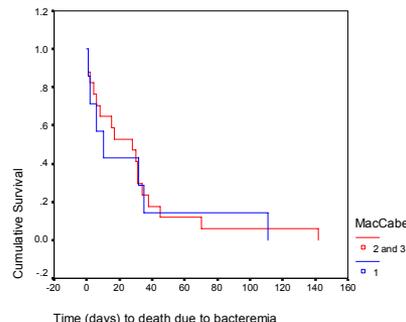


Figure 2: Survival in cohorts stratified by the dichotomized MacCabe classification



RESULTS (continued)

Table 3: Risk factors for death in patients with *P. aeruginosa* nosocomial bloodstream infection, considering Charlson WCI

Risk factor	Univariate Analysis*		Multivariate Analysis*	
	OR	P	OR	P
Hematologic failure	12.4	0.001	7.5	0.001
Cardiovascular failure	5.5	0.003	5.7	0.003
Charlson WCI ≥3	4.46	0.014	3.6	0.014

*Variables not significant in univariate analysis are not shown

Table 4: Risk factors for death in patients with *P. aeruginosa* nosocomial bloodstream infection, considering MacCabe classification

Risk factor	Univariate Analysis*		Multivariate Analysis*	
	OR	P	OR	P
Hematologic failure	12.4	0.001	6.6	0.001
Cardiovascular failure	5.5	0.003	6.1	0.003
MacCabe 2 and 3	3.96	0.013	3.7	0.013

*Variables not significant in univariate analysis are not shown

CONCLUSIONS

- The development of hematologic and cardiovascular failure were independent predictors of death.
- Neither the Charlson index nor the MacCabe classification predicted mortality in patients with *P. aeruginosa* nBSI, suggesting that underlying disease processes are less important than the development of BSI-associated end organ dysfunction in predicting survival.



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