

# Clinical Outcomes Associated With Procalcitonin Algorithms to Guide Antibiotic Therapy in Respiratory Tract Infections

**Clinical Question** In patients with respiratory tract infection, is measurement of procalcitonin to guide antibiotic prescriptions associated with reduced antibiotic exposure without increases in all-cause mortality or treatment failure?

**Bottom Line** The measurement of procalcitonin to guide initiation and duration of antibiotic treatment in patients with respiratory tract infections of varying severity is associated with lower antibiotic exposure without increasing all-cause mortality or treatment failure.

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THE MOST FREQUENT INDICATION for antibiotic use is respiratory tract infections, ranging in severity from self-limited acute bronchitis to life-threatening bacterial pneumonia.<sup>1</sup> Antibiotic prescriptions for nonpneumonia respiratory tract infections and prolonged courses for pneumonia contribute to antibiotic overuse, which potentially leads to the development of antibiotic resistance and increased risk of *Clostridium difficile* infection.

Procalcitonin, a biomarker for bacterial infections, is released ubiquitously in response to bacterial toxins and bacteria-specific proinflammatory mediators.<sup>2</sup> Previous trials investigated the effects of procalcitonin al-

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**Table.** Summary of Results for Measuring Procalcitonin to Guide Antibiotic Use in Respiratory Tract Infections<sup>a</sup>

	No. of Trials	Procalcitonin Group (n)	Control Group (n)	Adjusted Odds Ratio or Mean Difference (95% CI)
All patients with respiratory tract infections	14	(n = 2085)	(n = 2126)	
Mortality, No. (%)		118 (5.7)	134 (6.3)	0.94 (0.71 to 1.23)
Treatment failure, No. (%)		398 (19.1)	466 (21.9)	0.82 (0.71 to 0.97) <sup>c</sup>
Total antibiotic exposure, median (IQR), d		4 (0 to 8)	8 (5 to 12)	-3.47 (-3.78 to -3.17) <sup>b</sup>
Patients treated in primary care	2	(n = 507)	(n = 501)	
Mortality, No. (%)		0	1 (0.2)	NA
Treatment failure, No. (%)		159 (31.4)	164 (32.7)	0.95 (0.73 to 1.24)
Total antibiotic exposure, median (IQR), d		0 (0 to 0)	6 (0 to 7)	-3.06 (-3.48 to -2.65) <sup>b</sup>
Patients treated in the emergency department	7	(n = 1291)	(n = 1314)	
Mortality, No. (%)		61 (4.7)	59 (4.5)	1.03 (0.7 to 1.5)
Treatment failure, No. (%)		182 (14.1)	228 (17.4)	0.76 (0.61 to 0.95) <sup>c</sup>
Total antibiotic exposure, median (IQR), d		5 (0 to 8)	9 (5 to 12)	-2.96 (-3.38 to -2.54) <sup>b</sup>
Patients treated in the intensive care unit	5	(n = 287)	(n = 311)	
Mortality, No. (%)		57 (19.9)	74 (23.8)	0.84 (0.54 to 1.31)
Treatment failure, No. (%)		57 (19.9)	74 (23.8)	0.84 (0.54 to 1.31)
Total antibiotic exposure, median (IQR), d		8 (5 to 15)	12 (8 to 18)	-3.21 (-4.32 to -2.10) <sup>b</sup>
Patients with community-acquired pneumonia	12	(n = 999)	(n = 1028)	
Mortality, No. (%)		92 (9.2)	111 (10.8)	0.89 (0.64 to 1.23)
Treatment failure, No. (%)		190 (19.0)	240 (23.4)	0.77 (0.62 to 0.96) <sup>c</sup>
Total antibiotic exposure, median (IQR), d		6 (4 to 10)	10 (8 to 14)	-3.98 (-4.44 to -3.52) <sup>b</sup>
Patients with exacerbation of COPD	6	(n = 288)	(n = 296)	
Mortality, No. (%)		9 (3.1)	8 (2.7)	1.15 (0.43 to 3.09)
Treatment failure, No. (%)		35 (13.7)	45 (15.2)	0.75 (0.46 to 1.22)
Total antibiotic exposure, median (IQR), d		0 (0 to 6)	7 (0 to 10)	-3.03 (-3.76 to -2.3) <sup>b</sup>

Abbreviations: COPD, chronic obstructive pulmonary disease; IQR, interquartile range; NA, not available.

<sup>a</sup>Source: Schuetz P, Müller B, Christ-Crain M, et al. Procalcitonin to initiate or discontinue antibiotics in acute respiratory tract infections. *Cochrane Database Syst Rev*. 2012;9:CD007498. doi:10.1002/14651858.CD007498.pub2

<sup>b</sup>P < .01.

<sup>c</sup>P < .05.

**Evidence Profile**

No. of randomized trials: 14

Study years: 2004-2011

No. of patients: 4211 with respiratory infections of varying severity, including upper respiratory tract infections, bronchitis, exacerbation of chronic obstructive pulmonary disease (COPD), community-acquired pneumonia (CAP), and ventilator-associated pneumonia

Men: 2282 (54%) Women: 1929 (46%)

Race/ethnicity: Unavailable

Age, mean (SD): 59.8 (19.2) years

Settings: Inpatient and outpatient, medical and surgical intensive care units (ICUs)

Countries: United States, Germany, Switzerland, France, Denmark, China

Comparison: Antibiotic therapy with vs without guidance from procalcitonin levels

Primary outcomes: (1) All-cause 30-day mortality and (2) treatment failure within 30 days defined as death, ICU admission, hospitalization, respiratory infection-specific complications, recurrent or worsening infection, and patients reporting any symptoms of ongoing respiratory tract infection at 30 days

Secondary outcome: Total antibiotic exposure at 30 days

gorithms for guiding antibiotic use.<sup>3</sup> These algorithms recommended antibiotic prescription only if procalcitonin levels are elevated and recommended antibiotic cessation when procalcitonin levels normalize. Yet the safety and efficacy of procalcitonin algorithms have not been conclusively demonstrated.

**SUMMARY OF FINDINGS**

There was no increase in all-cause mortality associated with procalcitonin testing for any clinical setting or any type of respiratory tract infection (TABLE). Treatment failure was lower for procalcitonin-guided patients treated in the emergency department (odds ratio [OR], 0.76 [95% CI, 0.61-0.95]) and in patients with CAP (OR, 0.77 [95% CI,

0.62-0.96]). Patients for whom procalcitonin was used to guide antibiotic therapy had lower antibiotic exposure, which was mainly due to lower antibiotic prescription rates for exacerbation of COPD (48% vs 73%) and bronchitis (24% vs 66%). Patients with measurement of procalcitonin also had shorter antibiotic courses in the emergency department (median, 7 days [IQR, 4-10] vs 10 days [IQR, 7-13]), in the ICU (median, 8 days [IQR, 5-15] vs 12 days [IQR, 8-18]), and with a diagnosis of CAP (median, 7 days [IQR, 5-10] vs 10 days [IQR, 8-14]).

**COMMENT**

Among 14 trials and 4211 patients with respiratory tract infections, measurement of procalcitonin to guide antibiotic therapy was not associated with increased mortality or treatment failure.<sup>4</sup> Results suggest that using procalcitonin to guide antibiotic therapy is associated with fewer antibiotic prescriptions in primary care settings and in patients with less severe infections, and with shorter courses of antibiotics for patients evaluated in the emergency department or the critical care setting. The lower risk of treatment failure associated with procalcitonin testing may in part relate to lower antibiotic exposure, but this needs further exploration.

**Limitations**

The randomized trials were not blinded in regard to allocation and outcome assessment. Adherence to procalcitonin algorithms was variable and lower in the critical care setting (between 47% and 81%). Included trials were mostly conducted in Europe (78%). Two trials were conducted in China and one multinational trial was conducted in the United States. Immunosuppressed patients, children, and patients with non-respiratory infections were not included.

**Comparison of Findings With Current Guidelines**

There are currently no practice guidelines in the United States regarding

measurement of procalcitonin to guide antibiotic therapy. The 2011 European Respiratory Society guidelines recommended using procalcitonin to guide antibiotic treatment duration in CAP, even in severe cases.<sup>5</sup> The 2012 international Surviving Sepsis Campaign guidelines emphasize procalcitonin for stopping antibiotics.

**Areas in Need of Future Study**

Further studies are needed to determine the safety of procalcitonin algorithms in critical care settings and for guiding treatment of nonrespiratory infections. Future trials should evaluate the cost-effectiveness of procalcitonin algorithms by considering country-specific costs of procalcitonin measurements and potential savings and health benefits by reducing antibiotic prescription rates.<sup>6,7</sup>

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