

## **Outcomes and risk factors for mortality in community-onset bacteremia caused by extended-spectrum beta-lactamase-producing *Escherichia coli*, with a special emphasis on antimicrobial therapy.**

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### **Abstract**

#### **BACKGROUND:**

Although extended-spectrum beta-lactamase (ESBL)-producing *Escherichia coli* has emerged as a significant pathogen, there is little information regarding treatment outcomes in community-onset bacteremia due to ESBL *E. coli*. The purpose of this study was to evaluate treatment outcomes of community-onset bacteremia caused by ESBL-producing *E. coli* and the factors associated with mortality.

#### **METHODS:**

A retrospective cohort study was performed, including 92 adult patients with community-onset bacteremia caused by ESBL-producing *E. coli*.

#### **RESULTS:**

The 30-day mortality rate was 10.9% (10/92). Independent risk factors for mortality were underlying liver disease and severity of illness (e.g., high Pitt bacteremia score, the presence of severe sepsis or septic shock;  $p < 0.05$ ). Mortality in patients receiving inappropriate initial antimicrobial therapy was not significantly higher than mortality in those receiving appropriate empirical antimicrobial therapy (10.9 vs 10.7%;  $p = 0.975$ ), if antimicrobial therapy was adjusted appropriately according to susceptibility results. Carbapenems, piperacillin/tazobactam, fluoroquinolones, and amikacin were the most effective antibiotics for community-onset bacteremia caused by ESBL-producing *E. coli*, although susceptibility profiles confirmed that alternatives to carbapenems are limited. Of 68 isolates in which the ESBLs and their molecular relationships were studied, all isolates produced ESBLs from the CTX-M family (CTX-M-14, 30 isolates; CTX-M-15, 22; and other CTX-M, 16).

#### **CONCLUSIONS:**

In patients with community-onset bacteremia caused by ESBL-producing *E. coli*, severe sepsis and underlying liver disease were significantly associated with mortality, and a delay in appropriate antimicrobial therapy was not associated with a higher mortality if therapy was adjusted appropriately according to the susceptibility results.

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