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The supplementation of L-carnitine in critically ill patients with sepsis

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ADMINISTRATIVE INFORMATION

Support - None.

Review Stage at time of this submission - Completed but not published.

Conflicts of interest - None declared.

INPLASY registration number: INPLASY202460086

Amendments - This protocol was registered with the International Platform of Registered Systematic Review and Meta-Analysis Protocols (INPLASY) on 23 June 2024 and was last updated on 23 June 2024.

INTRODUCTION

Review question / Objective L-carnitine may reduce mortality in critically ill patients with sepsis, but the conclusion is inconsistent in different studies. We conducted a meta-analysis to evaluate the effect of L-carnitine compliance on mortality in patients with sepsis.

Condition being studied 28-day mortality and 12month mortality.

METHODS

Participant or population Adult patients diagnosed with sepsis.

Intervention L-carnitine or placebo.

Comparator Placebo.

Study designs to be included The search strategy was RCTs.

Eligibility criteria (1) Adult patients diagnosed with sepsis according to Sepsis 3.0. (2) Patients taking L-carnitine or placebo. (3) Outcome indicators: 28-day mortality and 12-month mortality.

Information sources We will search the references in the included trials and personal files. We will request advice from experts in the field. In addition, we will search associated articles from meetings, and contacted the authors of included trials, if need.

Main outcome(s) 28-day mortality and 12-month mortality.

Quality assessment / Risk of bias analysis We evaluated the methodological quality of the individual studies using the Cochrane risk of bias tool for RCTs.

Strategy of data synthesis We will consider using the number of participants and deaths between different groups for analysis.

Subgroup analysis None.

Sensitivity analysis China.

Country(ies) involved China.

Keywords L-carnitine; critical ill; sepsis.

Contributions of each author

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