



**Epinephrine for in-hospital LUCAS-CPR:  
a predictor of outcome?**

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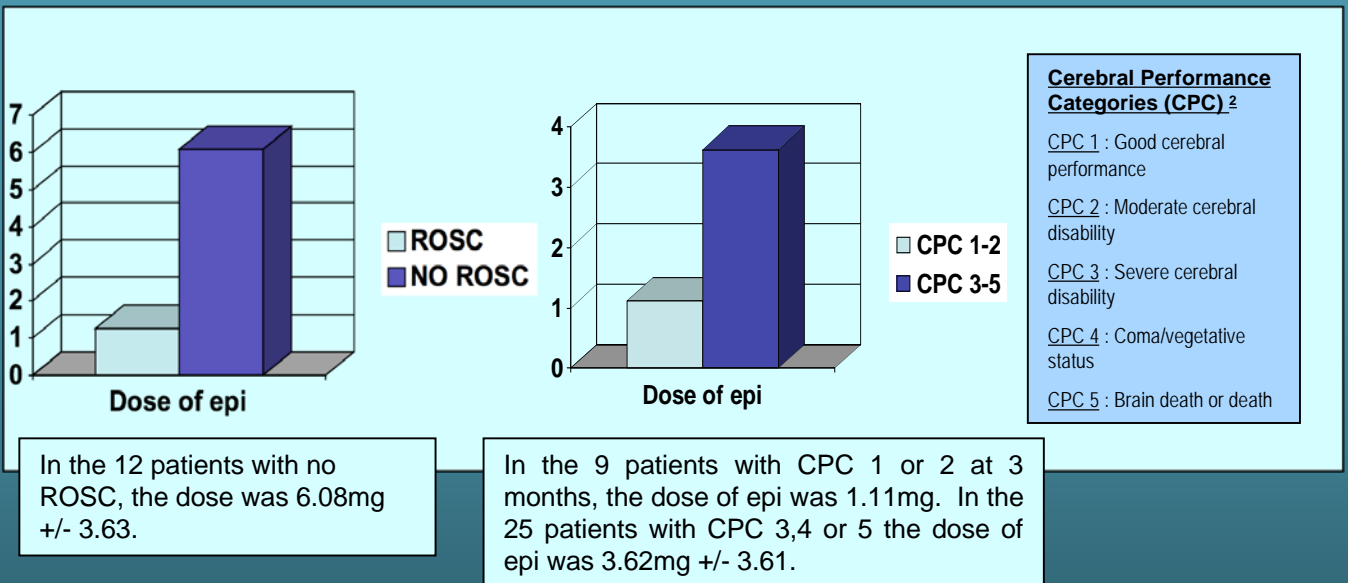


**Introduction:** The role of epinephrine in cardiac arrest remains debated<sup>1</sup>. We compare the doses of epinephrine administered in patients with good versus bad outcome after in-hospital LUCAS-CPR.



**Methods:** From February until June 2006, LUCAS-CPR was used for all cases of adult in-hospital arrest, after arrival of the in-hospital emergency team. Outcome parameters such as ROSC, Cerebral Performance Categories<sup>2</sup> ( CPC ) at 3 months and administered doses of epinephrine were recorded. CPC 1 or 2 at 3 months were considered good outcome. Epinephrine was administered per milligram to all patients during CPR every 3 to 5 minutes according to the guidelines 2005 and at the discretion of the attending physician. Results are presented as means +/- standard deviation.

**Results:** 35 patients received in hospital LUCAS-CPR. In one patient with no ROSC, the dose of epinephrine remains unknown. In the 22 patients with ROSC, 1.25mg +/- 1.25 of epinephrine was used. In the 12 patients with no ROSC, the dose was 6.08mg +/- 3.63. In the 9 patients with CPC 1 or 2 at 3 months, the dose of epi was 1.11mg +/- 0.78. None of these patients received more than 2 mg of epinephrine. In the 25 patients with CPC 3,4 or 5 the dose of epi was 3.62mg +/- 3.61.



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In the 9 patients with CPC 1 or 2 at 3 months, the dose of epi was 1.11mg. In the 25 patients with CPC 3,4 or 5 the dose of epi was 3.62mg +/- 3.61.

**Conclusion:** The total dose of epinephrine used is inversely related to outcome. No good outcome was achieved in patients who needed more than 2 mg of epinephrine during in hospital LUCAS-CPR.

**References:**  
 1. European Resuscitation Council Guidelines for Resuscitation 2005; Resuscitation ( 2005 ) 67S; S47  
 2. The Brain Resuscitation Clinical Trial II Study Group. A randomized clinical trial of calcium entry blocker administration to comatose survivors of cardiac arrest: design, methods, and patient characteristics. Control Clinical Trials 1991; 12: 525-45