High levels of atrial natriuretic peptide and copeptin and mortality risk

[Asociación de valores elevados de péptido natriurético auricular y copeptina con riesgo de mortalidad]

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Objective To determine whether high levels of mid-regional pro-atrial natriuretic peptide (MR-proANP), copeptin, and procalcitonin (PCT) plasma concentrations are associated with increased mortality risk. Methods Prospective observational study including 254 critically ill children. MR-proANP, copeptin and PCT were compared between children with high (Group A; n = 33) and low (Group B; n = 221) mortality risk, and between patients with failure of more than 1 organ (Group 1; n = 71) and less than 2 (Group 2; n = 183). Results Median (range) of MR-proANP, copeptin, and PCT levels in group A vs B were, respectively: 209.4 (30.5?1415.8) vs. 75.0 (14.6?867.2) pmol/L (P<.001); 104.4 (7.4?460.9) vs. 26.6 (0.00?613.1) pmol/L (P<.001), and 7.8 (0.3?552.0) vs. 0.3 (0.02?107.0) ng/mL (P<.001). The area under the curve (AUC) for the differentiation of group A and B was 0.764 (95% CI: 0.674-0.854) for MR-proANP; 0.735 (0.642-0.827) for copeptin, and 0.842 (0.744-0.941) for PCT, with no statistical differences. The AUCs for the differentiation of group 1 and 2 were: 0.837 (0.784-0.891) for MR-proANP, 0.735 (0.666-0.804) for copeptin, and 0.804 (0.715-0.892) for PCT, with statistical differences between MR-proANP and copeptin, P=.01. Conclusions High levels of MR-proANP, copeptin and PCT were associated with increased mortality risk scores. MR-proANP showed a higher association than copeptin with number of organs in failure.

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Atrial natriuretic peptide
Copeptin
Critically ill children
Mortality risk
Organ failure
Procalcitonin
Prognosis
atrial natriuretic factor
biological marker
calcitonin
copeptin
glycopeptide
adolescent
blood
child
critical illness
female
human
infant
male
mortality
multiple organ failure
newborn
preschool child
prognosis

risk assessment
Adolescent
Atrial Natriuretic Factor
Biomarkers
Calcitonin
Child
Child, Preschool
Critical Illness
Female
Glycopeptides
Humans
Infant
Infant, Newborn
Male
Multiple Organ Failure
Prognosis
Prospective Studies
Risk Assessment

prospective study