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Objective To describe our preliminary experience with Levosimendan during the last 4 years, a new calcium-sensitizing agent in critically unwell infants and children with severe heart failure.

Design Retrospective cohort analysis.

Setting Pediatric cardiology intensive care unit.

Patients 8 children aged 2.5 months to 13 yrs (median age 44 months) with severe myocardial dysfunction secondary to endstage heart failure who were inotropic dependend (requiring at least one catecholamine).

Interventions A single dose (continuous intravenous infusion over 24 hrs) of Levosimendan was given under continuous hemodynamic monitoring in our intensive care unit.

Six children received a single dose, two children received two doses.

Echocardiographic assessments of ventricular function were made before and 3–5 days after Levosimendan infusion.

Measurements and Main Results Heart rate, systolic pressure, diastolic pressure, mean blood pressure, shortening fraction, the dose of inotrope at the beginning of levosimendan infusion, at 24 hours and 36 hours, ECG result 24 hour after levosimendan infusion.

Conclusions Levosimendan appeared to be a safe and efficacious drug when given to children with uncompensated end-stage heart failure in this size-limited sample. It warrants formal prospective large-cohort evaluation and multicenter trial to determine its safety profile and clinical application in the pediatric population.

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Background and Aims To determine whether there is an association between platelet counts and patent ductus arteriosus (PDA) incidence and/or closure in preterm newborns.

Methods Premature infants with hemodynamically significant PDA (n=154) and a control group without PDA (n=207) who were hospitalized in the NICU were eligible. Platelet counts and other platelet indices including mean platelet volume (MPV) and platelet distribution width (PDW) were obtained before treatment and after documented ductal closure. Pre and posttreatment cerebral TOI and SaO2 were recorded.

Results Fourteen newborns were included, abdominal recordings were available in 13. Mean BW and GA were 1089 g and 29 weeks respectively. No change was observed in cerebral or abdominal tissue oxygenation and oxygen extraction before and after medical closure of PDA. Pre and posttreatment cerebral TOI values (median and range) were 67.17 (50.9–89.1) and 64.35 (54.9–87.4) P<0.05, and pre and posttreatment cerebral FTOE values were 0.3 (0.03–0.45) and 0.29 (0.05–0.42) P<0.05 respectively. Pre and posttreatment abdominal TOI values were (median and range) 53.9 (40.1–62.9) and 50.29 (39.2–78.5) P=0.7, pre and posttreatment abdominal FTOE values were 0.44 (0.32–0.59) and 0.46 (0.2–0.6) P<0.05 respectively.

Conclusion Results of this small group may suggest that cerebral and abdominal tissue oxygenation is preserved during hemodynamically significant PDA, however more detailed studies are warranted.


Background and Aim Patent ductus arteriosus (PDA) is a frequent problem in pretermers known to have significant effects on organ perfusion. The aim of this study was to investigate the difference between cerebral and abdominal tissue oxygenation index (TOI) measured by near infrared spectroscopy (NIRS) before and after treatment of hemodynamically significant PDA in preterm newborns.

Methods Cerebral and abdominal TOI were recorded by NIRS (NIRO 200 Hamamatsu, Japan) in preterm newborns with hemodynamically significant PDA requiring ibuprofen treatment. Newborns with congenital anomalies were excluded. 20 minute recordings were obtained before treatment and after documented ductal closure by echocardiography as well as real time oxygen saturation (SaO2) monitoring by pulse oxymetry. Fractional tissue oxygen extraction (FTEO) was calculated using TOI and SaO2.

Results Median gestational age and birth weight of the infants were 28 (range 26–29) weeks and 1060 (range 892–1250) gr respectively. No change was observed in cerebral or abdominal tissue oxygenation and oxygen extraction before and after medical closure of PDA. Pre and posttreatment cerebral TOI values (median and range) were 67.17 (50.9–89.1) and 64.35 (54.9–87.4) P<0.05, and pre and posttreatment cerebral FTOE values were 0.3 (0.03–0.45) and 0.29 (0.05–0.42) P<0.05 respectively. Pre and posttreatment abdominal TOI values were (median and range) 53.9 (40.1–62.9) and 50.29 (39.2–78.5) P=0.7, pre and posttreatment abdominal FTOE values were 0.44 (0.32–0.59) and 0.46 (0.2–0.6) P<0.05 respectively.

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Background and Aim In this retrospective cohort study all infants born at a gestational age of less than 32 weeks were evaluated. Maternal, fetal and infant factors associated with prenatal and perinatal hypoxia-ischemia were related to BNP levels after birth. Pathologic examination of the placenta was routinely performed.


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