MEETING ABSTRACTS

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Basic Science

P1
Bone morphogenetic protein-2 and leptin but not endothelin-1 induce osteochondrogenesis through increasing oxidative stress in vascular smooth muscle cells
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Introduction Vascular calcification is a regulated process, which associates with coronary artery disease (CAD) and occurs through an increase in transcription factor expression such as RUNX2, MSX2 and alkaline phosphatase (ALP), then inducing calcium deposition. Bone morphogenetic protein-2 (BMP2) is a potent osteochondrogenic mediator, which is expressed in CAD. Endothelin-1 (ET1) and leptin have a role in regulating inflammation and CAD. We hypothesized that BMP2, leptin or both increase ROS formation in C57BL/6 vascular smooth muscle cells (SMC), stimulating osteochondrogenic differentiation. We also investigated the effect of ET1 in SMC osteochondrogenesis. Our objectives were to investigate ROS production in SMC after BMP2 (50 ng/ml) and/or leptin (10 ng/ml) incubation for 6 hours; and to assess osteochondrogenic gene expression and calcification of SMC stimulated with BMP2, leptin or ET1 (10 nM).

Methods We assessed 2-hydroxyethidium, more specific for superoxide, and ethidium which reflects hydrogen peroxide through HPLC analysis in SMC after stimulation. SMC cells were incubated with these stimuli for 48 to 96 hours and RUNX2, MSX2, ALP mRNA and protein expression were assessed using qPCR and western blotting. We quantified SMC calcification after 14 days of stimulation through Alizarin Red staining.

Results The results are shown as mean ± SD and were statistically significant when pHydrogen peroxide and superoxide production increased both in BMP2 and in leptin-incubated SMC (3.77 ± 0.32 and 3.26 ± 0.26) versus control (n = 6); pBMP2 and leptin alone increased SMC calcification (1.25 ± 0.08 and 1.28 ± 0.14) versus control after 14 days (n = 6); pET1 alone did not stimulate osteochondrogenic mRNA expression vs. control.

Conclusion We showed that BMP2 and leptin increased ROS formation in SMC, which stimulated osteochondrogenic mRNA/protein expression to induce SMC calcification. ET1 alone did not increase osteochondrogenesis in SMC.

P2
Effects of rapid repetition of a vascular occlusion test on near-infrared spectroscopy-derived variables in healthy subjects and in critically ill patients
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Introduction Transient ischemia modifies cellular metabolism and microvascular physiology in order to limit damage from future hypoxic episodes, a phenomenon called preconditioning. Near-infrared spectroscopy (NIRS) is a non-invasive technique that, when coupled to a vascular occlusion test (VOT), provides an indirect measurement of muscle oxygen consumption (VO2) and microvascular reactivity. We hypothesized that: rapid repetition of a VOT may alter VOT-induced NIRS-derived variables and these changes could reflect preconditioning; and these alterations would be different in healthy volunteers and critically ill patients.

Methods Continuous non-invasive measurements of thenar tissue oxygen saturation (StO2) were performed using NIRS technology (InSpectra 650; Hutchinson, USA). VOTs were performed by inflating a cuff to 50 mmHg above the systolic pressure for 3 minutes. In a group of healthy volunteers, the VOT was repeated after 5 minutes on day 1, after 15 minutes on day 2 and after 30 minutes on day 3. In a group of critically ill patients, the VOT was repeated after 5 minutes. For each VOT, we calculated the StO2 desaturation slope (DescSlope), StO2 resaturation slope (AscSlope) and the NIRS VO2 as the DescSlope × mean total hemoglobin index over the occlusion time. All statistical analyses were performed using SPSS 19.0 (IBM, USA).

Results Twenty-one healthy volunteers (age 29 ± 6 years, heart rate 71 ± 6 bpm, mean arterial pressure 82 ± 6 mmHg) and 18 critically ill patients (age 59 ± 14 years, APACHE II score 21 ± 9, norepinephrine use in 10/18, ICU mortality 22%) were included. In the healthy volunteers, repetition of the VOT was associated with a decrease in the DescSlope and in NIRS VO2. This effect was not observed in the critically ill patients (Tables 1 and 2).

Table 1 (abstract P2). Effects of a repeat VOT on VOT-induced NIRS-derived variables in healthy volunteers

<table>
<thead>
<tr>
<th>Interval</th>
<th>Variable</th>
<th>First</th>
<th>Second</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>5 minutes</td>
<td>AscSlope</td>
<td>4.2 (3.4 to 4.9)</td>
<td>4 (3.3 to 5)</td>
<td>0.298</td>
</tr>
<tr>
<td></td>
<td>DescSlope</td>
<td>12 (9.2 to 14.3)</td>
<td>9 (8.5 to 10.8)</td>
<td>&gt;0.001</td>
</tr>
<tr>
<td></td>
<td>NIRS VO2</td>
<td>151 (132 to 171)</td>
<td>131 (118 to 146)</td>
<td>0.001</td>
</tr>
<tr>
<td>15 minutes</td>
<td>AscSlope</td>
<td>4 (3.2 to 5.2)</td>
<td>4.1 (3.4 to 5)</td>
<td>0.676</td>
</tr>
<tr>
<td></td>
<td>DescSlope</td>
<td>10.3 (9.6 to 11.3)</td>
<td>9.4 (8.3 to 10.2)</td>
<td>0.003</td>
</tr>
<tr>
<td></td>
<td>NIRS VO2</td>
<td>153 (141 to 165)</td>
<td>141 (120 to 146)</td>
<td>0.005</td>
</tr>
<tr>
<td>30 minutes</td>
<td>AscSlope</td>
<td>4.2 (3.6 to 5.3)</td>
<td>3.4 (3.1 to 4.8)</td>
<td>0.006</td>
</tr>
<tr>
<td></td>
<td>DescSlope</td>
<td>10.9 (9.5 to 12.6)</td>
<td>9 (7.4 to 10.4)</td>
<td>&gt;0.001</td>
</tr>
<tr>
<td></td>
<td>NIRS VO2</td>
<td>157 (122 to 171)</td>
<td>132 (112 to 152)</td>
<td>&gt;0.001</td>
</tr>
</tbody>
</table>

Table 2 (abstract P2). Effects of a repeat VOT on VOT-induced NIRS-derived variables in critically ill patients

<table>
<thead>
<tr>
<th>Interval</th>
<th>Variable</th>
<th>First</th>
<th>Second</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>5 minutes</td>
<td>AscSlope</td>
<td>3.6 (2.7 to 4)</td>
<td>3.4 (2.8 to 4.6)</td>
<td>0.065</td>
</tr>
<tr>
<td></td>
<td>DescSlope</td>
<td>10 (8.4 to 11.6)</td>
<td>10.5 (8 to 11.8)</td>
<td>0.774</td>
</tr>
<tr>
<td></td>
<td>NIRS VO2</td>
<td>103 (74 to 156)</td>
<td>108 (73 to 140)</td>
<td>0.442</td>
</tr>
</tbody>
</table>
Conclusion Rapid repetition of a VOT alters VOT-induced NIRS-derived variables in healthy volunteers but not in critically ill patients. If these alterations reflect preconditioning, our results suggest that this phenomenon may be altered in critically ill patients.

P3 Low incidence of delirium in patients followed by physiotherapists in the ICU
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Introduction Delirium is an acute temporary and fluctuating mental-organic syndrome, characterized by a global impairment of cognitive function, reduced level of consciousness, attentional deficits and altered sleep–wake cycle, and changes in arousal (hyperactive, hypoactive, or mixed). The Confusion Assessment Method (CAM and CAM ICU) [1] is a diagnostic assessment instrument for delirium and can lead physiotherapeutic treatment, aiming to optimize patient’s recovery, reinforce the importance of preventive and therapeutic measures, and appraise the multidisciplinary treatment approach in this severe complicating syndrome. Delirium is present in 20 to 40% of ICU patients. So far we have no data regarding the incidence of delirium at the ICU in patients followed by physiotherapists. The objective of this study was to verify the incidence of delirium through the CAM ICU instrument in ICU patients followed by physiotherapists.

Methods Trained and capacitated physiotherapists applied the CAM ICU in patients admitted to Albert Einstein Jewish Hospital ICU, older than 18 years old and with 24 hours physiotherapy assistance per day. The content and level of consciousness investigation was performed daily in all physiotherapy sessions, which allowed characterizing the need to apply the CAM ICU. These data were evaluated twice a week during a 30-day period through medical charts.

Results During the study period, 226 patients were evaluated by physiotherapists, median age 70 years old (range 21 to 92), and 60% were male. The clinical admission diagnoses were: 36% sepsis, 20% cardiac, 19% neurologic, 11% respiratory, 4% orthopedic, 3% liver and gastric patients, and 2% vascular. The mean Simplified Acute Physiology Score (SAPS 3) was 45 points. We found a delirium incidence of 7% (n = 16), of these patients 25% were under mechanical ventilation 16 hours or more. Delirium occurred in 85% of patients, 14% liver patients, 8% respiratory patients and 2% cardiac patients. We found no correlation of delirium with clinical admission diagnosis (P = 0.76; r = –0.054).

Conclusion There was a low incidence of delirium in ICU patients followed by physiotherapists. This may be due to physiotherapy and multi-professional team interventions performed earlier.

Reference

P4 Relationship between clinical and cytokines profile of brain-death donors
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Introduction Brain death induces a massive inflammatory response. The majority of transplants are derived from donors who suffered from brain injury. The possible relation of clinical profile and cytokines in donors has been poorly explored. The objectives of this study were to analyze clinical characteristics of brain-dead donors and the correlation with cytokine profile in the ICU of a unique tertiary-care hospital.

Methods We evaluated 120 consecutive potential brain-dead organ donors (mean age 34.9 years, 74.2% males) between July 2007 and June 2008. Plasma cytokines (TNF, IL-2, IL-4, IL-5, IL-6, IL-8, IL-10, IFNγ) were measured in 40 donors immediately after criteria for brain death (or confirmatory tests) and after obtaining consent from families. Cytokines were assessed by cytometric bead array in the plasma and all laboratory personnel were blinded to clinical information.

Results The main cause of brain death was cerebral trauma (80%) and cerebral vascular accidents. The use of vasoactive agents was 90.6%. The median time of stay in the ICU was 2 days and the mean of organs transplanted was 2.2. Data (mean pg/ml) of cytokines were: IL-2: 3.32; IL-4: 2.63; IL-5: 11.4; IL-10: 59; IFN, 9.72; and TNF, 2.32. In 35% of donors IL-6 was above 5,000 pg/ml and in 15% IL-8 was below the detection limit of analysis. We did not find correlation (nonparametric statistical tests) between cytokines and gender, age, and laboratory tests of our organ donors. Pearson correlation between IL-6 and TNF was 0.001. IL-2 and IL-4, IL-5, IL-10 and IFN γ presented Pearson correlation ≤0.00. See Table 1.

Table 1 (abstract P4). Cytokine levels

<table>
<thead>
<tr>
<th>Cytokine</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>IL-2</td>
<td>0.147</td>
</tr>
<tr>
<td>IL-4</td>
<td>0.044</td>
</tr>
<tr>
<td>IL-5</td>
<td>0.252</td>
</tr>
<tr>
<td>IL-8</td>
<td>0.764</td>
</tr>
<tr>
<td>IL-10 Th2</td>
<td>0.214</td>
</tr>
<tr>
<td>IFNγ</td>
<td>0.001</td>
</tr>
</tbody>
</table>

Conclusion Levels of proinflammatory and anti-inflammatory cytokines were increased in brain-dead donors and were correlated. There was no difference between cytokines and clinical and laboratory profiles.

References

Cardiology

P5 Development of a sleep quality questionnaire to assess sleep in the ICU: a polysomnography study
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Introduction Sleep is an important issue for the maintenance of cardiovascular homeostasis through heart rate and blood pressure modulation. Patients admitted in a coronary care unit (CCU) may exhibit a peculiar sleep pattern that is not fully understood. A feasible and cost-effective tool to analyze sleep in this scenario could bring important information for clinicians. The aim of this study was to evaluate sleep with a questionnaire developed specifically for the CCU and to establish correlations with polysomnography.

Methods Consecutive acute coronary syndrome patients admitted to a CCU between March 2011 and October 2012 were selected. The exclusion criteria were: hemodynamic instability, sedation, receiving vasoactive drugs, or under ventilation support. Patients were submitted to polysomnography in the first 36 hours after admission. A specific 18-questionnaire (Storti questionnaire) divided into diurnal and nocturnal sleep was developed according to experts’ skills. The Pittsburgh and the Storti questionnaires were applied immediately before the CCU discharge. Cronbach’s alpha test was used for internal questionnaire validation. Spearman and Kruskal–Wallis tests were used to analyze the correlation between polysomnography variables and questionnaire.

Results Ninety-nine patients (68% male; mean age 56 ± 10 years) were included. The mean BMI was 27 ± 5 kg/m². Arterial hypertension was observed in 52% of the sample; 17% had diabetes, 39% were smokers. The patients present a total sleep time of 265 ± 81 minutes during the polysomnography, sleep efficiency was 62 ± 18%, REM sleep was
Variables were compared by Pearson's chi-square test. Continuous data were calculated based on the change between the first BNP collected and the last one, indicating for clinical purposes. The BNP variation was calculated as the difference between 2004 and 2012 in a prospective single-center registry. A logistic regression model was used to test the association between BNP increase (categorized by an increase ≥200 pg/ml) and mortality, adjusted for age, gender, troponin levels, Killip classification, left ventricular ejection fraction (LVEF) and ST elevation AMI.

**Conclusion** The Storti questionnaire had a good correlation with sleep efficiency assessed by polysomnography. The majority of patients in CCH had a poor or regular sleep.

**P6** Serial brain natriuretic peptide strongly predicts in-hospital mortality in patients with acute myocardial infarction

AE Pereira Pesar, M Katz, C Pereira, AG Correa, AN Fava, M Franken, ACB Nunes, LM Forlenza, F Tarasoutchi, MR Makdisse

**Introduction** Brain natriuretic peptide (BNP) can be useful in risk stratification of patients with acute myocardial infarction (AMI). However, the value of serial in-hospital BNP assessment to predict mortality in this setting was poorly investigated before. Thus, the aim of this study was to evaluate the usefulness of serial BNP measurements to predict mortality in patients with AMI.

**Methods** Patients with AMI (n = 2,198) were consecutively enrolled between 2004 and 2012 in a prospective single-center registry. A subgroup analysis was performed in the patients submitted to serial BNP measurements as indicated for clinical purposes. The BNP variation was calculated as the difference between the first BNP collected (baseline) and the highest subsequent in-hospital BNP measurement. Baseline characteristics and the BNP variation were compared between patients who survived or died during the in-hospital period. Categorical variables were compared by Pearson's chi-square test. Continuous variables were compared using the Student t test or Mann–Whitney test. The logistic regression model was used to test the association between BNP increase (categorized by an increase ≥200 pg/ml) and mortality, adjusted for age, gender, troponin levels, Killip classification, left ventricular ejection fraction (LVEF) and ST elevation AMI. P <0.05 was considered statistically significant.

**Results** Serial BNP levels were determined in 280 patients (59% men, 78 ± 12 years, 33% ST elevation). The BNP increase >200 pg/ml was detected in 114 (41%) patients. All baseline clinical parameters (gender, age, diabetes, hypertension, Killip classification, LVEF and ST elevation) were similar between patients with or without BNP increase. Mortality was higher in patients with BNP increase (25% vs. 12%; P = 0.006). In the adjusted logistic regression model, only age (OR = 1.04; 95% CI = 1.002 to 1.08; P = 0.04) and BNP increase >200 pg/ml (OR = 3.9; 95% CI = 1.83 to 8.20; P <0.001) were independent predictors of in-hospital mortality.

**Conclusion** This study demonstrated that an in-hospital BNP increase >200 pg/ml is strongly and independently related to in-hospital mortality in patients with AMI. Thus, serial BNP testing may be useful to detect high-risk AMI patients.

**Epidemiology/Quality of Life/Administration**

**P7** Causes of ICU readmission and mortality: analysis of a 6-month period

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**Introduction** Patients readmitted to ICU have a higher mortality and longer ICU and hospital stay. Furthermore, the readmission rate is used as a quality indicator of critical care unit performance, because this index may reflect the adequacy of treatment. The objective was to evaluate the readmission rate of a tertiary public hospital during a 6-month period.

**Methods** We performed a retrospective analysis of all adult patients readmitted to a 20-bed mixed-case ICU between 1 September 2012 and 28 February 2013. The cases (readmission) were collected from clinical electronic information systems.

**Results** During this period 402 patients were admitted to the ICU. The mortality in the ICU was 24.6% and overall hospital mortality rate was 31.6%. The average SAPS 3 during readmission on the ICU was 52 with a predicted mortality of 34%. The readmission rate was 5.2%, ICU mortality was 23.8% and hospital mortality was 28.6%. The most frequent cause of readmission was nosocomial pneumonia (29%), neurologic causes (19%), sepsis (14%), administrative (14%), postoperative support (10%), metabolic disorders (10%) and cardiology events (5%). The patients were most commonly readmitted from the ward (33%), emergency department (14%), step-down unit (14%), operating theater (5%) and others (33%). The most common supportive therapies after readmission were mechanical ventilation in 38.1%, vasopressors in 28.6%, and renal support in 9.5%.

**Conclusion** The most common reason for ICU readmission in our unit was nosocomial pneumonia. The mortality of the readmitted patients was not superior to the predicted mortality for the overall cohort of patients.

**P8** Cost of the quantitative adequacy of nursing staff in the ICU

CP Guimarães1, PC Garcia2, FMT Fugulin1

1 Escola de Enfermagem da Universidade de São Paulo, SP, Brazil; 2 Hospital Universitário da Universidade de São Paulo, SP, Brazil

**Introduction** The high cost to maintain a complex structure such as the ICU has justified its strict control. Nevertheless, budgetary limitation and expenses abatement directly affect the outcome of a fully adequate nursing staff due to its high–percentage representation in human resources. Therefore, the lack of proper personnel interferes on many aspects of daily activities such as organization and safety of patients and staff, affecting also the assistance provided and the institutional goals. For that reason, study on costs and management of nursing personnel is paramount, as it can evidence the effects of an impaired scenario and the relation between cost and efficiency in a healthcare environment. The objectives were: to verify the average staff time required by a patient for a proper assistance or treatment; to calculate the actual average time and cost spent by the crew; and to estimate the average daily time and cost for a proper adult ICU’s activity.

**Methods** The research was based on a quantitative and descriptive data. The study took place in the Hospital Universitário da Universidade de São Paulo’s adult ICU block, from 1 January 2008 to 31 December 2009. Data concerning the average time of assistance given to the patients and requested by them were collected from the ICU’s management instruments. Data concerning personnel fees per hour were based on the ICU nursing staff’s wage bill, provided by the Finance Department.

**Results** The daily average time of required assistance is 16 hours. However, the actual daily average time of provided assistance is 14 hours, which poses a great disparity statistically. In 24 hours the average cost of given assistance per patient was R$715.79. On the other hand, adequate assistance would require R$805.66. The average cost per month to amend the actual scenario would be R$40,490.00, which corresponds to an increase of 17.16% over the existing outline’s budget.

**Conclusion** The literature review and the data suggest that although the adequacy of the nursing staff entails higher costs, it may contribute to improve the quality of care, reducing costs arising from possible negative outcomes in patients.

**References**


Factors associated with prolonged ICU stay: a retrospective analysis
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Introduction
Critically ill patients frequently stay on the ICU for prolonged periods. Prolonged ICU stay (PIS) is associated with increased costs, resource use and family burden. Nevertheless, risk factors at admission associated with prolonged ICU stay are only partially described. The objective was to evaluate factors associated with prolonged ICU stay on a mixed ICU.

Methods
Retrospective analysis of 3,257 patients admitted to a tertiary hospital in São Paulo, Brazil. Twenty-seven relevant variables that were clinically associated with prolonged (>14 days) were included on a univariate analysis. Variables included demographic data, reason for admission, type of admission (clinical, elective surgery, emergency surgery), previous status performance, presence of comorbidities, illness severity (assessed by SAPS III score), laboratorial data and need for organ support device (vasopressors, mechanical ventilation, dialysis) on the day of admission. A multivariate analysis was performed to identify variables independently associated with PIS.

Results
In total, 203 (6.3%) of the 3,257 patients admitted in the analyzed period stayed on the ICU for at least 14 days. Hospital mortality was higher in patients with PIS (49.7% vs. 9.8%; P < 0.01). On multivariate analysis, SAPS III (OR = 1.03, CI = 1.01 to 1.04), reduced status performance (dependency for one or more daily activities – OR = 1.71, CI = 1.18 to 2.46), bedridden status (OR = 1.91, CI = 1.08 to 3.38), emergency surgery (OR = 2.87, CI = 1.27 to 6.51), admission due to intracranial mass effect (OR = 4.46, CI = 1.16 to 17.04), admission from the ward (OR = 3.35, CI = 1.05 to 10.63) and hospital transfer (OR = 5.23, CI = 1.62 to 16.91) were independently associated with PIS. Age was not related to PIS. No comorbidity or organ support device was independently associated with PIS.

Conclusion
In a large database of critically ill patients, global illness severity, baseline status performance and emergency surgery were related to PIS. No comorbidity or need for organ support device was associated with PIS.

Acknowledgement
The authors would like to thank Dr Marcelo Park for helping with statistical analysis.

Feasibility of transitioning from APACHE II to SAPS III as prognostic model in a Brazilian general ICU
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Introduction
Prognostic models reflect the population characteristics of the countries which they originate from. The development of the Acute Physiology and Chronic Health Evaluation II (APACHE II) system was based on a cohort of patients in the United States, and it has been used in many ICUs around the world. Newer score systems were developed as the Simplified Acute Physiology Score III (SAPS III) that was developed and validated in a multicenter and multinational cohort study. Predictive models should be customized to fit in the case-mix where they will be used because the outcomes in the original databases and period from which the models were derived may be different from the databases of ICUs using the models. In the present study, we performed the external validation of two predictive models and directly compared their performance in an independent population of mixed critically ill patients in Brazil. The aim is to assess the feasibility of transitioning from APACHE II to SAPS III.

Methods
Data were retrospectively collected only for APACHE II during August 2011 and December 2011, and only for SAPS III during May 2012 and September 2012. From January 2012 to April 2012, during a period of calibration, the two scores were calculated in all patients admitted to the ICU and were collected for analysis. All ICU admissions were enrolled during the period analyzed. The exclusion criteria were: age <18 years, missing data, and not receiving ICU care. The calibration of the scores was tested using the Hosmer–Lemeshow goodness-of-fit procedure. The discriminatory ability of the models was assessed using receiver operating characteristic (ROC) curves and respective areas under curves (AUC). The standardized mortality ratio (SMR) was calculated using the models by dividing the number of observed deaths by the number of expected deaths. Confidence intervals of the SMR were computed to test the model’s uniformity-of-fit and were calculated using the proposed methods.

Results
A total of 3,333 ICU admissions were enrolled until the end of September 2012. The Hosmer–Lemeshow goodness-of-fit statistics supported model fit of all models for in-ICU mortality with the exception of APACHE II in patients in the calibration database undergoing elective surgery. For in-hospital mortality there is a worse fit of APACHE II in clinical patients during the first period and of SAPS III in patients in the calibration database undergoing elective surgery. The calibration curves for APACHE II and SAPS III shows overestimation of the risk of death in all ranges of predicted mortality. Discrimination, evaluated by the AUC, in general and clinical patients was best for SAPS III for in-ICU and in-hospital mortality. SMRs for the whole population were 0.27 (CI = 0.23 to 0.33) for APACHE II and 0.28 (CI = 0.22 to 0.36) for SAPS III. In the calibration database, the SMRs for APACHE II and SAPS III were 0.33 (CI = 0.22 -to 0.50) and 0.36 (CI = 0.25 to 0.55), respectively. For all models, the SMRs showed some variation across the spectrum of patients. The SMRs ranged from 0.24 to 0.46 for APACHE II, and 0.09 to 0.31 for SAPS III. In the calibration database, the SMRs ranged from 0.13 to 0.38 for APACHE II, and from 0.18 to 0.40 for SAPS III.

Conclusion
The external validation of two widely used prognostic models showed good discrimination and good calibration when applied to the same independent population of Brazilian ICU patients. The transition from APACHE II to SAPS III in this Brazilian ICU was feasible and in some scenarios the SAPS III had even better performance than APACHE II. In conclusion, we showed in a cohort of Brazilian patients from a tertiary hospital that SAPS III is the best prognostic score, with the highest discrimination and calibration power. The transition from an older score (APACHE II) to a newer one (SAPS III) is feasible in this scenario.
This authorizes us to question the routine practice of laboratory tests, resulted in normal values, with the aggravating factor of consecutivity.

**Conclusion**

Results were normal.

95% CI in parentheses. ACEI, angiotensin-converting enzyme inhibitor; ARB, angiotensin receptor blockers.

**Acknowledgements**

This project is funded by Hospital Totalcor and Amil Clinical Research.

P12

**Laboratory routine in the ICU: a practice to be abolished?**

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**Introduction**

The execution of laboratory tests is frequently requested for diagnosis and/or monitoring of critical patients. Their role as an aid, however, is diminished by unreasonable practices – resulting, unfortunately, in iatrogenesis and higher costs. In face of these omenous results, we tracked the laboratory tests performed routinely in an ICU.

**Methods**

We retrospectively analyzed the results of tests performed on a daily and consecutive basis during the months of May and June 2012. We deliberately restricted our analysis to the levels of creatinine, urea, sodium, and potassium, and to the prothrombin time (PT) and activated partial thromboplastin time (aPTT) tests. To infer their propriety, we compared each result with their respective normality references. In addition to comparing the cumulative rates of normal tests, we were also interested in the correlation of the results and the admission APACHE II score and time of hospitalization in the ICU, as well as in the volumes of blood drawn and in the test costs.

**Results**

Forty-eight patients (28 men) were studied, with a mean age of 46.6 ± 18.6 years, average APACHE II score of 16.5 ± 6.9 points, average length of stay in the ICU of 15.2 ± 11.7 days, and 33% fatality rate. A total of 3,622 tests were performed (90.6/deceased patient × 67.1/surviving patient), with a predominance of potassium (13.5%), sodium (13.3%), and creatinine (13.3%) levels, and complete blood count, (13.2%). We observed a linear correlation \( r = 0.81 \); \( P < 0.05 \) between the number of performed tests/patient and the respective length of stay in the ICU; no correlation was observed for the APACHE II score. The volume of blood drawn per patient per hospitalization varied between 10 and 525 ml, being higher for deceased patients (average of 103.5 ± 84.2). Of the total tests, 48.8% of the results were normal (43.9 tests/surviving patient × 44.9 tests/deceased patient), especially for potassium levels (9.8%) and aPTT (9.2%). Also, 31.8% of all results (43.9 tests/surviving patient × 44.9 tests/deceased patient), especially of 103.5 × 84.2). Of the total tests, 48.8% of the results were normal.

**Conclusion**

The collected data reveal that almost one-half of the tests in transusions) and their respective costs (approximately US$54,000 annually).

P13

**Nursing care time and quality indicators for the ICU: correlation analysis**

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**Introduction**

The objective of this exploratory, retrospective, quantitative study was to analyze the time spent by the nursing staff to assist patients in an adult ICU (AICU) of the University Hospital, University of São Paulo (UH-USP) and verify its correlation with quality care indicators.

**Methods**

This research started on 1 January 2008 until 31 December 2009. Data were collected from the administration tools used by the head nursing staff of the unit. Analysis of the average length of time in relation to the average length of time required by patients was performed using the paired \( t \) test. The correlation coefficient was used to verify the correlation of the average length of time care given to patients in the AICU with the quality indicator incidence.

**Results**

The average length of time regarding assistance for patients, in the analyzed period, accounted for 14 hours, of which 31% were performed by nurses and 69% by technicians/nursing assistants. The hospitalized patients required approximately 16 hours of care. The application of statistical tests showed that the differences found between the hours of assistance given by the nursing staff and those required by patients was significant (\( P < 0.001 \)), suggestive of the heavy workload for the nurses in the AICU. The correlational analysis between the length of time of nursing care given by nurses and the quality indicator incidence of accidental extubation evidences Pearson’s correlation coefficient (\( r = -0.454 \)), with \( P = 0.026 \), indicating negative linearity between variables, which allowed us to infer that the incidence of accidental extubation decreases with increasing nursing care time given by nurses.

**Conclusion**

The results revealed that the average hours of nursing care for patients of the AICU were lower than those recommended by the official Brazilian Agencies. The average time of care required by patients hospitalized in this unit was higher than that recommended by ANVISA and lower than that established by COFEN Resolution Nr. 293/04. This indicates that the quantitative assessment of nursing staff in ICUs requires prior knowledge of the users’ healthcare demands and not only the use of parameters indicated by official agencies, since this procedure may cause an overdimensioning or underdimensioning of nursing professionals. The results of this study showed the influence of nursing care time provided by nurses in the outcome of care given to patients assisted in the AICU. The addition of more evidence may help to demonstrate the impact of nursing care time in healthcare results and patient safety.

P14

**SAPS 3 as a predictor admission of surgical patients in the ICU**

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**Introduction**

Owing to lack of intensive care beds, patients undergoing intermediate-risk surgery are usually sent to the ward postoperatively. However, a part from this population evolves with complications requiring intensive care (ICU). The aim of the study was to evaluate the characteristics of surgical patients who were admitted to the ICU lately and to find predictors of the need for intensive care.

**Methods**

A prospective cohort study was performed in a tertiary hospital for 1 year. The study included patients with preoperative indication for ICU, but who at the end of surgery were taken to the
ward postoperatively because of good clinical surgery. We evaluated the need for the ICU in this group, the SAPS 3 score preoperatively, the ASA physical status, demographics, origin and service requestor, need for blood transfusions intraoperatively, surgery time and hospital mortality. Patients undergoing palliative surgery were excluded. Independent predictors of the need for intensive care were assessed using logistic regression, with the sensitivity and specificity studied by ROC curve. See Figure 1.

**Results** We included 100 patients aged 66.4 ± 14.7 years. The SAPS 3 score average was 38.5 ± 8.6, 71% had ASA 2, women constituted 66% of casuistic. Most surgery was elective, the most frequent gynecologic (30%) and orthopedic (28%) and neuraxial regional anesthesia (49%). Of all patients, 27% required ICU admission, average on the sixth day after surgery and 3.0% died. The SAPS 3 score average was higher (45.4 ± 7.8 vs. 35.9 ± 7.4, P < 0.001) and ASA 3 was more prevalent (40.7% vs. 8.2%, P = 0.001) in patients who required intensive care postoperatively. Furthermore, these patients had longer duration of surgery (4.2 ± 1.9 vs. 2.7 ± 1.5 hours, P < 0.001), higher prevalence of gastrointestinal surgery (14.8% vs. 5.5%, P = 0.03) and greater need for intraoperative transfusion (18.5% vs. 5.5%, P = 0.04). In these patients admitted to the ICU mortality was 11.1% versus 0.0%, P = 0.004. In multivariate analysis, we found the value of SAPS 3 as an independent factor in determining whether the patient would need the ICU; OR = 1.25; 95% CI = 1.1 to 1.4; P = 0.001, and even the time of surgery, OR = 3.33; 95% CI = 1.7 to 6.3; P = 0.002. The ROC curve was 0.87; 95% CI = 0.78 to 0.93 for the SAPS 3 discriminating need for intensive care, rather than ASA, ROC 0.64; 95% CI = 0.54 to 0.74.

**Conclusion** The identification of high-risk surgical patients is a difficult task, but essential for their proper treatment, surgery time together with the SAPS 3 seem to be useful tools in this differentiation and may help to better characterize this population.

---

**Table 1 (abstract P15). SAPS 3 based on categories of severity**

<table>
<thead>
<tr>
<th>Category</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
<th>5</th>
<th>6</th>
<th>7</th>
<th>8</th>
<th>9</th>
</tr>
</thead>
<tbody>
<tr>
<td>SAPS 3 points</td>
<td>0 to 24</td>
<td>25 to 34</td>
<td>35 to 44</td>
<td>45 to 54</td>
<td>55 to 64</td>
<td>65 to 74</td>
<td>75 to 84</td>
<td>85 to 94</td>
<td>&gt;95</td>
</tr>
</tbody>
</table>

---

**Table 2 (abstract P15). Data with SAPS 3, SRU and SMR**

<table>
<thead>
<tr>
<th>Category</th>
<th>SAPS 3, mean (SD)</th>
<th>Patients</th>
<th>Survivors</th>
<th>Σ days LOS (all patients)</th>
<th>SRU</th>
<th>SRU standardized</th>
<th>SMR</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>19.75 ± 4.08</td>
<td>112</td>
<td>111</td>
<td>161</td>
<td>1.45</td>
<td>0.03</td>
<td>1.1</td>
</tr>
<tr>
<td>2</td>
<td>30.36 ± 2.92</td>
<td>335</td>
<td>335</td>
<td>673</td>
<td>2.01</td>
<td>0.13</td>
<td>0</td>
</tr>
<tr>
<td>3</td>
<td>39.31 ± 2.77</td>
<td>350</td>
<td>346</td>
<td>823</td>
<td>2.37</td>
<td>0.16</td>
<td>0.13</td>
</tr>
<tr>
<td>4</td>
<td>49.18 ± 2.86</td>
<td>304</td>
<td>295</td>
<td>1,087</td>
<td>3.68</td>
<td>0.22</td>
<td>0.13</td>
</tr>
<tr>
<td>5</td>
<td>59.06 ± 2.79</td>
<td>202</td>
<td>174</td>
<td>1,155</td>
<td>6.64</td>
<td>0.23</td>
<td>0.3</td>
</tr>
<tr>
<td>6</td>
<td>68.77 ± 2.79</td>
<td>86</td>
<td>73</td>
<td>795</td>
<td>10.89</td>
<td>0.16</td>
<td>0.22</td>
</tr>
<tr>
<td>7</td>
<td>78.5 ± 2.45</td>
<td>34</td>
<td>18</td>
<td>251</td>
<td>13.94</td>
<td>0.05</td>
<td>0.56</td>
</tr>
<tr>
<td>8</td>
<td>88.54 ± 2.99</td>
<td>13</td>
<td>7</td>
<td>86</td>
<td>12.28</td>
<td>0.02</td>
<td>0.5</td>
</tr>
<tr>
<td>9</td>
<td>97.8 ± 4.21</td>
<td>5</td>
<td>2</td>
<td>16</td>
<td>8</td>
<td>0.003</td>
<td>0.62</td>
</tr>
<tr>
<td>Total</td>
<td>45.14 ± 15.9</td>
<td>1,441</td>
<td>1,361</td>
<td>5,047</td>
<td>3.72</td>
<td>0.13</td>
<td>0.24</td>
</tr>
</tbody>
</table>
References

P16
Use of drugs in the ICU and its iatrogenic potential
E de Castro Vieira, NM Vidal, GMA Fidelis, EC Santos, AA Peixoto Jr, FA Meneses
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Introduction Developed and administered for the purpose of benefit, pharmaceutical agents can cause harmful effects to the patient. We analyzed their use in a population of patients admitted to the ICU, trying to detect their potential exposure to drug–drug interactions.

Methods Prospective study of patients hospitalized for more than 48 hours, during the period from May to July 2012, with tracking of drug interactions according to Micromedex (Version 2.0).

Results We analyzed 50 patients with a mean age 45.4 ± 18.7 years, the majority were female (54%) and from the emergency unit (74%). The mean APACHE II score was 17.6 ± 7.3 points, the mean SOFA score (day 1) 7.3 ± 4.2 points, and the median length of stay 21 (IQR: 12.5 to 31.5) days. One hundred and three drugs were prescribed, predominantly antimicrobials (100% of patients) and analgesics (98% of patients). The average/day/patient of prescription drugs was 10 ± 2.6, and the average/day/patient drug interactions 2.7 ± 2.8. On exposure to drug interactions important and moderate risk occurred, respectively, in 78% and 86%, identifying the association contraindicated in 34% of patients of a positive correlation between length of stay in the ICU and exposure to important risk interactions (r = 0.53, P = 0.00006) and moderate risk interactions (r = 0.34, P = 0.013). Patients exposed to important risk interactions had greater severity at ICU admission for the APACHE II (18.5 ± 7.1 vs. 14.3 ± 6.9 points, P = 0.045) and SOFA (day 1) (7.8 ± 4 vs. 4.2 ± 3.9 points, P = 0.017).

Conclusion The high numbers of drug interactions with important risk incidents, especially in sick population, alert us to the necessity of knowledge by the intensivist for the use of drugs, due to the iatrogenic potential exacerbating the severity already underway.

References

P17
Analysis of the efficacy of an experimental expert system of medical prescription in reducing medical errors and excessive physician workload: a cross-sectional study
HH Shieh, ER Barreira, EJ Troster, SC Brasica, AC Ventura, PF Góes, I de COF Fernandes, DC de Souza, JC Fernandes, F Pereira das Chagas, R de Jesus, LO Zagne, FR Caino, AE Gilio, VHK Koch, S Fukugava, ER Barreira, EJ Troster
University Hospital of Universidade de São Paulo, Vila Iara, São Paulo, SP, Brazil

Introduction Deaths attributable to preventable medical errors (PME) in hospitals exceed those caused by well-known life-threatening conditions, such as motor vehicle accidents, breast cancer, and AIDS [1]. The Institute of Medicine estimates that as many as 98,000 deaths are caused by PME every year [1]. The risks derived from PME are even more severe when they affect critically ill patients, or include medications that must be adjusted for the patient’s body weight. Fatigue and work overload can represent a threat to the patients’ safety in pediatric ICUs [2]. Expert Systems (ES) [3], a branch of artificial intelligence, can be used to solve the problems related to medical prescription errors (MPE). Studies analyzing the role of ES in MPE are still lacking. The objective of this study was to compare the accuracy of an experimental ES with the written medical prescription.

Methods After signing an informed consent, pediatricians working in a university hospital were asked to write a medical prescription containing 10 different medications (maintenance fluids, adenosine, epinephrine, atropine, phenytoine, vancomycin, ceftazidine, amphotericin B, dobutamine, and fentanyl) for a hypothetical patient. The written medical prescription was compared with the ES prescription made by the same physicians, after a 2-minute training period. Statistical analysis was done using the χ², Fisher’s exact test, paired t test or Wilcoxon test (paired samples), whenever applicable. A significance level of 0.05 was used for all analyses.

Results Thirteen pediatric residents and seven attending physicians participated on the study; the mean time since medical graduation was 10.1 ± 9 years. Fifty-seven prescribing errors were detected on medical prescription (nine unreadable items, 23 omissions, 14 dilution errors and five velocity of infusion errors) in comparison with one error of duplication of medication in the ES prescription (P <0.001).

Conclusion The medical prescription of critically ill pediatric patients deserves special attention. The use of an experimental ES required a short training period and resulted in a significant decrease in prescribing errors and physicians’ workload. Nevertheless, this computerized approach is not error free, and double-checking must be performed by the prescriber physician.

References
1. To Err is Human: Building a Safer Health System. Institute of Medicine, 1999.

P18
Update of the pediatric hypertension graphic adjusted for gender and height percentiles: systolic blood pressure for boys, 1 to 17 years old
HH Shieh, AE Gilio, VHK Koch, DC Raulik, C Vranjac, S Fukugava, ER Barreira, EJ Troster
University Hospital of Universidade de São Paulo, Vila Iara, São Paulo, SP, Brazil

Introduction Hypertension is the most important preventable risk factor for premature death worldwide. It increases the risk of ischemic heart disease, strokes, peripheral vascular disease, and other cardiovascular diseases, including heart failure, aortic aneurysms, diffuse atherosclerosis, and pulmonary embolism. In childhood, hypertension can be determined according to a table adjusted for height, age and gender [1]. A graphic representation of pediatric hypertension was published in 1987 [2], and no graphic updates have been published since then. The objective of this study was to update the graphic representation of pediatric hypertension.

Methods We used a computerized calculation method to develop high-resolution graphics containing curves with 5,841 points each, to depict the main percentiles associated with high blood pressure for boys from 1 to 17 years old in the 50th percentile of height. Each point represents the calculation of the polynomial equation that includes the statistical processing of the last Report on Blood Pressure [1]. We also analyzed the effect of height on blood pressure in the 5th to 95th percentile range. Statistical functions generated by computerized program were used.

Results Six monotonic curves of systolic BP for boys representing the 50th, 75th, 90th, 95th, 99th, and 99th +5 percentiles were built (Figure 1). In relation to the table published by the NIH, we confirm use of approximation of the values in the published table by truncation. Considering a tolerance of 1 mmHg, the monotonic curve of adjustment of the values in the published table by truncation.
the 36.5th to 64.5th percentile of height, but needs maximal correction for the 95th percentile of height (+3.8 mmHg correction).

Conclusion
The adjustment of systolic BP for height is of little significance, and the updated graphic can be used to diagnose high systolic blood pressure for boys. Clinical studies are necessary to determine the systolic BP percentile that better represents clinically significant hypertension.

References

P19
Update of the pediatric hypertension graphic adjusted for gender and height percentiles: systolic blood pressure for girls, 1 to 17 years old
HH Shieh, AE Gilio, VHK Koch, DC Raulik, C Vranjac, S Fukugava, ER Barreira, EJ Troster
University Hospital of Universidade de São Paulo, Vila Iara, São Paulo, SP, Brazil

Introduction
Hypertension is the most important preventable risk factor for premature death worldwide. It increases the risk of ischemic heart disease, strokes, peripheral vascular disease, and other cardiovascular diseases, including heart failure, aortic aneurysms, diffuse atherosclerosis, and pulmonary embolism. In childhood, hypertension can be determined according to a table adjusted for height, age and gender [1]. A graphic representation of pediatric hypertension was published in 1987 [2], and no graphic updates have been published since then. The objective of this study was to update the graphic representation of pediatric hypertension.

Methods
We used a computerized calculation method to develop high-resolution graphics containing curves with 5,841 points each, to depict the main percentiles associated with high blood pressure for girls from 1 to 17 years old in the 50th percentile of height. Each point represents the calculation of the polynomial equation that includes the statistical processing of the last Report on Blood Pressure [1]. We also analyzed the effect of height on blood pressure in the 5th to 95th percentile range. Statistical functions generated by computerized program were used.

Results
Six monotonic curves of systolic BP for girls representing the 50th, 75th, 90th, 95th, 99th, and 99th +5 percentiles were built (Figure 1). In relation to the table published by the NIH, we confirm use of approximation of the values (rounding) in the published table by truncation. Considering a tolerance of 1 mmHg, the monotonic curve of adjustment for height of the SBP for girls does not need any correction in the 31.5th to 68.5th percentile of height, but needs maximal correction for the 5th percentile for height (+4.7 mmHg correction).

Conclusion
The correction of systolic BP for height is of little significance, and this updated graphic can be used to diagnose high systolic blood pressure for girls. Clinical studies are necessary to determine the systolic BP percentile that better represents clinically significant hypertension.

References

P20
Update of the pediatric hypotension graphic adjusted for gender and height percentiles: diastolic blood pressure for boys, 1 to 17 years old
HH Shieh, ER Barreira, A Bousso, AC Ventura, EJ Troster
University Hospital of Universidade de São Paulo, Vila Iara, São Paulo, SP, Brazil

Introduction
According to the National Heart, Lung, and Blood Institute of the National Institute of Health, hypotension refers to an abnormally low blood pressure (BP). In childhood, hypotension can be determined according to two different definitions: BP below the 5th percentile or below two standard deviations (SDs) of the mean for age and gender [1]. A graphic representation of pediatric hypotension was published in 1977 [2], and no updates have been published since then.
The objective of this study was to update the graphic representation of pediatric hypotension.

Methods We used a computerized calculation method to develop high-resolution graphics containing 5,841 points each, to depict the main percentiles associated with low BP for boys from 1 to 17 years old in the 50th percentile of height. Each point represents the calculation of the polynomial equation that includes the statistical processing of the last Report on Blood Pressure [3]. We also analyzed the effect of the adjustment for height of the DBP for boys does not need any correction in the in the 25.5th to 76th percentile of height.

Results Five monotonic curves of diastolic BP for boys representing the 50th, 25th, 10th, 5th, and 2.275th (−2SD) percentiles were built (Figure 1). Considering a tolerance of 1 mmHg, the monotonic curve of adjustment for height of the diastolic BP for boys does not need any correction in the 36.5th to 64.5th percentile of height. The correction of diastolic BP for height is of little significance, and this updated graphic can be used to diagnose low diastolic BP for boys. Clinical studies are necessary to determine the SBP percentile that better represents clinically significant hypotension.

Conclusion The adjustment of systolic BP for height is of little significance, and the updated graphic can be used to diagnose low systolic BP for boys. Clinical studies are necessary to determine the SBP percentile that better represents clinically significant hypotension.

References

Figure 1 (abstract P21). Update of chart for systolic blood pressure (SBP) based on the last Report on Blood Pressure in 2004 [3], for boys 1 to 17 years old (50th percentile of height). Considering a tolerance of 1 mmHg, the curve of adjustment for height of the SBP for boys does not need any correction in the in the 36.5th to 64.5th percentile of height.
systolic BP percentile that better represents clinically significant hypotension.

References

P22
Update of the pediatric hypotension graphic adjusted for gender and height percentiles: systolic blood pressure for girls, 1 to 17 years old
HH Shieh, ER Barreira, A Bousso, AC Ventura, EJ Troster
University Hospital of Universidade de São Paulo, Vila Iara, São Paulo, SP, Brazil

Introduction According to the National Heart, Lung, and Blood Institute of the National Institute of Health, hypotension refers to an abnormally low blood pressure (BP). In childhood, hypotension can be determined according to two different definitions: BP below the 5th percentile or below two standard deviations (SDs) of the mean for age and gender [1]. A graphic representation of pediatric hypotension was published in 1977 [2], and no updates have been published since then. The objective of this study was to update the graphic representation of pediatric hypotension.

Methods We used a computerized calculation method to develop high-resolution graphics containing curves with 5,841 points each, to depict the main percentiles associated with low BP for girls from 1 to 17 years old in the 50th percentile of height. Each point represents the calculation of the polynomial equation that includes the statistical processing of the last Report on Blood Pressure in 2004 [3]. We also analyzed the effect of height on BP from the 5th to 95th percentile. Statistical functions generated by computerized program were used.

Results Five monotonic curves of systolic BP for girls representing the 50th, 25th, 10th, 5th, and 2.275th (~2SD) percentiles were built (Figure 1). In relation to the table published by Haque and Zaritsky [3], the Bland–Altman analysis of the published female systolic BP compared with actual update shows a mean bias of ~0.61, with limits of agreement from ~2.12 to 0.9 mmHg, which confirms the use of approximation of the values in the published table [3]. Considering a tolerance of 1 mmHg, the monotonic curve of adjustment for height of the SBP for girls does not need any correction in the 31.5th to 68.5th percentile of height, but needs maximal correction for the 5th percentile for height (~4.7 mmHg correction).

Conclusion The correction of systolic BP for height is of little significance, and this updated graphic can be used to diagnose low systolic BP for girls. Clinical studies are necessary to determine the systolic BP percentile that better represents clinically significant hypotension.

References

P23
Update of the pediatric hypotension graphic adjusted for gender and height percentiles: diastolic blood pressure for girls, 1 to 17 years old
HH Shieh, ER Barreira, A Bousso, AC Ventura, EJ Troster
University Hospital of Universidade de São Paulo, Vila Iara, São Paulo, SP, Brazil

Introduction According to the National Heart, Lung, and Blood Institute of the National Institute of Health, hypotension refers to an abnormally low blood pressure (BP). In childhood, hypotension can be determined according to two different definitions: BP below the 5th percentile or below two standard deviations (SDs) of the mean for age and gender [1]. A graphic representation of pediatric hypotension was published in 1977 [2], and no updates have been published since then. The objective of this study was to update the graphic representation of pediatric hypotension.

Methods We used a computerized calculation method to develop high-resolution graphics containing curves with 5,841 points each, to depict the main percentiles associated with low BP for girls from 1 to 17 years old in the 50th percentile of height. Each point represents the calculation of the polynomial equation that includes the statistical processing of the last Report on Blood Pressure in 2004 [3]. We also analyzed the effect of height on BP from the 5th to 95th percentile. Statistical functions generated by computerized program were used.

Results Five monotonic curves of diastolic BP for girls representing the 50th, 25th, 10th, 5th, and 2.275th (~2SD) percentiles were built (Figure 1). Considering a tolerance of 1 mmHg, the monotonic curve of adjustment for height of the DBP for girls does not need any correction in the 16th to 78.5th percentile of height, but needs maximal correction for the 95th percentile of height (~2.08 mmHg correction).

Conclusion The correction of female diastolic BP for height is of minimal significance, and this updated graphic can be used to diagnose low diastolic BP for girls. Clinical studies are necessary to determine the diastolic BP percentile that better represents clinically significant hypotension.
3. Analysis using a fixed-effect model showed a reduction in norepinephrine requirement among patients receiving terlipressin or vasopressin infusion compared with control (standardized mean difference, –1.58 (95% CI, –1.73 to –1.44); P < 0.0001). Overall, vasopressin and terlipressin, as compared with norepinephrine, reduced mortality (relative risk (RR): 0.87 (0.77 to 0.99); P = 0.04). Vasopressin compared with norepinephrine decreased mortality in adult patients (RR: 0.87 (0.76 to 1.00); P = 0.05) and in patients with septic shock (42.5% vs. 49.2%, respectively; RR, 0.87 (95% CI, 0.75 to 1.00); P = 0.05; number needed to treat, 1 to 15). There was no difference in adverse events between the vasopressin and control groups (RR: 0.98 (0.65 to 1.47); P = 0.92).

Conclusion Vasopressin use in vasodilatory shock is safe, associated with reduced mortality, and facilitates weaning of catecholamines. In patients with septic shock, vasopressin compared with norepinephrine may also decrease mortality.

References

P24 Vasopressin and terlipressin in adult vasodilatory shock: a systematic review and meta-analysis of nine randomized controlled trials

VGM Pereira, A Serpa Neto, SO Cardoso, JA Manetta, DC Espósito, M de Oliveira Prado Pasqualucci

ABC Medical School (FMABC), Príncipe De Gales, Santo André, SP, Brazil


Introduction Catecholamines are the most used vasopressors in vasodilatory shock. However, the development of adrenergic hypotension and the subsequent loss of catecholamine pressor activity necessitate the search for other options. Our aim was to evaluate the effects of vasopressin and its analogue terlipressin compared with catecholamine infusion alone in vasodilatory shock.

Methods Systematic review and meta-analysis of publications between 1966 and 2011. The Medline and CENTRAL databases were searched for studies on vasopressin and terlipressin in critically ill patients. The meta-analysis was limited randomized controlled trials evaluating the use of vasopressin and/or terlipressin compared with catecholamine in adult patients with vasodilatory shock. The assessed outcomes were: overall survival, changes in the hemodynamic and biochemical variables, a decrease of catecholamine requirements, and adverse events.

Results Nine trials covering 998 participants were included. A meta-analysis using a fixed-effect model showed a reduction in norepinephrine...
Parenteral colistin for the treatment of severe infections by multidrug-resistant Gram-negative bacteria

C Gram, MT Tantia, CM Dantas de Maio Carrilho, JP Garcia, J Festi, LTO Cardoso, F Chiquetti, MM Kanheissa, CC Branco Lopes, DB Blum, V Anami, AR Ruiz, PA Rossatto

Hospital Universitário – Universidade Estadual de Londrina, Villa Operária, Londrina, PR, Brazil


Abstract withdrawn

Vancomycin dose adjustment in severe burn patients based on trough level for drug effectiveness against pathogens at 1 mg/l minimum inhibitory concentration

JM Silva Jr1, AM Oliveira1, EV Campos1, DS Gomez2, MC Ferreira2, CS Giraud1, CV Silva Jr1, SRL Santos1

Burn Unit/Plastic Surgery Division of Clinics Hospital, Medical School, University of Sao Paulo, Butantán, São Paulo, SP, Brazil; 1School of Pharmaceutical Sciences, University of Sao Paulo, Butantán, São Paulo, SP, Brazil


Introduction

Vancomycin is usually prescribed to severe burn patients with sepsis from the intensive care burn unit (ICBU) for control of hospital infection. The objective of this study was to evaluate the contribution of dose regimen adjusted in burn patients with renal function preserved based on drug plasma concentration at the trough and pharmacokinetic-pharmacodynamic (PK/PD) correlation.

Methods

Sixty severe burn patients with documented Gram-positive nosocomial infection from the ICBU were enrolled in a prospective cohort study, and the period of inclusion was 2 years; the protocol was approved by the hospital’s ethical committee. Patients of both genders (43 male/17 female) with preserved renal function were investigated (176 sets). The vancomycin dose regimen was initially 2 g daily for the control of infection in burn patients with sepsis. Pharmacotherapeutic follow-up was performed by a serial blood sample collection (2 ml each) for drug plasma measurements [1]. Drug effectiveness was based on the parameter AUC0–24/MIC >400 [2]; AUC0–24 was the area under the curve (plasma concentration vs. time) integrated up to 24 hours, and the minimum inhibitory concentration (MIC) from in vitro antimicrobial susceptibility testing performed in the hospital [2]. Dose adjustment was required and the daily dose was increased to reach trough levels >10 μg/ml and AUC/MIC >400.

Results

Characteristics of patients investigated were (mean ± SD): 38.9 ± 14.1 years; 70.0 ± 10.6 kg; 28.0 ± 19.0% total burn surface area. Thermal injury occurred in 51/60 patients, and inhalation injury occurred in 54.9% of them, versus electrical injury reported in 9/60 patients; renal function was monitored by serum creatinine (0.72 ± 0.29 mg/dl) and creatinine clearance (153.7 ± 70.4 ml/minute). Significant increase was chosen by comparison of the initial dose against adjusted dose according to trough levels >10 g/ml and also AUC/MIC >400 (Table 1).

Table 1 (abstract P29). Daily dose medians, trough and PK/PD data

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Initial dose</th>
<th>Adjusted dose</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Daily dose (mg/kg/day)</td>
<td>286</td>
<td>42.9</td>
<td>0.01</td>
</tr>
<tr>
<td>Trough levels (μg/ml)</td>
<td>7.1 (3.6 to 13.3)</td>
<td>16.0 (12.0 to 23.3)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>AUC/MIC (MIC: 1 mg/l)</td>
<td>436 (248 to 659)</td>
<td>648 (467 to 942)</td>
<td>0.02</td>
</tr>
</tbody>
</table>

Data presented as median (25 to 75% percentile).

Conclusion

The initial dose recommended for vancomycin must be increased according to trough levels and also AUC/MIC to achieve the PK/PD target in burn patients with preserved renal function.

Acknowledgement

Foundation for Research State of Sao Paulo/SP, Brazil (FAESP).

References

1. López KJV, Bentolucci DF, Vicente KM, Dell’Agua A, Santos SRCJ.

Acute kidney injury according to RIFLE criteria in an ICU: incidence and mortality impact


Unidade de Terapia Intensiva Adulto do Hospital Santa Luzia, Acaú Sul, Brasília, DF, Brazil


Introduction

Acute kidney injury (AKI) is a very common condition in hospitalized patients, especially in ICUs. It is also closely related to adverse patient outcomes, mortality rates as high as three-quarters and as many as 13% need of renal support after hospital discharge. A systematic review demonstrated a close correlation between AKI according to the RIFLE criteria and mortality. The objective of this study was to evaluate the incidence of AKI according to the RIFLE criteria and the impact of each category on mortality in an ICU.

Methods

A retrospective cohort study was conducted with patients admitted to the adult ICU of Hospital Santa Luzia, Brasilia, DF, Brazil, in the period of 6 months. Patients were categorized as Risk (R), Injury (I), Failure (F), or without AKI according to RIFLE criteria. Patients with a previous diagnosis of chronic kidney disease were excluded.

Results

A total of 626 patients were included. Average age was 60 ± 20 years, 326 were male (50.8%), APACHE II was 9 ± 6, 67.1% (n = 326) were nonsurgical, and the mortality rate was 12.3% (n = 77). According to RIFLE criteria, 148 had AKI. Eighty-three patients were classified as R (13.3%, mortality rate of 21.7%), 43 as I (6.9%, mortality rate of 33.5%), and 22 patients as F (3.5%, mortality rate of 54.5%). The relative risk (RR) of death in patients classified as R was 2.72 (95% CI: 1.26 to 4.09), I was 11.27 (95% CI: 5.81 to 21.83), and F was 9.91 (95%...
2. Uchino et al showed a causal role in neuronal death and ICH prognosis. The objective of response associated with hemoglobin byproducts seems to have nonsurvivors than in survivors (496.04×58.5 mg/dl, was 8.86 (IQR: 0 to 27.08). Plasmatic iron concentration was higher in patients) and in 28 days was 66.6% (10 patients). Median hematoma subarachnoid hemorrhage. Overall mortality in 7 days was 40% (six patients) and in 10 patients (IQR: 55 to 65), six patients (40%) were male. Median Glasgow Coma Score was 11.22 (95% CI: 6.57 to 19.17). Eight (1.3%) patients underwent renal replacement therapy during. ICU hospitalization, and mortality. In these patients was 75% (RR: 23.11, 95% CI 4.58 to 116.71). Significant difference was observed in the Kaplan–Meier survival curves of the patients with or without AKI at 28 days (P = 0.00). See Figure 1. Conclusion AKI according to RIFLE criteria was associated with an increased mortality for all categories, mainly in patients with criteria to injury and acute kidney failure, and notably those who needed renal replacement therapy.

References

P31 Analysis of cytokine profile and heme metabolism byproducts after hemorrhagic stroke
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Introduction Intracerebral hemorrhage is a fatal disease, accounting for about 15% of deaths from stroke. Local and systemic inflammatory response associated with hemoglobin byproducts seems to have a causal role in neuronal death and ICH prognosis. The objective of this study was to investigate the relationship of cytokine profile and hemoglobin degradation products with brain injury severity and prognosis.

Methods We developed a prospective cohort study conducted in three tertiary hospitals. All ICH patients with hemoventricle and external ventricular device (EVD) inserted who were admitted to the neurocritical care unit between 2008 and 2011 were included. We collected blood and cerebrospinal fluid (CSF) from the EVD on days 1, 2, 3, 5, and 7 after ICH for measurement of C-reactive protein, cytokines, heme, hemoglobin, cytometry, hemopexine, haptoglobin, enolase, and s-100B concentration. A multiplex analysis was performed to evaluate levels of 17 cytokines. CT scans were evaluated for hematoma and hemoventricle volume. Primary outcome was death in 7 days. This study was approved by the ethics committee of all participating institutions.

Results Fifteen patients were included. Median age was 59 years (IQR: 55 to 65), six patients (40%) were male. Median Glasgow Coma Score was 7 (6 to 9), APACHE II score was 22 (IQR: 15 to 25) and SAPS III was 43 (IQR: 32 to 53). Five patients had hemorrhagic stroke and 10 subarachnoid hemorrhage. Overall mortality in 7 days was 40% (six patients) and in 28 days was 66.6% (10 patients). Median hematoma volume was 10.53 ml (IQR: 5.42 to 31.75) and hemoventricle volume was 8.86 (IQR: 0 to 27.08). Plasmatic iron concentration was higher in nonsurvivors than in survivors (496.04×58.5 mg/dl, P = 0.05) 24 hours after the event. CSF cytometry and lymphocyte count was more increased in nonsurvivors than in survivors (WBC count: 247.5 × 3 cells/ml, P = 0.01 and lymphocyte count: 179×5 cells/ml, P = 0.01) 24 hours after the event. Plasmatic IL-2, IL-6, IL-8, and IL-12 levels are higher in nonsurvivors than in survivors. CSF IL-4 levels were higher in survivors and CSF IL-17 levels were higher in nonsurvivors than in survivors. There was no correlation between plasmatic or CSF levels of enolase and S100-B with mortality. Plasmatic hemopexine and haptoglobin levels also do not seem to be associated with survival.

Conclusion A systemic and local proinflammatory response is associated with higher mortality in patients with hemorrhagic stroke. CSF IL-4 higher concentrations are associated with better prognosis.

P32 Characterization of potential donors of multiple organs and tissues reported in Maranhão from 2009 to 2012
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Introduction Thousands of lives worldwide are improved annually with transplant and donation of organs and tissues. Human organ transplant was one of the greatest medical advances in the 20th century, with a success rate between 80 and 90%. The Brazilian National Transplantation System coordinates and regulates perhaps the largest public transplantation program worldwide. However, family refusal represents the biggest obstacle to the donation process. It is one of the main factors responsible for the shortage of organs and tissues for transplantation. The objectives of this study were to characterize potential donors reported to the Central of Notification, Caption and Distribution of Organs and Tissues in Maranhão (CNCDO-MA) between January 2009 and September 2012, to evaluate the frequency of potential donors converted into effective donors, and to identify the captured organs in effective donors.

Methods A descriptive, retrospective study and quantitative analysis from CNCDO-MA reports, using a standardized data collection form. The inclusion criteria was suspected brain death (BD), with at least the first clinical examination for BD protocol performed – potential donor, notified to CNCDO-MA between January 2009 and September 2012. The exclusion criteria were the notifications in which the cause of death was unknown or supplementary test results were inconsistent with BD. Results A total of 345 notifications were analyzed. The majority were male (63.2%), with an average age of 37.85 (± 18.47) years. The most incident causes of BD were traumatic brain injury (39.7%) and hemorrhagic stroke (36.2%). Arteriography was the predominant supplementary examination (45.5%) to complete the BD protocol. The potential donors converted into effective donors were 14.8%. The amount of collected organs in effective donors was 50 corneas, 29 kidneys, six hearts and one liver.

Conclusion The notification and donation of organs and tissues rates are lower in Maranhão than in Brazil. The integration of multidisciplinary teams is necessary for the responsibility of maintaining potential donors and mandatory BD notification, regardless of the chance of organ donations, which can improve the rates in our state.

P33 Comparison between APACHE II and POSSUM 2 scores in neurosurgical patients admitted to an ICU in Fortaleza, Brazil
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Introduction The need to stratify surgical risk is essential to assess the quality of care of patients undergoing intervention. The objective of this study was to compare APACHE II and POSSUM 2 severity scores in patients undergoing neurosurgery, in the immediate postoperative period.
Methods An observational prospective study with 155 patients admitted consecutively to the ICU of a tertiary hospital in Fortaleza, Brazil, from December 2011 to June 2012, after neurological intervention.

Results The population analyzed showed an average age of 48.0 ± 15.8 years, predominantly female (55.3%). Elective surgeries were more prevalent (92.1%), especially aneurysm clipping (14.6%) and resection of neoplasm (64.2%). At admission, 45 patients (27.6%) had at least one organ dysfunction. The APACHE II score mean was 9.7 ± 5.1, corresponding to the mean of predicted mortality of 14.0 ± 10.7%. The POSSUM 2 score showed a trend to be higher in patients that died, corresponding to the mean of predicted mortality of 14.0 ± 10.7%.

POSSUM 2 score showed a trend to be higher in patients that died, corresponding to the mean of predicted mortality of 14.0 ± 10.7%.

Figure 1: POSSUM 2 score and APACHE II score in patients that died.

Contraction/Relaxation

Introduction Delirium assessment is already a well-established practice in the ICU. Usually, these evaluations are represented as delirium incidence or delirium-free days without coma. We propose four derived variables to predict the outcome in delirium patients in order to identify the most accurate data.

Methods A prospective study took place in the ICU of the University Hospital Professor Edgard Santos, Salvador, Brazil during the period of January to March 2013. Adult patients were assessed twice daily for detection of delirium with the Confusion Assessment Method for the ICU. Those patients with less than 48 hours in the ICU were excluded. The derived variables were classified into four groups: Group 1 – days of delirium, Group 2 – delirium episodes; Group 3 – maximum time of consecutive positive delirium; Group 4 – delirium density (days of consecutive positive delirium/days of delirium). These variables were compared with the outcome – mortality – of patients with positive delirium during the ICU stay. SPSS 21 for Windows was used for statistical analyses.

Results Forty-five patients were analyzed, 16 of whom presented delirium. The mortality of delirium patients was 60%. Group 1: 1 day with delirium was associated with mortality of 60% compared with 50% with 2 days or more. Group 2: patients who had just one episode had 67% mortality compared with 50% mortality if they have two or more episodes. Group 3: mortality with a density >0.5 was 57% versus 50% mortality in the subgroup <0.5. Finally, Group 4: comparing the patients who died with the survival patients, we found 1.75 days of consecutive positive evaluation in the patients who died and 4 days in those who survived. None of the differences between these results for any of the four groups were statistically significant.

Conclusion Owing to the small population analyzed we could not conclude which was the best variable to predict delirium outcomes. None of the variables analyzed affected outcome when compared with just one positive evaluation during the ICU stay. New studies with a larger population are needed to identify the best variable.

P35 Evaluation of new variables to predict delirium outcome

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Introduction Delirium is generally managed by treating its underlying causes. However, symptomatic treatment may also be indicated. Although hypoactive delirium in critically ill patients is the most prevalent subtype of delirium, the effects of treatment with drugs specifically for this group are not well defined. The aim of this systematic review is to evaluate the role of pharmacological treatment in critically ill patients with hypoactive delirium.

Methods A systematic review was conducted, based on the PRISMA criteria, to identify articles on the pharmacological approach to hypoactive delirium in critically ill patients. First, a MEDLINE and Scielo database search was performed for articles published in the English language, involving patients in ICUs in which pharmacological therapy was used to treat delirium. Second, these studies were reevaluated to identify subtypes of delirium and the impact of the treatment.

Results The number of studies included in the qualitative synthesis was 18. One-half of them were clinical trials and the others were either letters or comments. However, only one study specified the treatment.
of hypoactive subtype delirium. The design of this study was a post-hoc analysis of a double-blind, randomized, placebo-controlled study that used quetiapine as an adjuvant therapy to haloperidol. This study suggested that quetiapine appears to have more rapid resolution of many delirium symptoms, including a hypoactive state. These results were not statistically significant. The other 17 studies do not address the subtype.

Conclusion There is poor evidence regarding the use of drugs for the management of hypoactive delirium. Not only the study design but the number of patients studied in the single trial is very limited, which affects the power of evidence. Double-blind, randomized, placebo controlled trials must be performed to guide the treatment and the management of hypoactive delirium.

Acknowledgements JPLMC and RA contributed equally to this work.

Reference

Nutrition/Metabolism

P37
Effect of N-acetylcysteine on serum thyroid hormone levels in nonthyroidal illness syndrome
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Introduction Nonthyroidal illness syndrome (NTIS) refers to changes in thyroid hormone levels affecting up to 75% of critically ill patients. Cytokines and oxidative stress have been implicated as causative factors, as they derange deiodinase reactions. Interestingly, the addition of N-acetylcysteine (NAC) in a cell model prevented the effect of IL-6 on deiodinases, probably through a mechanism that restores catalytic activity of the enzyme [1]. NTIS is an independent marker of poor prognosis during myocardial infarction (MI) [2]. Here, we investigate whether NAC administration would prevent the decrease in serum thyroid hormone levels observed in patients with MI.

Methods This was a randomized, multicenter clinical trial. Patients with MI within 12 hours of evolution who underwent primary percutaneous coronary intervention were eligible. Patients were randomized to receive NAC (1,200 mg, intravenous, every 12 hours) or placebo for 48 hours. Baseline characteristics, clinical history and serial blood samples for measurements of thyroid hormones were collected. Primary outcome was the variation of serum T3 levels. Statistical tests used included χ2 test analyses for proportions and Student’s t test analyses for means.

Results Sixty-seven patients were included. There were no differences between groups with respect to baseline clinical characteristics. When compared with baseline, the levels of serum T3 decreased in the placebo group at 12 hours of follow-up (98.6 ± 86.8 μg/dL, P = 0.001), as expected for the NTIS. Interestingly, the T3 descendent curve was attenuated in patients who were randomized to NAC (100.4 ± 96.9 μg/dL, P = 0.396) (Figure 1). Similar serum T3 levels were observed in both groups at 48 hours and on the 5th day. Serum TSH levels were virtually identical between the two groups. In the comparison between groups, the mean T3 serum levels were higher in the NAC treatment group at 12 hours than in the placebo group (P = 0.045).

Conclusion This is the first clinical trial designed to investigate the effect of NAC on NTIS in humans. We show that patients with myocardial infarction who received NAC treatment on admission had attenuation of the degree of T3 level decrease compared with those who received placebo. As expected, NAC did not interfere with central feedback, as TSH levels presented a similar rise in both groups, probably as a marker of recovery from acute illness. However, whether this effect is able to prevent mortality or to ameliorate patient outcome remains to be elucidated.

Acknowledgement Clinical Trials: NCT01501110.

References

Pneumology

P38
Association between use of lung-protective ventilation with lower tidal volumes and risk of acute lung injury, mortality, pulmonary infection, and atelectasis: a meta-analysis
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Introduction Lung-protective mechanical ventilation with the use of lower tidal volumes has been found to improve the outcome of patients with acute lung injury (ALI) or its more severe form acute respiratory distress syndrome. It has been suggested that use of lower tidal volumes also benefits patients not suffering from ALI. The objective of this study was to test the hypothesis that use of lower tidal volumes is associated with improved outcomes of patients without ALI.

Methods Data source A search of MEDLINE, CINAHL, Web of Science, and Cochrane Central Register of Controlled Trials up to August 2012. Study selection Eligible studies were those evaluating use of lower versus higher tidal volumes in patients without ALI at onset of mechanical ventilation and reporting lung injury development, overall mortality, pulmonary infection, atelectasis and biochemical alterations. Data extraction Three reviewers extracted data on study characteristics, methods, and outcomes. Disagreement was resolved by consensus.

Results Twenty articles (2,822 participants) were included. Meta-analysis using a fixed-effect model showed a decrease in lung injury development (risk ratio [RR], 0.33 [95% CI 0.23 to 0.47]; number needed to treat [NNT], 1 to 11), mortality (RR, 0.64 [95% CI 0.46 to 0.89]; NNT, 1 to 23) and pulmonary infection (RR, 0.52 [95% CI 0.33 to 0.82]; NNT, 1 to 26) in patients ventilated with lower tidal volumes. The results of lung injury development were similar when stratified by the type of study (randomized vs. nonrandomized), was significant only in randomized trials for pulmonary infection and only in nonrandomized trials for mortality. The hospital length of stay was lower in the protective ventilation group (6.91 ± 2.36 vs. 8.87 ± 2.93 days, respectively; standardized mean difference, 0.60 [95% CI 0.50 to 0.71]). Protective ventilation was associated with higher PaCO2 levels (41.05 ± 3.79 vs. 37.90 ± 4.19 mmHg, respectively; SMD, −0.47 [95% CI −0.59 to −0.34]), lower pH (7.37 ± 0.03 vs. 7.40 ± 0.04, respectively; SMD, 0.75 [95% CI 0.58 to 0.92]) but similar PaO2/FiO2 (304.40 ± 65.7 vs. 312.97 ± 68.13, respectively; SMD, 0.08 [95% CI 0.00 to 0.16]). The tidal volume gradient between the two arms did not influence significantly the final results.

Conclusion Among patients without ALI, protective ventilation with lower tidal volumes was associated with a better clinical outcome. Some of the limitations of our meta-analysis were the mixed setting of mechanical ventilation (ICU or operating room) and the duration of mechanical ventilation.
Comparison of CURB-65 and CRB-65 as predictors of death in community-acquired pneumonia in adults admitted to an ICU

Introduction
Community-acquired pneumonia is one of the most common causes of sepsis and ICU admissions. Patients with CAP who demand critical care had mortality rates of 25 to 50%. Thereby, the assessment of the severity is essential to guide the treatment. There are several severity scores for CAP and some of the most acknowledged are CURB-65 and CRB-65. The objective of this study was to evaluate the accuracy of CURB-65 and CRB-65 as predictors of death in patients with community-acquired pneumonia.

Methods
A prospective study during 6 months was conducted with patients diagnosed with CAP admitted to the ICU of the Hospital Santa Luzia, Asa Sul, Brasilia, DF, Brazil. Patients were stratified according to CURB-65 and CRB-65 (0 to 5) and CRB-65 (0 to 4) and their risk categorized as: low (CURB-65: 0 to 1 and CRB-65: 0), moderate (CURB-65: 2 and CRB-65: 1 to 2) and high (CURB-65: 3 to 5 and CRB-65: 3 to 4). The sensitivity, specificity, positive predictive value (PPV), negative predictive value (NPV), likelihood ratio positive (LR+), and likelihood ratio negative (LR−) were calculated. Validity and reliability were assessed with the Spearman correlation coefficient. Patients with chronic kidney failure and those submitted to mechanical ventilation at the time of admission were excluded.

Results
A total of 62 patients were included. Twenty-seven with low risk, 24 with moderate risk and 11 with high risk according to CURB-65 and their mortality rates were 7.4%, 8.3% and 54.5%, respectively. According to CRB-65, 11 were low risk, 44 moderate risk and seven high risk. The mortality on CRB-65 stratification was 0%, 15.9% and 54.5%, respectively. The mortality on CURB-65 stratification was 0%, 15.9%, and 42.9% for low, moderate and high risks, respectively. When we exchanged between patients with the ventilator circuit.

Conclusion
This prospective randomized study included all adult patients undergoing mechanical ventilation for more than 12 hours, admitted to a 37-bed general ICU in the period from October 2011 to January 2013. The study was approved by the Ethics Committee of Hospital São Domingos under number 026/ 2011. Group 1 used HME filters TWINSTAR 55 (Drager Medical, Germany). The filter was changed every 24 hours. Group 2 used heated humidifier MR 730 (Fisher and Paykel Healthcare Inc., Auckland, New Zealand). The humidifier was exchanged between patients with the ventilator circuit.

Results
One hundred and fifty-three patients were randomized. Two were excluded: excessive airway secretion forcing frequent exchanges of HME filter (G1); length of MV <12 hours (G2). One hundred and fifty-one patients were analyzed. The two groups were comparable with regard to demographic data and severity (APACHE IV and SOFA). The main indication of MV was hypoxemic respiratory failure (G1 = 34; G2 = 35). The duration of mechanical ventilation was comparable between the two groups (G1 = 10.6 ± 19.0 days; G2 = 12.0 ± 20.7 days; P = 0.65). The incidence of VAP also did not differ significantly between groups: G1 = 7 (9.2%), G2 = 10 (13.3%); P = 0.42. No endotracheal tube occlusion was identified.

Conclusion
This study, like several others, including a meta-analysis of the Cochrane Institute (2,236 patients), failed to show that HME filters are able to reduce the incidence of ventilator associated pneumonia, compared with heated humidifiers. We had no cases of obstruction of the endotracheal tube. There is evidence in the literature that an HME filter change at intervals greater than that we used are related to a higher incidence of artificial airway occlusion. The choice of the system of humidification of inspired air in patients on mechanical ventilation (heated humidifier filter or HME) should not have the purpose to reduce the incidence of VAP.
the current evidence for use of neuromuscular blocking agents (NMBA) in the early phase of ARDS.

Methods A systematic review and meta-analysis of publications between 1966 and 2012. The Medline and CENTRAL databases were searched for studies on NMBA in patients with ARDS. The meta-analysis was limited to randomized controlled trials; adult human patients with ARDS or acute lung injury; and use of any NMBA in one arm of the study compared with another arm without NMBA. The outcomes assessed were: overall mortality, ventilator-free days, time of mechanical ventilation, adverse events, and changes in gas exchange, in ventilator settings, and in respiratory mechanics.

Results Three randomized controlled trials covering 431 participants were included. Patients treated with NMBA showed less mortality (risk ratio, 0.71 (95% CI, 0.55 to 0.90); number needed to treat, 1 to 7), more ventilator-free days at day 28 (P = 0.02), higher PaO2, to FiO2 ratios (P = 0.004), and less barotraumas (P = 0.030). The incidence of critical illness neuromyopathy was similar (P = 0.540).

Conclusion The use of NMBA in the early phase of ARDS improves outcome.

P42

Profile of reintubated patients submitted to daily weaning screen and spontaneous breathing trial in a general ICU

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Introduction Mechanical ventilation (MV) weaning is the transition of artificial ventilation to spontaneous breathing of patients intubated for more than 24 hours. Reintubation may occur, even if the weaning process has been well conducted, in 13 to 19% of the extubated patients. Daily weaning screen and spontaneous breathing trial are widely used to evaluate patients ready to be weaned, although a reintubation risk may occur [1]. The objective of this study was to verify the profile of patients that failed the weaning process and needed to be reintubated.

Methods Mechanically ventilated patients submitted to our institutional MV weaning protocol from January to July 2012, who were extubated and failed extubation within a 48-hour period, were included in the study. The weaning protocol consisted of daily weaning screen and spontaneous breathing trial. Demographic data, MV time, ICU and hospital length of stay, causes of reintubation, and mortality rate were collected during the study period.

Results Two hundred patients were included, and 29 (14%) were reintubated. Of the reintubated patients, 59% were male, with a median age of 69 years (range of 24 to 94), mean Simplified Acute Physiology Score (SAPS 3) of 60 ± 11, mean MV time of 9 days ± 5, median ICU stay of 14 days (range of 5 to 30), and 46 days of hospital stay. Causes of reintubation were acute respiratory failure (38%), low level of consciousness associated with lack of airway protection (27%), and hemodynamic instability (14%). ICU discharge occurred in 70% of the patients, and 31% were tracheostomized due to dysphagia, low level of consciousness, or lack of airway protection. The ICU mortality rate was 30%. Only one tracheostomized patient died. Patients with ages ranging from 86 to 88 years had a higher incidence of low consciousness level. Patients that did not use noninvasive ventilation (NIV) after extubation were reintubated earlier than others (median of 20 hours, P < 0.02 and r = -0.551), although there was no correlation with the use of NIV with mortality or MV time.

Conclusion The use of daily screening and spontaneous breathing trial is associated with a low reintubation rate. Acute respiratory failure and a low level of consciousness were the most common causes of reintubation, and most patients were discharged from the ICU. NIV may prevent the need for reintubation. Patients with no perspective of short-term improvement in level of consciousness may be considered for a tracheostomy.

Reference


P43

Spontaneous breathing trial reduces mechanical ventilation weaning when compared with SmartCare™ ventilation

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Introduction Mechanical ventilation (MV) weaning is commonly performed using a spontaneous breathing trial (SBT) with pressure support ventilation after a daily weaning screen [1]. Recently there has been increased interest in automatic weaning trials, using respiratory rate, tidal volume and ETCO2 monitoring during SBT [1,2]. So far there is no clinical evidence comparing an automatic weaning trial with SBT. Our study’s objective was to compare MV and weaning times between Automatic Weaning Ventilation System (SmartCare™/PS) and SBT groups.

Methods A randomized, controlled study was performed in a general ICU. We enrolled adult patients who were ventilated for more than 24 hours. Patients were randomized either to the control or SmartCare™ group. All patients were ventilated with a Drager EvitaXL (Drager Medical, Lubeck, Germany) ventilator with SmartCare™/PS software version 1.1. The control group consisted of a daily weaning screen and SBT with pressure support ventilation; if patients tolerated SBT they were extubated. SmartCare™ group patients were also submitted to a daily weaning screen, after which they were ventilated with the SmartCare™/PS mode. We evaluated the MV and weaning time, maximum inspiratory pressure, maximum expiratory pressure, tidal volume and ETCO2 monitoring during SBT [1,2]. So far there is no clinical evidence comparing an automatic weaning trial with SBT. Our study’s objective was to compare MV and weaning times between Automatic Weaning Ventilation System (SmartCare™/PS) and SBT groups.

Methods A randomized, controlled study was performed in a general ICU. We enrolled adult patients who were ventilated for more than 24 hours. Patients were randomized either to the control or SmartCare™ group. All patients were ventilated with a Drager EvitaXL (Drager Medical, Lubeck, Germany) ventilator with SmartCare™/PS software version 1.1. The control group consisted of a daily weaning screen and SBT with pressure support ventilation; if patients tolerated SBT they were extubated. SmartCare™ group patients were also submitted to a daily weaning screen, after which they were ventilated with the SmartCare™/PS mode. We evaluated the MV and weaning time, maximum inspiratory pressure, maximum expiratory pressure, tidal volume and ETCO2 monitoring during SBT [1,2]. So far there is no clinical evidence comparing an automatic weaning trial with SBT. Our study’s objective was to compare MV and weaning times between Automatic Weaning Ventilation System (SmartCare™/PS) and SBT groups.

Results We evaluated a total of 70 patients (35 patients randomized in each group). There was no difference in age (P = 0.298) or gender (P = 0.08) between groups (Table 1). There was no difference in MV time between the control and SmartCare group (P = 0.534) (Table 1). Weaning duration was lower in the control group (Pf/VT, P = 0.414), use of NIV post extubation (P = 0.811) and re-intubation rate (P = 1.0) (Table 1).

Table 1 (abstract P43). Characteristics of patients between SmartCare™ and control groups

<table>
<thead>
<tr>
<th></th>
<th>SmartCare™</th>
<th>Control</th>
<th>P value*</th>
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<tbody>
<tr>
<td>Idade</td>
<td>60 (46 to 77)</td>
<td>65 (57 to 81)</td>
<td>0.298</td>
</tr>
<tr>
<td>Gender, male</td>
<td>23 (65)</td>
<td>16 (45)</td>
<td>0.07</td>
</tr>
<tr>
<td>MP</td>
<td>45 (40 to 53)</td>
<td>40 (36 to 50)</td>
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</tr>
<tr>
<td>MEP</td>
<td>40 (30 to 59)</td>
<td>40 (21 to 44)</td>
<td>0.059</td>
</tr>
<tr>
<td>VC</td>
<td>1,200 (900 to 1,850)</td>
<td>1,000 (500 to 1,600)</td>
<td>0.834</td>
</tr>
<tr>
<td>f/VT</td>
<td>35 (24 to 55)</td>
<td>40 (26 to 68)</td>
<td>0.414</td>
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<tr>
<td>MV time</td>
<td>4 (2 to 6)</td>
<td>3 (2 to 7)</td>
<td>0.534</td>
</tr>
<tr>
<td>Weaning time</td>
<td>110 (80 to 120)</td>
<td>60 (50 to 80)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Use of NIV post extubation</td>
<td>18 (51.4)</td>
<td>16 (45.7)</td>
<td>0.811</td>
</tr>
<tr>
<td>Reintubation rate</td>
<td>2 (5.7)</td>
<td>2 (5.7)</td>
<td>1.00</td>
</tr>
</tbody>
</table>

Data presented as median (interquartile range) or n (%). f/VT, respiratory rate to tidal volume ratio; MEP, maximum expiratory pressure; MP, maximum inspiratory pressure; MV, mechanical ventilation; NIV, noninvasive ventilation; VC, vital capacity. *P value significant <0.05.

Conclusion SBT showed a reduction in weaning time when compared with the SmartCare™/PS group, although there was no impact on total MV time and reintubation rate.

References

Impact of acute brain dysfunction on the outcomes of mechanically ventilated cancer patients

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Introduction Delirium and coma are a frequent source of morbidity for ICU patients. Several factors are associated with the prognosis of mechanically ventilated (MV) cancer patients, but no studies have evaluated delirium and coma (acute brain dysfunction). The present study evaluated the frequency and impact of acute brain dysfunction on hospital mortality.

Methods The study was performed at the National Cancer Institute, Rio de Janeiro, Brazil. We prospectively enrolled patients ventilated >48 hours with a diagnosis of cancer. The presence of acute brain dysfunction was assessed daily during the first 14 days of ICU care using the CAM-ICU. Patients were followed until hospital discharge. Univariate and multivariate analysis were performed to evaluate factors associated with hospital mortality.

Results A total 170 patients were included. Seventy-three percent had solid tumors, age 65 (53 to 72 (median, IQR 25 to 75%)) years. Admission SAPS II score was 54 (46 to 63) points and SOFA score was 7 (6 to 91) points. Median duration of MV was 13 (6 to 21) days and ICU stay was 14 (7.5 to 22) days. ICU mortality was 54% and hospital mortality was 66%. Acute brain dysfunction was diagnosed in 161 patients (95%). Nonsurvivors had a higher frequency of acute brain dysfunction (110 (97.3%) vs. 51 (89.4%)), P = 0.06. Survivors had more delirium/coma-free days (4 (1.5 to 6) vs. 1 (0 to 2), P = 0.771 (0.681 to 0.873), P < 0.001).

Conclusion Acute brain dysfunction in MV cancer patients is frequent and independently predictive of increased hospital mortality. Future studies should investigate means of preventing or mitigating acute brain dysfunction as they may have a significant impact on clinical outcomes.

Acknowledgements Supported by the National Cancer Institute, CNPq and FAPERJ.

Sepsis

Comparative analysis of survival between older and nonolder severe sepsis and septic shock resuscitated patients

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Introduction Advanced age has been associated with increased mortality in severe sepsis and septic shock patients [1,2]. However, the impact of early resuscitation following the Surviving Sepsis Campaign Guidelines in this population of patients is unclear. The objective of this study was to compare the in-hospital mortality between older (EP) and nonolder (N-EP) resuscitated patients according to the Surviving Sepsis Campaign Guidelines.

Methods A retrospective observational study. All patients with severe sepsis and septic shock admitted to the ICU between January 2006 and March 2012 were studied. Comparisons were performed between older (≥65 years) and nonolder patients (<65 years).

Results A total of 913 patients with severe sepsis and septic shock were included in this analysis. Older patients accounted for 63% (573/913) of patients and nonolder patients for 37% (340/913) of patients. The median (IQR) age was 80 years (73 to 85) and 51 years (40 to 59) for EP and N-EP, respectively. The incidence of severe sepsis (43% vs. 44%) and septic shock (57% vs. 56%) did not differ between the EP and N-EP groups (P = 0.78). EP patients had a higher median (IQR) APACHE II score (23 (18 to 28) than N-EP patients (19 (16 to 24), P < 0.001), although the median number of organ dysfunctions (3 vs. 2 for EP and N-EP, respectively, P = 0.57) did not differ between the groups. EP patients were more likely to have hypertension (51% vs. 29%, P < 0.001), diabetes (33% vs. 24%, P = 0.02), ischemic heart disease (16% vs. 7%, P < 0.001) and chronic renal failure (8.5% vs. 4.2%, P < 0.03) when compared with N-EP patients. Solid organ transplantation (24% vs. 4%, P < 0.001) and liver cirrhosis (17% vs. 5%, P < 0.001) were more frequently in N-EP patients. There was no significant between-group difference in the in-hospital mortality (33% in the EP group and 28% in the N-EP group; odds ratio, 1.27; 95% CI, 0.94 to 1.70; P = 0.12) (Figure 1). The length of hospital stay (14 (7 to 29) vs. 12 (6 to 21) days (median (IQR)), P = 0.001) was significantly higher in EP patients compared with the N-EP patients.

Conclusion In this population of severe sepsis and septic shock patients, the early resuscitation of older patients was not associated with increased mortality. However, prospective studies addressing the long-term impact of the resuscitation maneuvers on outcomes are necessary.

References

Effect of vasoactive drugs on the response of the baroreceptor and regulation of heart rate variability in patients with septic shock

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Introduction We evaluated the action of vasoactive drugs on baroreceptor regulation and the interaction of this in regulation of heart rate variability (HRV) in patients with septic shock.

Methods A prospective observational study of patients with severe sepsis or septic shock. Collected data were analyzed retrospectively, separating patients according to their clinical evolution: survival or death. We monitored troponin I, vasoactive drug dose, HRV and blood pressure variability.

Results The study included 31 patients, of whom 14 died. Increased troponin levels were related to an increased risk of mortality. The alpha index of HRV low frequency and high frequency indicates that the interaction of baroreceptor in the regulation of heart rate variability with changes in blood pressure had a significant reduction in patients with septic shock and death. The use of dobutamine in patients with
septic shock correlates well with troponin levels \((r = 0.77)\), while for norepinephrine the correlation was poor \((r = 0.53)\). The use of dobutamine showed a negative correlation with the low-frequency alpha index (index that assesses the integration of baroreceptor); on the other hand, norepinephrine showed a positive correlation. See Table 1 and Figure 1.

Table 1 (abstract P46)

| Patients, n | 30 |
| Gender, female | 12 (40%) |
| Age (years) | 64 ± 5 |
| Nonsurvivors | 14 (47%) |

Infectious site

| Lung | 18 (60%) |
| Abdominal tract | 5 (15%) |
| Urinary tract | 4 (14%) |
| Blood | 2 (8%) |
| Nonidentified | 1 (4%) |
| Gram-negative | 22 (76%) |

Conclusion Patients with septic shock have impaired baroreceptor function, and this is correlated with progression to death. Dobutamine is related to higher levels of cardiac damage and higher doses of dobutamine interfere with the responsiveness of the baroreceptor. Moreover, norepinephrine has a positive effect on the integration of baroreceptor.

P47

Erythrocyte selenium concentration is a predictor of mortality in patients with septic shock

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Introduction Severe sepsis and septic shock are major healthcare problems, affecting millions of people around the world each year, and increasing in incidence [1]. During sepsis there is increased oxidative stress, with reduced body stores of selenium (Se) and lower activity of glutathione peroxidase (GPx1), with patient outcomes of multiple organ failure and death [2]. The objective of this study was to determine the influence of Se concentration in plasma, erythrocytes and erythrocyte GPx1 activity on the length of hospital stay, length of ICU stay and ICU mortality in patients with septic shock.
Methods This prospective study included all patients with septic shock on admission or during ICU stay, over the age of 18, admitted to one of the three ICUs of the Botucatu Medical School, from January to August 2012. Demographic information, clinical evaluation and blood samples were taken within the first 72 hours of the patient’s admission or within 72 hours after septic shock diagnosis for laboratory analysis, GPx activity, plasma and erythrocyte Se concentration. Categorical variables were analyzed by chi-squared or Fisher exact test. Continuous variables were analyzed by Student’s t test. For length of ICU or hospital stay, prediction multiple linear regression was used. For mortality prediction, multiple logistic regression was performed. The level of significance was set at 5%.

Results We evaluated 110 patients with a mean age of 58 ± 16 years and 63% were male. The median length of ICU stay and hospital stay was 9 (5 to 15) and 11 (11 to 34) days, respectively, and the ICU mortality rate was 55%. Patients had an average plasma and erythrocyte Se concentration of 23.37 ± 8.99 and 32.83 ± 11.89 g/l, respectively, and the median GPx1 activity was 30.56 (23.88 to 38.41) U/g Hb. All patients had Se deficiency; however, only 25% had reduced GPx activity. There was no association of plasma and erythrocyte Se concentration and GPx activity with the length of hospital and ICU stay. Higher values of APACHE II, albumin and creatinine were associated with higher erythrocyte Se concentration. Regarding mortality, it was associated with higher lactate, urea, APACHE II and SOFA scores and lower values of albumin and length of hospital and ICU stay. In multiple logistic regression analysis adjusted for age, gender, and APACHE II, the erythrocyte Se concentration was a predictor of mortality in patients with septic shock.

Conclusion The erythrocyte Se concentration is a predictor of mortality in patients with septic shock.

Acknowledgement Supported by CAPES.

References

P48
Evaluation of recognition and signalling receptors on the peripheral blood cells of septic patients and their correlation with clinical outcomes
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Introduction
Monocytes and neutrophils play a key role in host defence by sensing and destroying microorganisms [1]. We evaluated the expression of cellular receptors implicated in pathogen recognition, cell activation and migration on both cell types during sepsis. Blood samples were collected from 77 septic patients (SP) at admission (D0), from 45 patients after 7 days of therapy (D7) and from 40 healthy volunteers (HV).

Results
The expression of CD14 on monocytes and of CD11b and CXCR2 on neutrophils from SP was lower than that from HV. Conversely, the expression of TLR5 on monocytes and neutrophils was higher in SP when compared with HV. The expression of TLR2 on the surface of neutrophils and that of TLR5 on monocytes and neutrophils of SP was lower at D7 than at D0. In addition, the SP that survived showed reduced expression of TLR2 and TLR4 on the surface of neutrophils at D7 compared with D0 (Figure 1). The expression of CXCR2 for surviving patients was higher at follow-up when compared with baseline (Figure 1).

Conclusion
The expression of recognition and cell signalling receptors is differentially regulated between SP and HV depending on the receptor being evaluated. However, despite these changes, it is likely that the functional changes in monocytes and neutrophils that are observed during sepsis are not directly linked to the modulation of the expression of TLRs [2].

References

P49
Gene expression in peripheral mononuclear cells from septic patients secondary to community-acquired pneumonia: patterns of gene expression and outcomes
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Introduction
Sepsis is defined as a systemic inflammatory response secondary to a proven or suspected infection. Mechanisms governing this inflammatory response have been shown to be complex and dynamic, involving cross-talk among diverse signaling pathways. However, current knowledge on mechanisms underlying sepsis is far from providing a complete picture of the syndrome, justifying additional efforts that might add to this scenario. Microarray-based expression profiling is a powerful approach for the investigation of complex clinical conditions such as sepsis: the analysis of gene transcription at the genome level potentially avoids results derived from biased assumptions. In this study we investigate whole-genome...
gene expression profiles of mononuclear cells from survivor and non-survivor septic patients. **Methods** Blood samples were collected at the time of sepsis diagnosis and 7 days later, allowing us to evaluate the role of biological processes or genes possibly involved in patient recovery. Aiming to circumvent, at least partially, the heterogeneity of septic patients, we included only patients admitted with sepsis caused by community-acquired pneumonia. Global gene expression profiling allowed us to characterize early sepsis, as compared with healthy individuals. **Results** Our results corroborate literature reports on inflammation response in the early stages of sepsis but highlight great heterogeneity in gene expression during the onset of sepsis. Differences in oxidative stress seem to be associated with clinical outcome, since significant differences in the expression profile of related genes were observed between survivors and nonsurvivors. However, our results substantiate current knowledge supporting that sepsis syndrome development is indeed multifaceted. Although the initial infection of enrolled patients was pneumonia, with no sign of organ failure at the time of diagnosis, 7 days later gene expression profiles seemed to be characteristic for each patient, with no clear pattern of development. This result could be associated with the underlying health status of each one of them, with complications due to sepsis itself as well as with distinct timing for response to treatment.

**Conclusion** At this point we conclude that studies should focus on studying sepsis at multiple time points, aiming to capture common, although possibly separated in time when patients are considered, biological processes associated with recovery.

**PS0 Georeferencing of sepsis in São Paulo**
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**Introduction** Morbidity and lethality in sepsis varies according to demographic and socioeconomic factors, which can be linked to geographic distribution of populations. The objective of this study is to relate sepsis deaths with geographic distribution in the city of São Paulo, Brazil.

**Methods** Death certificates for sepsis and sepsis-related infections (pneumonia, urinary tract infections, meningitis, skin and soft tissue infections, peritonitis, multiple organ failure) from 2004 to 2009 in the city of São Paulo, Brazil, were searched. Patients were distributed according to sex, age, main cause of death, secondary cause of death, residence address, and death location. The Human Development Index (HDI) was used to compare city districts. Health institutions were identified as public or private administered according to Ministry of Health registration.

**Results** The number of deaths increased with age, but there is no difference between sexes for the whole population studied. However, in younger age groups (up to 18 years and from 18 to 64 years) deaths were more frequent for males than females (53.9 ± 46.1% and 61 ± 39%, respectively). In the age group older than 65 years there were 46.3% of deaths in males and 53.7% for females. Deaths occurred homogeneously in all city districts, but its frequency was larger in districts with higher HDI. Death location was 93.73% in hospitals and 5.47% in dwellings. From the identifiable institutions where deaths occurred, 52.4% were public, 46.4% were private and 1.2% were nonidentifiable. The mean age at death was 79% higher in public institutions compared with private ones. There was no visible difference in death distribution according to age and residence address. Comparing patients’ residence address with death location showed a concentration near the closest hospital, without difference between private or public hospitals. There was a higher proportion of deaths for the age group older than 65 years in private hospitals. For the age group between 19 and 64 years, the proportion of deaths was slightly higher in public hospitals. The main cause of death was pneumonia, followed by sepsis, multiple organ failure, intra-abdominal infections, meningitis, skin and soft tissue infections and urinary tract infections. Distributing main death causes by age groups, patients older than 65 years died more over pneumonia, sepsis and multiple organ failure. Patients up to 18 years died more from urinary tract infections and meningitis. The age group between 19 and 64 years was distributed fairly amongst all death causes.

**Conclusion** Georeferencing is a potent tool for epidemiological studies. Deaths occurred homogeneously in all city districts, but the frequency was larger in districts with higher HDI. As expected, older patients died in greater numbers, affected by respiratory tract infections, sepsis and multiple organ failure. Deaths occurred mainly in the public health system.

**PS1 Lactate as a prognostic marker in patients with severe sepsis or septic shock admitted to the ICU**
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**Introduction** Severe sepsis and septic shock are conditions associated with high morbimortality worldwide despite improvements in its management in the last two decades. It is of great interest for intensive care physicians to have the opportunity to use biomarkers for prediction of severity of disease and prognostication in a way that the more severely ill the patient, the more aggressive the treatment should be. Lactate is globally available benchside. The presence of elevated lactate levels are associated with increased mortality in distinct critically ill patient populations. Although lactate levels above 4 mmol/l are the classical trigger for early-goal directed therapy, there are some studies showing that even normal or slightly elevated lactate levels are associated with worse outcome. The objective of this study is to evaluate stratified lactate serum levels at admission as a predictor of 28-day mortality.

**Methods** A retrospective cohort study where data were collected from electronic charts of adult patients diagnosed with severe sepsis or septic shock admitted to our ICU between July 2005 and December 2010. Data collected included the following: social and demographic, presence of organic dysfunction at admission, initial lactate level, APACHE II score, compliance with the institutional sepsis protocol, use of mechanical ventilation and 28-day mortality. Lactate levels were expressed in mg/dl and stratified into four quartiles: (1) normal, <14.4 mg/dl; (2) mild elevated, 14.5 to 28 mg/dl; (3) intermediate elevated, 28.1 to 36 mg/dl; and (4) high elevated, >36 mg/dl. The study protocol was approved by the local ethics committee. Categorical variables were expressed as frequencies and/or percentages and continuous as mean ± standard deviation. For proportions and means comparisons, the chi-square and Student t tests were used, respectively. Statistical significance was set as P <0.05. A univariate analysis was performed and statistically significant variables were included in a logistic regression model. The SPSS® software (IBM Corporation, USA) was used to perform the tests.

**Results** A total of 760 patients were included, with mean age of 67.2 ± 18.66 years and 57.9% male. Mean APACHE II score was 21 and 46.4% were in mechanical ventilation and 64.7% were using vasopressors. Global mortality was 36.7%. An intermediate compliance with the institutional sepsis protocol was observed. The best stratum to predict mortality in our study was the high elevated lactate level group (P <0.0001).

**Conclusion** Lactate levels are one of the most used biomarkers in sepsis. When their level is superior to 36 mg/dl patients are at highest risk of mortality and an aggressive resuscitation strategy shall be warranted in these patients.

**PS2 Procalcitonin as a prognostic biomarker of severe sepsis and septic shock**
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**Introduction** Procalcitonin (PCT), the precursor peptide of calcitonin, has extremely low levels in healthy subjects. In response to bacterial infectious stimulation, PCT serum levels rise substantially. Recently PCT has been used as a biomarker for prognosis of severe sepsis and septic shock. Some studies have shown that isolated high levels do not predict outcome. Encouraging results were obtained with the...
evaluation of serum PCT levels. The objective of this study was to evaluate the tendency of the plasma concentration and clearance of PCT as biomarkers for prognosis of patients with severe sepsis and septic shock, compared with another early prognosis marker, the number of SIRS criteria at sepsis diagnosis.

**Methods** We conducted a prospective, observational, cohort study, with patients with severe sepsis and septic shock. The serum procalcitonin was determined at diagnosis of sepsis and after 24 and 48 hours. Demographic data, APACHE IV score, SOFA score on arrival, number of SIRS criteria at diagnosis, site of infection and microbiological results were recorded.

**Results** Twenty-eight patients were included, 19 clinical and nine surgical. In 13 patients (46.4%) the source of sepsis was pulmonary, abdominal in seven cases (25.0%), urinary in five cases (17.9%) and soft tissue in three cases (10.7%). Fifteen patients had severe sepsis and 13 septic shock. Overall mortality was 17.9% (five patients), three with septic shock. Twenty-eight PCT determinations were performed at sepsis diagnosis, 27 after 24 hours and 26 after 48 hours. The initial concentration was not significantly different between the survivor and nonsurvivor groups, but the differences between the two groups after 24 and 48 hours were statistically significant. There was no difference in the number of SIRS criteria. The 24-hour procalcitonin clearance proved to be significantly higher in the group of survivors (~3.0 vs. ~300.0, P = 0.028). Although the 48-hour procalcitonin clearance was shown to be higher in the group of survivors when compared with nonsurvivors, the difference did not reach statistical significance.

**Conclusion** Persistently high PCT concentrations in plasma, as well as reduced 24-hour PCT clearance, were associated with a significant increase in mortality in patients with severe sepsis and septic shock.

**P53**

**Serum C-reactive protein concentration in early abdominal and pulmonary sepsis**

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**Introduction** The objective of this study was to evaluate the C-reactive protein serum levels in patients with pulmonary and abdominal sepsis during the first 5 days of sepsis progression.

**Methods** The present investigation was a retrospective cohort study conducted at the university hospital with 345 patients who were admitted to the ICU and diagnosed with sepsis of pulmonary or abdominal origin. Serum C-reactive protein concentrations were measured by the turbidimetric immunoassay. For analysis of C-reactive protein, day 1 was defined as the day on which the patient was clinically diagnosed with sepsis.

**Results** Thirty-four patients with sepsis (9.8%), 114 patients with severe sepsis (33.0%), and 197 patients with septic shock (57.2%) were evaluated. The age of the patients was 56.4 ± 19.8 years. The serum C-reactive protein concentrations were higher on the day of sepsis diagnosis in the group with abdominal infection compared with the group with pulmonary sepsis (17.8 ± 10.1 mg/dl vs. 14.9 ± 11.1 mg/dl; P = 0.025) and remained significantly higher during the first 5 days of sepsis progression. The values of the area under the ROC curve, sensitivity, specificity, and best cutoff points are listed in Table 1.

**Table 1 (abstract P53). Values of the area under the ROC curve, sensitivity, specificity, and best cutoff points**

<table>
<thead>
<tr>
<th>ROC curve</th>
<th>AUC (95% CI)</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>Cutoff (mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CRP day 1</td>
<td>0.53 (0.43 to 0.62)</td>
<td>0.26</td>
<td>0.88</td>
<td>6.12</td>
</tr>
<tr>
<td>CRP day 2</td>
<td>0.51 (0.41 to 0.60)</td>
<td>0.67</td>
<td>0.42</td>
<td>18.6</td>
</tr>
<tr>
<td>CRP day 3</td>
<td>0.52 (0.43 to 0.61)</td>
<td>0.57</td>
<td>0.56</td>
<td>13.6</td>
</tr>
<tr>
<td>CRP day 4</td>
<td>0.51 (0.41 to 0.60)</td>
<td>0.42</td>
<td>0.65</td>
<td>14.2</td>
</tr>
<tr>
<td>CRP day 5</td>
<td>0.59 (0.48 to 0.68)</td>
<td>0.58</td>
<td>0.62</td>
<td>9.10</td>
</tr>
</tbody>
</table>

AUC, area under the ROC curve; CI, confidence interval; CRP, C-reactive protein; ROC, receiver operator characteristic.

**Conclusion** The accuracy of C-reactive protein for the differential diagnosis of pulmonary and abdominal sepsis is limited, although significantly higher serum levels were observed in patients with abdominal sepsis.

**P54**

**Serum pro-metalloproteinase 9 is a predictor of length of ICU and hospital stay in patients with septic shock**

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**Introduction** The matrix metalloproteinases (MMPs) participate in fundamental processes, such as cell proliferation, differentiation, adhesion, migration, angiogenesis, apoptosis, and inflammation [1]. The increased expression of MMPs suggests that these proteases may influence the pathogenesis of endotoxia in sepsis [2]. The objective of this study was to evaluate the serum activity of MMP-2 and MMP-9, length of hospital stay, length of ICU stay and mortality in patients with septic shock.

**Methods** This prospective study included all patients with septic shock on admission or during ICU stay, over the age of 18 years, admitted to the ICU from March to July 2012. Demographic information, clinical evaluation and blood samples were taken within the first 72 hours after the patient’s admission or within 72 hours after septic shock diagnosis for laboratory analysis and MMP-2 and MMP-9 activity. The activity of MMPs was performed byzymography. The level of significance was set at 5%.

**Results** We evaluated 67 patients with a mean age of 56 ± 15 years, 66% male, the median length of ICU and hospital stay was 9 (4 to 15) and 16 (10 to 29) days, respectively, and the ICU mortality rate was 61%. Higher values of APACHE II, SOFA, lactate and urea, and lower values of albumin, length of ICU and hospital stay were associated with ICU mortality. In univariate analysis, serum activity of MMP-2 and MMP-9 were not associated with length of ICU, hospital stay and mortality in septic shock patients. However, in the regression model analysis when adjusted for sex, age, lactate and APACHE II, the activity of pro-MMP-9 was negatively associated with the length of ICU (coefficient: −0.016; P = 0.028) and hospital (coefficient: −0.015; P = 0.041) stay, but was not associated with ICU mortality (OR: 1.051; 95% CI: 0.950 to 1.163; P = 0.332).

**Conclusion** Serum activity of pro-MMP-9 was negatively associated with length of ICU and hospital stay in patients with septic shock.

**Acknowledgement** Supported by CAPES.

**References**


**P55**

**Serum thiamin concentration is negatively correlated with lactate levels in survivors of septic shock**

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**Introduction** Thiamine deficiency can be present in 20% of patients in the ICU. This deficiency is considered to be an uncommon source of lactic acidosis in septic patients. An elevated serum lactate level is associated with morbidity and mortality [1]. Increased levels of thiamine increase the activity of glutathione peroxidase (GPx), a major component of the cellular antioxidant system [2]. The objective of this study was to determine the influence of serum thiamine concentrations on lactate levels, GPx activity, length of hospital stay, length of ICU stay and ICU mortality in patients with septic shock.

**Methods** This prospective study included all patients with septic shock on admission or during ICU stay, over the age of 18, admitted to one of the three ICUs of the Botucatu Medical School from January to August.
2012. Demographic information, clinical evaluation and blood samples were taken within the first 72 hours of the patient's admission or within 72 hours after septic shock diagnosis for laboratory analysis, serum thiamine and GPx activity determination. The level of significance was set at 5%.

Results One hundred and eight consecutive patients were evaluated. The mean age was 57.5 ± 16.0 years, 63% were male and 54.6% died in the ICU. The frequency of thiamine deficiency was 71.3%. Neither the serum thiamine concentration nor the erythrocyte GPx activity was associated with mortality in septic shock patients. Thiamine levels were also not associated with GPx activity ($r = 0.141$, $P = 0.165$). In addition, thiamine levels were not associated with serum lactate in the 108 patients with septic shock ($r = –0.074$, $P = 0.444$). However, vitamin B1 levels were negatively associated with lactate in patients who survived ($r = –0.311$, $P = 0.029$). In the regression model analysis, vitamin B1 levels were not associated with ICU mortality or with the length of the ICU or hospital stay.

Conclusion Thiamine deficiency is common in septic shock patients. Furthermore, thiamine was negatively associated with lactate levels in survivors of septic shock.

Acknowledgement Supported by CAPES.

References

P56
Tissular perfusion influence on central, mixed and atrial venous oxygen saturations
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¹Hospital São Paulo, Universidade Federal de São Paulo, SP, Brazil; ²University of São Paulo, Universidade Federal de São Paulo, SP, Brazil

Introduction Even though there has been quite a discussion on whether venous oxygen saturations are useful to guide treatment during initial resuscitation of sepsis, using mixed and central venous oxygen saturations as goals is still advised in the Surviving Sepsis Campaign under strong recommendation but a low level of evidence (1C). According to these guidelines, $\text{SvO}_2 < 65\%$ or $\text{SvO}_2 < 70\%$ demands treatment. In addition, there is no consensus whether these variables are interchangeable. The objective of this study was to evaluate the influence of tissular perfusion on the correlation between the central venous ($\text{SvO}_2$), the mixed venous ($\text{SmO}_2$) and the atrial oxygen saturations ($\text{SvO}_2$) by the analysis of arterial lactate.

Table 1 (abstract P56). Baseline

<table>
<thead>
<tr>
<th>Variable</th>
<th>Group 1 ($n=37$)</th>
<th>Group 2 ($n=28$)</th>
<th>$P$ value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>66 (54 to 69)</td>
<td>65 (54 to 69)</td>
<td>0.171</td>
</tr>
<tr>
<td>Male (%)</td>
<td>35.1 (13)</td>
<td>85.7 (24)</td>
<td>0.000</td>
</tr>
<tr>
<td>APACHE II</td>
<td>18 (17 to 22)</td>
<td>24 (20 to 29)</td>
<td>0.016</td>
</tr>
<tr>
<td>SOFA admission</td>
<td>6 (5 to 10)</td>
<td>10.5 (7 to 12.75)</td>
<td>0.007</td>
</tr>
<tr>
<td>SOFA sample</td>
<td>9 (7 to 11)</td>
<td>11 (9.25 to 16)</td>
<td>0.007</td>
</tr>
<tr>
<td>$\text{SvO}_2$ (%)</td>
<td>72 (68.5 to 76.5)</td>
<td>70.5 (67 to 73)</td>
<td>0.182</td>
</tr>
<tr>
<td>$\text{SmO}_2$ (%)</td>
<td>81 (76 to 85)</td>
<td>77 (72 to 80)</td>
<td>0.016</td>
</tr>
<tr>
<td>$\text{SvO}_2$ (%)</td>
<td>77.9 ± 8.6</td>
<td>74.3 ± 9.7</td>
<td>0.274</td>
</tr>
<tr>
<td>SaO (%)</td>
<td>98 (97 to 98.5)</td>
<td>95 (93 to 96)</td>
<td>0.000</td>
</tr>
</tbody>
</table>

Results expressed as % ($n$) or mean ± standard deviation or median (25 to 75% percentiles). APACHE, Acute Physiologic Chronic Health Evaluation; SOFA, Sequential Organ Failure Assessment.

Table 2 (abstract P56). Spearman correlation ($r$)

<table>
<thead>
<tr>
<th>Variable</th>
<th>Total ($n=65$)</th>
<th>Group 1 ($n=37$)</th>
<th>Group 2 ($n=28$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\text{SvO}_2 \times \text{SvO}_2$</td>
<td>0.74*</td>
<td>0.66*</td>
<td>0.83*</td>
</tr>
<tr>
<td>$\text{SvO}_2 \times \text{SmO}_2$</td>
<td>0.68*</td>
<td>0.60*</td>
<td>0.82*</td>
</tr>
<tr>
<td>$\text{SvO}_2 \times \text{SvO}_2$</td>
<td>0.72*</td>
<td>0.63*</td>
<td>0.85*</td>
</tr>
</tbody>
</table>

* $P<0.05$.
Methods A prospective observational study; the populations from three ICUs of the Hospital São Paulo were evaluated from October 2011 to November 2012 and patients diagnosed with severe sepsis or septic shock monitored by pulmonary artery catheter (PAC) were included. Hyperlactatemia was defined as an arterial lactate value >28 mg/dl and the correct location of the PAC was confirmed by chest radiography and pulmonary artery pressure tracings. For the statistical analysis, samples were allocated into two groups: normal lactate levels (Group 1) and hyperlactatemia (Group 2). Results were expressed in mean ± standard deviation or median (25 to 75% percentiles) or percentages.

Results Twenty-one patients were included; altogether, 65 paired blood samples were obtained (Table 1). A higher correlation between the venous oxygen saturations was found in the hyperlactatemia group (Table 2). APACHE II and SOFA scores were higher among these individuals (Table 1). SvCO₂ and SvO₂ were shown not to be acceptable surrogates by the analysis of the Bland–Altman plots, but bias and limits of agreement were narrower in Group 1 (Figures 1 and 2).

Conclusion In patients with hyperlactatemia, a global tissular perfusion marker, venous oxygen saturations presented a higher correlation and narrower bias and limits of agreement, suggesting, perhaps, that under high arterial lactate levels there is a generalized hypoperfusion that reflects not only on the SvO₂, but also on the SvCO₂. There was no agreement between those variables either.

References
Role of hypothermia in the immediate postoperative period on mortality in a surgical ICU

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Introduction

Surgical patients are submitted to numerous factors that may cause postoperative hypothermia, such as a cool operating room environment, cold intravenous fluids and blood, cold antiseptic skin preparations and anesthetic-induced impairment of thermoregulatory control. Hypothermia may increase susceptibility to surgical wound infection, length of stay, intraoperative blood loss, morbid cardiac events, postoperative shivering, coagulopathy and also altered duration of drug action. The objective of this study was to evaluate the impact of hypothermia at ICU admission on hospital length of stay and mortality in a surgical ICU.

Methods

A prospective cohort study conducted on patients admitted to the ICU of Hospital Santa Luzia, Brasília, Brazil, during the period of 1 year. Hypothermia was defined as axillary temperature inferior to 35.5°C (95.9°F). Patients were divided into groups with hypothermia (HG) and without hypothermia (NG).

Results

A total of 484 patients were enrolled. Mean age was 59 ± 16 years and 52.5% were male. Seventy-eight patients (16.1%) were submitted to emergency surgeries. Mean APACHE II score was 8 ± 5, mean SAPS II was 16 ± 10. Twenty-four patients (5%) had hypothermia at the time of ICU admission. The general mortality rate at 7 days, 28 days and hospital mortality was 0.8% (n = 4), 1.9% (n = 9) and 4.1% (n = 20), respectively. Patients with hypothermia had higher APACHE II score (15 ± 12 vs. 8 ± 4, P = 0.00) and SAPS II (25 ± 17 vs. 26 ± 10, P = 0.00). There was no difference regarding the age of the groups and the hospital length of stay (4 ± 6 days in NG vs. 5 ± 12 days in HG, P = 0.76). However, the group with hypothermia had higher mortality rates (20% vs. 4.3%, P = 0.00). The relative risk for hospital mortality in patients with hypothermia at ICU admission was 4.6 (95% CI: 1.02 to 29.88).

Conclusion

Few patients were hypothermic at the time of ICU admission in the immediate postoperative period. This may reflect the effectiveness of perioperative warming of the patients. Hypothermia at ICU admission was associated with greater severity scores and increased hospital mortality in this sample of surgical patients studied.

References


Cite abstracts in this supplement using the relevant abstract number, e.g.: Santana AR, et al.: Role of hypothermia in the immediate postoperative period on mortality in a surgical ICU [abstract]. Critical Care 2013, 17(Suppl 3):P59.