local infiltration of ropivacaine, a day after knee joint arthroplasty operations with spinal anesthesia.

**Introduction:** Inadequately chosen postoperative anesthesia method after knee joint arthroplasty surgery might cause prolonged hospitalization period, readmissions due to pain and overall increased cost of care.

**Methods:** In 2016 a prospective research was conducted in Vilnius University Hospital Santaros Clinics. 25 patients undergoing knee joint arthroplasty surgery with spinal anesthesia were enrolled in the study. Group1 – local soft tissue ropivacaine infiltration anesthesia around the knee (n = 13; dose 300 mg); Group2 – intrathecal morphine sulfate analgesia (n = 12; dose 0.1–0.2 mg). Pain intensity (using VAS) at rest and in motion, patient’s satisfaction and side effects - nausea, vomiting, itch, urinary retention – were assessed at time intervals – 1, 2, 4, 6, 12, 18, 24 h postoperatively.

**Results:** In the first 12 h mean values of VAS were 1.8 ± 2.6/1.4 ± 1.7 in Group1 and Group2 accordingly. After 12 h period a downturn occurred and values were 1.7 ± 1.1/1.1 ± 1.5, respectively (p > 0.05). Examining pain in motion 12h after the surgery pain intensity values were 2.5 ± 2.7/3.3 ± 2.7 and after 24 h in both groups pain intensity was 3.2 ± 1.5/3.6 ± 2.1, resp. (p > 0.05). Zero episodes of nausea/vomiting were registered in Group1, while 58.3% (n = 7) of Group2 patients experienced nausea and 5 of them also vomited. Even 66.7% (n = 8) patients in Group2 had itch while none patients of Group1 indicated this side effect. It was difficult to assess urinary retention as 30.8% (n = 4) Group1 and 66.7% (n = 8) Group2 patients were catheterized prior surgery. Finally, satisfaction level of both groups were evaluated very similarly: 8.2 ± 1.7/8.2 ± 1.3 (p > 0.05).

**Conclusion:** VAS values at rest were very similar in both groups, but pain relief efficiency compared to the intensity of pain during movement was better with local ropivacaine infiltration, also patients with ropivacaine analgesia experienced no side effects.

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**PS072**

**Chronic subdural hematoma in aging population – How the age influence the outcome after surgical treatment**

Uładzislau Ulasavets *, Evelina Grzywna

Universytet Jagielloński – Collegium Medicum, Poland

*E-mail address: vladvlasgeo@icloud.com* (U. Ulasavets).

**Aim:** The aim of our work is to examine how the age influence the outcome after surgical treatment of chronic subdural hematoma.

**Introduction:** Chronic subdural hematoma (CSDH) is a common condition, characterized by the collection of hemolysed blood between dura and arachnoid mater of the brain surrounded by two pathological hematoma membranes – internal and external. The number of CSIDH incidence increases with age and it is why more attention should be directed for surgical treatment in elder patients group.

**Methods:** Data on management and outcomes for patients with CSIDH were collected retrospectively from years 2014–2017 and investigated using statistic methods. The study group was divided into two subgroups according to the age: <75 years and ≥75 years old. Age, gender, comorbidities, neurological status on admission and at discharge, pre-/postoperative epilepsy, surgical technique were investigated.

**Results:** We analyzed 257 patients with a diagnosis CSIDH. Analyzed subgroups have not differ significantly except the gender and concomitant diseases according to the Chi2 and exact Fisher tests. We found craniotomy in patients ≥75 years old increases the risk of postoperative epilepsy comparing to the bur-hole (logistic regression analysis: 9.8 [95% CI: 1.9–49.8], p = .006), same as the internal hematoma membrane removal during surgery (logistic regression analysis: 10.3 [95% CI: 2.0–52.15], p = .005). These dependencies do not occur in the younger age group. Type of treatment have not influenced the mRS in patients younger than 75 years old. In elder patients reoperation and removal of the internal membrane of the hematoma worsened outcome measured in mRS (logistic regression analysis: 5.5 [95% CI: 1.4–20.90], p = .013 and 3.1 [95% CI: 1.4–7.2], p = .007).

**Conclusion:** Craniotomy and internal membrane removal increase the risk of epilepsy in elder CSIDH patients. Reoperation

(P = 1.08, n = 1228) and study (P < 1.08, n = 100). The differences in socio-demographic factors between control and study group were not statistically significant. Data were analysed using chi-squared test, independent sample 2-tailed T-test and logistic regression. p value < 0.05 was statistically significant.

**Results:** In study group was observed statistically significant increased risk of delivery provided by cesarean section (OR = 1.8; p = 0.015), preterm delivery (OR = 2.91; p = 0.0001), birth weight > 2500 g (OR = 5.87; p < 0.00001) and APGAR score < 7 in 1st (OR = 6.56; p < 0.0001), 3rd (OR = 7.04; p < 0.0001) and 5th (OR = 5.4; p = 0.017) minute after delivery, compared to control group. Moreover, low CPR was associated with lower incidence of foetus birth weight within normal limits (OR = 0.37; p < 0.0001) and on-term delivery (OR = 0.61; p = 0.0001).

**Conclusion:** Detection of low value of CPR in every case should be alarming signal for obstetrician. Normal CPR appears to suggest better foetal tolerance to the stress of labour. CPR may be used to stratify the risk of pregnancy before labour.

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**PS134**

**The role of cerebroplacental ratio in prediction of neonatal outcomes and route of delivery**

P. Janas *, A. Staroń, G. Wilczyńska, M. Brzozowska

Jagiellonian University Medical College, Cracow, Poland

*E-mail address: przemyslaw.janas@gmail.com* (P. Janas).

**Aim:** The aim of our study was to check the appropriability of cerebroplacental ratio (CPR) measured within 48h before delivery in prediction of route of delivery and adverse neonatal outcomes.

**Introduction:** The cerebroplacental ratio is an important obstetric ultrasound tool used for assessment of foetal oxygenation. It is also a valuable predictor of adverse pregnancy outcomes. CPR is calculated by dividing the Doppler pulsatile indices of the middle cerebral artery (MCA) and the umbilical artery (UA).

**Methods:** The retrospective study included 1328 pregnant women who gave birth in Department of Obstetrics and Gynecology Jagiellonian University Medical College, Cracow, Poland. Main inclusion criteria were: singleton pregnancy and the interval between ultrasound examination and delivery within 48 h. Exclusion criteria consisted: active labour, multiple pregnancy, preeclampsia, foetal growth restriction and evidence of intrauterine infection. CPR value lower than 1.08 was classified as pathological. Participants were divided into 2 groups: control (CPR ≥ 1.08, n = 1228) and study (CPR < 1.08, n = 100). The differences in socio-demographic factors between control and study group were not statistically significant. Data were analysed using chi-squared test, independent sample 2-tailed T-test and logistic regression. p value < 0.05 was statistically significant.

**Results:** In study group was observed statistically significant increased risk of delivery provided by cesarean section (OR = 1.8; p = 0.015), preterm delivery (OR = 2.91; p = 0.0001), birth weight > 2500 g (OR = 5.87; p < 0.00001) and APGAR score < 7 in 1st (OR = 6.56; p < 0.0001), 3rd (OR = 7.04; p < 0.0001) and 5th (OR = 5.4; p = 0.017) minute after delivery, compared to control group. Moreover, low CPR was associated with lower incidence of foetus birth weight within normal limits (OR = 0.37; p < 0.0001) and on-term delivery (OR = 0.61; p = 0.0001).

**Conclusion:** Detection of low value of CPR in every case should be alarming signal for obstetrician. Normal CPR appears to suggest better foetal tolerance to the stress of labour. CPR may be used to stratify the risk of pregnancy before labour.

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