

Comparison of basic regional cerebral oxygen saturation values in patients of different ages: a pilot study

Journal of International Medical Research

48(8) 1–9

© The Author(s) 2020


Article reuse guidelines:

sagepub.com/journals-permissions

DOI: 10.1177/0300060520936868

journals.sagepub.com/home/imr



Chaohui Lian¹, Peng Li^{1,2}, Ning Wang¹, Yi Lu¹
and Wangning Shangguan¹ 

Abstract

Objective: To explore the basic values of regional cerebral oxygen saturation (rSO₂) among different age groups.

Methods: One hundred twenty patients who were scheduled for elective surgery aged 0 to 80 years (American Society of Anesthesiologists [ASA] physical status I or II) or neonates just after birth via cesarean section were enrolled and divided into the following six groups: infant (0 month and ≤12 months), toddler (>1 and ≤3 years old), preschool (>3 and ≤6 years old), school age (>6 and ≤18 years old), adult (>18 and ≤65 years old), and elderly (>65 and ≤80 years old) groups. There were 20 patients in each group.

Results: The basic values of rSO₂ in infant, toddler, preschool, school age, adults, and elderly groups were 70.41% ± 4.66%, 72.43% ± 3.81%, 70.77% ± 3.27%, 70.62% ± 2.20%, 69.76% ± 6.02%, and 62.69% ± 3.14%, respectively. The basic value in the elderly group was lower compared with other five groups. There was no significant difference among infant, toddler, preschool age, school age, and adult groups.

Conclusions: The basic value of rSO₂ in elderly patients is lower. Age is an important factor that affects the underlying value of rSO₂.

Keywords

Spectroscopy, near-infrared, oximetry, age groups, pulse oximetry, cerebral hypoxia–ischemia

Date received: 18 February 2020; accepted: 3 June 2020

¹Department of Anesthesiology and Perioperative Medicine, The Second Affiliated Hospital and Yuying Children's Hospital of Wenzhou Medical University, Wenzhou, China

²Department of Anesthesiology, The First Affiliated Hospital of Wenzhou Medical University, Wenzhou, China

Corresponding author:

Wangning ShangGuan, Department of Anesthesiology and Perioperative Medicine, The Second Affiliated Hospital and Yuying Children's Hospital of Wenzhou Medical University, 109 West Xueyuan Road, Wenzhou 325027, China.

Email: sgwning@163.com



Introduction

Regional cerebral oxygen saturation (rSO₂) is a non-invasive monitoring technique that uses near-infrared spectroscopy (NIRS) to measure oxygen saturation of superficial brain cortex regions, which are among the most vulnerable to ischemic-hypoxic injury. NIRS is sensitive to hypoxia in tissues and organs. Even if the oxygenation of tissues and organs is slightly changed, such as pulse oximetry (SpO₂) that is still within the normal range, the rSO₂ value could be significantly changed.¹ The more obvious the decrease in rSO₂, the more serious hypoxia there is in the brain tissue, which could be used to judge the prognosis.² A study showed that the cerebral hypoxia-ischemia threshold of rSO₂ for functional impairment was 33% to 44%.³ Yao et al.⁴ reported that patients with a nadir rSO₂ <35% or areas of rSO₂ <40% for more than 10 minutes had significantly higher incidences of postoperative cognitive impairment. Most studies⁵⁻⁷ set the threshold to 50%, and they showed that rSO₂ <50% is related to adverse outcomes. rSO₂ monitoring could provide a bedside method that continuously and conveniently monitors “real-time” oxygen consumption, and that also monitors important tissues and organs, especially the brain, kidneys, and intestines.⁸ It is, therefore, increasingly used as a complement to traditional monitoring in clinical practice especially in major surgeries that seriously affect the cerebral circulation. However, there is currently no uniform standard for its basic value, and there is no consensus on the influencing factors of the value of rSO₂.

It has been reported that the rSO₂ value was negatively correlated with the patient's age, positively correlated with hemoglobin concentration, and also related to type of equipment and sensor location, but not related to patient weight, height, head size, or gender.⁹ Another study found that the

basic value of rSO₂ was not related to patient age, gender, and body weight.¹⁰ Therefore, the purpose of this study was to explore the basic values of rSO₂ in different age groups of patients who were scheduled for elective surgery and neonates who were just born after cesarean section, and to provide a reference for the clinical application of rSO₂. We hypothesized that different age groups have different basic values of rSO₂.

Methods

This study was registered at www.chictr.org.cn, registration number ChiCTR1800017406. The study protocol was reviewed and approved by the Ethics Committee of the Second Affiliated Hospital of Wenzhou Medical University (No. 2018-10). Informed consent was provided by all enrolled patients or their relatives (legally authorized guardian). The inclusion criteria included the following: obtained informed consent from the patient or guardian; patients aged 0 to 80 years; American Society of Anesthesiologists (ASA) physical status I to II (newborn Apgar score ≥7 points); SpO₂ ≥97%; and normal body temperature. The exclusion criteria including the following: mental developmental disorders; hypertension, heart disease, and diabetes; history of head injury, epilepsy, and psychosis; those with cerebrovascular disease or cranial CT or MRI showing cerebrovascular disease; history of lung infection or pulmonary insufficiency; the patient could not cooperate; or the cerebral oxygen monitoring site could not be revealed.

All patients who were scheduled for elective surgery or neonates just after birth via cesarean section were divided into six groups based on their age, as follows: infant group (0 month and ≤12 months); toddler group (>1 and ≤3 years old); preschool group (>3 and ≤6 years old); school age group (>6 and ≤18 years old); adult

group (>18 and ≤65 years old); and elderly group (>65 and ≤80 years old). All the subjects were assigned to a well-ventilated and smoke-free room, and kept in a resting state for more than 15 minutes. A Somanetics INVOS 5100C (Medtronic Inc., Minneapolis, MN, USA) cerebral oxygen monitor and appropriate sensor based on the patient's weight was applied (an adult sensor for body weight ≥40 kg, a pediatric sensor for body weight <40 kg, or a baby special sensor for infants and neonates). The sensors were placed at the right and left frontal regions, 1 cm above the patients' eyebrows, and the edges of the sensors on both sides were 1 cm from the median line. To avoid problems that are possibly caused by ambient lighting, sensors were covered with an opaque plastic covering. The SpO₂ probe was fixed on the right index finger. The average value of the rSO₂ from the left and right sides was taken as the basic value. The values of rSO₂, SpO₂, and pulse rate (PR) at 1 minute (T1), 3 minutes (T3), and 5 minutes (T5) after starting the monitor was recorded. The hemoglobin concentrations of all subjects were obtained before the tests.

The SPSS 24.0 (IBM Corp., Armonk, NY, USA) statistical software package was used for data analysis. After the normality test and the homogeneity test of variance, the measurement data with normal distribution were expressed as the mean ± standard deviation (SD), and the q test was used for comparison between groups (S-N-Ka). The intra-group comparison was analyzed by repeated measures analysis of variance. $P < 0.05$ was considered to be statistically significant.

Results

The study was conducted from August to December 2018. There were 131 patients who were assessed for eligibility, and eight patients were excluded because they did not

meet the study entry criteria, they refused to participate, or their hemoglobin concentration levels before the study could not be obtained, and three infants were non-cooperative. Thus, 120 patients were enrolled with 20 patients in each group, as described in Figure 1. All 120 patients completed the study. The basic hemoglobin concentration in infant, toddler, preschool, school age, adult, and elderly patient groups were 126.5 ± 10.3 (116–159) g/L, 123.5 ± 6.4 (110–138) g/L, 124.3 ± 8.5 (111–140) g/L, 125.0 ± 9.7 (113–151) g/L, 123.4 ± 10.2 (111–152) g/L, and 121.8 ± 10.0 (109–142) g/L, respectively. There was no significant difference in the hemoglobin concentration among the six groups.

For each group, the values of SpO₂ and PR were not significantly different at T1, T3, and T5 time points (Table 1). The values of rSO₂ had wide ranges among the six groups, from 54.65% to 80.90%. The basic values of rSO₂ in infant, toddler, preschool, school age, adult, and elderly patient groups were $70.41\% \pm 4.66\%$, $72.43\% \pm 3.81\%$, $70.77\% \pm 3.27\%$, $70.62\% \pm 2.20\%$, $69.76\% \pm 6.02\%$, and $62.69\% \pm 3.14\%$, respectively (Table 2). The basic value of rSO₂ in the elderly group was lower compared with the other five groups ($P < 0.05$). There was no significant difference among infants, toddlers, preschool children, school age children, and adults. Additionally, the maximum difference of basic rSO₂ value was 14.5% (62.5%–77.0%) in the infant group, 12.4% (66.6%–79.0%) in the toddler group, 13.6% (63.8%–77.4%) in the preschool group, 8.4% (65.5%–73.9%) in the school age group, 22.4% (58.5%–80.9%) in the adult group, and 16.1% (54.6%–70.7%) in the elderly group.

Discussion

The main outcome of this current study was that the values of rSO₂ had a wide range among the six groups, from 54.7% to

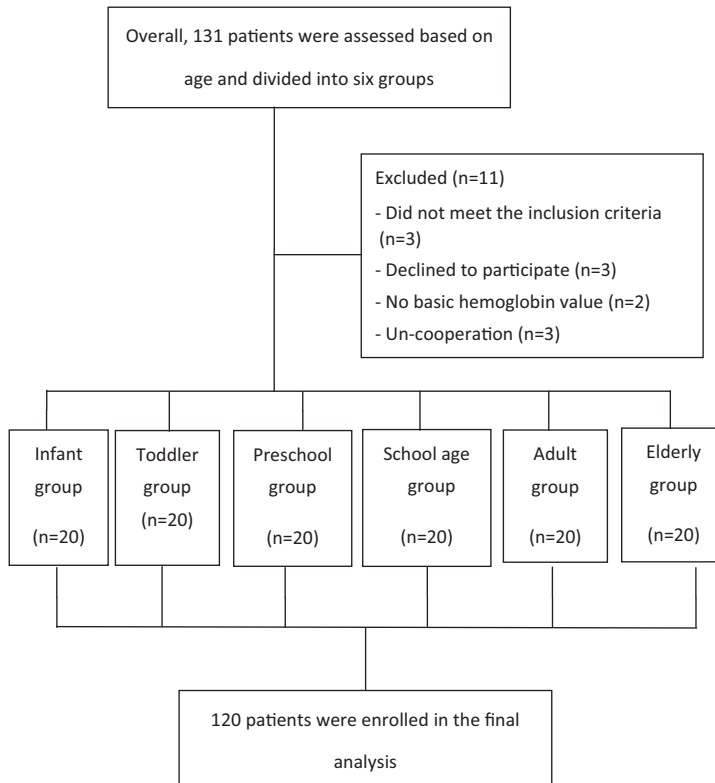


Figure 1. The flowchart of the study.

Table 1. SpO₂ and PR among six age groups at three time points.

| Groups | SpO ₂ (%) | | | PR (bpm) | | |
|------------|----------------------|----------|----------|----------|----------|----------|
| | T1 | T3 | T5 | T1 | T3 | T5 |
| Infant | 99 ± 0.5 | 99 ± 0.8 | 99 ± 0.6 | 132 ± 11 | 129 ± 15 | 134 ± 11 |
| Toddler | 99 ± 0.6 | 99 ± 1.0 | 99 ± 0.5 | 110 ± 5 | 109 ± 7 | 108 ± 9 |
| Preschool | 99 ± 0.4 | 99 ± 0.8 | 99 ± 0.7 | 90 ± 5 | 89 ± 4 | 87 ± 9 |
| School age | 99 ± 0.7 | 99 ± 0.4 | 99 ± 0.6 | 80 ± 3 | 82 ± 4 | 81 ± 2 |
| Adult | 99 ± 0.8 | 99 ± 1.0 | 99 ± 0.4 | 67 ± 9 | 66 ± 9 | 69 ± 5 |
| Elderly | 99 ± 0.8 | 99 ± 0.4 | 99 ± 0.7 | 73 ± 4 | 70 ± 8 | 71 ± 9 |

Data are presented as the mean ± SD; n = 20.

SpO₂, pulse oximetry; PR, pulse rate; SD, standard deviation; bpm, beats per minute.

80.9%. Additionally, the basic value of rSO₂ in the elderly group was lower, while there was no significant difference among other five groups.

It has been reported that when the sensor position moved 1 cm lateral from the mid-line, the rSO₂ value was significantly reduced.⁹ However, when the sensor was

Table 2. The values of rSO₂ among six age groups at three time points.

| Groups | rSO ₂ (%) | | | |
|------------|----------------------|----------------|----------------|-------------------------|
| | T1 (mean ± SD) | T3 (mean ± SD) | T5 (mean ± SD) | Mean ± SD (Min–Max) |
| Infant | 70.4 ± 4.6 | 70.4 ± 4.7 | 70.4 ± 4.7 | 70.4 ± 4.7 (62.5–77.0) |
| Toddler | 72.4 ± 3.8 | 72.4 ± 3.8 | 72.4 ± 3.8 | 72.4 ± 3.8 (66.6–79.0) |
| Preschool | 70.8 ± 3.3 | 70.8 ± 3.3 | 70.8 ± 3.2 | 70.8 ± 3.3 (63.8–77.5) |
| School age | 70.6 ± 2.2 | 70.6 ± 2.2 | 70.7 ± 2.2 | 70.6 ± 2.2 (65.6–74.0) |
| Adult | 69.7 ± 5.9 | 69.9 ± 6.1 | 69.7 ± 6.1 | 69.8 ± 6.0 (58.5–80.9) |
| Elderly | 62.7 ± 3.2* | 62.7 ± 3.1* | 62.7 ± 3.1* | 62.7 ± 3.1 (54.7–70.8)* |

*P < 0.05 vs. infant, toddler, preschool, school age, adult groups.

Data are presented as a percent (%; n = 20).

rSO₂, regional cerebral oxygen saturation; SD, standard deviation; Min, minimum; Max, maximum.

placed at the center of the forehead, the rSO₂ value was significantly higher compared with other parts. This phenomenon is consistent with our study; when the position of the sensor was not fixed, the value of rSO₂ changed rapidly. In this study, the position of the sensors was uniformly placed 1 cm above the eyebrows on both sides, and the edges of the sensors on both sides were 1 cm from the center line of the forehead. Thus, the test error caused by different devices and different sensor positions was effectively avoided.

Torella et al.¹¹ reported that the rSO₂ value was closely related to the hemoglobin concentration, and the basic value of rSO₂ increased when the hemoglobin concentration increased. Lassnigg et al.¹² demonstrated a strong correlation between cerebral oxygenation and hemoglobin concentration during hemodilution by cardiopulmonary bypass. Liem et al.¹³ also reported that blood transfusion increased cerebral oxygenation in premature infants with anemia, while the opposite changes were observed during hemodilution. In this study, the hemoglobin concentration among the six groups showed no statistical difference, and thus, the interference of hemoglobin concentration on rSO₂ could be eliminated. However, there are subtypes of hemoglobin. It is speculated that the

content of hemoglobin subtypes may differ in different age groups, which affects the basic values of cerebral oxygen saturation in all age groups. Therefore, further research is required to confirm the effect of hemoglobin type on rSO₂ values.

Currently, there is no uniform standard for the basic values of rSO₂ in different age groups, and the influencing factors on the basic values of rSO₂ remain unclear, although related rSO₂ studies have been reported.^{9,10,14–17} rSO₂ monitoring reflects the mixed oxygen saturation of the local brain tissue, which consists of about 70% venous blood, 20% arterial blood, and 10% capillary blood. The reference value of arterial oxygen saturation in normal people ranged from 95% to 98%, and the venous oxygen saturation ranged from 60% to 85%. Theoretically, the normal base value of rSO₂ should be about 70%,^{18–20} which is consistent with our findings. Ružman et al.²¹ showed that the mean basal values of rSO₂ on the left and right sides were 71.42 ± 7.92% and 72.98 ± 7.94%, respectively, which is much closer to our results, but they did not find any correlation with age. Alderliesten et al.²² reported that the average rSO₂ in neonates was approximately 65% at admission, and it increased with gestational age (1% per week). The average basal values of rSO₂ were slightly higher

than our results. Increasing of age (months) and different sensors might be the reasons for the diversity of their results compared with our study. There are many commercially available NIRS devices on the market, and numerous studies^{23–27} have compared the differences between them. The related results showed that there were significant differences between the devices in how they respond to changes in oxygen saturation, so caution should be used when applying evidence that was generated with one manufacturer's device to all devices. Even if the same device is used, the values measured by different probes may be different. The value measured with a neonatal probe was reported to be 10% higher compared with the adult probe. However, the monitoring probe was selected according to the manufacturer's recommendation and common clinical practice to observe the differences in the normal basic values of rSO₂ between different ages rather than to assess the value differences for using an adult probe in children or a neonatal probe in adults.

The current study showed that the basic value of rSO₂ in the elderly group was lower compared with the other groups, while there was no difference among other five groups. Kishi et al.⁹ measured the basic values of rSO₂ in 111 patients between 7 to 89 years old under general anesthesia. The results showed that the basic values were related to the patient's age, hemoglobin concentration, and the position of the sensors, but they were not significantly associated with the patient's height, weight, gender, and head circumference. However, the enrolled patients were mainly limited to adults and the elderly. Our study also confirmed that the basic values of rSO₂ were related to the age. A previous small-sample study by Hock et al.²⁸ also reported that the basic values of rSO₂ were significantly correlated with age in awake subjects of different ages.

The mechanisms of this correlation remain unclear. The possible explanations are suggested as follows: first, there are age-associated decreases in cerebral blood flow, cerebral blood volume, cerebral metabolic rate of oxygen, and cerebral hemoglobin oxygenation response during brain activation, and brain atrophy is gradually increased in elderly people. This leads to a decrease in brain volume and the number of brain cells and accompanies cerebrovascular degeneration,²⁹ which will have an important impact on the basic values, resulting in a low rSO₂ value in elderly people. Second, with increasing age, the central nervous system myelin sheath is gradually prolonged (especially in the elderly), resulting in a significant extension of the length of the intracranial near-infrared light path.³⁰ The duration from when the rSO₂ detector probe emits near-infrared light to when it receives the reflected near-infrared light is significantly prolonged and the energy attenuation is significantly increased, resulting in a decrease in the measured value of rSO₂. This may be a major factor leading to the low basic values of rSO₂ in elderly people. Finally, the configuration of hemoglobin changes with age, giving rise to a decrease in cerebral hemoglobin oxygenation.³¹ The total amount of hemoglobin is equal to the sum of oxyhemoglobin and deoxyhemoglobin. With aging, the amount of oxyhemoglobin decreases and the amount of deoxyhemoglobin gradually increases, leading to a decline in the oxygen carrying capacity of cerebral hemoglobin, thereby decreasing the basic value of rSO₂ in elderly people. However, the validity of these explanations requires further experimental evidence, and more clinical studies are warranted.

The rSO₂ value is susceptible to fluctuation that is caused by external factors. One study showed that the mental state, posture, external sounds, light, and environmental ventilation conditions may have an effect

on the value of rSO_2 .³² Another study found that the rSO_2 value was significantly increased when the subject's brain was doing a calculation, and the value was significantly reduced when the carotid artery was lightly pressed.³³ Our study also observed that the values of rSO_2 changed rapidly and the sensitivity was high. In this study, the maximum difference of basic rSO_2 value was 14.5% (62.5%–77.0%) in the infant group, 12.4% (66.6%–79.0%) in the toddler group, 13.6% (63.8%–77.4%) in the preschool group, 8.4% (65.5%–73.9%) in the school age group, 22.4% (58.5%–80.9%) in the adult group, and 16.1% (54.6%–70.7%) in the elderly group. Therefore, the external conditions should be uniformly defined during the measurement of the basic rSO_2 value.

There were some limitations in our study. First, although 120 ASA grade I to II patients of all ages were included in this study, this was a pilot investigation study and the sample size in our study was small. The selected subjects were also hospitalized in the same hospital, and there might have some selection bias. Thus, a large multi-center study is required to further confirm this result. Second, the Somanetics INVOS 5100C has good repeatability, but with the extension of the measurement duration, the fluctuation range of rSO_2 may increase.³⁴ Thus, a measurement period of 5 minutes may not be long enough. Finally, the value measured using the neonatal probe is 10% higher compared with the adult probe. However, the values obtained using the same probe in different age groups were not measured, which may partially limit the clinical application of the experimental results.

In summary, this study provides the normal basic values of rSO_2 under Somanetics INVOS 5100C monitoring among all age groups (from newborn to elderly). This could show whether the

patient's rSO_2 value is in a reasonable range. Additionally, the baseline value of rSO_2 in ASA grade I to II elderly patients (>65 and ≤80 years old) tended to be lower compared with other age groups, and there was no significant difference among the other five age groups. Thus, age is an important factor that affects the underlying rSO_2 value.

Acknowledgements

The authors thank the Zhejiang Provincial Public Welfare Technology Application Research Foundation of China (2018ZD033).

Declaration of conflicting interest

The authors declare that there is no conflict of interest.

Funding

The research was funded by Zhejiang Provincial Public Welfare Technology Application Research Foundation of China (2018ZD033).

ORCID iD

Wangning Shangguan  <https://orcid.org/0000-0002-8088-1180>

References

1. Tobias JD. Cerebral oximetry monitoring with near infrared spectroscopy detects alterations in oxygenation before pulse oximetry. *J Intensive Care Med* 2008; 23: 384–388.
2. Jöbsis FF. Noninvasive, infrared monitoring of cerebral and myocardial oxygen sufficiency and circulatory parameters. *Science* 1977; 198: 1264–1267.
3. Kurth CD, Levy WJ and McCann J. Near-infrared spectroscopy cerebral oxygen saturation thresholds for hypoxia-ischemia in piglets. *J Cereb Blood Flow Metab* 2002; 22: 335–341.
4. Yao FSF, Tseng CCA, Ho CYA, et al. Cerebral oxygen desaturation is associated with early postoperative neuropsychological dysfunction in patients undergoing cardiac

- surgery. *J Cardiothorac Vasc Anesth* 2004; 18: 552–558.
5. Slater JP, Guarino T, Stack J, et al. Cerebral oxygen desaturation predicts cognitive decline and longer hospital stay after cardiac surgery. *Ann Thorac Surg* 2009; 87: 36–44; discussion 44–35.
 6. Jo YY, Shim JK, Soh S, et al. Association between cerebral oxygen saturation with outcome in cardiac surgery: brain as an index organ. *J Clin Med* 2020; 9: 840.
 7. Alderliesten T, Lemmers PM, Van Haastert IC, et al. Hypotension in preterm neonates: low blood pressure alone does not affect neurodevelopmental outcome. *J Pediatr* 2014; 164: 986–991.
 8. Biedrzycka A and Lango R. Tissue oximetry in anaesthesia and intensive care. *Anaesthesiol Intensive Ther* 2016; 48: 41–48.
 9. Kishi K, Kawaguchi M, Yoshitani K, et al. Influence of patient variables and sensor location on regional cerebral oxygen saturation measured by INVOS 4100 near-infrared spectrophotometers. *J Neurosurg Anesthesiol* 2003; 15: 302–306.
 10. Schoen J, Heringlake M, Tiemeyer C, et al. The association between gender, cognitive function, markers of inflammation and preoperative cerebral oxygen saturation in a cohort of preoperative cardiac surgery patients. *Appl Cardiopulm Pathophysiol* 2011; 15: 62–70.
 11. Torella F, Cowley R, Thorniley MS, et al. Monitoring blood loss with near infrared spectroscopy. *Comp Biochem Physiol A Mol Integr Physiol* 2002; 132: 199–203.
 12. Lassnigg A, Hiesmayr M, Keznickl P, et al. Cerebral oxygenation during cardiopulmonary bypass measured by near-infrared spectroscopy: effects of hemodilution, temperature, and flow. *J Cardiothorac Vasc Anesth* 1999; 13: 544–548.
 13. Liem KD, Hopman JC, Oeseburg B, et al. The effect of blood transfusion and haemodilution on cerebral oxygenation and haemodynamics in newborn infants investigated by near infrared spectrophotometry. *Eur J Pediatr* 1997; 156: 305–310.
 14. Mohandas BS, Jagadeesh AM and Vikram SB. Impact of monitoring cerebral oxygen saturation on the outcome of patients undergoing open heart surgery. *Ann Card Anaesth* 2013; 16: 102–106.
 15. Conforti A, Giliberti P, Landolfo F, et al. Effects of ventilation modalities on near-infrared spectroscopy in surgically corrected CDH infants. *J Pediatr Surg* 2016; 51: 349–353.
 16. Heringlake M, Garbers C, Kabler JH, et al. Preoperative cerebral oxygen saturation and clinical outcomes in cardiac surgery. *Anesthesiology* 2011; 114: 58–69.
 17. Pellicer A, Greisen G, Benders M, et al. The SafeBoosC phase II randomised clinical trial: a treatment guideline for targeted near-infrared-derived cerebral tissue oxygenation versus standard treatment in extremely preterm infants. *Neonatology* 2013; 104: 171–178.
 18. Cetin M, Birbicer H, Hallioglu O, et al. Comparative study between the effects of dexmedetomidine and propofol on cerebral oxygenation during sedation at pediatric cardiac catheterization. *Ann Card Anaesth* 2016; 19: 20–24.
 19. Yoshitani K, Kawaguchi M, Tatsumi K, et al. A comparison of the INVOS 4100 and the NIRO 300 near-infrared spectrophotometers. *Anesth Analg* 2002; 94: 586–590.
 20. Beese U, Langer H, Lang W, et al. Comparison of near-infrared spectroscopy and somatosensory evoked potentials for the detection of cerebral ischemia during carotid endarterectomy. *Stroke* 1998; 29: 2032–2037.
 21. Ruzman T, Mraović B, Šimurina T, et al. Transcranial cerebral oxymetric monitoring reduces brain hypoxia in obese and elderly patients undergoing general anesthesia for laparoscopic cholecystectomy. *Surg Laparosc Endosc Percutan Tech* 2017; 27: 248–252.
 22. Alderliesten T, Dix L, Baerts W, et al. Reference values of regional cerebral oxygen saturation during the first 3 days of life in preterm neonates. *Pediatr Res* 2016; 79: 55–64.
 23. Bickler PE, Feiner JR and Rollins MD. Factors affecting the performance of 5 cerebral oximeters during hypoxia in healthy volunteers. *Anesth Analg* 2013; 117: 813–823.
 24. Pisano A, Galdieri N, Iovino TP, et al. Direct comparison between cerebral

- oximetry by INVOS(TM) and EQUANOX (TM) during cardiac surgery: a pilot study. *Heart Lung Vessel* 2014; 6: 197–203.
25. Cournoyer A, Denault A, Cossette S, et al. Reproducibility, interchangeability of measures, time to measure stabilization, and reference values of two tissue oximeters in healthy volunteers. *J Biomed Opt* 2016; 21: 97003.
 26. Douds MT, Straub EJ, Kent AC, et al. A systematic review of cerebral oxygenation-monitoring devices in cardiac surgery. *Perfusion* 2014; 29: 545–552.
 27. Tomlin KL, Neitenbach AM and Borg U. Detection of critical cerebral desaturation thresholds by three regional oximeters during hypoxia: a pilot study in healthy volunteers. *BMC Anesthesiol* 2017; 17: 6.
 28. Hock C, Müller-Spahn F, Schuh-Hofer S, et al. Age dependency of changes in cerebral hemoglobin oxygenation during brain activation: a near-infrared spectroscopy study. *J Cereb Blood Flow Metab* 1995; 15: 1103–1108.
 29. Leenders KL, Perani D, Lammertsma AA, et al. Cerebral blood flow, blood volume and oxygen utilization. Normal values and effect of age. *Brain* 1990; 113: 27–47.
 30. Duncan A, Meek JH, Clemence M, et al. Measurement of cranial optical path length as a function of age using phase resolved near infrared spectroscopy. *Pediatr Res* 1996; 39: 889–894.
 31. Mehagnoul-Schipper DJ, Van Der Kallen BF, Colier WN, et al. Simultaneous measurements of cerebral oxygenation changes during brain activation by near-infrared spectroscopy and functional magnetic resonance imaging in healthy young and elderly subjects. *Hum Brain Mapp* 2002; 16: 14–23.
 32. Abdul-Khaliq H, Troitzsch D, Berger F, et al. Regional transcranial oximetry with near infrared spectroscopy (NIRS) in comparison with measuring oxygen saturation in the jugular bulb in infants and children for monitoring cerebral oxygenation. *Biomed Tech (Berl)* 2000; 45: 328–332.
 33. Gottlieb EA and Mossad EB. Limitations of cerebral oxygenation monitoring by near-infrared spectroscopy in children with cyanotic congenital heart disease and profound polycythemia. *J Cardiothorac Vasc Anesth* 2014; 28: 347–349.
 34. Pocivalnik M, Pichler G, Zotter H, et al. Regional tissue oxygen saturation: comparability and reproducibility of different devices. *J Biomed Opt* 2011; 16: 057004.