Cerebral tissue oxygenation index in very premature infants

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Aim: To describe normal values of the cerebral tissue oxygenation index (TOI) in premature infants.

Methods: TOI was measured by spatially resolved spectroscopy in preterm infants on the first 3 days of life. Infants with an abnormal cranial ultrasound were excluded. Other simultaneously measured variables were PaO₂, PaCO₂, pH, mean arterial blood pressure, heart rate, haemoglobin, glycaemia, and peripheral oxygen saturation.

Results: Fifteen patients with a median postmenstral age of 28 weeks were measured. There was a significant increase in median TOI over the first 3 days of life: 57% on day 1, 66.1% on day 2, and 76.1% on day 3. Multiple regression analysis showed no correlation between TOI and postmenstral age, peripheral oxygen saturation, mean arterial blood pressure, PaO₂, PaCO₂, and haemoglobin concentration.

Conclusion: Cerebral TOI increases significantly in the first 3 days of life in premature babies. This increase probably reflects the increase in cerebral blood flow at this time.

PATIENTS AND METHODS
Fifteen patients with a postmenstral age of less than 31 weeks were included. The median postmenstral age was 28 weeks (range 25–30). A brain ultrasound was performed on all patients before measurements were started. Exclusion criteria were an abnormal brain ultrasound before the TOI measurement and severe pulmonary hypertension as evidenced by echocardiography and/or congenital malformations.

An NIRO 300 (Hamamatsu Phototonics K.K., Tokyo, Japan) was used for spatially resolved spectroscopy. The optode was placed at the right frontoparietal side with the sensors at 4 cm distance. All patients were measured within 6 hours of birth for at least 30 minutes. The second and third measurements were performed 24 and 48 hours later. The specific variable measured was TOI.

Spatially resolved spectroscopy is a new method, using near infrared spectroscopy (NIRS), for measuring cerebral haemoglobin oxygen saturation. The tissue oxygenation index (TOI) is measured with a light detector with three sensors placed at different distances from the near infrared light source. If light with a given intensity is sent into tissue, it is attenuated because of “scatter” loss and “absorption” loss. If the distance between the light source and the sensor is large enough, the isotropy of scatter distribution becomes so homogeneous that the scatter loss is the same at the three sensors. Therefore any differences in intensity measured at the three different sensors can be interpreted as differences in absorption loss. Thus the local absorption change can be seen as a function of the distance between the light source and the three sensors. With this information, an oxygenation index can be calculated using a previously reported algorithm. Several research groups have measured TOI in healthy adults with different NIRS instruments, and no differences in TOI values between the right and left forehead have been found. To our knowledge, no normal cerebral TOI values in the first days of life have been reported for premature babies (less than 30 weeks gestation). In this study, we measured TOI on days 1, 2, and 3 to obtain “normal” values and to investigate changes with postnatal age.

RESULTS
Fifteen patients were studied during the first 3 days of life. The mean (SD) birth weight was 1053 (395) g, and the mean (SD)
head circumference was 24.8 (2.5) cm. All patients had a normal ultrasound of the brain before measurement on day 1. Further follow up of the ultrasound on days 3, 5, and 7 showed an intraventricular haemorrhage grade 1 or II in seven of the 15 patients. However, no posthaemorrhagic hydrocephalus or intraventricular haemorrhage grade III or IV was noted. One patient developed cystic periventricular leucomalacia three weeks later. Seven patients developed retinopathy of prematurity grade III, and three of these patients needed cryotherapy. Psychomotor follow up was normal in 10 patients at the age of 6 months.

Figure 1 shows the TOI values. The median TOI was 57% (95% CI 54 to 65.7%) on day 1, 66.1% (61.9 to 82.2%) on day 2, and 76.1% (67.1 to 80.1%) on day 3. The increase from day 1 to day 2 and from day 1 to day 3 was significant (p<0.05).

Table 1 shows the different variables. To test the effect of oxygenation, analysis of variance was performed to find an eventual difference in peripheral oxygen saturation for the three different subgroups (Kolmogorov-Smirnov test for normal distribution, p>0.2). Analysis of variance showed a significant difference (p<0.001) between TOI and postnatal age. A Student-Newman-Keuls test for all pairwise comparisons showed that there was a significant increase in TOI from day 1 to day 2 and from day 1 to day 3, but not from day 2 to day 3.

Table 1 shows the different variables. To test the effect of oxygenation, analysis of variance was performed to find an eventual difference in peripheral oxygen saturation for the different days. No significant difference was found (p = 0.44). A significant (p = 0.003) increase in blood pressure from day 1 to day 3 was found. The increase in blood pressure from day 1 to day 2 was not significant. There was also a significant difference in PaCO2 on the different days. Mean PaCO2 on day 1 was 31 mm Hg, on day 2 45 mm Hg, and on day 3 36.5 mm Hg. This is a significant difference (p < 0.001) between the three days, but there was an increase from day 1 to 2 and a decrease from day 2 to 3. There was no significant difference in Pao2, bicarbonate, and pH on the different days. Haemoglobin concentration was 14.1 g/dl on day 1, 13 g/dl on day 2, and 14.1 g/dl on day 3. The percentage of fetal haemoglobin was only measured in eight patients. It was 86.7% on day 1, 88.5% on day 2, and 69.7% on day 3. This significant decrease on day 3 is explained partly by the fact that these patients had received a blood transfusion.

Multiple regression analysis showed an increase in TOI only from day 1 to day 3 (p < 0.05). No correlation between TOI and blood pressure, PaCO2, peripheral oxygen saturation, haemoglobin concentration, fetal haemoglobin, glycaemia or postmenstrual age was found with multiple regression analysis. No correlation was found between TOI or the standard deviation of TOI and head circumference.

No relation was found between TOI and intraventricular bleeding, periventricular leucomalacia, retinopathy, or abnormal psychomotor follow up.

**DISCUSSION**

NIRS is a non-invasive method for measuring oxygenated and deoxygenated haemoglobin and derived values of brain oxygenation, cerebral blood flow, and cerebral blood volume. Until now, it has been used only in research because it is very sensitive to movement artefacts. Furthermore, it does not provide absolute values, only values relative to the starting point in a continuous way. TOI, in contrast, is an absolute value and can be measured on different occasions in the same patient. Although in this study TOI was determined in very premature babies and their head circumference was small, very stable TOI values with a mean standard deviation of 2.2% were obtained from measurements taken over at least 30 minutes.

Whether TOI mainly reflects cerebral venous saturation is still under discussion. Several studies report TOI values obtained in healthy adult volunteers.10–14 Quaresima et al compared TOI with cerebral venous oxygen saturation, measured by NIRS (NIRO 300, Hamamatsu), and concluded that TOI mainly reflected saturation of the intracranial venous compartment. Two studies using different NIRS instruments also found a correlation, but between TOI and jugular bulb oximetry.15 Other studies comparing jugular bulb oximetry with TOI did not find any correlation.11–15 Al-Rawi et al measured TOI in 60 patients undergoing endarterectomy. They found a significant correlation between TOI and flow velocity, measured by transcranial Doppler of the ipsilateral middle cerebral artery. The change in TOI was predominantly associated with internal carotid artery clamping, and a change during external carotid artery clamping was only seen if there was also a change in blood pressure. The sensitivity of TOI to intracranial and extracranial changes, when there were no pressure changes or an extracranial to intracranial anastomosis, was 87.5% and 0% respectively. They conclude that TOI predominantly measures intracranial changes. Another important finding was that a decrease in TOI is much more...
cerebral blood flow, and cerebral venous oxyhaemoglobin saturation, cerebral oxygen consumption, and cerebral fractional oxygen extraction can be performed with NIRS.\textsuperscript{12,13} However, measurement of cerebral venous oxyhaemoglobin saturation is still difficult and cannot be performed continuously. TOI is a non-invasive parameter for measurement of cerebral oxygenation and may be useful for continuous monitoring of venous oxyhaemoglobin saturation. Although several validation studies have been carried out in adults, further studies in animals need to be performed to study the relation between TOI, venous jugular saturation, and cerebral blood flow. In neonates, further studies are needed to determine the relation between TOI and venous oxyhaemoglobin saturation as measured by partial jugular venous occlusion. Further clinical studies should attempt to elucidate the variation in TOI over several days. These longer lasting measurements may be able to reflect the relation between periods of low cerebral oxygenation and neurological complications such as periventricular leukomalacia. As the measurement of TOI is less sensitive to movement artefacts, these studies may result in a new way to detect and prevent severe cerebral ischaemia by continuous monitoring of cerebral oxygenation.

In conclusion, measurement of TOI in premature infants is a new non-invasive method for measuring aspects of cerebral brain oxygenation. An increase in TOI over the first 3 days of life was found in 15 premature babies. This increase in TOI may reflect an increase in cerebral blood flow during this time.

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