Abstract

The measurement of saturation of venous blood as it drains out of brain by sampling it from the jugular bulb provides us with an estimate of cerebral oxygenation, cerebral blood flow and cerebral metabolic requirement. Arterio‑jugular venous difference of the oxygen content (AVDO₂) and jugular venous oxygen saturation (SjVO₂) values per se helps clinicians in identifying the impairment of cerebral oxygenation due to various factors thereby prompting implementation of corrective measures and the prevention of secondary injury to the brain due to ischaemia. SjVO₂ values are also used for prognostication of patients after traumatic brain injury and in other clinical situations. Sampling and measuring SjVO₂ intermittently or continuously using fibreoptic oximetry requires the tip of the catheter to be placed in the jugular bulb, which is a relatively simple bedside procedure. In the review below we have discussed the relevant anatomy, physiology, techniques, clinical applications and pitfalls of performing jugular venous oximetry as a tool for measurement of cerebral oxygenation.

Key words: Cerebral oxygenation, jugular bulb, jugular venous oximetry

INTRODUCTION

Minimising secondary injury to the brain after a primary insult forms the backbone of intervention in neuroanaesthesia and neurointensive care. However, in order to achieve this it is of absolute importance to identify the insult as promptly as possible so that appropriate intervention can be instituted. The salvageable portion of the brain, that is, the penumbra requires a constant supply of oxygen so that it does not undergo irreversible brain damage. Jugular bulb oximetry is one such monitor, which provides information about the status of cerebral oxygenation and helps in guiding the therapy to prevent secondary injury due to various factors.

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CLINICAL ANATOMY

Blood from the brain drains mainly through the venous sinuses. Almost whole of the blood comes into the right and left sigmoid sinus, which continue downwards as respective IJVs after passing through the jugular foramen. The jugular bulb is the dilatation of the jugular vein as it exits from the base of the skull. The tip of the catheter is placed in the jugular bulb for withdrawal of a sample or continuous monitoring of SjVO$_2$ [Figure 1]. The jugular bulb contains the blood, which is being drained out from both sides of the brain, out of which around 70% is from the same hemisphere and 30% is from the opposite hemisphere. Most of the patients will have a dominant side of the venous drainage and in the majority of them it will usually be right. The side with major venous drainage, as seen on the cerebral angiogram, gives an idea about the side of dominance.

The comparison of jugular foramen size using computed tomography scan can also be used to predict the side of the predominant drainage, with the larger size of foramen corresponding to the side of the major drainage. Ultrasonography can also be used to compare the size of the veins with the vein having larger diameter being the one with dominant drainage. The patients in whom the intracranial pressure (ICP) is being monitored, the dominant vein can be determined by the one with a higher rise in ICP following manual compression of each side. It is assumed that compressing the side with more drainage will cause a greater increase in ICP as a larger portion of cerebral outflow will be occluded.

MONITORING SITE

Patients sustaining an injury to both the hemispheres of the brain should have a catheter placed in the jugular bulb on the side of major drainage, which in most of the cases is generally right. However, if the patient has a focal injury, then there are no clear cut guidelines regarding the side on which the catheter should be placed. As more than two-third of the blood from the brain is drained through the ipsilateral IJV, it may be reasonable to cannulate the IJV on the side of the lesion, even if it is not the side with dominant drainage.

TECHNIQUES OF MEASUREMENT

A catheter that is used to monitor the SjVO$_2$ is almost same as the one, which is used to monitor the central venous pressure. The catheter is inserted in the IJV in a retrograde direction and the tip is advanced up to the jugular bulb. Intermittent samples may be drawn and SjVO$_2$ measured. It is important to note that the rate of aspiration of blood for measurement should be <2 ml/min to avoid contamination from extracranial vessels. Fibre optic technology has allowed the development of in vivo catheters wherein reflectance oximetry using such catheters allows continuous SjVO$_2$ monitoring, thus obviating the need for repeated sampling and measurement.

Oxyhaemoglobin has a unique light absorption spectrum and this forms the basis of fibreoptic oximetry. Catheters that are used to measure SjVO$_2$ contains two optical fibres, one of which is used to direct the light into the blood and the other one is used to transmit the reflected light to a sensor. The sensor measures the absorbed light which is reflected at different wavelengths. The SjVO$_2$ values are therefore a percentage of oxygenated haemoglobin to the total haemoglobin. These catheters using a dual wavelength of light (Edslab, Baxter-Edwards system) require that the haemoglobin concentration of the patient to be manually entered. The SjVO$_2$ values are thus dependent on the values entered. Alternatively, the catheters which use three wavelengths (Opticath Oximetrix, Abbott Critical Care System) calculate the haemoglobin concentration from the absorption spectrum itself and hence can be used for a real time, continuous monitoring of SjVO$_2$.

CATHETER INSERTION

Puncture site for insertion is similar to that used for IJV cannulation. However, the needle followed by guidewire and subsequently the catheter are all advanced in a direction towards the skull [Figure 1]. To prevent injury, it is important that the tip of the guidewire is J-shaped and is advanced only 3–4 cm beyond the site of needle insertion. The catheter is advanced over the guide wire in a cephalad direction till there is a feeling of resistance at the jugular bulb. The catheter is subsequently withdrawn for about 0.5–1 cm so that that the tip does not lodge against the roof of the jugular bulb. This also reduces the risk of injury.
to the bulb and prevents occlusion of the catheter tip. Markings on the guidewire may also be used to help us in determining the distance to which the catheter should be inserted [Figure 2]. It should be 1 cm lesser than the length as measured from the point at which the catheter enters the neck up to the mastoid process. Skull and neck radiography can be used to confirm the correct placement of the catheter.[16] On a lateral radiograph of the neck, the tip of the catheter should be at the level of mastoid process and just medial to it and also above the lower border of C1 vertebrae, as the aspiration of blood from this position will reduce the chance of contamination from extracranial vessels [Figure 3]. Alternatively, fluoroscopy can also be used intraoperatively for quick confirmation of the position of the catheter tip.

**PRINCIPLE OF JUGULAR VENOUS OXIMETRY**

Jugular venous saturation indirectly gives us an idea of the use of oxygen by the brain. To put it in simpler terms when the demand for oxygen is more, the brain extracts a greater amount of oxygen, resulting in decreased jugular bulb oxygen saturation. When cerebral oxygen supply exceeds demand, the extraction is less and saturation of oxygen in the venous blood increases. If the CBF reduces too much, then a point is reached when the neurons of the brain cannot tolerate the reduction of blood flow. After this stage is reached, the consumption of oxygen reduces leading to anaerobic metabolism and production of lactate. The oxygen saturation in jugular bulb is related to the CMRO\textsubscript{2} and CBF as represented by the Ficks equation:

\[
\text{CMRO}_2 = \text{CBF} \times (\text{CaO}_2 - \text{CjO}_2)
\]

\[
\text{CaO}_2 \text{ (Arterial oxygen content)} = \text{SaO}_2 \times 1.34 \times \text{Hb} + 0.0031 \times \text{PaO}_2
\]

\[
\text{CjO}_2 \text{ (Jugular venous oxygen content)} = \text{SjO}_2 \times 1.34 \times \text{Hb} + 0.0031 \times \text{PjO}_2
\]

The contribution from dissolved oxygen, as we know, is very small and can safely be neglected. The difference between arterial oxygen content and oxygen content of venous blood is represented as (\text{CaO}_2 - \text{CjO}_2) or AVDO\textsubscript{2} (arterio-venous oxygen difference). If the above equation is rearranged then:

\[
\text{AVDO}_2 = \text{CMRO}_2/\text{CBF}
\]

Normal AVDO\textsubscript{2} values are between 4 ml/dl and 8 ml/dl.[17,18] Presuming that if CMRO\textsubscript{2} does not change then any change in AVDO\textsubscript{2} should be because of the change in CBF. If AVDO\textsubscript{2} is <4 ml/dl of blood, it is probably because the oxygen supply is in excess of the demand (i.e., luxury perfusion or infarction). An AVDO\textsubscript{2} value more than 8 ml/dl of blood points towards the demand being more than the supply (i.e., ischaemia). If CMRO\textsubscript{2} increases but the CBF does not increase concomitantly, then there will be enhanced extraction of oxygen from the blood. There is a proportional decrease in oxygen content, as well as oxygen saturation, of the venous blood coming out of the brain with eventual widening of AVDO\textsubscript{2}. In a healthy brain flow-metabolism coupling is preserved and SjVO\textsubscript{2} ranges between 55% and 75%.[2] However in pathological conditions, the coupling between CMRO\textsubscript{2} and CBF is lost and in circumstances where hypoperfusion is not accompanied by proportional reduction in CMRO\textsubscript{2}, the brain will extract a greater proportion of arterial oxygen and SjVO\textsubscript{2} will be observed to fall. Values below 54% imply that oxygen supply may be critically low for the metabolic demand and the brain is at risk for ischaemic injury.[19] Value more than 75% is representative of cerebral hyperaemia or decreased oxygen extraction as in cerebral infarction.

![Figure 2: Estimation of insertion length using a guidewire](image-url)

![Figure 3: Correct positioning of catheter tip seen on X-ray of neck and skull (lateral view)](image-url)
DETERMINANTS OF JUGULAR VENOUS OXYGEN SATURATION

A reduction in SjVO₂ may be due to fall in supply of oxygen or an increase in usage of oxygen by the brain whereas an increase in SjVO₂ values may be seen due to increase in oxygen delivery or a reduction in oxygen consumption:

**Decreased jugular venous oxygen saturation (<50%)**
- Due to reduced supply of oxygen: Cerebral vasoconstriction, hypocapnia, vasospasm, hypotension, anaemia and sepsis
- Due to increased oxygen requirement: Increased cerebral metabolism, which include causes such as fever, agitation, inadequate sedation, pain and seizures.

**Increased jugular venous oxygen saturation (>75%)**
- Increased supply of oxygen: Cerebral vasodilation, hypercapnia, hypertension
- Reduced oxygen requirement: Deep sedation, coma, hypothermia, cerebral infarction, brain death.

**CLINICAL APPLICATIONS**

**Traumatic brain injury**

In patients who have sustained a traumatic brain injury, SjVO₂ monitoring provides a means to identify ischaemia, which may be due to either systemic or intracranial causes. The goal is to maintain a SjVO₂ value above 50% as we know that the values lesser than this suggest a relative failure of oxygen supply compared with demand and episodes of such desaturation have demonstrated to predict poor outcome after head injury. Whenever hyperventilation is to be instituted, SjVO₂ monitoring is a valuable aid in guiding its optimal usage without causing cerebral hypoperfusion. As per the brain trauma foundation guidelines for the management of traumatic brain injury, prophylactic hyperventilation (<25 mm Hg PaCO₂) is not recommended, more so in the first 24 h after injury. If other measures to reduce ICP have failed, then hyperventilation is to be used only as a temporary measure. It is vital to monitor SjVO₂ or brain tissue oxygen tension (PBrO₂) while hyperventilating this group of patients. SjVO₂ monitoring can also be used to guide administration of fluids and level of oxygenation and optimise the cerebral perfusion pressure.

**Lactate oxygen index (LOI)** can also be used to predict outcome in traumatic brain injury patients. Its normal values are <0.03 and values >0.08 are seen in patients with failing oxygen extraction, leading to failure of aerobic metabolism and increased production of lactic acid in the brain especially in the first 24 h after head injury. It is calculated by the formula mentioned below:

\[
\text{LOI} = -\frac{\text{AVDL}}{2.24} \times \frac{\text{AVDO₂}}{\text{AVDL}}
\]

AVDL = Arterial to jugular difference for lactate
AVDO₂ = Arterial to jugular difference in oxygen content.

**Neurosurgery**

SjVO₂ monitoring has been extensively used by Matta et al., in neurosurgical patients. They demonstrated that the SjVO₂ catheter could be placed quickly and can detect frequent critical episodes of SjVO₂ desaturation that would otherwise have been undetected and untreated. During intracranial aneurysm surgery, SjVO₂ monitoring has been used to determine the minimal blood pressure that should be maintained to avoid hypoperfusion and also to measure LOI. Intraoperative LOI >0.08 during surgery for aneurysm clipping has been associated with a poor outcome. SjVO₂ monitoring can also be used in any neurosurgical procedure to help in deciding the appropriate mean arterial pressure and PaCO₂ values to ensure adequate cerebral oxygenation. Our own experience suggests that measurement of SjVO₂ provides important information regarding cerebral oxygenation and is a potential means to optimize cerebral perfusion.
in patients undergoing clipping of ruptured cerebral aneurysm is feasible and can help in guiding the perioperative anaesthetic management.[32]

Cardiac surgery
Neurologic dysfunction is seen quite commonly after cardiac surgery especially in those patients who undergo cardiopulmonary bypass.[33] Most of the episodes of SjVO₂ desaturation are seen during the period of rewarming after hypothermic cardiopulmonary bypass. SjVO₂ desaturation can result in increased incidence of post-operative cognitive dysfunction.[34–36] It would be prudent to monitor SjVO₂ during cardiac surgery to prevent these episodes of desaturation and improve the cognitive outcome.[37]

Cardiac arrest
SjVO₂ values are higher in the non-survivors than the survivors of cardiac arrest (values of 80% vs. 67%) probably due to the inability of dead neurons to utilise the supplied oxygen.[38] Moreover, SjVO₂ values more than the mixed venous oxygen saturation are also associated with poor outcome.[39] However, these findings are still controversial and have been countered by studies, which did not find any difference in the SjVO₂ values in between survivors and non-survivors of cardiac arrest.[40]

Research applications
Jugular venous oximetry can be used to guide research to evaluate the effect of various anaesthetic agents on CBF. In a recently concluded study comparing propofol and desflurane in patients undergoing clipping of aneurysmal neck after subarachnoid haemorrhage, it was observed that use of desflurane was associated with increase in SjVO₂ desaturation can result in increased incidence of post-operative cognitive dysfunction.[35–36] It would be prudent to monitor SjVO₂ during cardiac surgery to prevent these episodes of desaturation and improve the cognitive outcome.[37]

DRAWBACKS AND LIMITATIONS
• It is an invasive procedure requiring the insertion of the catheter retrograde into the jugular vein. Cannulating the vein may fail at times due to anatomical variations and abnormalities. Carotid puncture and subsequent haematoma formation can also occur
• Malpositioning of the catheter tip can occur which may give erroneous readings if the position of the tip is not confirmed prior to the measurement of SjVO₂ values
• Mechanical complications of injury to the vessel wall or jugular bulb may occur. Infectious complications have also been reported though the incidence of such complications is very less[41]

CONCLUSIONS
Jugular venous oximetry is the oldest technology for the measurement of cerebral oxygenation and hence becomes a natural benchmark against which newer modalities of monitoring are evaluated. Despite its limitations, it is
a relatively low-cost monitor and the fact that it can be inserted at the bedside to assess the adequacy of cerebral oxygenation makes it quite attractive. Its role becomes even more important when hyperventilation or blood pressure directed management has to be instituted for patients with traumatic brain injury and other intracranial pathologies. As a part of multimodality monitoring which includes ICP monitoring, monitoring of CBF by transcranial Doppler and assessing the adequacy of cerebral oxygen delivery at the target area by brain tissue oxygen measurement (PbO2) and cerebral microdialysis catheters, SjVO2 monitor functions as a tool for global cerebral oxygenation and helps to guide various goal-directed therapies in the management of patients with various intracranial pathologies. It subserves as an important tool for research related to various interventions, which can influence CBF and cerebral oxygenation. Hence in the absence of other viable alternatives to measure the adequacy of cerebral oxygenation, Jugular venous oximetry will continue to thrive as a useful monitoring modality for this purpose.

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Conflicts of interest
There are no conflicts of interest.

REFERENCES


