

## Tissue oxygenation monitoring (StO<sub>2</sub>)

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Note: *Select one best option*

1. Tissue oxygenation monitors utilise;
  - a. infrared light
  - b. near infrared light
  - c. ultra violet light
  - d. All of the above
  - e. None of the above
2. Tissue oxygenation monitor works;
  - a. Invasively
  - b. Non-invasively
  - c. Minimally invasively
  - d. With remote sensing
  - e. None of the above
3. Tissue oxygenation monitor has been proved to be superior than SpO<sub>2</sub> monitors in;
  - a. Resuscitation of trauma patients
  - b. Septic shock
  - c. Hypothermic patients
  - d. All of the above
  - e. None of the above
4. Serious hypoperfusion (inadequate blood flow) is indicated by StO<sub>2</sub> levels of;
  - a. 70%
  - b. 75%
  - c. 80%
  - d. 85%
  - e. 90%
5. The most sensitive site for probe application in StO<sub>2</sub> monitor is;
  - a. Index finger
  - b. Thumb
  - c. Thenar eminence
  - d. Forearm
  - e. Ear lobe
6. Peripheral vasoconstriction effects more the measurement of;
  - a. SpO<sub>2</sub>
  - b. NIBP
  - c. CVP
  - d. StO<sub>2</sub>
  - e. All of the above
7. The following statement regarding StO<sub>2</sub> is true;
  - a. It is the quantification of the ratio of oxygenated hemoglobin to total hemoglobin in the microcirculation of a volume of illuminated tissue and is an absolute number
  - b. Its principle of detecting oxygen saturation is similar to measuring oxygen saturation by pulse oximetry
  - c. It is a valid measure of hemoglobin oxygen saturation
  - d. a and c only
  - e. All of the above
8. One important difference between SpO<sub>2</sub> and StO<sub>2</sub> is;
  - a. Unlike SpO<sub>2</sub>, StO<sub>2</sub> is a measure of oxygen saturation in the microcirculation where oxygen is exchanged with tissue and is therefore a local measure
  - b. StO<sub>2</sub> uses near infrared light, whereas SpO<sub>2</sub> uses infrared light
  - c. Measurements of SpO<sub>2</sub> and StO<sub>2</sub> both will not change with changes in the local conditions of supply and consumption in the tissue
  - d. None of the above
  - e. All of the above
9. The need for transfusion is best predicted by;
  - a. blood pressure
  - b. arterial lactate
  - c. tachycardia
  - d. StO<sub>2</sub>
  - e. hemoglobin concentration
10. StO<sub>2</sub> has been shown to possess a high “negative predictive value” (NPV) in several clinical trials in trauma patients of;
  - a. a 91% chance of not developing multiple organ dysfunction
  - b. a 96% chance of survival
  - c. a 93% chance of receiving blood transfusion
  - d. None of the above
  - e. All of the above

### ANSWERS:

1: b; StO<sub>2</sub> employs the use of near infrared light, which is transmitted through the skin, into the underlying tissues and takes measurements of the light, based on a patented algorithm, after it has achieved greater levels of penetration into the tissues.

2: b; It is a non-invasive technique.

3: d; Various researchers have now confirmed that StO<sub>2</sub> is more sensitive index than SpO<sub>2</sub> in all of these conditions.

4: b; Thenar StO<sub>2</sub> values have been shown by Crookes BA and colleagues to be: normals, 87 +/- 6%; no shock, 83 +/- 10%; mild shock, 83 +/- 10%; moderate shock, 80 +/- 12%; and severe shock, 45 +/- 26%. 75% is usually accepted as the cut-off point for hypoperfusion) in trauma patients, and that StO<sub>2</sub> levels above 75% indicates adequate perfusion.

5: d; A recent study by Bezemer R and colleagues has shown that forearm StO<sub>2</sub> is a more sensitive parameter to hemodynamic changes than thenar StO<sub>2</sub> for the detection of central hypovolemia and hypovolemic shock in (trauma) patients and that the depth at which StO<sub>2</sub> is measured is of minor influence.

6: d; SpO<sub>2</sub> requires a pulsatile flow, while StO<sub>2</sub> readings do not. NIBP and CVP are also effected by many different pathologic conditions of the liver, heart and kidneys and are poor predictors of the tissue perfusion.

7: d; StO<sub>2</sub> is the quantification of the ratio of oxygenated hemoglobin to total hemoglobin in the microcirculation of a volume of illuminated tissue and is an absolute number. The measurement of StO<sub>2</sub> is taken with a noninvasive, fiber optic light that illuminates tissues below a sensor placed on the skin. StO<sub>2</sub> correlates well with other accepted means of measuring oxygen saturation and the results of these studies demonstrate that StO<sub>2</sub> is a valid measure of hemoglobin oxygen saturation. StO<sub>2</sub> is a measure of oxygen saturation in the microcirculation where oxygen is exchanged with tissue and is therefore a local measure. Pulse oximetry, which also uses near infrared light, measures the systemic oxygen saturation of arterial blood.

8: a; StO<sub>2</sub> measurements differ from the SpO<sub>2</sub> measurements provided by pulse oximetry. StO<sub>2</sub> is a measure of oxygen saturation in the microcirculation where oxygen is exchanged with tissue and is therefore a local measure. Pulse oximetry, which also uses near infrared light, measures the systemic oxygen saturation of arterial blood. Measurements of StO<sub>2</sub> will therefore change with changes in the local conditions of supply and consumption in the tissue, and SpO<sub>2</sub> will not.

9: d; A recent clinical study involving 26 trauma patients at risk for hemorrhagic shock. Results from this study showed that of patients who required a transfusion within 24 hours of arrival in the ED, 88% had a minimum StO<sub>2</sub> below 70% in the first hour of arrival in the ED,\* and of those who did not require a transfusion, only 22% had StO<sub>2</sub> values that dropped below 70% for the first hour. The need for transfusion was not predicted by conventional indicators of hypoperfusion and shock (blood pressure, arterial lactate, tachycardia, base deficit), nor by hemoglobin concentration.

10: e; StO<sub>2</sub> has been shown to possess a high NPV in several clinical trials in trauma patients. The first of these showed that, in patients believed to be at significant risk of shock, those who maintained an StO<sub>2</sub> at 75% or greater in the first hour of arrival in the ED had a 91% chance of not developing multiple organ dysfunction, and a 96% chance of survival. The clinical study by Smith et al.<sup>12</sup> described above had a similar conclusion specifically with regard to the need for transfusion in a high risk trauma population. The probability of not requiring a transfusion within 24 hours of arrival at the ED was 93% if the StO<sub>2</sub> did not drop below their designated threshold of 70% for a period of one minute or more in the first hour.

### REFERENCES

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