

# Neuro Quiz 49: Monitoring the Brain



START

*THIS QUIZ IS BEING  
PUBLISHED ON BEHALF OF THE  
EDUCATION COMMITTEE OF  
THE SNACC*

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# 1. Which of the following statements about Intra-Cranial Pressure (ICP) monitoring is **TRUE**?

- A. The transducer of the intra-parenchymal pressure monitor is placed at the level of the tragus
- B. The intra-ventricular pressure monitor does not provide a global measurement of ICP
- C. ICP- guided therapy improves outcomes in patients with traumatic brain injury
- D. ICP monitoring is indicated in brain injured patients with GCS  $\leq 8$  even if the CT scan is normal

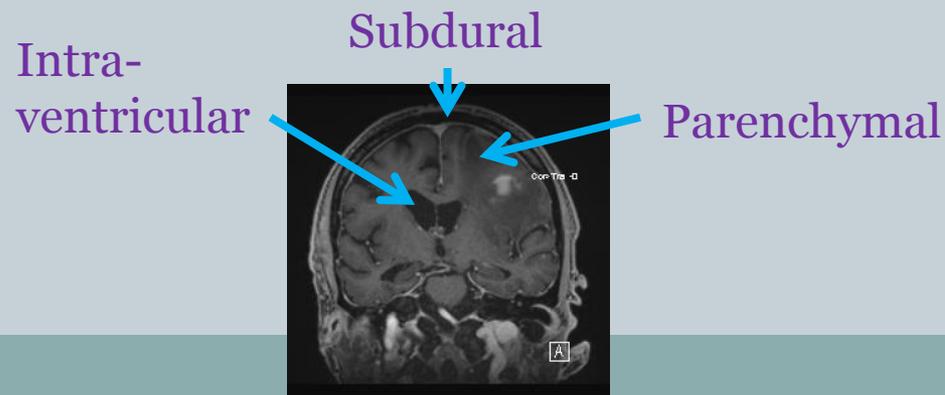
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## 1A. The transducer of the intra-parenchymal pressure monitor is placed at the level of the tragus



- **Intra-parenchymal** ICP monitors have micro-transducers located at the tip of the catheters and their position cannot be changed.
- The miniature transducer technology varies with different manufacturers, e.g. Codman has a semiconductor strain gauge attached to a thin diaphragm. Any change in ICP distorts the membrane and changes the resistance of the strain gauge which is measured by a Wheatstone bridge and displayed as ICP
- These transducers are zeroed before insertion and cannot be recalibrated *unlike* **the Intra-ventricular monitors**

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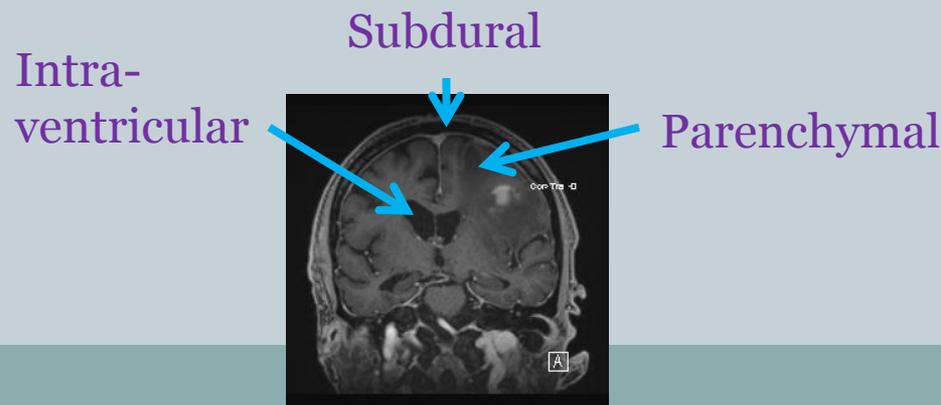
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## 1B. The intra-ventricular pressure monitor does not provide a global measurement of ICP



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- The **intra-ventricular catheter** is usually inserted into the lateral ventricles and it measures the **global ICP**
- The transducer should be kept at the level to the tragus
- When recording the cerebral perfusion pressure (CPP), the transducer measuring the arterial pressure should also be placed at the level of the tragus



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# 1C. ICP- guided therapy improves outcomes in patients with traumatic brain injury



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- Although, ICP monitoring is the standard of care for all patients with severe brain injury, there is no Class I evidence suggesting that ICP-guided therapy improves outcomes in such patients

Global Neurotrauma Research Group. A trial of intracranial-pressure monitoring in traumatic brain injury. *N Engl J Med* 2012;367:2471-81.

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1D. ICP monitoring is indicated in brain injured patients with GCS  $\leq 8$  even if the CT scan is normal



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- This statement is correct
- ICP monitoring allows early detection of an expanding lesion and the CPP
- Cerebrovascular Pressure reactivity (PRx) index is a correlation of consecutive values of ICP and arterial pressure
  - A positive PRx suggests impaired autoregulation
  - A negative value reflects normal autoregulation
  - PRx can be used to estimate optimal CPP levels for individual patients

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## 2. Which of the statements about Intra-cranial pressure (ICP) monitoring is **TRUE**?

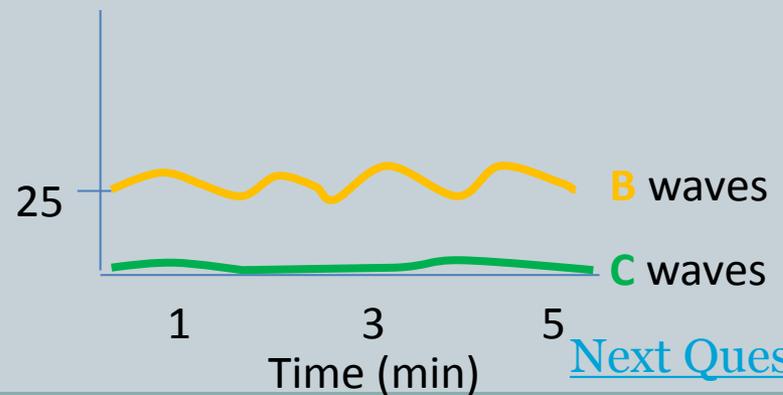
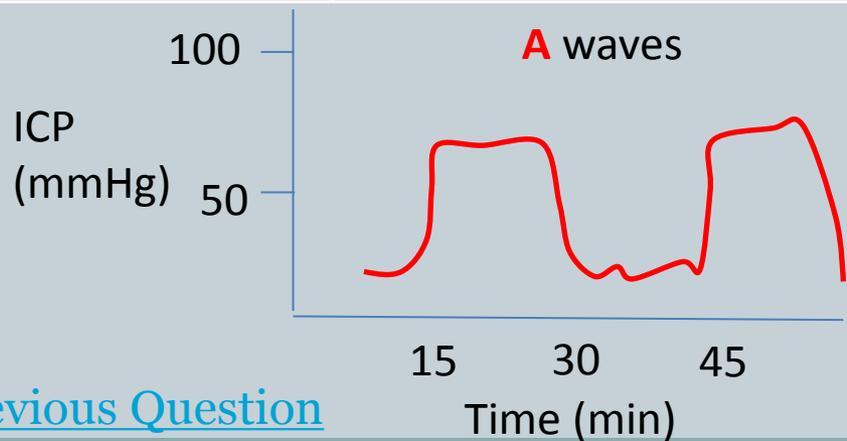
- A. Lundberg type C wave indicates a poor prognosis in a patient with brain injury
- B. The normal ICP in a 6 month old child is 10 mmHg
- C. The normal ICP tracing is pulsatile
- D. To calculate the cerebral perfusion pressure, the transducers measuring the MAP and the ICP should be zeroed at the level of the patient's heart

# 2A. Lundberg type C wave indicates a poor prognosis in a patient with brain injury



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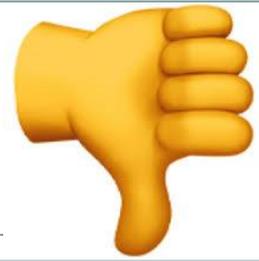
| Lundberg | Features                                 | Clinical                                     |
|----------|--|--|
| <b>A</b> | Plateau shaped; 50-100 mmHg; 5-20min     | Pathological, Very high brain impedance      |
| <b>B</b> | Rhythmic oscillations; <50 mmHg; 1-2 min | High brain impedance                         |
| <b>C</b> | Rhythmic oscillations; <20 mmHg; 4-8min  | Normal, synchronous with arterial pulsations |



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2B. The normal ICP in a 6 month old child is 10 mmHg



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- This is incorrect
- The normal ICP is 3-4 mmHg up to 1 year of age and 10-15 mmHg in adults

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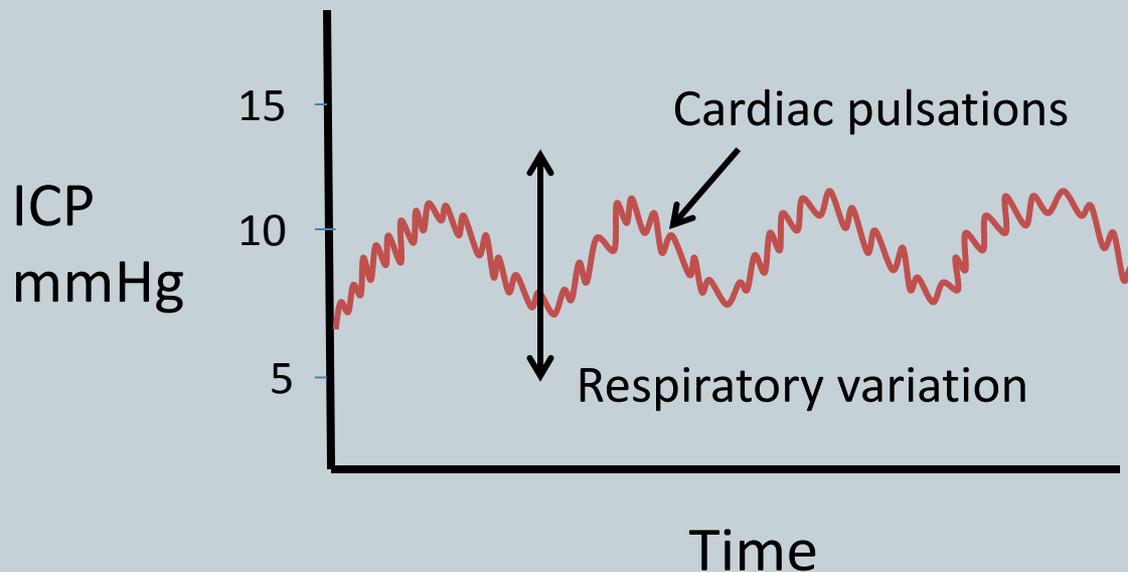
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## 2C. The normal ICP tracing is pulsatile



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- Dynamic tracing of the ICP reflects the cardiac pulsations and the respiratory variations



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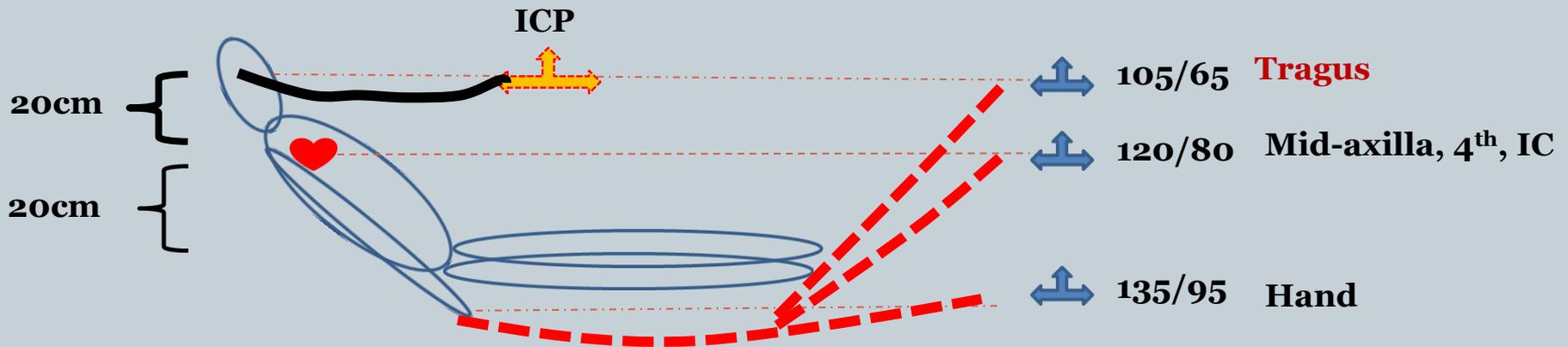
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2D. To calculate the cerebral perfusion pressure, the transducers measuring the MAP and the ICP should be zeroed at the level of the patient's heart



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- This is incorrect
- The transducers measuring the arterial pressure and the intraventricular ICP monitor should be at the level of the Circle of Willis, which corresponds to the **tragus**.



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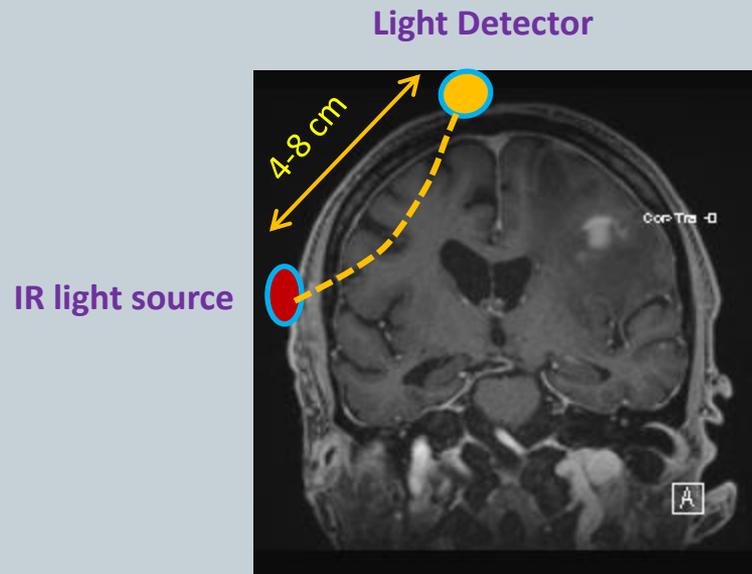
### 3. Which of the following statements about measuring Cerebral Oxygenation is **TRUE**?

- A. Near Infrared spectroscopy gives a good assessment of global cerebral oxygenation
- B. The catheter tip of the Jugular venous oxygen saturation ( $S_{jv}O_2$ ) monitor should be at the level of C1/C2 spine
- C. Normal brain tissue oxygen pressure  $P_{br}O_2$  is  $<15\text{mmHg}$
- D. Brain tissue oxygen is measured by aspirating tissue fluid and analyzing it in a standard lab

## 3A. Near Infrared spectroscopy gives a good assessment of global cerebral oxygenation



- This is incorrect as the NIRS is a non-invasive method to measure **regional** cerebral oxygenation [Try Again](#)
- Infrared light (700-1000nm) is able to penetrate skin, bone and brain tissue and is absorbed by HbO<sub>2</sub> & Hb



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3B. The catheter tip of the Jugular venous oxygen saturation (SjvO<sub>2</sub>) monitor should be at the level of C<sub>1</sub>/C<sub>2</sub> spine



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- The SjvO<sub>2</sub> catheter is inserted into the internal jugular and passed cephalad to reach the jugular bulb and confirmed with X-ray



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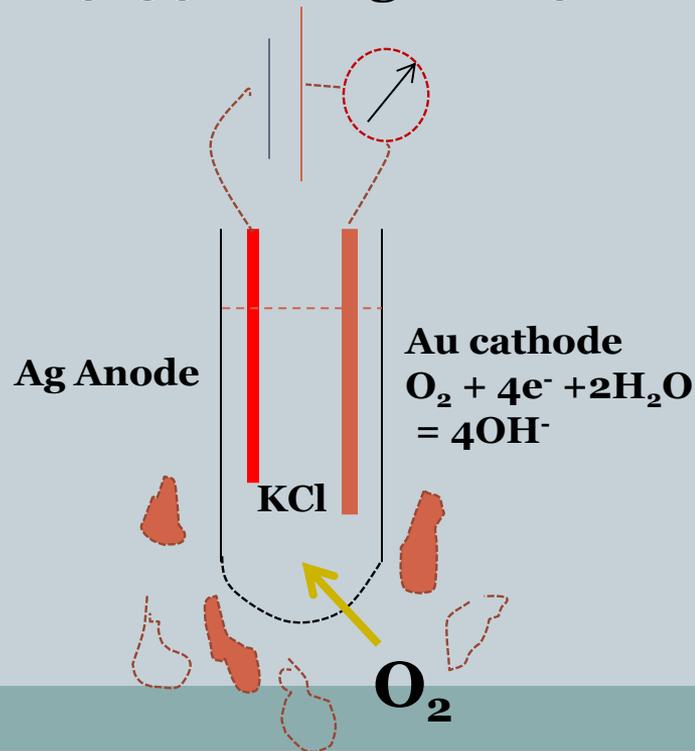
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# 3C. Normal brain tissue oxygen pressure is <15 mmHg



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- Micro-catheters with a Polarographic electrode incorporated into its tip are placed into the brain tissue to measure  $P_{br}O_2$
- Normal  $P_{br}O_2$  is 25-35mmHg & <15 mmHg suggests local ischemia



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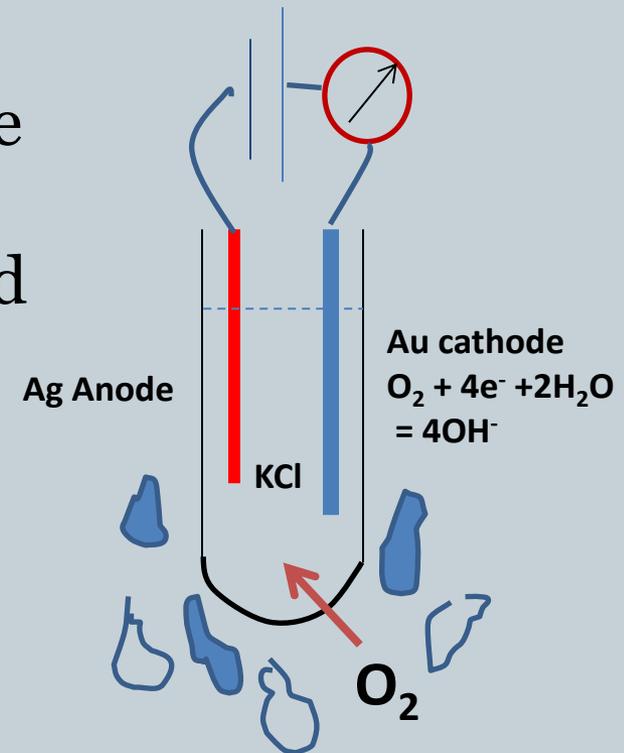
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### 3D. Brain tissue oxygen is measured by aspirating tissue fluid and analyzed in a standard lab



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- This is incorrect
- Micro-catheters with a Polarographic electrode incorporated into its tip are placed into the brain tissue to measure  $P_{br}O_2$  directly
- pH electrodes can also be incorporated to measure pH and  $PCO_2$  levels
- Normal values
  - $P_{br}O_2$  – 25-35 mmHg
  - $P_{br}CO_2$  – 40-70 mmHg
  - pH – 7.05-7.25



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#### 4. Which of the following statements about monitoring Cerebral Blood Flow (CBF) is **FALSE**?

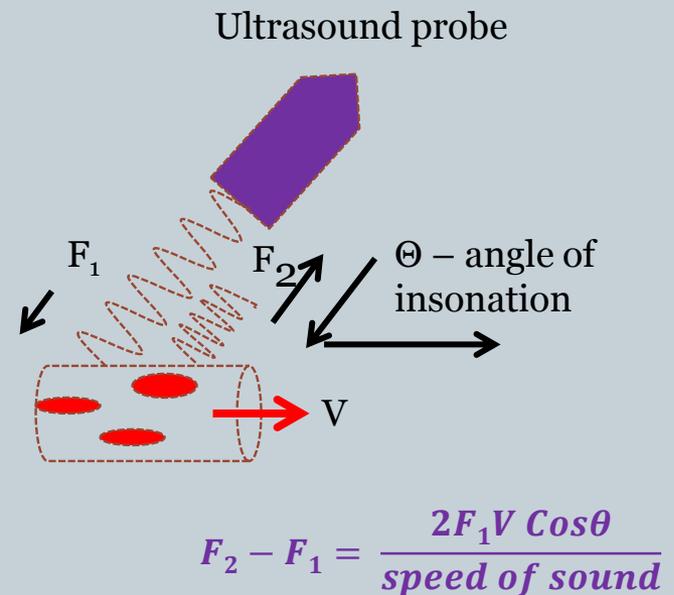
- A. Transcranial Doppler (TCD) study is reliable for monitoring vasospasm after SAH
- B. TCD can be used to estimate the ICP by measuring the pulsatility index
- C. Xenon-enhanced CT scan can be used to quantify CBF
- D. Measuring CBF by CT perfusion scan is time consuming and clinically unreliable

## 4A. Transcranial Doppler (TCD) study is reliable for monitoring vasospasm after subarachnoid hemorrhage



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- This is a correct statement
- A perceived change in frequency when a sound wave is reflected off a moving object is **Doppler Effect**, and the change depends on the velocity of the moving object
- TCD is used to monitor vasospasm after SAH. A flow velocity in the MCA of  $>120$  cm/s with a Lindegaard index of 3-6 is highly suggestive of vasospasm
- $$\text{Lindegaard Index} = \frac{\text{FV in MCA}}{\text{FV in Int Carotid}}$$



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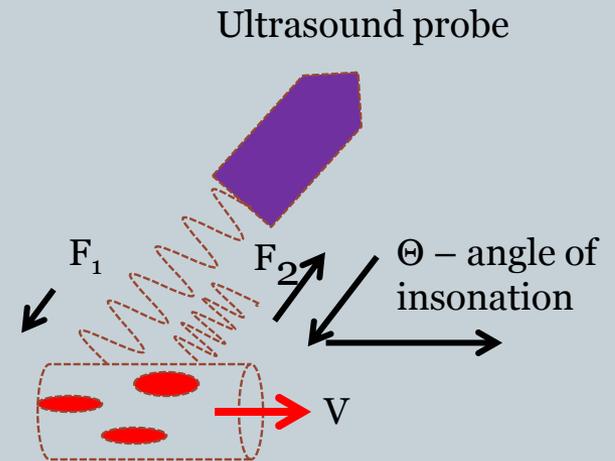
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## 4B. TCD can be used to estimate the ICP by measuring the pulsatility index



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- $Pulsatility\ Index\ (PI) = \frac{Systolic\ FV - Diastolic\ FV}{Mean\ Flow\ velocity\ (FV)}$
- There is a strong correlation between ICP & PI
- Surg Neurol. 2004;62:45-51
- TCD can also detect micro-emboli and intraoperative cerebral perfusion during carotid surgery



$$F_2 - F_1 = \frac{2F_1V \cos\theta}{\text{speed of sound}}$$

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## 4C. Xenon-enhanced CT scan can be used to quantify CBF



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- Xenon, being highly lipid soluble, can readily cross the blood-brain barrier and enhance the CT scans
- After a baseline CT scan the patient breathes xenon till it equilibrates
- The xenon is then discontinued and serial scans are performed to analyze the washout of xenon which is used to quantify the CBF

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## 4D. Measuring CBF by CT perfusion scan is time consuming and clinically unreliable



- The second part of this statement is incorrect
- CBF can be measured **accurately** by CT perfusion, especially in acute stroke and SAH to delineate the area of potentially reversible ischemic penumbra from the infarcted area
- After administering a contrast dye scan slices at the level of the basal ganglia are taken to visualize the anterior, middle and posterior cerebral artery territories
- These methods are expensive, time-consuming and put the patient at risk of contrast agents related problems and also transportation to a remote facility

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5. Which of the following statements regarding monitoring the brain metabolism using the Cerebral Micro-dialysis is **TRUE**?

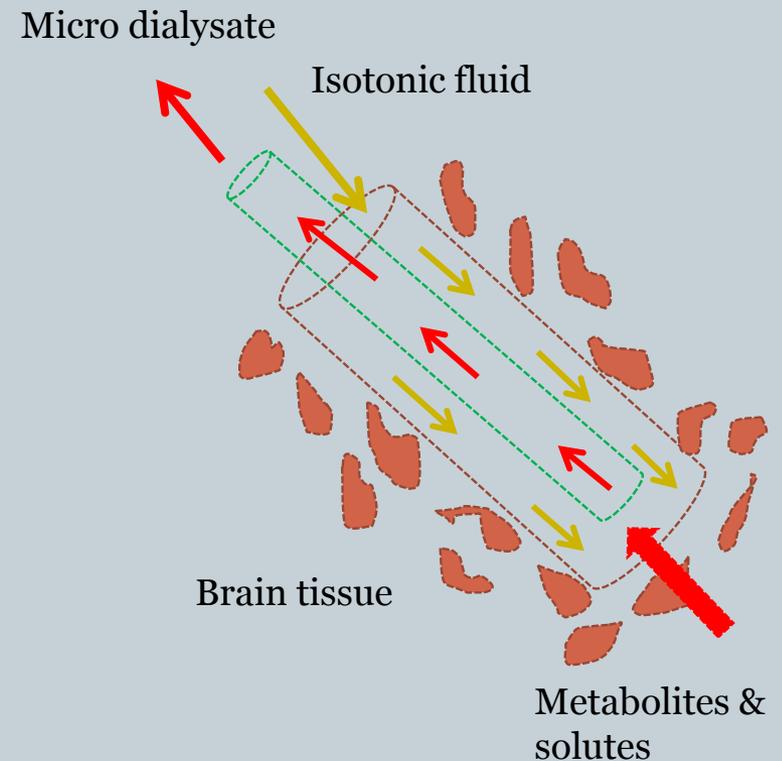
- A. The brain tissue fluid is directly aspirated and the concentration of the metabolites measured
- B. It is used as a test to confirm secondary brain injury after it is evident on other monitors
- C. A high Lactate-Pyruvate ratio indicates cerebral ischemia
- D. A rise in glucose in micro-dialysate 2-3 days after a brain injury indicates cell death

## 5A. The brain tissue fluid is directly aspirated and the concentration of the metabolites measured



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- This is incorrect
- The micro-dialysis probe is essentially a coaxial catheter with a semipermeable dialysis membrane lining its tip
- Through the outer channel, fluid, isotonic to the brain extracellular fluid, is pumped at  $0.3\mu\text{L}/\text{min}$  and aspirated back through the inner tube
- The dialysis membrane at the tip allows diffusion of water and solutes from the interstitial fluid into the catheter along its concentration gradient



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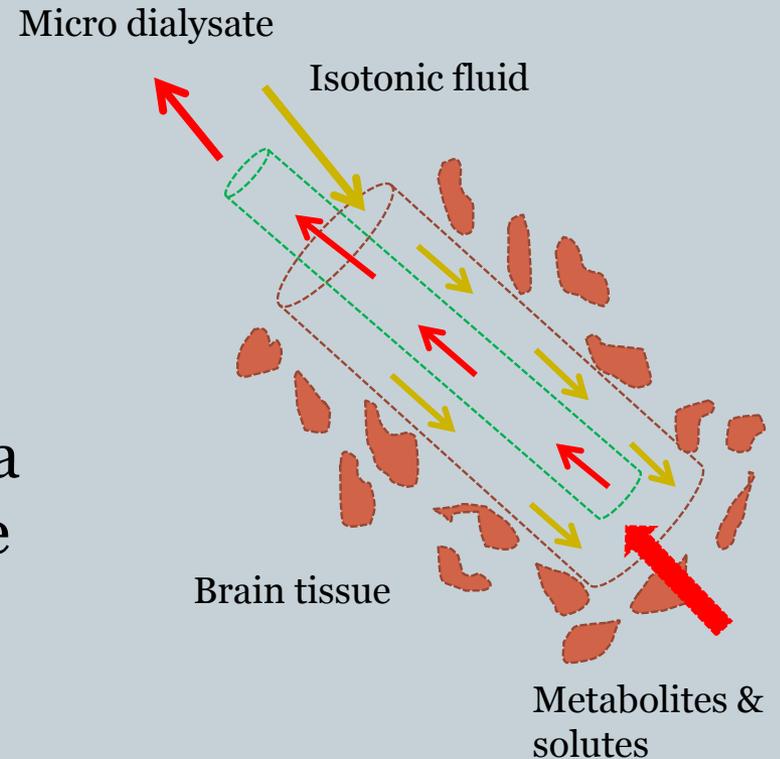
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5B. It is used as a test to confirm secondary brain injury after it is evident on other monitors



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- Cerebral micro-dialysis can detect changes in the metabolism at the cellular level **before** changes are detected in other monitors for brain physiology
- The micro-dialysis probe is essentially a coaxial catheter with a semipermeable dialysis membrane lining its tip that allows diffusion of cellular metabolites



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## 5C. A high Lactate-Pyruvate ratio indicates cerebral ischemia



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Metabolites and solutes measured by cerebral micro-dialysis

- Energy related metabolites – glucose, lactate, pyruvate
  - **Markers of secondary brain ischemia**
    - ✦ Glucose  $<1.5$  mmol/L
    - ✦ Raised lactate to pyruvate ratio ( $>20$ )
- Neurotransmitters – glutamate, aspartate
  - High levels are seen in secondary cerebral ischemia.
- Cellular degradation markers – glycerol, potassium
  - Glycerol is produced by degradation of the phospholipids from dead cells. High levels have been measured after severe TBI and also secondary ischemia.
- Exogenous – drugs

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5D. A rise in glucose in the micro-dialysate 2-3 days after a brain injury indicates cell death



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This is incorrect.

A rise in **glycerol** indicates cell death

Glucose level **decreases** with brain ischemia

Metabolites and solutes measured by cerebral micro-dialysis

- Energy related metabolites – glucose, lactate, pyruvate
  - ✦ Markers of secondary brain ischemia
    - Glucose <1.5 mmol/L
    - Raised lactate to pyruvate ratio (>20)
- Neurotransmitters – glutamate, aspartate
  - ✦ High levels are seen in secondary cerebral ischemia.
- Cellular degradation markers – glycerol, potassium
  - ✦ Glycerol is produced by degradation of the phospholipids from dead cells. High levels have been measured after severe TBI and also secondary ischemia.
- Exogenous – drugs

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[References](#)

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