



Transcranial Doppler (TCD) Ultrasonography and its Clinical Application- A Review and Update



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Abstract

Transcranial Doppler (TCD) ultrasonography is a bedside, non-invasive duplex study of intracranial blood flow dynamic through some selected region or window in the cranial region. TCD provides information about the physiology of blood flow in the basal intracranial major arteries by measuring flow velocity and pulsatility in different segment of these arteries. The physiologic data obtained from the cerebrovascular hemodynamic are complimentary to anatomical data obtained from various imaging techniques and it can be used as bedside monitor to assess the cerebrovascular changes in response to any intervention. Common application of TCD includes detection of vasospasm after subarachnoid haemorrhage, vasocclusion in sick cell disease, cerebral strokes, intracranial pressure monitoring, intraoperatively monitoring to detect micro-embolism, early detection of brain death and cerebral auto regulatory testing. The review focus on basic principles of TCD, scanning techniques and described application of TCD in some of the common clinical conditions.

Keywords: Transcranial doppler ultrasonography; Cerebral blood flow; Cerebral vasospasm; Sick cell disease; Intracranial pressure; Brain death; Cerebral auto regulation

Introduction

Transcranial Doppler (TCD) ultrasonography is a non-invasive technique that utilised a pulsed low-frequency ultrasonic transducer of low frequency through some selected region (window) in the cranial region to enable recording to flow velocity pattern in intracranial arteries. Unlike the cerebral angiography which provides an anatomical configuration of both extra and intracranial arteries and their smaller branches, TCD provides information about the physiology of blood flow in the basal intracranial major arteries by measuring flow velocity and pulsatility in different segment of these arteries [1,2]. TCD being relatively inexpensive, non-invasive, reputable, portability and provides real time information that makes it more practical and convenience as bed side monitoring in intensive care setting for cerebral blood flow (CBF) velocity [3-5]. The physiologic data obtained from the cerebrovascular hemodynamic are complimentary to anatomical data obtained from various imaging techniques and it can be used as bedside monitor to assess the cerebrovascular changes in response to any intervention [4,5].

Aaslid et al. [6] in the year 1982 first demonstrated the technique of using pulsed Doppler ultrasound (without imaging facility) through transtemporal approach to measure the cerebral blood flow velocities [6]. The technique they used was non-

duplex (non-imaging) or blind as there was no direct grey scale sonographic visualisation of cerebral arteries, the identity of the vessel being interrogated was determined by standard criteria that included transducer orientation, depth of the sample volume and direction of blood flow. This technique is still in use by many clinicians, however, inability to direct visualize target vessels and to perform angle correction, may have led to errors in the velocities recorded with this technique. The Power-motion mode Doppler or M-mode Doppler was described by Moehring [7] in 2002 with 33 sampling gates to detect cerebral blood flow and cerebral embolism [7]. Modern TCD involves use of spectral Doppler and colour doppler in addition to grey-scale tissue imaging that allows direct visualization of the major intra cranial arteries, allowing quick, accurate identification of arteries and their flow -velocity dynamics.

Basic Principle

TCD ultrasonography, like all Doppler imaging studies, is based on two main principles; the Doppler effect and the Bernoulli principle. Sound wave with certain frequency when strikes a moving object, the frequency of the reflected sound wave is changes. The differences in the frequency between emitted and reflected waves is called "Doppler shift frequency" that is directly proportional to the speed of the moving object (blood flow in TCD). If the flow is

towards the probe, results in shift to a higher frequency or positive Doppler shift and if the flow is away from the probe results in Doppler shift to lower frequency or negative Doppler shift. Since blood flow within a vessel is laminar, the Doppler signal obtained actually represents a spectral display of mixture of different Doppler shift from individual red blood cells [4].

The relationship between flow velocity and Doppler shift frequency is express by following formula

$$\text{Reflector speed (cm/s)} = \frac{(\text{Doppler Shift}) \times \text{propagation speed}}{2 \times \text{Incident frequency} \times \cos(\theta)}$$

The propagation speed a sound wave in a particular medium, theta θ is the angle of insonation relative to the direction of the blood vessel. The larger the angle of insonation, the larger is the cosine of the angle, and greater error in the measurement. If the angle is 0, (angle of insonation parallel to flow). We achieve the most accurate flow measurement since the cosine of 0 is 1. It is important to keep the angle below 30 degrees to keep the error below 15% (Figure 1).

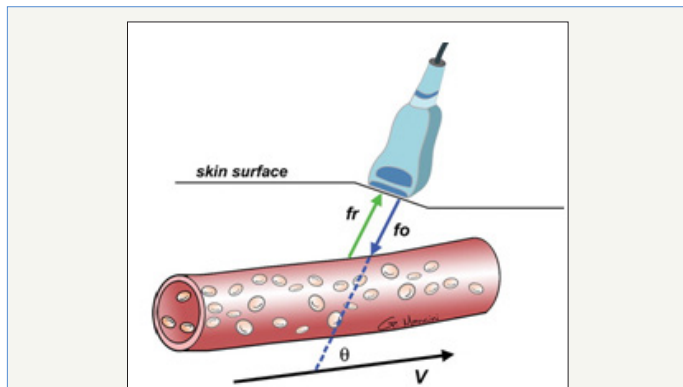


Figure 1: Diagram shows the Doppler effect. Sound waves emitted at a specific frequency (f_o) are reflected off moving red blood cells and back to the transducer at a higher or lower frequency (f_r). The difference in frequencies, known as the Doppler shift, can be used to calculate the blood flow velocity (V) and direction. θ = angle between the incident ultrasound beam and the direction of blood flow.

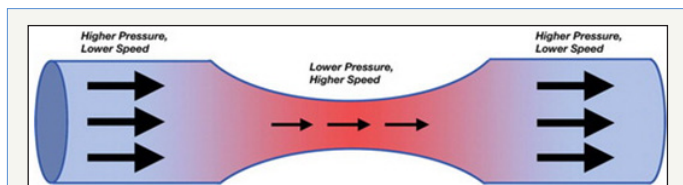


Figure 2: Diagram of the Bernoulli principle shows that as fluid flows from a conduit or vessel of greater diameter to one of lesser diameter, the velocity of flow increases and the pressure decreases to allow the same volume of fluid to pass through the narrower area.

The Bernoulli principle described the flow in terms of velocity and pressure of liquid (blood) through different size of conduit (vessels). Blood flow velocity increases and pressure decreases

when blood flow from a greater to lesser diameter (vasospasm), this is to allow the same volume of blood pass through narrow area. The changes in blood velocity (detection of vasospasm) can be estimated using Doppler Effect and Doppler equation (Figure 2).

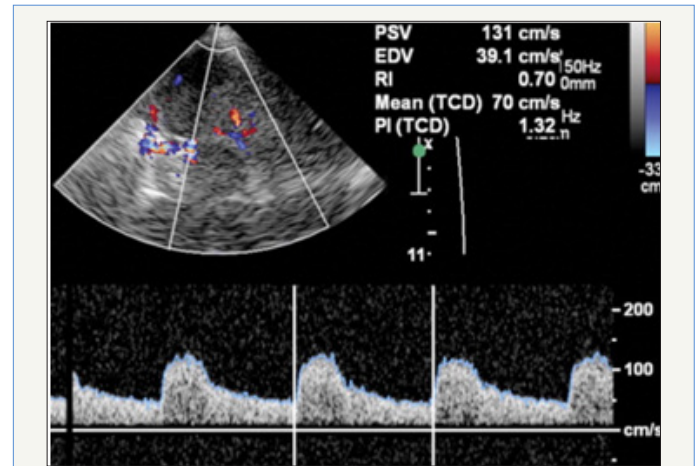


Figure 3: Duplex Doppler image and spectral waveform show the velocity of blood flowing within the PCA. The PSV, EDV, MFV, and resistive index (RI) are automatically calculated from the waveform. PI = pulsatility index, TCD = transcranial Doppler.

TCD relies on pulse wave Doppler to image a particular vessel and a received echo generates an electrical impulse which is processed electronically to produce a spectral waveform [8,9]. A spectral Doppler waveform is visual display of flow-velocities within a specific area of blood vessels as a time-velocity curve. Spectral analysis can thus be use to estimate not only the blood flow velocity, it can also used to calculate the peak systolic (PSV) and end diastolic (EDV) velocities, systolic upstroke or acceleration time, pulsatility index (PI), resistive index (RI) and time averaged mean maximum velocity (Vmean) (Figure 3). TCD devices are equipped with larger sample volume unlike the normal Pulse Wave Doppler (PWD) for other uses; this is to obtain better quality of signal in noisy background also. Standard TCD is recorded bilaterally using PWD for at least 10 cardiac cycles [10].

Image Technique

The TCD ultrasonography examination is performed by using phase array probe of 1-2MHz .The higher frequency waves have lower penetration and not able to penetrate the through the skull bone in adults and hence are not useful for intracranial scanning [8,9]. Based on anatomical location of the major cerebral arteries and ease of ultra sound penetration into brain though thinner region of skull has been identified, termed as acoustic window [8,9].

Four acoustic windows are used for TCD

Transtemporal, transformational or sub occipital, transorbital and submandibular (retromandibular). Each window specifically target particular portion of major branch of the circle of Willis, which are first optimally scan using grey scale and colour flow imaging; once a particular vessel is identified, spectral waveform are obtained using pulse wave Doppler [3,8-10]. The most

commonly used acoustic window is transtemporal window, located above the zygomatic ridge between the lateral canthus of the eye and auricular pinna, through this window all three major arteries (anterior, middle and posterior cerebral arteries) forming the circle of Willis as well as terminal portion of Internal carotid artery can be insonated (Figure 4). once probe is placed in the temporal window, identification of sphenoid bone, through the butterfly wing sign guide the visualization of the middle cerebral artery (MCA) in almost all patients which usually lies at depth of 50-65mm indicated by red colour signal as flow is towards the probe, with fine adjustment and following the flow pattern, the branching of MCA can be identified. The internal carotid artery (ICA) bifurcation observed at about 55-70mm in adults and is represented by bidirectional flow (red and blue in colour doppler), the Anterior cerebral artery (ACA) flow is away from the probe (blue) (Figure 4).

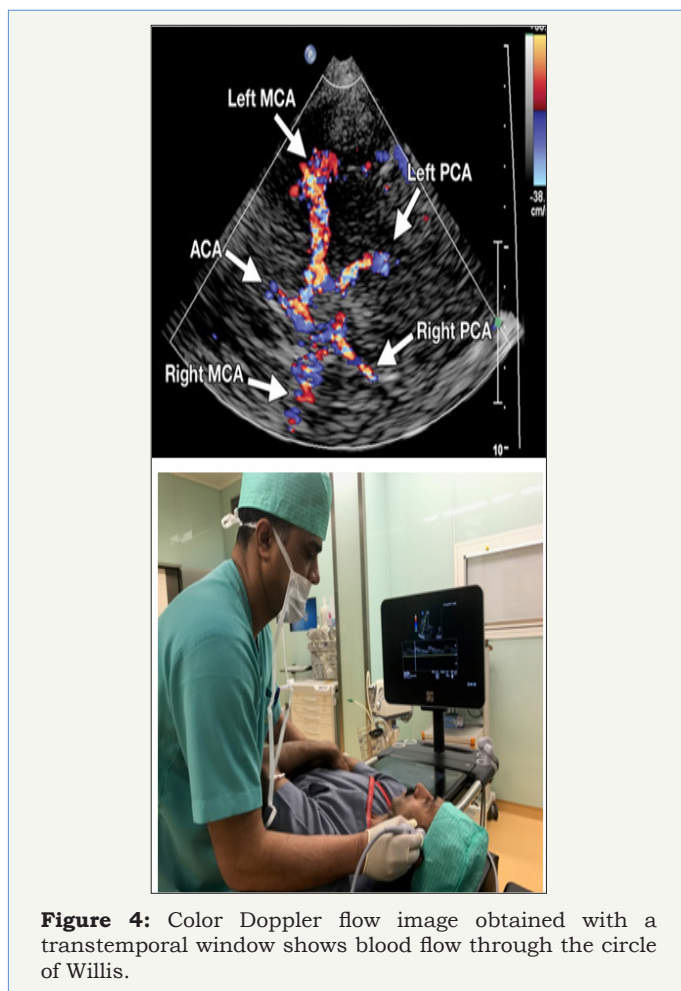


Figure 4: Color Doppler flow image obtained with a transtemporal window shows blood flow through the circle of Willis.

Once signal from MCA and ACA are obtained, the ultrasound probe is slowly oriented posteriorly by 10-30 degrees, initially there will be a flow gap before signals from the posterior cerebral artery (PCA) is obtained at 55-70mm. Two segment of PCA are obtained, P1 represented by flow towards the probe and P2 away from the probe.

Transformational or sub occipital approach involve placing the transducer in sub occipital region in a flex neck, in mid line below

the occipital, medial to the mastoid process and the probe is angled or tilted towards the bridge of the nose or contra lateral eye to insonate portion of the vertebral artery (VA) at a depth of 50-75mm (flow away from the probe) and further upward and medial tilting of probe is done to obtained from the basilar artery (BA) at a depth of 75-110mm (Figure 5).

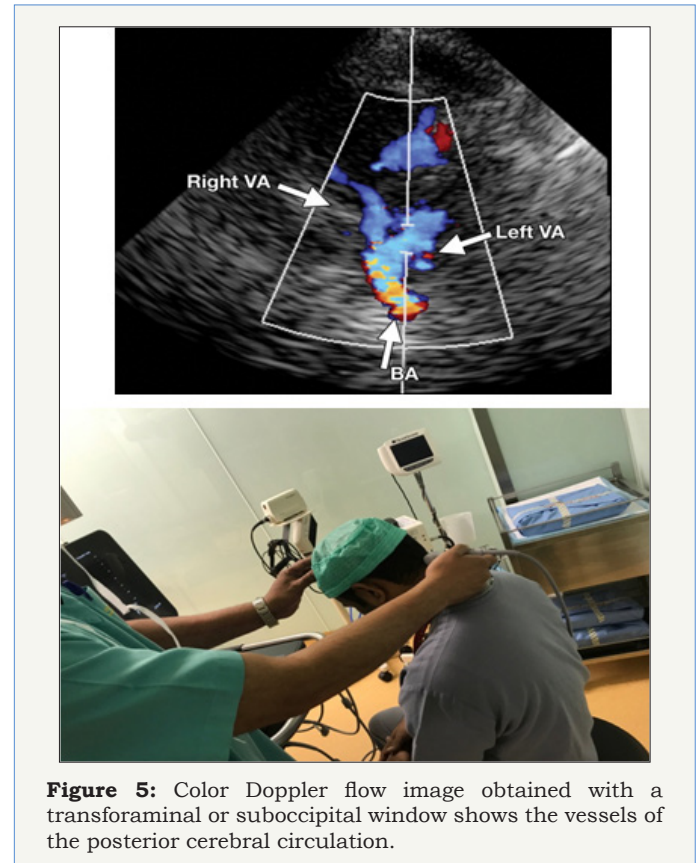


Figure 5: Color Doppler flow image obtained with a transforaminal or suboccipital window shows the vessels of the posterior cerebral circulation.



Figure 6: Submandibular and Transorbital windows.

Transorbital approach involve placing ultrasonographic probe gently on the closed eye lid to visualize the ophthalmic artery and the cavernous portion of the internal carotid artery (carotid siphon). The probe is angled slightly medially and upward, at around 60 mm ophthalmic artery can be traced followed by trace from the carotid siphon. Submandibular window is at just below the angle of jaw and can be used to locate the distal extra cranial portion of internal carotid artery in the neck (Figure 6).

Target artery is identified depending on the appropriate acoustic window, probe orientation and angle, depth of insonation and direction of flow, in difficult scenario when it is not possible to differentiate the anterior from posterior artery, dynamic maneuvers such as direct carotid compression or vibration and resulting waveform changes are recorded [3,9].

TCD Indices

Peak Systolic Velocity (PSV) -The maximum value of flow velocity in systole, at the apex of the waveform.

End Systolic Velocity (EDV)-The velocity measured at end diastole, usually at the lowest point before a new waveform begins. The central parameter in TCD is the mean flow velocity (MFV) which can be calculated similar to that of mean arterial pressure (MAP), the MFV is equal to 1/3rd PSV + 2/3rd EDV. (MFV= {PSV+EDV x 2}/3). The MFV can be influenced by number of physiological factors [11-13] (Table 1).

Table 1: Physiological Factors influencing MFV.

Physiologic Variable	Change in MFV
Age	Increases up to 6-10 years of age then decreases
Sex	Higher MFV in women than men
Pregnancy	Decreased in the 3rd trimester
PCO2	Increases with increasing PCO2
Mean arterial Pressure (MAP)	Increases with increasing MAP(CBF autoregulates between CPP 50-150mmHg)
Haematocrit	Increases with decreasing haematocrit

Common pathological factors increasing MFV are vasospasm, stenosis, hyper dynamic flow, whereas, hypotension, decrease in cerebral blood flow, intracranial pressure or brain stem death [11]

other important TCD indices are:

Gosling’s pulsatility index (PI)

This is the most commonly used TCD parameter to determine the flow resistance. PI is the measure of pulsatility of blood flow that reflects the resistance to blood flow. It is the difference between systolid and diastolic flow velocity divided by mean velocity. It is calculated as $PI=(PSV-EDV)/MFV$. Normal value-0.5-1.19. Downstream obstruction will increase the PI and proximal obstruction or stenosis may decrease the PI due to down stream vasodilatation [14,15]. Also it is affected by change in Intra Cranial Pressure (ICP) in a linear way and a progressive rise in PI

may reflect rise in ICP [16,17]. Patients with Arterio-Venous (A-V) malformation will have reduced PI as the resistance to flow suddenly reduced due to distal venous flow [18].

Pourcelot resistivity index (RI)

It is similar to pulsatility index. It measures the resistance to blood flow distal to the site of measurement. It is calculated as $RI= (PSV-EDV)/PSV$; value greater than 0.8 indicates increased in downstream resistance [11].

Lindegaard ratio (LR)

Its a ration of mean flow velocity in intracranial to extra cranial carotid artery and calculated as MFV in Middle cerebral artery/MFV in extra cranial internal carotid artery [19]. The LR differentiate between hyper dynamic flow from vasospasm. LR more than 3 indicates cerebral vasospasm. A high LR ratio indicates increased in intracranial flow due to vasospasm and abnormally low LR indicates hyper dynamic flows like in A-V malformation [20]. A modified LR ratio for intracranial bsilar and extra cranial vertebral artery can also be cam also be used to calculate vasospasm in posterior circulation [21].

Micro embolic signal (MES) detection

TCD a useful monitor intraoperatively for carotid and other cardiac surgery where there is high risk of micro embolism [22,23].

Abnormal spectrum waveforms

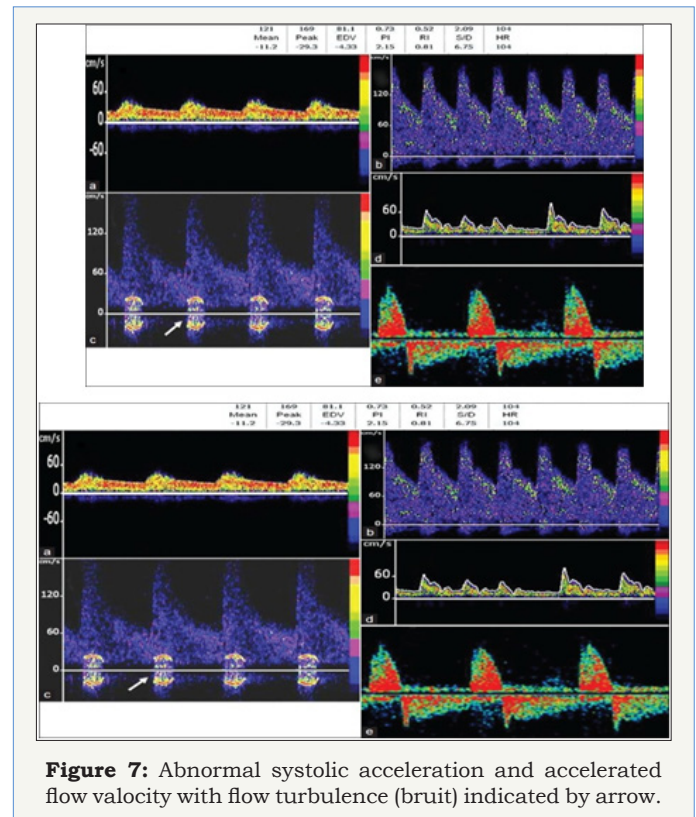


Figure 7: Abnormal systolic acceleration and accelerated flow velocity with flow turbulence (bruit) indicated by arrow.

Delayed systolic flow acceleration or flattened systolic upstroke and slow diastolic deceleration with end diastolic velocity (EDV) above 50% of peak systolic velocity (PSV) is hallmark of stenosis

proximal to the site of PWD this is known as blunted flow, and is due to compensatory vasodilatation distal to the stenosis (Figure 7).

Patients with focal intracerebral stenosis/ vasospasm- there will be focal flow acceleration, with increasing stenosis; the increased in the flow creates flow turbulence, represented by bruit, which is seen as symmetrical artefact on either side of baseline.

Table 2: Common clinical applications of TCD.

Cerebro-Vascular Ischemic Diseases	Perioperative/ Periprocedural Monitoring	Neuromonitoring In ICU	Others
Sickle cell disease	Cerebral thrombolysis in acute stroke	Vasospasm after subarachnoid haemorrhage	Pharmacologic vasomotor testing
Right to left cardiac shunts	Carotid endarterectomy	Raised intracranial pressure	Cerebral pressure auto regulation
Intra and extra-cranial arterial steno-occlusive disease	Carotid angioplasty and stenting	Head injury	Liver failure/Hepatic encephalopathy
Arteriovenous malformations and fistulas	cardiac surgeries to detect micro emboli.	Cerebral circulatory arrest and brain death	Preeclampsia
		Intra cerebral aneurysm and parenchymal hematoma detection	

Monitoring of vasospasm in subarachnoid haemorrhage (SAH)

TCD was initially invented for monitoring of cerebral vasospasm after SAH [6,7]. The basis of using TCD in vasospasm is on Bournellis principle, the narrowing of vessel will lead to increase in velocity of blood flow, even though absolute flow decreased. TCD is a reliable, cost effective tool for monitoring of vasospasm after SAH delayed cerebral vasospasm is one of the common complication after SAH, can occur in upto 70% between 4-17 days, which can leads to delayed ischemic deficit in up to 25% patients [24-26]. There are reports of early vasospasm within 48 hours in up to 13% cases, [27-32] and very delayed vasospasm even after 20 days, morbidity and mortality are considered to increased significantly, longer the period of vasospasm [33]. Use of TCD can be used to early detection of even asymptomatic vasospasm, follow progression of vasospasm and can be used to monitor the effect of treatment or intervention for vasospasm [34-39].

Suggested protocol for TCD in SAH by Sharma et al [38]:

First baseline TCD as soon as possible after the diagnosis of SAH. Flow velocity to be obtained from both extra cranial and intracranial artery to calculate the LR index.

Daily or more frequently if needed evaluation by TCD for early detection of vasospasm.

Standard diagnostic criteria to be used for diagnosis of Vasospasm (Table 2).

If TCD suggested of vasospasm, to initiate treatment for vasospasm and TCD re-evaluation every 6-8 hours.

Information regarding ICP if available should be taken into account during interpretation of TCD.

Clinical Application of TCD

TCD was initially used to diagnose vasospasm following sub-arachnoid haemorrhage. However, a clinical use of TCD has expanded greatly over the decades and it has emerged as bedside, non-invasive tool for evaluating cerebral artery patency, detection of stenosis, embolism, flow pattern and even as alternative monitoring of ICP. Common clinical applications of TCD can be summarised as categorical groups (Table 2).

In doubtful finding, other diagnostic modality (imaging) should be employed.

TCD in comparison to angiography has good sensitivity and specificity in diagnosis vasospasm for both MCA and VA. [35,40,41] Mean flow velocity (MFV) of MCA greater than 120 cm/sec has 99% specificity and 67% sensitivity to identify angiographic vasospasm of 25% or more [41]. The positive predictive value (PPV) increases to 98% when MFV exceed more than 200 cm/sec. Whereas, when the MFV less than 12 cm/sec has 94% negative predictive value (NPV) [42]. Therefore, MFV of more than 200 cm/s or less than 120 cm/sec could predict both present and absence of MCA vasospasm, respectively. Unlike the sensitivity and specificity of even in mild vasospasm of MCA, TCD is little weaker in detecting less than 50% narrowing in VA and BA [35,40,43,44].

Similarly, ACA and PCA vasospasm has lower specificity and sensitivity in mild to moderate vasospasm unlike the MCA vasospasm [45]. It is daily changes in MFV that is more important than the absolute value; an increase in MFV is highly predictive of vasospasm. MFV rise of 50cm/sec or more within 24 hours or MFV increases of more than 65cm/s per day from day 3-7 indicate development of vasospasm and high risk for delayed cerebral ischemia [46,47].

Though the TCD is a valid diagnostic tool to identify and to monitor the development of intra cerebral vasospasm after SAH. The interpretation of TCD finding should be correlated with clinical assessment and other diagnostic tools like computed angiography should be used for diagnosis of vasospasm after SAH instead of the single independent test [47,48].

Acute ischemic/embolic stroke

TCD can accurately identify acute occlusions of intracranial arteries with high, specificity, sensitivity, PPV, and NPV of greater

than 90% for MCA and 70-90% for ICA siphon, VA and BA [49]. TCD has been compared with Magnetic resonance angiography (MRA) and Computed tomography angiography (CTA) in early evaluation in acute ischemic strokes. It has been used to assess steno-occlusive pathology of terminal ICA, ICA siphon and MCA. TCD is almost 100% specific and 93% sensitive for identifying the MCA ischemia, whereas MRA had 74% specificity and only 46% sensitivity in the assessment of intracerebral arteries. Moreover, TCD can be used in emergency department as bed side tool and provide real time information about cerebral blood flow (CBF) [50-52].

In addition to diagnostic tool, TCD has more prognostic value in strokes. A complete arterial occlusion at presentation predicted worse neurologic outcome, disability and death after 90 days, [53,54] a normal TCD finding predicted early neurological improvement [47,55]. Unlike other imaging modality, the advantages of TCD is that it can be used serially to detect any cerebral hemodynamic changes that would otherwise go undetected by a single MRA, CTA studies [56].

TCD is also useful to assess the hemodynamic changes before and after thrombolysis; recanalisation within 6 hours of symptoms onset or within 30 min of thrombolysis is associated with marked clinical improvement and functional independence at 3 month [57,58]. TCD can also be useful tool to detect early reocclusion after thrombolysis.

TCD in Sickle-Cell Disease

Patients with sickle cell disease (SCD) (homozygous) carry a significant risk of cerebral strokes, 11% all SCD children developed strokes before the age of 20 years and majority of these strokes involve occlusion or stenosis of the distal portion of ICA or proximal portion of MCA [38]. Pathophysiologies involve increasing circulation of irreversibly sickled cells and their adherence to the vascular endothelium which leads to vaso-occlusion and ischemia [59].

Table 3: STOP classification for risk stratification and treatment strategy [64].

Group	TAMM Velocity	Action
Normal	<170cm/s	A repeat assessment is indicated.
Conditional	170-200cm/s in the MCA and/or terminal ICA	In children with no previous records, a repeat TCD is planned in about 2 weeks
Abnormal	>200cm/s in the MCA and/or terminal ICA	Urgent blood transfusion is arranged.

TCD is a useful tool in assessment of children with SCD to identify the risk of stroke, need of blood transfusion. Elevated blood flows in SCD are related to anemia, cerebral vasodilatation caused by tissue hypoxia or vessel stenosis. Blood transfusion correct some of these abnormalities to an extent and reduces cerebral blood flow velocity and thereby the risk of stroke [60,61]. Detection of time average MFV exceeding 200cm/s on two separate occasion is

the main risk factor for the first-ever stroke [62,63]. This can be preventable up to 90% by timely transfusion blood [63]. Stroke prevention in sickle cell disease (STOP) trial has developed a protocol for TCD screening in children with SCD. The STOP protocol is different than routine clinical TCD examination, where they uses time-average mean of the maximum (TAMM) velocities of ICA and or MCA recorded on two separate occasions at least 2 weeks apart [64]. The detail description of STOP protocol is beyond the scope of this chapter (Table 3).

Traumatic brain injury (TBI) and raised ICP

TBI have spectra of effects on brain and cerebral circulation, immediately after trauma, there may be hypo perfusion (day 0) followed by hyperaemia (day1-3), there may be delayed vasospasm and raised ICP after the initial phase is over [65]. TCD as a non-invasive tool can be used to early detection of cerebral hemodynamic changes, it also provide long term prognostic information. TCD derived pulsatility index (PI) provides useful information about ICP and CPP, avoiding complications of invasive monitoring [16,17,66] Recently Wakerley et al. [67], conducted a study to define the relationship of PI and ICP, they performed TCD measurement and simultaneously Lumbar puncture for ICP measurement. TCD-PI has acceptable accuracy in differentiating patients with ICP more than 20cm of cerebrospinal fluid (CSF). Serial measurement over short period of time are reliable than single measurement [67]. There is no standardisation technique or formula for absolutely quantifying the ICP/ CPP by TCD method, hence at present, TCD is reserved for assessing serial change, rather than absolute value.

TCD monitoring for spontaneous emboli

Emboli from carotid plaque or from cardiac origin (left atrial thrombus or right to Left shunt) can produce cerebral ischemia by blocking regional perfusion. Extended monitoring of intracranial artery distal to embolic origin site can detect spontaneous micro-emboli called Micro-embolic signals (MES).

The International Cerebral Hemodynamic Society described MES as [38]

Random occurrence during the cardiac cycle.

Brief duration (usually <0.1 s).

High intensity (>3 dB over background).

Primarily unidirectional signal (if fast Fourier transform is used).

Audible component (chirp, whistle, or pop).

Patients with history of cerebrovascular ischemia or high grade carotid stenosis can also undergo TCD monitoring for MES [68-70]. MES can also be used to assess the treatment effectiveness after initiating antiplatelet drug therapy in carotid stenosis [71,72].

Right to left (R-L) shunt detection

Patent foramen ovale can be present in approximately 25% of general population and these patients are at risk of paradoxical embolism [73]. TCD bubble test can be done to detect R-L shunt

in patients with cerebral ischemia due to suspected paradoxical embolism. TCD is considered to be superior to transesophageal echocardiography as it has not only higher sensitivity and specificity for detection of R-L shunt, it can quantify its functional -potential for embolism [74,75].

TCD bubble test involve injection of small quantity of micro bubbles, produced by agitation with saline and patient own blood through antecubital vein and simultaneous TCD scanning for detection of MES in one MCA. The test is repeated with change in body posture.

The R-L shunt can be quantified by international consensus criteria [76]

Grade 0: No MES.

Grade 1: MES count 1-10.

Grade 2: MES count 11-30.

Grade 3: MES count 31-100.

Grade 4: MES count 101-300.

Grade 5: MES count more than 300.

TCD in brain death: Brain stem dysfunction and death is a clinical diagnosis based on clinical evaluation and various provocative test and electroencephalography. However, the diagnosis of brain stem death may be delayed considerably in condition like use of sedative and muscle relaxant, metabolic derangement, hypothermia. TCD may be alternative confirmatory test in such situation. This phenomenon could have serious implications in cases where organ preservation is needed in preparing for possible organ transplant [77-80]. TCD per cannot be used on its own to diagnose brain death since this is a clinical diagnosis. TCD can be used to help detecting the cerebral circulatory changes. A prolonged (more than 30 min) absence of cerebral perfusion or cerebral circulatory arrest is detected using one of the following criteria [81].

Reverberating (oscillating) flow pattern- an oscillating waveform- with reversal of equal flow pattern in systole and diastole- zero net flow.

Small systolic spike less than 200ms duration of less than 50cm/sec peak systolic velocity with absent diastolic flow.

Absence flow signals in intracranial circulation with typical flow signal in extra cranial circulation.

Cerebral auto-regulation or vasomotor reactivity

Cerebral auto regulation refers to the maintenance of cerebral blood flow (CBF) despite changes in cerebral perfusion pressure over the range of 50-150mm/hg [82]. Impairment in auto regulatory response has been demonstrated in varieties of brain insult such as strokes, traumatic brain injury, menigo-encephalitis, carotid disease etc. Impaired cerebral auto regulation or vasomotor response has prognostic value. TCD can be used to assess the various aspects of CBF regulation in response to change in blood pressure, circulating level of vasomotor in a dynamic way [82-84]. A pressure reactivity

index Prx and CBF related index has been introduced to assess the cerebral auto regulation [85] but none of the index has been considered as gold standard [86]. Changes in MFV in MCA is used as surrogate for changes in CBF. Many different non-pharmacologic techniques such as direct carotid artery compression, Valsalva manoeuvre, application of PEEP, head up tilting, negative pressure to lower body portion etc were used to induced pressure changes [87-90].

The TCD assessment of cerebral auto regulation is still an area of clinical research given the high temporal resolution, convenience and non-invasiveness of the technique. Significant auto regulatory impairment in TBI and strokes are common and are associated with poor neurologic outcome. However, the role of auto regulatory assessment in carotid stenosis, syncope or extra cranial pathology is less clear.

Conclusion

TCD being non-invasive, portability and provide real time information and high temporal resolution have promoted its use as bedside neuromonitoring tool. Majority of its use are early detection of impaired cerebral flow dynamic and it helped in initiation of preventive measures in SAH, SCD, TBI and Strokes or even preparation of organ retrieval in impending brain death. Advanced application of TCD includes evaluating various mechanism of cerebral ischemia. TCD helps in planning and monitoring the cerebral hemodynamic, effectiveness of therapy as well as aids in establishing the prognosis. The main limitation of TCD is high operator dependency and it require some technical expertise and training to standardise its application.

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