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Review Article

Role of anaesthetist in antepartum haemorrhage- A complete review

Suchitra Malhotra¹, Reena Mahajan², Kiran Bhatia³, Mohinder Kumar⁴, Prachi Renjhen⁵ ^{1,2,3}Department of Anaesthesia ,SHKM Govt Medical College, Nalhar, Mewat ⁴Department of Surgery, MMIMSR Mullana, Ambala ⁵Department of Obstetrics and Gynaecology, SHKM Govt Medical College Nalhar, Mewat

*Corresponding author

Suchitra Malhotra Email: <u>suchitramalhotra@yahoo.co.in</u>

Abstract: Maternal mortality is a matter of great concern in developed and developing nations. More than 25% of maternal deaths can be attributed to haemorrhage. Availability, of trained birth attendant and emergency medical services are the main objectives for achieving the Millennium Development Goal 2015 of reducing maternal mortality rate to 109/100,000 from the present 190. One of the major shortcomings for providing high quality emergency obstetric care is a serious shortage of specialists such as obstetricians and anaesthetists at various levels in rural setting. Anaesthetists, perhaps the only perioperative physicians in labour room and operating room with special training in resuscitation and critical care get involved early in management of these bleeding patients. The differences in management options between antepartum haemorrhage (APH) and postpartum haemorrhage (PPH) are that there are two individuals to care for in APH and that delivery of the fetus and placenta will help to arrest the bleeding. The greatest threat in APH is not to the mother but her fetus. Thus a multidisciplinary approach and team effort involving the obstetrician, anaesthetist, haematologist neonatologist, midwifery and paramedic operation theatre staff can go a long way in averting impending death and disability. In this guideline we shall discuss the latest trends in management of APH so that mothers and babies lives and not in jeopardy.

Keywords: placenta praevia; accreta; abruption;. Uterine rupture; anaesthetic management.

INTRODUCTION

Maternal mortality is a matter of great concern in developed and developing nations[1]. More than 25% of maternal deaths can be attributed to haemorrhage[2.3]. Availability, of trained birth attendant and emergency medical services are the main objectives for achieving the Millennium Development Goal 2015 of reducing maternal mortality rate to 109/100,000 from the present 190[4,5].

Antepartum haemorrhage (APH) is bleeding from the genital tract after 24 weeks gestation up to and including delivery of the fetus [6,7] Incidence of APH is 2-5% of all pregnancies [7,10] Main causes are

- 1. Placenta praevia
- 2. Placenta abruption
- 3. Uterine rupture
- 4. Vasa praevia

Blood loss>1000ml and/or presence of shock

Major complications of APH are perinatal mortality [8] preterm labour and PPH [1,2,,4,10,] Current data suggests that placenta praevia and placental abruption are responsible for perinatal mortality of 2.3% and12% respectively 1 [1,12,13].

Table-1: Definition	
MINOR HAEMORRHAGE	Blood loss<50ml
MAJOR HAEMORRHAGE	Blood loss 50-1000ml ,no shock

A recumbent pregnant woman can maintain a normal pulse and blood pressure until she has lost one

MASSIVE HAEMORRHAGE

third of her blood volume due to relative hemodilution and high cardiac output. Therefore haemodynamic status may not correspond with apparent blood loss in obstetric haemorrhage

As visual blood loss estimation does not accurately judge the blood loss[14], other reliable methods such as blood collection drapes for vaginal deliveries and weighing swabs ,floor spills and suction bottles are preferred. A laparotomy sponge if 50% soaked with blood gives an estimate of 25ml blood loss and ,in case it is dribbling with blood the estimated loss is 100ml.[15]. Pictorial guides based on experience at Queen Charlotte's Hospital, London.is a useful aid for staff working in labour room to measure the bloodloss[16,17].Multidisciplinary observations of estimated blood loss revealed that these were grossly miscalculated by 30-50%[14].

Accurate estimation of blood loss along with measurement of vital signs can be used to classify haemorrhagic shock into 4 different stages. Royal College of Obstetricians and Gynaecologists. RCOG recommends MEOWS(maternal obstetric early warning score) based on above clinical parameters for early identification of the shock[10].

Tuble 2: Clubbilleution of Dirock				
Classification of shock	STAGE 1	STAGE 2	STAGE 3	STAGE 4
BLOOD LOSS	<15%	15-30%	30-40%	>40%
HEART RATE	<100/min	>100/min	>120/min	>140/min
SYSTOLIC BLOOD	normal	normal	decreased	decreased
PRESSURE				
RESPIRATORY RATE	14-20/min	20-30/min	30-40/min	>40/min
MENTAL STATUS	Slightly anxious	Mildly anxious	Anxious, confused	Confused
				lethargic

Table 2: Classification of Shock

Successful resuscitation of bleeding obstetric patients requires a TEAM(together everybody achieves maximum) approach. A multidisciplinary effort, including obstetricians, nurses, anesthesiologists, haematologists neonatologist paramedical staff should be organized early in the course of obstetric hemorrhage, involving other services as necessary in order to achieve good outcome.

Important signs and symptoms of massive obstetric haemorrhage are: [19]

- Rising pulse rate,
- Pallor,
- Fall in blood pressure,
- Altered conscious level
- Rising respiratory rate,
- Decreased urine output and fetal demise. In APH signs of fetal jeopardy may occur much before the onset of maternal shock

Role of Anaesthetist [20]

- Resuscitation of the bleeding patient-- ABC,s
- Estimation of blood loss•
- Fluid & blood product replacement
- Anaesthesia Induction a & maintenance
- Drug administration
- Sending lab investigations and bedside tests viz haemecue, arterial blood gases, FIBTEM A5 assay
- Monitoring including invasive e.g CVP,ARTERIAL LINE

Delivery room staff are trained Major Obstetric Haemorrhage protocol after 1000 ml blood loss. This

ensures multidisciplinary involvement early in the management of haemorrhage. This will allow obstetrician to search for and control the source of bleeding while the anaesthetist manages hemodynamic and pharmacologic aspect of the patient, blood and fluid replacement, laboratory testing, and ordering and procuring blood products instead of one person (usually the obstetrician) trying to deal with all these issues at the same time. Arterial access with a radial arterial line should be strongly considered early in any haemorrhage that appears to be significant or on-going. This allows repeated sampling to assess haemoglobin and platelet concentrations, coagulation function, arterial blood gases , pH, and ionized calcium levels during on-going rapid transfusion.

CVP monitoring preferrably ultrasound guided by senior anaesthetist early during resuscitation reduces incidence of complications. Central line, is useful for both high-volume infusion and for administration of vasoactive drugs, for example, epinephrine [21].

In cases of major or massive haemorrhage, four units of blood should be crossmatched and a series of lab investigations should be sent for immediately Thee include blood count and coagulation screen . Urea, electrolytes and liver function tests .Latest development is the near patient testing like Haemcue and FIBTEM A5 ASSAY.

Placenta praevia

Placenta is implanted partially or wholly in lower uterine segment. It can cause intractable haemorrhage. Trans vaginal ultrasound (TVS) has redefined the degrees of placenta praevia – Oppenheimer[23] classified placenta praevia according to relationship of placental edge to internal os and risk

of APH and need for caesarean delivery.

DISTANCE OF PLACENTAL EDGE FROM INT OS	RISK
0-10mm	Caesarean section(CS) and high risk of bleeding
11-20mm	Low risk of CS and bleeding
>20mm	CS not indicated

 Table 3: OPPENHEIMER,S CLASSIFICATION OF PLACENTA PRAEVIA

Advances in TVS have made DOUBLE SET UP examination (i.e. vaginal exam with all personnel ready for immediate caesarean section) nearly obsolete in modern obstetric practice. However it may be needed in morbidly obese patient. TVS probe should not be inserted>3cm into the vagina otherwise it may precipitate bleeding.

Anaesthetic management.

Active labour, persistent bleeding, a mature fetus or non-reassuring fetal status necessitate prompt delivery by caesarean section

- In the pre-operative visit the anaesthesiologist should assess the airway, blood loss, vital signs, senior help should be available, consent for blood transfusion, post-operative ICU care should be obtained as well as possibility of hysterectomy
- While shifting to operation theatre ., the patient should be given oxygen and placed in left lateral position to reduce aorto-caval compression .
- Blood 4 units should be ordered in the operation theatre
- 2 wide bore cannulae should. be inserted
- Invasive monitoring is useful but it should not delay resuscitation.
- All fluid that is administered during resuscitation should be warmed preferrably Rapid infusion devices must be used when available.
- Nowadays neuraxial anaesthesia is preferred in patients without active bleeding, coagulopathy or intravascular volume deficit[24]
- The advantages of regional technique are that mother and baby are awake ,the relative problems of general anaesthesia (GA) like tachycardia are avoided , pain relief is better and less number of transfusions are needed
- On the contrary as surgeon has to proceed quickly, the motor blockade must be effective, if haemorrhage is severe compensatory mechanisms being blocked hypotension is more difficult to control. At this juncture many times GA may be required .This is a difficult situation to handle for the patient, obstetrician,

as well as the anaesthesiologist intubating the patient.

- Normally these patients remain at risk of increased blood loss as the obstetrician may injure anteriorly located placenta during uterine incision. Secondly after delivery lower uterine segment does not contract well, thirdly patients with placenta praevia are at increased risk of placenta accreta especially with previous caesarean delivery.
- Most anesthesiologists prefer providing regional anesthesia for caesarean delivery, others prefer G A under all circumstances[25,26]
- It is more practical to prefer GA in cases of difficult intubation and previous caesarean with present anterior placenta praevia
- Oyelese and Smullian stated that orientation of placenta to the anterior uterine wall and patient status are factors that should be considered when choosing the anaesthetic for parturient with placenta previa [27]

Placenta praevia with accreta

When an anteriorly located placenta praevia presents in a mother who has a uterine scar the possibility of placenta accreta should be considered. Incidence of placenta accreta is rising because of increased caesarean delivery rate. The risk increases from 24% with one previous caesarean section to 67% in women with 3 or more prior caesareans[28]. Placenta accreta is noted at the time of delivery or CS with difficulty in separating the placenta from uterine wall . MRI may be useful in diagnosis. Unfortunately, many times placenta accreta occurs in unanticipated patients who are not prepared for hysterectomy.

- When placenta accreta is suspected traditional management is caesarean hysterectomy [27,29] with availability of blood and blood components;
- Conservative treatment to preserve fertility is caesarean delivery and leaving the placenta in situ[30,31] This is accompanied by bilateral embolization of uterine arteries and parenteral methotrexate..
- Hospitals without adequate blood bank supplies should consider transferring patients with placenta accreta to a tertiary care facility where 24 hour in house obstetric consultant, anaesthesia consultant,

gynaecologist, fully stocked blood bank, oncologist, urologist ,and radiologist as well as facility for cell salvage are available

- Estimated blood loss is>2000ml in 66% cases.
- Interventional radiologist inserts balloon occlusive devices in both internal iliac arteries prior to surgery. Normally balloons are inflated after delivery so as to provide a less bloody surgical field.
- Fetal bradycardia has been observed so it should ideally be placed in operation theatre to allow rapid delivery and avoid dislodgement
- Intraoperative possibility of infection exists while patient is being shifted from operating room to radiology department. The transfer of parturient can be challenging for paramedical staff and anaesthetist monitoring the bleeding patient
- The advantages of balloon occlusion of internal iliac arteries are that: it is a minimally invasive technique, it involves very less fluoroscopy time, and it does not jeopardize fetal blood supply as the inflation of balloon is done after clamping of umbilical cord.
- If foetal distress develops in the interventional radiology suite, then it would be mandatory to carry out the caesarean section there itself. Such a situation is fraught with problems. General anaesthesia is the technique of choice in these patients. It is better to defer neuraxial block because of chances of hypotension and coagulopathy due to massive blood loss. Besides, these patients have to be anti-coagulated with heparin to prevent thrombosis of the intra-vascular balloon and the sheath.

Royal college of obstetricians and gynaecologists(RCOG) and the Royal college of medicine (RCM) set up an expert working group to develop a care bundle for placenta praevia accreta. called placenta praevia care bundle(PPCB) in February 2010[22]. There are six elements of this bundle .viz

- Consultant obstetrician planned and directly supervising delivery
- Consultant anaesthetist planned and directly supervising anaesthetic at delivery
- Blood and blood products available

Multidisciplinary involvement in pre-op planning
Discussion and consent includes possible interventions (such as hysterectomy, leaving the placenta in place, cell salvage and intervention radiology)

• Local availability of a level 2 critical care bed.

It is vital to normalize haemoglobin and coagulation status and identify the placental site by sonography before surgery. Active management of the third stage of labour is recommended, as this is associated with lower bloodloss. The Obstetric and Anaesthetic consultant should be informed on admission of patient.

Anaesthetic management

- Anesthesia management of the hemorrhage consists of blood and fluid replacement, guided by assessment of the amount of blood loss along with heart rate, systemic blood pressure, central venous pressure and urine output[35].
- Most patients who present for routine planned cesarean sections are anesthetized using neuraxial techniques[32]. Use of neuraxial anesthesia has the added benefit that newborn will be awake.even though incision delivery interval will be prolonged which could be harmful for the baby if GA was the anaesthetic technique In a classic study of cesarean hysterectomy at five institutions from the 1980s. 32% of planned cesarean hysterectomies were performed under regional anesthesia. There was no difference in intraoperative blood loss or hypotension, and no one required induction of general anesthesia[33]
- Gather anesthesia team: It is advisable to have more than one anaesthetist present except in case of emergency.
- Establish at minumum two large-bore (14-16 G) peripheral IVs
- Place an arterial line and central line,
- Rapid infusion and , warming devices
- Procure blood and blood products
- Have "Point-of-Care" electrolyte, haemoglobin, and blood gas analysis device, rotational thromboelastometry ROTEM
- Have vasopressors (epinephrine, norepinephrine, Dopamine, dobutamine) immediately available
- Have all available uterotonics
- Order 10 Upacked red cells PRBCs, 10 Ufresh frozen plasma FFP, 10 U platelets to start, with 10 more each available in the blood bank (double the number if known percreta)[20].

A combined spinal epidural (CSE), regional anaesthesia is technique of choice in elective cases .with moderate risk However, general anaesthesia is warranted if the airway appears difficult, there is significant thrombocytopenia , pre-existing coagulopathy, or if the patient refuses regional anaesthesia. Airway alterations during a caesarean hysterectomy under regional anaesthesia that can make intubation more difficult have been reported, presumably secondary to fluid resuscitation and transfusion[35]. Transfusion Therapy, Research with resuscitation in trauma using massive transfusion has shown an improved outcome by using a 1:1:1 ratio of packed red blood cells, fresh frozen plasma, and platelets[34].Red blood cell transfusion is required when 30–40% of the blood volume is lost

The common goals for transfusion in the obstetric patient is to achieve

- Haemoglobin >8 g/dl, with PRBC,s
- Platelet count $>75 \times 109/l$, with platelets
- Prothrombin time (PT) <1.5 \times mean control ,with FFP
- Fibrinogen >1.0 g/l, with cryoprecipitate or fibrinogen concentrate
- FIBTEM A5 >12with cryoprecipitate or fibrinogen concentrate

Whole blood, a cheaper alternative replaces many coagulation factors, and its plasma expands blood volume. Platelet therapy (6 -10U at a time) is usually considered as the total packed red blood cells(PRBC) units used approaches 10 U,.

. Cell salvage may be particularly useful in cases where homologous blood use is not possible, such as for Jehovah's Witnesses. Its use in obstetrics has been restricted, mainly due to concerns related to reinfusion of fetal red blood cells and amniotic fluid, and the difficulty of predicting which patients will require transfusion.

Recombinant factor VIIa has been considered a last resort when conventional treatment has failed before proceeding for hysterectomy[36]. For treatment of severe postpartum hemorrhage, it is important to avoid hypothermia and hypocalcemia to maintain normal hemostasis[34]. Tranexamic acid could be considered for prophylaxis after cord clamp Dose is 1 g intravenously, followed by another 1 g if additional bleeding occurs[37].

ABRUPTIO PLACENTA is defined as premature separation of a normally situated placenta. Incidence is 0.6-1% of all births. Maternal haemorrhage may be revealed (80%) or concealed (20%).Fetal compromise occurs because of loss of placental surface area for feto- maternal exchange of oxygen and nutrients.

Table 5. COWILLICATIONS of abruption [30, 57].		
MATERNAL	FETAL	
Disseminated intravascular coagulation	Prematurity	
Haemorrhagic shock	Intrauterine growth restriction	
Death	Intrauterine fetal death	

Table 5: COMPLICATIONS of abruption [38, 39].

- The classical presentation is vaginal bleeding ,uterine tenderness and increased uterine activity[40].
- The diagnosis of abruption is primarily clinical. USG is highly specific for abruption(96%), but is not very sensitive.(24%)[39,44,45].
- Blood loss if often underestimated and the amount visible may only be a portion of the total volume of the haemorrhage as the haemorrhage can be concealed, therefore clinicians should also observe for signs of maternal clinical shock and fetal compromise or demise
- One third of coagulopathies in pregnancy are attributable to abruption and coagulopathy is associated with fetal demise[43].
- In abruption placenta the extrinsic pathway triggered by tissue destruction is activated. Thromboplastin is liberated from the placenta leading to consumption of clotting factors and platelets Fibrinolytic system gets activated in response to fibrin deposition. Plasminogen in turn converts to plasmin causing lysis forming fibrinogen, fibrin monomer, fibrin polymer and fibrinogen degradation products-FDP. These in turn cause bleeding.

Plasma fibrinogen levels are diagnostic in severe haemorrhage. Rapid detection and treatment of hypofibrinogenaemia is essential to stabilize the situation. The use of near-patient testing of coagulation using ROTEM allows monitoring of trends of coagulopathy[46,47].

Anaesthetic management

- If mother develops hemodynamic instability or coagulopathy or fetal status becomes nonreassuring urgent Category 1 caeserean section may be done under general anaesthesia[40,41].
- 2 intravenous lines(16G) are established
- Blood is analysed for haemoglobin estimation platelet count, blood grouping and crossmatching (4 or more units) and clotting profile (including a measure of fibrinolysis, such as Fibrin Degradation Products [FDPs] or Ddimer)fibrinogen level ,prothrombin time(PT)/Activated partial thromboplastin time((aPTT),arterial blood gases(ABG), base line electroloytes, creatinine, and ROTEM
- If any evidence of Disseminated Intravascular Coagulopathy (DIC) and/or fibrinolysis is found, fresh frozen plasma platelet

packs,,cryoprecipitate, fibrinogen concentrate, tranexaemic acid,recombinant factor seven a should be ordered as well.

- Regularly monitor volume loss, Consider CVP/arterial line, Non-invasive BP, Pulse oximetry
- Renal function: monitor urine output hourly report volume <30 mL/hr ,
- Fetal heart rate
- Intrauterine fetal resuscitation while being transferred to Operation theatre(SPOILT)
- S-Syntocinon off
- P-Position full left lateral
- O- Oxygen
- I- Iv infusion of 1 L crystalloid
- L- Low blood pressure, correct with vasopressor
- T-Tocolysis to stop contractions
- Surgical team should then be scrubbed with the woman"painted and draped" prior to induction.. When fetal bradycardia occurs it becomes mandatory to extract the baby as soon as possible by caeserean section within a span of at least 20 minutes in order to reduce neonatal mortality as well as incidence of cerebral palsy.
- Rapid sequence intubation with Ketamine and Etomidate are induction agents of choice in volume depleted. For maintenance,adding a volatile agent helps to reduce awareness,however the concentration should be less than 0.5 MAC, to prevent uterine

atony.. Nitrous oxide is better avoided in fetal distress

- Volume resuscitation is commenced with crystalloid (2L) and colloid (1.5L).
- Oxytocin should be given to prevent uterine atony
- Tranexaemic acid 1gm IV is given when uterotonics fail to control bleeding.
- Selective embolisation of these vessels may lead to cessation of bleeding.
- In cases that fail to respond to these conservative methods, hysterectomy may be necessary

RCOG Green-top Guideline No. 52 states that: 'While results of coagulation studies are awaited, up to 1 litre (equivalent to 4 units) of FFP and 10 units of cryoprecipitate (two packs) may be given empirically in uncontrollable bleeding 47]. Early the face of empirical FFP might be justified if significant consumption is likely as in placental abruption ,however the FIBTEM A5 result being available within 10 minutes[48] it is more logical to give FFP after knowing FIBTEM A5 levels. If the FIBTEM A5 is ≥ 16 mm (equivalent to a fibrinogen level of about 3 g.l_1[49])no coagulation factors are required .If between 12-16 defrost FFPand transfuse if ongoing bleeding In case FIBTEM A5 is less than 12 which roughly corresponds to a fibrinogen level of 2.2gm%[49],the available source of fibrinogen either cryoprecipitate or fibrinogen concentrate may be given.

FIBTEM A5 LEVEL	APPROX	MANAGEMENT
	FIBRINOGEN	
>16	3gm/l	No coagulation factors required
12-16	2-3gm/l	Defrost FFP and transfuse if active
		bleeding
<12	2.2gm/l	Fibrinogen concentrate or cryoprecipitate
6	1gm/l	Fibrinogen concentrate or cryoprecipitate

Table-6: Fibrinogen levels and management

Indication for platelets: In abruption consumptive coagulopathy can cause thrombocytopaenia, Platelets are infused if the count <75 x 10[9]l_1 ,low EXTEM but normal FIBTEM levels[49], more than 10 units blood given[49].

If EXTEM monitors coagulation via the extrinsic pathway,.FIBTEM monitors clot firmness after blocking platelet contribution to it . Fibrinogen concentrate[50-53], available as a powder for reconstitution, is especially useful in resource limited areas and gives a reliable and quick increase in fibrinogen levels thereby averting coagulopathy. In order to raise the fibrinogen level by 1 g/l, 30 ml/kg of FFP needs to be given, compared with 3 ml/kg of cryoprecipitate.and 60mg/kg of fibrinogen concentrate[54]. Recombinant FVII a is a controversial option to be given as last resort just before hysterectomy.

If there is fetal death, then the abruption is large and indicates significant maternal blood loss. and a coagulopathy is possible in 30%cases.[55]There is likely to have been up to 1500ml of haemorrhage. It is important to give up to 2 units of blood as soon as possible The on-call consultant obstetrician and anaesthetist should be informed The incidence of coagulopathy is very low if the foetal demise has happened within 2 weeks of presentation. Anaesthetist should assess the patient for possibility of coagulopathy and or sepsis, especially before administering regional anaesthesia.

A coagulopathic patient may present for vaginal delivery after fetal demise,.In this case intravenous patient controlled opiod analgesia should be given.

After Care:

All women following major APH require intensive monitoring for at least first 24 hours. It is important to remember that thrombo-embolic disease (TED)[56] is still one of the commonest cause of maternal morbidity .Stockings is the bare minimum that can be offered.

UTERINE RUPTURE

It is one of the most dreaded complications of childbirth with potentially grave consequences to both mother and fetus. Occurrence is - 0.4-0.6% of all deliveries.

Prognosis

4.2% maternal mortality[57]46% perinatal mortality[57] In half of the cases, uterine rupture occurs at the previous lower segment cesarean section (LSCS) scar .The rupture of classical caesarean scar is related with more severe morbidity and mortality as the anterior uterine wall is highly vascular and may include some part of placenta[58]. The overall incidence of uterine rupture following caesarean section is 1in100[57].

INCIDENCE OF UTERINE	%
RUPTURE	
After first LSCS	0.79%,
second LSCS	0.9-1.8%
short interdelivery interval	1.3-4.8%,
-induction of labor with oxytocin	0.7%
prostaglandin induction	2.4%

 Table 7: Incidence of uterine rupture

Women with previous LSCS scar and having induction of labor are more prone to rupture than those who deliver spontaneously [59, 60].

Warning signs of impending rupture [60,61,59]

- Uterine contraction>5 in 10 minutes.lasting >60-90seconds
- Fetal heart showing tachycardia,bradycardia lasting >10 minutes
- Tenderness over scar.
- Maternal tachycardia
- Bandl,s ring formed

Patient may present with

- abdominal pain,
- abnormal fetal heart rate
- vaginal bleeding,
- uterine hypertonia,
- cessation of labour,
- hypotension,
- loss of fetal station and
- change in fetal presentation
- Breakthrough pain of neuraxial labor analgesia may also indicate rupture[60].

Abnormal fetal heart rate pattern on CTG including bradycardia, recurrent, variable or late deceleration is the most diagnostic feature [59,60,62]. There can be sudden fetal death (10-80%)[62], or hypoxia and acidosis. Maternal mortality (0-30%) is due to shock, DIC and sepsis .Neonatal outcome following a uterine rupture will depend primarily on the speed with which the C-section is performed.[63] Every minute matters.[85]. It is not necessary that fetal hypoxia will be avoided if the "30-minute decision-to-incision rule" is followed.[64] Studies suggest that babies born within 17 minutes of complete anoxia or severe hypoxia will survive neurologically intact, while babies born after 17 minutes may have severe damage, or will not survive at all.[65,83]

Anaesthetic management

The ASA guidelines recommend that neuraxial techniques are no more contraindicated in patients attempting vaginal birth after ceaserean(VBAC)[66] Chestnut stated that the pain of uterine rupture being continuous in nature may not be relieved by epidural analgesia "[67].

The anesthesiologist should be informed before labour of all VBAC patients so that he/she may perform a pre-anesthetic evaluation and be prepared in case an emergency arises[66,84].

Chestnut's textbook states that "epidural analgesia is a necessary requirement of a successful VBAC program . analgesic doses should be given. Epidural analgesia does not delay the diagnosis of uterine rupture or decrease the likelihood of successful VBAC."[68]. The anesthesiologist should alert the obstetrician if the patient has excessive analgesic requirements[69], suggesting the need for an evaluation for uterine rupture. Anesthesiologists should be actively involved, [84]. In the process of labour. Repeat cesarean section (C-section),has a higher morbidity and mortality in comparison to a successful VBAC.A failed VBAC is associated with a small but significant risk of uterine rupture that can result in death or serious injury to both the mother and the infant.[66,81].

A patient should not be offered a VBAC in a facility where practitioners capable of performing cesarean sections are not available 24x7[66]. The definitive treatment is hysterectomy in case rupture occurs .General anaesthesia is preferred because of fetal compromise.Pre existing epidural may be topped up in stable patients.Invasive monitoring is helpful. Give prophylactic antibiotics.

A rapid response institutional protocol should be developed[70].

Vasa praevia

In this condition fetal blood vessels traverse the lower uterine segment in advance of the presenting part[71,72].It presents as painless vaginal bleeding at the time of spontaneous or artificial rupture of membranes.Since blood loss is fetal in origin fetal demise or shock may occur rapidly [72,73,74].

Fetal blood loss can be confirmed by the haemoglobin alkaline denaturation test (Apts test)[76] or Kleihauer-Betke[75]test. All women with a history of a low lying placenta should have a colour Doppler ultrasound done before term to confirm location of placenta[77,78,79] An antenatal diagnosis requires elective caesarean section [80] Prompt management by neonatologist and volume replacement with O negative blood may improve outcome.

Antepartum haemorrhage irrespective of its cause carries significant risk of adverse perinatal outcome. Ante natal diagnosis of placenta praevia, accreta along with prompt and effective measures have reduced the maternal complications Changing trends in management of abruption with point of care devices like FIBTEM A5assay and availability of fibrinogen concentrate has revolutionalized the patient care in coagulopathy. Above all is a multi disciplinary team effort involving the experienced anaesthetist at an early stage to achieve desired outcome.

REFERENCES

- 1. Maternal Mortality in 2005:estimates developed by WHO, UNICEF, UNFPA and the World Bank. World Health Organisation.
- 2. WHO Report 2005: Make every mother and child count. Department of Health.
- 3. Saving Mothers 2005–2007. Fourth Report on Confidential Enquiries into Maternal Deaths in South Africa. Pretoria: Department of Health; 2009.
- 4. Joint Statement: Prevention and treatment of post-partum haemorrhage new advances for low resource settings. Inter- national

Confederation of Midwives and International Federation of Gynecologists and Obstetricians. www.figo.org.

- 5. The Millenium Development goal report 2009 United Nations 2009.
- 6. Mayer D, Spielman FJ, Bell EA. Antepartum and postpartum hemorrhage. Chestnut DH, editor. Obstetric anesthesia. Principles and practice. 3rd edition. Philadelphia: Elsevier Mosby; 2004: 662–82.
- Antepartum Haemorrhage (APH) King Edward Memorial Hospital for Women Clinical Guidelines: Obstetrics & Midwifery Perth, Western Australia 2015 (B2.3)
- 8. Crafter H, J B. Common problems associated with early and advanced pregnancy. In: Marshall J, Raynor M, editors. Myles textbook for midwives. 16th ed. Edinburgh: Churchill Livingstone Elsevier; 2014. p. 221-42.
- Camann WR, Biehl DH. Antepartum and postpartum hemorrhage. In: Hughes SC, Levinson G, Rosen MA, editors. Shnider and Levinson's anesthesia for obstetrics. 4th edition. Philadelphia: Lippincott Williams & Wilkins; 2002:361–71.
- 10. Royal College of Obstetricians and Gynaecologists. Antepartum Haemorrhage. Green-top Guideline No 63. 2011.
- Calleja-Agius J, Custo R, Brincat MP, Calleja N. Placental abruption and placenta praevia. Eur Clin Obstet Gynaecol 2006; 2:121–7
- 12. Ananth C.V., and Wilcox A.J. Placental abruption and perinatal mortality in the United States. Am J Epidemiol 2001; 153: 332-337.
- Tikkanen M., Luukkaala T., Gissler M., et al: Decreasing perinatal mortality in placental abruption. Acta Obstet Gynecol Scand 2013; 92: 298-305.
- Moussa HA, Alfirevic Z. Treatment of primary postpartum haemorrhage. Cochrane Database of Systematic Reviews 2007, Issue 1 Art. No. CD003249. DOI: 10.1002/14651858.CD003249.pub2.

15. Schorn MN. Measurement of blood loss review of the literature. J Midwifery Womens Health, 2010; 55 (1), 20–27.

- 16. Bose P, Regan F, Paterson-Brown S. Improving the accuracy of estimated blood loss at obstetric haemorrhage using clinical reconstructions. BJOG. 2006; 113(8) :919 -924
- 17. Patel A, Goudar SS, Geller SE et al. Drape estimation vs. visual assessment for estimating postpartum hemorrhage. Int. J. Gynaecol. Obstet. 93(3), 220–224 (2006).
- Royal College of Obstetricians and Gynaecologists. Placenta Praevia, Placenta Praevia Accreta and Vasa Praevia: Diagnosis

and Management. Green-top Guideline No. 27.London: RCOG; 2011.

- 19. Nottingham university hospitals NHS Trust Guideline for the Management of Antepartum Haemorrhage May 2014.
- 20. George Gallos, , Imre Redai, Richard M. Smiley, The Role of the Anesthesiologist in Management of Obstetric Hemorrhage Semin Perinatol 33:116-123 c 2009 Elsevier Inc.
- Confidential Enquiry into Maternal and Child Health. Saving Mothers Lives 2003–2005. Seventh Report on Confidential Enquiries into Maternal Deaths in the United Kingdom. London: CEMACH; 2006 [www.cemach.org.uk/getattachment/927cf18a-735a-47a0-9200-cdea103781c7/Saving-Mothers--Lives-2003-2005_full.aspxy.8
- 22. Royal College of Obstetricians and Gynaecologists. Placenta praevia after caesarean section care bundle February 2010.
- 23. Oppenheimer L, Holmes P, Simpson N, Dabrowski A. Diagnosis of low-lying placenta: can migration in the third trimester predict outcome? Ultrasound Obstet Gynecol 2001;18:100–2.
- 24. Bonner SM, Haynes SR, Ryall D. The anaesthetic management of Caesarean section for placenta previa: a questionnaire survey. Anaesthesia. 1995; 50:992-4.
- 25. Peel WJ. A survey of the anaesthetic management of patients presenting for Caesarean section with high risk obstetric conditions. Int J Obstet Anaesth. 1996; 5:219-20.
- 26. Plumer MH, Rottman R. How anesthesiologists practice obstetric anesthesia. practicing Responses of obstetric anesthesiologists at the 1993 meeting of the for Obstetric Society Anesthesia and Perinatology. Reg Anesth. 1996; 21:49-60.
- 27. Oyelese Y, Smullian J. Placenta previa, placenta accrete, and vasa previa. Obstet Gynecol. 2006; 107:927-41.
- Miller DA, Chollet JA, Goodwin TM. Clinical risk factors for placenta previaplacenta accrete. Am J Obstet Gynecol. 1997; 177(1):210 – 4.
- 29. Ted Hunter MD, Simcha Kleiman MO. Obtetric K. Anaesthesia for Caesarean hysterectomy in a patient with preoperative diagnosis of placenta excreta with invasion of the urinary bladder . Can J Anaesth. 1996; 4(3):246-51
- 30. Tan CH, Tay KH, Sheah K, Kwek K, Wong K, Tan HK, et al. Perioperative endovascular internal iliac artery occlusion balloon placement in management of placenta accreta. Am J Roentgenol 2007;189:1158-63.

- 31. Bodner LJ, Nosher JL, Gribbin C, Siegel RL, Beale S, Scorza W. Balloon assisted occlusion of the internal iliac arteries in patients with placenta accreta/ percreta. Cardiovasc Intervent Radiol 2006;29:354-61
- 32. Reitman E, Devine PC, Laifer-Narin SL, Flood P. Case scenario: perioperative management of a multigravida at 34-week gestation diagnosed with abnormal placentation. Anesthesiology 2011 Oct; 115(4): 852-7.
- Chestnut DH, Dewan DM, Redick LF, Caton D, Spielman FJ: Anesthetic management for obstetric hysterectomy: A multi-institutional study. Anesthesiology 1989 Apr; 70(4):607– 10.
- Kristen C. Sihler and Lena M. Napolitano. Complications of Massive Transfusion MD CHEST 2010; 137(1): 209 – 220.
- 35. Bhavani-Shankar K, Lynch EP, Datta S: Airway changes during Cesarean hysterectomy. Can J Anesth 47:338-341, 2000 27.
- Ahonen J, Jokela R. Recombinant factor VIIa for life-threatening post-partum haemorrhage. Br J Anaesth 2005 May; 94(5):592–5
- 37. Gungorduk K, Yıldırım G, Asıcıogʻlu O, Gungorduk OC, Sudolmus S, Ark C. Efficacy of intravenous tranexamic acid in reducing blood loss after elective cesarean section: A prospective, randomized, double-blind, placebo-controlled study. Am J Perinatol 2011; 28:223–40.
- Oyelese Y, Ananth C.V. Placental Abruption. Obstet Gynecol 2006; 108: pp.1005-1016
- Ananth C.V., Wilcox A.J. Placental abruption and perinatal mortality in the United States. Am J Epidemiol 2001; 153: pp. 332-337
- Hurd W.W., Miodovnik M., Hertzberg V., and Lavin J.P.: Selective management of abruptio placentae: a prospective study. Obstet Gynecol 1983; 61: 467-473
- 41. Tikkanen M., Luukkaala T., Gissler M., et al: Decreasing perinatal mortality in placental abruption. Acta Obstet Gynecol Scand 2013; 92: pp. 298-305
- 42. Kayani S.I., Walkinshaw S.A., and Preston C.:Pregnancy outcome in severe placental abruption. BJOG 2003; 110: pp. 679-683.
- 43. Hall DR. Abruptio Placentae and Disseminated Intravascular Coagulopathy. Seminars in Perinatology. 2009;33:189-95.
- Glantz C., and Purnell L.: Clinical utility of sonography in the diagnosis and treatment of placental abruption. J Ultrasound Med 2002; 21: pp.837-840
- 45. Kikutani M., Ishihara K., and Araki T.: Value of ultrasonography in the diagnosis of

Available online at http://saspublisher.com/sjams/

placental abruption. J Nippon Med Sch 2003; 70: pp. 227-233

- 46. McNamara H , et al. Coagulopathy and placental abruption: changing management with ROTEM-guided fibrinogen concentrate therapy. Int J Obstet Anesth. 2015 May; 24(2): 174-9.
- 47. Kozek-Langenecker SA. Perioperative coagulation monitoring. Best Pract Res Clin Anaesthesiol. 2010;24(1):27-40
- 48. Huissoud C, Carrabin N, Audibert F, et al. Bedside assessment of fibrinogen level in postpartum haemorrhage by thrombelastometry. International Journal of Obstetrics and Gynaecology 2009; 116: 1097– 102
- Mallaiah S, Barclay P, Harrod I, Chevannes C, Bhalla A. Introduction of an algorithm for ROTEM-guided fibrinogen concentrate administration in major obstetric haemorrhage. Anaesthesia 2014 Oct 7; doi:10.1111/anae.12859
- 50. Ahmed S, Harrity C, Johnson S, et al. The efficacy of fibrinogen concentrate compared with cryoprecipitate in majorobstetric haemorrhage an observational study. Transfusion Medicine 2012; 22: 344–9
- 51. Bell SF, Rayment R, Collins PW, Collis RE. The use of fibrinogen concentrate to correct hypofibrinogenaemia rapidly during obstetric haemorrhage. International Journal of Obstetric Anesthesia 2010; 19: 218–23.
- 52. Glover NJ, Collis RE, Collins P. Fibrinogen concentrate use during major obstetric haemorrhage. Anaesthesia 2010; 65:1229–30.
- 53. Gollop ND, Chilcott J, Benton A, Rayment R, Jones J, Collins PW. National audit of the use of fibrinogen concentrate to correct hypofibrinogenaemia. Transfusion Medicine 2012; 22: 350–5
- 54. Dyer RA, Vorster AD, Arcache MJ, Vasco M. New trends in the management of postpartum haemorrhage. South African Journal of Anaesthesia and Analgesia, 2014;20(1):44 – 7
- 55. Maslow, Breen, Sarna, Soni, Watkins,Oriol. Prevalence of coagulation abnormalities associated with intrauterine foetal death. Canadian J Anaesthesia,1996;43(12):1237-43.
- 56. Sachdeva A, Dalton M, Amaragiri S, Lees T. Graduated compression stockings for prevention of deep vein thrombosis (Review). Cochrane Database of Systematic Reviews. 2014 Dec 17;12CD001484.
- 57. Landon MB, Hauth JC, Leveno KJ, et al; Maternal and perinatal outcomes associated with a trial of labour after prior caesarean delivery. N Engl J Med. 2004Dec 16;351(25):2581-9.

- 58. Smith J, Mertz, H, Merrill, D. "Identifying risk factors for uterine rupture." Clin Perinatol 2008;35: 85–99.
- 59. Craver Pryor E, Mertz H, Beaver B, et al. "Intrapartum predictors of uterine rupture." Am J Perinatol 2007; 24:317–322.
- Vaginal Birth after Previous Cesarean Delivery. Washington, DC: ACOG; August 2010
- 61. Grobman WA, Lai Y, Landon MB, et al; Prediction of uterine rupture associated with attempted vaginal birth after caesarean delivery. Am J Obstet Gynecol. 2008 Jul;199(1):30.
- 62. Smith GC, Pell JP, Pasupathy D, et al; Factors predisposing to perinatal death related to uterine rupture during attempted vaginal birth after caesarean section: retrospective cohort study. BMJ. 2004 Aug 14;329(7462):359-60.
- 63. Toppenberg K, Block W. "Uterine Rupture: What Family Physicians Need to Know. Am Fam Physician 2002 Sept; 66(5): 823–8.
- 64. Zakowski M. "Obstetric anesthesiology: What's new, what's old and what's standard?" CSA Bulletin; 60:87–96.
- Leung, et al. "Uterine Rupture after Previous Cesarean Delivery: Maternal and Fetal Consequences." Am J Obstet Gynecol 1993;169: 945–950.
- 66. American Society of Anesthesiologists, Practice Guidelines for Obstetric Anesthesia. Anesthesiology 2007; 106:843–863
- 67. Chestnut D. "Vaginal Birth After Cesarean Section," in Obstetric Anesthesia: Principles and Practice, 3rd ed., Chestnut D (ed), 2004 (Philadelphia: Mosby Elsevier), p. 403.
- 68. Chestnut D. "Vaginal Birth After Cesarean Delivery," in Obstetric Anesthesia: Principles and Practice, 4th ed., Chestnut D (ed), 2009 (Philadelphia: Mosby Elsevier), p. 383
- 69. Cahill A, Odibo A, Allsworth J, et al. "Frequent epidural dosing as a marker for impending uterine rupture in patients who attempt vaginal birth after cesarean delivery." Am J Obstet Gynecol 2010 Apr;202(4):355.
- Catanzarite V, Almryde K, Bombard A. "Grand Rounds: OB Team Stat: Developinga Better L&D Rapid Response Team." Contemporary OB/GYN 2008 September; 1–7.
- 71. Catanzarite V, Maida C, Thomas W, Mendoza A, Stanco L, Piacquadio KM. Prenatal sonographic diagnosis of vasa previa: ultrasound findings and obstetric outcome in ten cases. Ultrasound Obstet Gynecol 2001 Aug;18(2):109–15.
- 72. Stafford IP, Neumann DE, Jarrell H. Abnormal placental structure and vasa previa:

Available online at http://saspublisher.com/sjams/

confirmation of the relationship. J Ultrasound Med 2004;23:1521–2.

- Antoine C, Young BK, Silverman F, Greco MA, Alvarez SP. Sinusoidal fetal heart rate pattern with vasa praevia in twin pregnancy. J Reprod Med 1982;27:295–300.
- 74. Cordero D R, Helfgott A W, Landy H J, Reik R F, Medina C, O'Sullivan M J. A nonhemorrhagic manifestation of vasa previa: a clinico pathological case report. Obstet Gynecol 1993;82:698–700
- 75. Kleihauer E, Braun H, Bethe K. [Demonstration of fetal hemoglobin in erythrocytes of a blood smear]. KlinWochenschr 1957;35:637–8. Article in German.
- 76. Apt L, Downey Jr WS. Melena neonatorum: the swallowed blood syndrome; a simple test for the differentiation of adult and fetal hemoglobin in bloody stools. J Paediatr 1955;47:6–12.
- 77. Oyelese Y, Catanzarite V, Prefumo F, Lashley S, Schachter M, Tovbin Y, et al. Vasa previa: the impact of prenatal diagnosis on outcomes. Obstet Gynecol, 2004;103:937–42.
- 78. Oyelese Y, Smulian JC. Placenta previa, placenta accreta, and vasa previa. Obstet Gynecol 2006;107:927–41.
- 79. Fung TY, Lau TK. Poor perinatal outcome associated with vasa previa: is it preventable? A report of three cases and review of the literature. Ultrasound Obstet Gynecol 1998;12:430–3.
- National Institute for Clinical Excellence. Clinical Guideline 13:Caesarean section. London: National Institute for Clinical Excellence; 2004[http://www.gserve.nice.org.uk/nicemedia /pdf/CG013NICEguideline.pdf].
- Greene, MF. "Vaginal delivery after cesarean section—is the risk acceptable?" N Engl J Med 2001; 345:54–55.
- 82. Task Force on Neonatal Encephalopathy and Cerebral Palsy, Staff American College of Obstetricians and Gynecologists with American College of Obstetricians and Gynecologists, Practice Bulletin Number 106. Washington, DC: ACOG; July 2009.
- Birth after caesarean Green Top Guidelines No 45. RCOG 2007. Caesarean section.
- 84. Managing the Risk of Uterine Rupture During a Trial of Labor After Cesarean. Section By NORCAL Mutual Insurance Company. September 2011.