Evaluation of Risk Factors and Outcome of Haemorrhagic Stroke in Neonates

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Abstract

Background: Few data regarding causes and outcomes of hemorrhagic stroke in late preterm and term neonates are available so we were trying to identify them in this study.

Methods: This prospective case control study was carried out at Sohag University Hospitals, during the period from 1/1/2015 to 31/12/2015 at the department of pediatrics and neonatal care unit at Sohag University. A total of 50 neonates term and late preterm neonates (≥34 weeks gestation) and ≤28 days of life had met the inclusion criteria were 23 males and 27 females.

Results: As regard outcome in our study, 8% had grade I, 14 % had grade II, 32% had grade III, 12% had grade IV, 14.5% had subdural hemorrhage and 20% had subarachnoid hemorrhage. So most patients have grade III intracranial hemorrhage and subarachnoid hemorrhage. According to the severity, in our study patients who had mild intracranial hemorrhage represented 48% and patients with moderate intracranial hemorrhage represented 30% and patients with severe intracranial hemorrhage represented 22%.

Conclusion: IVH is very rarely reported in full-term and late preterm neonates and may occur in these children with a variety of risk factors, in our study we tried to identify these risk factors and also the outcome and we found that the most frequent associated risk factors were absence of vitamin K administration followed by prolonged labour then neonatal sepsis.

Introduction

Hemorrhagic stroke (HS) defined as intraventricular, intraparenchymal intraventricular hemorrhage is common and usually caused by germinal matrix hemorrhage. While it is less common in late preterm (34-37) wks and term neonates. Much less is known about risk factors and outcome of HS in late preterm (34-37) weeks and term neonates (1).

Incidence of symptomatic HS in term neonates has recently been estimated at 0.17/1000 live births (2). Hemorrhage location often dictates the clinical course. Symptoms can include seizures or may be non-specific like apnea, respiratory distress, fever or poor feeding (3).

Brouwer AJ have reported impairment in (9-57%) of these children though the higher estimate comes from a study that included neonates with isolated subdural hemorrhage. Most studies report mortality rates from 1% to 25 % (4). Intracranial hemorrhage in term newborn has been increasingly recognized but the occurrence and risk factors in late preterm are still unclear (5).

Several risk factors have been reported in term newborns with ICH. Maternal risk factors causing ICH in the first neonatal week includes usage of drugs (such as aspirin, cocaine), pregnancy-induced hypertension, placental abruption, placental alloimmunization and autoimmune disorders. Major perinatal risk factor are birth trauma, low Apgar score, resuscitation at birth, thrombocytopenia, breastfed infants who received no vitamin K, inherited coagulopathy, disseminated intravascular coagulopathy, increased cerebral venous pressure, prolonged labor, unassisted vaginal delivery, forceps delivery and suction cup (6).

Aim of the work: The work aimed to identify the risk factors and outcome of hemorrhagic stroke in neonates ≥34 weeks of gestational age and ≤28 days of life in Sohage University Hospital.

Patients & Methods:

Type of the Study: prospective hospital based study.

Place of the Study: Neonatal intensive care unit (N.I.C.U) and Neuropediatric Outpatient Clinic at Sohage University Hospital.
Discussion

The present study performed a clinical analysis and reviewing the data of 50 term and late preterm neonates (≥34 weeks gestation) and ≤28 days having intracranial hemorrhage by neuroradiological findings (CT or MRI Brain or transcranial U/S) with special consideration for the risk factors and neurological out come. The study included a total of 50 neonates having intracranial hemorrhage with age range between 34wks of gestational age to less than 28 days of life. The median age was 12 days and there were 23 males (46%) and 27 females (54%). The male to female ratio was (1:1.2). This ratio was close to the study of Bruno, et al., 2014 where male to female ratio was (1:1). The mean age incidence was 13.56 days which was higher than that of study of Bruno, which was 7 days (7).

Concerning the median gestational age of the studied population is 37 wk range from 34wks to 39wks, which is less than that of Burno et al., that show median gestational age 39.7wks, range from 38wks to 40.7wks (7) and it was also less than that of Afsharkhaset al., 2015 (38 wks)(8).
Intracranial hemorrhage in late preterm and full term neonates is multifactorial, in our study cases had two or more risk factors but the most associated risk factors were prolonged labor (28%) and patients who did not administered vitamin K (30%). As regard risk factors of intracranial hemorrhage, mode of delivery is one of the most important factors that had been implicated as a cause of intracranial hemorrhage. In our study 50% had been delivered by normal vaginal delivery and the other 50% has been delivered by cesarean section which was different from the result of Bruno et al., in which 72% had been delivered by normal vaginal delivery and 28% by cesarean section (7). Instrumental use during delivery was used in 3% of our patients.

Towner et al., reviewed that the rate of intracranial hemorrhage was higher among infants delivered by vacuum extraction, forceps, or cesarean section than among infants delivered vaginally without instrumentation. The increased risk of intracranial hemorrhage among infants delivered by vacuum extraction or forceps, however, may be the result of common risk factors during abnormal labor (9). In our study, Apgar score mean at 5 minutes was 8.8 in contrast to the study of Jahwar et al., the mean Apgar scores at 5 minutes for case patients was 6.0. Also, Jahwar et al., reported that among 11 cases of term neonates, intracranial hemorrhage was evidenced in all 3 of the infants who had 5-minute Apgar score of 7 or less also reported that low Apgar scores, with and without requirement for resuscitation at birth, are risk factors for intracranial hemorrhage (10). A significant clinical problem, which is generally regarded as one of the risk factors of IVH, is asphyxia, a secondary consequence to an interruption of placental blood flow and decreased blood flow to the brain and impairment of cerebral autoregulation. Treatment with therapeutic hypothermia in neonates with hypoxic ischemic encephalopathy may prevent the development of severe brain injuries and improve the long-term outcome. On the other hand, it places newborns at greater risk of IVH by causing fluctuations of the cerebral blood flow, depressed cardiac function, hypotension, and changes in coagulation cascade (11).

In our study, foetal distress is recorded in 6% of our cases which is lower than the study of Bruno et al., in which patients with respiratory distress reported in 21% of cases (7).

Patients who did not administered vitamin K was 30% of cases this result was close to Hubbardet al., that reported that vitamin K is essential for the final carboxylation of coagulation factors II, VII, IX, and X (12). Vitamin K deficiency can lead to hemorrhagic disease of the newborn, which is rare in the United States because of routine administration of vitamin K but thereare reports of intracranial hemorrhage with this deficiency. It is most common in exclusively breast-fed infants who received no vitamin K after birth, and also in infants with mothers taking various antiepileptic drugs, analgesics or street drugs (13).

Neonatal sepsis had been reported in 6% of cases of our study, sepsis can lead to intracranial hemorrhage by several ways such as thrombocytopenia and disseminated intravascular coagulopathy. Disseminated intravascular coagulation can also lead to severe thrombocytopenia and intracranial hemorrhage in the term newborn, but this occurs in the sick newborn, usually in the neonatal intensive care unit setting (14).

In our study, the most frequent presentation was convulsions (50%) followed by pallor (20%) and poor suckling (22%). Indeed, intracranial hemorrhage is a common cause of neonatal seizures. The seizure occurrence, type, and severity are variable in term newborns. Seizure may occur soon after birth or after an extended period of normality days after birth. Neonatal seizures may prove difficult to differentiate from normal movement, or may have variable presentations, ranging from a subtle seizure such as apnea to a more apparent seizure to status epilepticus (15).

Apnea was one the most important clinical manifestations it represented 16%. Apnea of any duration associated with significant tachycardia or bradycardia in a newborn who is otherwise apparently well should prompt the physician to consider intracranial abnormalities such as intracranial hemorrhage Temporal lobe hemorrhage with apneic seizures in term neonates although uncommon, has been reported (16).
On the other hand, apnea with bradycardia may be a manifestation of elevated intracranial pressure (17).

Concerning hematological investigations, 54% of patients had prolonged PT and aPTT and 24% had thrombocytopenia which was near the study of Afsharkhas et al., and also near to the study of (Afsharkhas et al, 2015) in which 50% of neonates had coagulation profile disorders including DIC (20%), sinusovous thrombosis (16.7%) and thrombocytopenia (13.3%) of cases (8,10).

Sandberg et al., Identified 11 term newborns with spontaneous intraparenchymal hemorrhage; coagulopathy was noted in 3 of the 11. Thrombocytopenia is the most common condition leading to intracranial hemorrhage in term newborns. Thrombocytopenia has multiple causes; it may be drug induced, infectious, genetic, or immune-related, or due to disseminated intravascular coagulation or placental insufficiency. The risk for intracranial hemorrhage increases with severity of thrombocytopenia. Most cases occur with platelet counts of less than 30,000/mm3. Neonatal alloimmune thrombocytopenia is a particularly important risk for intracranial hemorrhage: 7-26% of affected neonates develop intracranial hemorrhage (15).

Intracranial haemorrhage can be classified according to the degree as follow: grade I (the mildest form of IVH that bleeding was limited to the lining of the ventricles), grade II (the blood does spilled into the ventricles, but there was no enlargement or swelling), grade III (the ventricles had become enlarged and are full of blood), and grade IV (Blood spilled out from the ventricles into the surrounding brain). In our study 8% had grade I, 14% had grade II, 32% had grade III, 12% had grade IV, 14.5% had subdural hemorrhage and 20% had subarachnoid hemorrhage. So most patients had grade III intracranial hemorrhage followed by subarachnoid hemorrhage 20%, this was different from the study of (Afsharkhas et al., 2015) in which most patients had grade I (33%) followed by grade II (20%) followed by grade IV (15%) (8).

Also intracranial hemorrhage can be classified according to the severity into mild intracranial hemorrhage which involves only one compartment or one lobe, with a maximum midline shift of 0.5 cm, or as intraventricular hemorrhage in only one ventricle with no hydrocephalus. Moderate intracranial hemorrhage is defined as hemorrhage involving only one lobe and compartment with midline shift or as intraventricular hemorrhage of more than one ventricle but no hydrocephalus; when two or more lobes are involved, midline shift is inconsequential. Severe intracranial hemorrhage is defined as hemorrhage in more than one lobe and in more than one compartment or as intraventricular hemorrhage with hydrocephalus. In our study grade I is considered mild, grade II moderate, and grade III & IV severe. In our study mild intracranial hemorrhage represent 48% and patients with moderate intracranial hemorrhage represents 30% and patients with severe intracranial hemorrhage represents 22% so most patients in our study had mild intracranial hemorrhage and the least had severe intracranial hemorrhage this is close to the study of Afsharkhas et al., in which most patients had mild intracranial hemorrhage (33%) followed by moderate intracranial hemorrhage which represented (30%) followed by severe intracranial hemorrhage which represented (22%) (8).

As regard the outcome; there is no single predictor of intracranial hemorrhage outcome in newborns. However, the extent (compartmental, lobar, or both), the severity, and the etiology of intracranial hemorrhage may influence the prognosis. In addition, low gestational age (i.e., prematurity), early occurrence of recurrent seizures or status epilepticus, and the need for multiple anticonvulsants to control seizures were associated with poor outcome (18). Afsharkhas et al., also reported that the consequences of IVH are dependent to the grade of bleeding. So that, mortality and sequelae due to IVH, ranged from 5% in grade I to 90% in grade IV, respectively (8). The majority of newborns with spontaneous subarachnoid hemorrhage will make a complete recovery. Disability is more likely to occur in a newborn with frontal lobe hemorrhage or when multiple intracranial compartments are involved and this was confirmed with our study that report that all patients with subarachnoid hemorrhage had no
neurological abnormalities in the follow up and patients who had grade III intracranial hemorrhage associated with the highest incidence of neurological abnormalities (71.43%).

33% of patients had abnormal neurological outcome in the follow up at the age of six months in the form of hypotonia, spasticity, hydrocephalus, apathy and poor activity. The incidence of neurological abnormality in our study was lower than that of Bruno et al., in which nearly half of survivors demonstrated a neurologic deficit at the last follow-up, and approximately 56% of subjects required physical, occupational, and/or speech therapy services (7). Hydrocephalus in our study had been reported in 6 cases that required surgical intervention.

Summary and conclusion:
This study carried out at Neonatal Care Unit and Pediatric Neurology Clinic at Sohag University Hospitals, during the period from 1/1/2015 to 31/12/2015. The study included a total of 50 neonates meeting the inclusion criteria having intracranial hemorrhage with age range (≥34 weeks gestation and ≤28 days of life). The median age was 12 days and there were 23 males (46%) and 27 females (54%). The male to female ratio was (1:1.2).

As regard outcome in our study, 8% had grade I, 14% had grade II, 32% had grade III, 12% had grade IV, 14.5% had subdural hemorrhage and 20% had subarachnoid hemorrhage. So most patients have grade III intracranial hemorrhage and subarachnoid hemorrhage. All newborns with subarachnoid hemorrhage showed a complete recovery. Disability is more likely to occur in a newborn with frontal lobe hemorrhage or when multiple intracranial compartments are involved and this was confirmed in our study which showed that all patients with subarachnoid hemorrhage had no neurological abnormalities in the follow up and patients who had grade III intracranial hemorrhage associated with the highest incidence of associated neurological abnormalities (71.43%). 33% of patients had abnormal neurological outcome in the follow up at the age of six months in the form hypotonia, spasticity, hydrocephalus, apathy and poor activity. These patients required physical, occupational, and/or speech therapy services.

Hydrocephalus has reported in 6 cases in our study that required surgical intervention.

References


