Chicken or Egg: Which Came First?

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Abstract

Cerebral venous sinus thrombosis (CVST) is quite an unusual diagnosis in pregnancy and puerperium. It can present with a wide range of symptoms and signs, and a high index of suspicion is required to diagnose this disorder. The diagnosis is challenging and could if left undiagnosed, result in life-threatening complications. Yet its management is simple with excellent prognosis, with an early diagnosis and timely intervention. We report a case of CVST in a young pregnant patient who presented with features of raised intracranial tension and loss of consciousness, after several weeks of hyperemesis gravidarum. She was treated with intravenous fluid resuscitation, low molecular weight heparin, steroids, and anti-cerebral edema measures. She subsequently made excellent recovery and is under follow-up.

Keywords: Cerebral venous sinus thrombosis; Hyperemesis gravidarum; Magnetic resonance venogram; Pregnancy complications

Introduction

Cerebral venous sinus thrombosis (CVST) is a unique condition, which is rarely seen with a highly variable clinical presentation and diverse etiologies. Though its actual incidence is not exactly known, it is estimated to be 5/1,000,000. These account for 15-20% of young strokes. The condition of "occlusion of veins that drain" was first published in 1820s and the first case of dural sinus thrombosis described by Ribes from France. The knowledge of the subject has grown tremendously since the original review by Tonelle et al in 1829. Sir Charles Symonds was the first to describe the association with benign intracranial hypertension as “otic hydrocephalus”. There are several possible etiologies though idioptic in up to 30% of the patients. CVST is quite an unusual diagnosis in pregnancy and puerperium. It can present with a wide range of symptoms and signs, and a high index of suspicion is required to diagnose this disorder. It is often confused with eclampsia, especially if it develops on a background of preeclampsia. The diagnosis of CVST is challenging and could if left undiagnosed, result in life-threatening complications. Yet its management is simple with excellent prognosis, with an early diagnosis and timely intervention. In this paper, we aim to present a case of sinus thrombosis as a rare complication of an undertreated hyperemesis gravidarum (HG), with a good outcome with aggressive medical treatment.

Case Report

A 26-year-old primigravida presented in the 10th week of pregnancy, admitted obtunded with hemiparesis and delirium in the context of HG for which she was undergoing treatment in a local hospital. She did not have any significant past medical history and denied use of any over the counter medication/herbal remedies or contraceptive pills.

On examination, the patient had bilateral papilledema, aphasia, right hemiplegia, in addition to being delirious, rapidly deteriorating to coma. There was no obvious nystagmus, anisocoria, neck stiffness or skin rash. Emergency magnetic resonance angiogram (MRA) and magnetic resonance venogram (MRV) showed extensive thrombosis involving superior/inferior sagittal veins, bilateral transverse and left sigmoid sinuses (Fig. 1). There was in addition hemorrhagic infarct in bilateral fronto parietal, cortical and suboccipital regions with moderate mass effect and diffuse venous infarcts in the right frontal cortical and subcortical regions. Bloods showed minimal leucocytosis and prerenal failure (possibly from dehydration), hypokalemia (from hyperemesis) and minimally raised ESR. Vasculitic screen (including lupus anticoagulant) and thrombophilia screen were negative.

Patient was commenced on intravenous fluids, subcutane-
ous enoxaparin twice daily, intravenous thiamine, levetiracetam, methyl prednisolone and anti-cerebral edema measures. Ultrasound of abdomen showed a 10-week-old viable intrauterine pregnancy. She had an uneventful full term normal vaginal delivery. Her symptoms improved and she was discharged on day 9 with minimal neurologic deficit on enoxaparin (at her request instead of oral anticoagulant), acetazolamide (for 8 weeks), levetiracetam and outpatient physiotherapy and rehabilitation. Repeat MRI of brain in May 2012 showed encephalomalacia in left parietal and right parietooccipital regions and no evidence of sinus thrombosis (Fig. 2).

Discussion

New onset of a neurologic syndrome in pregnancy could be the manifestation of flare up of a preexisting illness, initial presentation of a disease unique to this group or a pregnancy unrelated condition [1]. The various neurologic conditions occurring in pregnancy and peripartum period include eclampsia (specific to pregnancy), ischemic stroke, intracerebral hemorrhage, post partum angiopathy, posterior reversible encephalopathy syndrome (PRES), Wernicke-Korsakoff’s syndrome (WKS), amniotic fluid embolism, pituitary apoplexy and CVST [1]. The latter is a well-recognized, though uncommon diagnosis to inherited prothrombotic tendencies such as factor V Leiden mutation, protein C and S and anti-thrombin III deficiencies. Ten percent to 15% of CVST are due to inherited prothrombotic tendencies such as factor V Leiden mutation, protein C and S and anti-thrombin III deficiencies. Infections of head, face and middle ear leading to CVST are less common because of modern broad spectrum antibiotic treatment [11].

HG is a severe form of morning sickness, with a multifactorial neurohormonal basis, affecting 1:200 pregnancies and is the most common indication for hospital admission during the first trimester [12]. HG as a cause of cortical sinus thrombosis has been reported only once in literature [13]. CVST occurring in the first trimester is rare with mortality as high as 30% and 20% having residual weakness. The mechanism includes severe dehydration contributing to hyperviscosity that aggravates the physiologically hypercoagulable milieu (secondary to hyperestrogenism) [13, 14]. The other risk factors include cesarean section, traumatic delivery, anemia, hyper homocysteinemia and CSF hypotension secondary to leak following an epidural anesthesia. The diagnosis of CVST in this setting is often delayed due to symptoms of raised intracranial tension being attributed to hyperemesis, until focal neurologic deficit sets in. The main differential for an acute neurologial complication in the setting of HG is WKS that could also present with coma in severe cases [14]. The triad of ataxia, confusion and ophthalmoplegia and response to IV thiamine should confirm WKS [15-17]. The condition is due to poor nutritional intake, hyperemesis and increased metabolic demand [15, 18].

Neuroimaging remains the gold standard investigation [4, 9, 11, 19, 20]. A CT scan is useful for the diagnosis of infarcts or hemorrhages and to rule out other intracranial pathology, but its results can be entirely normal [20]. The most sensitive
imaging modality is MRV especially with gradient spin-echo sequence [21]. The combination of abnormal signal and a corresponding absence of flow in a cerebral venous sinus is suggestive of CVST [22]. MRI in WKS would show symmetrical increase in signal intensity in the mesencephalic tegmentum, mammillary body and medial thalamus [15, 17]. Since the diagnosis cannot be made with certainty until MR, such neurologic syndromes should be treated with intravenous hydration (to reduce viscosity and progressive thrombosis), especially if there is clinical and biochemical evidence of dehydration from the hyperemesis, in addition to intravenous thiamine, the treatment of choice for potential WKS. Treatment modalities for CVST are anticoagulation, thrombolytic therapy and in some cases surgical thrombectomy [9, 10]. Anticoagulation with low molecular weight heparin is the treatment of choice followed by oral warfarin for a 6 - 12 months period with repeat imaging to establish the status of recanalization. There are no randomized trials giving us management guidelines. In our patient warfarin was avoided as she was pregnant and then discharged on enoxaparin at her request. Prognosis is quite variable though better than CVST occurring outside of pregnancy and in men due to unknown factors. The outcome can vary from complete recovery to death; however, prospective studies have reported independent survival rates of approximately 80% [23]. Mortality is primarily due to intracerebral hemorrhage. The risk of recurrent CVST in future pregnancies and puerperium is low [24].

In conclusion, CVST may present with a wide range of symptoms and signs. Diagnosis can be confirmed by MRV in most cases. It is a life-threatening condition if left undiagnosed, but remains a potentially reversible neurologic emergency if managed timely and appropriately. It should be considered in every woman with neurological symptoms in pregnancy and puerperium. Early diagnosis and timely intervention can save lives and prevent debilitating consequences. This case also highlights the need for aggressive fluid replacement in patients with HG, often intravenous and potentially needing parenteral nutrition, to prevent devastating neurologic sequelae in this select group of pregnant women and thus minimize the morbidity and mortality associated with the condition. The awareness has to be raised amongst midwives, obstetricians, emergency physicians and the general public alike since we strongly believe the egg came first and not the chicken. Such patients should be cared for in a center that has the facility for multidisciplinary input, involving neurologists, neurosurgeons, interventional radiologists, high risk obstetricians and critical care in addition to dedicated neurorehabilitation. With improved public and medical awareness of the condition, increasing availability and accessibility to MRI and ongoing research in the field leading to better understanding, the outcomes would improve further.

Conflict of Interest

No conflict of interest was reported.

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