Case report

SUPERIOR SAGITTAL SINUS THROMBOSIS IN CHILDHOOD

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Abstract

Cerebral venous sinus thrombosis (CVST) is presence of a blood clot in the dural venous sinuses. This is a rare, but dangerous condition. CSVT is characterized by a highly variable clinical spectrum, difficult diagnosis, variable etiologies and prognosis. The International Study on Cerebral Vein and Dural Sinus Thrombosis (ISCVT) determined the frequency of the sites of SCVT. The aim of this case study was to show the clinical presentation, the examination we made, the therapy that was ordinated and the outcome of the treatment. Case study: A 14-year-old male teenager was admitted to the University Clinic for Neurosurgery in Skopje with GCS 10, accompanied with tonic-clonic epileptic seizures. CT examinations by systems were made, and during the time of recording the patient was given Dormicum 2mg overall dose. CT scan of the brain showed hyperdense zones formation around superior sagittal sinus (SSS), the rest of the medical finding was normal. We ran laboratory tests and the test for hemostasis showed deviation from the normal range. With the ordained therapy the clinical condition of the patient drastically improved. He was discharged home 12 days after the admission. He was given a recommendation for further check-ups by a transfusologist and regular visits to our clinic. Occlusion of the cerebral veins and dural venous sinuses may occur on the basis of local trauma, neoplasm or infection. Primary aseptic thrombosis may involve either cerebral veins or dural venous sinuses, or both in combination. The clinical picture and the prognosis of intracranial venous thrombosis probably depend largely on the location, extent, and rapidity of development of the venous occlusion. Conclusion: Superior sagittal sinus thrombosis is a condition that can be manifested with diverse and many symptoms and signs, which often can start unexpectedly and can be life-threatening.
Introduction

Cerebral venous sinus thrombosis (CVST) is the presence of a blood clot in the dural venous sinuses, which drain blood from the brain. This is a rare, but dangerous condition, affecting predominantly younger people, occurring with an incidence of 3-4 cases/million/year\(^1\) and for children 7 per million\(^2\). Women tend to be at an increased risk particularly between the ages of 20 - 35, mainly due to the use of oral contraceptive pills and postpartum period\(^1\). Also, one of the biggest factors for cerebral venous thrombosis in childhood is obesity. CSVT is characterized by a highly variable clinical spectrum, difficult diagnosis, variable etiologies and prognosis. It is an interesting data that in poor countries there is an association with the puerperium, with no clear arguments, but probably related to factors such as inappropriate perinatal care, metabolic conditions, and infections associated to childbirth\(^2\). Clinically nine in ten people with sinus thrombosis have a headache; this tends to worsen over the period of several days. In 40% of acute clinical presentation seizures may be present. These mostly affect only one part of the body and unilateral (occurring on one side), but occasionally the seizures are generalized and rarely can lead to status epilepticus (persistent or recurrent seizure activity for a long period of time). In the elderly, many of the abovementioned symptoms may not occur. Common symptoms in the elderly with this condition are otherwise unexplained changes in the mental status and a depressed level of consciousness. The International Study on Cerebral Vein and Dural Si-
nus Thrombosis (ISCVT) determined the frequency of the sites of SCVT as follows: transverse sinus 86%, superior sagittal sinus 62%, straight sinus 18%, cortical veins 17%, jugular veins 12%, vein of Galen and internal brain veins 11%\(^2\). In this article we report a case of a 14-year—old patient who developed a superior sagittal sinus thrombosis.

The aim of this case report was to show the clinical presentation, the examination we made, the therapy that was ordinated and the outcome of the treatment.

Case presentation

A 14-year-old male teenager, weighing 55kg, was admitted to the University Clinic for Neurosurgery in Skopje. During the check-up the patient was unresponsive, weakly reactive on motoric senses, with eyes open wide, pupils were symmetrically and narrow with movable eyeballs. The condition of consciousness when admitted was GCS 10, accompanied with tonic-clonic epileptic seizures which stopped after giving amp. Diazepam 10 mg i.m. Measured values of TA during admission equaled 160/100 mmHg, oxygen saturation O298%. From the anamnestic data received from the patient’s family we understood that the patient was found in a lying position on the house yard, without any sign of fall or trauma, uncontactable, with occasionally present epileptic seizures. By further talk with the family we did not get information about any previous diseases, no history for trauma suffered by the patient in the past, no
information for family positive anamnesis. We were also given data that the patient did not consume alcohol or cigarettes, nor he was taking any medicines or drugs. Due to the presence of the frequent epileptic seizures right after the patient check-up, the anesthesiologist was called and CT examinations by systems were made, and during the time of recording the patient was given Dormicum 2 mg overall dose. The computer findings showed the neck, thorax, abdomen and pelvis in pristine condition. The CT scan of the brain showed hyperdense zones formation around superior sagittal sinus. The rest of the medical finding was normal.

Because of the turbulent clinical condition of the patient, which had started from “full health”, we decided to make a list of examinations to discover the reason for this condition. Right after the admission to our clinic, the presence of medications and narcotics in blood was tested, and the results were negative. Complete laboratory examinations were also made, where except slightly increased values of WBC and increased values of CRP (27.4 mg/L), there were no other deviations from the normal ranges. We ran tests of hemostasis which showed a deviation from the normal range for the values of activated partial thromboplastin time (23.2 seconds), where the normal range is 27.9-37.7 seconds. We detected increased D-dimer values (4311ng/mL), the normal range being 0-500ng/mL. The following therapy was given: Tabl. Tegretol 2x200 mg, Sol. Manitol 20% 3x100 ml, amp. Nevaxon 2g, amp. Dexason 2x1, amp. Clexane (Enoxaparin) s.c 2x40 I.E, but Dexason administration was interrupted on the second day of hospitalization. Because there
was not a clear clue for the cause of capturing the consciousness accompanied with epileptic seizures, and the presence of an expansive process in the brain was excluded, MRI of the brain with and without contrast was made, but the finding was normal. In the following days of hospitalization, we continued with the anticoagulant, antibiotic, and antiepileptic and antiedematous therapy (mentioned above) and laboratory examinations were regularly made showing that WBC and CRP values were normalized.

The clinical condition of the patient drastically improved and he was reacting well on the given therapy. On the tests of hemostasis, 4 days after the admission, there were improvements of the activated partial thromboplastin time, which was 26.28 seconds, and of the D-dimer values which were 1035 ngr/mL. After 3 days, a new examination of hemostasis was conducted where activated partial thromboplastin time was in the normal range and the values of the D-dimer were 944.5 ngr/mL. On day 11 after the admission, the tests of hemostasis were again in the limits of normal, and the values of D-dimer decreased to 788.4 ngr/mL. The patient recovered pretty well and he was discharged home 12 days after the admission. He was given a recommendation for further check-ups by a transfusiologist and to make hemostasis controls. Also, he was advised to pay regular check-ups to our Clinic.

Discussion

Occlusion of the cerebral veins and dural venous sinuses may occur on the basis of local trauma, neoplasm or infection. Primary aseptic thrombosis may involve either cerebral veins or dural venous sinuses, or both in combination. Several authors have described some of the main angiographic changes in spontaneous thrombosis of the superior sagittal sinus. Arteriovenous shunting is a well-known phenomenon in cerebral infarction due to arterial occlusion. It seems reasonable to attribute the arteriovenous shunting which may occur in the presence of intracranial venous thrombosis to a similar process, namely the frequent pathologic finding of cerebral infarction. However, the pronounced, very local arteriovenous shunting into the patent part of the thrombosed cerebral vein (with associated collateral venous drainage) raises the possibility of some other mechanism. The clinical picture and the prognosis of intracranial venous thrombosis probably depend largely on the location, extent, and rapidity of development of the venous occlusion. The classical signs and symptoms of dural venous sinus thrombosis are not difficult to be recognized, but we suspect that cerebral venous thrombosis, particularly when localized and unaccompanied by sinus thrombosis, occurs much more frequently. Intracranial venous thrombosis should be suspected especially in patients with “stroke” of unexplained etiology. As illustrated by our case, papilledema may not be present, particularly during the early stages of the disease.

Conclusion

Superior sagittal sinus thrombosis is a condition that can be manifested with diverse and many symptoms and signs, which often can start un-
expectedly and can be life-threatening. Beside the rarity of this type of thrombosis, even more exceptional was this case of a male teenager. From the list of examinations that we conducted, we succeeded to discover that all of this was regarding to this specific condition, but we did not manage to discover the reasons that prompted this clinical condition. This might has developed to idiopathic thrombosis.

References


