Bacterial meningitis in neonates and infants – the sonographic picture

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Abstract

Bacterial meningitis is a major diagnostic and therapeutic problem among children and neonates, with severe, rapidly progressing course and potentially life-threatening complications. Early antibacterial treatment is essential for the patient’s favorable prognosis. Cerebral imaging plays an important role in the diagnostic process alongside physical examination and laboratory tests. Magnetic resonance imaging is the gold standard for diagnosing bacterial meningitis. Because of limited availability of magnetic resonance imaging, cranial ultrasound is the first imaging procedure to be performed (if the anterior fontanelle is preserved providing an adequate acoustic window). The safety and reliability of ultrasound examination, possibility to perform the examination at bedside without the need for sedation make cranial ultrasound a useful tool both for preliminary diagnostic investigation and for the monitoring of both treatment and long-term complications. Sonographic findings in patients with bacterial meningitis and possible complications are diverse. Changes can be seen on the surface of the brain, in the extra-axial space, in the ventricular system and in brain tissue. In some cases they can also be visible in the lumbosacral segment of the spinal cord. This paper presents ultrasound characteristics of lesions associated with bacterial meningitis in neonates and infants, based on the authors’ own material and data from the available literature.

Keywords

cranial ultrasound, neuroultrasound, meningitis, pediatrics, neonates

Introduction

Bacterial meningitis (BM) is a serious condition, presenting a major diagnostic and therapeutic challenge in everyday practice. Amongst the pediatric population, neonates (up to 4 weeks of age) and infants are particularly susceptible. The course of infection in this group is severe and usually systemic in nature. The lack of characteristic initial symptoms and rapid progression cause the mortality rate to exceed 50% (1,2).

Early diagnosis and initiation of correct treatment are key aspects for the patient’s prognosis. Apart from laboratory tests, imaging procedures are also important for the diagnostic process. Sonography should be especially appreciated among the available methods as the first modality used in diagnosing neonates and infants with suspected BM. Cranial ultrasound (CU) is also useful for monitoring the therapeutic process and the detection of early and late complications of BM.

This article presents the possibilities of using CU in the preliminary diagnostic investigation of BM and treatment monitoring in neonates and infants based on the authors’ own material and a review of the available literature.
Epidemiology

*Neisseria meningitidis* is the most common etiological factor for BM in the European countries. The incidence rate for neonates and infants is 20/100,000\(^3\). Data for the Polish population indicate that the most common etiological factor for BM in neonates are *Escherichia coli* and *Streptococcus agalactiae*, while in infants *Neisseria meningitidis* and *Streptococcus pneumoniae*. The combined prevalence of BM in the analyzed group is 15.71/100,000 (data from 2011)\(^2\).

Pathophysiology

BM is usually blood-borne, and most often associated with a systemic infection. From the clinical standpoint, it is important to confirm or exclude sepsis since it substantially compromises the patient’s prognosis. Initially bacteria accumulate in the lumen of choroid plexus capillaries located in the cerebral ventricles and in the small and medium-sized vessels of the subarachnoid space, where they induce an inflammatory response. Developing vasculitis leads to the inflammation of the arachnoid, dura mater and pia mater.

As a result of disruption to the blood-brain barrier, bacteria and inflammatory mediators (acute-phase proteins, leukotrienes, prostaglandins, among others) penetrate to the cerebrospinal fluid (CSF), thus exacerbating inflammation. In the acute phase cerebral vessels become dilated with increasing blood flow due to the influence of bacterial toxins. In later phases blood flow is reduced, which is associated with impaired vascular autoregulation\(^4\). The cytotoxic activity of oxygen radicals and proteolytic enzymes causes excessive sodium and water accumulation. This leads to impaired recirculation of CSF, dilation of the ventricular system and increased intracranial pressure, leading to brain edema\(^1,4–9\).

Sepsis can damage the endothelium of cerebral capillaries. This results in the formation of parietal thrombi. Severe cases involve diffuse cerebral venous thrombosis (affecting both superficial vessels such as superior sagittal sinus perforator vessels and deep veins such as the vein of Galen) or thrombosis in the confluence of sinuses\(^1,8\).

Vascular occlusion results in the development of infarction foci in deep structures (e.g. in the periventricular area) and in the subcortical regions, which leads to brain tissue damage. This process is accelerated by proteolytic enzymes which are stimulated by an inflammatory response.

Radiological abnormalities caused by encephalopathy are diverse, including focal hypoxic-ischemic lesions, areas of leukomalacia, hydrocephalus as well as focal and diffuse loss of brain tissue\(^5–10\).

Diagnostic imaging

Diagnostic imaging of BM involves the use of cranial ultrasound, computed tomography (CT) and magnetic res-
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onance imaging (MRI). CT and MRI are high-resolution modalities allowing full evaluation of cerebral structures, including the assessment of the posterior cranial fossa. Elaborate possibilities of vascular imaging in MRI and CT allow for early detection of thrombosis foci and fresh infarction in the deep structures of the brain, which are often not identifiable in conventional sonographic and color Doppler imaging. In addition, some lesions located in the extra-axial space, particularly in the temporal and parieto-occipital area may not be identifiable in CU. According to the current guidelines, MRI is the reference examination, the gold standard of diagnostic imaging in children. Performing CT and MRI requires sedation, which is associated with the risk of additional complications. Moreover, CT uses ionizing radiation, which is another risk factor. It is worth emphasizing that there is often no possibility of performing CT or MRI in some healthcare centers in Poland without transporting the patient to a different hospital or radiology unit. This delays the achievement of a clinical response. In the light of those facts CU seems to be the best alternative since it is easily accessible, can be performed at the point of care and is not associated with any adverse reactions. In the available literature many authors recommend CU as the first-line examination for diagnosing pediatric patients with suspected BM if a preserved anterior fontanelle provides an adequate acoustic window.

Changes on the brain surface

1. Dilated sulci with elevated echogenicity on the brain surface
The most common sonographic symptom associated with BM, which is found in approximately 60% of patients. Increased echogenicity is the consequence of the accumulation of inflammatory exudate in this area (Fig. 1).

2. Meningeal thickening
An enhancement in the form of a hyperechoic line can be visible on the surface of the cerebral gyri, which is formed by the pia mater and the adjacent arachnoid (the leptomeninges); this line becomes thickened in BM. Measurement is taken in the coronal plane through the anterior fontanelle, with the foramina of Monro visualized. The reference value of the meninx thickness is 0.7 mm, whereas thickening is identified with the value of 1.3 mm or higher (Fig. 2). It is also possible to measure sulcus thickness between the frontal gyri; the reference thickness of the sulcus is 1.2 mm and thickening is identified when this value is 2 mm or higher.

Ultrasound

Cranial ultrasound imaging can be unaltered in patients with uncomplicated BM. Ultrasound abnormalities are observed in approximately 65% of cases in the acute phase of the infection. In neonates and infants with severe neurological symptoms sonographic abnormalities are observed in up to 100% of patients.

There is a wide spectrum of possible abnormalities, which can be categorized depending on the location: pathologic changes visible on the surface of the brain, in the ventricular system, in deep structures – brain tissue, and in the lumbosacral segment of the vertebral canal.

Fig. 2. Thickening of the pia mater and the adjacent arachnoid – the leptomeninges layer

Fig. 3. A. Color Doppler imaging shows increased flow in the superficial cerebral vessels. In the superior sagittal sinus flow constriction associated with the presence of a parietal thrombus (arrows) is seen. B. Imaging through the anterior fontanelle in the longitudinal plane. One of the perforator veins is dilated with a surrounding hyperechoic extra-axial echo. A vessel partly covered by the color Doppler gate with a visible ostium into the superior sagittal sinus (star) is seen.
3. Dilation of cerebral vessels in the arachnoid
Dilation can be observed in the superficial cerebral veins, perforator vessels and the superior sagittal sinus, among others, which are easily accessible to imaging through the anterior fontanelle (Fig. 3).

4. Enlargement of extra-axial fluid spaces
The enlargement of the subdural and/or subarachnoid space is caused by the presence of inflammatory exudate (Fig. 4).

5. Closed fluid cavities
Closed fluid cavities in the extra-axial space (subdural, subarachnoid) take the form of hygromas or empyemas. It is also possible to visualize closed fluid spaces associated with the dissection of the dura mater itself. A hygroma filled with sterile fluid content is visible as a homogeneously anechoic area (Fig. 5). The presence of hygromas usually does not alter the prognosis\(^\text{15}\) – fewer than 5% become infected and turn into empyemas. In such cases CU shows higher or mixed echogenicity. Sometimes echogenic debris forms in the lumen of empyemas which by gravitation forms a pat-
tern with fluid levels on the brain surface and on the bottom of gyri. High amount of protein and fibrin in the infected fluid leads to their precipitation, organizing conglomerates of hyperechoic strands with smaller fluid cavities (Fig. 6). From the clinical standpoint, those CU findings are important because the fluid cavities may act as a bacterial reservoir and at the same time be the cause of therapeutic failures due to significantly limited drug penetration. In such cases it is necessary to administer a drug directly into the lesion's surroundings by subdural puncture (Fig. 7).

6. Cerebral thrombosis
Echogenic parietal thrombi are usually visualized with cranial ultrasound in the superior sagittal sinus as well as in larger deep veins (e.g. the vein of Galen). Color-coded Doppler imaging shows a characteristic impression of the blood flow shape being influenced by the thrombus. In cases of disseminated intravascular coagulation a substantial portion of the venous system can be affected, including the confluence of sinuses (Fig. 8).

Abnormalities of the ventricular system
1. Ventricular system dilation
Ventricular system dilation can be found in approximately 30% of patients with BM. Sonographic hallmarks of ventricular dilation are diverse, can be symmetrical, involv-
Changes can be inconspicuous, such as rounded impressions in the anterior horns of the lateral ventricles, or very severe and classified as hydrocephalus. The most sensitive sonographic sign of hydrocephalus is the dilation of the third ventricle or the temporal horns of the lateral ventricles (11) (Fig. 9, 10).

2. Elevated CSF echogenicity in the ventricles with visible fluid levels

3. Dilation of choroid plexuses with increased echogenicity

4. Ventriculitis
Sonographic signs of ventriculitis include: thickened and irregular contours of the ventricular system, increased ventricular lining echogenicity, increased periventricular white matter echogenicity, irregular contours of choroid plexuses, increased CSF echogenicity inside the ventricles, intraventricular debris and compartmentalized ventricular lumen containing fluid cavities (Fig. 11).

Brain tissue abnormalities

1. Brain edema
One of the most severe complications of BM is brain edema. Its sonographic appearance involves increased echogenicity of sulci and brain tissue. Advanced brain edema presents as obsfuscation of the boundaries between gyri and sulci as well as compression and deformation of the ventricular system and the remaining fluid cavities. Color Doppler imaging reveals an increased resistance index (RI) (> 0.8) measured in afferent arteries, e.g. in the anterior cerebral artery (9) (Fig. 12).

2. Hypoxic-ischemic foci
Pathologic changes in this group are diverse. They may be seen as homogeneous hyperechoic areas or, in advanced disease, as heterogeneous malacia lesions with fluid spaces (Fig. 13, 14).

3. Subcortical infarction foci
Infarction foci develop in brain tissue as a result of thrombosis. In the acute phase they are initially visible as areas of increased echogenicity with evident obsfuscation of the echostucture. As a result of ischemia, local necrosis develops and echogenicity decreases in the center of the hyperechoic area with restricted vascular flow, which is visible in color Doppler imaging. After the acute phase recedes, leukomalacia lesions develop in the affected subcortical area (Fig. 10).
Imaging of the lumbosacral spine

Ultrasound imaging of the vertebral canal in the lumbosacral area is used primarily when developmental spinal cord pathology is suspected. It is considered to be the procedure of choice in neonates and infants. The ossification process of vertebral arches and spinous processes lasts until 12 months of age, which provides an adequate acoustic window for ultrasound examination. In older patients the utility of ultrasound examination of this area is limited. Imaging is performed with both convex and linear transducers. The examination is usually used for: the evaluation of spinal cord anatomy with the analysis of potential developmental variants (filar cyst, terminal ventricle), assessment of structural mobility in the vertebral canal (to exclude signs of spinal cord impaction) and examination of soft tissue and osseous structures surrounding the spinal cord in order to exclude potential developmental pathologies (myelomeningocele, spina bifida, skin sinuses/fistulas).

Changes associated with BM observed on the brain surface can also be partially visible in the lumbosacral segment of the spinal cord. This is of high clinical importance for making the diagnosis, planning lumbosacral puncture and predicting long-term complications. The presence of echogenic debris in the subdural space increases the risk of progressive hydrocephalus (Fig. 15).

Conclusion

Cranial ultrasound is a useful diagnostic method when bacterial meningitis is suspected in neonates and infants. The spectrum of characteristic signs visualized in BM using conventional imaging and Doppler imaging allows for quick preliminary diagnosis and initiation of treatment, which has a significant impact on the patient’s prognosis. The possibility of brain imaging without any
adverse reactions and performing the procedure at the point of care makes cranial ultrasound an ideal tool not only for the preliminary diagnosis of patients with suspected BM, but also for treatment monitoring and identification of long-term complications of the disease.

Conflict of interest

The authors do not report any financial or personal affiliations to persons or organizations that could negatively affect the content of or claim to have rights to this publication.

References

1. Albrecht P, Hryniewicz W, Kuch A, Przyjałkowski W, Skoczynska A, Szenborn L: Rekomendacje postępowania w zakażeniach bakteryjnych ośrodkowego układu nerwowego. Narodowy Instytut Leków, Warszawa 2011.
2. Skoczynska A, Wasko I, Kuch A, Kadłubowski M, Gołębiewska A, Foryś M et al.: A decade of invasive meningococcal disease surveillance in Poland. PLoS One 2013; 8: e71943.
3. Harrison LH, Trotter CL, Ramsay ME: Global epidemiology of meningococcal disease. Vaccine 2009; 27 (Suppl. 2): B51–B63.
4. Ashwal S, Stringer W, Tomasi L, Schneider S, Thompson J, Perkin R: Cerebral blood flow and carbon dioxide reactivity in children with bacterial meningitis. J Pediatr 1990; 117: 523–530.
5. Berman PH, Banker BQ: Neonatal meningitis. A clinical and pathological study of 29 cases. Pediatrics 1966; 38: 6–24.
6. Tunkel AR, Scheld WM: Pathogenesis and pathophysiology of bacterial meningitis. Annu Rev Med 1993; 44: 103–120.
7. Pfister HW, Fontana A, Täuber MG, Tomasz A, Scheld WM: Mechanisms of brain injury in bacterial meningitis: Workshop summary. Clin Infect Dis 1994; 19: 463–479.
8. DiNubile MJ, Boom WH, Southwick FS: Septic cortical thrombophlebitis. J Infect Dis 1990; 161: 1216–1220.
9. Tunkel AR, Scheld WM: Alterations of the blood-brain barrier in bacterial meningitis: in vivo and in vitro models. Pediatr Infect Dis J 1989; 8: 911–913.
10. Babcock DS, Han BK: Sonographic recognition of gyral infarction in meningitis. AJR Am J Roentgenol 1985; 144: 833–836.
11. Hughes DC, Raghavan A, Mordekar SR, Griffiths PD, Connolly DJ: Role of imaging in the diagnosis of acute bacterial meningitis and its complications. Postgrad Med J 2010; 86: 478–485.
12. Yikilmaz A, Taylor GA: Sonographic findings in bacterial meningitis in neonates and young infants. Pediatric Radiol 2008; 38: 129–137.
13. Mahajan R, Lodha A, Anand R, Patwari AK, Anand VK, Garg DP: Cranial sonography in bacterial meningitis. Indian Pediatr 1995; 32: 989–993.
14. Jéquier S, Jéquier JC: Sonographic nomogram of the leptomeninges (pia-glia plate) and its usefulness for evaluating bacterial meningitis in infants. AJNR Am J Neuroradiol 1999; 20: 1359-1364.
15. Barkovich AJ: Infections of Central Nervous System. In: Barkovich AJ (ed.): Pediatric Neuroimaging. Lippincott Williams & Wilkins, Philadelphia 2008: 801–868.
16. Rudas G, Almássy Z, Papp B, Varga E, Méder U, Taylor GA: Echodense spinal subarachnoid space in neonates with progressive ventricular dilatation: a marker of noncommunicating hydrocephalus. AJR Am J Roentgenol 1998; 171: 1119–1121.