Asymmetric optic nerve sheath diameter as an outcome factor following cranioplasty in ‘syndrome of the trephined’ patients

Assimetria do diâmetro da bainha do nervo óptico como fator prognóstico após cranioplastia em pacientes portadores da ‘Síndrome do Trefinado’

ABSTRACT

Background: Decompressive craniectomy (DC) is gaining an increasing role in the neurosurgical treatment of intractable intracranial hypertension, but not without complications. A rare complication is the “syndrome of the trephined” (ST). It occurs when the forces of gravity overwhelm intracranial pressures, leading the brain to become sunken. Methods: We followed-up 5 patients submitted to DC and diagnosed with ST. All were submitted to brain MRI to calculate the optic nerve sheath diameter (ONSD). Our goal was to determine the usefulness of asymmetric ONSD as an outcome factor after cranioplasty. Results: Only two patients presented an asymmetric ONSD, being ONSD larger at the site of craniectomy. Surprisingly these patients had a marked neurological improvement after cranioplasty. They became independent a week after and statistically earlier than others. Conclusion: It is presumed that the presence of an asymmetric ONSD in trephined patients is an independent factor of good outcome after cranioplasty.

Keywords: decompressive craniectomy, syndrome of the trephined.

INTRODUCTION

Decompressive craniectomy is gaining an increasing role in the neurosurgical treatment of intractable intracranial hypertension in patients with head injury, acute stroke, and severe brain edema. Though technically straightforward, the procedure is not without significant complications. Herniation through the cranial defect, seizures and subdural effusion are common early complications after decompressive craniectomy, whilst hydrocephalus is the most common late complication. Others rare late complications are sinking skin flaps, paradoxical herniation and external brain tamponade.

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During the late phases of recovery, patients may develop a new cognitive, neurological, or psychological deficit termed “syndrome of the trephined” (ST)\(^1\). This syndrome clinically may include dizziness, tinnitus, orthostatic headaches\(^1\), central sleep apnea\(^1\), dysarthria, limb rigidity, parkinsonian levodopa-resistant tremor, diplopia (abducens nerve palsy)\(^2,11\), reversible monoparesis\(^1\), fatigue and fluctuating levels of consciousness up to coma states\(^2,11,16\). It may occur following a unilateral craniectomy or a bilateral bifrontal craniectomy\(^1\).

Increasing the intracranial pressure by placing the patient into Trendelenburg position or increasing hydration may reverse the symptoms\(^3,4\). However up to now there is no clinical or radiological signal responsible to the diagnosis of the “syndrome of trephined” (ST).

In some patients harboring the ST, sinking skin flaps, paradoxical herniation and external brain tamponade may be seen\(^1\). Paradoxical herniation consists of brain midline shift up to the contralateral side from the craniectomy, easily visualized by head CT or brain MRI, and may suggest the diagnosis.

Cerebral blood flow changes, the effect of the atmospheric pressure on the brain, as well as cerebrospinal fluid (CSF) hydrodynamic changes have been postulated as the possible reasons for this syndrome\(^6,7\). A moderate increase in venous outflow as well as a two-fold increase in cranio-caudal cerebrospinal fluid systolic flow velocity has already been measured after the skull closure, reaffirming the hydrodynamic hypothesis\(^6\).

### METHODS AND OBJECTIVE

We followed-up 5 patients submitted to decompressive craniectomy, secondary to head injury or acute stroke, up to 6 months postoperatively, among whom was noted altered levels of consciousness, sinking skin flap at the site of craniectomy, or paradoxical herniation. All patients were operated from 2008 to 2012 at our hospital by the same surgical crew.

All trephined patients who died or evolved without symptoms or signs compatible to ST were excluded from this patient case series. Epidemiological data, motive to decompression, time to decompressive craniectomy, symptoms and signs compatible to ST, and time to cranioplasty may be seen in Table I.

All patients included in this study were routinely submitted to laboratory essays, radiographic exams and video-electroencephalography to exclude others causes of neurological impairment. All were submitted to brain MRI scan, with small coronal fat-suppression cuts, in order to calculate the optic nerve sheath diameter, before and after cranioplasty. Cranioplasty was routinely planned after 3 months from decompression.

Our goal was to determine the usefulness of the asymmetric optic nerve sheath diameter in craniectomy patients as a diagnostic signal and as a good outcome factor after cranioplasty.

### RESULTS

Among these 5 patients, all have developed neurological deterioration after unilateral decompressive craniectomy at a mean follow-up of 6 months, including orthostatic headache, diplopia, contralateral hemiparesis, gait ataxia, and excessive sleepiness up to periods of minimal consciousness. All patients had their symptoms reversed by placing then into cranio-caudal cerebrospinal fluid systolic flow velocity has already been measured after the skull closure, reaffirming the hydrodynamic hypothesis.
Trendelenburg position. At ectoscopy all patients presented sinking skin flap at the site of craniectomy. On radiologic examination all presented paradoxical herniation.

After ruling out all other possible disorders, all patients have been presumably diagnosed with ST.

However, on MRI scanning only two patients (patients B and E) presented an asymmetric optic nerve sheath diameter (ONSD), being the ONSD larger at the side of craniectomy. Surprisingly these two patients had a marked neurological improvement after cranioplasty, with normalization of the ONSD and disappearance of the brain shift a day after procedure (Figure 2), becoming independent a week after procedure.

The others three patients (patients A, C and D), who had a symmetric ONSD at MRI, have also improved their consciousness level, but in a very slowly way, becoming independent after a few months.

Patients harboring the ST with symmetric ONSD needed statistically more time to become independent after cranioplasty than patients with asymmetric ONSD (mean time of 38 days vs 8 days, t-test, p<0.05).

**Table 2.** Optic nerve sheath diameter (ONSD) at the side of cranioplasty from patients harboring the “syndrome of the trephined”, pre and post cranioplasty, and Glasgow Outcome Scale (GOS) one week and two months after cranioplasty

<table>
<thead>
<tr>
<th>Patient</th>
<th>ONSD pre</th>
<th>ONSD post</th>
<th>GOS 1-w</th>
<th>GOS 2-m</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>5.6mm</td>
<td>5.6mm</td>
<td>3</td>
<td>4</td>
</tr>
<tr>
<td>B</td>
<td>6.8mm</td>
<td>5.8mm</td>
<td>5</td>
<td>5</td>
</tr>
<tr>
<td>C</td>
<td>4.8mm</td>
<td>4.8mm</td>
<td>4</td>
<td>5</td>
</tr>
<tr>
<td>D</td>
<td>5.4mm</td>
<td>5.4mm</td>
<td>3</td>
<td>5</td>
</tr>
<tr>
<td>E</td>
<td>6.4mm</td>
<td>5.6mm</td>
<td>5</td>
<td>5</td>
</tr>
</tbody>
</table>

Abbreviations: ONSD=optic nerve sheath diameter; GOS=Glasgow Outcome Scale; w=week; m=months.

**Figure 1:** A-B) Axial T1-weighted MRI, FLAIR sequence, showing sinking skin flap at the left side of craniectomy, paradoxical herniation to the right side, and subfalcine herniation; (C) Axial T1 MRI, cerebral sulci effacement at the left side; (D-E) Axial T2-weighted MRI, asymmetry of the optic nerve sheath diameter (ONSD), being larger at the left side (6.8mm vs right side (5.8mm)); (F) Coronal fat-suppressed T2-weighted MRI, asymmetric ONSD.

**Figure 2:** Axial brain CT one day after methyl methacrylate cranioplasty of the trephined patient from Figure 1. (A-B) Disappearance of the brain shift and subfalcine herniation; (C) Recovery from the cerebral sulci effacement.

**Discussion**

It is well known the usefulness of decompressive craniectomy in the treatment of refractory intracranial pressure, but not without significant complications. One of the harmful complications is surely the “syndrome of the trephined”. Up to now there was no diagnostic tool or prognostic factor capable to make a tailored diagnosis and to predict a good outcome.

The measurement of the ONSD is an important indirect method to suppose or diagnose increased levels of intracranial pressure. It is a cheap, non-invasive, fast and reliable method of diagnosis, with the same accuracy of other expensive methods (dandy-walker, encephalography, angio-encephalography). It is an excellent tool to evaluate shunt function in hydrocephalus and to diagnose benign intracranial hypertension. However, it is not a perfect method. The ONSD is not a very specific test and is not able to differentiate between the normal and the pathological ONSD. For example, it is possible to have a normal ONSD in a pathological condition like subarachnoid hemorrhage, and vice versa. The ONSD is also affected by other factors like age, gender, and height. For these reasons, the ONSD should be used in combination with other methods to make a diagnosis.

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pressure (ICP). The ultrasonographic visualization of the optic nerve is feasible, costless, and world-wide applied to diagnose intracranial hypertension5,8.

By its time, the use of MRI to detect raised ICP is a more recent method. It seems to bring more precision to the measurement of the ONSD. ONSD is strongly related to ICP, a finding that reflects distension of the nerve sheath during increases in CSF pressure. On T2-weighted sequences, CSF exhibits a high signal (white), whilst fat and gray matter appear as light grey18. The perioptic CSF is surrounded by orbital fat. Contrast between CSF and orbital fat can be improved with fat suppression, increasing the image resolution for the ONSD measurement10,18.

The most useful clinical message derived from current data may be summarized as followed: an ONSD less than 5.3mm is unlikely to be associated with raised ICP; an ONSD above 5.82mm indicates that the probability of raised ICP is 90%10,18.

In our series of trephined patients, those that presented ONSD asymmetry (patients B and E) had at the side of craniectomy an ONSD greater than 6mm (strongly suggesting raised ICP), and had their diameter decreased to normal values one day after cranioplasty, see Table II. Surprisingly those two patients evolved better than patients without ONSD asymmetry, becoming independent (Glasgow Outcome Scale 5) just one week after cranioplasty (Table 2).

That ONSD asymmetry may be explained by local changes of the CSF hydrodynamics, resulting in distension of the nerve sheath ipsilateral to the craniectomy side. This distension by its turn may reflect that the intracranial pressures finally have been overwhelmed by the forces of atmospheric pressure and gravity. Correction of this distension just after cranioplasty may indicate the recovery from the CSF hydrodynamics and normalization of intracranial pressures from a cranium converted back to a ‘closed box’ system.

CONCLUSION

The “syndrome of trephined” is a rare late complication after decompressive craniectomy. It is presumed that the presence of an asymmetric ONSD at MRI of trephined patients is an independent factor of good outcome after cranioplasty.

REFERENCES

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