Clinical spectrum of callosum corpus splenium lesions: subdiagnostics of a common entity

Abstract

The corpus callosum is the largest white matter commissure of the brain that communicates both cerebral hemispheres; and can be altered by multiples etiologies; some of them, with a pathophysiological mechanism not elucidated. There are some reports, about splenium callosum lesions, associated to nonspecific clinical condition, but with a common characteristic: disturbance of consciousness and wakefulness as the predominant symptom; although, also have been described seizures, headache or athetosis in some reports. Independent of previous features, this type of lesions, may be evidenced in asymptomatic patients, as a radiological finding. We present three cases of patients with these radiological features, in two of them which a common finding, was a uremic syndrome that does not required chronic dialysis. We concluded that in patients with transient impairment of consciousness of unclear origin, a callosum splenium lesion would to be suspect, especially if they are associated to uremic syndrome. Also, that these types of findings could be more frequent than expected, so more studies are necessary to clarify the physiopathology.

Keywords: callosum corpus lesions, consciousness disturbance, splenium callosum lesions, mild encephalitis/encephalopathy

Introduction

The callosum corpus (CC) is the largest white matter commissure of the brain that communicates both hemispheres. Anatomically, can be divided in 3 portions: anterior, middle and posterior; two first that depends of anterior circulation; and the splenius of the corpus callosum of posterior respectively. The most frequent causes of injury of CC are cerebrovascular stroke, cranial brain trauma, central nervous system infections, alcohol, epilepsy, multiple sclerosis and MERS syndrome (mild encephalitis/encephalopathy with a reversible isolated splenium callosum corpus lesion). Pekala et al. suggest to include patients who are or have undergone oncologic radiotherapy of the head and neck.

The MERS’s clinic is variable. Confusion or alteration of consciousness is the most frequent clinical sign. Another study considers fever and headache as a predominant symptomatology. Both mention dysarthria, seizures, hemiparesis and associated ataxias. And recently, athetosis was described by one of the authors. It’s important to consider that in elderly patients, it is possible to find a silent hypodense lesion in the splenius of the corpus callosum as radiological findings; as well it has been mentioned in children; probably directly related to the immaturity and involution of the central nervous system related to a fragility of the blood-brain barrier (BBB). We communicate 3 patients hospitalized in our Neurology Unit (NU) between July and October 2017, due to confusional syndrome in two of them, and seizures in the remainder; and in which, neuroimaging studies showed a unique hypodensity in the splenius of the corpus callosum.

Case presentation

Case 1

Male, of 65 years old, with diabetes mellitus, and hypertension both with poor treatment; suspended smoking and occasional alcohol consumption who has a emergency department (ED) consultation by 10 days of fluctuating confusional syndrome associated with headache, psychomotor agitation and gait instability. Is initially evaluated in the ED, with blood pressure level (231/122 mmHg), hemoglucotest (448 mg/dl); and physical examination that evidenced a arousal patient, time and place disoriented, and without focal neurological deficits. Laboratory tests showed a serum creatinine level in 3.29, blood urea nitrogen 44, glycemia 393, Ph 7.33, HCO3, 19, Na 142, K 4.3. Computer tomography (CT) was informed as a microangiopathy of the supratentorial white matter and a left thalamic sequela stroke. In the NU, the patient evolves with more disturbance of consciousness and acute renal failure that required hemodialysis (HD) and evolve with a septicemia of pulmonary origin. Studies were complemented with echocardiogram (ECHOC), lumbar puncture (LP) and electroencephalogram (EEG); all they normal. Because the persistence of the symptoms, new CT shows a hypodensity in the splenius of the corpus callosum, and in a second look this finding was present in the first scanner (Figure 1) (Table 1); and a diagnosis of MERS syndrome was made.

Case 2

A 60-year-old woman with diabetes mellitus type 2, hypertension, chronic obstructive pulmonary disease, and suspended smoking habit; with the previous antecedent of a recent hospitalization for...
digestive tract hemorrhage, and paroxystic atrial fibrillation in oral anticoagulant treatment with warfarin. One month later of this previous clinical antecedent, the patients come to ED by focal seizures auto-limited, without loss of consciousness; which was followed by left face arm paresis that reverted after hours. A Todd’s phenomenon was diagnosed in ED. EEG normal, laboratory tests on the ED show a hematocrit in 26%, a hemoglobin in 7.5, a platelet in 782,000, INR 1.29; and Na 135 mEq, and K 2.9 mEq. All corrective treatment of this findings, was start in ED. CT did show a mild calcified atheromatosis in both carotid siphons in computed tomography angiography; and a hypodensity in the splenius of the corpus callosum (Figure 2), (Table 1). The diagnosis of MERS was made.

### Table 1 Our clinical cases, co morbidities and relevant data

<table>
<thead>
<tr>
<th>Case</th>
<th>Sex/Age</th>
<th>Comorbidities</th>
<th>Symptomatology</th>
<th>Neurological examination in El</th>
<th>Exams (highlight)</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>65 - Male</td>
<td>Diabetes Mellitus, Hypertension</td>
<td>Consciousness disturbance, Headache</td>
<td>High Stood Pressure 231/122 mmHg, Capillary glycemic 448 mg/dl</td>
<td>Cr 2.9 mg/dl, BUN 44 mg/dl</td>
<td>Acute renal failure, Pulmonary Infection</td>
</tr>
<tr>
<td>2</td>
<td>60 - Female</td>
<td>Diabetes Mellitus, Hypertension</td>
<td>Seizures, Todd's paresis</td>
<td>Attentive, Todd's phenomenon</td>
<td>ECG (-), ECHOC (-), EEG (-), Hto 20%, Hb 7.5 nigicell, Platelets 782,000 nig/111, INIt 1.29</td>
<td>Blood Transfusion, Asymptomatic</td>
</tr>
<tr>
<td>3</td>
<td>64 - Female</td>
<td>Hypertension</td>
<td>Consciousness disturbance, Psychomotor Agitation</td>
<td>Agitation, Left Hemiplegia</td>
<td>EEG: encephalopathy pattern, ECG (-)</td>
<td>Good response to benzodiazepines and quetiapine, Neurorehabilitation</td>
</tr>
</tbody>
</table>

HD, hemodialysis; ECG, electrocardiogram; ECHOC, echocardiogram; LP, lumbar puncture; EEG, electroencephalogram; (-), None pathologic event

![Figure 1 Sagittal image showing hypodense lesion in the splenium callosum corpus.](image)

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Clinical spectrum of callosum corpus splenium lesions: subdiagnostics of a common entity

A 64-year-old female; with hypertension and smoking habit of 15 cigarettes per day; did consultation to the ED by a disturbance of consciousness of 24 to 36 hours with severe psychomotor agitation. The first ED evaluation described a vigil patient, with dysarthria, delusional ideas and left hemiplegia. Is evaluated by psychiatry in ED, whom request CT without acute lesion and serum laboratory that show hematocrit in 29%, red blood cell count in 3’200’000, electrolytes and creatinkinasa normal; but creatinine in 6 and BUN in 56. A diagnosis of acute over chronic renal disease is made. In ED, patient requires high doses of benzodiazepines for agitation, by this reason, is referral to intensive care unit, where it is sedated with propofol. It evolves with difficulty to arousal, with incoherent language, and delusional ideas. A treatment whith Quetiapine is initiates at dose of 300mg per day, with reversion of the symptoms. A LP was made, with cerebrospinal fluid (CSF) normal; and a EEG was performed, that shows encephalopathic pattern. Is studied again with cerebral tomography, which shows hypodense lesion in splenius of the corpus callosum (Figure 3) (Table1). The patient is transferred to the NU, whit clinical improvement; and in a posterior cerebral tomography the lesion was not present.

Discussion

Previous studies, classify the splenial callosum lesions topographically in isolated and multiple lesions. The first of them are most frequent and with best prognosis. Because the clinic is non-specific, and the etiology is unclear; it is expected that the pathophysiological mechanism of this type of lesion are not elucidated yet. There are many hypotheses; like intramielinic edema, axonal damage, hyponatremia, hyperglycemia, oxidative stress, malnutrition among others.1,3–7

In cases where there is an associated systemic infection; it’s postulated that the pathophysiological mechanism would be to a myelinic axonal damage, produced either by the antigen itself, or due the antibodies reaction against the antigen that alters the permeability of the BBB generating a focal edema.1,3 Considering that our case patient 01 was infected, this could have been one of the mechanisms that can caused the hypodense in the CC, as suggest Yuan et al.3

Also, the literature mentions some cases attributed to antiepileptic drug toxicity.1,7,8 In consequence the manifestations of LHECC are variable; and is difficult to evaluate because the large number of comorbidities. Also these type of lesions can be found in asymptomatic individuals. Polster,6 found these lesions in 3 asymptomatic individuals, who were to undergo neurosurgical processes due to seizures refractory to medical treatment. Pekala,4 found these
lesions in asymptomatic, elderly individuals with no relevant morbid antecedents. Clinical cases 1 and 3, both lesions were described in the evaluation of a confusional syndrome; while case 2 in a focal seizure, followed by self-limited abrupt left hemiparesis. In both patient: 2 and 3; there were found focal neurologic deficit at the ED evaluation (Figure 3).

It is necessary to clarify that the lesions in the first instance were not identifiable in any of our cases; probably due to the poor image resolution that the CT vs MRI used in the majority of previous reports; and despite this considerations; other authors agree on the difficulty of evaluating the splenius lesion of the corpus callosum even using MRI.1-4,8-10

Another conditions in our patients; was the many comorbidities associated, with a neurological examination without neurological focus that would to delay the diagnosis. In the cases 01 and 03, there was an acute renal disease, and the needs of hemodialysis in one of them; this finding, could presume if this type of lesions would be secondary to uremia. This is a question that would require more studies of series of patients with chronic kidney disease, hemodialysis and uremic syndrome; but has been previously reported in patients with hemolytic-uremic syndrome.11 In none of our patients, a chronic diayals was indicated.

We propose that a LHECC has a clinical relevance when is concomitant to a confusional syndrome or disturbance of consciousness, as Min-Keun Park1 and Junliang2 and Juan postulate in their respective publications;3,4 and was evident in cases 01 and 03.

<table>
<thead>
<tr>
<th>Table 2 Description of Lesions involving the corpus callosum at CT</th>
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<tbody>
<tr>
<td>Glioblastoma</td>
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<td>Multiforme</td>
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<tr>
<td>Lymphoma</td>
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<td>Meningioma</td>
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<tr>
<td>Metastasis</td>
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<td>Multiple Sclerosis</td>
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<tr>
<td>Head Trauma</td>
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<tr>
<td>Hypoxemic Ischemic</td>
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<tr>
<td>Encephalopathy</td>
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<tr>
<td>Marchiafava - Bigman</td>
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<tr>
<td>Arteriosvenous</td>
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<tr>
<td>Malformation</td>
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</table>

On the other hand; like case 02 show, the clinically presentation of Todd phenomenon without unconsciousness forced to do differential diagnosis with a transient ischemic attack, especially in the context of a probably hypoperfusion with severe active anemia, thrombocytosis reactive or non-anticoagulated paroxystic atrial fibrillation as our patient had.

Due to the increasingly frequent neuroimaging of higher and better resolution, this type of lesions is expecte to be identified more frequently. Previous studies4,9 show that these lesions are more frequent than expected. Chrysikopoulos8 performed a retrospective study of 352 patients with neuroimaging; and did found 28 cases with lesions in the corpus callosum, in greater proportion in the splenium. Pekala,4 nevertheless questions why the splenium callosum lesions are being reported in more frequency nowadays and not before despite the neuroimaging are available years ago?.

Finally; we also mentioned the importance of evaluating the images not only in axial plane, but also sagittal to be able to show structures and lesions that are easily altered by artefacts or simply not evidence at the axial one. Because our NU doesn’t have MRI, we affirm due the high frequency found in consecutive months, that the splenius lesions of the corpus callosum are more common than expected, and should be a good practice try find them patients with the characteristics described above and reported in the literature; and keep in mind the differential diagnoses of corpus callosum lesions evidenced in CT (Table 2). These can be of variable etiology and have different presentations.13

**Conclusion**

The LHECC, has multifactorial etiologies, without a clear mechanism, but usually, they are presented in relation to important comorbidities of the patients, mainly chronic non-transmissible diseases with poor metabolic control; and could be secondary to a uremic syndrome. Not all lesions in the splenius of the corpus callosum are clinically significant; so it is important to correlate them with the symptomatology; especially if there are found in relation to consciousness disturbance and confusion; and in these cases, keep in mind to evaluate the sagittal sections of the cerebral neuroimaging. Finally, and as other reports conclude, we also propose that these lesions are much more frequent than previously described in the literature.

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To the authorities of our hospital, who supported our initiative.
To our families who always believe in us.

**Conflict of interest**

The authors declare not to have any financial interest or any conflict of interest in the realization of this publication.

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Patient consent form

All patients signed an informed consent form to authorize the use of the dates in this publication, assuring them the confidentiality of their identities.

References


