Dysembryoplastic Neuroepithelial Tumor (DNT): Morphological and Immunohistochemical Features

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\textbf{Abstract.} Dysembryoplastic neuroepithelial tumor is a rare mixed neuronal-glial tumor. The recognition and correct diagnosis of dysembryoplastic neuroepithelial tumor is important because this tumor is curable by excision. The records of the Pathology Department at King Faisal Specialist Hospital and Research Center, Jeddah, Saudi Arabia, were reviewed for cases that were histologically diagnosed as dysembryoplastic neuroepithelial tumor between 2000 and 2007; five cases were found. The clinical, radiological, histological and immunohistochemical findings were reviewed. The ages ranged between 9 and 25 years. There were 2 males and 3 females. All patients were diagnosed with epilepsy. Three patients had temporal tumor and the other two had frontal tumors. Radiological evaluation showed non-enhancing mass without mass effect or vasogenic edema. All five cases were characteristically composed of small round oligodendroglia-like cells, astrocytes and mature neurones in varying proportions. Immunocytochemistry for glial fibrillary acidic protein and neuronal markers (neuron-specific enolase; and synaptophysin) highlights the astrocytic and neuronal components. The recognition of this tumor is very important because it is a surgically curable lesion with excellent prognosis. Pathologists need to be familiar with the characteristic histopathological findings to avoid unnecessary overtreatment by radiotherapy or chemotherapy.

\textbf{Keywords:} Dysembryoplastic neuroepithelial tumor, Gliomas, Neural-glial tumor.
Introduction

Dysembryoplastic neuroepithelial tumors (DNT) are rare mixed neuronal-glial tumors that should be differentiated from low-grade gliomas. They are usually supratentorial tumors occurring in children and young adult males are more frequently affected\textsuperscript{[1]}. The recognition and correct diagnosis of DNT is important because this tumor is curable by excision. In the recent WHO classification, DNT are considered a frequent cause of drug-resistant epilepsy\textsuperscript{[1]}. The incidence in specimens resected from patients treated surgically for long-standing epilepsy varies, ranging from 12.6 to 56.3\% in the large series\textsuperscript{[2,3]}.

Methods

The records of the Pathology Department at King Faisal Specialist Hospital and Research Center (KFSH&RC), Jeddah, Saudi Arabia (KSA), were reviewed for cases that were histologically diagnosed as DNT. The search covered the period between 2000 and 2007. KFSH&RC is the main hospital in KSA for surgical treatment of epilepsy. Five cases were found. The clinical, radiological, histological and immunohistochemical findings were reviewed. Immunocytochemistry study was performed for glial fibrillary acidic protein (GFAP) (Dako, dilution: 1/100), neuron-specific enolase (Cell marque, dilution 1/250), synaptophysin (Cell marque, dilution 1/500), and neurofilament (Cell marque, dilution 1/200).

Results

The ages ranged between 9 and 25 years. There were 2 males and 3 females. All patients were diagnosed with epilepsy. Three patients had the tumor in the temporal lobe and two in the frontal lobe. Radiological evaluation using magnetic resonance imaging (MRI) and computerised tomography (CT scan) showed non-enhancing tumors without mass effect or vasogenic edema. Calcification was identified in two cases. The clinical presentation and radiological findings are summarised in Table 1.
Table 1. Summary of the clinical and MRI results of the patients.

<table>
<thead>
<tr>
<th>Patients</th>
<th>Age/sex</th>
<th>Clinical presentation</th>
<th>Radiology</th>
<th>Follow-up period</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>12Y/F</td>
<td>Epilepsy with right temporal tumor</td>
<td>Tumor measures 3.5 cm mass with cystic component and without enhancement. No mass effect or vasogenic edema.</td>
<td>7 years. Patient developed a new lesion 2 years after initial diagnosis with similar histology. She had no other recurrence over the last 5 years</td>
</tr>
<tr>
<td>2</td>
<td>25Y/M</td>
<td>Epilepsy with right frontal tumor</td>
<td>Tumor measures 4.2 cm mass without enhancement with calcification. No mass effect or vasogenic edema. It has hyperintensity on T2.</td>
<td>3½ years No recurrence</td>
</tr>
<tr>
<td>3</td>
<td>9Y/F</td>
<td>Epilepsy with left frontal tumor</td>
<td>Tumor measures 5 cm without enhancement. There is calcification and scalloping effect on the inner table. No mass effect or vasogenic edema. It has hyperintensity on T2.</td>
<td>1½ years No recurrence</td>
</tr>
<tr>
<td>4</td>
<td>14Y/F</td>
<td>Epilepsy with right temporal tumor</td>
<td>Tumor measures 5.3 cm mass with cystic component and without enhancement. There is calcification No mass effect or vasogenic edema.</td>
<td>2 years. No recurrence</td>
</tr>
<tr>
<td>5</td>
<td>15Y/M</td>
<td>Epilepsy with left temporal tumor</td>
<td>Tumor measures 3.4 cm mass without enhancement. No mass effect or vasogenic edema. It has hyperintensity on T2.</td>
<td>1 year. No recurrence</td>
</tr>
</tbody>
</table>

The cases were characteristically composed of small round oligodendroglial-like cells (OLC), astrocytes and mature neurones in varying proportions (Fig. 1a-c). Immunocytochemistry for GFAP and neuronal markers (neuron-specific enolase, synaptophysin (Fig. 1d) and neurofilament) confirm the diagnosis and highlight the astrocytic and
neuronal components. The response to surgery was excellent. One patient presented with a new lesion 2 years later with similar histological features; none of the other tumors have recurred, and the control of seizures remained good.

Fig. 1. a) Section from one of the tumors reveals mixed cells with mucoid areas. (Hematoxylen and Eosin stain, original power ×100). b) A higher power reveals mainly small round oligodendroglia-like cells (OLC) with perinuclear halos (Hematoxylen and Eosin stain, original power ×200). c) A higher power reveals a mixed pattern composed of oligodendroglia-like cells (OLC), astrocytes and scattered mature neurones (Hematoxylen and Eosin stain, original power ×400). d) Immunohistochemistry stain for synaptophysin reveals positive staining in neuronal cells.

Discussion

King Faisal Specialist Hospital and Research Center in Jeddah is considered one of the main hospitals in KSA for surgery of epilepsy. Annually more than 30 surgical procedures are performed at this institution for the treatment of epilepsy. We recognized five cases of histologically proven DNT among those resections performed for treatment of epilepsy. DNT is now a well-known, seizure-producing entity that occurs characteristically in the cerebral cortex of children and young adults\(^{[4-12]}\). Daumas-Duport et al.\(^{[9]}\) described DNT as having the
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following features: (1) intracortical location; (2) partially nodular structure; (3) mixed glial and neuronal composition, always with the so-called 'specific' glioneuronal element and finally, (4) the tumors were usually associated with some form of cortical dysplasia in the adjacent cortex. Additional reports have confirmed these features\cite{4,6,7}. Our five patients have the above features. DNT, despite having a heterogeneous morphological appearance, are unified by their clinical presentation of drug-resistant epilepsy often with the onset in childhood\cite{1}. In about 90% of cases, the first seizure occurs before 20 years of age\cite{1}. Patients may present with seizures in adulthood. Four of the five patients were children below the age of 15 years; all of them presented with epilepsy that was drug resistant. The fifth patient was a 25 year old adult. The tumor has been reported in association with neurofibromatosis Type 1\cite{13}. No recurrence was noted despite incomplete removal of the tumor in 71% of the cases\cite{14}. Honavar et al.\cite{15} reported a follow-up for up to 15 years, and confirmed the indolent biological behaviour of this tumor. All of our patients showed the tumor in the cerebral cortex. DNT show a predilection for the temporal lobe\cite{1}. Although most DNT occur in the cerebral cortex, reports are appearing about DNT in other locations, including the pons\cite{16}, thalamus\cite{16,17}, basal ganglia\cite{16,18,19}, cerebellum\cite{16,20,21}, and third ventricle\cite{16}. Of these “ectopic” locations, the basal ganglia/lateral ventricle has been the most common site\cite{12}. Although the glial component is usually that of an astrocytoma, oligodendroglioma may represent the glial component or small foci of oligodendroglioma may be seen within a more typical neoplasm\cite{22,23}. Although there is variation in markers used and results obtained by the different groups, others have also demonstrated evidence of multiple paths of differentiation\cite{5-9,16}. The proliferative markers indicated low proliferative profile in DNT\cite{5-7,9,24}. Cytologically, the principal tumor cells of the DNT are similar to those of the oligodendroglioma\cite{12}. Both tumors may also share a similar microcystic architecture and delicate vascular network. The differentiation between them is very difficult in frozen material and in small biopsies, which lack the multinodular pattern of DNT. In addition, the presence of isolated neurons within the DNT can be interpreted as evidence of an infiltrating neoplasm (i.e., oligodendroglioma or fibrillary astrocytoma)\cite{12}. Perineuronal satellitosis, frequently a prominent feature in oligodendroglioma, is scant, if present at all, in the DNT\cite{12}. In the context of the classic morphological
appearance, usually immunohistochemistry stains are not required for the diagnosis; they have been of only marginal use in establishing the diagnosis\textsuperscript{[12]}. The inconsistent immunoreactivity of GFAP in the OLCs contributed to misinterpretation as a glial neoplasm\textsuperscript{[12]}. Granular perivascular immunoreactivity for synaptophysin in tumor neuropil, although a finding in DNT\textsuperscript{[7,24]}, is nonspecific and often focal. Immunohistochemistry is useful in eliminating certain entities with characteristic immunohistochemical profiles from the differential diagnosis such as central neurocytoma and clear cell ependymomas. Central neurocytoma is diffusely and strongly positive for synaptophysin that is not a feature of DNT. Likewise, immunoreactivity for GFAP in the pseudorosettes of clear cell ependymomas distinguishes ependymomas from DNT. Fortunately, given the difficulties in histologic diagnosis, the radiological features of these lesions are distinctive\textsuperscript{[12]}. In conclusion DNT are extremely slow growing masses, as evidenced by their ability to deform the overlying skull. Total or even subtotal resection has reportedly been curative; however the rare incidence of tumor recurrence necessitates long-term surveillance by neuroimaging. Although the characteristic clinical, radiological and histopathological features are well documented, full understanding of the nature of this lesion and its pathogenesis and molecular characteristics awaits further clinical experience. The recognition of this tumor is very important because it is a surgically curable lesion. Pathologists need to be familiar with the characteristic histopathological findings to avoid unnecessary overtreatment by radiotherapy or chemotherapy.

References


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ورم الخلايا العصبية ذات التشويه الجنيني التكوين: الوصف النسيجي والصبغات المناعية

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الخلايا العصبية ذات التشويه الجنيني التكوين

يعتبر من الأورام النادرة. التشخيص النسيجي الصحيح لهذه الأورام مهم جداً، وذلك لأنها تعالج تماماً بالاستئصال الجراحي فقط. تم مراجعة سجلات قسم علم الأمراض بمستشفى الملك فيصل التخصصي ومركز الأبحاث - جدة - المملكة العربية السعودية، لكل حالات ورم الخلايا العصبية ذات التشويه الجنيني التكوين، وتم حصر خمس حالات من هذا الورم، وذلك خلال الفترة ما بين عام 2002 إلى 2007. تم مراجعة الصفات الإكلينيكية، الإشعاعية، النسيجي والصبغات المناعية. تم تشخيص خمس حالات من هذه الأورام النادرة. أعمار المرضى تراوحت ما بين 9 و 25 سنة، وكان هناك رجلان وثلاث نساء. كل المرضى أشتكوا من الصرع، في مريض واحد كان وجود الورم في الفقرين الصدغي والمرضى الآخرين في الفص الجبهي. الأشعة أظهرت وجود الورم غير المميز وغير تأثير كتلي أو أديم. الفحص النسيجي في كل الحالات أظهر أن الورم يتكون من خلايا مشابهة لنسج غسينات الدبق العصبي الدقيقة، الخلايا العصبية النجمية وخلايا الوحدة العصبية. الاستجابة للجراحة كانت ممتازة. التعرف على هذا الورم

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مهمًا، وذلك لأنه يتم معالجته تمامًا بالجراحة. يجب على أطباء علم الأمراض أن يكونوا على دراية تامة بالصفات النسيجية لهذا الورم، وذلك لتجنب العلاج الإشعاعي والكيميائي بدون حاجة.