Psychosis and Dandy-Walker syndrome: a case report and review of the literature

Alejandro Porras Segovia,1 Margarita Guerrero-Jiménez 2, Carmen Maura Carrillo de Albornoz Calahorro,2 Luis Gutierrez-Rojas 3

ABSTRACT
Dandy-Walker syndrome (DWS) is a group of brain malformations which sometimes present with psychotic symptoms. We present the case of a patient diagnosed with Dandy-Walker variant who presented with schizophrenia-like psychosis. A man in his 30s was admitted to an acute psychiatric unit presenting with persecutory delusions, auditory hallucinations and violent behaviour. The MRI performed showed the typical alterations of Dandy-Walker variant: vermian hypoplasia and cystic dilatation of the fourth ventricle. He also suffered from mild intellectual disability. After being treated with olanzapine 10 mg/d for a month, his psychotic symptoms greatly improved and he was discharged. In conclusion, DWS may cause psychosis through a dysfunction in the circuit connecting prefrontal, thalamic and cerebellar areas. The association between these two conditions may contribute to the understanding of the aetopathogenesis of schizophrenia.

INTRODUCTION
Dandy-Walker syndrome (DWS) is a group of brain structural abnormalities that includes four subtypes: Dandy-Walker malformation, mega cisterna magna, posterior fossa arachnoid cyst and Dandy-Walker variant.1

Dandy-Walker variant is the mildest form and is characterised by vermian hypoplasia and cystic dilatation of the fourth ventricle. Its clinical presentation typically consists of intellectual disability and epileptic seizures, although some patients may remain asymptomatic. Diagnosis is based on neuroimaging tests, mainly MRI. There is currently no curative treatment and, consequently, therapeutic strategies focus on its symptoms and complications.2

Occasionally, psychotic symptoms and other psychiatric disorders have been reported to occur in patients with DWS.3-18 However, the prevalence and clinical features of this comorbidity have not been established yet, and the literature about the topic is scarce. This is not the only neurological condition that presents with psychiatric comorbidities: several brain alterations, both structural and pathophysiological, can result in psychiatric symptoms.19 Moreover, neurodevelopmental factors have been found to be involved in the aetiology of schizophrenia,20-22 and the cerebellum appears to play an important role in this process.23-26

CASE PRESENTATION
A man in his 30s was admitted to an acute psychiatric unit after presenting with delusions and violent behaviour.

The patient had been diagnosed with Dandy-Walker variant as a child, manifesting as mild cognitive impairment and epileptic seizures. Electroencephalogram had revealed the presence of a rolandic focus, and MRI studies had shown vermian hypoplasia and dilatation of the fourth ventricle. He was treated with valproate for several years until seizures stopped around the age of 15 years. He lived with his mother, had no friends and worked as a janitor in a primary school.

A year prior to his admission, he started to feel suspicious of his neighbours and thought the children in the school where he worked made fun of him. He occasionally manifested aggressive behaviour, insulting and threatening his neighbours. The day before his admission, at the age of 32, the patient stepped into a school class wielding a knife and threatened the children and their teacher. He was arrested but was declared legally unaccountable after forensic psychiatric evaluation.
To review the association between DWS and psychosis, we carried out a search on the MEDLINE database using the terms [(Psychosis OR Schizophrenia) AND Dandy-Walker], in all fields, without filter restrictions. The initial search generated 17 articles, of which 12 were included in the review, while the other 3 were discarded due lack of relevance or unavailability. Searching the references of these articles, we found three additional studies. A total of 15 articles, reporting 19 patients, were included in the review.

Fifty-three percent of the patients were female. The mean age of onset of psychosis was 23 years. Dandy-Walker variant was the subtype more frequently associated with psychosis. Additional psychiatric comorbidities, such as obsessive-compulsive disorder, post-traumatic stress disorder or mania, were described in some of the cases. Atypical antipsychotics were the main treatment for psychosis in most of the cases reviewed. Complete results are shown in table 1.

A coincidental association between DWS and psychosis cannot be ruled out. However, the high number of cases of comorbidity reported, considering the rareness of the condition, suggests a causal relationship.

DWS may cause psychosis via its cerebellar alterations, which may explain why Dandy-Walker variant, characterised by vermian hypoplasia, is the subtype more strongly associated with psychosis, despite being the milder condition of the group. Some studies have shown a smaller vermian volume and other cerebellar alterations in patients with schizophrenia. According to the model proposed by Andreasen et al, psychotic symptoms may be the result of a dysfunction in the circuit connecting prefrontal, thalamic and cerebellar areas, a neurodevelopmentally derived ‘misconnection syndrome’ that would result in ‘cognitive dysmetry’, which has been described as a difficulty in prioritising, processing, coordinating and responding to information. According to this, a ‘poor mental coordination’ may explain both the cognitive deficits of schizophrenia and the wide range of symptoms. They opined that a decrease in the Purkinje cell size and decreased excitatory input to them from the granule cells have major implications that explain cerebellar and corticocerebellar-thalamic-cortical circuit dysfunction in schizophrenia and related abnormalities in symptoms and cognition. We think that these findings could predispose the occurrence of the other mentioned psychiatric disorders (table 1).

In conclusion, this case report provides further evidence linking DWS with psychosis. Considering these findings, some clinical features seem to be common in this kind of psychosis: earlier age of onset, higher prevalence of treatment resistance and frequent comorbid intellectual disability. One of the challenges that this condition poses is that of differentiating which symptoms are related to psychosis and which are caused by the primary neurological condition, especially in the case of intellectual disability, since it can often

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**DISCUSSION**

This case illustrates the association between Dandy-Walker variant and schizophrenia-like psychosis.

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**Figure 1** Midsagittal magnetic resonance T1-weighted scan shows cystic dilatation of the fourth ventricle and vermian hypoplasia.

He was transferred to the emergency room and later admitted to a psychiatric emergency ward.

At admission, the patient presented with persecutory delusions, claiming that the schoolchildren and other townspeople were planning to end his life and defended his actions as self-defence. He also presented with auditory hallucinations, claiming he could hear the voices from the deceased with whom he talked daily. He did not display thought disorder, but some limitations in abstract thinking and a mild intellectual disability (Wechsler Adult Intelligence Scale III score of 65) were evident on examination.

The patient gave written informed consent to all diagnostic procedures and allowed us to use the available data for scientific reports. An MRI was performed, showing the typical alterations of Dandy-Walker variant. No other alterations were found (figure 1).

He was treated with olanzapine 10 mg/d. Apart from mild sedation, which decreased after the first week, he tolerated the medication. His psychotic symptoms greatly improved a month after admission, with full remission of auditory hallucinations and a reduction from 87 to 51 in the Positive and Negative Syndrome Scale (PANSS). His delusions persisted but greatly decreased in intensity. He was discharged from the hospital and his care was transferred to his local mental health outpatient team.

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<table>
<thead>
<tr>
<th>Study</th>
<th>Patient's sex and age (years)</th>
<th>Dandy-Walker subtype</th>
<th>Psychotic symptoms</th>
<th>Other psychiatric comorbidities</th>
<th>Age of onset of psychosis</th>
<th>Last antipsychotic medication used</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Kvitvik Aune et al⁹</td>
<td>Male, 22</td>
<td>Dandy-Walker variant</td>
<td>Auditory and visual hallucinations, persecutory delusions, blunted affect</td>
<td>Borderline intelligence (IQ=79)</td>
<td>12</td>
<td>Risperidone 2mg/24 hours</td>
<td>Remission of positive symptoms, negative symptoms remained</td>
</tr>
<tr>
<td>Blaettner et al⁹</td>
<td>Male, 19</td>
<td>Dandy-Walker variant</td>
<td>Persecutory delusions, delusions of reference</td>
<td>OCD</td>
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<td>Partial remission</td>
</tr>
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<td>Buonaguro et al⁹</td>
<td>Female, 29</td>
<td>DWS</td>
<td>Persecutory delusions, blunted affect, agitation episodes</td>
<td></td>
<td>20</td>
<td>Haloperidol 4mg/24 hours</td>
<td>No response to treatment</td>
</tr>
<tr>
<td>Ferentinos et al⁹</td>
<td>Female, 21</td>
<td>Mega cisterna magna</td>
<td>Auditory hallucinations, persecutory delusions</td>
<td>Intellectual disability</td>
<td>18</td>
<td>Amisulpiride 1200mg/24 hours</td>
<td>57% decrease in PANSS score (after treatment resistance with previous antipsychotics)</td>
</tr>
<tr>
<td>Gan et al⁹</td>
<td>Female, 15</td>
<td>Dandy-Walker malformation</td>
<td>Auditory hallucinations, delusions of persecution and control, abulia</td>
<td>Intellectual disability</td>
<td>11</td>
<td>Risperidone 4mg/24 hours</td>
<td>39–76 decrease in the Scale for the Assessment of Positive Symptoms score, negative symptoms remained</td>
</tr>
<tr>
<td>Gan et al⁹</td>
<td>Male, 13</td>
<td>Dandy-Walker variant</td>
<td>Delusions of reference and persecution, blunted affect</td>
<td>Obsessive worries and school phobia</td>
<td>13</td>
<td>Quetiapine 400mg/24 hours</td>
<td>50% reduction in BPRS (from 43 to 25) but recurrence because of discontinuation of the treatment</td>
</tr>
<tr>
<td>Gan et al⁹</td>
<td>Male, 45</td>
<td>Mega cisterna magna</td>
<td>Auditory hallucinations, delusions of persecution and jealousy</td>
<td>Borderline intelligence and memory impairment</td>
<td>32</td>
<td>Risperidone 7mg/24 hours</td>
<td>34–56 decrease in BPRS, 4–6 decrease in the Clinical Global Impression of Severity scale core</td>
</tr>
<tr>
<td>Gan et al⁹</td>
<td>Male, 20</td>
<td>Posterior fossa arachnoid cyst</td>
<td>Delusions, stereotyped thinking</td>
<td>Borderline intelligence (IQ=69), hypomanic episode</td>
<td>20</td>
<td>Olanzapine 20mg/24 hours</td>
<td>Full remission</td>
</tr>
<tr>
<td>Kumar et al⁹</td>
<td>Female, 37</td>
<td>Mega cisterna magna</td>
<td>Auditory and visual hallucinations, Catatonia.</td>
<td></td>
<td>31</td>
<td>Risperidone 4mg/24 hours</td>
<td>Partial remission</td>
</tr>
<tr>
<td>Langarica nd Peralta⁹</td>
<td>Female, 52</td>
<td>Mega cisterna magna</td>
<td>Persecutory and reference delusions</td>
<td></td>
<td>52</td>
<td>Olanzapine 7.5mg/24 hours</td>
<td>Significant clinical improvement</td>
</tr>
<tr>
<td>Mauritz et al¹⁰</td>
<td>Female, 47</td>
<td>Dandy-Walker malformation</td>
<td></td>
<td>PTSD</td>
<td>47</td>
<td>Quetiapine 600mg – 1000mg/24 hours</td>
<td>Full remission</td>
</tr>
<tr>
<td>Pandurangi et al¹¹</td>
<td>Male, 26</td>
<td>Megacisterna magna</td>
<td>Delusions of grandiosity</td>
<td>Mania</td>
<td>26</td>
<td>Risperidone 4mg/24 hours</td>
<td>Significant clinical improvement, delirious and urinary incontinence as secondary effects</td>
</tr>
<tr>
<td>Pandurangi et al¹¹</td>
<td>Male, 20</td>
<td>Megacisterna magna</td>
<td>Catatonia</td>
<td></td>
<td>18</td>
<td>Risperidone 3mg/24 hours</td>
<td>Discontinuation of treatment due to acute dystonia</td>
</tr>
</tbody>
</table>

Continued
be mistaken for negative symptoms. Additionally, the management of non-psychiatric complications of DWS can improve the patient’s general well-being.

Despite the body of evidence linking schizophrenia to brain abnormalities, current Diagnostic and Statistical Manual of Mental Disorders, Fifth Edition, diagnostic criteria for schizophrenia exclude this diagnosis when symptoms are considered to be caused by a neurological condition. Thus, we must refer to this entity as schizophrenia-like psychosis, even if the rest of the criteria for schizophrenia are met. As research advances and the aetiology of schizophrenia become clearer, the boundary between primary and secondary schizophrenia may become blurrier. Further research is needed to establish the prevalence and characteristics of psychotic symptoms in patients with DWS. This association may contribute to the understanding of the aetiopathogenesis of schizophrenia.

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**REFERENCES**


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