Mania as a Possible Prodrome to Dementia

SIR: There has been much debate over whether late onset depression may be a prodrome to dementia. ¹ Kessing and Fleming² have also suggested a link between affective disorders and dementia. The authors could not find a report in the literature that described mania preceding the diagnosis of dementia. Below we describe an elderly patient who was diagnosed with dementia following a solitary episode of mania.

Case Report

"Mr. V" is a 76-year-old Caucasian man who presented to our facility with an isolated manic episode prior to obtaining a definitive diagnosis of dementia. Besides a selflimited episode of depression that occurred in his thirties, he had no other notable past psychiatric history. The patient was reported to be euthymic and relatively cognitively intact until he underwent the removal of a basal cell carcinoma for which he received a 2 to 3 week course of corticosteroids. Approximately 6 weeks following the discontinuation of the corticosteroids, Mr. V exhibited symptoms of mania, which included insomnia, hyperreligiosity, hypersexuality, hyperverbosity, irritability, and increased cleaning behavior which led to his first psychiatric hospital admission. Upon admission to the hospital, Mr. V's Mini-Mental State Examination (MMSE) was 21/30. He was treated with a vast array of mood stabilizers and his mania eventually abated 2 months into his hospital stay. Following the aforementioned manic episode, the patient then became

apathetic while in a euthymic state. He would sleep the majority of the day and night and express no desire to participate in activities. His apathy was mood congruent and there was no indication of any affective disturbance. The patient would also frequently confabulate his activities and experiences. Neuropsychological testing revealed impairments in verbal and visual learning and memory, executive functioning, and complex aural comprehension. Object naming and word finding tasks were preserved, suggesting a diagnosis of Alzheimer's dementia. His MMSE at this time was 23/30. A full medical work-up was also conducted and revealed no vitamin deficiencies, thyroid dysfunction, or source of infection. His EEG was normal and a magnetic resonance image of his brain revealed small vessel ischemic changes. Furthermore, a single photon emission computed tomography scan revealed hypoperfusion of both frontal lobes and mild bilateral temporal lobe perfusion; both of which were deemed nondiagnostic findings.

Comment

The presentation of the above patient suggests that affective disorders in general may precede the diagnosis of dementia. Such a prodrome may not be confined to depression and may, in fact, include mania.

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References

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- dementia? Int J Geriatr Psychiatry 2002; 17:997–1005
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A Rare Case of Epilepsy in a 16-Year-Old Girl With Fallot Tetralogy Attributed to CNS Heterotopia and Pachygyria

SIR: We report a case of a 16year-old girl who was admitted to our clinic because of partial seizures. The clinical neurological examination and the neuropsychological tests were normal. The laboratory tests (i.e., CBC, blood biochemistry, urine tests, immunologic assay tests) were normal. The CSF analysis and cultures, as well as the CSF protein electrophoresis, were normal. Cerebral computed tomography (CT) was normal and ECG showed RBBB (right bundle branch block) and LAH (left anterior hemiblock). However, the EEG revealed abundant generalized slow wave activity with occasional focal and diffuse spikes and spike wave activity.

The brain magnetic resonance imaging (MRI) scan revealed subcortical heterotopia of the gray matter extending across the right ventricle. More specifically, along the surface of the occipital horn of the right ventricle and at its posterior part as well as at the right temporal horn, there was an irregularly lobulated mass of gray matter that extended into the hemispheric white matter giving signals similar to the cerebral cortex. There were also similar