Alzheimer Abnormalities of the Amygdala With Klüver-Bucy Syndrome Symptoms

An Amygdaloid Variant of Alzheimer Disease

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Background: Neurofibrillary tangles and β-amyloid plaques have been observed in the amygdala in Alzheimer disease. A disproportionate abundance of this abnormality in the amygdala may cause behavioral symptoms similar to Klüver-Bucy syndrome.

Objectives: To describe an atypical behavioral presentation of Alzheimer disease and to review the literature on the subject.

Design: Case study.

Setting: Outpatient specialty clinic.

Patient: A 70-year-old man with progressive behavioral symptoms of hyperorality, hypersexuality, hypermetamorphosis, visual agnosia, hyperphagia, and apathy who died at age 77 of asphyxiation on a foreign object.

Main Outcome Measures: Clinical symptomatology, brain imaging, and neuropathology.

Results: The pathologic diagnosis was Alzheimer disease with abundant tangles and plaques in the lateral amygdala.

Conclusions: This case represents a variant of Alzheimer disease with prominent amygdala abnormalities and a Klüver-Bucy phenotype that was misdiagnosed as frontotemporal dementia. Clinical and imaging findings that may aid in accurate diagnosis are reviewed.

theophylline, allopurinol, and indomethacin. There was no history of alcohol, tobacco, or other substance use. His sister and father had unspecified late-onset dementia.

On examination, he was observed to be obese (107.55 kg, 167.64 cm tall) and “jocular and inappropriate.” His Mini-Mental State Examination score was 30 of 30. Results of his neurologic examination were normal except for symmetrical hyporeflexia, marginally impaired graphesthesia, and a right palmenatal sign. Neuropsychological testing revealed impairment in object naming and mild impairment in acquisition, with otherwise normal scores (Table). Basic chemistries, complete blood cell count, thyrotropin level, vitamin B12 level, and microhemagglutination Treponema pallidum level were normal. Brain magnetic resonance imaging revealed a 0.7 × 2-cm meningioma adjacent to the left sylvian fissure and very mild subcortical punctate white matter changes; otherwise, the results were interpreted as normal. Retrospective review of these images suggested bilateral parietal atrophy (Figure 1). The initial clinical impression was mild frontotemporal dementia (FTD).

During the next 2 years, he reported that he began having more prominent sexual fantasies about other women while also becoming increasingly suspicious that his wife was cheating on him; he was prescribed paroxetine mesylate. His Mini-Mental State Examination score remained at 28 of 30; however, the neuropsychological studies began to show clear deficits in delayed recall and language with spared visuoperceptive ability (Table). Positron emission tomography (PET) demonstrated hypometabolism in the bilateral parietal and anterotemporal regions (Figure 2). The possibility of AD was considered, but the leading diagnosis remained FTD. From the ages of 74 to 77 years, he had a substantial decline in cognitive function, and his Mini-Mental State Examination score declined to 7. He displayed increasing hyperorality, and his wife noted that he would put “any” object in his mouth, including dog food, candles, adhesive bandages, and his wedding ring. His appetite seemed insatiable. He would reach into the toilet to manipulate fecal matter. When asked by his wife to get a rake, he brought a screwdriver. He became progressively withdrawn and apathetic. During the last year of his life, he required assistance with most

**Table. Neuropsychological Test Results for the Patient at Ages 70 to 74 Years**

<table>
<thead>
<tr>
<th>Neuropsychological Test</th>
<th>70</th>
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<td>29</td>
<td>28</td>
<td>27</td>
<td>18</td>
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<td>Attention</td>
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<td>Digit Span score</td>
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<td>14 (101)</td>
<td>13 (97)</td>
<td>9 (82)</td>
<td>9 (82)</td>
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<tr>
<td>Language</td>
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</tr>
<tr>
<td>Boston Naming Test score</td>
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<td>39 (55)</td>
<td>31 (&lt;55)</td>
<td>23 (&lt;55)</td>
<td>16 (&lt;55)</td>
</tr>
<tr>
<td>Memory score</td>
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<tr>
<td>Logical Memory I</td>
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<td>7 (71)</td>
<td>4 (65)</td>
<td>3 (63)</td>
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<td></td>
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<td></td>
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<tr>
<td>Block design score</td>
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<td>27 (115)</td>
<td>20 (105)</td>
<td>12 (90)</td>
<td>16 (100)</td>
</tr>
</tbody>
</table>

*Abbreviations: MMSE, Mini-Mental State Examination; NA, not applicable.

a Standard scores are given in parentheses (mean [SD], 100 [15]).

**Figure 1.** Brain magnetic resonance images of the patient at age 72 years showing slightly enlarged biparietal sulci and the absence of frontal or temporal lobar atrophy.
activities of daily living. He died at age 77 years of asphyxiation on several adhesive bandages.

The pathologic diagnosis was AD with abundant neurofibrillary tangles and moderate neuritic plaques throughout the brain (Braak and Braak stage VI and a CERAD [Consortium to Establish a Registry for Alzheimer’s Disease] rating of “frequent”). The brain weight was 1400 g. The distribution of tangles and plaques was similar to that encountered in sporadic AD with this staging; however, the degree of amygdala abnormality was atypical. Neurofibrillary tangles were abundant in the lateral amygdala (approximately 150 per 1.6 mm²) (Figure 3 and Figure 4). This number of tangles is greater than 4+ (>50 tangles per 1.6 mm²) using the Kromer Vogt rating system for tangles in the amygdala in AD.² There were neuritic plaques in the lateral, laterobasal, and accessory basal amygdala ranging from approximately 10 to 20 per 1.6 mm² (Figure 5); 2+ Kromer Vogt rating for plaques in the amygdala.² The amygdala showed moderate neuronal depletion and gliosis. There was also evidence of grade II cerebral amyloid angiopathy. Genetic testing showed that the patient had apolipoprotein E 3/3; presenilin genotyping was not performed.

**COMMENT**

Described by Klüver and Bucy in the late 1930s, this behavioral syndrome is associated with bilateral amygdala lesions and is characterized by the following cluster of symptoms: hypersexuality, hyperorality, hypermetamorphosis (excessive exploration of visual stimuli), visual agnosia, apathy, and withdrawal.⁶ The amygdala processes sensory information for emotional valence; there-
fore, amygdala lesions cause abnormal emotional responses, resulting in aggressiveness, fearlessness, or apathy.2 Lesion studies3 in rats demonstrate that efferent projections from the posterodorsal amygdala to the hypothalamus lead to hyperphagia, obesity, and sexual dysregulation. Projections between the visual cortex and the amygdala have been mapped in nonhuman primates.4 Injury to these projections likely underlies the visual agnosia and hypermetamorphosis observed in patients with amygdala lesions.

Sourander and Sjogren10 described KB syndrome symptoms in patients with AD. Morris et al11 suggested that excessive eating in some patients with dementia may be a form of KB syndrome. Burns and colleagues12 examined 178 patients with AD to quantify the number of KB syndrome symptoms: 29.3% had 1, 24.7% had 2, 17.2% had 3, 8.6% had 4, and 0.6% had all the KB syndrome symptoms.

Neuronal loss in the medial, central, and cortical nuclei of the amygdala has been observed in AD.13,14 Amyloid plaques and neurofibrillary tangles are consistently present in the amygdala in AD.3,5 The present patient had 150 tangles per 1.6 mm² in the lateral amygdala. By comparison, 14 other AD brains with Braak and Braak stage VI were studied and were found to have 5 to 90 tangles per 1.6 mm² in the amygdala. Kromer Vogt et al2 investigated the amygdala abnormalities of 20 patients with AD and demonstrated a significant amount of tangles and neuritic plaques in the cortical nucleus and accessory basal nucleus of the amygdala, which have projections with the hippocampus; however, the lateral nuclei of the amygdala were relatively spared (1 patient had a 4+ tangle score and 1 had a 3+ plaque score). Compared with patients with typical AD described by Kromer Vogt et al,2 the patient described herein had significant tangle and plaque density in the lateral amygdala. Hayman and colleagues15 also described a patient with KB syndrome after lateral amygdala lesions.

This is the first published study of this amygdaloid variant of AD with serial neuropsychological testing, magnetic resonance imaging, PET, and pathologic analysis. A frontal variant of AD with prominent behavioral symptoms has been described.16 The frontal variant is characterized by an atypical distribution of AD abnormalities in the frontal lobes that may be more difficult to distinguish from FTD. In contrast, PET of this patient revealed the typical pattern of posterior hypometabolism. This tool might be a useful diagnostic aid. Behavioral variants of AD are primarily a diagnostic challenge because several of the symptoms overlap with FTD.17 Cognitive-behavioral symptoms in AD often respond to acetylcholinesterase inhibitors, and misdiagnosis of FTD might preclude this intervention and other potential disease-modifying agents for AD that are currently being investigated. Also, FTD has more rapid progression, with mean survival of 4.2 years from initial evaluation compared with 6 years for AD.18

This patient’s prominent behavioral symptoms led to the incorrect diagnosis of FTD. There were, however, a variety of noteworthy signs and imaging clues that pointed toward the correct diagnosis of AD. The patient noted memory dysfunction, and delayed memory scores were very poor relatively early in the illness. Age at presentation was late, and the course was relatively slow. Magnetic resonance imaging showed evidence of biparietal atrophy. The PET demonstrated biparietal hypometabolism consistent with AD. The specificity and sensitivity of PET, used to differentiate between AD and FTD, were recently shown to be superior to those of clinical assessment alone.10 Furthermore, the positive finding of temporoparietal hypometabolism is strongly associated with AD abnormalities even in the presence of atypical features.20

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Author Contributions: All authors had full access to all of the data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis. Study concept and design: Kile, Olichney, and DeCarli. Acquisition of data: Kile, Ellis, and Farias. Analysis and interpretation of data: Kile, Olichney, and Farias. Drafting of the manuscript: Kile and Farias. Critical revision of the manuscript for important intellectual content: Kile, Ellis, Olichney, and DeCarli. Administrative, technical, and material support: Kile and Farias. Study supervision: Ellis, Olichney, Farias, and DeCarli.

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REFERENCES


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