Left Frontotemporal Hypoperfusion Is Associated With Aggression in Patients With Dementia

Nobutsugu Hirono, MD; Michael S. Mega, MD, PhD; Ivo D. Dinov, PhD; Fred Mishkin, MD; Jeffrey L. Cummings, MD

Background: Aggressive behavior is common in patients with dementia. Temporolimbic and prefrontal cortices lesions can produce pathological aggression; however, involvement of these structures has not been established in aggressive patients with dementia.

Objective: To study the relation between regional brain perfusion and aggressive behavior in patients with dementia.

Methods: We compared the pattern of regional cerebral perfusion determined with technetium Tc 99m-labeled hexamethylpropelene amineoxime single photon emission computed tomography in 2 groups of 10 patients with dementia with and without aggression, that were comparable for demographic factors, severity of cognitive impairments, and other behavioral symptoms as measured by the Neuropsychiatric Inventory.

Results: Patients with aggression revealed significant ($P < .001$) hypoperfusion in the left anterior temporal cortex; additional bilateral dorsofrontal and right parietal cortex were also found to be significantly hypoperfused.

Conclusion: These results indicated an association between aggression and decreased perfusion in the left anterior temporal cortex.

Arch Neurol. 2000;57:861-866

Behavioral symptoms manifest in patients with dementia and have a major influence on quality of life and social affiliations for patients and caregivers. Among these behavioral problems, aggressive outbursts are frequent and may pose the greatest caregiver distress. Agitation or aggression are reported in 12% to 65% of patients with Alzheimer disease (AD) or other dementias. Aggressive behavior causes serious burden and depression in caregivers, and increases the likelihood of hospitalization, early institutionalization, and abuse of patients. In the nursing home, aggressive behavior is associated with a need for increased staff supervision, and the use of physical restraints. Moreover, aggressive behaviors may have severe adverse consequences, including injury, which may lead to the death of patients or others. Therefore, evaluation and management of aggressive behaviors are of considerable importance in the care of patients with dementia.

Although functional factors, such as pathological rearing, psychological trauma, and social stress, are important in the occurrence of aggression, regional brain lesions can produce pathological aggression in animals and humans. Among these regions, the temporolimbic and prefrontal cortices have been most frequently implicated. Aggressive verbalization and rage occur with stimulation of the amygdala and hippocampus in animals. Although destruction of the amygdala in monkeys and humans usually results in placidity, Rosvold et al reported that amygdalectomy in submissive monkeys increased aggression, suggesting that amygdalar dysfunction modifies aggressive behavior in a variety of ways. Recently, Zagrodzka et al reported that unilateral damage to the central nucleus of the amygdala alone increased the expression of aggressive behavior in the cat. Patients with temporal lobe epilepsy may have increased aggression during interictal periods in addition to ictal and postictal aggressive outbursts. Furthermore, violent patients with focal lesions are most likely to have focal temporal abnormalities on electroencephalography and structural abnormalities in temporal lobes. Functional neuroimaging studies using positron emission tomography...
**PATIENTS AND METHODS**

**PATIENTS**

From a pool of 198 patients with dementia initially seen at the University of California Los Angeles (UCLA) Alzheimer's Disease Center, 20 outpatients who met all clinical criteria described below, comparable across demographic and behavioral domains and divided equally into those with and without agitation/aggression, were studied. All patients had acquired persistent decline involving at least 3 of the following domains: language, memory, visuospatial skills, cognition (ie, calculation, abstraction, judgment, and others), and emotion or personality. All patients with AD met National Institute of Neurological and Communicative Disorders and Stroke–Alzheimer’s Disease and Related Disorders Association criteria for probable or possible AD. All patients with vascular dementia met National Institute of Neurological Disorders and Stroke–Alzheimer’s Disease and Related Disorders Association International Workshop criteria for probable or possible vascular dementia; patients with large cortical strokes were excluded from the study. All patients with dementia who had Lewy bodies met the Consortium for Dementia with Lewy Bodies criteria for probable dementia with Lewy bodies. Diagnostic evaluation for all patients included complete medical history, physical and neurological examination, magnetic resonance imaging or computed tomographic imaging, technetium Tc 99m–labeled hexamethylpropylene amine oxime–SPECT imaging of the brain, and electroencephalography, and routine blood tests (including thyroid, vitamin B12 level, and serological test for syphilis), and had caregivers who were willing to be interviewed. Exclusion criteria for all patients were delirium, history of alcohol or other substance abuse, history of head trauma with loss of consciousness, and history of psychiatric disorder preceding the onset of symptoms of dementia. Patients were physically well without pain or other identifiable precipitants of behavioral disturbances. Cognitive assessment included the Mini-Mental State Examination, the Consortium to Establish a Registry for Alzheimer’s Disease 10-word memory test, the modified 15-item Boston Naming Test, the Consortium to Establish a Registry for Alzheimer’s Disease visuospatial task (copying a circle, diamond, overlapping rectangles, and cube), and a verbal fluency task (animal name generation in 1 minute). The 20 patients chosen were representative of the 198 patient pool having a similar sex, age, Mini-Mental State Examination score, and educational level of the larger group; the pool was assembled from all patients seen in the UCLA Alzheimer’s Disease Center who had cognitive, behavioral, laboratory, structural, and functional (SPECT) imaging.

**BEHAVIORAL ASSESSMENT**

Caregivers were interviewed with the NPI following procedures previously described in which screening questions for each behavior were posed first. The caregiver was asked if the behavior represented a change from that exhibited by the patient prior to the onset of the dementia and was present during the past month. If a positive response was obtained from the screening questions, then the behavioral domain was explored with scripted questions focusing on specific features of the behavioral disturbance. The caregiver then rated the behaviors; scores from 1 to 4 (4 being the most frequent) were obtained for the frequency and 1 to 3 (with 3 being the most severe) for the severity of each behavior (a composite score for each domain was the product of the frequency and severity subscores; maximum attainable score = 12). The NPI has been shown to be valid and reliable; raters receive specific training in NPI administration and are retrained periodically to prevent drift. The 10 domains assessed by the NPI are delusions, hallucinations, agitation/aggression, depression, anxiety, euphoria, apathy, disinhibition, irritability, and abnormal motor output. The cognitive assessment was done concurrently with the NPI.

**SINGLE PHOTON EMISSION COMPUTED TOMOGRAPHY**

For all patients, an intravenous line was placed and a 15-minute period was allowed to pass for patients to regain a quiet comfortable state before the intravenous administration of 1110 MBq of technetium Tc 99m–labeled hexamethylpropylene amine oxime (Ceretec; Amersham, Arlington Heights, Ill). Room lights were dimmed and quiet was observed to minimize patient visual and auditory stimuli during the brain uptake phase for another 15 minutes following...
behavioral symptoms captured by the NPI, including delusions, hallucinations, disinhibition, irritability, apathy, anxiety, and depression. We hypothesized that patients with dementia with significant agitation/aggression would have significantly lower anterior temporal and lateral orbitofrontal perfusion on SPECT analysis compared with patients without agitation/aggression.

Table 1 and Table 2 show the demographic features, psychotropic medications, Mini-Mental State Examination (all other cognitive indices assessed by our battery were comparable between groups), and behavioral profiles of the 2 groups. Of the 10 aggressive patients with dementia 8 were diagnosed as having AD, and 2 were diagnosed as having vascular dementia. Of the 10 nonaggressive patients with dementia 8 were diagnosed as having AD, 1 was diagnosed as having dementia with Lewy bodies, and 1 was diagnosed as having normal pressure hydrocephalus. Agitation/aggression was the only behavior assessed by the NPI that was statistically significantly (P<.001) different between the 2 groups. The Figure and Table 3 show the Talairach and Tournoux atlas location of the peak significance for regions with significantly lower perfusion in the 10 patients with dementia who had high aggression scores compared with the 10 nonaggressive patients. Patients with aggression revealed statistically significant (P<.001) hypoperfusion in the left anterior temporal cortex (Brodmann areas 20, 21, and 38); the difference in the orbitofrontal cortices was not significant. Besides the regions we hy-
pothesized would be affected in aggressive patients with dementia, patients with aggression also revealed significant ($P<.001$) hypoperfusion in the right and left superior frontal cortices (Brodmann area 9) and the right superior parietal area (Brodmann area 7) with between-group differences after Bonferroni correction. Possible confounding effects of medication on cerebral perfusion were controlled for by composing the groups to have an equal percentage of patients with neuroleptic and antidepressant use in both groups.

### COMMENT

Several studies have suggested that aggressive behavior is associated with demographic factors including male gender, and increasing cognitive loss in patients with dementia. Therefore, we minimized the confounding effects of these variables: age, sex, education level, psychotropic medications, Mini-Mental State Examination scores, and cognitive performance, all of which were similar between the patients groups with and without aggressive behavior. Moreover, because aggression has been associated with other neuropsychiatric manifestations including delusions, hallucinations, irritability, and depression, we also ensured that the neuropsychiatric symptoms in the 2 groups were comparable. This approach aids identification of the brain regions that, when dysfunctional, may be specific to aggressive behavior.

Generalizability of this study is limited by the small sample size. The study demonstrated a clear association between aggressive behavior and hypoperfusion in the left anterior temporal cortex but failed to demonstrate the expected involvement of the orbitofrontal cortices in patients with dementia. This failure might be caused by the lack of statistical power due to the small sample size.

The anterior temporal cortex has reciprocal connections with the amygdaloid complex. The amygdala has primary symptom of brain dysfunction and others view it as a symptom of psychopathology. In this study, multiple cortical sensory systems; visceral brain areas including brainstem and hypothalamus; basal ganglia; and other limbic structures such as parahippocampal gyrus, insula, and cingulate cortices. Anterior temporal structures have been posited to play an important role in associating sensory experiences with emotion. Dysfunction of these areas has been implicated in causing abnormal emotional associations to external sensory stimuli, and in provoking inappropriate aggressive acts to trivial or misinterpreted stimuli.

In addition to involvement of the left anterior temporal cortex, we found significant regional hypoperfusion in the right and left dorsolateral frontal and right superior parietal regions. Within the prefrontal regions, the disturbances of orbitofrontal cortices have been linked most often to aggression; however, several studies have shown the involvement of the dorsolateral frontal region in aggression. Dorsolateral frontal regions are considered important to critical thinking and planning, and reduced metabolic activity in these regions may lead to misinterpretation of environmental and social situations, which might be related to aggressive behavior. Raine et al described significantly decreased glucose metabolism in both superior parietal regions in murderers as compared with normal controls. These findings also are compatible with the fact that aggressive behaviors are more frequent in patients with frontotemporal dementia, who usually have dysfunction of dorsolateral frontal regions as well as of anterior temporal structures, as compared with patients with AD. The superior parietal lobes are critical for sensorimotor integration providing subjective knowledge of both the world and one's own body. Dysfunction of the superior parietal lobes may cause the sensory information processing deficits and abnormal assessments of both the world and one's own body resulting in abnormal emotional responses.

This study bears importantly on the understanding of agitation in dementias. Agitation encompasses many behaviors including aggressive behavior assessed by the NPI. The pathogenesis of agitation in dementia is controversial with some investigators considering it a primary symptom of brain dysfunction and others viewing it as a symptom of psychopathology. In this study,
CONCLUSIONS

We prospectively studied aggressive behaviors in patients with dementia and the relevant regional cerebral hypoperfusion by using SPECT and an established method of neuropsychiatric assessment in dementia. We demonstrated an association between aggression and decreased perfusion in the left anterior temporal cortex. Our results support previous reports of anterior temporal involvement in aggressive behaviors, irrespective of disease origin. This finding might predict the development of aggression and the effectiveness of therapeutic interventions used for the patients with dysfunction in this region.

Accepted for publication December 22, 1999.

This investigation was supported by Career Development Award K08AG100784 from the National Institute on Aging, National Institutes of Health, Bethesda, Md (Dr Mega); Alzheimer’s Disease Research Center grant P50 AG16570 from the National Institute on Aging (Dr Cummings); an Alzheimer’s Disease Research Center of California grant, Los Angeles (Dr Cummings); and the Sidell-Kagan Foundation, Los Angeles (Drs Mega and Cummings).

Corresponding author: Nobutsugu Hirono, MD, Division of Clinical Neuroscience, Hyogo Institute for Aging Brain and Cognitive Disorders, 520 Saiisho-ko, Himeji 670-0981, Japan (e-mail: hirono@hiabcd.go.jp).

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