

Periodontal Health Condition in Patients With Alzheimer's Disease

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Abstract

Objective: To compare periodontal health status in individuals with and without Alzheimer's disease (AD). **Methods:** A total of 58 individuals with AD and 60 cognitively normal (ND) adult individuals, ranging in age from 50 to 80 years, were assessed for periodontal health status. Individuals with AD were further divided as mild, moderate, and severe, based on degree of cognitive impairment as evaluated using Mini-Mental State Examination. Gingival index (GI), plaque index (PI), probing depth (PD), clinical attachment level (CAL), and percentage of bleeding sites (%BOP) were evaluated. **Results:** All the evaluated periodontal parameters were higher in individuals with AD than that in ND individuals, and the periodontal status deteriorated with the progression of AD. There were significant differences in mean GI, PI, PD, CAL, and %BOP between all the groups. **Conclusion:** The periodontal health status of individuals with AD deteriorates with disease progression and was closely related to their cognitive function.

Keywords

Alzheimer's disease, dementia, chronic periodontitis, oral health

Introduction

Alzheimer's disease (AD) is the most common type of dementia affecting the elderly people. Alzheimer's disease is characterized by progressive degeneration in the central and peripheral neurological system¹ and presents with progressive memory loss and cognitive impairments.² Memory loss involves not only difficulty in remembering recent events but also impairments in holding information in mind over short periods of time.^{3,4} Alzheimer's disease is thought to progress uniformly from the earliest signs of impaired memory to severe cognitive loss, terminating inevitably in complete incapacity and death.⁵

According to World Alzheimer's report 2010, there are an estimated 35.6 million people with dementia worldwide. This number will nearly double every 20 years and much of the increase will be in developing countries. About 58% of people with dementia live in developing countries. The fastest growth in the elderly population is taking place in China, India, and their South Asian and Western Pacific neighbors. Alzheimer's disease mainly affects people older than 65 years, with a greater predilection for women, although there is a growing awareness of cases that start before the age of 65.⁶

Patients with AD experience limitations in medical and social conditions which may also complicate overall personal and medical care.⁷ Deficiencies in the care of these elderly patients may result in several systemic and intraoral health

problems including malnutrition, hygiene defects, pressure ulcers, and respiratory disturbances.^{8,9} Impaired cognition may severely affect daily activities, and poor oral hygiene execution may lead to caries and eventual loss of teeth.¹⁰⁻¹² Chronic periodontitis (CP) is a peripheral, infectious disease and is one of the leading cause for tooth loss.¹³ According to the Third National Health and Nutrition Examination Survey (NHANES

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III), gingival bleeding and loss of periodontal attachment (indexes of periodontal disease) were significantly associated with lower cognitive function and are thought to be a risk factor for AD.^{14,15} Clinical CP has been linked to other systemic inflammatory conditions such as cardiovascular disease,¹⁶ diabetes mellitus,¹⁷ and other neurodegenerative conditions like Parkinson's disease.¹⁸

To the best of our knowledge, there are no studies evaluating the periodontal health status in individuals with AD and assessing the effect of progression of AD on periodontal health in Asian population. With growing concerns regarding increasing incidences of AD and related dementia and its effect on periodontal health, this study aimed to evaluate the periodontal health status in individuals with AD, in order to find the influence of progression of AD on CP.

Materials and Methods

Study Population

This was a cross-sectional study, conducted from November 2011 to May 2012, in which 58 individuals with AD and 60 nondemented (ND) or cognitively normal adult individuals, ranging in age from 50 to 80 years, were selected from the Department of Neurology, National Institute of Mental Health and Neurosciences (NIMHANS), Bangalore, India. Ethical approval for the study was obtained from institutional ethical committee, Government Dental College and Research Institute, Bangalore.

Inclusion and Exclusion Criteria

The diagnosis of ND individuals and individuals with AD was done by medical/neurological evaluations, psychiatric behavioral assessments, and neuropsychological evaluation. All participants received extensive diagnostic screening to rule out confounding medical, neurological, and psychiatric conditions. The neurological tests carried out to confirm the diagnosis and progression of AD include magnetic resonance imaging and computed tomography scan. These neuroimaging modalities help in proper diagnosis of AD and differential diagnosis of non-AD brain lesions. Individuals with AD and ND individuals were free from other systemic conditions known to affect cognition. Individuals with psychiatric or behavioral conditions (eg, depression or substance abuse) or taking medications that affect cognition (eg, benzodiazepines) were excluded. The minimum number of scorable natural teeth required was 12. Edentulous individuals or those with <12 natural teeth present were excluded from the study.

The diagnosis of AD was in accordance with the National Institute of Neurological and Communicative Disorders and Stroke and the Alzheimer's Disease and Related Disorders Association (NINCDS-ADRDA) workgroup recommendations.¹⁹ The degree of cognitive impairment was evaluated using Mini-Mental State Examination (MMSE). The MMSE is a 30-point scale for assessing attention, orientation, registration, and calculation. The MMSE scores were classified using

the system developed by Mungas.²⁰ Those participants scoring 26 or more (out of 30) were categorized as within normal cognitive range, those scoring from 21 to 25 had mild dementia, those scoring from 11 to 20 had moderate dementia, and those scoring 10 or less had severe dementia. Accordingly individuals were classified into 4 groups: group 1: normal individuals; group 2: mild AD, n = 22 (13 females and 9 males); group 3: moderate AD, n = 18 (10 females and 8 males); and group 4: severe AD, n = 18 (9 females and 9 males). Written informed consent was obtained from all individuals. In severe cases, consent was obtained from the caregiver or any relative accompanying the individuals.

Periodontal Examination

A full-mouth comprehensive examination of periodontal condition was done for all individuals. Individuals were clinically evaluated in the outpatient section of Neurology department, NIMHANS. Patients with severe AD were examined in the outpatient section or respective hospitalized wards of NIMHANS with due help from the caretakers. Two examiners (SSM and SPS) blinded to the objectives of the study performed all dental examinations under the supervision of chief examiner ARP. Several calibration exercises were conducted before the start of the study and throughout the study period to control and minimize examiner drift. The examiners were calibrated on all periodontal measures used including plaque index (PI), gingival index (GI), and probing measures. Examiners calculated absolute agreement and relative agreement for probing measures in 10 patients and converted the continuous probing measures into categorical entities to calculate the κ -index value. For categorical probing measures, the κ -index values for intraexaminer agreement ranged from 0.81 to 0.95, which was higher than κ -index value for interexaminer agreement that ranged from 0.64 to 0.78.

The examination was done using a mouth mirror and a standardized periodontal probe (UNC 15 periodontal probe, Hu Friedy, Illinois) by the same examiner who was blinded to the neurological state of the patient. The periodontal parameters were recorded at 6 sites per tooth, namely mesiobuccal, buccal, distobuccal, mesiolingual, lingual, and distolingual. The parameters evaluated were (1) PI,^{21,22} (2) GI,²³ (3) probing depth (PD) taken from the gingival margin to the bottom of the pocket, (4) clinical attachment level (CAL) measured to the nearest millimeter from the cemento-enamel junction to the deepest probeable point,²⁴ and (5) percentage bleeding on probing (% BOP) sites. The BOP was scored positive if a site bled immediately after pocket probing or if a site bled at completion of the probing of a jaw quadrant.

Statistical Analysis

Statistical analysis was performed with statistical software (SPSS version 10.5, SPSS, Chicago, Illinois). The values of different parameters collected are expressed as mean \pm standard deviation (SD). Unpaired *t* test was used to compare

Table 1. Demographic and Neurological Characteristics of Individuals With AD and ND Individuals.

	AD Group (n = 58)	ND Group (n = 60)	P Value
Age (mean ± SD)	65.2 ± 7.3	64.5 ± 9.4	NS
Females	32/58 = 55.2%	34/60 = 56.7%	NS
Males	26/58 = 44.8%	26/60 = 43.3%	NS
Num of teeth present	15.8 ± 3.6	16.2 ± 4.2	NS
Oral hygiene status			
Good	2/58 = 3%	8/60 = 13%	NS
Fair	16/58 = 27.5%	18/60 = 30%	NS
Poor	40/58 = 69%	34/60 = 57%	NS
MMSE scores	14.2 ± 8.4	28.5 ± 1.2	<.001 ^a

Abbreviations: NS, not significant; AD, Alzheimer's disease; ND, nondemented; SD, standard deviation; Num, number.

^aStatistically significant at ($P < .05$); unpaired t test.

Table 2. Clinical Parameters (mean ± SD) and P Values.

Parameter	Group 1	Group 2	Group 3	Group 4	P Value
PI	1.37 ± 0.29	1.96 ± 0.18	2.62 ± 0.12	3.47 ± 0.27	<.01 ^a
GI	0.64 ± 0.21	1.15 ± 0.21	1.68 ± 0.22	2.31 ± 0.26	<.01 ^a
PPD	2.39 ± 0.5	3.18 ± 0.35	3.99 ± 0.32	5.02 ± 0.56	<.01 ^a
CAL	2.76 ± 0.55	3.58 ± 0.37	4.52 ± 0.38	5.58 ± 0.58	<.01 ^a
%BOP sites	29.17 ± 5.43	37.09 ± 5.24	55.44 ± 7	67 ± 12.36	<.01 ^a

Abbreviations: ANOVA, analysis of variance; GI, gingival index; PI, plaque index; PPD, probing pocket depth; CAL, clinical attachment level; %BOP, percentage bleeding on probing; SD, standard deviation.

^aStatistically significant at ($P < .05$); single factor ANOVA.

Table 3. Intergroup Relation for Clinical Parameters (Mean ± SD).

	Group 1 Vs Group 2	Group 1 Vs Group 3	Group 1 Vs Group 4	Group 2 Vs Group 3	Group 2 Vs Group 4	Group 3 Vs Group 4
PI	0.59 ± 0.08 ^a	1.25 ± 0.07 ^a	2.10 ± 0.11 ^a	0.66 ± 0.07 ^a	1.51 ± 0.11 ^a	0.85 ± 0.10 ^a
GI	0.51 ± 0.07 ^a	1.04 ± 0.08 ^a	1.67 ± 0.10 ^a	0.53 ± 0.10 ^a	1.16 ± 0.11 ^a	0.63 ± 0.11 ^a
PPD	0.80 ± 0.14 ^a	1.60 ± 0.14 ^a	2.64 ± 0.21 ^a	0.81 ± 0.15 ^a	1.84 ± 0.22 ^a	1.03 ± 0.22 ^a
CAL	0.82 ± 0.15 ^a	1.76 ± 0.16 ^a	2.82 ± 0.22 ^a	0.94 ± 0.17 ^a	2.00 ± 0.22 ^a	1.06 ± 0.23 ^a
%BOP	7.92 ± 1.87 ^a	26.28 ± 2.54 ^a	37.83 ± 4.24 ^a	18.35 ± 2.82 ^a	29.91 ± 4.41 ^a	11.56 ± 4.7 ^a

Abbreviations: GI, gingival index; PI, plaque index; PPD, probing pocket depth; CAL, clinical attachment level; %BOP, percentage bleeding on probing; SD, standard deviation.

^aStatistically significant at ($P < .05$); unpaired t test.

demographic characteristics and different clinical parameters between 2 groups. Single factor analysis of variance was performed to examine the differences between the 4 groups for different parameters; conventional P values $<.05$ were regarded as statistically significant. The sample size was determined in order to achieve study power of greater than 80%.

Results

Basic demographic and neurological features of the individuals in the study (AD) and control (ND) groups are presented in Table 1. Mean ± SD for ages of the individuals in the study (65.2 ± 7.3 years) and control groups (64.5 ± 9.4 years) were similar, and there were no differences found between individuals with AD and ND individuals with respect to gender. There were not much significant differences in the number of teeth present in individuals with AD (15.8 ± 3.6) and ND (16.2 ± 4.2) individuals and in the oral hygiene status between these groups. Significant differences

were found in MMSE scores between individuals with AD and ND individuals. The mean MMSE score for AD was 14.2 ± 8.4 when compared to 28.5 ± 1.2 for ND individuals.

Table 2 shows periodontal parameters assessed for individuals with AD and ND individuals, while the intergroup relation for the periodontal parameters is summarized in Table 3. Individuals with AD showed higher values of periodontal parameters when compared to ND individuals, while the periodontal condition worsened as the disease level progressed from mild to severe, as illustrated by increase in the parametric values from group 1 to group 4. Individuals with AD showed higher GI and PI values, higher values of PD, higher loss of attachment, and greater percentage of bleeding sites. The mean values for GI and PI also showed statistically significant difference between individuals with AD and ND individuals ($P < .01$), and the difference reached to a level of statistical significance from group 1 to group 4, when intergroup comparisons were considered. The mean PD and CAL values were

higher in individuals with AD and showed a statistically significant difference between individuals with AD and ND individuals ($P < .01$), while there was a significant increase in values from group 1 to group 4. The mean %BOP sites were much higher in individuals with AD when compared to ND individuals, and there was a significant difference between them ($P < .01$). The intergroup comparisons for %BOP sites also showed significant increase from group 1 to group 4.

Discussion

The results of the current study showed that individuals with AD showed a greater amount of periodontal breakdown when compared to the ND individuals. The mean values for all periodontal parameters (PD, CAL, GI, PI, and %BOP sites) were much higher in AD groups when compared to ND individuals. Moreover, the periodontal status of individuals with AD deteriorates as the disease progressed from mild to severe stage. Alzheimer's disease is a well-known disease, related to altered cognitive function which is thought to alter daily activities. Cognitive and motor deficits are accompanied by a gradual inability to perform adequate oral hygiene.²⁵ Impaired cognition, apathy, and apraxia in the middle stages of the disorder are liable for a disinterest and an inability to execute proper oral hygiene techniques, permitting food debris and plaque to remain undisturbed in the proximal and cervical areas of the residual dentition.²⁶ This can be considered as the major factor for the deteriorated periodontal health status in individuals with AD. There are few studies suggesting declined oral health status in individuals with AD and related dementia. A study by Syrjälä et al²⁷ showed that individuals with AD and those with other types of dementia had an increased likelihood of having carious teeth, teeth with deep periodontal pockets, and poor oral and denture hygiene, compared with nondemented persons. Another study by Chalmers et al²⁸ showed that individuals with dementia had significantly higher experiences of oral diseases and conditions at baseline and 1 year compared to participants without dementia: decreased use of dentures, increased prevalence of denture-related oral mucosal lesions, increased plaque accumulation, increased prevalence and experiences of coronal and root caries, and increased numbers of decayed retained tooth roots. These higher experiences of oral diseases and conditions were related to dementia severity. Individuals with AD have been shown to have poor oral hygiene by Ship.²⁹ In a large population survey that included over 6000 individuals (NHANES III) with a broad age range (20-59 and ≥ 70), associations were found between measures of periodontal disease (extent of bleeding on probing and extent of attachment loss ≥ 3 mm) and cognitive function.^{13,14} In accordance with these studies, our results suggest a declined periodontal health status in individuals with AD.

The results of the present study are in accordance with few other studies in which there was deterioration in oral health status of patients with AD. In a study by Machado et al,³⁰ patients with mild, moderate, and severe AD demonstrated poor oral health status, but there was no association between severity of their disease and the presence of oral health problems. In

another study, oral condition in patients with mild dementia was found to be poor than that in patients with normal cognitive function, thus suggesting that an alteration in cognitive function may negatively influence oral health status.³¹ Moreover, Lee et al,³² demonstrated that impaired cognitive function may adversely affect oral health status and overall quality of life. Thus, as the cognitive function declines from mild to severe AD, periodontal condition was found to deteriorate in the current study.

Tooth loss is considered as one of the risk factors associated with higher incidence of dementia.³³ Impaired cognition in AD may severely affect daily activities, and poor oral hygiene execution may lead to caries,¹⁰⁻¹² periodontitis,²⁷⁻²⁹ and eventual loss of teeth. A study reported in monozygotic twins suggested a strong association between tooth loss and the presence of AD.³⁴ Another study, examining oral health in a longitudinal study of aging in Catholic Nuns, reported that a lower number of teeth increased the risk of dementia.³³ Tooth loss may occur due to multiple factors including CP.¹³ Other causes may include carious lesions, trauma, and extractions due to prosthetic/orthodontic indications or impacted teeth. Periodontitis is the single major risk factor for tooth loss, and moderate to severe periodontitis may further complicate status of AD and related dementia. Thus, the results showed increased periodontal destruction in individuals with AD and also increase in severity of periodontal breakdown with progression of AD, supportive to this concept. Thus, it can be stated that individuals with AD have a declined periodontal status, and progression of AD severely affects the periodontal status. When treating individuals with AD, oral health care providers must develop timely, preventive, and therapeutic strategies compatible with the patient's physical and cognitive ability for prevention, early detection, and prompt treatment of periodontal disease. This was a preliminary study which took into consideration the clinical findings and the uncontrolled variables, including socioeconomic status, education, changes in oral hygiene regimens as dementia progresses, dietary changes, and access to dental care, need evaluation and could be considered in further studies. Further longitudinal, multicentered studies with larger sample sizes are required in future to confirm these findings and also to find a possible association between AD and CP.

Conclusion

Periodontal health status of the individuals with AD is closely related to their cognitive function. Cognitive and motor deficits are accompanied by a gradual inability to perform adequate oral hygiene. Individuals with AD show a greater extent of periodontal destruction which increases as the disease progression occurs. Thus, the individuals with AD require special attention toward overall oral health care, and the individuals and especially the caregivers must be educated regarding the same.

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Authors' Note

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Declaration of Conflicting Interests

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References

- Whitehouse PJ, Maurer K, Ballenger JF. *Concepts of Alzheimer Disease: Biological, Clinical, and Cultural Perspectives*. 1st ed. Baltimore: The Johns Hopkins University Press; 2000:83-103.
- Salmon D, Bondi M. Neuropsychological assessment of dementia. *Annu Rev Psychol*. 2009;60:257-282.
- Gazzaley A, Sheridan M, Cooney J, D'Esposito M. Age-related deficits in component processes of working memory. *Neuropsychology*. 2007;21(5):532-539.
- Conway AR, Kane MJ, Bunting MF, Hambrick DZ, Wilhelm O, Engle RW. Working memory span tasks: a methodological review and user's guide. *Psychon Bull Rev*. 2005;12(5):769-786.
- Small SA, Mayeux R. Alzheimer's disease and related dementias. In: Rowland LP, ed. *Merritt's Neurology*. Philadelphia, PA: Lippincott Williams & Wilkins, 2000:633.
- World Alzheimer's Report 2010, Alzheimer's disease International.
- World Health Organization. *Neurological Disorders: Public Health Challenges*. Switzerland: World Health Organization; 2006:204-207.
- Kim JM, Stewart R, Prince M, et al. Dental health, nutritional status and recent-onset dementia in a Korean community population. *Int J Geriatr Psychiatry*. 2007;22(9):850-855.
- Gambassi G, Landi F, Lapane KL, Sgadari A, Mor V, Bernabei R. Predictors of mortality in patients with Alzheimer's disease living in nursing homes. *J Neurol Neurosurg Psychiatry*. 1999;67(1):59-65.
- Chalmers JM, Carter KD, Fuss JM, Spencer AJ, Hodge CP. Caries experience in existing and new nursing home residents in Adelaide, Australia. *Gerodontology*. 2002;19(1):30-40.
- Ellefsen B, Holm-Pedersen P, Morse DE, Schroll M, Andersen BB, Waldemar G. Caries prevalence in older persons with and without dementia. *J Am Geriatr Soc*. 2008;56(1):59-67.
- Syrjala AM, Ylostalo P, Sulkava R, Knuutila M. Relationship between cognitive impairment and oral health: results of the health 2000 health examination survey in Finland. *Acta Odontol Scand*. 2007;65(2):103-108.
- Al-Shammari KF, Al-Khabbaz AK, Al-Ansari JM, Neiva R, Wang HL. Risk indicators for tooth loss due to periodontal disease. *J Periodontol*. 2005;76(11):1910-1918.
- Noble JM, Borrell LN, Papapanou PN, Elkind M, Scarmeas N, Wright C. Periodontitis is associated with cognitive impairment among older adults: analysis of NHANES-III. *J Neurol Neurosurg Psychiatry*. 2009;80(11):1206-1211.
- Stewart R, Sabbah W, Tsakos G, D'Alto F, Watt RG. Oral health and cognitive function in the Third National Health and Nutrition Examination Survey (NHANES III). *Psychosom Med*. 2008;70(8):936-941.
- Buhlin K, Gustafsson A, Pockley AG, Frostegard J, Klinge B. Risk factors for cardiovascular disease in patients with periodontitis. *Eur Heart J*. 2003;24(23):2099-2107.
- Lakschevitz F, Aboodi G, Tenenbaum H, Glogauer M. Diabetes and periodontal diseases: interplay and links. *Curr Diabetes Rev*. 2011;7(6):433-439.
- Schwarz J, Heimhülger E, Storch A. Increased periodontal pathology in Parkinson's disease. *J Neurol*. 2006;253(5):608-611.
- National Institute of Neurological and Communicative Disorders and Stroke and the Alzheimer's Disease and Related Disorders Association (NINCDS-ADRDA) workgroup recommendations, 1988.
- Mungas D. In-office mental status testing: a practical guide. *Geriatrics*. 1991;46(7):54-66.
- Quigley GA, Hein JW. Comparative cleansing efficiency of manual and power brushing. *J Am Dent Assoc*. 1962;65:26-29.
- Turesky S, Gilmore ND, Glickman I. Reduced plaque formation by chloromethyl analogue of vitamin C. *J Periodontol*. 1970;41(1):41-43.
- Loe H, Silness J. Periodontal diseases in pregnancy: prevalence and severity. *Acta Odontol Scand*. 1963;21:533-551.
- Glavind L, Loe H. Errors in the clinical assessment of periodontal destruction. *J Periodontol Res*. 1967;2(3):180-184.
- Gitto CA, Moroni MJ, Terezhalmay GT, Sandu S. The patient with Alzheimer's disease. *Quintessence Int*. 2001;32(3):221-231.
- Hatipoglu MG, Kabay SC, Güven G. The clinical evaluation of the oral status in Alzheimer-type dementia patients. *Gerodontology*. 2011;28(4):302-306.
- Syrjälä AM, Ylostalo P, Ruoppi P, et al. Dementia and oral health among subjects aged 75 years or older. *Gerodontology*. 2012;29(1):36-42.
- Chalmers JM, Carter KD, Spencer AJ. Oral diseases and conditions in community-living older adults with and without dementia. *Spec Care Dentist*. 2003;23(1):7-17.
- Ship JA. Oral health of patients with Alzheimer's disease. *J Am Dent Assoc*. 1992;123(1):53-58.
- Machado MC, Lopes GH, Marchini L. Oral health of Alzheimer's patients in São José dos Campos, Brazil. *Geriatr Gerontol Int*. 2012;12(2):265-270.
- Sumi Y, Ozawa N, Michiwaki Y, Washimi Y, Toba K. Oral conditions and oral management approaches in mild dementia patients [in Japanese]. *Nihon Ronen Igakkai Zasshi*. 2012;49(1):90-98.
- Lee KH, Wu B, Plassman BL. Cognitive function and oral health-related quality of life in older adults. *J Am Geriatr Soc*. 2013;61(9):1602-1607.
- Stein PS, Desrosiers M, Donegan SJ, Yepes JF, Kryscio RJ. Tooth loss, dementia and neuropathology in the Nun Study. *J Am Dent Assoc*. 2007;138(10):1314-1322.
- Gatz MJ, Fratiglioni L, Johansson B, Berg S, Reynolds CA, Pedersen NL. Potentially modifiable risk factors for dementia in identical twins. *Alzheimers Dement*. 2006;2(2):110-117.