Cytological and cytometric analysis of oral mucosa in patients with Alzheimer’s and Parkinson’s disease

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Background: Alzheimer’s disease (AD) and Parkinson’s disease (PD) are the two most common neurodegenerative diseases. Recent studies have sought to identify precursor symptoms of AD and PD that occur before the onset of the disease. We evaluated changes in the oral mucosa of patients with AD and PD using a stereological method.

Patients and methods: The study included 29 patients with AD, 30 patients with idiopathic PD, and 30 healthy volunteers. Brush biopsies were obtained from all participants, and the nucleator method was used to estimate the volume of cells obtained from the buccal mucosa.

Results: Cytomorphometric analysis revealed that the nuclear volume was 484.39 ± 171.10 µm³ in the AD group, 509.71 ± 132.26 µm³ in PD patients, and 509.30 ± 100.21 µm³ in the control group. The cytoplasmic volume was 115,456.60 ± 30,664.98 µm³ in the AD group, 103,097.93 ± 25,034.65 µm³ in PD patients, and 109,528.45 ± 28,381.43 µm³ in the control group. The nuclear and cytoplasmic volumes were not significantly different among groups (P > 0.05).

Conclusion: The cytomorphic analysis revealed no significant differences in the cytoplasmic and nuclear volumes of buccal cells obtained from patients with AD and PD and healthy volunteers.

Keywords: Alzheimer’s disease, Parkinson’s disease, exfoliative cytology, stereology, oral mucosa

Introduction

Alzheimer’s disease (AD), the most common cause of dementia, is characterized by an insidious onset and progressive impairment of short-term memory and other cognitive disorders linked to the degeneration of neurons in the limbic system and neocortical areas. 1

Parkinson’s disease (PD) is the second most common neurodegenerative disease after AD. It is characterized by movement disorders such as rigidity, bradykinesia, postural imbalance, and unilateral resting tremor caused by the degeneration of neuromelanin-containing neurons in the ventral region of the substantia nigra pars compacta and locus coeruleus in the brain stem. 2 Both diseases are chronic and progressive, and in the advanced stages, patients are completely dependent on caregivers, imposing a serious social and economic burden. 3,4

AD and PD are diagnosed clinically based on detailed medical histories and thorough neurological examinations. Definitive diagnoses can only be made post mortem or by invasive biopsies to identify disease-specific pathological changes. 5,6

Several recent studies have investigated various precursors of AD and PD to identify individuals at risk before the onset of symptoms. For instance, olfactory
disorders, rapid-eye-movement sleep behavior disorder, and constipation are thought to be precursor symptoms of PD. Moreover, gastrointestinal tract biopsies have been studied for specific markers of PD. Although several studies have investigated changes in nasal mucosa, fibroblasts, and erythrocytes to identify precursors of AD, no specific marker of the disease has been found.

Recent studies have examined oral mucosa cells for precursors of neurological diseases because the cells are derived from ectodermal tissue and are thought to be embryologically related to the central nervous system.

A previous study found that gene expression of the tau protein was increased in the oral epithelial cells of patients with AD. In another study, it was determined that telomere analysis of buccal cells differed between AD stages. However, the morphological and cytological structure of the oral mucosa has only been analyzed in one study. In a cytological analysis of the oral mucosa of 10 AD patients, de Oliveira et al. found an increase in the nucleus area of the buccal mucosa and cell groups in the tongue and a decrease in the nuclear/cyttoplasmic area ratio. However, the authors noted that the changes they observed could have been caused by diabetes mellitus (DM) or B12, folic acid, and iron deficiencies, indicating that further study was needed to clarify their findings.

Exfoliative cytology is a readily used, noninvasive, inexpensive, and rapid procedure based on the microscopic evaluation of fixed epithelial cells.

Several methods may be used to investigate exfoliated cells; however, stereology is one of the most reliable methods for objective morphometric and three-dimensional analyses.

We aimed to investigate whether the oral exfoliative cytology can be a tool for early diagnosis of AD and PD.

Patients and methods

The study included 29 patients with AD diagnosed according to the National Institute of Neurological and Communicative Disorders and Stroke (NINCDS) and Alzheimer’s Disease and Related Disorders Association (ARDA) criteria, 30 patients with idiopathic PD diagnosed according to the Parkinson Society Brain Bank criteria, and 30 healthy volunteers. All patients were diagnosed at the Kafkas University Faculty of Medicine Neurology Clinic. The study was approved by the Kafkas University Local Ethics Committee and was performed in accordance with the Declaration of Helsinki. All participants provided written informed consent.

Patients who smoked; were alcohol dependent; had iron, B12, or folate deficiencies; had DM; had undergone chemotherapy or radiotherapy; or had an oral mucosa infection were excluded from the study.

The modified Hoehn and Yahr scale was used to identify the stage, and the Unified Parkinson’s Disease Rating Scale (UPDRS) was used to determine the severity of idiopathic PD. The Mini-Mental State Examination (MMSE) and Clinical Dementia Scale were used to assess the patients with AD.

Oral mucosa samples were taken from all participants after they underwent a detailed neurological examination. The oral mucosa was dried with a gauze swab; then, smears were taken from the clinically healthy buccal mucosa using a cotton swab and transferred to glass slides. The samples were immediately fixed in 95% ethyl alcohol, stained using the Papanicolaou method, and examined under a light microscope.

Stereological estimation of cytoplasmic and nuclear volumes

The samples were placed under a microscope with a motor-driven stage and cells were projected onto a monitor via a camera at ×200 magnification. A minimum of 200 randomly selected epithelial cells per patient were measured. The nuclear (NV) and cytoplasmic (CV) volumes were evaluated for each cell using Stereologist software (MicroBrightField, Williston, VT, USA).

The volume of cells obtained from the buccal mucosa samples was estimated using the nucleator method according to the following formula:

\[ V_{(n)} = \frac{4\pi}{3} l^n \]

where \( V_{(n)} \) is the number-weighted volume, \( l \) is the length of intercepts, and \( n \) is the number of nucleator estimates (Figure 1).

Statistical analysis

The Statistical Package for the Social Sciences version 22.0 for Windows (IBM Corporation, Armonk, NY, USA) was used to conduct the statistical tests. The normal distribution of data was assessed using the Kolmogorov–Smirnov test, and nonparametric data were assessed using the Mann–Whitney U-test. Differences among subgroups were evaluated using the Kruskal–Wallis test. \( P<0.05 \) was deemed to indicate statistical significance.

Results

The study included 29 patients with AD (16 males, 13 females), 30 patients with idiopathic PD (16 males, 14 females),
alterations in oral mucosa in patients with AD and PD

and 30 healthy controls (15 males, 15 females). The AD patient ages ranged from 60 to 89 years (mean, 78.93±7.47 years), the PD patients were aged 62–85 years (mean, 76.52±7.29 years), and the healthy controls were aged 60–88 years (mean, 79.49±8.13 years). The age and sex of the participants were not significantly different among groups (P>0.05).

In the AD group, the mean MMSE score was 14.32±4.23, and the mean Clinical Dementia Scale score was 2.59±0.68.

Figure 1 (A, B) Histologic views of exfoliative cell samples. (C) Stereological formulas.
The mean modified Hoehn and Yahr and UPDRS scores in the idiopathic PD patients were 2.41±0.53 and 101.23±8.21, respectively. The sociodemographic data of the patients are shown in Table 1.

Thirteen of the 29 patients were not using any medication for the AD due to different reasons; five of them were not using medication due to various side effects, and eight of them were newly diagnosed patients (these patients had AH symptoms for at least 1 year); on the other hand, 16 of the patients were using different medications; 14 of them were using memantine-donepezil and two of them were using donepezil. There were no statistically significant differences in NV or CV between the groups with and without drug use. However, according to MMSE, the patients were grouped as mild (group 1), moderate (group 2), or severe (group 3). Since the number of severe stage patients was insufficient, they could not be included in the statistics. No significant differences among groups (AD patients, 115,456.60±30,664.98 µm³; PD patients, 103,097.93±25,034.65 µm³; and controls, 109,528.45±28,381.43 µm³; P>0.05).

### Discussion

Our microscopic and cytomorphometric analyses of buccal cells in patients with AD and PD, the two most common neurodegenerative diseases, and healthy volunteers revealed no differences in NV or CV among groups.

Exfoliative cytology has been used for the screening and the early diagnosis of malignancies for several years. Buccal mucosa cells, in particular, are readily obtained using this inexpensive, noninvasive procedure, which is well tolerated by patients. Stereological assessment of the buccal cells provides objective information based on mathematical and statistical principles. Thus, recent studies have investigated cytological changes in oral mucosa cell-associated diseases other than cancer.

Keles et al found a significant increase in NV and CV values in exfoliated buccal mucosa cells after renal transplant (RT). Given the increased risk of malignancy in RT recipients, the authors concluded that the observed changes may have been related to the malignant transformation of oral mucosa cells.

Previous cytomorphometric analyses performed in patients with DM have revealed an increase in NV and the nuclear/cytoplasmic area ratio in oral mucosa cells.

<table>
<thead>
<tr>
<th>Sociodemographic data</th>
<th>AD patients (n=29)</th>
<th>PD patients (n=30)</th>
<th>Controls (n=30)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (mean years)</td>
<td>78.93±7.47</td>
<td>76.52±7.29</td>
<td>79.49±8.13</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>Sex (male/female)</td>
<td>16/13</td>
<td>16/14</td>
<td>15/15</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>MMSE</td>
<td>14.32±4.23</td>
<td>–</td>
<td>–</td>
<td></td>
</tr>
<tr>
<td>CDS</td>
<td>2.59±0.68</td>
<td>–</td>
<td>–</td>
<td></td>
</tr>
<tr>
<td>Modified Hoehn and Yahr score</td>
<td>–</td>
<td>2.41±0.53</td>
<td>–</td>
<td></td>
</tr>
<tr>
<td>UPDRS score</td>
<td>–</td>
<td>101.23±8.21</td>
<td>–</td>
<td></td>
</tr>
<tr>
<td>Time of disease (mean years)</td>
<td>8.4±2.41</td>
<td>9.3±3.2</td>
<td>–</td>
<td>0.05</td>
</tr>
</tbody>
</table>

**Abbreviations:** AD, Alzheimer’s disease; CDS, Clinical Dementia Scale; MMSE, Mini-Mental State Examination; PD, Parkinson’s disease; UPDRS, Unified Parkinson’s Disease Rating Scale.

<table>
<thead>
<tr>
<th>NV and CV values according to AD stages</th>
<th>AD group 1 (n=14)</th>
<th>AD group 2 (n=13)</th>
<th>Controls (n=30)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>NV</td>
<td>482.19±215.10</td>
<td>470.39±122.75</td>
<td>509.30±100.21</td>
<td>0.889</td>
</tr>
<tr>
<td>CV</td>
<td>100,423.20±40,224.48</td>
<td>115,212.60±35,552.90</td>
<td>109,528.45±28,381.43</td>
<td>0.669</td>
</tr>
</tbody>
</table>

**Notes:** According to MMSE, the patients were grouped as mild (group 1), moderate (group 2), or severe (group 3). Since the number of severe stage patients was insufficient, they could not be included in the statistics.

**Abbreviations:** AD, Alzheimer’s disease; CV, cytoplasmic volume; NV, nuclear volume.
This finding may be explained by nuclear hypertrophy, which is an adaptive response to the cellular damage caused by DM.

Buccal mucosa cells are derived from ectodermal tissue during the embryological period of development, similar to brain and skin epithelial cells. Thus, it may be that buccal mucosa cells reflect changes in brain tissue. Accordingly, we searched the literature for previous cytological and cytometric analyses of buccal mucosa cells in patients with AD. de Oliveira et al performed a cytornorphometric analysis of buccal mucosa cells in 10 AD patients and nine age-matched control patients. They found a significant reduction in intermediate cells (P<0.05) and in the nuclear/cytoplasmic area ratio (P<0.0001) in AD patients, compared with the control patients. However, the authors acknowledged that their sample size was small, and the changes observed could have been caused by type 2 DM or deficiencies in vitamin B12, folic acid, and iron. To our knowledge, no such study has been conducted in patients with PD.

Our sample size was larger than that of de Oliveira et al, and we excluded participants with DM or vitamin B12, folic acid, and iron deficiencies; nevertheless, we found no significant differences in buccal cell cytology among the AD, PD, and control groups.

**Limitation**

However, a limitation of our study is that we did not perform analyses according to buccal cell types.

**Conclusion**

In our study, we could not reach a conclusion that oral cytological evaluation helped early diagnosis. Our findings suggest that further study is needed to clarify the effect of neurodegenerative diseases on buccal mucosa cells, and whether such changes are useful clinical predictors of neurodegenerative disease.

**References**


