Two cases of non-fluent variant primary progressive aphasia with different pathological diagnoses

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ABSTRACT

Non-fluent variant primary progressive aphasia (nfvPPA), a subtype of frontotemporal lobar dementia syndrome, has been proven to have various pathological diagnoses. A 63-year-old woman and 71-year-old man separately visited our clinic for language dysfunction. Both patients showed non-fluent speech. The female patient showed personality change accompanied by language dysfunction, while the male patient had parkinsonian symptoms such as bradykinesia and cogwheel rigidity. Both patients were clinically diagnosed with nfvPPA. Several years after the first visit, the patients died, and a brain autopsy was performed. On postmortem examination, the female patient was pathologically diagnosed with Pick’s disease, while the male patient was diagnosed with progressive supranuclear palsy. Our report suggests that nfvPPA patients might show distinct clinical features depending on underlying pathologies.

Keywords: Autopsy; Pick disease of the brain; Primary progressive nonfluent aphasia; Supranuclear palsy, progressive
INTRODUCTION

Frontotemporal lobar dementia (FTD) syndromes include a behavioral variant (bvFTD) and two subtypes of primary progressive aphasia [1,2]. These syndromes are united by overlapping clinical and anatomical features and their link to the underlying frontotemporal lobar degeneration pathology (FTLD) [1,2]. FTLD is characterized by neuronal and glial inclusions immunoreactive for hyperphosphorylated tau (FTLD-tau), or TAR DNA-binding protein of 43 kDa (TDP-43, FTLD-TDP) [3,4]. Despite attempts to associate FTD clinical syndromes with specific FTLD neuropathological diagnoses, no such correlation has yet been proven invariant [5-7]. In particular, non-fluent variant primary progressive aphasia (nfvPPA) has been proven to have various pathological diagnoses. Over 60% of nfvPPA cases show FTLD-tau, including Pick's disease (PiD), progressive supranuclear palsy (PSP), and corticobasal degeneration. Most of the remaining cases show TDP type A and Alzheimer's disease pathology [6,8]. Herein, we report two patients clinically diagnosed with nfvPPA that displayed different pathological findings at autopsy.

CASE REPORTS

The collection of the data was conducted as set forth in the Declaration of Helsinki. This study was approved by the Institutional Review Board of Samsung Medical (IRB Rile No. 2016-11-111) and we obtained informed consent from the participant and their next of kin (daughter and spouse).

Case 1

A 63-year-old right-handed woman visited our clinic due to language dysfunction and personality change. She had started to show effortful, non-fluent speech and a decreased amount of speech 1 year prior to her first visit. After 1 year, she showed abulia and blankly watched TV all day. She occasionally got upset for no reason. There was no familial history of dementia. She had hypertension and hyperlipidemia, and was on hemodialysis for chronic renal failure.

On neurological examination, the patient showed motor impersistence and grasp reflex on the bilateral upper extremities. She scored 22 out of 30 on the Mini-Mental State Examination (MMSE). There were no signs of Parkinsonism. Detailed neuropsychological tests revealed that her attention, language, and verbal memory were decreased; especially her frontal executive function, which was severely decreased.

Table 1. Western Aphasia Battery (WAB) results

| Category of K-WAB (highest possible score) | Case 1 2013.06.15 | Case 2 2009.12.14 | Case 2 2011.12.07 |
|-------------------------------------------|------------------|------------------|------------------|
| Spontaneous speech (20)                   | 8                | 17               | 14               |
| Information content (10)                  | 3.5              | 9                | 7                |
| Fluency (10)                              | 4.5              | 8                | 7                |
| Comprehension (10)                        | 6.05             | 8.75             | 7.25             |
| Yes/No question (3)                       | 1.65             | 2.85             | 2.55             |
| Auditory word recognition (3)             | 2.6              | 2.75             | 2                |
| Sequential commands (4)                   | 1.8              | 3.15             | 2.55             |
| Repetition (10)                           | 5.3              | 8.5              | 7.2              |
| Naming (10)                               | 4.3              | 5.6              | 7.1              |
| Confrontation naming (6)                  | 4.3              | 8.4              | 5.1              |
| Generative naming (2)                     | 0                | 0.8              | 0.4              |
| Sentence completion (1)                   | 0                | 1                | 0.6              |
| Responsive speech (1)                     | 0                | 1                | 1                |
| Reading (10)                              | 8.2              | 4.7              |                  |
| Writing (10)                              | 7.6              | 8.3              |                  |

On the Western Aphasia Battery (WAB), she demonstrated non-fluent speech and a lack of verbal response. The average number of word segments per utterance was one; word comprehension was mildly decreased, but complex sentence comprehension was remarkably decreased, confrontational naming ability was severely decreased, and repetition was...
mildly decreased. In particular, she showed word-finding difficulty and delayed word retrieval. There was no effortful speech, agrammatism, or dysarthria. These results indicated a moderate degree of Broca’s aphasia (Table 1). Her fluid-attenuated inversion recovery (FLAIR) images showed bilateral frontal and anterior temporal atrophy and periventricular white matter hyperintensities in the anterior and posterior portion. On sagittal T1-weighted images, there is no significant midbrain atrophy. Gradient echo images showing strictly multiple lobar microbleeds and cortical superficial siderosis in the left parietal cortex. [18F]-fluorodeoxyglucose positron emission tomography showing moderate hypometabolism in the bilateral frontal cortex which is more severe on the left.

Fig. 1. Brain images of a 63-year old woman (A) and 71-year old man (B). (A) Axial fluid-attenuated inversion recovery (FLAIR) magnetic resonance imaging (MRI) showing bilateral frontal and anterior temporal atrophy. (B) Axial FLAIR MRI images showing diffuse brain atrophy and periventricular white matter hyperintensities in the anterior and posterior portion. On sagittal T1-weighted images, there is no significant midbrain atrophy. Gradient echo images showing strictly multiple lobar microbleeds and cortical superficial siderosis in the left parietal cortex. [18F]-fluorodeoxyglucose positron emission tomography showing moderate hypometabolism in the bilateral frontal cortex which is more severe on the left.

Postmortem examination showed a decreased brain weight of 988 g (standard for age: 1,360 g). Gross examination revealed moderate to severe gyral atrophy in the bilateral frontal lobes, and mild gyral atrophy in the bilateral parietal and medial temporal lobes (Fig. 2A). Histological examination revealed microvacuolation with severe neuronal loss and gliosis in the frontal cortex layers 2 and 3 (Fig. 2B). Mild neuronal loss and gliosis were present in the parietal cortex layer 2, temporal cortex layer 2, amygdala, basal nucleus, basal ganglia, and dentate nucleus. Tau-positive intraneuronal cytoplasmic inclusions (Pick bodies), identified by tau and 3-repeat tau stains, were located in the frontal, parietal, and temporal cortex layers. Pick bodies were also found in the dentate gyrus, hippocampus, subiculum, basal ganglia, thalamus, periaqueductal gray matter, locus coeruleus, and pontine nucleus (Fig. 2C). In the absence of TDP-43 pathology...
and α-synucleinopathy, a pathological diagnosis of FTLD-tau, PiD, was rendered.

**Case 2**

A 71-year-old right-handed man visited our clinic because of language dysfunction, left hand clumsiness, and memory impairment. He had started to show slow speech velocity, stuttering, and word-finding difficulty half a year prior to his first visit. There was no familial history of dementia. He was suffering from hypertension.

On neurological examination, the patient showed ideomotor apraxia and poor performance in fist-edge-palm. He also experienced bradykinesia in the left upper limb. He scored 25 out of 30 on MMSE. Detailed neuropsychological tests revealed that his attention, language, verbal memory, and frontal executive function, except visuospatial function and visual memory, were decreased. On WAB, he was overall fluent, but showed frequent hesitation, halting, repetition of syllables, phonemic paraphasia, and word-finding difficulty. Word comprehension, repetition, and naming were slightly decreased. These results indicate a mild degree of anomic aphasia (Table 1). His apolipoprotein E genotype was ε3/ε3.

On FLAIR images, diffuse brain atrophy and periventricular white matter hyperintensities in the anterior and posterior portions were seen. On gradient echo images, strictly multiple lobar microbleeds and cortical superficial siderosis in the left parietal cortex were observed, suggesting probable cerebral amyloid angiopathy (CAA). [18F]-fluorodeoxyglucose positron emission tomography (FDG-PET) revealed moderate hypometabolism in the bilateral frontal cortex, which was more severe on the left side (Fig. 3). Based on his clinical symptoms, WAB, detailed neuropsychological tests, brain

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**Fig. 2.** Postmortem examination of a 63-year-old woman. (A) Severe gyral atrophy in bilateral frontal lobes with mild gyral atrophy of bilateral parietal and medial temporal lobes (scale bar=10 cm). (B) Microvacuolation, neuronal loss, and gliosis in frontal cortex layer 2 and 3 (H&E stain, ×400; scale bar=100 μm). (C) Tau-positive intracytoplasmic inclusion in dentate gyrus (Tau [left], 3-repeat Tau stain [mid], and 4-repeat Tau [right], ×400; scale bar=100 μm).
MRI, and FDG-PET, the patient was clinically diagnosed with nfvPPA. Three years after his first visit, he showed cogwheel rigidity in the bilateral upper and lower limbs. However, the patient did not show axial rigidity, postural instability and vertical gaze limitation. The overall language function worsened, and he demonstrated phonemic paraphasia, neurogenic stuttering, and perseveration. He did not show dysarthria or apraxia of speech. These results indicate a moderate degree of anomic aphasia (Table 1). Four years after his first visit, he showed impatience and repetitive motor behaviors, such as tearing papers. Six years after his first visit, he was unable to walk anymore and died of pneumonia at the age of 77 years. After his death, his body underwent a brain autopsy at the Samsung Medical Center.

Postmortem examination showed a decreased brain weight of 940 g (standard for age: 1,300 g). Gross examination revealed marked atrophy of the bilateral precentral gyrus with moderate atrophy of the bilateral frontal and superior temporal gyri (Fig. 3A). On microscopic examination, tufted astrocytes were identified in the middle frontal gyrus, inferior parietal lobule, and occipital lobe. Globoid tangles and pretangles were observed in the temporal lobe, putamen, subthalamic nucleus, pons, and medulla oblongata (Fig. 3C). β-Amyloid plaques with amyloid angiopathy were present in the occipital cortex, amygdala, hippocampus, and cerebellum. Atherosclerosis in the vessels of the circle of Willis and hemosiderin pigmentation in the left inferior parietal lobule, suggestive of old subarachnoid hemorrhage, were noted. Microinfarcts were identified in the left frontal, temporal, and parietal lobes. Overall, the patient was diagnosed with FT-
Non-fluent primary progressive aphasia

LD-tau, PSP with Alzheimer’s disease (Braak Stage II), CAA, and cerebrovascular disease.

DISCUSSION

Here, we report two clinically diagnosed nfvPPA cases. Both patients showed decreased fluency and effortful and decreased speech. As the disease progressed, they showed not only language dysfunction but also other symptoms, such as behavioral changes or motor dysfunction. However, they demonstrated showed several distinguished characteristics according to the pathological diagnosis. The patient pathologically diagnosed with PiD showed early age at onset and initial personality change accompanied by language dysfunction. On the other hand, the patient with PSP showed late age at onset and initial parkinsonian features accompanied by language dysfunction.

Patients with nfvPPA might show distinct clinical features depending on the underlying pathologies. Clinico-pathological association between nfvPPA clinical syndrome and FTLD neuropathological diagnosis is not specific; nfvPPA is associated with multiple pathologies, including tauopathy, TDP type A pathology, and Alzheimer’s disease pathology [6,8]. The first case was pathologically diagnosed as PiD. PiD, a rare subtype of FTLD-tau, is characterized by right dominant frontotemporal atrophy and clinical features of bvFTD. However, a few patients with PiD have showed left dominant frontal and peri-Sylvian atrophy and clinical features of nfvPPA. Gross neuropathological findings of PiD show circumscribed lobar atrophy, and microscopic findings are characterized by neuronal loss, gliosis, and argyrophilic and round intraneuronal inclusions (Pick’s body). Pick’s body is composed of a tau protein containing three microtubule-binding repeats (3R) [9].

PSP primarily presents with atypical parkinsonism with axial rigidity, frequent falls, postural instability, and progressive gaze palsy [9]. In particular, 80% to 90% of clinically suspected progressive supranuclear palsy syndrome (PSPS) show PSP pathology. However, approximately 10% of PSP presents the clinical features of nfvPPA, similar to our second case [6]. The cardinal neuropathological findings consisted of tufted astrocytes, globose neurofibrillary tangles, and oligodendroglial coiled bodies. These astrocytic, neuronal, and oligodendroglial lesions of PSP are mainly composed of tau protein containing four microtubule-binding repeats (4R) [9].

In conclusion, clinically diagnosed nfvPPA patients are expected to have various pathological diagnoses. Patients with nfvPPA might show distinct clinical features depending on the underlying pathologies.

CONFLICTS OF INTEREST

No potential conflict of interest relevant to this article was reported. Sang Won Seo has been editorial board of P&FM since December 2017. He was not involved in the review process of this case report.

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