



# Relationship Between Swallowing Function and Maximum Phonation Time in Patients With Parkinsonism

Eu Jeong Ko, MD<sup>1</sup>, Minji Chae, MD<sup>1</sup>, Sung-Rae Cho, MD, PhD<sup>1,2,3,4</sup>

<sup>1</sup>Department of Rehabilitation Medicine and Research Institute of Rehabilitation Medicine,

<sup>2</sup>Brain Korea 21 Plus Project for Medical Science, <sup>3</sup>Avison Biomedical Research Center,

<sup>4</sup>Rehabilitation Institute of Neuromuscular Disease, Yonsei University College of Medicine, Seoul, Korea

**Objective** To identify the relationship between maximum phonation time (MPT) and swallowing function, as well as the elements of swallowing, in order to provide a rationale for speech therapy in patients with Parkinsonism manifesting dysphagia.

**Methods** Thirty patients with Parkinsonism who underwent speech evaluation and videofluoroscopic swallowing study (VFSS) were recruited. The MPT, the longest periods of sustained pronunciation of /aa/, was evaluated. The VFSS was evaluated using Penetration Aspiration Scale (PAS), National Institutes of Health-Swallowing Safety Scale (NIH-SSS), and Videofluoroscopic Dysphagia Scale (VDS). The relationship between dysphagia scales and MPT was analyzed using Pearson correlation. The difference in VDS variables between subgroups (Parkinson disease or Parkinsonian syndrome, independent or dependent ambulation, and normal or abnormal MPT) and the difference in MPT between subgroups based on the VDS variables were analyzed using the independent t-test.

**Results** Bolus formation and laryngeal elevation functions were significantly higher in the normal MPT group compared with the impaired group. In the VDS variables, patients with intact bolus formation, oral transit time, pharyngeal swallow triggering, and laryngeal elevation showed significantly longer MPTs compared with the impaired groups. In addition, MPT was significantly correlated with the VDS and modestly correlated with the NIH-SSS, but not the PAS, suggesting that phonatory function is related to the oropharyngeal swallowing function, but not directly to the aspiration itself.

**Conclusion** The correlation between MPT and several swallowing-related elements was identified, indicating an interactive correlation between swallowing and phonation. This result justifies voice therapy as a treatment for dysphagia in patients with Parkinsonism.

**Keywords** Parkinson disease, Parkinsonian disorders, Deglutition disorders, Phonation

Received July 26, 2017; Accepted September 18, 2017

Corresponding author: Sung-Rae Cho

Department of Rehabilitation Medicine and Research Institute of Rehabilitation Medicine, Yonsei University College of Medicine, Rehabilitation Hospital 5th Floor, 50-1, Yonsei-ro, Seodaemun-gu, Seoul 03722, Korea. Tel: +82-2-2228-3715, Fax: +82-2-363-2795, E-mail: srcho918@yuhs.ac  
ORCID: Eu Jeong Ko (<http://orcid.org/0000-0001-8796-9655>); Minji Chae (<http://orcid.org/0000-0002-0590-2113>); Sung-Rae Cho (<http://orcid.org/0000-0003-1429-2684>).

© This is an open-access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (<http://creativecommons.org/licenses/by-nc/4.0>) which permits unrestricted noncommercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

Copyright © 2018 by Korean Academy of Rehabilitation Medicine

## INTRODUCTION

Parkinson disease (PD) is a complex, progressive, neurodegenerative disorder resulting in a wide range of deficits [1,2]. Speech and swallowing function are impaired, even in the early stages, significantly affecting health and the quality of life [3].

The cause of dysphagia in patients with PD is not clearly understood; however, a dysfunction involving the central pattern generator for swallowing in the brainstem and degeneration of the substantia nigra are possible causes that contribute to damaged non-dopaminergic neural networks [4,5]. Such damage leads to oral and pharyngeal phase dysfunction, which is characterized by abnormal bolus formation, repetitive tongue pumping, delayed swallowing reflex, decreased elevation of the hyolaryngeal complex, residues in both the vallecular and pyriform sinuses, and limited pharyngeal peristalsis [6].

Dysphagia not only affects the nutritional status of patients, but also their social interaction, susceptibility to fatigue, quality of life, and many other aspects. It may also lead to aspiration pneumonia, which is a major cause of death in patients with PD [1,4,7].

Conventional treatments such as bolus modification, postural and airway protective maneuvers, and pharmacological interventions, as well as other treatments such as expiratory muscle strength training, thermal-tactile stimulation, electrical stimulation, logopedic dysphagia treatment, and surgical intervention have been used to manage dysphagia in patients with PD. However, the long-term effects have yet to be verified, while a few studies have shown short-term benefits [4,8].

Since organs related to swallowing and speech are structurally and neurologically linked, speech therapy has been used to treat dysphagia without any evidence of sustained long-term benefits [4,9].

This study was conducted in patients with PD and Parkinsonian syndrome (PS) to identify the relationship between the maximum phonation time (MPT) and swallowing function, as well as the elements of swallowing, to provide a rationale for the use of voice therapy for the treatment of dysphagia.

## MATERIALS AND METHODS

### Subjects

Thirty patients diagnosed with PD or PS, who visited the clinic of the Department of Rehabilitation Medicine from January 2007 to December 2016 and underwent speech evaluation and videofluoroscopic swallowing study (VFSS), were included in this study. VFSS and speech evaluation were conducted simultaneously on eligible and selected patients. Patients with secondary Parkinsonism related to drugs, infection, or hydrocephalus were excluded. Patients with other neurological deficits and medical diseases, such as stroke or severe cardiopulmonary disease that may affect dysphagia were also excluded.

Of the 30 patients, 10 manifested PD and 20 had PS. The 20 patients with PS included 15 manifesting multiple system atrophy, 2 cases of dementia with Lewy body, and 1 with corticobasal degeneration. Of the 30 patients, the MPT was found normal in 9 and abnormal in 21. The baseline characteristics, including ambulatory and functional status and dysphagia scale scores, are listed in Table 1. This study was approved by the Institutional Review Board of Yonsei University Health System (No. 4-2014-0528).

### Methods

Speech evaluation and VFSS were conducted to compare swallowing and speech. The speech evaluation was conducted by speech therapists in the form of one-on-one sessions in the treatment units. The MPT was measured with the pronunciation of /aa/ as the primary value. The MPT represent the maximum time (in seconds) for which an individual can sustain a vowel sound when uttered on one deep breath at a reasonably comfortable pitch and loudness. MPT is used to indirectly evaluate the laryngeal function and vocal cord vibration [10,11]. It was widely used since it is easier to access than the other parameters of evaluation. Compared with other acoustic parameters, MPT is relatively easy to evaluate regardless of the patient's function or cognition. It can be used in older patients and patients with Parkinsonism. Furthermore, MPT is user-friendly since the patients or the caregivers can evaluate the changes themselves. In addition, its relationship with aspiration, which is the most important factor in dysphagia, was observed in previous studies [12].

**Table 1.** Baseline demographic and clinical characteristics of patients

| Variable             | PD<br>(n=10) | PS<br>(n=20) | Total<br>(n=30) |
|----------------------|--------------|--------------|-----------------|
| Age (yr)             | 72.1±9.4     | 67.4±11.6    | 69.0±11.0       |
| Sex                  |              |              |                 |
| Male                 | 5 (16.7)     | 11 (36.7)    | 16 (53.3)       |
| Female               | 5 (16.7)     | 9 (30.0)     | 14 (46.7)       |
| Ambulation           |              |              |                 |
| Dependent W/C        | 5 (16.7)     | 9 (0.3)      | 14 (46.7)       |
| Independent W/C      | 1 (3.3)      | 3 (10.0)     | 4 (13.3)        |
| Standing alone       | 0 (0)        | 4 (13.3)     | 4 (13.3)        |
| Walking with aid     | 1 (3.3)      | 1 (3.3)      | 2 (6.7)         |
| ID walking (indoor)  | 2 (6.7)      | 1 (3.3)      | 3 (10.0)        |
| ID walking (outdoor) | 1 (3.3)      | 2 (6.7)      | 3 (10.0)        |
| H&Y stage            | 3.8±1.5      | 3.8±1.4      | 3.8±1.4         |
| MBI                  | 26.0±16.4    | 31.7±21.2    | 28.8±20.2       |
| FIM                  | 53.4±18.0    | 53.5±18.9    | 55.7±14.8       |
| NIH-SSS              | 3.2±2.8      | 2.6±2.0      | 2.8±2.3         |
| PAS                  | 4.0±3.2      | 4.5±2.9      | 4.3±2.9         |
| VDS                  | 34.5±24.1    | 28.5±13.7    | 30.5±17.7       |

Values are presented as mean±standard deviation or number (%).

PD, Parkinson's disease; PS, Parkinsonian syndrome; W/C, wheelchair; ID, independent; H&Y, Hoehn and Yahr; MBI, Modified Barthel Index; FIM, Functional Independence Measure; NIH-SSS, National Institutes of Health-Swallowing Safety Scale; PAS, Penetration Aspiration Scale; VDS, Videofluoroscopic Dysphagia Scale.

MPT testing requires deep inhalation and a relaxed voice without stopping. The normal standard for MPT was based on the normal standards for age and sex as recommended by 'Neurologic Speech-Language Disorders' [13].

VFSS was performed by experienced physiatrists in a fluoroscopy/radiography room. The subjects sat laterally in a posture that was appropriate for observation of the anatomical structure and function. The subjects swallowed 5 mL and 15 mL of 12% semisolids followed by 5 mL and 15 mL of liquid, in the stated order, which were all mixed with barium sulfate solution. The test was recorded on video, which was evaluated by two physiatrists. The Penetration Aspiration Scale (PAS), National Institutes of Health-Swallowing Safety Scale (NIH-SSS), and Videofluoroscopic Dysphagia Scale (VDS) scores were evaluated.

### Video fluoroscopic swallowing study analysis

#### National Institutes of Health-Swallowing Safety Scale

The NIH-SSS is a numerical scale that quantifies swallowing safety based on the observations derived from the VFSS, including food residue, laryngeal penetration, aspiration response, maximal esophageal entry, and multiple swallows. Higher scores indicate greater impairment [14].

#### Penetration aspiration scale

The PAS is a numerical scale that quantifies the degree of penetration and aspiration observed during VFSS. This 8-point scale is used to assess the depth of the passage of a bolus into the airway and the patient's response. Higher scores indicate greater impairment [15].

#### Videofluoroscopic dysphagia scale

The VDS for oropharyngeal functional assessment during swallowing is based on the following 14 items: lip closure, bolus formation, mastication, apraxia, tongue-to-palate contact, premature bolus loss, oral transit time, pharyngeal swallow triggering, vallecular residue, laryngeal elevation, pyriform sinus residue, coating of pharyngeal wall, pharyngeal transit time, and aspiration [16]. The scale has a maximum score of 100. Higher scores indicate poor swallowing function.

### Statistical analysis

Data were imported into SPSS version 20.0 (IBM, Armonk, NY, USA) for analysis. Statistical analysis of VDS variables across subgroups was performed using an independent t-test. The differences in MPT between the subgroups, which were classified based on the VDS variables, were compared using the independent t-test. To determine the correlation between dysphagia scales and MPT, Pearson correlation coefficients were calculated; values above 0.6 were considered strong and those ranging between 0.4 and 0.6 were moderate. A p-value <0.05 was considered statistically significant.

## RESULTS

### Baseline demographic and clinical characteristics

VFSS and speech evaluation were performed in 30 patients (10 PD and 20 PS). The 30 patients included 16 males and 14 females. Sixteen patients were ambulatory

(with or without device) and 14 were non-ambulatory. No statistically significant difference was observed in the Hoehn and Yahr (H&Y) stages, Modified Barthel Index, and Functional Independence Measures between patients diagnosed with PD and PS (Table 1).

### Comparison of VDS variables among subgroups

The patients were divided into the following groups: PD or PS, independent or dependent ambulation, and normal or abnormal MPT. These groups were compared to determine differences in their VDS variables. The results showed significant differences in bolus formation ( $0.0\pm 0.0$  vs.  $0.7\pm 1.3$ ;  $p=0.021$ ) and laryngeal elevation ( $0.0\pm 0.0$  vs.  $1.7\pm 3.6$ ;  $p=0.042$ ) between normal and abnormal MPT groups (Table 2).

### Comparison of mean MPT in the VDS subgroups

Among the 30 patients, 22 (73.3%) showed premature bolus loss. Eight (26.7%) had intact oral transit time. Six (20.0%) exhibited intact swallowing reflex. Nineteen (63.3%) had vallecular residue. Twenty-six (86.7%) showed complete laryngeal elevation. Seventeen (56.7%)

carried pyriform sinus residue. Twenty (66.7%) exhibited pharyngeal wall coating. Twenty-seven (90.0%) had an intact pharyngeal transit time. Thirteen (43.3%) showed aspiration.

Comparison of these VDS subgroups revealed significant differences in MPT among those assigned to groups based on bolus formation ( $7.3\pm 4.9$  vs.  $3.3\pm 2.0$ ;  $p=0.009$ ), oral transit time ( $9.5\pm 5.4$  vs.  $5.6\pm 4.2$ ;  $p=0.047$ ), pharyngeal swallow triggering ( $12.4\pm 4.6$  vs.  $5.2\pm 3.6$ ;  $p<0.001$ ), and laryngeal elevation ( $7.3\pm 4.7$  vs.  $2.1\pm 2.4$ ;  $p=0.040$ ) (Fig. 1).

### Correlation of MPT with VDS, PAS, NIH-SSS scores

A correlation analysis was conducted between the MPT and the dysphagia scales. Oral phase of VDS was defined as the sum of lip closure, bolus formation, mastication, apraxia, tongue-to-palate contact, premature bolus loss, and oral transit time. Pharyngeal phase of VDS was defined as the sum of pharyngeal swallowing triggering, vallecular residue, laryngeal elevation, pyriform sinus residue, coating of the pharyngeal wall, and pharyngeal transit time. The MPT was moderately correlated with total score of VDS ( $r=-0.459$ ,  $p=0.011$ ), oral phase score

**Table 2.** Comparison of VDS variables among subgroups

| VDS variable             | PD<br>(n=10) | PS<br>(n=20) | Independent<br>ambulation<br>(n=16) | Dependent<br>ambulation<br>(n=14) | Normal<br>MPT<br>(n=9) | Abnormal<br>MPT<br>(n=21) | Total<br>(n=30) |
|--------------------------|--------------|--------------|-------------------------------------|-----------------------------------|------------------------|---------------------------|-----------------|
| Lip closure              | 0.0±0.0      | 0.0±0.0      | 0.0±0.0                             | 0.0±0.0                           | 0.0±0.0                | 0.0±0.0                   | 0.0±0.0         |
| Bolus formation          | 0.6±1.3      | 0.5±1.1      | 0.4±1.0                             | 0.6±1.3                           | 0.0±0.0                | 0.7±1.3*                  | 0.5±1.1         |
| Mastication              | 0.0±0.0      | 0.0±0.0      | 0.0±0.0                             | 0.0±0.0                           | 0.0±0.0                | 0.0±0.0                   | 0.0±0.0         |
| Apraxia                  | 0.0±0.0      | 0.0±0.0      | 0.0±0.0                             | 0.0±0.0                           | 0.0±0.0                | 0.0±0.0                   | 0.0±0.0         |
| Tongue to palate contact | 0.5±1.6      | 0.0±0.0      | 0.0±0.0                             | 0.4±1.3                           | 0.0±0.0                | 0.2±1.1                   | 0.2±0.9         |
| Premature bolus loss     | 1.5±1.7      | 2.5±1.4      | 2.5±1.8                             | 1.8±1.3                           | 1.5±1.7                | 2.5±1.5                   | 2.2±1.6         |
| Oral transit time        | 2.4±1.3      | 2.1±1.4      | 2.3±1.3                             | 2.1±1.4                           | 1.7±1.6                | 2.4±1.2                   | 2.2±1.3         |
| PST                      | 3.2±2.2      | 3.8±1.6      | 3.7±1.8                             | 3.5±1.9                           | 2.5±2.4                | 4.1±1.4                   | 3.6±1.8         |
| Vallecular residue       | 4.0±2.5      | 2.8±2.8      | 3.4±2.8                             | 3.0±2.7                           | 3.1±3.0                | 3.2±2.6                   | 3.2±2.7         |
| Laryngeal elevation      | 2.7±4.3      | 0.5±2.0      | 1.1±3.1                             | 1.3±3.3                           | 0.0±0.0                | 1.7±3.6*                  | 1.2±3.1         |
| Pyriform sinus residue   | 5.9±4.3      | 3.2±4.2      | 4.2±4.5                             | 3.9±4.3                           | 3.0±4.5                | 4.5±4.3                   | 4.1±4.3         |
| CPW                      | 7.2±3.8      | 5.4±4.5      | 6.2±4.3                             | 5.8±4.5                           | 5.0±4.7                | 6.4±4.2                   | 6.0±4.3         |
| Pharyngeal transit time  | 1.2±2.5      | 0.3±1.3      | 0.8±2.0                             | 0.4±1.6                           | 0.0±0.0                | 0.9±2.2                   | 0.6±1.8         |
| Aspiration               | 5.4±6.0      | 7.5±4.7      | 7.5±5.1                             | 6.0±5.3                           | 5.3±4.7                | 7.4±5.3                   | 6.8±5.2         |
| Total                    | 34.5±24.1    | 28.5±13.7    | 31.9±17.9                           | 28.9±17.9                         | 22.1±14.3              | 34.1±18.0                 | 30.5±17.7       |

Values are presented as mean±standard deviation.

VDS, Videofluoroscopic Dysphagia Scale; PD, Parkinson's disease; PS, Parkinsonian syndrome; MPT, maximal phonation time; PST, pharyngeal swallow triggering; CPW, coating of pharyngeal wall.

\* $p<0.05$ .

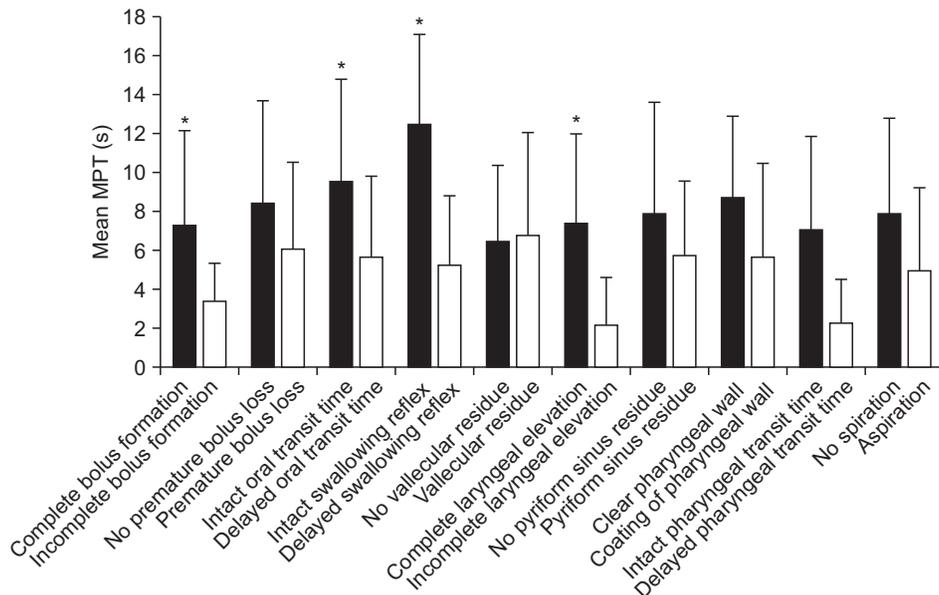


Fig. 1. Comparison of maximal phonation time (MPT) in the Videofluoroscopic Dysphagia Scale subgroups. \*p<0.05 by independent t-test.

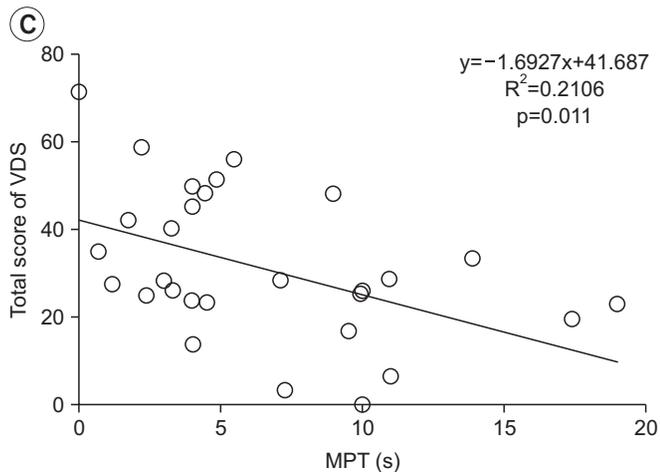
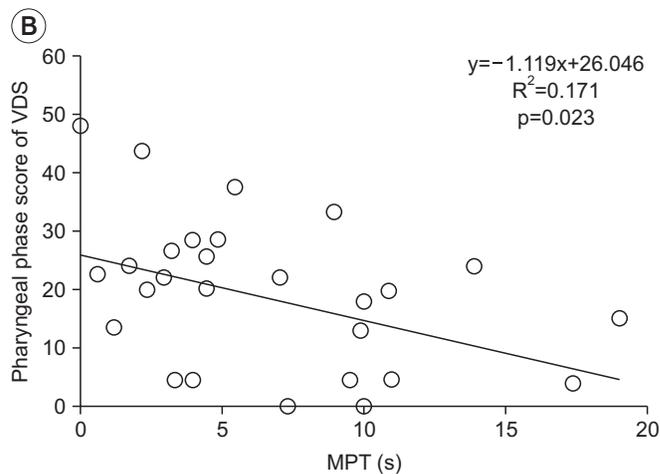
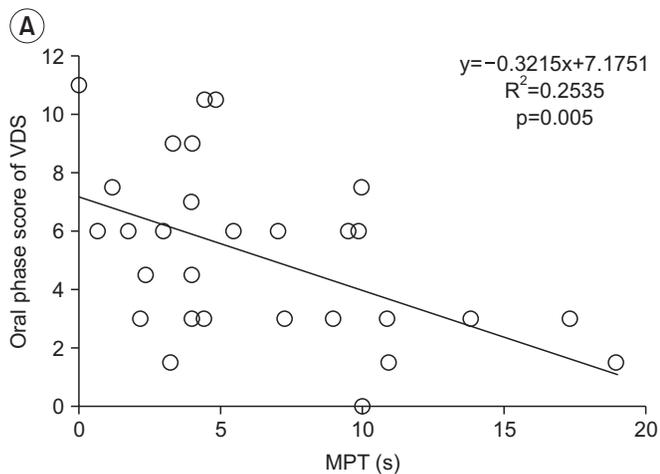
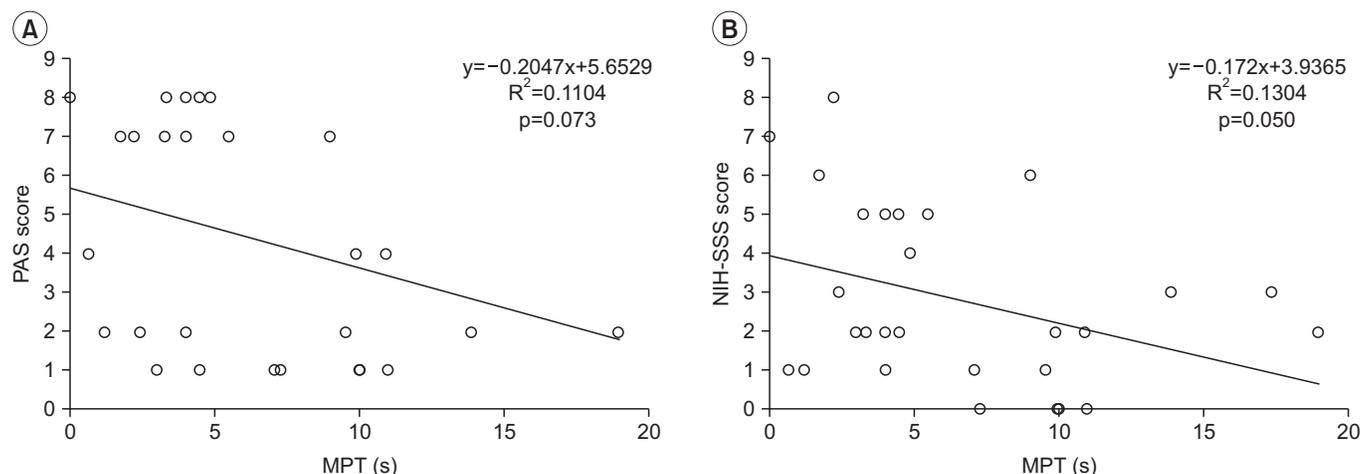


Fig. 2. Correlation of maximal phonation time (MPT) with oral phase score of VDS (A), pharyngeal phase score of VDS (B), and total score of VDS (C). VDS, Videofluoroscopic Dysphagia Scale.



**Fig. 3.** Correlation of maximal phonation time (MPT) with PAS (A) and NIH-SSS scores (B). PAS, Penetration Aspiration Scale; NIH-SSS, National Institutes of Health-Swallowing Safety Scale.

of VDS ( $r = -0.504$ ,  $p = 0.005$ ), and pharyngeal phase score of VDS ( $r = -0.413$ ,  $p = 0.023$ ) (Fig. 2). It also showed a weak correlation with the NIH-SSS ( $r = -0.361$ ,  $p = 0.050$ ), but not the PAS (Fig. 3).

## DISCUSSION

In this study, bolus formation and the laryngeal elevation functions were significantly higher in the normal MPT group compared with the impaired group. When the VDS variable subgroups were also compared, those with intact bolus formation, oral transit time, pharyngeal swallow triggering, and laryngeal elevation showed significantly longer MPTs compared with the impaired groups, indicating an interactive correlation between swallowing and phonation.

In addition, the MPT correlated with oropharyngeal motor function, such as tongue movement (bolus formation, oral transit time), laryngeal elevation, and pharyngeal swallow triggering. This function may be attributed to an increased bolus volume with appropriate tongue movement when swallowing, leading to increased stimulation of sensory receptors that trigger a pharyngeal swallowing reflex [6,17].

VDS and NIH-SSS correlated with MPT in patients with PD and PS, which provides additional evidence that oropharyngeal dysphagia correlates with phonatory dysfunction. There was no correlation between PAS and MPT, a weak correlation between NIH-SSS and MPT, and a moderate correlation between VDS and MPT. The PAS fo-

cuses on the presence of aspiration, whereas the NIH-SSS indicates remnant peristalsis, and other aspects of the pharyngeal phase. The VDS additionally indicates oral phase elements, showing comprehensive oropharyngeal dysfunction in patients suggesting that phonatory function is related to the quality of oropharyngeal swallowing function, but not directly to the aspiration itself.

Swallowing is an organized sensorimotor process within a complex neural network involving automatic and volitional systems [18]. In patients with PD, dopaminergic drugs had no significant effect on non-motor symptoms [4] and other dysphagia treatments have therefore been attempted. Lee Silverman Voice Treatment (LSVT) resulted in improved tongue movement during swallowing, and swallowing efficiency reduced oral residue and oral transit time [17]. Expiratory muscle strength training in patients with PD for 4 weeks also resulted in short-term improvement in PAS scores. The increase in duration and displacement of hyoid movement led to a wider and longer opening of the upper esophageal sphincter, improving the coordination of airway protection and bolus flow through the oropharynx [18,19]. In another study, a significant improvement in the laryngeal elevation time was reported after 8 weeks of group therapeutic singing, which presumably protected the airway from foreign material for a longer period [20].

These studies did not directly target swallowing. However, these types of training can recruit muscles that are involved in swallowing and stimulate similar central and peripheral neural control elements during training,

and are therefore, considered to improve dysphagia [21]. Similarly, our study revealed several swallowing elements related to MPT. Since MPT is improved by training, voice therapy is a potential treatment for dysphagia. The aforementioned studies were typically conducted exclusively in patients with idiopathic PD and with relatively mild H&Y stages. However, patients with PS and those at relatively advanced H&Y stage were also included in this study suggesting that a wider range of patients than previously demonstrated stand to benefit from voice therapy. PS is generally less responsive or not responsive to dopaminergic drug therapies than PD. Therefore, alternative treatment methods are more important.

In previous studies, MPT was used to indirectly evaluate laryngeal function and vocal fold vibration efficiencies. Instead of solely targeting MPT for training, MPT was included as a part of voice therapy, similar to LSVT, or indirect training using an expiratory muscle strength training program [10,11,17]. Since this study indicates that an increase in MPT leads to an improvement of swallowing symptoms, training that is designed and conducted to target MPT will directly confirm the treatment effects.

In addition to MPT, diverse acoustic parameters such as intensity, jitter, shimmer, harmonics-to-noise ratio, voice onset time, sequential motion rate, and alternating motion rate can be measured to evaluate the voice. In this study, MPT was based on the subject's characteristics, ease of evaluation and parameter representation. In future studies, the relationship between different parameters and swallowing function needs to be investigated in a large group.

Overall, there is a need for large-scale, well-designed, randomized, and controlled studies in order to correlate MPT with swallowing-related elements, as well as determine the short- and long-term effects of voice therapy on dysphagia, by comparing the results before and after treatment.

## CONFLICT OF INTEREST

No potential conflict of interest relevant to this article was reported.

## ACKNOWLEDGMENTS

This research was supported by grants from the Min-

istry of Health and Welfare, Republic of Korea (No. HI16C1012 and No. HI16C1176).

## REFERENCES

1. Plowman-Prine EK, Sapienza CM, Okun MS, Pollock SL, Jacobson C, Wu SS, et al. The relationship between quality of life and swallowing in Parkinson's disease. *Mov Disord* 2009;24:1352-8.
2. Kalia LV, Lang AE. Parkinson's disease. *Lancet* 2015; 386:896-912.
3. Potulska A, Friedman A, Krolicki L, Spychala A. Swallowing disorders in Parkinson's disease. *Parkinsonism Relat Disord* 2003;9:349-53.
4. van Hooren MR, Baijens LW, Voskuilen S, Oosterloo M, Kremer B. Treatment effects for dysphagia in Parkinson's disease: a systematic review. *Parkinsonism Relat Disord* 2014;20:800-7.
5. Suttrup I, Warnecke T. Dysphagia in Parkinson's disease. *Dysphagia* 2016;31:24-32.
6. Regan J, Walshe M, Tobin WO. Immediate effects of thermal-tactile stimulation on timing of swallow in idiopathic Parkinson's disease. *Dysphagia* 2010;25:207-15.
7. van Hooren MR, Baijens LW, Vos R, Pilz W, Kuijpers LM, Kremer B, et al. Voice- and swallow-related quality of life in idiopathic Parkinson's disease. *Laryngoscope* 2016;126:408-14.
8. Speyer R, Baijens L, Heijnen M, Zwijnenberg I. Effects of therapy in oropharyngeal dysphagia by speech and language therapists: a systematic review. *Dysphagia* 2010;25:40-65.
9. Stegemoller EL, Radig H, Hibbing P, Wingate J, Sapienza C. Effects of singing on voice, respiratory control and quality of life in persons with Parkinson's disease. *Disabil Rehabil* 2017;39:594-600.
10. Maslan J, Leng X, Rees C, Blalock D, Butler SG. Maximum phonation time in healthy older adults. *J Voice* 2011;25:709-13.
11. Speyer R. Effects of voice therapy: a systematic review. *J Voice* 2008;22:565-80.
12. Valim MA, Santos RS, Macedo Filho ED, da Silva Abdulmassih EM, Serrato MR. The relationship between the maximum time for phonation, fundamental frequency and protection of the lower airways in patients with neurological dysphagia. *Int Arch Otorhinolaryn-*

- gol 2007;11:260-6.
13. Kim H. Neurologic speech-language disorders. Seoul: Sigmaph; 2012.
  14. Belafsky PC, Kuhn MA. The clinician's guide to swallowing fluoroscopy. New York: Springer; 2014. p. 59-68.
  15. Rosenbek JC, Robbins JA, Roecker EB, Coyle JL, Wood JL. A penetration-aspiration scale. *Dysphagia* 1996; 11:93-8.
  16. Han TR, Paik NJ, Park JW, Kwon BS. The prediction of persistent dysphagia beyond six months after stroke. *Dysphagia* 2008;23:59-64.
  17. El Sharkawi A, Ramig L, Logemann JA, Pauloski BR, Rademaker AW, Smith CH, et al. Swallowing and voice effects of Lee Silverman Voice Treatment (LSVT): a pilot study. *J Neurol Neurosurg Psychiatry* 2002;72:31-6.
  18. Troche MS, Okun MS, Rosenbek JC, Musson N, Fernandez HH, Rodriguez R, et al. Aspiration and swallowing in Parkinson disease and rehabilitation with EMST: a randomized trial. *Neurology* 2010;75:1912-9.
  19. Pitts T, Bolser D, Rosenbek J, Troche M, Okun MS, Sapienza C. Impact of expiratory muscle strength training on voluntary cough and swallow function in Parkinson disease. *Chest* 2009;135:1301-8.
  20. Stegemoller EL, Hibbing P, Radig H, Wingate J. Therapeutic singing as an early intervention for swallowing in persons with Parkinson's disease. *Complement Ther Med* 2017;31:127-33.
  21. Russell JA, Ciucci MR, Connor NP, Schallert T. Targeted exercise therapy for voice and swallow in persons with Parkinson's disease. *Brain Res* 2010;1341:3-11.