Oropharyngeal swallow physiology and swallowing-related quality of life in underweight patients with concomitant advanced chronic obstructive pulmonary disease

This article was published in the following Dove Press journal:
International Journal of COPD

Kendrea L Garand1
Charlie Strange2
Luca Paoletti2
Theresa Hopkins-Rossabi3
Bonnie Martin-Harris3

1Department of Speech Pathology and Audiology, College of Allied Health Professions, University of South Alabama, Mobile, AL 36688, USA; 2Division of Pulmonary, Critical Care, and Sleep Medicine, College of Medicine, Medical University of South Carolina, Charleston, SC 29425, USA; 3Department of Communication Sciences and Disorders, School of Communication, Northwestern University, Evanston, IL 60208, USA

Background: Swallowing impairment (dysphagia) has been associated with COPD and may contribute to exacerbations of this chronic and progressive disease. Further, risk of mortality increases with concomitant presence of cachexia in the COPD population. The purpose of this prospective study was to depict oropharyngeal swallowing physiology in underweight patients with stable but advanced-stage COPD.

Patients and methods: Ten underweight patients with stable but advanced COPD underwent a modified barium swallow study. Analysis of oropharyngeal swallowing function was completed using the standardized Modified Barium Swallow Impairment Profile and the Penetration–Aspiration Scale. Scores from the Dysphagia Handicap Index and 10-item Eating Assessment Tool were collected to assess patient perception of swallowing difficulty. Findings were compared to age- and sex-matched healthy controls.

Results: Significantly higher MBSImP oral total scores \((P=0.007)\) were observed in COPD patients compared to matched controls, but no difference was observed in pharyngeal total scores \((P=0.105)\). Patients with COPD had significantly higher maximum PAS scores compared with controls \((P=0.030)\). There was no significant difference in EAT-10 or DHI scores between patients with COPD and controls \((P=0.41\) and \(P=0.08\), respectively).

Conclusion: Underweight patients with severe but stable COPD present with dysphagia that may not be recognized by the patient. Further investigation is needed to elucidate the interaction between the respiratory–swallowing systems, how muscular weakness may contribute to swallowing impairment, and responsiveness to swallowing treatment.

Keywords: COPD, swallowing impairment, MBSImP, dysphagia, underweight

Introduction
A chronic inflammatory lung disease, COPD is the third leading cause of death in the United States.1 Exacerbations of symptoms are typical and can be severe during disease progression, whereas contributing to repeated, costly hospitalizations and early mortality.2 The estimated medical cost of COPD in 2012 was over 32 billion dollars3 and is expected to rise, given frequent hospital readmissions within 30 days after initial discharge.4 COPD disease severity and clinical manifestations vary greatly between patients and research cohorts that have studied the incidence and type of associated oropharyngeal swallowing impairment. Further, assessment methods and metrics to investigate oropharyngeal swallowing mechanism and dysfunction have varied. One study
reported the incidence of dysphagia associated with COPD was 85%, with laryngeal penetration and tracheal aspiration observed in 28% and 42% of subjects, respectively.6 Further, complications secondary to swallowing impairments, such as aspiration pneumonia, dehydration, and airway obstruction, have been shown to contribute to disease exacerbations.6 The negative effects of exacerbation on patient’s quality of life, physical and physiological well-being, and early mortality2 highlight the potential impact that dysphagia may have on the natural course of COPD.

Ghannouchi et al completed a systematic literature review revealing differences between COPD and healthy age-matched controls that were observed on videofluoroscopic imaging.7 For example, prolonged airway closure and a lower laryngeal position were observed in COPD patients with moderate to very severe disease.8 It was postulated that the prolonged airway closure represents an underlying protective response to reduce the risk of airway invasion.9 Other observations include prolonged transit times for solid foods, longer laryngeal vestibule closure times during liquid swallows,9 delayed swallow initiation,4 and oropharyngeal residue.10 Additionally, the rate of tracheal aspiration of thin liquids observed on videofluoroscopy in COPD patients was higher, compared to age-matched controls (25% vs 0.1%, respectively).11 Robinson et al found a significantly higher risk of aspiration on bedside swallow evaluations in a COPD group (44%) compared to a non-COPD hospitalized group (29%).12 Although the non-COPD patients’ videofluoroscopic results were not reported, 17% of the COPD patients aspirated on videofluoroscopy.12 An additional study using fiberoptic endoscopic evaluation of swallowing revealed that the majority of patients with COPD presented with pharyngeal residue (90%), and 25% of the study sample had bolus airway invasion.10

In addition to oropharyngeal motor impairment, altered upper airway sensory mechanics in COPD may further contribute to penetration/aspiration. Evidence suggests that COPD patients have reduced laryngopharyngeal sensitivity, resulting in decreased measures to clear pharyngeal residue.13 Moreover, the COPD patient may be less likely to engage in productive coughing to expel ingested material from the airway.

Furthermore, dysphagia is known to have devastating effects on nutritional status, with approximately 25% of patients with COPD developing cachexia – a wasting syndrome associated with a 50% reduction in median survival.14 To date, the authors are not aware of any study that investigates the concomitant effects of low body mass index (BMI) on swallowing in patients with COPD.

The purpose of the present prospective cohort study was to describe oropharyngeal swallowing physiology in patients with advanced-stage COPD (Global Initiative for Chronic Obstructive Lung Disease [GOLD] stages 3 and 4) with concomitant low BMI using standardized, validated, and reproducible metrics of swallowing impairment and bolus airway invasion. This is the only investigation known to the authors that attempts to define dysphagia in a population of COPD patients who were also underweight. Prior research has reported on BMI but not limited inclusion to those patients presenting with low BMI.9,16 Further, we detailed oral intake, patient-reported symptoms of swallowing impairment, and the impact of dysphagia on quality of life. These findings were compared to age- and sex-matched healthy controls retrieved from a large normative database.

Patients and methods

Each patient underwent a baseline modified barium swallow study (MBSS) as part of a clinical trial evaluating the effects of an investigational nutritive agent to treat low BMI in 12 patients with advanced but stable COPD. Key eligibility criteria included: 1) ages 30–80 years; 2) GOLD Stage 3 or Stage 4; and 3) low weight, defined as BMI below 22 kg/m². Exclusion criteria were: 1) current tobacco, alcohol, or illicit drug use; 2) unstable cardiovascular disease; 3) cognitive dysfunction; 4) non-COPD lung disease; 5) history of organ transplant; 6) history of lung cancer or any metastatic cancer; 7) HIV/AIDS; 8) taking an immunosuppressive agent; and 9) taking greater than 20 mg of prednisone or equivalent corticosteroid.

Swallowing-related data from healthy controls were extracted from a large normative database as part of a separate study investigating effects of aging on swallowing function. A healthy participant was defined as an individual without current or history of dysphagia, upper aerodigestive tract surgical procedures, hiatal hernia, pulmonary disease (eg, COPD), head and neck cancer, and/or neurological disease (eg, stroke, Parkinson’s disease). Further, he/she was able to drink thin liquids and eat solids as part of a regular diet and did not have a known allergy or dietary restriction for food or barium contrast materials used in study procedures. For each COPD patient, a maximum of four age-matched (±2 years) and sex-matched controls were extracted.

All informed consent procedures were in accordance with the ethical standards of the responsible committee on human experimentation. Written informed consent was obtained from all study patients and healthy controls. This study received Institutional Review Board Approval from
Medical University of South Carolina (IRBs# Pro0027289 [COPD] and Pro00011566 [healthy controls]). This study was registered as a clinical trial with the ClinicalTrials.gov Identifier of NCT02067377.

Videofluoroscopic imaging was obtained using continuous fluoroscopy, and high-resolution images were recorded at 30 frames (Digital Swallowing Workstation Model 7100, Kay Elemetrics Corp.) Standardized consistencies of commercially prepared barium (Varibar®) of increasing volume (5 mL, single-cup swallow, and sequential swallows of liquid) and viscosity (thin, nectar- and honey-thickened liquids, pudding, and cookie) were provided for a maximum of 12 swallowing tasks in two viewing planes (ten swallows lateral and two swallows anteroposterior) following the standardized Modified Barium Swallow Impairment Profile©™ (MBSImP) approach. The MBSImP measures 17 validated and reliable components of swallowing physiology across three functional domains: oral, pharyngeal, and esophageal.

The presence and degree of airway invasion were measured using the validated Penetration–Aspiration Scale (PAS) – an eight-point ordinal scale. A score of 1 represents the absence of airway invasion, and a score of 8 represents an absent attempt to react to or expectorate aspirated material (ie, silent aspiration). Scores of 3 and greater are considered impaired, on the basis of normative data. PAS was derived for each swallowing task.

The Functional Oral Intake Scale (FOIS) was used to document the patients’ oral intake level. The FOIS consists of seven levels to assess diet level of safe oral intake, with Level 1 representing no oral intake and Level 7 representing total oral intake without restrictions. Additionally, each patient completed the Dysphagia Handicap Index (DHI)2 and 10-item Eating Assessment Tool (EAT-10). The DHI measures the handicapping effects of dysphagia on emotional, functional, and physical aspects of the lives of dysphagic patients. Each of the 25 statements distributed across the three domains (emotional, functional, and physical) are rated as “never”, “sometimes”, or “always” occurring. Higher scores indicate more severe impairment. The DHI was validated in a heterogeneous dysphagic patient sample (N=77), using a control of non-dysphagic patients for clinical validity. The EAT-10 measures symptom-specific outcomes of dysphagia using ten items rated on a Likert scale of 0 (no problem) to 4 (severe problem), which was validated using a large cohort of voice and swallowing patients. The EAT-10 was validated using a large cohort of voice and swallowing patients (N=235), with a score of score of 3 and greater indicating abnormal swallowing function based on published normative data.

COPD assessments included spirometry before and after bronchodilator, plethysmographic lung volumes to record the residual volume/total lung capacity (RV/TLC) ratio. A 6MWD to define the distance walked by standardized protocol. The St Georges Respiratory Questionnaire (SGRQ) was administered to define COPD-specific quality of life by total score. The SGRQ has been used since 1991 and has been validated for correlations between symptoms and activity levels.

Data collection

MBSImP component scores of 1 and greater were considered impaired for purposes of analysis. However, lip closure, tongue base retraction, and pharyngeal residue were deemed impaired if the score was greater than 2 (scores of 1 on these components could be considered normal variance). Descriptive measures were calculated for MBSImP component and composite scores (oral and pharyngeal totals), PAS, DHI, and EAT-10. Mann–Whitney U tests were calculated to compare composite scores between patients with COPD to sex- and age-matched (±2 years) healthy, non-dysphagic controls that were collected as part of a large normative study (n=195) separate from the current project. After all the eligible control matches were selected, a maximum of four were randomly chosen for a total of 37 control subjects because one COPD subject only had one matching healthy control. Regression analysis was conducted between measures of OT and PT scores and the following COPD physiologic...
Results

Results of ten patients (five females) with a mean age of 62 (±8) years are presented. Two patients were removed from analysis because of previous anterior cervical surgeries which may impact swallowing function. All patients had obstructive lung disease, with most classified as Stage 4 (90%). Thirty-seven adults (20 females) comprised the control group, with a mean age of 61 (±8) years. Further demographic and clinical characteristics are described in Table 1.

MBSImP component and composite scores

All patients demonstrated impairment in oral residue, initiation of pharyngeal swallow, tongue base retraction, pharyngeal residue, and esophageal clearance. Moreover, the majority of patients demonstrated impairment in bolus preparation/mastication and pharyngoesophageal segment opening (70% and 90%, respectively; Table 2).

The mean (±SD) OT and PT scores were 7.7 (±2.6) and 7.9 (±3.0), respectively (Table 3). When composite scores of the patients with COPD were compared with matched controls, there was a significant difference in OT scores ($P=0.007$) but not PT scores ($P=0.105$; Table 3).

Table 1 Patient demographics. Data presented as mean ± SD unless otherwise reported

<table>
<thead>
<tr>
<th>Variables</th>
<th>COPD (N=10)</th>
<th>Healthy controls* (N=37)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>61.8±8.2</td>
<td>61.0±7.8</td>
</tr>
<tr>
<td></td>
<td>Range</td>
<td>53–76</td>
</tr>
<tr>
<td>Sex frequency (female %)</td>
<td>5 (50.0%)</td>
<td>20 (54.1%)</td>
</tr>
<tr>
<td>Racial category frequency (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>White/Caucasian</td>
<td>9 (90%)</td>
<td>32 (86.5%)</td>
</tr>
<tr>
<td>Black/African American</td>
<td>1 (10%)</td>
<td>4 (10.8%)</td>
</tr>
<tr>
<td>Ethnicity category frequency (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Non-Hispanic/Non-Latino</td>
<td>10 (100%)</td>
<td>36 (97.3%)</td>
</tr>
<tr>
<td>BMI</td>
<td>18.5±1.4</td>
<td>–</td>
</tr>
<tr>
<td></td>
<td>Range</td>
<td>16.6–20.5</td>
</tr>
<tr>
<td>Pulmonary function</td>
<td></td>
<td></td>
</tr>
<tr>
<td>FEV₁ % predicted</td>
<td>27.9±7.8</td>
<td>–</td>
</tr>
<tr>
<td>FVC % predicted</td>
<td>70.0±10.9</td>
<td>–</td>
</tr>
<tr>
<td>FEV₁/FVC</td>
<td>30.6±6.8</td>
<td>–</td>
</tr>
<tr>
<td>TLC % predicted</td>
<td>125.1±31.9</td>
<td>–</td>
</tr>
<tr>
<td>RV % predicted</td>
<td>202.3±76.1</td>
<td>–</td>
</tr>
<tr>
<td>RV/TLC</td>
<td>58.4±13.4</td>
<td>–</td>
</tr>
<tr>
<td>Hand grip strength</td>
<td>59.6±20.3</td>
<td>–</td>
</tr>
</tbody>
</table>

Note: Healthy controls did not self-report a diagnosis of COPD, pulmonary, or neurologic disease, or history of head/neck or cardiovascular cancer or surgery. Abbreviations: BMI, body mass index; FEV₁, forced expiratory volume in the first second; FVC, forced vital capacity; TLC, total lung capacity; RV, residual volume.

PAS

The mean PAS score (±SD) for all COPD patients across swallowing tasks was 1.3 (±0.9), with a range of 1–8 (Tables 3 and 4). Highest PAS scores were observed on the sequential thin liquid (worst score of 4) and sequential nectar-thickened liquid (worst score of 8) swallowing tasks. Patients with COPD had significantly higher maximum (worst) PAS scores compared with controls ($P=0.030$) (Tables 3 and 4). One COPD patient was once observed to have silent aspiration (PAS=8) during the sequential nectar-thickened liquid swallowing task only. Reduced laryngeal elevation and laryngeal vestibular closure contributed to the bolus entering the airway without clearance, which resulted in the residue falling below the true vocal folds on a subsequent swallow without patient attempt to expectorate the aspirated material.

FOIS

All patients had a FOIS level of 7, which indicates they reported a total oral diet of all consistencies without any restrictions.

EAT-10

The EAT-10 mean (±SD) total score was 2.5 (±3.2), with a range from 0 to 8 (Table 3). Forty-four percent of patients with COPD had a total score of 3 or greater as compared with 19% of healthy controls (n=7). There was no significant difference in EAT-10 total scores between patients with COPD and controls ($P=0.413$).

DHI

The mean (±SD) total DHI score was 12.2 (±11.1), with a range from 2 to 36 (Table 3). Summary statistics across the physical, functional, and emotional domains are provided in Table 3. Comparisons between patients with COPD and controls (n=5) did not reveal a significant difference in DHI scores ($P=0.075$).

COPD metrics

Correlations between the OT score trended with the RV/TLC ratio – a measure of hyperinflation and 6MWD distance – an important functional outcome in COPD. Further, the RV/TLC trended with the component score bolus preparation/mastication – a critical component of efficient bolus preparation and airway protection. PT scores trended with the SGRQ to suggest that both oral and pharyngeal components of swallowing are dysfunctional in severe COPD patients who perceive they have no swallowing problems (Figure 1).
The purpose of this cohort study was to describe oropharyngeal swallowing physiology in patients with stable, advanced-stage COPD and were underweight (BMI <22). Oropharyngeal swallowing impairment was observed in all COPD patients. Oral impairment and PAS scores were significantly worse from those of age- and sex-matched healthy controls. Swallowing impairment, despite the minimal airway invasion observed in this cohort, contributes to increased aspiration and nutritional risks. The patients included in this sample were stable; however, during an acute exacerbation, these patients may not be able to protect their airway as efficiently. Further, if there is a progression of dysphagia severity, it may contribute to COPD severity and exacerbations. Moreover, increasing dysphagia severity may further exacerbate nutritional impairments in patients who are already underweight (low BMI), with detrimental effects on the body, including further loss of muscle mass and gastrointestinal dysfunction. The latter is of interest because impairment in esophageal clearance in the upright position was observed in all COPD patients but only half of the healthy control subjects.

Interestingly, patients with COPD have no specific neurological or structural etiology known to account for dysphagia. However, all study patients exhibited delayed initiation of the pharyngeal swallow and oropharyngeal residue, which may indicate muscular weakness and/or reduced sensation of upper aerodigestive tract structures involved in swallowing – specifically, the tongue and pharynx. For example, the majority of patients exhibited decreased bolus preparation, and all patients demonstrated reduced tongue base retraction, suggesting lingual weakness that further supports findings from previous studies. Generalized weakness present in the COPD population has been reported to be related, at least in part, to poor nutrition and reduced exercise tolerance. Further, specific sarcopenia (which affects 15% of patients with stable COPD) may occur in diffuse muscle groups and can affect muscles used during swallowing. In addition, discoordination of respiration with impaired swallowing has been implicated in patients with stable and acute exacerbations of COPD.
pulmonary function and impaired swallowing increase aspiration risk in patients with COPD.

Another contributor of dysphagia may be the reduction in oropharyngeal sensation. Similar to results by Good-Fratturelli et al who found delayed initiation of the pharyngeal swallow in 60% of patients, we found delayed swallowing in all patients in this COPD cohort. Notably, the pharyngeal swallow was not triggered until the leading

<table>
<thead>
<tr>
<th>Swallowing task</th>
<th>PAS=1 or 2</th>
<th>PAS=3–5</th>
<th>PAS=6–8</th>
</tr>
</thead>
<tbody>
<tr>
<td>5 mL thin 2</td>
<td>10 (100%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Cup thin</td>
<td>10 (100%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Sequential thin</td>
<td>8 (80%)</td>
<td>2 (20%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>5 mL nectar</td>
<td>10 (100%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Cup nectar</td>
<td>10 (100%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Sequential nectar</td>
<td>9 (90%)</td>
<td>0 (0%)</td>
<td>1 (10%)</td>
</tr>
<tr>
<td>Honey</td>
<td>10 (100%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Pudding</td>
<td>10 (100%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
</tr>
<tr>
<td>Cookie</td>
<td>10 (100%)</td>
<td>0 (0%)</td>
<td>0 (0%)</td>
</tr>
</tbody>
</table>

Notes: *Data from Robbins.° PAS scores of 1 or 2 are frequently observed in healthy adults and, thus, are considered “normal”.
Abbreviation: PAS, Penetration–Aspiration Scale.

Figure 1 Analysis of COPD metrics oral total scores on the MBSImP are associated with measures of hyperinflation.
Notes: Oral total scores are associated with measures of hyperinflation (A) and reflected in bolus preparation scores (B). Both oral and pharyngeal total scores are associated with measures of COPD severity including 6MWD (C) and SGRQ (D).
Abbreviations: RV/TLC, residual volume/total lung capacity; SGRQ, St Georges Respiratory Questionnaire; MBSImP, Modified Barium Swallow Impairment Profile.
edge of the bolus was well into the pharynx in 60% of our patients on larger sequential swallowing tasks. Patients with COPD had consistently higher impairment rates for initiation of pharyngeal swallow as compared to matched controls, although differences on comparison were found to be non-significant. Two possible explanations for this result include the small sample size and the normal variation on this swallowing component observed in aging adults. For example, it has been shown that healthy older adults may initiate pharyngeal swallowing later when the leading edge of the bolus is advanced into the airway without aspiration occurrence.\(^{31}\) Interestingly, however, delayed swallow initiation was also observed in 88% of patients studied with recent exacerbation of COPD by Kobayashi et al.\(^{32}\) It may be that swallow delay may be confounded by other impaired airway protection mechanisms, such as decreased sensation, in patients with COPD. For example, Clayton et al reported decreased laryngeal sensation in patients with COPD, together with abnormal pharyngeal residue; the latter was observed in our cohort.\(^{33}\) The high incidence of oropharyngeal residue and delayed initiation of the swallow may in part result from overall reduced sensation. It has been proposed that the impact of corticosteroids, current or past smoking behavior, and oxygen therapy may impact pharyngeal and laryngeal mucosal sensory receptors of the pharyngeal and laryngeal mucosa in this population, and could then lead to decreased sensation.\(^{13}\)

Impairment in laryngeal elevation was observed in almost half of the studied patients. Furthermore, those patients with decreased laryngeal elevation frequently presented with incomplete laryngeal vestibule closure, which further places patients at risk for airway invasion at the height of the swallow (when the vestibule should be fully closed and shielded from the bolus flowing through the pharynx). These findings support the investigation by Mokhlesi et al who also observed a lowered laryngeal position at this point in the swallow in patients with stable COPD.\(^{8}\)

A higher incidence of laryngeal penetration and tracheal aspiration in COPD patients has been reported in previous studies, with the suggestion that ingested material in the airway may increase exacerbations in COPD.\(^{6}\) Good-Fratturelli et al reported 28% and 42% penetration and aspiration rates, respectively,\(^{5}\) similar to other studies.\(^{10-12}\) Although only one patient had one instance of silent aspiration, outside of the clinical context of the MBSS, this physiologic impairment during daily eating and drinking may well result in trace, low-grade aspiration and, therefore, confound pulmonary status. The patients’ perception of swallow function was incongruent with the nature and severity of the swallowing impairment observed. Their self-reported symptomatology related to swallowing impairment (EAT-10) and subsequent impact on quality of life (DHI) was minimal, with no difference observed between patient scores and matched healthy controls. Further, the patients in this cohort indicated no dietary restrictions (FOIS Level 7). Reduced pharyngeal sensation may account, in part, for the patient’s limited insight into swallowing impairment. Further, reduced laryngopharyngeal sensation places the patient at increased risk for aspiration and aspiration-related pulmonary complications. The mismatch between patient’s perceived and actual swallowing impairment and the nature of the swallowing impairments may signal a need for swallowing assessment and subsequent management in this patient population.

**Study limitations**

The primary limitation of this study includes the small patient sample representing a subset of COPD patients. The original sample size included 12 patients, but two were excluded secondary to history of anterior cervical disectomy fusion (ACDF) spinal surgeries because of the known swallowing impairment that may be result from disruption to the sensory end organs and muscles of the pharynx. Because this analysis was specifically investigating effects of COPD and underweight on swallowing function, we did not want potential confounders influencing results. Further, an MBSS represents only a few minutes in time. Fatigue is known to greatly impact patients with COPD when completing basic activities of daily living – including eating and drinking.\(^{33}\) It follows that swallowing function could likely further deteriorate outside the context of a clinical assessment and over the course of a full meal, leading to increased risk of aspiration. Next, certain measures (ie, BMI, hand grip strength, and spirometry) were not available in the normative dataset and, thus, we were not able to further analyze comparisons between the patient group and their matched controls. Lastly, we did not have data available for advanced-stage but stable COPD patients with normal weight for comparison. This would have strengthened clinical implications for the contribution of low BMI on swallowing function.

**Conclusion**

The results of this study suggest that patients with severe but stable COPD with concomitant underweight and dysphagia may require a multisystem approach for defining and
rehabilitating dysphagia. Although the evidence from this current study revealed the presence of significant swallow impairment in underweight patients with severe COPD, there remains the lack of a clear cause and effect. This investigation included patients with a BMI below 22 kg/m². Future research should include patients with COPD who present with both higher and lower BMI to determine if factors such as overall body mass, rather than COPD status, impacts swallow function.

Swallowing impairment in COPD is likely a multifactorial issue and influenced by muscular weakness, alteration in oropharyngeal and laryngeal sensation, and mismatch between patient perception and actual presence and severity of dysphagia. Moreover, patients with COPD have been shown to have disruption in the coordinative processes between respiration and swallowing. Discoordination between the respiratory cessation and swallowing initiation has been shown to contribute to dysphagia and aspiration. Further investigations that directly examine aberrations in respiratory swallowing coordination and oropharyngeal muscle function on the type and severity of dysphagia are warranted. Such studies will lead to the discovery of potential behavioral interventions that target the impaired mechanisms contributing to disruptions in swallowing safety and efficiency in patients with COPD.

Acknowledgments

The authors would like to acknowledge Julie Blair, MA, CCC-SLP, BCS-S, Kate Davidson, MS, CCC-SLP, and R Jordan Hazelwood, PhD, CCC-SLP, BCS-S, CBIS, for their assistance with data collection and analysis of videofluoroscopic examinations of oropharyngeal swallowing in this patient sample.

This work was partially supported by Entera Health, Inc. (grant no. EH5220; principal investigator [PI]: Paoletti), Veterans Affairs Career Development Award-1 (VA CDA-1) R&D (grant no. 1IK1RX001628-01A1; PI: Garand), the National Institute on Deafness and Other Communication Disorders of the National Institutes of Health (grant no. K24DC12801; PI: Martin-Harris), and the South Carolina Clinical & Translational Research (SCTR) Institute, with an academic home at the Medical University of South Carolina, NIH/NCATS (grant no. TL1 TR000061; PI: Brady; Project PI: Garand), and the American Speech-Language-Hearing Foundation (PI: Garand).

This paper was presented as a scientific paper presentation at the 24th Annual Meeting of the Dysphagia Research Society, February 25–27, 2016, Tucson, AZ, USA. Link: https://c.vymcdn.com/sites/dysphagiaresearch.sitrym.com/resource/resm//Meeting_Files/DRS_2016_Annual_Mtg_and_Post.pdf

The paper’s abstract was published in Dysphagia. 2016;Vol 31(6):803–804.

This paper was also presented as a Technical research presentation at the 2016 American Speech-Language-Hearing Association (ASHA) Annual Convention, November 17–19, 2016; Philadelphia, PA, USA. (ASHA is not an openly accessible published abstract).

Disclosure

CS has received grants and non-financial support from the Alpha-1 Foundation, CSL Behring, and Pulmonx; grants, personal fees, and non-financial support from Grifols; grants from Inhibrix, Novartis, Pulmonx, and Shire; and non-financial support from Abeeona and personal fees and non-financial support from AstraZeneca, all outside the submitted work. BMH receives copyright royalties from Northern Speech Services and Medical University of South Carolina and is the principal investigator (NIH/NIDCD, 1K24DC12801) in a research study and mentoring of swallowing impairment and respiratory–swallow coordination (2013–2018). KLG, THR, and LP report no conflicts of interest in this work.

References