Pathophysiology and treatment of patients with globus sensation —from the viewpoint of esophageal motility dysfunction—

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Abstract

“Globus sensation” is often described as the sensation of a lump in the throat associated with dry swallowing or the need for dry swallowing, which disappears completely during eating or drinking and for which no organic cause can be established. Due to the uncertain etiology of “globus sensation”, it remains difficult to establish standard treatment strategies for affected patients. Lately most attention has been focused on gastroesophageal reflux disease and several reports have indicated that there is a close relationship between esophageal acid reflux and globus sensation. Nowadays, empirical therapy with a high dose of a proton pump inhibitor (PPI) is considered to be indicated for patients with globus sensation, after excluding organic diseases such as pharyngeal cancer, Zenker’s diverticulum, or thyroid enlargement. If patients are nonresponsive to PPI therapy, evaluation of esophageal motility should be done. In our recent study, 47.9% had abnormal esophageal motility, with the most common esophageal motility abnormality being an ineffective esophageal motility in PPI-resistant patients with globus sensation. This suggests that prokinetics alone or adding prokinetics to PPI should be the treatment to be considered, although few studies have investigated the efficacy of prokinetics in the treatment of patients with globus sensation. If patients without any esophageal motility dysfunctions are nonresponsive to PPI therapy, either cognitive-behavioral therapy, anti-depressants, or gabapentin could be helpful, although further well-designed, randomized controlled large-scale studies will be necessary to determine the effectiveness of each treatment strategy on patients with globus sensation.

Key words: globus sensation, gastroesophageal reflux disease, proton pump inhibitor, esophageal manometry

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Abbreviations used: GERD, Gastroesophageal reflux disease; GI, gastrointestinal; PPI, proton pump inhibitor; UES, upper esophageal sphincter.

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Introduction

It is approximately 2500 years since Hippocrates first noted globus pharyngeus (1). In 1707, Purcell (2) was the first to accurately describe the condition; he believed that globus resulted from pressure on the thyroid cartilage due to contraction of the strap muscles of the neck. In the past, globus was described as “globus hystericus” because of its frequent association with menopause or psychogenic factors. However, Malcomson (3) coined the more accurate term “globus pharyngeus” in 1968 after discovering that most patients experiencing globus did not have a hysterical personality.

“Globus sensation” is often described as the sensation of a lump in the throat associated with dry swallowing or the need for dry swallowing, which disappears completely during eating or drinking and for which no organic cause can be established. It is a common condition that accounts for approximately 4% of new referrals to ear, nose and throat clinics, and it is reported by up to 46% of apparently healthy individuals, with a peak incidence in middle age (4, 5). Although the prevalence of patients with globus sensation was high, the natural history of patients with globus sensation has not been fully elucidated. In the past, Rowley et al. showed that during the mean follow-up period of 7 years and 7 months, 55% of patients with globus sensation were asymptomatic and 45% of patients had persistent symptoms, while no patient developed upper gastrointestinal malignancy (6). Despite the benign nature of the condition, the likelihood of long-term symptom persistence often leads to a reduction in their quality of life, due to the lack of a highly effective single treatment, attributable to the fact that its pathogenesis has been poorly understood.

Gastroesophageal reflux disease (GERD) is a condition that develops when reflux of gastric contents causes troublesome symptoms and/or complications (7). GERD has now been considered the most common gastrointestinal (GI) disease worldwide. Subsequent to this trend, burgeoning of clinical entities attributed to GERD has also occurred. These include many ear, nose and throat, pulmonary and allergic symptoms, which are collectively termed extra-esophageal reflux disease (8). Extra-esophageal reflux is thought to be one of main factors in the pathogenesis of globus sensation (9, 10).

In this review, we will first review potential causes and treatments for globus sensation, and then we will introduce our data about the efficacy of PPI therapy for Japanese GERD patients complaining of globus sensation and to elucidate the characteristics of esophageal motility in patients showing resistance to PPI therapy.

Potential causes of globus sensation

Until now, many causes of globus sensation have been suggested, and it has been linked to several conditions from organic diseases such as pharyngeal cancer (11), Zenker’s diverticulum (12), or thyroid enlargement (13) to non-organic diseases such as functional diseases. Table 1 shows potential causes of globus sensation which have been reported in previous papers.

Organic diseases causing globus sensation

First of all, excluding organic diseases is clinically important. A recent study by Nagano et al. (14) showed that 36.5% of patients with globus sensation had esophageal diseases in the absence of an aberrant otorhinolaryngological region, which suggests that consideration of esophageal disorders is important in patients with globus sensation without laryngeal and hypopharyngeal findings. In our study (15), among the 350 patients complaining of globus sensation, 13 patients (3.7%) had endoscopic evidence of organic disease, including 10 patients with esophageal cancer, 2 patients with pharyngeal cancer, and 1 patient with Zenker’s diverticulum

Table 1. Potential causes of globus sensation

Organic diseases
Gastroesophageal reflux disease
Pharyngeal inflammatory causes including: pharyngitis, tonsillitis and chronic sinusitis
Upper aerodigestive malignancy
Hypertrophy of the base of the tongue
Retroverted epiglottis
Thyroid diseases
Cervical heterotopic gastric mucosa
Rare laryngopharyngeal tumors
Dysfunction of salivary secretion
Non-organic diseases
Abnormal upper esophageal sphincter function
Esophageal motor disorders
Psychological factors and stress

Fig. 1. Disposition of patients with globus sensation. From Tsutsui H et al. (15). Reproduction by copyright permission of Informa UK Ltd.

(Fig. 1). Based on these findings, otolaryngologists who generally do not use an upper GI endoscopy should remember that infection and malignant tumor cannot be ruled out by routine examination by otolaryngologists and such cases should be referred to a facility or department at which an upper GI endoscopy can be performed, when the symptom of globus sensation does not improve by any medication.

On the other hand, several recent papers have shown that globus sensation has also been linked to the presence of cervical heterotopic gastric mucosa, from which acid secretion might cause the symptom of globus sensation (16–18). Furthermore, Alagozlu et al. revealed that *Helicobacter pylori* infection of the cervical heterotopic gastric mucosa might be related to its symptom, although further studies will be necessary to confirm this result (19).

Association between globus sensation and GERD

Although there is still considerable debate about the causative role of GERD patients complaining of globus sensation, gastroesophageal reflux has been suggested to be a major etiology of this symptom, potentially accounting for 23%–68% of patients with globus sensation (20, 21). A recent population-based study also showed that typical symptoms of GERD such as heartburn or regurgitation were associated with globus sensation (odds ratio: 3.9; 95% confidence interval: 1.5–9.7) (22). In addition, it was reported that proximal acid exposure occurred in half of all patients with laryngeal symptoms (including globus sensation) or abnormal laryngoscopic findings, versus no GERD patients without laryngeal symptoms (23).

Currently, two main mechanisms are now thought to underlie the occurrence of globus sensation associated with acid reflux. First, acidic reflux into the esophagus may stimulate a vagal reflex that leads to the patient noting a globus sensation. Second, if refluxate passes into the larynx, direct contact with gastric acid and pepsin could cause mucosal injury that leads to the patient noting a globus sensation (9). Although both of these mechanisms are related to GERD, proton pump inhibitor (PPI) therapy often achieves disappointing results.

Association between globus sensation and esophageal motility

It has been reported that there is a high prevalence of esophageal motor abnormalities, including upper esophageal sphincter (UES) dysfunction (24), in patients who complain of globus sensation resistant to PPI therapy without any organic diseases, although the evidence obtained has been inconsistent.

1) Abnormal upper esophageal sphincter function

Abnormal UES function has been suggested to be one of the important causes of globus sensation (25–28). Elevated UES pressure has been found to be significantly more frequent in patients with globus sensation than in controls (28% vs 3%), suggesting that hypertensive UES is a crucial background factor for globus sensation (29). Another study to evaluate the effectiveness of injection of botulinum toxin into the cricopharyngeal muscle in patients with both globus sensation and high UES pressure showed that a decrease in UES pressure by botulinum toxin led to resolution of the globus sensation (25). In a recent study of high-resolution manometry in patients with globus sensation, normal controls, and GERD patients without globus sensation, hyperdynamic respiratory UES pressure changes were most prevalent in patients reporting globus sensation (28). However, results regarding to the UES function in patients with globus sensation were inconsistent because of a small number of subjects enrolled and differences in modalities used in each study (30–32).

2) Esophageal motor disorders

While there is some variation among previous studies, patients with globus sensation have been reported to have some kind of esophageal motor disorder, suggesting that esophageal motor disorders are a possible cause of, or a contributing factor in the development of globus sensation (33–36).

Moser et al. (37) noted that esophageal motor disorders might, before giving rise to dysphagia, be sensed more vaguely and induce the globus sensation. Esophageal manometry has revealed abnormalities in as many as 67% of patients with globus sensation, with nonspecific esophageal motility disorder being the most frequent finding (35). Our study to evaluate the characteristics of esophageal motility dysfunction in patients with globus sensation resistant to PPI treatment (15) showed that of the 47.9% of them that had abnormal esophageal motility, 66.4% had ineffective esophageal motility (Fig. 2), 14.4% had achalasia, 9.6% had diffuse esophageal spasm, 8% had nutcracker esophagus, and 1.6% had hypertensive lower esophageal sphincter. This suggests that there were significant differences of UES pressure and esophageal body peristalsis between the patients

Fig. 2. Typical multichannel intraluminal impedance and manometry (MII-EM) image of patients with ineffective esophageal motility dysfunction: This is the case of a proton pump inhibitor (PPI)-resistant GERD patient complaining of globus sensation. The MII-EM shows that peristaltic dysfunction in the lower esophagus causes incomplete bolus transit. From Tsutsui H et al. (15). Reproduction by copyright permission of Informa UK Ltd.

Table 2. Comparison of esophageal function parameters between healthy controls and the proton pump inhibitor-resistant GERD patients complaining of globus sensation groups

	Healthy controls (<i>n</i> =42)	PPI-resistant LPRD (<i>n</i> =98)	<i>P</i> -Value
Peak UES pressure after swallowing (mmHg)	346.5.1 ± 81.2	288.1 ± 97.4	0.0054
Resting UES pressure (mmHg)	71.1 ± 31.0	52.0 ± 20.3	0.0007
UES relaxation duration (s)	0.5 ± 0.1	0.5 ± 0.1	N.S.
LES pressure (mmHg)	16.7 ± 6.0	18.8 ± 10.4	N.S.
Residual LES pressure (mmHg)	2.2 ± 2.7	5.4 ± 4.7	0.0003
Amplitude (20 cm above LES) (mmHg)	39.1 ± 16.8	33.9 ± 16.0	N.S.
Amplitude (15 cm above LES) (mmHg)	50.3 ± 21.8	38.7 ± 20.0	0.0049
Amplitude (10 cm above LES) (mmHg)	63.0 ± 19.0	53.2 ± 24.7	0.0328
Amplitude (5 cm above LES) (mmHg)	78.0 ± 26.8	64.4 ± 29.5	0.0151
Distal esophageal amplitude (mmHg)	71.7 ± 19.9	59.6 ± 25.4	0.0084
Velocity (cm/s)	4.8 ± 1.8	3.9 ± 1.8	0.0181
Peristaltic contractions (%)	90.5 ± 10.8	71.8 ± 31.7	<0.0001
Complete bolus transit (%)	83.5 ± 10.9	53.6 ± 28.9	<0.0001

Abbreviations: PPI = proton pump inhibitor, LPRD = laryngopharyngeal reflux disease, UES = upper esophageal sphincter, LES = lower esophageal sphincter. From Tsutsui H et al. (15). Reproduction by copyright permission of Informa UK Ltd.

with PPI-resistant GERD patients complaining of globus sensation and healthy controls matched for age and sex (Table 2).

Recently, one mechanism that was newly proposed is that globus sensation might in part be associated with excessive laryngeal and pharyngeal tension, although there are several points to address regarding this question (38).

To validate the etiological significance of esophageal motility disorder in patients with globus sensation, further studies will be necessary to investigate whether the treatment of esophageal motility dysfunction would resolve globus sensation.

Visceral hypersensitivity

Visceral hypersensitivity has been identified in a number of functional gastrointestinal disorders including non-cardiac chest pain (39) functional dyspepsia (40) and irritable bowel syndrome (41). Chen et al. investigated the esophageal visceral perception in patients with globus sensation by using an esophageal balloon distension and electrical stimulation. They found patients with globus sensation demonstrate esophageal visceral hypersensitivity to mechanical distension: the differential responses to stretch and electrical stimuli may indicate that the hypersensitivity is a peripheral, rather than central phenomenon (42).

Psychological factors and social stress

It has been reported that several psychological problems or social stress have often been considered to cause or trigger globus sensation (43). In the past, several studies have shown that higher levels of psychological distress including anxiety or depression in patients with globus sensation (44), although a few data showed that there were no differences in the psychological states between patients with globus sensation and normal controls (34). A recent study showed that globus patients with laryngopharyngeal reflux (LPR) exhibited weaker psychological symptoms than non-LPR globus patients, and globus patients who did not respond to PPI had significantly higher anxiety scores (45), indicating a close relationship between psychological states and globus sensation.

Treatment

There have been no evidence-based treatment concepts available at this point in time, because of the few controlled trials for treatment of globus sensation. At present, there has been no single effective treatment for its symptoms. Therefore, the mainstays of its treatment are explanation and reassurance as for other functional gastrointestinal diseases (46). After explanation and reassurance, several possible treatment options based on the causes of globus sensation should be provided, such as anti-reflux therapy, therapy for improving gastrointestinal dysmotility including UES, treatment for decreasing the perception of globus sensation (visceral hypersensitivity), antidepressants and others.

Acid suppressive therapy

Since GERD is considered a major cause of globus sensation (47, 48), it seems reasonable that acid suppressive therapy such as PPI treatment should be the first-line therapy for patients with globus sensation after excluding organic diseases. Current evidence shows that the clinical response to PPI in patients with globus sensation is variable (8, 49) and that the symptom improves more slowly than typical GERD symptoms such as heartburn or regurgitation following acid-suppression therapy (20). Therefore, it is now widely accepted that patients with extra-esophageal GERD including globus sensation require more aggressive and more prolonged therapy than those with typical GERD (50). Empirical twice-daily therapy with PPI for at least 3 months is recommended; this can be extended for a maximum period of 6 months (51, 52). Our recent study that investigated the effectiveness of double doses of PPI (rabeprazole sodium at 20 mg daily for 4 weeks) in GERD patients with globus sensation showed that the response rate with PPI was 44.4% and that PPI-responsive patients had

significantly higher scores than the PPI-resistant patients for reflux-related symptoms (15). Our results suggested that the more acid reflux is involved in the etiology of patients with globus sensation, the more effective PPI becomes even though the duration of administration of PPI is relatively short. A recent study by Jeon et al. also showed the same results as ours, namely that the presence of reflux symptoms and short symptom duration were independent predictors of responsiveness to 4-week PPI treatment in patients with globus sensation (53).

However, recent studies have suggested some potential negative sequelae to long-term, high-dose PPI management (54). Therefore, it is necessary to remember that patients with globus sensation require more aggressive and more prolonged PPI therapy than those with typical GERD, and at the same time, prescribing a high dose of PPI for too long might lead to several side effects. Our data showed that 55.6% of GERD patients with globus sensation were resistant to PPI therapy, among whom 47.9% had abnormal esophageal motility with the most common esophageal abnormality being ineffective esophageal motility (15), suggesting that half of the patients with globus sensation basically are not related to acid reflux. We should change from PPI alone to another treatment such as prokinetics alone or adding prokinetics to PPI while monitoring its effectiveness in resolving patients' symptoms.

Therapy for improving gastrointestinal dysmotility including UES

A recent study suggested that stimulation of gastric emptying or esophageal clearance in addition to inhibition of gastric acid secretion may be an effective treatment for LPR. Ezzat et al. (55) reported that adding prokinetics, such as cisapride and itopride to PPIs to treat LPR reduced the recurrence of symptoms including globus sensation. Another recent study showed that rikkunshito, a Japanese herbal medicine, improved globus sensation in patients with PPI-refractory LPR, in part, because of stimulation of gastric emptying, suggesting that rikkunshito is an effective treatment for PPI-refractory LPR including globus sensation (21).

Furthermore, speech therapy has been successfully used to treat patients with persistent globus sensation to relieve vocal tract discomfort and tension (38). Recently Khalil et al. compared the efficacy of speech therapy to reassurance by a nurse practitioner for globus sensation, and found a significant improvement in the speech therapy group compared to controls (56). However, it is not clear whether there is a specific effect from the speech therapy or if the improvement is due to increased reassurance.

Treatment for decreasing the perception of globus sensation (visceral hypersensitivity)

Cognitive-behavioral therapy or antidepressants were considered as second-line or third-line therapy for patients with globus sensation.

1) Cognitive-behavioral therapy

It has been reported that a cognitive-behavioral therapy was the best treatment for a variety of somatoform disorders and medically unexplained symptoms (57). Therefore cognitive-behavioral therapy is now considered to be an effective therapy for patients with globus sensation, because globus sensation is the fourth most common symptom of somatization disorder after vomiting, aphonia, and pain in the extremities (58). Recently, Kiebles et al. also revealed that hypnotically assisted relaxation therapy could provide a substantial improvement in globus sensation irrespective of its cause, while esophageal motility including UES was unaffected (59). From their results of the combination of high responsiveness to hypnotically assisted relaxation therapy and no apparent impact on physiology, they speculated that globus sensation might be centrally mediated, although further studies will be necessary to address this question.

2) Antidepressants

The visceral hypersensitivity is considered one of features of globus sensation (42), and might be modulated by antidepressant therapy. A close relationship has been reported between psychological states including depression and globus sensation. A small series of anti-depressants have also been found to be beneficial for some globus patients with concomitant psychiatric disorders, such as panic, somatization, major depression, and agoraphobia (60, 61).

Based on these findings, cognitive-behavioral therapy or antidepressants should be considered as second-line or third-line therapy for patients with globus sensation.

3) Gabapentin which is frequently used for neuropathic pain, migraine headaches, and nystagmus, or both

It is now recognized that neuralgia can cause cough and other head and neck symptoms including globus sensation (62–64). It has also been suggested that the globus sensation might actually be a peripherally-mediated phenomenon (42). Kirch et al. (65) investigated the effectiveness of treatment of globus sensation with PPI and/or gabapentin which is frequently used off-label for neuropathic pain, migraine headaches, and nystagmus, or both. In this study, 77% of patients with globus sensation could be helped by treating reflux or neuralgia. According to this result, they concluded that a trial of gabapentin should be considered for patients who do not respond or only partially respond to reflux management. They speculated that it might be a potential neurogenic cause for globus sensation because acid reflux irritates nerve endings in the esophagus, pharynx, or larynx.

Other treatments

There have been several other treatments in addition to the above therapies. First, thyroidectomy in patients with thyroid disorder (66) or partial epiglottectomy in selected cases whose retroverted epiglottis made contact with the tongue base (67, 68), were both reported to significantly relieve the globus symptom.

Second, recent several studies showed that an ablation of cervical heterotopic gastric mucosa by argon plasma coagulation had shown some promise in improving chronic globus symptoms (69, 70). Therefore, an ablation of cervical heterotopic gastric mucosa by argon plasma coagulation is considered to be one of the precious alternative therapies for patients with globus sensation resistant to any medical treatments.

Closing remarks

There is no standard treatment strategy for patients with globus sensation despite its common clinical symptom. It is partly because its etiology remains uncertain. Globus sensation has been known as one of the multifactorial diseases, and recent several studies show that GERD is a major cause of globus sensation as interests tend to focus on GERD. However, efficacy of PPI for patients with globus sensation is limited to clinical situations, suggesting that its etiology might not be always related to gastric acid reflux. Our recent study demonstrated that 4 weeks of double doses of PPI therapy was effective in only 44.4% patients with globus sensation and suggested that esophageal motility dysfunction (including UES dysfunction) plays an important role in the pathogenesis of PPI-resistant GERD patients with globus sensation. Although there have been appeared several promising therapies for these patients, no large studies to investigate its efficacy have been reported. In the future, well-designed, randomized controlled studies will be needed to definitively determine the effect of several treatment strategies on globus sensation, as well as to ascertain the etiology of globus sensation *via* large-scale studies.

Conflict of interest

The authors have no conflict of interest.

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