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CLINICAL TRIAL REPORT

Aural stimulation with capsaicin ointment improved swallowing function in elderly patients with dysphagia: a randomized, placebo-controlled, double-blind, comparative study

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Objective: The aim of this study was to assess whether aural stimulation with ointment containing capsaicin improves swallowing function in elderly patients with dysphagia. **Study design:** A randomized, placebo-controlled, double-blind, comparative study. **Settings:** Secondary hospital.

Patients and methods: Twenty elderly dysphagic patients with a history of cerebrovascular disorder or Parkinson's disease were randomly divided into two groups: 10 receiving aural stimulation with 0.025% capsaicin ointment and 10 stimulated with placebo. The ointments were applied to the external auditory canal with a cotton swab. Then, swallowing of a bolus of blue-dyed water was recorded using transnasal videoendoscopy, and the swallowing function was evaluated according to both endoscopic swallowing scoring and Sensory-Motor-Reflex-Clearance (SMRC) scale.

Results: The sum of endoscopic swallowing scores was significantly decreased 30 and 60 min after a single administration in patients treated with capsaicin, but not with placebo. Reflex score, but not Sensory, Motion and Clearance scores, of the SMRC scale was significantly increased 5, 30 and 60 min after single administration in patients treated with capsaicin, but not with placebo. No patient showed signs of adverse effects.

Conclusion: As capsaicin is an agonist of the transient receptor potential vanilloid 1 (TRPV1), these findings suggest that improvement of the swallowing function, especially glottal closure and cough reflexes, in elderly dysphagic patients was due to TRPV1-mediated aural stimulation of vagal Arnold's nerve with capsaicin, but not with a nonspecific mechanical stimulation with a cotton swab.

Keywords: dysphagia, capsaicin, cough reflex, videoendoscopy, SMRC scale, external auditory canal

Introduction

Dysphagia affects elderly and neurological patients, increasing the risk for aspiration pneumonia.¹ Patients with dysphagia show difficulty in moving a bolus safely from the oral cavity to the esophagus, which allows it to penetrate into the airways. Thus, dysphagia can cause severe complications such as aspiration pneumonia, respiratory infection and malnutrition or death.

There is evidence that dysphagia is related to oropharyngolaryngeal sensory deficits, and that the therapeutic sensory stimuli on the oropharynx compensate for the loss of oropharyngolaryngeal sensitivity in elderly dysphagic patients.² In fact, it was reported that vagal sensory stimulation in the pharynx with a capsaicin pastille

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© 2017 Kondo et al. This work is published and licensed by Dove Medical Press Limited. The full terms of this license are available at https://www.dovepress.com/terms.php and incorporate the Creative Commons Attribution — Non Commercial (unported, v3.0) License (http://creative.commons.org/licenses/by-nc/3.0/). By accessing the work you hereby accept the Terms. Non-commercial uses of the work are permitted without any further permission from Dove Medical Press Limited, provided the work is properly attributed. For permission for commercial use of this work, please see paragraphs 4.2 and 5 of our Terms (https://www.dovepress.com/terms.php). improves the delay in swallowing function in elderly patients with dysphagia.³ However, oral administration of capsaicin pastille has the risk of causing aspiration in these patients. Accordingly, we then proposed the hypothesis that ectopic stimulation of the Arnold's branch of the vagus in the external auditory canal with capsaicin also improved swallowing function. In our previous study, we indeed showed that aural stimulation with capsaicin ointment improved swallowing function in elderly patients with dysphagia, suggesting that aural stimulation with capsaicin ointment may be a safer therapy for dysphagia.⁴

Capsaicin is the main component of red pepper. It is an agonist of transient receptor potential vanilloid 1 (TRPV1)5 and activates TRPV1 in peripheral sensory C-fibers. As a subset of neurons in the vagal nodose ganglion expresses TRPV1,6 our previous study suggested that ointment containing capsaicin activates TRPV1 in vagal Arnold's nerve of the external auditory canal, resulting in improvement of the swallowing function.⁴ However, we cannot exclude the possibility that a nonspecific mechanical stimulation of the external auditory canal with a cotton swab rather than the TRPV1-mediated stimulation with capsaicin improves swallowing function in dysphagic patients. To exclude this possibility, we designed a randomized, placebo-controlled, double-blind, comparative study to compare the effects of ointment containing capsaicin on swallowing function to those of placebo in elderly patients with dysphagia.

Patients and methods Study design

This study was a double-blind, randomized controlled, comparative study. It was conducted at Kochi National Hospital from July 2014 to March 2015. It was approved by the Committees for Medical Ethics of Kochi National Hospital and Tokushima University Hospital, and a written informed consent was obtained from each patient prior to the study. The Clinical Trial registration number of this study is UMIN000013443.

Subjects

Twenty elderly patients with dysphagia were enrolled in this study. The sample size was calculated based on the following parameters: α =0.05, β =0.20, power =80%, estimated difference =1.5 and SD=1.0. The calculation determined that the minimal size for each group was eight patients. Accordingly, 20 patients were enrolled and divided into two groups: 10 patients in the capsaicin group and 10 in the placebo group. Their baseline clinical characteristics are shown in Table 1. Patients complained of choking upon

swallowing water or food, but had no obstructive lesion in the pharyngolarynx as confirmed by endoscopy. They suffered from an old cerebrovascular disorder or Parkinson's disease, and no patients had an active ear disease such as otitis externa or myringitis.

Intervention

The patients had a baseline videoendoscopic evaluation of the swallowing function using both endoscopic swallowing scoring and Sensory–Motor–Reflex–Clearance (SMRC) scale. They were then divided into two groups by computerized randomization: a capsaicin group (group C) and a placebo group (group P). Their swallowing function was evaluated by transnasal videoendoscopy 5, 30 and 60 min after a single application of 0.5 g of 0.025% capsaicin or placebo ointment to the right external auditory canal with a cotton swab under an otoscope (Figure 1). The randomization and supply of ointment were performed by a blinded pharmacist who was not involved in the study. Throughout the study, the patients were observed to evaluate the occurrence of any adverse effects such as otalgia or headache.

Capsaicin ointment and placebo

Based on the *Japanese Pharmacopoeia* (16th edition) published by the Ministry of Health, Labor and Welfare of Japan, 0.025% capsaicin ointment was prepared according to the protocol of Japanese Drug Preparation of Hospital Pharmacy (4th edition) as follows: 25 mg of capsaicin (Sigma-Aldrich Co., St Louis, MO, USA) was dissolved in 500 μ L of 100% ethanol (Wako Pure Chemical Industries, Ltd., Osaka, Japan) and the solution was then mixed with 100 g hydrophilic ointment. Placebo was only hydrophilic ointment of the same color and viscosity as the capsaicin ointment.

Table I Patient characteristics

Charasteristic	Capsaicin (n=10)	Placebo (n=10)	P-value
Age, years ^a	80.4±9.5	80.1±5.9	NS*
Sex: M/F, n	10/0	9/1	NS**
Pathology: Ce/Pa, n	9/1	8/2	NS**
Duration after onset, months ^b	15.5	20.0	NS*
ESS ^{a,c}	5.3±1.4	5.4±1.4	NS*
SMRC scale ^a			
Sensory score	2.5±0.5	2.8±0.4	NS*
Motion score	1.9±0.3	1.8±0.4	NS*
Reflex score	1.5±0.5	1.6±0.5	NS*
Clearance score	2.0±0	1.7±0.5	NS*

Notes: ^aMean \pm SD, ^bmedian, ^cESS, total score of endoscopic swallowing scoring. ^{*}Mann–Whitney U test, ^{**}Fisher's exact test.

Abbreviations: Ce, cerebrovascular disease; NS, not significant; Pa, Parkinson's disease; SMRC, Sensory–Motor–Reflex–Clearance.



Figure I Patient flowchart.

Videoendoscopy

The standard protocol of videoendoscopic evaluation of swallowing proposed by the Oto-Rhino-Laryngological Society of Japan was used.⁷ Accordingly, patients were seated facing an otolaryngologist. Water was dyed with blue food coloring for ease of visualization and given to the patient in a bolus of 3 mL. Swallowing of the blue-dyed water was recorded by the video rhinolaryngoscope system with a flexible fiber optic endoscope of 3.1 mm diameter (VNL-100S[®]; Pentax, Tokyo, Japan). The video images of swallowing were evaluated using both endoscopic swallowing scoring and SMRC scale by another otolaryngologist blinded to clinical data and independent from the examiner.

Evaluation of the swallowing function

Endoscopic swallowing scoring was used to evaluate the total swallowing function with videoendoscopy (Table 2).^{8,9} The endoscopic swallowing scoring consists of four swallowing components: 1) salivary pooling in vallecula and pyriform sinuses, 2) the response of glottal closure reflex induced by touching the epiglottis with endoscope, 3) location of the bolus at the time of swallowing onset assessed by endoscopic whiteout and 4) the extent of pharyngeal clearance after swallowing of blue-dyed water. Each item was scored on a scale of 0–3 as shown in Table 2, and the sum of each score was used as an index of total swallowing function. Clinically, the full score is 12, and a score of more than 7 indicates a serious risk for aspiration, while a score of 7 or less during endoscopic swallowing evaluation with test jelly reliably predicts the ability to eat pureed diets.⁹

The swallowing function consists of four subfunctions: sensory initiation of swallowing reflex, motion to hold a bolus in the oral cavity and induce laryngeal elevation, glottal closure and cough reflexes and pharyngeal clearance of a bolus into the esophagus.^{10,11} Accordingly, we used the SMRC scale to evaluate the four subfunctions of the swallowing function with videoendoscopy separately: 1) Sensory: initiation of swallowing reflex as assessed by endoscopic whiteout; 2) Motion: holding bolus in the oral cavity and inducing laryngeal elevation according to instructions; 3) Reflex: glottal closure and cough reflexes induced by touching epiglottis or arytenoids with endoscope and 4) Clearance: pharyngeal clearance of the bolus after

Table 2 Endoscopic swallowing scoring

I. Salivary pooling in vallecula and pyriform sinuses

- 0: No pooling
- I: Pooling at only vallecula
- 2: Pooling in vallecula and pyriform sinuses and no penetration into the larynx
- 3: Pooling in vallecula and pyriform sinuses and penetration into the larynx
- The response of glottal closure reflex induced by touching epiglottis with endoscope
 - 0: Marked reflex by one touch
 - I: Slow and/or weak reflex by one touch
 - 2: Reflex by two or three touches
 - 3: No reflex despite three touches
- The location of the bolus at the time of swallow onset assessed by endoscopic whiteout
 - 0: Pharyngeal
 - I: Vallecula
 - 2: Pyriform sinuses
 - 3: No swallowing
- 4. The extent of pharyngeal clearance after swallowing of blue-dyed water 0: No residues
 - Pharyngeal residues remain, but are absent after swallowing is attempted two or three times
 - 2: Pharyngeal residues remain, but do not penetrate into larynx
- 3: Pharyngeal residues remain and penetrate into larynx

Table 3 SMRC scale

I. Sei wh	nsory: Initiation of swallowing reflex as assessed by endoscopic niteout
3:	Bolus at vallecula
2:	Bolus at pyriform sinuses
1:	Bolus retention at bottom of pyriform sinuses
0:	No swallowing initiation
2. Mo	otion: holding bolus in oral cavity and inducing laryngeal elevation
aco	cording to instructions
2:	Both holding bolus in oral cavity and inducing laryngeal elevation
1:	Either holding bolus in oral cavity or inducing laryngeal elevation
0:	Neither of them
3. Re	flex: glottal closure and cough reflexes induced by touching
epi	iglottis or arytenoids with endoscope
3:	Brisk cough reflex
2:	Brisk glottal closure reflex with weak or delayed cough reflex
1: '	Weak or delayed glottal closure and cough reflexes
0:	Neither of them
4. Cle	earance: pharyngeal clearance of bolus after swallowing
3:	No bolus and saliva residues after swallowing once or twice
2:	No bolus residue after swallowing once or twice, but saliva residue
1:	No bolus residue after swallowing three times or more
0:	Bolus residue even after swallowing three times or more

Abbreviation: SMRC, Sensory–Motor–Reflex–Clearance.

swallowing (Table 3).¹² Each function except Motion was scored on a scale of 0-3, and Motion was scored on a scale of 0-2 as shown in Table 3. SMRC scale scores were used as an index of each swallowing subfunction. Clinically, Score 0 of any function indicates no oral feeding because of the high risk for aspiration. Score 1 or 2 of any function (1 of Motion) indicates the need for intervention such as changing the form of food and swallowing training. In our recent study, we also showed a significant correlation between cough reflex sensitivity to citric acid in the cough reflex test and Reflex score with videoendoscopy in elderly patients with dysphagia (manuscript in preparation).

Statistical analysis

The primary outcome was improvement of the sum of endoscopic swallowing scores and each score of SMRC scale in patients with dysphagia after aural stimulation with capsaicin ointment or placebo. Comparison of baseline clinical characteristics between groups C and P was carried out by Mann–Whitney U test and Fisher's exact test. Changes in endoscopic swallowing scoring and SMRC scale after stimulation with capsaicin ointment or placebo were analyzed by Friedman test and Shirley–Williams post hoc test, and $P \leq 0.05$ was considered statistically significant. Statistical analysis was done using Microsoft[®] Excel 2010 Japanese version (Microsoft Japan, Tokyo, Japan) and Statcel version 3 (OMS Publishing Inc., Saitama, Japan).

Results

Baseline clinical characteristics of each group are shown in Table 1. There were no statistical differences in age, sex, pathology, duration after onset, total score of endoscopic swallowing scoring or SMRC scale between groups C and P.

Changes in the sum of endoscopic swallowing scores after single aural administration of capsaicin ointment or placebo are shown in Figure 2. The endoscopic swallowing scores of group C were significantly decreased 30 and 60 min after single aural administration of capsaicin ointment (P=0.05). But the endoscopic swallowing scores of group P were unchanged after single aural administration of placebo. Mean changes in endoscopic swallowing scores from baseline (pre) to post-5, -30 and -60 min in groups C and P are shown in Figure 3. The decrease in endoscopic swallowing scores of group C was significantly greater than that of group P at post-30 and -60 min (P<0.05).



Figure 2 Changes in endoscopic swallowing scores after single aural administration in capsaicin and placebo groups. Notes: *P=0.003 in Friedman test; P=0.05 with Shirley–Williams post hoc test. Data shown as mean ± SD. n=10. Abbreviation: NS, not significant.



Figure 3 Mean changes in endoscopic swallowing scores from baseline (pre) to post-5, -30 and -60 min in capsaicin (black) and placebo (gray) groups. Note: *P<0.05 in Mann–Whitney U test.

Abbreviations: NS, not significant; ESS, endoscopic swallowing scoring.

Sensory, Motion, Reflex and Clearance scores of SMRC scale after single aural administration of capsaicin ointment or placebo are shown in Figures 4 and 5. Reflex scores, but not Sensory, Motion and Clearance scores, of group C were significantly increased 5, 30 and 60 min after single aural administration of capsaicin ointment (P=0.05). However, any scores of SMRC scale in group P were unchanged after a single aural administration with placebo. Mean changes in Reflex scores from baseline (pre) to post-5, -30 and -60 min

in groups C and P are shown in Figure 6. The increase of the Reflex score in group C was significantly higher than that of group P at post-30 and -60 min (P<0.05). No adverse effects including otalgia, headache, otitis externa and myringitis were induced in any patient up to 120 min after administration.

Discussion

In this study, the sum of endoscopic swallowing scores was improved up to 60 min after the stimulation of the external auditory canal with 0.025% capsaicin ointment in elderly patients with dysphagia. Moreover, this study demonstrates that among the four subfunctions of swallowing function, only the Reflex score of glottal closure and cough reflexes was improved by stimulation with capsaicin ointment. These effects were not observed in the placebo group, indicating that it was not a nonspecific mechanical stimulation, but the capsaicin ointment applied to the external auditory canal that improved their swallowing function, especially glottal closure and cough reflexes. Furthermore, nonresponders in group C suggest individual differences in the effects of capsaicin ointment. But the nonspecific mechanical stimulation of cotton swab might improve the laryngeal reflex in some responders in group P.

Accordingly, it is suggested that capsaicin-TRPV1mediated stimulation of the Arnold's branch of the vagus in



 $\label{eq:states} \begin{array}{l} \mbox{Figure 4} Changes in SMRC scale after single aural administration in capsaicin group. \\ \mbox{Notes: } {}^{*}P{=}0.003 \mbox{ in Friedman test; } {}^{*}P{=}0.05 \mbox{ with Shirley-Williams post hoc test. Data shown as mean \pm SD, n=10. \\ \mbox{Abbreviations: NS, not significant; SMRC, Sensory-Motor-Reflex-Clearance.} \end{array}$



Figure 5 Changes in SMRC scale after single aural administration in placebo group. Note: Data shown as mean \pm SD, n=10. Abbreviations: NS, not significant; SMRC, Sensory–Motor–Reflex–Clearance.

the external auditory canal improved swallowing function in these elderly dysphagic patients. Indeed, the auricular branch of the vagus, the Arnold's nerve, is distributed to the external auditory canal, and its stimulation triggers cough reflex.¹³ This is frequently encountered by otolaryngologists in the management of the external auditory canal, such as ear syringing.¹⁴ As the stimulation of sensory C-fiber branches of the vagus in the laryngotracheal mucosa with capsaicin induces cough reflex,¹⁵ it is suggested that stimulation of



Figure 6 Mean changes in reflex scores from baseline (pre) to post-5, -30 and -60 min in capsaicin (black) and placebo (gray) groups. Note: *P < 0.05 in Mann–Whitney U test. Abbreviation: NS, not significant. Arnold's nerve, a branch of the vagus distributed to the external auditory canal, with capsaicin improved glottal closure and cough reflexes in dysphagic patients. On the other hand, it was reported that oral administration of a pastille containing capsaicin improved the delay in swallowing function in patients with dysphagia.³ Oral administration of capsaicin stimulates the sensory C-fibers of the trigeminal, glossopharyngeal and vagal nerves in the oral and pharyngolaryngeal mucosa, but aural stimulation with capsaicin stimulates only the Arnold's branch of the vagus. It is, therefore, suggested that oral administration of capsaicin improves the delay in swallowing function, while aural stimulation improves only the Reflex score of glottal closure and cough reflexes, but not Sensory, Motion and Clearance scores in dysphagic patients.

Glottal closure and cough reflexes are important airway protective mechanisms against aspiration. It was shown that sensitivity of the cough reflex was reduced in patients with aspiration pneumonia.¹⁶ Angiotensin-converting enzyme inhibitors have been shown to reduce the prevalence of aspiration pneumonia in post-stroke patients, because angiotensin-converting enzyme inhibitor induces cough.^{17–19} This study demonstrated that the aural stimulation with capsaicin improved glottal closure and cough reflexes in elderly

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patients with dysphagia, and future studies are necessary to investigate if the aural stimulation with capsaicin could have similar application to prevent aspiration pneumonia. Chronic exposure to high-dose capsaicin causes long-term functional impairment of sensory fibers due to desensitization of TRPV1 and depletion of neuropeptides such as substance P (capsaicin defunctionalization).²⁰ However, in our previous study, repeated alternative application with ointment containing 0.025% capsaicin to each external auditory canal once a day for 7 days still improved swallowing function in patients with dysphagia.⁴ Therefore, once-a-day alternative application to each external auditory canal with low-dose capsaicin may be a useful method to prevent capsaicin defunctionalization, although it is still uncertain how long the effects of single application with capsaicin ointment last. Furthermore, multicenter studies are required to explore whether repeated aural stimulation with 0.025% capsaicin ointment in such a way to prevent capsaicin defunctionalization really reduces the incidence of aspiration pneumonia in patients with dysphagia.

There are some limitations to this study. There was a preponderance of males. The endoscopic swallowing scoring and SMRC scale are subjective evaluations.²¹ Further studies are needed to objectively evaluate the swallowing function and cough reflex using videofluoroscopy²² and cough reflex test.²³ Aural stimulation with capsaicin ointment is effective in patients with nonobstructive, but not obstructive dysphagia.

In conclusion, this study demonstrated that the topical application of an ointment containing 0.025% capsaicin to the external auditory canal improves swallowing function, especially glottal closure and cough reflexes, in elderly patients with dysphagia. Therefore, repeated aural stimulation with capsaicin ointment may be a safe and promising intervention to prevent aspiration pneumonia in elderly dysphagic patients due to improvement of glottal closure and cough reflexes via Arnold's ear-cough reflex.

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Author contributions

EK designed the study, collected data and wrote a part of the manuscript. OJ designed the study and analyzed data. SN collected data. HO analyzed, interpreted data and wrote a part of the manuscript. IK interpreted data. NT supervised the study

and corrected the manuscript. All authors contributed toward data analysis, drafting and revising the paper and agree to be accountable for all aspects of the work.

Disclosure

The authors report no conflicts of interest in this work.

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