Examining the relationship between sepsis and oropharyngeal dysphagia in hospitalised elderly patients: a retrospective cohort study

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<th>Journal</th>
<th>Frontline Gastroenterology</th>
</tr>
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<tr>
<td>Manuscript ID</td>
<td>flgastro-2018-100994.R1</td>
</tr>
<tr>
<td>Article Type</td>
<td>Research</td>
</tr>
<tr>
<td>Date Submitted by the Author</td>
<td>04-May-2018</td>
</tr>
<tr>
<td>Complete List of Authors</td>
<td>Sasegbon, Ayodele; Pennine Acute Hospitals NHS Trust, Gastroenterology; Hamdy, Shaheen; University of Manchester, Gastroenterology; Salford Royal NHS Foundation Trust, Gastroenterology</td>
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<tr>
<td>Keywords</td>
<td>SEPSIS, DYSPHAGIA, ELDERLY</td>
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Examining the relationship between sepsis and oro-pharyngeal dysphagia in hospitalised elderly patients: a retrospective cohort study

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Key Words: Critical Illness; Dysphagia; Elderly; Sepsis; Swallowing
SEPSIS AND ORO-PHARYNGEAL DYSPHAGIA IN HOSPITALISED ELDERLY PATIENTS

Abstract
Introduction Elderly people are recognised to be at increased risk of oropharyngeal
dysphagia (OPD), the causes of which are multifactorial. Our aim was to identify if sepsis is
associated with OPD in the elderly during hospitalisation in the absence of known other risk
factors for OPD.

Methods A hospital electronic database was searched for elderly patients (≥65 years)
referred for assessment for suspected dysphagia between March 2013 and 2014. Exclusion
criteria were age < 65 years, pre-existing OPD or acute OPD secondary to: acute intracranial
event, space occupying lesion or trauma. Data were collected on factors including: age; sex;
co-morbidities; existing OPD; sepsis; microbiology; recovery of OPD and medication. Sepsis
was defined as evidence of a systemic inflammatory response syndrome with a clinical
susicion of infection.

Results A total of 301 of 1761 screened patients referred for dysphagia assessment met the
inclusion criteria. The prevalence of sepsis and subsequent OPD was 16% (51/301). The
mean age was 83 years (median 81 years). The commonest co-morbidity was dementia
(31%). The majority (84%) failed to recover swallowing during their hospital stay, 12% had
complications of aspiration and 35% died. The commonest source of sepsis was from the
chest (55%). Other factors contributing to the risk for dysphagia included delirium (22%) and
neuroactive medication (41%). However, 10% of patients had sepsis and subsequent OPD
without other identified risk factors.

Conclusion The prevalence of sepsis and subsequent dysphagia is significant and should be
taken into account in any elderly person in hospital with new onset OPD without other
predisposing risk factors.
SEPSIS AND ORO-PHARYNGEAL DYSPHAGIA IN HOSPITALISED ELDERLY PATIENTS

1. What is known?

No studies have been performed looking at the association between sepsis and oropharyngeal dysphagia in ward based elderly patients. Moreover, limited research has been done in this field with only one study published which showed an association between sepsis and oropharyngeal dysphagia in septic patients on an intensive care unit.

2. New findings?

We have demonstrated an independent association between acute sepsis and oropharyngeal dysphagia in elderly ward based patients.

3. How may it impact on clinical practice?

The implication is that sepsis should be taken into account as a cause of oropharyngeal dysphagia in any elderly person in hospital with new onset OPD without other predisposing risk factors.
SEPSIS AND ORO-PHARYNGEAL DYSPHAGIA IN HOSPITALISED ELDERLY PATIENTS

Introduction

Swallowing is a fundamental process for enabling life, allowing food and fluid to be ingested safely and efficiently thereby maintaining normal physiological and biochemical functions. Furthermore the enjoyment of eating and drinking plays an important part in an individual’s perception of their quality of life (1, 2). Dysphagia is defined as difficult/disordered swallowing (3-5). Various diseases can cause disruption to normal swallowing resulting in dysphagia. Dysphagia has the potential to cause malnutrition, dehydration and aspiration pneumonia, all of which lead to significant morbidity and mortality (6).

The process of aging affects the way the oral cavity and pharynx function. However, these changes are a natural part of the aging process and are not considered pathological. Age related changes to the process of swallowing are collectively termed presbyphagia (2, 7). As part of the aging process there is also breakdown of muscle fibres and a resultant decrease in skeletal muscle mass; otherwise categorised as sarcopenia. Studies have also shown an age related decrease in tongue and pharyngeal musculature (8, 9) alongside a prolongation in the pharyngeal phase of swallowing in healthy elderly individuals (2, 5). These changes mean older individuals have a reduced swallowing reserve and are more prone to developing dysphagia.

Sepsis is a very common cause of mortality, hospitalisation and admission to intensive care units. 4.7% of deaths of deaths in the UK can be attributed to sepsis (10). Additionally sepsis is the cause of 27% of admissions to intensive care units (11). Older individuals are more prone to developing sepsis than their younger counterparts. Despite this, only one study has looked at critical illness, sepsis and OPD, prospectively investigating two groups of ICU patients, with and without sepsis (12). In that study, patients were evaluated using
fibrescopic endoscopic evaluation of swallowing (FEES) and questionnaires. Nineteen of 30 patients in the sepsis group had dysphagia compared to 7 of 30 in the non-sepsis group. At four months, mortality in the sepsis group was 57% compared to 20% in the non-sepsis group. It should be noted that because critical illness and increasing age have been shown to increase the risk of OPD, the presence of critical illness could be considered a confounding factor in this study.

Thus, given the lack of data concerning the role of sepsis in the aetiology of swallowing dysfunction our study aim was to quantify the prevalence of OPD in older hospitalised individuals (age ≥ 65), associated with sepsis alone, without the presence of any other known risk factors of OPD. Our hypothesis was that sepsis would act as an independent predictor of dysphagia in this setting.

**Methods**

**Patient Recruitment**

Patients were initially identified from the speech and language therapy (SALT) referral database in a single centre teaching hospital in the north of England. Subsequent data were drawn from analysis of the medical notes of individual patients using an electronic patient record system (Allscripts: Sunrise EPR. Salford, UK).

The study analysed referral data over a one year time frame from March 2013 to 2014. Patients’ clinical notes were scrutinised and compared to the research inclusion and exclusion criteria. Inclusion criteria were: Patients ≥ 65 years of age; a documented diagnosis of OPD by the SALT team during this admission; sepsis and a course of antibiotic
SEPSIS AND ORO-PHARYNGEAL DYSPHAGIA IN HOSPITALISED ELDERLY PATIENTS

therapy. Exclusion criteria were: patients < 65 years of age; significant documented pre-existing OPD secondary to any cause or acute OPD secondary to: stroke, head trauma; space occupying brain lesion or neurosurgical intervention. Patients were identified for the study using a sequential multistep process as illustrated in Figure 1. Ethical approval for the study was granted by the London Bromley research ethics committee (REC number 15/LO/1413).

Information Retrieval

A large amount of information was retrieved for each patient. This can be divided into: Demographics, pre-existing clinical information, dysphagia, Infection, and recovery.

Regarding demographics, all patients had the following information recorded: age, sex, location of referral, admission and discharge body mass index (BMI). Admission BMIs were recorded within an arbitrary time period of 7 days from admission and discharge BMIs within 7 days of discharge. For pre-existing clinical information, patients had their co-morbidities and medications (prior to and during admission) recorded. For dysphagia, pre-existing OPD was recorded if present, the presence or absence of a baseline SALT assessment was recorded; when OPD was suspected, the method of OPD diagnosis and any suspicion or diagnosis of aspiration pneumonia also recorded. For infection, any diagnoses of infection were recorded along with: presence or absence of SIRS criteria, location, if antibiotic treatment was initiated and if sepsis was confirmed by positive microbiology. Finally under recovery, the length of time taken to recovery of swallowing function – if any occurred – was recorded in addition to mortality.
SEPSIS AND ORO-PHARYNGEAL DYSPHAGIA IN HOSPITALISED ELDERLY PATIENTS

Definitions

Definition of Sepsis

The decision was made to define sepsis according to the SIRS based 2001 International sepsis criteria (figure 2) (13).

Definition of Dysphagia

For the purposes of this study, dysphagia was defined as documented diagnoses of dysphagia by the SALT team in patients’ clinical notes. This included clinical diagnoses as well as diagnoses supported by videofluoroscopy (VFS) and FEES.

This method of defining OPD was adopted because SALT staff have two formal roles in a hospital setting: the assessment and management of OPD and the assessment and management of speech disorders. Therefore, they have the most expertise in the clinical diagnosis and further assessment of OPD. Clinical assessment of OPD by trained professionals has been shown to be robust with 42-90% sensitivity and 59-91% specificity (14).

The SALT referral database was used as the basis of patient recruitment in this study. Referrals are sent using the electronic patient record to the shared SALT staff email. Any member of clinical staff can send a referral. Patients are placed nil by mouth until they are formally assessed by the SALT team.

Definition of “Elderly” Patients

In this study elderly patients were defined as being ≥ 65 years of age. This definition is in line with the world health organisation definition of an elderly individual (15).
SEPSIS AND ORO-PHARYNGEAL DYSPHAGIA IN HOSPITALISED ELDERLY PATIENTS

Data Analysis

As this was a retrospective cohort observational study, we calculated the number of cases of OPD subsequent to sepsis in elderly individuals. Prevalence was calculated using the software package SPSS. Results are reported as percentiles and medians with interquartile range (IQR) unless otherwise stated.

Results

Patient screening and recruitment

1761 patients were referred for dysphagia assessment during the study period. On initial screening of the referral database, exactly 301 met the inclusion criteria and were therefore retained for further analysis (Figure 1).

Patient demographics and dysphagia prevalence

The prevalence of sepsis and subsequent OPD, within this group of ≥65 year old, non-ICU patients without pre-existing OPD was 16% (51/301 patients). The other 250 patients were not diagnosed as having OPD when formally assessed by SALT despite the suspicion of the nursing staff. The mean age of the dysphagic group was 83 years with a median of 81 (IQR 77-91) years. There was a preponderance of males (59%). Of the OPD group, 5 (10%) had no other known risk factors of OPD (other than age).

Method of Diagnosis

OPD was diagnosed by clinical assessment in 47 (92%) patients. VFS was used to diagnose 3 (6%) patients and FEES in 1 (2%) patient.
Location of Referrals

46 (90%) patients were referred to the SALT team from medical wards. 5 (10%) patients were referred from the medical high dependency unit.

Types of Sepsis

Types of sepsis included: 28 chest (55%); 11 mixed (22%); 6 urological (12%); 2 biliary (4%); 1 cellulitis (2%); 1 intra-abdominal (2%); 1 gastroenteritis (2%) and 1 unknown (2%).

Confirmatory microbiology including sputum, urine, stool and blood cultures was positive in 15 (29%) patients.

Mortality and Aspiration

Six patients (12%) had suspected complications of aspiration and 18 (35%) died during their admission.

Contributing Factors

Other factors potentially contributing to the risk for dysphagia included delirium (22%) or confusion (29%) and intake of medication potentially contributing to OPD (41%). These included any neuroleptic agents or medications with a potential to impair an individual’s level of consciousness for example: opiates, benzodiazepines or other hypnotic agents.

Co-morbidities

The co-morbidities are shown in Table 1. Of the 51 patients with sepsis and subsequent dysphagia, the most common co-morbidities (defined as a prevalence of over 10%) were:

Dementia from all causes (31%), chronic obstructive pulmonary disease (COPD) (24%), cancer (20%), chronic kidney disease (16%), atrial fibrillation (14%), Parkinson’s disease/Parkinsonism (14%), diabetes (14%), ischaemic heart disease (12%) and heart failure (10%).
# SEPSIS AND ORO-PHARYNGEAL DYSPHAGIA IN HOSPITALISED ELDERLY PATIENTS

Table 1:

<table>
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<td>Chronic obstructive pulmonary disease</td>
<td>12</td>
</tr>
<tr>
<td>Cancer</td>
<td>10</td>
</tr>
<tr>
<td>Unspecified dementia</td>
<td>9</td>
</tr>
<tr>
<td>Chronic kidney disease</td>
<td>8</td>
</tr>
<tr>
<td>Atrial fibrillation</td>
<td>7</td>
</tr>
<tr>
<td>Type 2 diabetes mellitus</td>
<td>7</td>
</tr>
<tr>
<td>Parkinson's disease/Parkinsonism</td>
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</tr>
<tr>
<td>Ischaemic heart disease</td>
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</tr>
<tr>
<td>Cardiac failure</td>
<td>5</td>
</tr>
<tr>
<td>Vascular dementia</td>
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<tr>
<td>Previous stroke (without any baseline OPD)</td>
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</tr>
<tr>
<td>Epilepsy</td>
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<tr>
<td>Bronchiectasis</td>
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<tr>
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<td>Akinetic rigid syndrome</td>
<td>1</td>
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<tr>
<td>Oesophagitis</td>
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</table>
Recovery of Swallowing

Eight (16%) patients recovered their swallowing to their pre-sepsis baseline. This was defined as any alteration in their ability to eat and drink prior to admission as documented in their medical notes and clinic letters. Forty three patients (84%) failed to recover their swallowing to their baseline prior to being discharged or passing away. Twenty five patients who survived their admission, were discharged with dietary modification. Within this group, 6 exhibited signs of partial recovery of their swallowing function as determined by repeated SALT assessments prior to discharge. The time to partial or complete recovery of swallowing function ranged from 2 to 30 days.

Discussion

This retrospective cohort study was designed to quantify the prevalence of sepsis induced OPD in an older hospitalised population. Any association observed will provide additional evidence that will prompt further work in this under-studied area of medicine.

Our study found that the prevalence of sepsis associated dysphagia was 16%. Of these around 10% or one in 10 patients did not have any other known OPD associated risk factors other than sepsis. This is suggestive of sepsis having a true independent association with OPD in elderly patients. This could be due to the reduced swallowing reserves in this patient group which leaves them at a greater risk of decompensation. One potential explanation for a decompensation could be the effects of sepsis on oropharyngeal muscle function resulting in a slowing of oropharyngeal bolus transit. Skeletal and oropharyngeal muscles waste in a similar fashion with age making it more prone to dysfunction when sepsis is present. Age related muscle wasting is thought to explain some of the increased risk of dysphagia in
SEPSIS AND ORO-PHARYNGEAL DYSPHAGIA IN HOSPITALISED ELDERLY PATIENTS

elderly patients. Tongue wasting and weakness with aging has been shown to slow bolus
transit and increase the risk of OPD in healthy older people. Additionally concurrent
diseases of the elderly, which cause slowed oropharyngeal bolus transit, increase the risk of
OPD (16).

Another potential explanation for sepsis causing OPD is the recognised effect that sepsis has
on neuronal activity (17, 18). This (critical illness) polyneuropathy could potentially affect
the highly co-ordinated sequence of neuronal activity needed to cause the sequential
contraction of swallowing musculature. Any muscular incoordination would increase the risk
of OPD. Such incoordination may be particularly marked in elderly patients with a reduced
swallowing reserve. It is important to note, however, that the precise effect of sepsis on
neuronal firing remains poorly understood.

Fifty one of 301 (or 16%) patients without strokes, head trauma and brain lesions,
developed sepsis and subsequent OPD. These patients did not have OPD prior to arriving in
hospital despite the fact that the majority of them had other OPD associated risk factors
such as: opiate medications, COPD, antipsychotic medication use and chronic neurologic
illness. This implies that these factors in isolation were not sufficient to cause OPD. Instead
sepsis plus those factors appeared to act in synergy to provoke swallowing decompensation.

Interestingly, 14 of the 33 patients who did not die during their admission exhibited full or
partial recovery of OPD following the resolution of their sepsis. It can be seen that the
majority (19/33) of surviving patients with sepsis and subsequent OPD did not show any
signs of recovery of their swallowing function and were discharged with dietary
modification. No patients were documented to have undergone swallowing rehabilitation.

As swallowing rehabilitation has been shown by several studies to improve swallowing
SEPSIS AND ORO-PHARYNGEAL DYSPHAGIA IN HOSPITALISED ELDERLY PATIENTS

function (19,20) the lack of rehabilitation in this study may have contributed to the low number of patients who recovered their baseline swallowing function. Further work will be necessary to highlight the observation that sepsis related dysphagia may not be being aggressively managed in the hospital setting.

Limitations

There are several limitations of this study which need to be considered. The study was retrospective in nature. As a result although it provided a picture of what may be occurring, it does not allow incidence to be calculated. It must also be noted that although there is evidence of an association between sepsis and OPD in elderly patients, this does not imply causality. 12 of the 51 patients with sepsis and subsequent OPD had pre-existing COPD. A study by Cvejic et al showed patients with exacerbations of COPD have a significant delay in their swallowing reflex (21). This finding is thought to be due to a change between the coordination of swallowing and respiration. This is a possible causative factor for the presence of OPD in patients with COPD. However, it is interesting to note that none of the patients with pre-existing COPD who developed sepsis and subsequent OPD had any documented episodes of OPD prior to their sepsis.

Lastly no baseline swallowing investigations were done on admission to objectively confirm the absence of any swallowing dysfunction prior to a diagnosis of sepsis and subsequently a diagnosis of OPD. No patients fulfilling the study inclusion criteria had a baseline swallowing assessment performed on admission to hospital. This is not the case for patients admitted to hospital with suspected strokes (22). This means patients are commenced on unmodified fluids and food on admission unless they have a pre-existing diagnosis of OPD or known
dietary modifications. This lack of initial assessment means potential problems are identified late. It must also be noted that OPD does not always manifest as overt swallowing dysfunction and aspiration. OPD can also cause micro-aspirations which are more difficult to detect (23). Indeed, 28 patients had chest sepsis and in this patient group there is a possibility that silent aspiration preceded their chest sepsis. However, careful review of their notes was used to identify and exclude any patients where any pre-existing dysphagia was documented. While a potential limitation, it is important to note an association between sepsis and OPD was observed in cases of non-chest sepsis.

Conclusion

This study provides new data on how sepsis may be associated with OPD in elderly, non-ICU patients. Around 1 in 6 non-stroke, non-trauma patients referred to the SALT team for assessment had sepsis in association with OPD. Of these, 10% developed sepsis and OPD without any other OPD associated risk factors. The association between sepsis and OPD should be taken into account in any new onset aspiration event in older hospitalised patients.
SEPSIS AND ORO-PHARYNGEAL DYSPHAGIA IN HOSPITALISED ELDERLY PATIENTS

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Competing Interest

Competing Interest: None declared.

Funding

No funding was received for this study.

Contributorship Statement

AS planned the study, collected the data, wrote and submitted the paper. SH helped plan the study and write the paper.
References

22. NICE. Stroke and transient ischaemic attack in over 16s: diagnosis and initial management https://www.nice.org.uk/guidance/cg68: NICE; 2008

Figure Captions
Figure 1: Flow diagram illustrating the multistep process of patient recruitment.
Figure 2: SIRS criteria

Table Captions
Table 1: Table containing co-morbidities of patients with sepsis and subsequent OPD.

*None of the above patients irrespective of their co-morbidity had pre-existing OPD prior to their admission to hospital.
Flow diagram illustrating the multistep process of patient recruitment

175x203mm (300 x 300 DPI)
Severe sepsis
Sepsis and organ dysfunction

Sepsis
2 SIRS criteria and suspicion/evidence of infection

SIRS
Temperature < 36C or > 38C
White cell count < 4000 or > 12000 cells/ul
Heart rate > 90 beats per minute
Respiratory rate > 20 breaths per minute

SIRS criteria

131x85mm (300 x 300 DPI)