# A comparison of normal saline mouth rinse and mouth rinse based on Salvia officinalis in palliative care: A randomized controlled trial

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	Total sample (n=88)	Normal Saline (n=44)	Salvie officinalis (n=44)	P-value
Age, years				0.728
Mean (SD)	63.9 (10.6)	63.5 (11.8)	64.3 (9.4)	
Range	29 – 84	29 – 84	45 – 83	
Sex, % (n)				0.151
Male	27% (24)	20% (9)	34% (15)	
Female	73% (64)	80% (35)	66% (29)	
Karnofsky score	(n=83)	(n=43)	(n=40)	0.504
Mean (SD)	52.1 (16.9)	53.3 (18.4)	50.8 (17.2)	
Range	20 – 80	20 – 80	20 – 80	
Primary diagnosis				0.325
Gastrointestinal cancer	26% (23)	29% (13)	23% (10)	
Lung cancer	17% (15)	16% (7)	18% (8)	
Gynecologic cancer	16% (14)	23% (10)	9% (4)	
Prostate cancer	3% (3)	5% (2)	2% (1)	
Breast cancer	13% (11)	9% (4)	16% (7)	
Other cancer	25% (22)	18% (8)	32% (14)	
Head/Neck	8% (7)	7% (3)	9% (4)	
Number of medications	(n=85)	(n=44)	(n=41)	0.161
Mean (SD)	11.4 (4.2)	10.8 (4.6)	12.1 (3.6)	
Range	4 – 26	4 – 26	4 – 20	

Table 1: Sample demographic and clinical characteristics.

### Background

There are few clinical studies evaluating the effect of interventions within oral palliative care. A mouth rinse solution made from the herb Salvia officinalis (SO) has been used in our unit for many years to improve patients' oral health.

### Aim

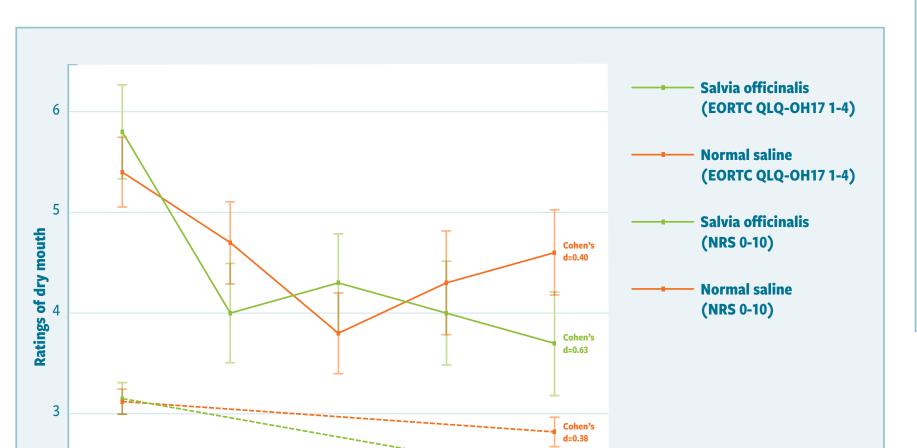
This study examines the efficacy of a SO-based herbal mouth rinse in comparison with normal saline (NS) mouth rinse on oral health and oral symptoms.

**Patient-reported oral** 

symptoms at baseline.

# Method

This study was a block-randomized controlled trial. Data were collected in an inpatient hospice unit before and after a 4-day oral hygiene intervention with either SO (n=44) or NS (n=44). Numerical Rating Scales (NRS 0-10) and 12 items taken from the European Organisation for Research and Treatment of Cancer Quality of Life Questionnaire-Oral Health 17 (EORTC QLQ-OH17) were used to measure oral symptoms. An oral examination was performed before and after the intervention.



Day 4

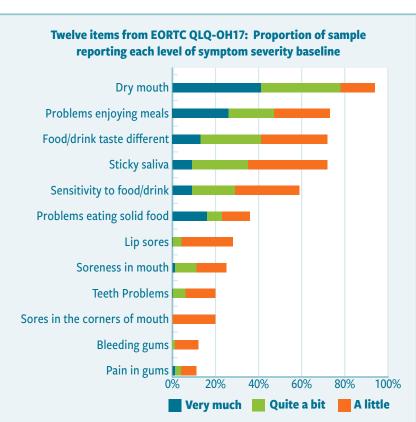


Figure 2:
Changes in EORTC QLQ-OH17 and
NRS ratings based on dry mouth
by treatment group.

Day 1

Pre-interv

Day 2

Day 3

# **Results**

Of the 538 patients admitted to the unit, 88 were included, and 73 (83%) completed the study. Of the 159 who met the eligibility criteria only 22 decline to participate. There was no group differences at baseline (Table 1). At baseline 78% reported dry mouth on the EORTC QLQ-OH17 questionnaire, and 80% rated dry mouth  $\geq$  4 on the NRS. Overall symptom scores (from the EORTC QLQ-OH17) improved in both

groups (p<.001), with dry mouth ratings showing particular improvement in the SO group (p=0.036; Cohen's d=0.75). NRS ratings of dry mouth also improved in the SO group (p=0.045; Cohen's d=0.63)(Figure 2). OMAS erythema scores improved for the NS group only. Plaque on the tongue (p=0.003), teeth (p<0.001) and sign of dry mouth (p=0.001) improved in both groups.

d=0.75

# **Conclusion**

This study detected no significant differences between Salvia officinalis and normal saline rinses in the treatment of oral symptoms, but showed that systematic assessment and oral care improved oral wellbeing and reduced oral discomfort.

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