

Good quality epidemiological studies of oro-facial pain needed

What is the prevalence and associated risk factors of oro-facial pain (OFP)?

Macfarlane TV, Glenny AM, Worthington HV. *Systematic review of population-based epidemiological studies of oro-facial pain.* *J. Dent.* 2001; 29(7):451–67

Data sources MEDLINE, Embase, Cinahl, Scientific Citation Index and Health CD databases. Reference lists of relevant and the journals Pain and "Community Dentistry and Oral Epidemiology" were handsearched.

Study selection Observational studies (cross-sectional surveys, cohorts case-control studies) and previous literature reviews of adult populations over 18 year of age were included. Oro-facial pain (OFP) was defined as pain located in the face, dental pain was excluded.

Data extraction and synthesis A standardised checklist was used to assess the methodological quality of each study by two reviewers before an attempt was made to summarise the results. The median quality score was 70% of the maximum attainable score. Due to methodological issues, it was not possible to pool the data on the prevalence of OFP.

Results 59 studies were evaluated. Age, gender and psychological factors were found to be associated with OFP, however there was not enough information on other factors such as local mechanical and comorbidities to draw any reliable conclusions. None of the factors fully fulfilled criteria for causality (Table 1).

Conclusions There is a need for good quality epidemiological studies of adequate sample size of OFP in the general population. To enable comprehensive examination of the aetiology of OFP, it is necessary to address a broad range of factors, eg demography and life-style, local mechanical factors, medical history and psychological factors. Data on

potential confounders and effect modifiers should also be collected and adjusted for in the statistical analysis.

Commentary

This systematic review had 4 main objectives; to describe and compare performance of several electronic data bases for identification of epidemiological studies on oro-facial pain (OFP); to identify and assess quality of population-based epidemiological studies of OFP; to determine the prevalence of OFP in different adult populations and to describe factors which are associated with OFP.

The review follows an excellent methodology for defining search strategy and initial inclusion criteria. The approach to determining quality is well defined with reported inter-examiner and intra-examiner levels of agreement. It is interesting to note that many of the check list items had low levels of agreement and less than half had good levels of agreement. Moderate agreement ($\kappa = 0.6$) between two observers for classification of papers into subgroups was reported. There was also a wide range in level of agreement between the two examiners for the component items of the quality checklists with more than half of the items having κ values below 0.7. This suggests that the validity of the quality assessment instrument is suspect, or calibration of the examiners was inadequate. Even the level of agreement between observers for classification of the studies by design was relatively low.

The first two objectives of this study were met. The study reported the sensitivity of electronic data bases in searching for epidemiological OFP studies. The quality of existing epidemiological and review articles were evaluated with an objective instrument.

However, the authors do not justify why they chose to include narrative reviews. Narrative reviews are subject to selection bias and therefore should not have been considered.

The final two stated objectives of this systematic review are not met. Unfortunately, although the authors completed a quality assessment on the selected articles, all articles were included in the results and ultimately the discussion. The authors should have planned a cut-off score below which the study would be rejected as being of inadequate quality. It is impossible to draw conclusions regarding prevalence of OFP in different populations and it is impossible to determine the factors associated with OFP from this data, when poor studies were not eliminated. The reader is left to attempt to judge the quality of the individual studies without adequate information on each specific study. Although according to the quality scores, some studies were of good quality, the good studies were not highlighted in this publication. Good evidence may exist for specific issues, but unfortunately the reader does not have the ability to interpret what specific evidence was properly collected and of good quality. There was a substantial time interval between completion of the search (1998) and acceptance for publication (2001) as a result this literature review is somewhat outdated. The abstract is misleading as the objective stated there was to "determine the prevalence and associated risk factors of oro-facial pain", yet the bulk of the article addresses the overall quality of existing research. The result being that this study does not allow conclusions regarding the prevalence or risk factors and is only

Table 1. Percentage of studies reporting factors which are possibly associated with OFP.

	Cross-sectional	Cohort
Age	68	33
Gender	73	67
Socio-economic characteristics	24	50
Education	11	17
Geographic area	11	—
Income/social class	11	17
Race/ethnicity	5	—
Marital status	5	—
Psychosocial factors	11	50
General state of health	11	17
Pain other than OFP	16	17
Headache	24	50
Denture status	27	—
Number of teeth	19	17
Parafunction	14	33
Joint clicking	16	17
Other	16	17

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helpful in providing some general guidance for designing future epidemiological OFP research.

Practice point

- This study does not allow conclusions regarding the prevalence or risk factors.

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Key to evidence graphic used in Evidence-based Dentistry with examples of study designs for Therapy/prevention/aetiology and harm. Study designs for evidence levels will be different for prognosis/diagnosis etc. (See Evidence-based Dentistry 2003;4:p17-18)

Evidence Graphic	Evidence-level	Therapy/Prevention /Aetiology/Harm
 3A 2C 2B 2A 1B 1A	1A	SR (with homogeneity ^a) of RCTs
 3A 2C 2B 2A 1B 1A	1B	Individual RCT with narrow confidence interval (CI)
 3A 2C 2B 2A 1B 1A	2B	SR (with homogeneity ^a) of cohort studies
 3A 2C 2B 2A 1B 1A	2C	Individual cohort studies (including low quality RCT eg < 80% follow-up)
 3A 2C 2B 2A 1B 1A	3A	SR (with homogeneity ^a) of Case-Controlled Studies

^a By homogeneity we mean a systematic review free of worrisome variations (heterogeneity) in the direction and degree of results between individual studies. Not all systematic reviews with statistically significant heterogeneity need be worrisome, and not all worrisome heterogeneity need be statistically significant.