

**A systematic review of the patient reported outcome (PRO) instruments used in clinical trials in head and neck surgery.**

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## **Abstract**

### *Introduction*

There is a growing number of clinical trials in patients with head and neck cancer. Although not often the primary outcome, patient reported outcomes (PROs) are now an important component. The aim of this structured review was to identify and report the characteristics of the questionnaires used and summarise the findings in the literature.

### *Materials and methods*

A search of several online databases was devised using the following key terms: head and neck oncology, head and neck surgery, reconstruction, clinical trials patient-reported outcomes, questionnaires, Quality of Life (QoL), validated instruments, and patient satisfaction. Information was collected relating to the topic of the paper, sample size, selection criteria, the main advantages and disadvantages of the PRO used, and if the tool was used in conjunction with any other.

### *Results*

1342 papers were screened, of which 54 articles eligible; across these papers, 22 questionnaires were identified. The primary reason for utilising a tool was its relevance to the focus of the paper; including features such as xerostomia, pain, swallowing to name a few.

### *Discussion*

We recommend that outcome measures for clinical trials should be chosen in relation to the following criteria: appropriateness, reliability, validity, responsiveness, precision, interpretability, acceptability and feasibility; to allow the patient experience to be the focus of the primary outcome. Clinical trials use validated questionnaires but the PRO measures

tended not to be the focus of the trial. There is merit in future clinical trials having PRO measures as the primary outcome and designed around an explicit hypothesis.

## **Introduction**

Measurement of patient QoL is imperative as UK head and neck cancer incidence is increasing.<sup>1</sup> The associated debilitating physiological and psychological morbidities may thus become more prevalent<sup>2</sup>. Side effects of treatment and functional difficulties can exacerbate emotional distress, depression, and self-esteem issues. The scope of randomised controlled trials in head and neck cancer is treatment; many compare toxicity of different treatment regimens or provide ideal interventions for side effects such as xerostomia. However, there is no universal instrument that is sufficiently robust to compare QoL and morbidity in patients followed up after initial surgical resection, chemotherapy and radiotherapy<sup>3</sup>.

Clinicians find it difficult to determine precisely what physical and emotional trauma affects each patient after initial head and neck cancer management, and to what severity. However, it is these very effects that affect adherence, compliance, morbidity and mortality<sup>4</sup>. It is here that patient-related outcomes become paramount. This review aims to summarise the literature in respect to the PROM used, the focus of the research and the key clinical findings.

## **Materials and methods**

A search strategy was devised using the following key terms: head and neck oncology, head and neck surgery, reconstruction, clinical trials, patient-reported outcomes, questionnaires, QoL, validated instruments, and patient satisfaction. The following databases were examined: Handle-on-qol, Medline, Ebase (Excerpta Medica), HAPI (Health and Psychosocial Instruments), Science Citation Index/Social Sciences Citation Index, Ovid Evidence-Based Medicine databases and PsychINFO.

Only manuscripts written in English were included. All instruments included in the review were identified as PROMs measuring head and neck-related QoL and/or satisfaction that had undergone development and validation with head and neck cancer patients. For the appraisal of the psychometric and operational performance of the instruments we looked for evidence of criteria as in Table 1. PRISMA guidance was considered in the search and presentation of the results<sup>5</sup>. A total of 2072 papers were identified describing QoL (QOL) measures. From an evaluation of the abstracts and available full text, 54 relevant papers were closely examined (Figure 1).

## **Results**

From 54 papers<sup>6-59</sup>, the authors found 22 QoL measures, which satisfied our inclusion criteria (Figure 1). In many studies the authors used more than one instrument. The most common tool used in the search was the EORTC QLQ C30, with its use in 18 of the 54 papers. The second most common was the EORTC QLQ-H&N35 tool which was used in 14 papers. The above two instruments are intended to be utilised together, which was the case in 11 papers.

The primary reason for utilisation of a tool was for its ability to be general or specific when measuring QoL (Table 2). Papers favoured tools that were well-validated, easy to use, and those that had a focus on functional and psychological aspects of QoL. Conversely, disadvantages were deemed to be a lack of relevance to the focus of the paper, low levels of completion, and those that required large numbers for statistical significance. Table 3 summarises the main foci of the clinical trials.

## **Discussion**

In the included papers, PROMs are generally utilised as a secondary outcome to quantify the effect of a treatment modality, rather than as a primary outcome in relation to the patient experience. In undertaking this review, the authors are aware of its limitations: by including papers only written in English, there is exclusion of a potentially large number of studies that could give further insight into how PROMs are utilised. In addition, there were restrictions in obtaining further papers by an inability to access the full text in our searches. The authors recommend that outcome measures for clinical trials should be chosen in relation to the criteria stated in Table 1. Analysis of the papers allows us to determine the key focus of the trial and why each PRO instrument was utilised.

### General Instruments

EQ-5D assesses mobility, self-care, usual activities, pain and anxiety. The instrument was developed for clinical and economic evaluation of healthcare and was designed for use alongside condition-specific instruments<sup>59</sup>. The instrument was used by one study<sup>19</sup>, assessing PET-CT surveillance versus neck dissection in advanced head and neck cancer. EQ-5D was used along with EORTC QLQ-C30, H&N35 and MD Anderson Dysphagia Index (Table 4). Mehanna et al<sup>19</sup> commented on the utilisation of EQ-5D in deriving quality adjusted life years to assess economic viability between the treatment groups.

COOP-WONCA Functional status charts assess general functional status aimed at primary care; the domains include: physical fitness, feelings, daily activities, social activities, change in health and overall health. COOP-WONCA charts were used by Van Bokhorst et al<sup>22</sup> in their trial assessing perioperative enteral nutrition and QoL of severely malnourished patients. The COOP-WONCA charts are not specific for cancer and so were used along with EORTC

QLQ-C30. Van Bokhorst concluded that the COOP-WONCA charts were not sensitive enough to pick up significant changes in QoL<sup>22</sup>.

### Cancer-specific Instruments

EORTC QLQ C30 consists of a general QoL questionnaire composed of functional scales, symptom scales, a global health status and QoL status. The questionnaire was acceptable to patients with 60% completing it in less than 30 minutes<sup>61</sup>. Eighteen randomised control trials used the EORTC QLQ C30, more than any other measure (Table 4). Eleven studies used EORTC QLQ C30 with the QLQ-H&N35 as is intended. The focus of most papers which utilised EORTC QLQ C30 was different chemotherapy medications and their effectiveness.

The Spitzer QoL index covers 5 domains: activity, daily living, health, support and outlook. The instrument was used by two studies in the review, Robert et al<sup>49</sup> were assessing a novel chemotherapy regime and Elliot et al<sup>50</sup> were measuring the effectiveness of a medication in preventing radiation mucositis. It was commented that if one question in the index is not answered the results cannot be interpreted which could be an issue in smaller studies; in addition, the index may be subject to patient and reviewer bias<sup>50</sup>.

The Rotterdam Symptom checklist consists of four main scales: physical symptom distress, psychological distress, activity level, and overall valuation of life. Griffiths et al<sup>51</sup> were the only trial to use this instrument. The paper assessed QoL in patients on the continuous hyper-fractionated accelerated radiotherapy randomised trial, which showed there was no clear difference in QoL compared to conventional radiotherapy. The authors modified the checklist by adding four domains: cough, coughing up blood, hoarseness and restlessness. Griffiths<sup>51</sup> used the RSCL alongside the Hospital Anxiety and Depression Scale.



### Head and Neck Cancer-specific Instruments

The EORTC Head and Neck cancer module (QLQ-H&N35) is a module which is designed to be used in conjunction with the EORTC QLQ C30. This consists of scales including pain, swallowing, senses, speech, social eating, social contact and sexuality. 11 papers used the EORTC QLQ C30 in conjunction with QLQ-H&N35, 7 used EORTC QLQ C30 without the Head and neck module and 3 used the QLQ-H&N35 without the EORTC QLQ C30 (Table 4). The focus the papers was in the comparison of different treatment regimes, particularly chemotherapeutic agents which was the case for 7 of the 11 papers.

The QoL Radiation Therapy Instrument (QoL-RTI) is designed specifically for radiation therapy. The instrument was used by one study in our search which was assessing the effect of a novel radiotherapy regime. The regime showed QoL returned to baseline after 1 month of treatment and it had acceptable toxicity. The instrument was acceptable to patients with 90% completing both pre-treatment and end of treatment questionnaires, but compliance reduced between month 3 and month 12 post treatment<sup>58</sup>.

The University of Washington QoL questionnaire is a head and neck-specific instrument. 9 papers used the UoWQoL (Table 4), two of which used it in conjunction with the Neck Dissection Impairment Index and the University of Michigan Xerostomia Questionnaire. The purpose of most the papers was to assess the effectiveness of different radioprotective regimes during radiotherapy; however, most papers saw that there was still a decrease in QoL despite intervention. The authors of these papers praised the UoWQoL as a general health measure but commented that it was insensitive to changes regarding xerostomia. Owen et al<sup>34</sup> found a lack of compliance to completion.

The ROTG modified University of Washington Head and Neck Symptom Questionnaire is essentially the same as the UoWQoL but focuses more on the effects of radiation in the head and neck<sup>38</sup>. It is used by three studies assessing the symptoms of patients undergoing radiotherapy who found that there is a negative change in QoL scores following treatment, particularly in relation to mucositis and xerostomia<sup>38,39,40</sup> (Table 4).

The University of Michigan Xerostomia-related QoL scale is specific to mouth and throat dryness. Four of the papers included used this instrument, all of which measured QoL in different radiotherapy techniques; those with parotid-sparing had an increase in QoL compared to other regimes (Table 4).

The Neck Dissection impairment index (NDII) is designed to assess function, particularly related to the shoulder, following neck dissection. This instrument was used in two studies to assess the effects of different treatments, including exercises and TENS, on QoL. There was found to be no statistical significance between treatments in either study. Both studies used the index along with more general head and neck indices for an overall view of QoL. The NDII was praised for its simplicity and specificity to neck dissection related QoL (Table 4).

The Performance Status Scale for Head and Neck Cancer is a clinician-rated instrument. The three domains are understandable speech, normalcy of diet and eating in public. Three separate studies used the PSS-HN, one in conjunction with H&N35 and EORTC QLQ-C30 and one in conjunction with the MD Anderson Dysphagia inventory (Table 4). The studies showed that chemoradiotherapy causes a deterioration in QoL but that there is no difference between different chemoradiotherapy regimes.

Functional Assessment of Cancer Therapy-Head and Neck is a self-reported instrument; the domains are physical, social, emotional, and functional well-being. Five studies in this review used FACT-HN, reporting it useful for functional assessment; the papers evaluated the difference in efficacy and QoL between different chemoradiotherapy regimes. There was found to be no difference in QoL between treatment and control groups in all the studies. Simon et al<sup>38</sup> reported low completion rates (60%) in their study comparing Gefitinib with methotrexate. Most studies reported the FACT-HN's utility in measuring functional performance for patients with head and neck cancer. (Table 4)

The FHNSI-10 is aimed at patients with refractory, recurrent or advanced disease. It is a subset of the FACT H&N and was designed to capture physical symptoms of disease. Two studies used the FHNSI-10; Stewart et al<sup>43</sup> utilised it in conjunction with the Functional Assessment of Cancer Therapy H&N instrument, as they reported that FHNSI-10 is ineffective for assessing general QoL. Again, the studies were assessing different chemotherapy regimens; no mention was made of patient acceptability<sup>70</sup>.

The MD Anderson Dysphagia Inventory (MDADI) is a self-administered instrument for assessing dysphagia; it includes items grouped into domains of dysphagia; global, emotional, functional and physical. The inventory was used by two studies in our review both of which praised it for its specific use with swallowing, but also used a more general measure alongside (Table 4). Hutcheson et al<sup>45</sup> used the study to assess QoL following chemoradiotherapy which was found to decrease.

The Head and Neck Radiotherapy Questionnaire is directed towards physical symptoms of skin, throat, stomatitis, digestion, energy and psychosocial for head and neck radiotherapy patients. All four of the studies that used this instrument were related to the prevention of xerostomia in patients undergoing radiotherapy; most radioprotective agents did not have a significant effect on QoL. Ringash et al complimented the disease specificity and ease of completion, however they also comment that data from healthy individuals is not available and while the score can remain the same the patients may have swapped pre-treatment problems with post-treatment problems<sup>73</sup>.

HNQoL is a validated QoL instrument divided into four domains: eating and swallowing, communication, head and neck pain and emotional wellbeing. The only study to utilise this instrument identified QoL between patients who underwent chemoradiotherapy plus a neck dissection versus chemotherapy alone; no difference in QoL was found. Donatelli-Lassig et al<sup>59</sup> determined that the disease specificity of the HNQoL was a useful feature.

Head and Neck Cancer inventory (HNCI) is a reliable validated health status instrument. Lazarus et al<sup>55</sup> utilised this study in assessing swallowing and tongue strength exercises in patients who underwent primary chemotherapy with or without adjuvant chemotherapy. Those who underwent exercises had an increase in QoL.

### Miscellaneous Instruments

The Hospital Anxiety and Depression Scale is an instrument with two domains (anxiety and depression)<sup>75</sup>. Four studies in the sample used the HADS which focussed on whether implementation of coping strategies improved QoL, which was found to be the case. All the

studies commented that it was a useful psychological test, but Scheifke et al<sup>9</sup> commented the need to use another instrument for a more rounded assessment.

The Dermatology life quality index is a dermatology-specific QoL instrument<sup>76</sup>. It was utilised by one trial in the search which was based on skin symptoms associated with etuximab. The trial also used the EORTC QLQ-C30 (Table 4).

The Hearing Handicap Inventory for the Elderly is a specific instrument for assessing QoL issues due to hearing loss. The instrument is valid, reliable and easy to use<sup>77</sup>. Schultz et al<sup>57</sup> were the only study to use this instrument, utilising it for hearing loss associated with head and neck cancer, showing that patients undergoing radiotherapy are more likely to have hearing related problems.

The Modified WHO performance status scale is a five-point scale based on ability to work and is not a true QoL instrument<sup>78</sup>; it was utilised by Correy et al<sup>56</sup> in the assessment of PEG vs NG tubes, where there was found to be no significant difference in QoL.

## **Conclusion**

There have been a variety of questionnaires used in clinical trials following H&N cancer; these have tended to be secondary outcomes. It is important when focusing on patient reported outcomes to include a validated questionnaire that is optimal to the hypothesis being tested between arms of the trial. It is worthwhile considering more than one questionnaire and to be as specific as possible in selection; in addition, anchoring a PROM with an objective measurement will be beneficial in ensuring the patient experience as the primary outcome. Although underlying issues have been widely discussed, three of our criteria:

appropriateness, precision and interpretability, are not always included in lists of desirable properties of instruments. The remaining five criteria are widely cited and identified in the same or similar terminology as in this review (Table 1).

In this review, the main areas where PROs were used in clinical trials were for evaluating the differences between different chemotherapy medications, the differences in techniques for preventing xerostomia, and between different radiotherapy regimes, amongst others. For the main, there was little difference in QoL between different treatment regimens; those papers that focussed on specific patient experience measures such as implementation of coping strategies, found an increase in QoL. In the future, trials can be broadened to include PROs as the primary outcome with an explicit hypothesis.

**Conflict of interest:** The authors have no conflict of interest to declare.

## References

1. Rogers SN, Semple C, Babb M, Humphris G. QoL considerations in head and neck cancer: United Kingdom National Multidisciplinary Guidelines. *J Laryngol Otol*. 2016 May;**130**(S2):S49-S52.
2. Rogers SN, Heseltine N, Flexen J, Winstanley HR, Cole-Hawkins H, Kanatas A. Structured review of papers reporting specific functions in patients with cancer of the head and neck: 2006 - 2013. *Br J Oral Maxillofac Surg*. 2016 Jul;**54**(6):e45-51.
3. Kamisetty A, Magennis P, Mayland C, Jack B, Lowe D, Rogers SN. Where do patients treated for oral cancer die? A 20-year cohort study 1992-2011. *Br J Oral Maxillofac Surg*. 2015 Dec;**53**(10):1015-20.

4. Kanatas A, Singh P, Lowe D, Rogers SN. How will I be after my operation for oral cancer? *Br J Oral Maxillofac Surg.* 2015 Jul;**53**(6):538-45.
5. Moher D, Liberati A, Tetzlaff J, Altman DG, The PRISMA Group (2009). Preferred reporting items for systematic reviews and meta-analyses: The PRISMA statement. *Plos Med* **6**(7):e1000097.
6. Duncan G et al. QoL, mucositis, and xerostomia from radiotherapy for head and neck cancers: a report from the NCIC CTG HN2 randomised trial of an antimicrobial lozenge to prevent mucositis. *Head Neck* 2005; **27**:421–8.
7. Rivera F et al. Cetuximab in metastatic or recurrent head and neck cancer: the EXTREME trial. *Expert Rev. Anticancer Ther.* 2009; **9**:1421-8.
8. Machiels J et al. Afatinib versus methotrexate as second-line treatment in patients with recurrent or metastatic squamous-cell carcinoma of the head and neck progressing on or after platinum-based therapy (LUX-Head & Neck 1): an open-label, randomised phase 3 trial. *Lancet Oncol* 2015; **16**:583-94.
9. Potthoff K et al. Randomized controlled trial to evaluate the effects of ethyl-2-cyanoacrylate on pain intensity and QoL in head and neck cancer patients suffering from cetuximab-induced rhagades during radioimmunotherapy: the support trial. *BMC Cancer* 2014; **14**:270.
10. Bottomley A et al. An International Phase 3 Trial in Head and Neck Cancer: QoL and Symptom Results EORTC 24954 on Behalf of the EORTC Head and Neck and the EORTC Radiation Oncology Group. *Cancer* 2014; **120**:390-8.
11. Van Herpen CML et al. Short-term health-related QoL and symptom control with docetaxel, cisplatin, 5-fluorouracil and cisplatin (TPF), 5-fluorouracil (PF) for induction in unresectable locoregionally advanced head and neck cancer patients (EORTC 24971/TAX 323). *British Journal of Cancer* 2010; **103**:1173-81.

12. Machiels J et al. Rationale and design of LUX-Head & Neck 1: a randomised, Phase III trial of afatinib versus methotrexate in patients with recurrent and/or metastatic head and neck squamous cell carcinoma who progressed after platinum-based therapy. *BMC Cancer* 2014; **14**:473.
13. Mesía R et al. QoL of patients receiving platinum-based chemotherapy plus cetuximab first line for recurrent and/or metastatic squamous cell carcinoma of the head and neck. *Annals of Oncology* 2010; **21**:1967–73.
14. Schiefke F et al. Function, post-operative morbidity, and QoL after cervical sentinel node biopsy and after selective neck dissection. *Head Neck* 2009; **31**:503-12.
15. Ackerstaff A et al. First-year QoL assessment of an intra-arterial (RADPLAT) versus intravenous chemoradiation phase III trial. *Head Neck* 2009; **31**:77–84.
16. Vilela L et al. Comparison of Psychosocial Outcomes in Head and Neck Cancer Patients Receiving a Coping Strategies intervention and Control Subjects Receiving No Intervention. *The Journal of Otolaryngology* 2006; **35**:88-96.
17. Heukelom J et al. Adaptive and innovative Radiation Treatment FOR improving Cancer treatment outcomE (ARTFORCE); a randomized controlled phase II trial for individualized treatment of head and neck cancer. *BMC Cancer* 2013; **13**:84.
18. Uster A et al. Influence of a nutritional intervention on dietary intake and QoL in cancer patients: A randomized controlled trial. *Nutrition* 2013; **29**:1342-9.
19. Mehanna H et al. PET-CT Surveillance versus Neck Dissection in Advanced Head and Neck Cancer. *Engl J Med* 2016; **15**:374.
20. Teguh D et al. Early hyperbaric oxygen therapy for reducing radiotherapy side effects: early results of a randomised trial in oropharyngeal and nasopharyngeal cancer. *Int. J. Radiation Oncology Biol. Phys.* 2009; **75**:711-6.



21. Fang F et al. QoL and survival outcome for patients with nasopharyngeal carcinoma receiving three-dimensional conformal radiotherapy vs intensity-modulated radiotherapy-a longitudinal study. *Int. J. Radiation Oncology Biol. Phys.* 2008; **72**:356-64.
22. Van Bokorst-De Van Der Schueren MAE et al. Perioperative enteral nutrition and QoL of severely malnourished head and neck cancer patients: a randomized clinical trial. *Clinical Nutrition* 2000; **19**: 437-44.
23. Myers E et al. QoL effects of psychosocial intervention in patients with head and neck cancer. *Otolaryngology* 1999; **120**: 507-16.
24. Van Rij CM et al. Parotid gland sparing IMRT for head and neck cancer improves xerostomia related QoL. *Radiation Oncology* 2008; **3**:41.
25. Bower W et al. Mode of treatment affects QoL in head and neck cancer survivors: Implications for holistic care. *Acta Oto-Laryngologica* 2010; **130**:1185-92.
26. Chang T et al. Xerostomia in Long-term Survivors of Aggressive Non-Hodgkin's Lymphoma of Waldeyer's Ring A Potential Role for Parotid-Sparing Techniques? *Am J Clin Oncol* 2009; **32**: 145-9.
27. Scrimger R et al. Correlation Between Saliva Production and QoL Measurements in Head and Neck Cancer Patients Treated With Intensity-Modulated Radiotherapy. *Am J Clin Oncol* 2007; **30**:271-7.
28. Warde P et al. A phase II study of Biotene in the treatment of postradiation xerostomia in patients with head and neck cancer. *Support Care Cancer* 2000; **8**:203-8.
29. Lin A et al. QoL after parotid-sparing IMRT for head and neck cancer: a prospective longitudinal study. *Int. J. Radiation Oncology Biol. Phys.* 2003; **57**:61-70
30. Parikh S et al. A double blind randomised trial of IIB or not IIB neck dissections on

- electromyography, clinical examination, and questionnaire-based outcomes: a feasibility study. *British Journal of Oral and Maxillofacial Surgery* 2012; 50: 394–403.
31. Lydiatt W et al. A Randomized, Placebo-Controlled Trial of Citalopram for the Prevention of Major Depression During Treatment for Head and Neck Cancer. *Arch Otolaryngol Head Neck Surg.* 2008; **134**:528-35.
32. Brennan S et al. Prospective trial to evaluate staged neck dissection or elective neck radiotherapy in patients with CT-staged T1-2 N0 squamous cell carcinoma of the oral tongue. *Head Neck* 2009; **32**:191-8.
33. Oton-Leite AF et al. Effect of intraoral low-level laser therapy on QoL of patients with head and neck cancer undergoing radiotherapy. *Head Neck* 2012; 398-404.
34. Owen P et al. Radiofrequency Ablation of Advanced Head and Neck Cancer. *Arch Otolaryngol Head Neck Surg.* 2011;**137**:493-8.
35. Jha N et al. Submandibular salivary gland transfer prevents radiation-induced xerostomia. *Int. J. Radiation Oncology Biol. Phys.* 2000; **46**:7-11.
36. Johnson D et al. Assessment of QoL and oral function of patients participating in a phase II study of radioprotection of oral and pharyngeal mucosa by the prostaglandin E1 analog misoprostol (RTOG 96-07). *Int. J. Radiation Oncology Biol. Phys.* 2002; **54**:1455-9.
37. Jha N et al. Prevention of radiation induced xerostomia by surgical transfer of submandibular salivary gland into the submental space. *Radiotherapy and Oncology* 2003; **66**: 283–2.
38. Hoffman K et al. The impact of concurrent granulocyte–macrophage colony stimulating factor on QoL in head and neck cancer patients: results of the randomized, placebo controlled Radiation. Therapy Oncology Group 9901 trial. *Qual Life Res*

- 2014; **23**:1841-58.
39. Heron D et al. Stereotactic body radiotherapy for recurrent squamous cell carcinoma of the head and neck: results of a phase I dose-escalation trial. *Int. J. Radiation Oncology Biol. Phys.* 2009; **75**:1493-500.
40. Fisher J et al. Phase III QoL study results: impact on patients' QoL to reducing xerostomia after radiotherapy for head and neck cancer – RTOG 97-09. *Int. J. Radiation Oncology Biol. Phys.* 2003; **56**:832-6.
41. McNeely M et al. Effect of Exercise on Upper Extremity Pain and Dysfunction in Head and Neck Cancer Survivors A Randomized Controlled Trial. *Cancer* 2008; **113**:214–22.
42. Kushwaha V et al. Gefitinib, Methotrexate and Methotrexate plus 5-Fluorouracil as palliative treatment in recurrent head and neck squamous cell carcinoma. *Cancer Biology & Therapy* 2015; **16**:346-51.
43. Stewart S et al. Phase III Study of Gefitinib 250 Compared With Intravenous Methotrexate for Recurrent Squamous Cell Carcinoma of the Head and Neck. *J Clin Oncol* 2009; **27**:1864-71.
44. Mittal B et al. Effect of induction chemotherapy on swallow physiology and saliva production in patients with head and neck cancer: A pilot study. *Head Neck* 2015; **37**: 567-72.
45. Hutcheson K et al. Long-term functional and survival outcomes after induction chemotherapy and risk-based definitive therapy for locally advanced squamous cell carcinoma of the head and neck. *Head Neck* 2014; **36**: 474-80.
46. Rischin D et al. Tirapazamine, Cisplatin, and Radiation Versus Cisplatin and Radiation for Advanced Squamous Cell Carcinoma of the Head and Neck (TROG 02.02, HeadSTART): A Phase III Trial of the Trans-Tasman Radiation Oncology

- Group. *J Clin Oncol* 2010; **28**:2989-95.
47. Cohen E et al. High Survival and Organ Function Rates After Primary Chemoradiotherapy for Intermediate-Stage Squamous Cell Carcinoma of the Head and Neck Treated in a Multicenter Phase II Trial. *J Clin Oncol* 2006; **24**:3438-44.
48. Ringash J et al. Hyperfractionated, accelerated radiotherapy for locally advanced head and neck cancer: QoL in a prospective phase I/II trial. *Radiotherapy and Oncology* 2008; **87**: 181-2.
49. Robert F et al. A Phase II Study of Topotecan in Patients with Recurrent Head and Neck Cancer: Identification of an Active New Agent. *American Journal of Clinical Oncology* 1997; **20**:298-302.
50. Elliot E et al. Phase III Trial of an Emulsion Containing Trolamine for the Prevention of Radiation Dermatitis in Patients With Advanced Squamous Cell Carcinoma of the Head and Neck: Results of Radiation Therapy Oncology Group Trial 99-13. *J Clin Oncol* 2006; **24**:2092-7.
51. Griffiths GO et al. Physical and psychological symptoms of QoL in the CHART randomized trial in head and neck cancer: short-term and long-term patient reported symptoms. *Journal of Cancer* 1999; **81**:1196–1205.
52. Ringash J et al. Postradiotherapy QoL for head and neck cancer patients is independent of xerostomia. *Int. J. Radiation Oncology Biol. Phys.* 2005; **61**:1403-7.
53. Wong R et al. A phase I-II study in the use of acupuncture-like transcutaneous nerve stimulation in the treatment of radiation-induced xerostomia in head and neck cancer patients treated with radical radiotherapy. *Int. J. Radiation Oncology Biol. Phys.* 2003; **57**:472-80.
54. Warde P et al. A phase III placebo-controlled trial of oral pilocarpine in patients undergoing radiotherapy for head and neck cancer. *Int. J. Radiation Oncology Biol.*

- Phys.* 2002; **54**; 9-13.
55. Lazarus C et al. Effects of exercise on swallowing and tongue strength in patients with oral and oropharyngeal cancer treated with primary radiotherapy with or without chemotherapy. *International Journal of Oral and Maxillofacial Surgery* 2014; **43**:523-30.
56. Corry J et al. Randomized study of percutaneous endoscopic gastrostomy versus nasogastric tubes for enteral feeding in head and neck cancer patients treated with (chemo)radiation. *Journal of Medical Imaging and Radiation Oncology* 2008; **52**:503–10.
57. Schultz C et al. Hearing Loss and Complaint in Patients With Head and Neck Cancer Treated With Radiotherapy. *Arch Otolaryngol Head Neck Surg.* 2010; **136**:1065-9.
58. Maguire P et al. Phase II trial of hyperfractionated intensity-modulated radiation therapy and concurrent weekly cisplatin for stage III and IVa head and neck cancer. *Int. J. Radiation Oncology Biol. Phys.* 2011; **79**:1081-8.
59. Donatelli-Lassig A et al. The effect of neck dissection on QoL after chemoradiation. *Otolaryngology Head and Neck Surgery* 2008; 13.
60. Nelson EC, Landgraf JM, Hays RD, Wasson JH, Kirk JW. The functional status of patients. How can it be measured in physicians' offices? *Med Care.* 1990;28(12):1111–26.
61. Kristin Bjordal & Stein Kaasa (1992) Psychometric Validation of the EORTC Core QoL Questionnaire, 30-Item Version and A Diagnosis-Specific Module for Head and Neck Cancer Patients, *Acta Oncologica*, 31:3, 311-320, DOI: 10.3109/02841869209108178.
62. Spitzer, W., Dobson, A., Hall, J. (1981). Measuring the QoL of cancer patients: a concise QL-Index for use by physicians. *Journal of Chronic Diseases*, 34: 585-597.

63. Haes JD, Knippenberg FV, Neijt J. Measuring psychological and physical distress in cancer patients: structure and application of the Rotterdam Symptom Checklist. *British Journal of Cancer* 1990;62:1034–8. doi:10.1038/bjc.1990.434.
64. Bjordal K, Hammerlid E, Ahlner-Elmqvist M, de Graeff A, Boysen M, Evensen JF, Bjorklund A, de Leeuw JR, Fayers PM, Jannert M, et al. QoL in head and neck cancer patients: validation of the European Organization for Research and Treatment of Cancer QoL Questionnaire-H&N35. *J Clin Oncol.* 1999;17:1008–1019.
65. Johnson DJ, Casey L, Noriega B. A pilot study of patient QoL during radiation therapy treatment. *Qual Life Res.* 1994;3(4):267–272.
66. Hassan S J, Weymuller EA. Assessment of QoL in head and neck cancer patients. *Head Neck* 1993; 15: 485-496.
67. Taylor RJ, Chepeha JC, Teknos TN, et al. Development and validation of the neck dissection impairment index. *Arch Otolaryngol Head Neck Surg* 2002; 128:44–49.
68. List MA, Ritter-Sterr C, Lansky SB. A performance status scale for head and neck cancer patients. *Cancer* 1990; 97:25-32. 66:564-9.
69. M.A. List, L.L. D'Antonio, D.F. Cella, et al. The Performance Status Scale for Head and Neck Cancer Patients and the Functional Assessment of Cancer Therapy-Head and Neck Scale: A study of utility and validity *Cancer*, 77 (1996), pp. 2294–2301.
70. Yount S, List M, Du H, et al. A randomized validation study comparing embedded versus extracted FACT Head and Neck Symptom Index scores. *Qual Life Res.* 2007;16:1615–1626.
71. Chen AY, Frankowski R, Bishop-Leone J, et al. The development and validation of a dysphagia-specific quality-of-life questionnaire for patients with head and neck cancer: the M.D. Anderson dysphagia inventory. *Arch Otolaryngol Head Neck Surg.* 2001;127(7):870–876.

72. G.P. Browman, M.N. Levine, D.I. Hodson, et al The Head and Neck Radiotherapy Questionnaire: A morbidity/quality-of-life instrument for clinical trials of radiation therapy in locally advanced head and neck cancer *J Clin Oncol*, 11 (1993), pp. 863–872.
73. Ringash J et al. Postradiotherapy QoL for head and neck cancer patients is independent of xerostomia. *Int. J. Radiation Oncology Biol. Phys.* 2005; 61:1403-7.
74. Terrell JE, Nanavati KA, Esclamado RM, et al. Head and neck cancer-specific QoL: Instrument validation. *Arch Otolaryngol Head Neck Surg.* 1997;123:1125–32.
75. A.S. Zigmond, R.P. Snaith The Hospital Anxiety and Depression Scale *Acta Psychiatr Scand*, 67 (1983), pp. 361–370.
76. Finlay AY, Khan GK. Dermatology Life Quality Index (DLQI) – a simple practical measure for routine clinical use. *Clin Exp Dermatol.* 1994;19(3):210–216.
77. Ventry IM, Weinstein BE. The Hearing Handicap Inventory for the Elderly: A new tool. *Ear Hear.* 1982;3:128–134.
78. Oken MM, Creech RH, Tormey DC, et al. (1982). "Toxicity and response criteria of the Eastern Cooperative Oncology Group". *Am. J. Clin. Oncol.* 5 (6): 649–55.

Validity
Reliability
Responsiveness
Appropriateness
Precision
Interpretability
Acceptability
Feasibility

Table 1. Criteria required to be evidenced by the instruments

<b>Type of PRO instrument</b>	<b>Number of instruments</b>	<b>Names of instruments</b>
<b>General</b>	2	EQ-5D; COOP-WONCA
<b>Cancer Specific</b>	3	EORTC QLQ C30; Spitzer QoL; Rotterdam Symptom Checklist
<b>Head &amp; Neck Cancer Specific</b>	13	EORTC QLQ-H&N35; University of Washington QoL; ROTG Modified University of Washington QoL; University of Michigan Xerostomia Questionnaire; QoL-RTI; NDII; PSS H&N; MD Anderson Dysphagia Inventory; FACT H&N, FHNSI-10; H&N QoL; HNCI; H&N Radiotherapy Questionnaire



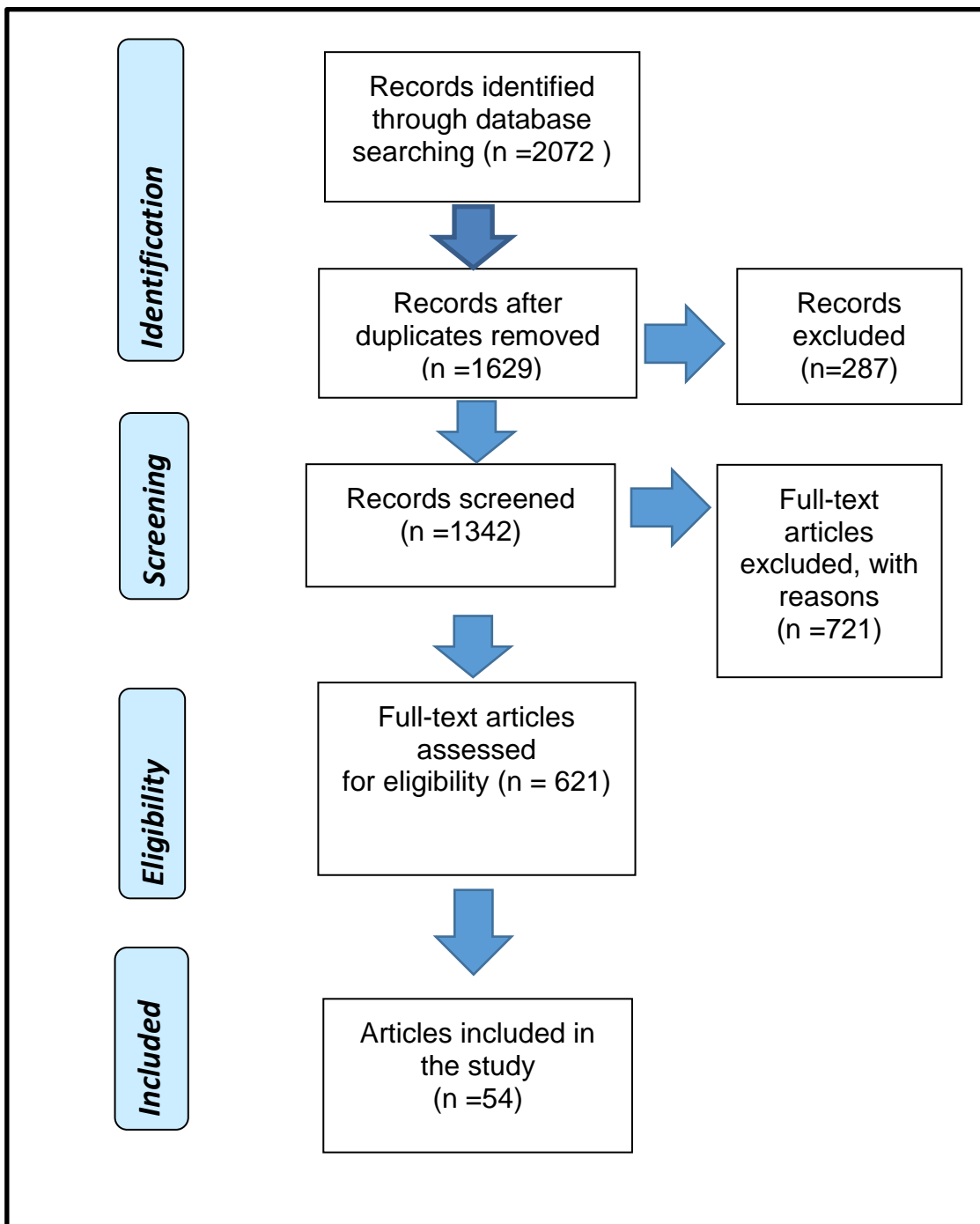
<b>Miscellaneous</b>	4	Hospital Anxiety & Depression Scale; Hearing Handicap Inventory; Dermatology Life Questionnaire; Modified WHO PS
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Table 2. Types of Patient Reported Outcome (PRO) instrument used

<b>Focus of the Trial</b>	<b>Number of Trials with this Focus</b>
Chemotherapy/ Chemoradiotherapy regimens	13
Xerostomia/ Mucositis	12
Radiotherapy regimens	11
Dietary intake	3
Psychosocial interventions	2
Pain medications	2
QoL	2
Coping strategies	2
PET CT vs SND for monitoring	1
Hyperbaric oxygen therapy	1
Utilising EMG studies	1
Shoulder exercises	1
Dermatitis	1
Hearing Loss	1
Swallowing	1

Table 3. Focus of clinical trials

Figure 1 Search results included in the review



Questionnaire	First Author/ Year	Focus of the RCT	Number of patients	Inclusion & Exclusion Criteria	Main Advantages	Main Disadvantages	Other Instruments
EORTC QLQ-C30	Duncan et al 2005 <sup>6</sup>	Xerostomia/Mucositis	138	Non-metastatic; had RT	Broad view of QoL Brief Validated	Not specific to oral cavity as needed Not H&N specific	
	Rivera et al 2009 <sup>7</sup>	Chemotherapy in metastatic disease	442	Recurrent or metastatic cancer	Cancer-specific Self-administered Multi-dimensional		HN35
	Machiels et al 2015 <sup>8</sup>	Chemotherapy in metastatic disease	483	Recurrent or metastatic cancer	Cancer-specific Validated		HN35
	Potthoff et al 2014 <sup>9</sup>	Medication for pain intensity	34	Cetuximab chemoradiotherapy rhagades	Cancer-specific Broad view of health Pt recorded		Dermatological Life Quality Index
	Bottomley et al 2013 <sup>10</sup>	Chemotherapy vs chemoradiotherapy	450	Neck nodes but no metastases	Broad view of health		H&N35
	Van Herpen et al 2010 <sup>11</sup>	Symptom control	358	Unresectable, advanced SCC	Robust Validated Frequently used in RCTs Functional & symptomatic scales	Not H&N specific	H&N35
	Machiels et al 2014 <sup>12</sup>	Chemotherapy in metastatic disease	474	Recurrent SCC not amenable to salvage	Good scale for measuring pain		H&N35
	Mesia et al 2010 <sup>13</sup>	Chemotherapy medication	442	Previously untreated advanced SCC	Cancer-specific Broad view of health	Lack of compliance to completion	H&N35
	Schiefke et al 2008 <sup>14</sup>	QoL after SND	49	Sentinel node bx or SND	Validated Reliable	May need large numbers for statistical significance	H&N35 Hospital Anxiety and Depression Scale

	Ackerstaff et al 2008 <sup>15</sup>	Chemotherapy medication	207	Ineligible for salvage surgery	Validated		H&N35
	Vilela et al 2005 <sup>16</sup>	Coping strategy therapy	101	Completed cancer treatment	Validated		Hospital Anxiety and Depression Scale
	Heukelom et al 2013 <sup>17</sup>	Chemoradiotherapy	268	T3/4 tumour new diagnosis	Validated		H&N35
	Uster et al 2013 <sup>18</sup>	Dietary intake post H&N cancer	58	Those who would benefit from nutritional support	Cancer-specific	Not specific to nutrition	
	Mehanna et al 2016 <sup>19</sup>	PET-CT in advanced SCC	564	N2 or N3 metastases	Validated Frequently used	Not H&N specific	EQ-5D Hi 35 MD Anderson dysphagia
	Teguh et al 2009 <sup>20</sup>	Hyperbaric oxygen	19	RT treatment tongue SCC	Validated		Performance Status Scale for Head and Neck Cancer H&N35
	Fang et al 2008 <sup>21</sup>	Different RT rxs	203	Requiring radical RT	Validated		H&N35
	Van bokhorst et al 2000 <sup>22</sup>	Enteral nutrition	49	Malnourished	Cancer-specific Validated		COOP-WONCA
	Myers et al 1999 <sup>23</sup>	Psychosocial intervention	47	H&N SCC, no previous mental health issue	Cancer-specific		Hospital Anxiety and Depression Scale
EORTC QLQ-H&N35	Rivera et al 2009 <sup>7</sup>	Chemotherapy in metastatic disease	442	Recurrent or metastatic cancer	H&N cancer-specific	Low completion May not be relevant for general QoL factors	EORTC QLQ-C30
	Machiels et al 2015 <sup>8</sup>	Chemotherapy in metastatic disease	483	Recurrent or metastatic cancer	Validated		EORTC QLQ-C30
	Bottomley et al 2013 <sup>10</sup>	Chemotherapy vs chemoradiotherapy	450	Neck nodes but no metastases	Designed for surgery, radiotherapy and chemoradiotherapy		EORTC QLQ-C30
	Van Herpen et al 2010 <sup>11</sup>	Symptom control	358	Unresectable, advanced SCC	Specific to H&N i.e. xerostomia etc, Good previous use in cancer clinical trials		EORTC QLQ-C30

	Machiels et al 2014 <sup>12</sup>	Chemotherapy in metastatic disease	474	Recurrent SCC not amenable to salvage	H&N specific		EORTC QLQ-C30
	Mesia et al 2010 <sup>13</sup>	Chemotherapy medication	442	Previously untreated advanced SCC	H&N specific		EORTC QLQ-C30
	Van Rij et al 2008 <sup>24</sup>	Xerostomia related to RT	192	For curative RT	Good for xerostomia		
	Bower et al 2009 <sup>25</sup>	Effects of treatment modalities for H&N	231	Any form of curative Rx	Validated H&N specific		
	Schiefke et al 2008 <sup>14</sup>	QoL after SND	49	Sentinel node bx or SND	Validated Reliable H&N specific	May need large numbers for statistical significance	EORTC QLQ-C30 Hospital Anxiety and Depression Scale
	Ackerstaff et al 2008 <sup>15</sup>	Chemotherapy medications	207	Ineligible for salvage surgery	Validated		EORTC QLQ-C30
	Heukelom et al 2013 <sup>17</sup>	Chemoradiotherapy	268	T3/4 tumour new diagnosis	Validated		EORTC QLQ-C30
	Mehanna et al 2016 <sup>19</sup>	PET-CT in advanced SCC	564	N2 or N3 metastases	H&N specific		MD Anderson Dysphagia Inventory
	Teguh et al 2009 <sup>20</sup>	Hyperbaric oxygen	19	RT treatment tongue SCC	H&N specific i.e swallowing		Performance Status Scale for Head and Neck Cancer EORTC QLQ-C30
	Fang et al 2008 <sup>21</sup>	Different RT regimes	203	Requiring radical RT	Validated H&N specific		EORTC QLQ-C30
University of Michigan Xerostomia Questionnaire	Chang et al 2009 <sup>26</sup>	Xerostomia	15	Disease free 1yr post-surgery; had RT	Simple Validated	Not suitable for less invasive RT treatments w/ lower doses	
	Scrimger et al 2007 <sup>27</sup>	Saliva Production	188	Bilateral RT to parotids	Specific to clinical question		UoWQoL
	Warde et al 2000 <sup>28</sup>	Xerostomia	28	RT; no anticholinergic medications	Simple		

	Lin et al 2003 <sup>29</sup>	Xerostomia	36	Post-parotid sparing RT	H&N specific Validated Specific to xerostomia	May not be as useful when sample size small	
University of Washington QoL	Scrimger et al 2007 <sup>27</sup>	Saliva Production	188	Bilateral RT to parotids	Well-rounded general health view	Insensitivity when related to question of xerostomia	University of Michigan Xerostomia Questionnaire
	Parikh et al 2011 <sup>30</sup>	Electromyographic studies	38	Selective neck dissection	Good for functional measures		Neck Dissection Impairment Index
	Lydiatt et al 2008 <sup>31</sup>	Depression medication	23	No pre-existing mental health condition	Self-administered Focuses on aspects of daily life		
	Brennan et al 2009 <sup>32</sup>	Staged vs elective neck treatment	25	T1-2 N0 SCC new diagnosis	General health view		
	Oton-Leite et al 2011 <sup>33</sup>	Laser therapy with RT	60	Undergoing RT salivary glands	Simple Brief Validated Easy to complete and interpret H&N cancer-specific		
	Owen et al 2011 <sup>34</sup>	Radio frequency ablation	21	Unresectable SCC or previously failed Rx	H&N specific	Lack of compliance to completion	
	Jha et al 2000 <sup>35</sup>	RT induced xerostomia	16	Eligible for RT	General health view	Not specific to xerostomia or RT	
	Johnson et al 2002 <sup>36</sup>	Radioprotection of mucosa	33	Resection + RT	Good for functional measures	heavily weighted on patient symptoms	
	Jha et al 2003 <sup>37</sup>	Xerostomia	76	Requiring RT	Good for functional measures		
RTOG-modified University of Washington QoL H&N Symptom	Hoffman et al 2014 <sup>38</sup>	GM-CSF effect of RT symptoms	114	No previous chemoradiotherapy	Self-administered Validated Specific to RT	Lack of compliance to completion	

	Heron et al 2009 <sup>39</sup>	Stereotactic body radiotherapy	25	Recurrent SCC	General health but with specific RT aspect		
	Fisher et al 2003 <sup>40</sup>	Xerostomia post RT	249	RT	Function specific Good for RT		
Neck Dissection Impairment Index	Parikh et al 2011 <sup>30</sup>	Electromyographic studies	38	Selective neck dissection	Simple		University of Washington QoL
	McNeely et al 2008 <sup>41</sup>	Exercise for shoulder pain	52	Neck dissection, shoulder dysfunction	Specific for neck dissection issues		Functional Assessment of Cancer Therapy H&N
FHNSI-10	Kushwaha et al 2015 <sup>42</sup>	Palliative chemotherapy medications	117	Pt ineligible for salvage surgery/RT/chemo	Specific for advanced/recurrent cancer Brevity		
	Stewart et al 2009 <sup>43</sup>	Methotrexate recurrent SCC	486	Recurrent SCC with RT not amenable to salvage	Good for symptomatic measures	Not useful for general health	Functional Assessment of Cancer Therapy H&N
Performance Status Scale for Head and Neck Cancer	Mittal et al 2015 <sup>44</sup>	Swallowing/ Saliva Production	13	Chemoradiotherapy	H&N cancer-specific		
	Hutcheson et al 2014 <sup>45</sup>	Chemotherapy in relation to swallowing	47	Untreated stage IV SCC	Disease specific, Involves another person i.e. the clinician	Involves another person i.e. the clinician	MD Anderson Dysphagia Inventory
	Teguh et al 2009 <sup>20</sup>	Hyperbaric oxygen	19	RT treatment tongue SCC	Good for functional activities such as swallowing		H&N35 EORTC QLQ-C30
Functional Assessment of Cancer Therapy H&N	Rischin et al 2010 <sup>46</sup>	Chemoradiotherapy effectiveness	850	Previously untreated advanced SCC	H&N cancer-specific Good for chemo/radiotherapy Rx options		
	Simon et al 2009 <sup>43</sup>	Methotrexate recurrent SCC	486	Recurrent SCC with RT not amenable to salvage	H&N cancer-specific	Low completion	FHNSI-10

	Cohen et al 2006 <sup>47</sup>	Chemoradiotherapy	53	Stage II or III SCC	Good focus on chemo and radiotherapy options		
	McNeely et al 2008 <sup>41</sup>	Exercise for shoulder pain	52	Neck dissection, shoulder dysfunction	Good for function & exercise		Neck Dissection Impairment Index
	Ringash et al 2008 <sup>48</sup>	RT	171	Locally advanced SCC III/IV	Good for function Good for RT		
MD Anderson Dysphagia Inventory	Hutcheson et al 2014 <sup>45</sup>	Chemotherapy in relation to swallowing	47	Untreated stage IV SCC	Specific to swallowing	Needs to be used in conjunction with another questionnaire for more rounded assessment	Performance Status Scale for Head and Neck Cancer
	Mehanna et al 2016 <sup>19</sup>	PET-CT in advanced SCC	564	N2 or N3 metastases	Another more specific dimension related to H&N		H&N35
Spitzer QoL Index	Robert et al 1997 <sup>49</sup>	Chemotherapy medications	26	Recurrent or metastatic cancer	Related to basic living tasks	Not H&N cancer specific	
	Elliot et al 2006 <sup>50</sup>	Prevention of radiation dermatitis	547	Stage III or IV SCC requiring RT	Validated Self-assessment	May be subject to reviewer or pt bias	Head and Neck Radiotherapy Questionnaire
Hospital Anxiety and Depression Scale	Griffiths et al 1999 <sup>51</sup>	Physical & Psychological symptoms	615	SCC treatment	Psychologically specific	Data needs to be analysed in several ways to ensure consistency	Rotterdam Symptom Checklist
	Schiefke et al 2008 <sup>14</sup>	QoL after SND	49	Sentinel node bx or SND	Psychologically specific	Needs to be used in conjunction with another questionnaire for more rounded assessment	EORTC QLQ-C30
	Vilela et al 2005 <sup>16</sup>	Coping strategy therapy	101	Completed cancer treatment	Psychologically specific		EORTC QLQ-C30
	Myers et al 1999 <sup>23</sup>	Psychosocial intervention	47	H&N SCC, no previous mental health issue	Psychologically specific		EORTC QLQ-C30
Head and Neck Radiotherapy Questionnaire	Elliot et al 2006 <sup>50</sup>	Prevention of radiation dermatitis	547	Stage III or IV SCC requiring RT	H&N cancer-specific RT specific	May be subject to reviewer or pt bias	Spitzer QoL Index



	Ringash et al 2005 <sup>52</sup>	Post-RT xerostomia	130	RT to parotids	Validated Cancer-specific RT specific Multi-dimensional Easy to complete	Questionnaire disease specific Uses summary scores so may not be adequate for xerostomia assessment Does not assess patient weighting of problem	
	Wong et al 2003 <sup>53</sup>	TENS post RT	37	Xerostomia post radial RT	Easy to complete		
	Warde et al 2002 <sup>54</sup>	Oral pilocarpine for RT	130	RT with inclusion of parotids	RT specific		
EQ-5D	Mehanna et al 2016 <sup>19</sup>	PET-CT in advanced SCC	564	N2 or N3 metastases	Good for cost effectiveness assessment		EORTC QLQ-C30 H&N35 MD Anderson dysphagia
Head and Neck Cancer Inventory	Lazarus et al 2013 <sup>55</sup>	Swallowing/ Tongue strength	23	Post-op SCC resection; no pre-existing dysphagia	Good for functional scores		
Modified WHO Performance Status Scale	Corry et al 2008 <sup>56</sup>	PEG vs NG	33	RT and needing enteral feeding	Specific to NG tubes Simple	Not related to cancer	
Dermatological Life Quality Index	Potthoff et al 2014 <sup>9</sup>	Medication for pain intensity	34	Cetuximab chemoradiotherapy rhagades	Specific to dermatology	Not related to H&N cancer	EORTC QLQ-C30
Rotterdam Symptom Checklist	Griffiths et al 1999 <sup>51</sup>	Physical & Psychological symptoms	615	SCC treatment	Cancer specific General view of health Pt recorded		Hospital Anxiety and Depression Scale
Hearing Handicap Inventory for the Elderly	Schultz et al 2010 <sup>57</sup>	Hearing loss of H&N cancer patients	141	Had RT	Specific to hearing loss	Not relevant for any other features of H&N cancer	
QoL-RTI	Maguire et al 2011 <sup>58</sup>	RT therapy in advanced SCC	30	Stage III or IV SCC requiring RT	Radiation and RT specific Good for swallowing measurements	Not related to other aspects of H&N cancer	

COOP-WONCA	Van bokhorst et al 2000 <sup>12</sup>	Enteral nutrition	49	Malnourished	Generic Practical Easy	Not specific to cancer or H&N	EORTC QLQ-C30
HNQoL	Donatelli-Lassig et al 2008 <sup>59</sup>	QoL post chemoradiotherapy	65	Inclusive of SND	Cancer-specific H&N specific		

Table 4. Patient Reported Outcome Measures used in papers relevant to Head & Neck Cancer.