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Authors

Rosko, Andrew
Birkeland, Andrew
Shuman, Andrew
et al.

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PET-CT Prediction of Occult Nodal Metastasis in Recurrent Laryngeal Cancer

Andrew Rosko, MD¹, Andrew Birkeland, MD¹, Andrew Shuman, MD¹, Mark Prince, MD¹, Carol Bradford, MD¹, Gregory Wolf, MD¹, Francis Worden, MD², Avraham Eisbruch, MD³, Ashok Srinivasan, MBBS⁴, Ka Kit Wong, MBBS⁵, and Matthew E. Spector, MD¹

¹Department of Otolaryngology-Head and Neck Surgery, University of Michigan Health Center, Ann Arbor, Michigan, USA

²Department of Internal Medicine, Division of Hematology and Oncology, University of Michigan Medical School, Ann Arbor, Michigan, USA

³Department of Radiation Oncology, University of Michigan Medical School, Ann Arbor, Michigan, USA

⁴Department of Radiology, Division of Neuroradiology, University of Michigan Medical School, Ann Arbor, Michigan, USA

⁵Department of Radiology, Division of Nuclear Medicine, University of Michigan Medical School, Ann Arbor, Michigan, USA

Abstract

Background—The objective of this study was to evaluate the predictive value of PET-CT in identifying occult nodal metastasis in clinically and radiographically N0 patients with recurrent laryngeal cancer undergoing salvage laryngectomy.

Methods—Retrospective review of 46 clinically and radiographically N0 patients with recurrent laryngeal cancer who underwent a PET-CT examination prior to salvage laryngectomy with neck dissection from January 1, 2002 to December 31, 2014 was performed.

Results—Two patients (16.7%) had true positive PET-CT results, while 10 patients (83.3%) had false negative scans, one patient (2.9%) had a false positive result and 33 patients (97.1%) had a true negative PET-CT. The sensitivity of PET-CT was 16.7% (95% CI, 3.5% – 46.0%) with a specificity of 97.1% (95% CI, 83.8% – 99.9%), positive predictive value (PPV) of 66.7% (95% CI, 20.2% – 94.4%) and negative predictive value (NPV) of 76.7% (95% CI, 62.1% – 87.0%).

Conclusions—PET-CT has poor sensitivity and NPV making PET-CT an imperfect predictor of nodal disease in recurrent laryngeal cancer.

Corresponding Author: Matthew E. Spector, M.D., University of Michigan Health System, Department of Otolaryngology-Head and Neck Surgery, 1500 E Medical Center Dr., 1904 TC, SPC 5312, Ann Arbor, MI 48109-5312, 734 936-3172 phone, 734 232-1007 fax, mspector@med.umich.edu.

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Keywords

PET-CT; Salvage Laryngectomy; Laryngeal Squamous Cell Carcinoma; Occult Nodal Metastasis; Recurrent Laryngeal Cancer

INTRODUCTION

Currently laryngeal squamous cell carcinoma (SCC) is routinely treated with radiation (XRT) or concurrent chemoradiation (CRT).¹⁻⁵ This approach has been adopted with the goal of maintaining speech and swallowing function while providing similar survival rates compared to primary surgery. Unfortunately, many of these patients develop recurrence and require salvage laryngectomy.⁶⁻⁸

The role of neck dissection during salvage laryngectomy is controversial. The management of patients with clinically apparent nodal disease is straightforward, however, there is no consensus regarding management of the clinically N0 neck. Additionally, neck dissection adds morbidity to total laryngectomy and is associated with a higher complication rate, including a higher rate of pharyngocutaneous fistula.⁹

The prevalence of occult nodal metastases in the N0 neck in recurrent laryngeal cancer has been reported from 4% to 20%.⁹⁻¹³ Supraglottic tumors and advanced T classification tumors have been associated with an even higher occult nodal metastasis rate.¹¹⁻¹³ When taken together, patients with T4 supraglottic tumors have a 50% risk of occult nodal disease.¹³ Efforts have been made to predict the likelihood of nodal metastasis in these patients. The staging of the neck prior to initial therapy has been shown to correlate with the risk of occult nodal disease in neck dissection specimens.¹¹ Pre-operative computed tomography (CT) and magnetic resonance imaging (MR) have been used to improve pre-operative staging, however these tests have poor sensitivity leading to an inadequate negative predictive value.¹⁴⁻¹⁶

2-[fluorine-18]fluoro-2-deoxy-d-glucose (FDG) positron-emission tomography, in combination with CT (PET-CT), is increasingly being used in the management of head and neck cancer patients, including initial staging, assessing response to therapy, detecting recurrence, and identifying unknown primary tumors.^{15,17-29} Additionally, PET-CT is increasingly used to improve the accuracy of staging in recurrent laryngeal cancer.³⁰ While the ability of PET-CT to detect nodal metastasis in other tumors has been addressed, the predictive value of PET-CT in detecting nodal disease in patients with recurrent laryngeal cancer has not been well studied.^{15,16,21,29-37} The objective of this study was to evaluate the predictive value of PET-CT in identifying nodal metastasis in patients with recurrent laryngeal carcinoma with a clinically and radiographically N0 neck. The overall goal is to assess the ability of PET-CT to discriminate the need for elective neck dissection.

MATERIALS AND METHODS

A retrospective review was performed of a cohort of patients who underwent salvage laryngectomy with neck dissection without clinical or radiographic (CT or MR) evidence of

nodal disease at the University of Michigan from January 1, 2002 to December 31, 2014. All patients had a PET-CT prior to surgery. Salvage laryngectomy was defined as surgery for persistent or recurrent laryngeal SCC after initial XRT or CRT. Patients were excluded if they had evidence of nodal disease on clinical exam or pre-operative CT or MR. Forty-six patients met inclusion criteria. Clinical and pathologic data was collected including initial stage and treatment of the primary tumor as well as the staging of the recurrent or persistent tumor. Patients were staged based on the 7th edition of the AJCC staging system.

Patients who underwent a PET-CT at the University of Michigan (91.3%, 42/46) and those who had a PET-CT at an outside institution (8.7%, 4/46) were included in this study. The PET-CT was performed per institutional protocol. At the University of Michigan, patients were fasted for > 4–6 h and had glucose levels < 250 mg/dL prior to imaging. Around sixty minutes following intravenous administration of 300 MBq (8 mCi) of FDG, sequential PET and CT imaging was performed on an integrated PET-CT scanner (Siemens Biograph T6; Siemens Medical Solutions, Hoffman Estates, IL, USA). Helical CT from skull vertex to mid-thigh was performed with 5 mm collimation (low dose CT parameters: 140kV, 80mA, tube rotation of 0.8 s per rotation, pitch of 3:1), followed immediately by whole body PET at multiple overlapping bed positions (area covered: skull vertex to mid-thigh, step-and-shoot mode, 3 mins per bed-position). Then a dedicated contrast-enhanced head and neck CT was performed with field of view 15 cm, commencing 40 seconds after intravenous injection of 100 ml volume of iopromide (Ultravist) at 1.5 ml/second, and co-registered to the whole body FDG PET dataset. Images were reviewed on a workstation (MedImage; MedView Pty, Canton, MI, USA) by 2 readers (one head and neck radiologist and one nuclear medicine physician) providing a single read per study.

Four of the PET-CT studies were performed at outside centers, with the imaging data transferred from compact disc to our local archive for review. This led to some heterogeneity in the imaging protocols used, as some outside centers performed dedicated head and neck PET-CT images acquired separately (in addition to the whole body PET-CT) whilst others performed only a whole body PET-CT (with a large field of view). Evidence of PET-CT positivity was determined by the official report of the reading radiologist at the time of the scan, on which patients' management was based. In addition, PET-CT images were reviewed by a single reader (author KKW, board certified in Nuclear Medicine) who was blinded to the results of the official report and final pathology, with interpretation compared to the initial clinical reads. There were no discrepancies between the official reports and the review by author KKW.

Cervical lymph nodes displaying FDG uptake above background were considered either suspicious or positive for regional nodal metastasis. PET-CT results were categorized as positive or negative. All nodes that were considered suspicious by the reading radiologist were categorized as positive. Final pathology results were collected and the nodal metastases were tabulated. For patients that underwent a unilateral neck dissection, only the dissected neck was included in the analysis.

Patient Population

Forty-six patients, 84.8% (39/46) men and 15.2% (7/46) women with a mean age of 62.1 ± 9.7 years were included in our cohort. At the time of salvage surgery 41.3% (19/46) of patients were current smokers and 58.7% (27/46) were former smokers. At the time of initial therapy, 47.8% (22/46) of tumors were located in the supraglottis, whereas 52.2% (24/46) were located in the glottis. The neck was initially staged as N0 in 71.1% (32/45) of patients, N1 in 13.3% (6/45) of patients, N2b in 6.7% (3/45) of patients and N2c in 8.9% (4/45) of patients. The initial staging was missing in 1 patient who was previously treated at an outside institution. Of the patients, 41.3% (19/46) were initially treated with XRT, 54.3% (25/46) were treated with CRT, and 4.3% (2/46) were treated with laser excision followed by adjuvant XRT or CRT. The mean interval to recurrence was 20.4 months. Of the recurrent primary tumors, 52.2% (24/46) were located in the supraglottis, 45.6% (21/46) were in the glottis, and 2.2% (1/46) were centered in the subglottis. Recurrent tumors were staged as T1 in 4.3% (2/46) of patients, T2 in 19.6% (9/46) of patients, T3 in 32.6% (15/46) of patients and T4 in 43.5% (20/46) of patients. Patient characteristics, staging and treatment are shown in Table 1.

Statistical Analysis

In our analysis we initially compared PET-CT positivity to pathologic node status on a per patient basis. We subsequently evaluated each neck specimen separately with the left and right necks being analyzed individually. After analyzing the cohort as a whole, subgroup analysis was performed by tumor subsite, T classification, and those patients without nodal metastasis (N0) prior to initial treatment (XRT or CRT). To calculate the sensitivity and specificity of PET-CT in predicting nodal metastasis, results of the PET-CT scans were compared to the final pathology (gold standard). Positive and negative predictive values were calculated. Ninety-five percent confidence intervals (95% CI) were calculated using the method described by Agresti and Coull.³⁸ All statistical analysis was performed using IBM SPSS (version 20) software (IBM Corporation, Armonk, New York) with consultation from the University of Michigan Center for Statistical Consultation And Research (CSCAR). This study was approved by the University of Michigan Internal Review Board (HUM00081554).

RESULTS

In our cohort of 46 patients, 3 (6.5%) patients had a positive PET-CT scan. Twelve patients (26.1%) were noted to have nodal metastasis on final pathology, with 2 (16.6%) true positive and 10 (83.3%) false negative PET-CTs. Of the 34 patients without nodal metastasis on final pathology there was one false positive scan (2.9%) and 33 (97.1%) true negative scans. (Table 2) The sensitivity of PET-CT when compared to the final pathology was 16.7% (95% CI, 3.5% – 46.0%) and the specificity was 97.1% (95% CI, 83.8% – 99.9%). In this cohort the positive predictive value was 66.7% (95% CI, 20.2% – 94.4%) and negative predictive value was 76.7% (95% CI, 62.1% – 87.0%).

We then evaluated the cohort looking at each neck specimen separately. Eight patients either had a previous unilateral neck dissection as part of their initial therapy (n=4) or only the ipsilateral neck was dissected (n=4) at the time of salvage laryngectomy, and thus there were

84 neck specimens in our analysis. In total, 15 (17.8%) of neck specimens contained nodal metastasis, 2 (13.3%) of which were detected by PET-CT. There were 69 neck specimens without nodal disease with only 2 (3.0%) having false positive scans. The false positive scans corresponded to one patient with a PET-CT suggestive of bilateral neck disease who had no pathologic lymph nodes on final pathology. The sensitivity of PET-CT when compared to the final pathology was 13.3% (95% CI, 24.8% – 39.1%) and the specificity was 97.1% (95% CI, 89.4% – 99.8%). In this analysis, the positive predictive value was 50.0% (95% CI, 15.0% – 85.0%) and negative predictive value was 83.8% (95% CI, 74.0% – 90.4%). (Table 3)

In a subgroup analysis, we evaluated the 32 patients classified as N0 prior to their initial therapy. In this group there were 10 (31.3%) patients with nodal metastasis on final pathology, 2 (20.0%) of which were identified on PET-CT with 8 false negative scans. Of the 22 patients without regional metastasis there was 1 (4.5%) false positive and 21 (95.4%) true negative PET-CT scans (Table 4). The sensitivity and specificity were 20.0% (95% CI, 4.6% – 52.1%) and 95.5% (95% CI, 76.5% – 99.9%) respectively. In this population this corresponds to a positive predictive value of 66.7% (95% CI, 20.2% – 94.3%) and negative predictive value of 72.4% (95% CI, 54.1% – 85.5%).

We evaluated our cohort of patients who were classified as N0 prior to their initial therapy looking at each neck specimen separately. There were 58 neck specimens in our analysis. There were 13 (22.4%) neck specimens with pathologically confirmed nodal metastasis, 2 (15.4%) of which were identified on PET-CT with 11 false negative scans. Of the 45 neck specimens without pathologically positive nodes, 43 (95.6%) had a negative PET-CT with 2 false positive scans (Table 5). The sensitivity and specificity were 15.4% (95% CI, 3.1% – 43.4%) and 95.6% (95% CI, 84.4% – 99.6%) respectively. The positive predictive value was 50.0% (95% CI, 15.0% – 85.0%) and the negative predictive value was 79.6% (95% CI, 66.9% – 88.4%).

In order to better understand the performance of PET-CT, we analyzed our cohort based on the recurrent tumor subsite and recurrent tumor T classification (Table 6). The small size of each group limits the analysis. As expected the NPV was highest in patients with low risk for nodal metastasis (T1 or T2 tumors and glottis primaries) and was lowest in patients who were at the highest risk for occult nodal disease (T3 and T4 tumors and supraglottic primaries). Although the sensitivity was slightly better in patients with T3 tumors compared to other T classifications, the differences were not statistically significant. This was due to the small number of true positive scans in the cohort.

The patients who had false negative PET-CT scans are of particular interest. Of the 10 patients in our overall cohort with false negative PET-CT scans, 6 patients had 1 occult node, 1 patient had 2 occult nodes, 1 patient had 3 occult nodes, 1 patient had 7 occult nodes and 1 patient had 29 occult nodes that were positive for metastatic squamous cell carcinoma on final pathology. (Table 7) Histopathologic lymph node tissue was available for review in 5 of the 10 patients. The average size of metastatic deposit in the lymph node was 3 mm (range 1–7mm). Representative images of the PET-CT from the patient with 29 occult nodes are

shown in Figure 1 and representative images of an occult node from this patient are shown in Figure 2.

DISCUSSION

PET-CT has proven to be a valuable tool in caring for patients with head and neck cancer. First described by Warburg in 1956, PET-CT capitalizes on the concept that malignant cells exhibit increased glucose utilization by upregulating glucose transporters (GLUT).³⁹ FDG, a radiopharmaceutical analog of glucose, is taken up by malignant cells and undergoes phosphorylation by hexokinase to FDG-6-phosphate; however, unlike glucose, FDG is trapped intracellularly as an index of the metabolic activity of tumor cells.³⁹ This technology has proven useful in initial staging, assessing treatment response, monitoring for recurrence, and identifying distant metastasis in patients with head and neck cancer.^{18,19,26,34,40,41} While PET-CT has been shown to have increased sensitivity when compared to CT or MR, the role of PET-CT in guiding treatment of the neck, especially in the salvage setting, is evolving.¹³⁻¹⁵

This is the largest study to evaluate the ability of PET-CT to detect nodal metastasis in patients with recurrent laryngeal carcinoma. In our study of clinically and radiographically (based on CT or MR) N0 patients, the sensitivity of PET-CT was 16.7% with a specificity of 97.1%. In our population, this yielded a negative predictive value of only 76.7%. When we compared our results to other reports in the literature, we noted a wide range of reported sensitivities and specificities. In these studies, the reported sensitivity ranges from 71% to 89% and specificity ranges from 82% to 100%.^{21, 32, 36, 40-41} In the largest meta-analysis of 1236 patients in 32 studies, PET-CT had pooled sensitivity of 79% (CI 72%–85%) in detecting cervical nodal metastases with a specificity of 86% (CI 83%–89%). Notably, PET-CT had an even lower sensitivity of 50% (CI 37%–63%) in detecting occult nodal metastasis in the clinically cN0 patient, though it did have a reasonable specificity of 87% (76%–93%).⁴²

The current study reports a sensitivity that is substantially lower than previous reports. This is likely due to the differences in study design. The previous studies included patients with head and neck cancer from all subsites, with laryngeal primaries making up a small proportion of the population. Additionally, the previous studies included patients who were previously untreated, while our study only included patients who were previously treated with XRT or CRT. This is important as chemotherapy and radiation alters the lymphatics with decreased vascularity of the residual nodes. Thus many of the pathologically positive nodes would be below the size detection limit of PET-CT (6–8 mm), which would in turn limit the sensitivity.⁴³ Perhaps most importantly, all other studies included patients with clinically or radiographically apparent nodes. Including patients with known nodal metastasis would increase the number of true positive scans, which in turn would increase the calculated sensitivity. This study design fails to answer the question regarding the ability of PET-CT to detect occult nodal disease. This phenomenon is demonstrated in the meta-analysis by Kyzas et al. as the sensitivity drops from 79% to 50% when patients with clinically evident nodes are excluded.⁴² The current study is only study that evaluating the utility of PET-CT to evaluate the neck prior to salvage laryngectomy in a cohort of patients previously treated

with CRT or XRT with no clinical or radiographic nodal metastasis prior to PET-CT. This difference in clinical design almost certainly accounts for the difference in sensitivity seen in our study.

The closest report to the current study was published by Gilbert et al. This study consisted of a review of 15 patients with SCC of the larynx who underwent elective neck dissection at the time of salvage laryngectomy. In their study, they reported a sensitivity of 70% (95% CI, 39% – 90%), specificity of 100% (95% CI, 51% – 100%) and negative predictive value of 63% (95% CI, 30% – 87%). Once again, this study included patients with and without clinically apparent nodal disease, which led to a higher sensitivity than the current study as described above. Based on these results, the authors concluded that the false negative rate is too high to defer neck dissection based solely on PET-CT results, and this conclusion is supported in our study.³⁰

By using the pre-test probability of occult nodal disease, the sensitivity and specificity of PET-CT from this study can be used to calculate the positive and negative predictive value of a particular patient population based on previously described methods⁴⁴ This allows the clinician to determine the post-test probability of occult nodal disease, and it is the post-test probability of occult disease that should drive the decision regarding whether or not a neck dissection should be performed. In patients with clinically apparent nodal disease at the time of the laryngectomy, the decision to proceed with neck dissection is straightforward, however, the decision becomes more complicated in the previously treated neck with no evidence of disease. In the original paper by Weiss et al. in 1994 decision analysis was used with the conclusion that the N0 neck should be treated if the risk of occult nodal metastasis is greater than 20%.⁴⁵ Ferlito et al. in their review of neck dissection in laryngeal cancer suggested that the neck should be treated if the risk of metastasis is 15%.⁴⁶

Careful consideration of the risk of occult nodal disease is important in deciding to perform a neck dissection as salvage laryngectomy is already associated with impaired wound healing and wound related complications. Furthermore, neck dissection is associated with increased morbidity, including a higher fistula rate.^{9,47-49} Unfortunately there are few remaining treatment modalities available to the patient undergoing salvage laryngectomy, and neck dissection becomes more difficult once a laryngectomy has been performed, especially if free tissue is used to reconstruct the pharynx. Thus identifying those patients at high risk of regional failure is important. In the present study, 26% of patients had occult cervical metastasis not detected on either clinical exam or CT. PET-CT failed to identify the majority of these patients as 23% of patients with a negative PET-CT had occult nodal disease. Thus, based on the low sensitivity, low negative predictive value PET-CT alone is an inadequate test to withhold neck dissection.

When we considered why PET-CT had an unacceptably low sensitivity, there was no evidence that the high rate of false-negative scans was related to patient preparation factors, modality of treatment (XRT versus CRT) or the timing of the PET-CT after treatment. It is believed that this was instead related to low volume disease in the previously treated neck with cancer deposits that were below the spatial resolution of PET-CT leading to partial volume effect and reduced sensitivity. On re-review of the histopathology in the patients who

had false negative PET-CT scans in our study, the average metastatic deposit was 3mm, confirming the low volume of disease. Post-treatment lymph nodes with limited tumor burden fall below the detection limit of the PET-CT leading to false negative scans.

There are limitations to our current study. While this is the largest study evaluating the ability of PET-CT to detect nodal disease prior to salvage laryngectomy the sample size is still relatively small. This retrospective study included PET-CT scans performed at our institution and also from outside centers leading to some heterogeneity in the imaging protocols used. Protocols performing dedicated head and neck PET-CT images acquired separately (in addition to the whole body PET-CT) could have improved conspicuity of small cervical nodes compared to protocols with whole body PET-CT acquired over a large field of view. However, excluding patients with outside imaging studies did not affect our sensitivity (18.2% vs 16.7%) and specificity (96.8% vs 97.1%) significantly. The retrospective nature of this study also limits the ability to draw conclusions from the dataset. Given these limitations, a prospective study of PET-CT in salvage laryngectomy is needed to fully address this question.

CONCLUSION

This study demonstrates that while PET-CT has a reasonable specificity and positive predictive value, it has inadequate sensitivity and negative predictive value. Based on these results, neck dissection should not be withheld solely on the basis of a negative PET-CT when performing salvage laryngectomy for recurrent laryngeal SCC.

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References

1. Urba S, Wolf G, Eisbruch A, et al. Single-cycle induction chemotherapy selects patients with advanced laryngeal cancer for combined chemoradiation: a new treatment paradigm. *J Clin Oncol.* 2006; 24:593–598. [PubMed: 16380415]
2. Lefebvre JL, Chevalier D, Luboinski B, et al. Larynx preservation in pyriform sinus cancer: preliminary results of a European Organization for Research and Treatment of Cancer phase III trial. EORTC Head and Neck Cancer Cooperative Group. *J Natl Cancer Inst.* 1996; 88:890–899. [PubMed: 8656441]
3. Forastiere AA, Goepfert H, Maor M, et al. Concurrent chemotherapy and radiotherapy for organ preservation in advanced laryngeal cancer. *N Engl J Med.* 2003; 349:2091–2098. [PubMed: 14645636]
4. Chone CT, Yonehara E, Martins JE, et al. Importance of anterior commissure in recurrence of early glottic cancer after laser endoscopic resection. *Arch Otolaryngol Head Neck Surg.* 2007; 133:882–887. [PubMed: 17875854]
5. Hakeem AH, Tubachi J, Pradhan SA. Significance of anterior commissure involvement in early glottic squamous cell carcinoma treated with trans-oral CO2 laser microsurgery. *Laryngoscope.* 2013; 123:1912–1917. [PubMed: 23606304]
6. Clark JR, de Almeida J, Gilbert R, et al. Primary and salvage (hypo)pharyngectomy: Analysis and outcome. *Head Neck.* 2006; 28:671–677. [PubMed: 16721745]

7. Paydarfar JA, Birkmeyer NJ. Complications in head and neck surgery: a meta-analysis of postlaryngectomy pharyngocutaneous fistula. *Arch Otolaryngol Head Neck Surg.* 2006; 132:67–72. [PubMed: 16415432]
8. Li M, Lorenz RR, Khan MJ, et al. Salvage laryngectomy in patients with recurrent laryngeal cancer in the setting of nonoperative treatment failure. *Otolaryngol Head Neck Surg.* 2013; 149:245–251. [PubMed: 23585149]
9. Bohannon IA, Desmond RA, Clemons L, et al. Management of the N0 neck in recurrent laryngeal squamous cell carcinoma. *Laryngoscope.* 2010; 120:58–61. [PubMed: 19877259]
10. Farrag TY, Lin FR, Cummings CW, et al. Neck management in patients undergoing postradiotherapy salvage laryngeal surgery for recurrent/persistent laryngeal cancer. *Laryngoscope.* 2006; 116:1864–1866. [PubMed: 17003711]
11. Wax MK, Touma BJ. Management of the N0 neck during salvage laryngectomy. *Laryngoscope.* 1999; 109:4–7. [PubMed: 9917031]
12. Yao M, Roebuck JC, Holsinger FC, et al. Elective neck dissection during salvage laryngectomy. *Am J Otolaryngol.* 2005; 26:388–392. [PubMed: 16275407]
13. Birkeland AC, Rosko AJ, Issa MR, et al. Occult Nodal Disease Prevalence and Distribution in Recurrent Laryngeal Cancer Requiring Salvage Laryngectomy. *Otolaryngol Head Neck Surg.* 2016; 154:473–479. [PubMed: 26884365]
14. Ng SH, Yen TC, Liao CT, et al. 18F-FDG PET and CT/MRI in oral cavity squamous cell carcinoma: a prospective study of 124 patients with histologic correlation. *J Nucl Med.* 2005; 46:1136–1143. [PubMed: 16000282]
15. Adams S, Baum RP, Stuckensen T, et al. Prospective comparison of 18F-FDG PET with conventional imaging modalities (CT, MRI, US) in lymph node staging of head and neck cancer. *Eur J Nucl Med.* 1998; 25:1255–1260. [PubMed: 9724374]
16. Braams JW, Pruijm J, Freling NJ, et al. Detection of lymph node metastases of squamous-cell cancer of the head and neck with FDG-PET and MRI. *J Nucl Med.* 1995; 36:211–216. [PubMed: 7830116]
17. Abgral R, Querellou S, Potard G, et al. Does 18F-FDG PET/CT improve the detection of posttreatment recurrence of head and neck squamous cell carcinoma in patients negative for disease on clinical follow-up? *J Nucl Med.* 2009; 50:24–29. [PubMed: 19091901]
18. Conessa C, Herve S, Fohrenbach H, et al. FDG-PET scan in local follow-up of irradiated head and neck squamous cell carcinomas. *Ann Otol Rhinol Laryngol.* 2004; 113:628–635. [PubMed: 15330142]
19. Dobert N, Kovacs AF, Menzel C, et al. The prognostic value of FDG PET in head and neck cancer. Correlation with histopathology. *Q J Nucl Med Mol Imaging.* 2005; 49:253–257. [PubMed: 16172571]
20. Hannah A, Scott AM, Tochon-Danguy H, et al. Evaluation of 18 F-fluorodeoxyglucose positron emission tomography and computed tomography with histopathologic correlation in the initial staging of head and neck cancer. *Ann Surg.* 2002; 236:208–217. [PubMed: 12170026]
21. Kim SY, Kim JS, Yi JS, et al. Evaluation of 18F-FDG PET/CT and CT/MRI with histopathologic correlation in patients undergoing salvage surgery for head and neck squamous cell carcinoma. *Ann Surg Oncol.* 2011; 18:2579–2584. [PubMed: 21409485]
22. Kitagawa Y, Nishizawa S, Sano K, et al. Prospective comparison of 18F-FDG PET with conventional imaging modalities (MRI, CT, and 67Ga scintigraphy) in assessment of combined intraarterial chemotherapy and radiotherapy for head and neck carcinoma. *J Nucl Med.* 2003; 44:198–206. [PubMed: 12571209]
23. Kitagawa Y, Sadato N, Azuma H, et al. FDG PET to evaluate combined intra-arterial chemotherapy and radiotherapy of head and neck neoplasms. *J Nucl Med.* 1999; 40:1132–1137. [PubMed: 10405132]
24. Kitagawa Y, Sano K, Nishizawa S, et al. FDG-PET for prediction of tumour aggressiveness and response to intra-arterial chemotherapy and radiotherapy in head and neck cancer. *Eur J Nucl Med Mol Imaging.* 2003; 30:63–71. [PubMed: 12483411]
25. Kresnik E, Mikosch P, Gallowitsch HJ, et al. Evaluation of head and neck cancer with 18F-FDG PET: a comparison with conventional methods. *Eur J Nucl Med.* 2001; 28:816–821.

26. Lowe VJ, Dunphy FR, Varvares M, et al. Evaluation of chemotherapy response in patients with advanced head and neck cancer using [F-18]fluorodeoxyglucose positron emission tomography. *Head Neck*. 1997; 19:666–674. [PubMed: 9406745]
27. Spector ME, Chinn SB, Rosko AJ, et al. Diagnostic modalities for distant metastasis in head and neck squamous cell carcinoma: are we changing life expectancy? *Laryngoscope*. 2012; 122:1507–1511. [PubMed: 22460441]
28. Terhaard CH, Bongers V, van Rijk PP, et al. F-18-fluoro-deoxy-glucose positron-emission tomography scanning in detection of local recurrence after radiotherapy for laryngeal/pharyngeal cancer. *Head Neck*. 2001; 23:933–941. [PubMed: 11754496]
29. Wong RJ, Lin DT, Schoder H, et al. Diagnostic and prognostic value of [(18)F]fluorodeoxyglucose positron emission tomography for recurrent head and neck squamous cell carcinoma. *J Clin Oncol*. 2002; 20:4199–4208. [PubMed: 12377963]
30. Brouwer J, de Bree R, Comans EF, et al. Improved detection of recurrent laryngeal tumor after radiotherapy using (18)FDG-PET as initial method. *Radiother Oncol*. 2008; 87:217–220. [PubMed: 18329117]
31. Gilbert MR, Branstetter BfT, Kim S. Utility of positron-emission tomography/computed tomography imaging in the management of the neck in recurrent laryngeal cancer. *Laryngoscope*. 2012; 122:821–825. [PubMed: 22344673]
32. Hyde NC, Prvulovich E, Newman L, et al. A new approach to pre-treatment assessment of the N0 neck in oral squamous cell carcinoma: the role of sentinel node biopsy and positron emission tomography. *Oral Oncol*. 2003; 39:350–360. [PubMed: 12676254]
33. Nahmias C, Carlson ER, Duncan LD, et al. Positron emission tomography/computerized tomography (PET/CT) scanning for preoperative staging of patients with oral/head and neck cancer. *J Oral Maxillofac Surg*. 2007; 65:2524–2535. [PubMed: 18022480]
34. Paulus P, Sambon A, Vivegnis D, et al. 18FDG-PET for the assessment of primary head and neck tumors: clinical, computed tomography, and histopathological correlation in 38 patients. *Laryngoscope*. 1998; 108:1578–1583. [PubMed: 9778305]
35. Stuckensen T, Kovacs AF, Adams S, et al. Staging of the neck in patients with oral cavity squamous cell carcinomas: a prospective comparison of PET, ultrasound, CT and MRI. *J Craniomaxillofac Surg*. 2000; 28:319–324. [PubMed: 11465137]
36. Wensing BM, Vogel WV, Marres HA, et al. FDG-PET in the clinically negative neck in oral squamous cell carcinoma. *Laryngoscope*. 2006; 116:809–813. [PubMed: 16652093]
37. Yamazaki Y, Saitoh M, Notani K, et al. Assessment of cervical lymph node metastases using FDG-PET in patients with head and neck cancer. *Ann Nucl Med*. 2008; 22:177–184. [PubMed: 18498032]
38. Agresti A, Coull BA. Approximate is better than “exact” for interval estimation of binomial proportions. *American Statistician*. 1998; 52:119–126.
39. Warburg O. On the origin of cancer cells. *Science*. 1956; 123:309–314. [PubMed: 13298683]
40. Ong SC, Schoder H, Lee NY, et al. Clinical utility of 18F-FDG PET/CT in assessing the neck after concurrent chemoradiotherapy for Locoregional advanced head and neck cancer. *J Nucl Med*. 2008; 49:532–540. [PubMed: 18344440]
41. Schoder H, Carlson DL, Kraus DH, et al. 18F-FDG PET/CT for detecting nodal metastases in patients with oral cancer staged N0 by clinical examination and CT/MRI. *J Nucl Med*. 2006; 47:755–762. [PubMed: 16644744]
42. Kyzas PA, Evangelou E, Denaxa-Kyza D, et al. 18F-fluorodeoxyglucose positron emission tomography to evaluate cervical node metastases in patients with head and neck squamous cell carcinoma: a meta-analysis. *J Natl Cancer Inst*. 2008; 100:712–720. [PubMed: 18477804]
43. Purohit BS, Ailianou A, Dulguerov N, et al. FDG-PET/CT pitfalls in oncological head and neck imaging. *Insights Imaging*. 2014; 5:585–602. [PubMed: 25154759]
44. Vecchio TJ. Predictive value of a single diagnostic test in unselected populations. *N Engl J Med*. 1966; 274:1171–1173. [PubMed: 5934954]
45. Weiss MH, Harrison LB, Isaacs RS. Use of decision analysis in planning a management strategy for the stage N0 neck. *Arch Otolaryngol Head Neck Surg*. 1994; 120:699–702. [PubMed: 8018319]

46. Ferlito A, Rinaldo A, Silver CE, et al. Neck dissection for laryngeal cancer. *J Am Coll Surg*. 2008; 207:587–593. [PubMed: 18926464]
47. Rutledge JW, Spencer H, Moreno MA. Predictors for Perioperative Outcomes following Total Laryngectomy: A University HealthSystem Consortium Discharge Database Study. *Otolaryngol Head Neck Surg*. 2014; 151:81–86. [PubMed: 24690762]
48. Weber RS, Berkey BA, Forastiere A, et al. Outcome of salvage total laryngectomy following organ preservation therapy: the Radiation Therapy Oncology Group trial 91–11. *Arch Otolaryngol Head Neck Surg*. 2003; 129:44–49. [PubMed: 12525193]
49. Graboyes EM, Yang Z, Kallogjeri D, et al. Patients undergoing total laryngectomy: an at-risk population for 30-day unplanned readmission. *JAMA Otolaryngol Head Neck Surg*. 2014; 140:1157–1165. [PubMed: 25144379]

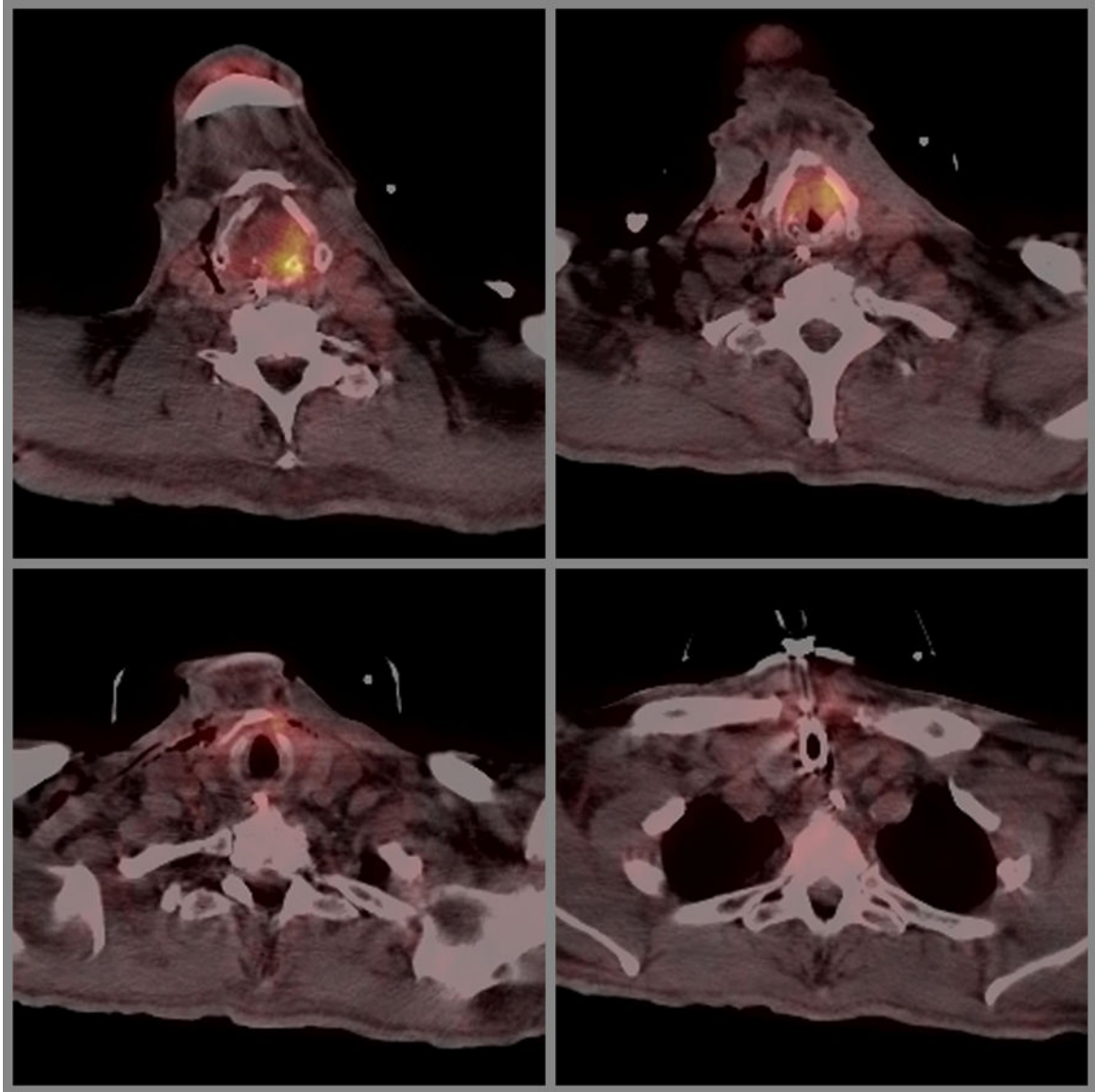


Figure 1. Representative images from a false negative PET-CT (patient number 3 in Table 6). Final pathology revealed 29 positive nodes.

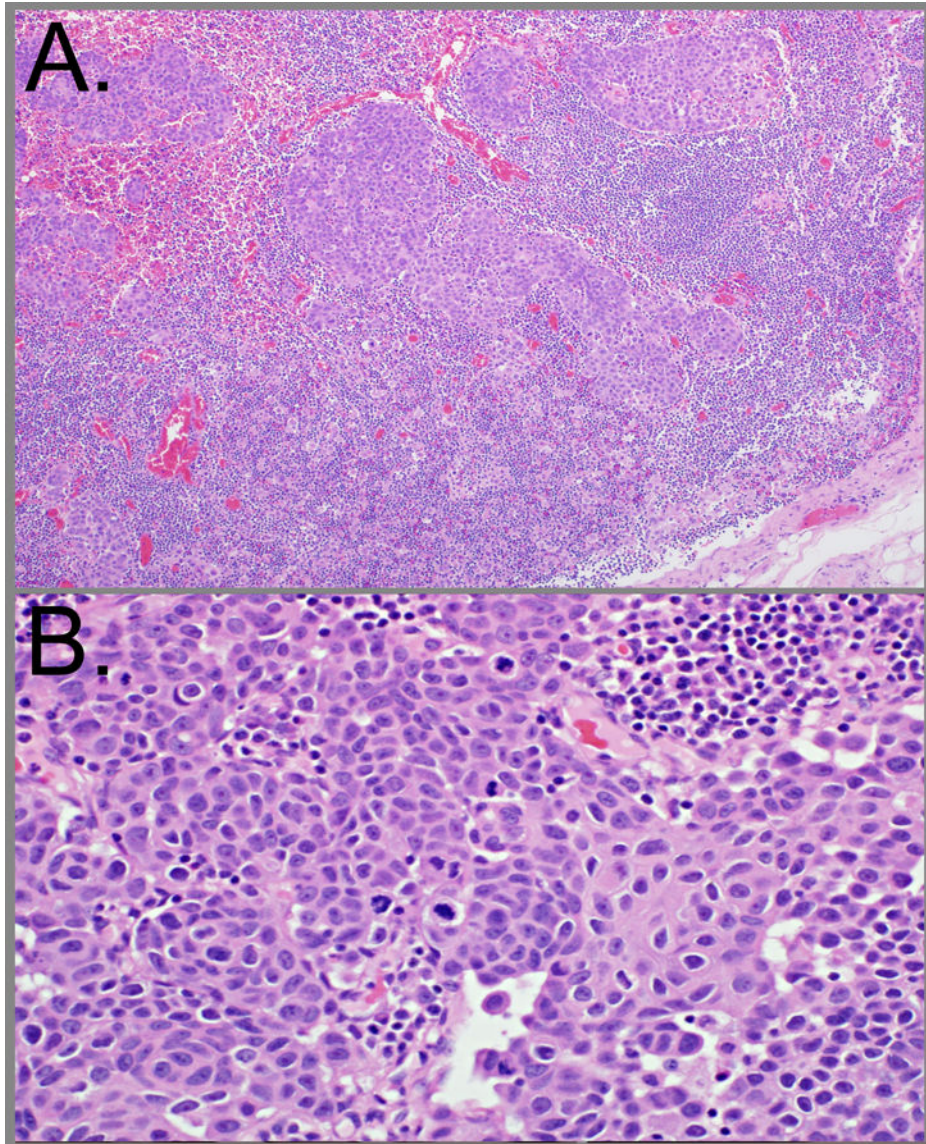


Figure 2. Representative images from the final pathology of an occult node from a patient with a false negative PET-CT (patient number 3 in Table 6, PET-CT shown in Figure 1). **A.** Low powered view (10×). **B.** High power view (40×).

Table 1

Baseline characteristics of the study cohort.

Characteristic	Value (n=46)
Age at Salvage Surgery (years), mean (SD)	62.1 (9.7)
Gender	
Male	84.8% (39)
Female	15.2% (7)
Tobacco Status at Salvage	
Current	41.3% (19)
Former	58.7% (27)
Never	0% (0)
Alcohol Status at Salvage	
Current *	15.9% (7)
Former *	11.4% (5)
Never *	72.7% (32)
Missing Data	3.8% (2)
Initial Tumor Subsite	
Supraglottic	47.8% (22)
Glottic	52.2% (24)
Subglottic	0% (0)
Initial Stage	
I *	20.0% (9)
II *	26.7% (12)
III *	33.3% (15)
IV *	20.0% (9)
Missing Data	2.2% (1)
Initial N Classification	
N0	71.1% (32)
N1	13.3% (6)
N2a	6.7% (3)
N2b	8.9% (4)
N2c	2.2 (1)
MD	
Initial Treatment	
XRT	41.3% (19)
CRT	54.3% (25)
Laser excision with XRT	2.2% (1)
Laser excision with CRT	2.2% (1)
Interval to Recurrence (months), mean (SD)	20.4 (20.0)
Recurrent/Persistent Tumor Subsite	
Supraglottic	52.2% (24)

Characteristic	Value (n=46)
Glottic	45.6% (21)
Subglottic	2.2% (1)
Recurrence T Classification	
T1	4.3% (2)
T2	19.6% (9)
T3	32.6% (15)
T4	43.5% (20)

Note, initial tumor designates the tumor characteristics and treatment prior to the diagnosis of recurrence or persistence. Recurrent tumor designates that tumor treated by salvage laryngectomy with neck dissection.

* Data shown as a percentage of the known conditions of the variable excluding those patients with missing data.

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Table 2

PET-CT test results compared to final pathology (gold standard) in the entire cohort.

PET-CT Evidence of Nodal Metastasis	Pathological Nodal Metastasis	
	YES	NO
YES	2	1
NO	10	33

Sensitivity 16.7% (95% CI, 3.5% – 46.0%)

Specificity 97.1% (95% CI, 83.8% – 99.9%)

Positive Predictive Value 66.7% (95% CI, 20.2% – 94.4%)

Negative Predictive Value 76.7% (95% CI, 62.1% – 87.0%)

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Table 3

PET-CT test results compared to final pathology (gold standard) in the entire cohort when evaluating the left and right neck separately.

PET-CT Evidence of Nodal Metastasis	Pathological Nodal Metastasis	
	YES	NO
YES	2	2
NO	13	67

Sensitivity 13.3% (95% CI, 24.8% – 39.1%)

Specificity 97.1% (95% CI, 89.4% – 99.8%)

Positive Predictive Value 50.0% (95% CI, 15.0% – 85.0%)

Negative Predictive Value 83.8% (95% CI, 74.0% – 90.4%)

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Table 4

PET-CT test results compared to final pathology (gold standard) in patients who were previously N0 prior to initial therapy.

PET-CT Evidence of Nodal Metastasis	Pathological Nodal Metastasis	
	YES	NO
YES	2	1
NO	8	21

Sensitivity 20.0% (95% CI, 4.6% – 52.1%)

Specificity 95.5% (95% CI, 76.5% – 99.9%)

Positive Predictive Value 66.7% (95% CI, 20.2% – 94.3%)

Negative Predictive Value 72.4% (95% CI, 54.1% – 85.5%)

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Table 5

PET-CT test results compared to final pathology (gold standard) in patients who were previously N0 prior to initial therapy when evaluating the left and right neck separately.

PET-CT Evidence of Nodal Metastasis	Pathological Nodal Metastasis	
	YES	NO
YES	2	2
NO	11	43

Sensitivity 15.4% (95% CI, 3.1% – 43.4%); Specificity 95.6% (95% CI, 84.4% – 99.6%); Positive Predictive Value 50.0% (95% CI, 15.0% – 85.0%); Negative Predictive Value 79.6% (95% CI, 66.9% – 88.4%)

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Table 6

PET-CT Performance based on recurrent tumor subsite and T classification

Variable	Sensitivity (95% CI)	Specificity (95% CI)	PPV (95% CI)	NPV (95% CI)
T1/T2 (n=11)	NA *	100% (71.2–100%)	NA *	100% (71.2–100%)
T3 (n=15)	33.3% (56.3–79.8%)	100% (71.2–100%)	100% (16.8–100%)	85.6% (58.8–97.2%)
T4 (n=20)	11.1% (0.0–45.7%)	90.1% (60.1–100%)	50.0% (9.5–90.6%)	55.6% (33.7–75.5%)
T3/T4 (n=35)	16.7% (3.5–46.0%)	95.7% (77.3–100%)	66.7% (20.2–94.4%)	68.8% (51.3–82.2%)
Supraglottis (n=24)	0% (0–37.2%)	100% (77.3–100%)	NA **	66.7% (46.6–82.2%)
Glottis (n=21)	50.0% (15.0–85.0%)	94.4% (72.4–100%)	66.7% (20.2–94.4%)	89.5% (67.4–98.3%)

* No patients with T2 tumors had positive nodes on final pathology

** There were no positive PET-CT scans

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Table 7

Number of positive nodes on final pathology in patients with false negative PET-CT

Patient Number	Nodes positive in Right Neck	Nodes positive in Left Neck	Total Positive Nodes	Largest Metastatic Deposit (mm)
1	1	0	1	md
2	4	3	7	md
3	20	9	29	7
4	1	1	2	1.5
5	0	1	1	md
6	0	1	1	md
7	NA *	1	1	1
8	1	NA *	1	1
9	NA *	3	3	md
10	0	1	1	3

* Unilateral neck dissection performed. NA – not applicable md – missing data

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