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should be clearly understood by clinicians before applying any of these methods.

Morbidity at 5 years	CHART N <sub>0</sub> T1-2			CHART N <sub>0</sub> T3-4		
	KM (1 <sup>st</sup> ) (SE)	KM (any) (SE)	CI (SE)	KM (1 <sup>st</sup> ) (SE)	KM (any) (SE)	CI (SE)
Dryness of mouth	56 (0.05)	55 (0.04)	0.46 (0.04)	0.55 (0.05)	52 (0.05)	0.37 (0.04)
Subcutaneous fibrosis and oedema Laryngeal oedema	0.23 (0.04) 0.49 (0.04)	0.32 (0.04) 0.50 (0.05)	0.19 (0.03) 0.42 (0.04)	0.28 (0.05) 0.55 (0.05)	0.43 (0.05) 52 (0.05)	0.18 (0.03) 39 (0.04)

022 POSTER

Patterns of local-regional recurrence in patients with head and neck cancer treated by parotid-sparing IMRT

Y. Nishimura<sup>1</sup>, K. Nakamatsu<sup>1</sup>, T. Shibata<sup>1</sup>, S. Kanamori<sup>1</sup>, R. Koike<sup>1</sup>, M. Suzuki<sup>2</sup>, K. Mori<sup>3</sup>. <sup>1</sup>Kinki University School of Medicine, Radiation Oncology, Osaka-Sayama, Japan; <sup>2</sup>Research Reactor Institute, Radiation Oncology Research Laboratory, Kumatori, Japan; <sup>3</sup>Kinki University School of Medicine, Otolaryngology – Head and Neck Surgery, Osaka-Sayama, Japan

**Background:** To analyze the patterns of local-regional recurrence in patients with head and neck cancer treated by parotid-sparing intensity-modulated radiation therapy (IMRT).

Methods: Forty-one patients with pharyngeal squamous cell carcinomas were treated by IMRT between 2000 and 2004. The mean age of the patients was 59 years old with a range from 35 to 81 years old. There were 15 nasopharyngeal cancers (NPC), 14 oropharyngeal cancers (OPC), and 12 hypopharyngeal cancers (HPC). Clinical stage (UICC, 2002) was stage I; 2, II; 10, III; 6, IVa; 20, or IVb; 3. For 20 patients with OPC or HPC, unilateral (n = 15) or bilateral (n = 5) neck dissection was performed before IMRT. All patients were treated with whole-neck RT to 46–50 Gy/23–25 fractions by IMRT, followed by boost IMRT to the high-risk clinical target volume to a total dose of 60 to 70 Gy in 30 to 35 fractions (median, 68 Gy). A median follow-up period was 17 months with a range of 3 to 47 months. Twenty-nine patients were treated with concurrent chemoradiotherapy using cisplatin (60–80 mg/m² x 2–3) for 13 NPCs or weekly docetaxel (15 mg/m²) for 16 OPCs or HPCs.

Results: By IMRT, mean doses to the contralateral and ipsilateral parotid glands could be reduced to 24.0±6.2 Gy and 30.3±6.6 Gy, respectively. Recurrence or persistent tumors in the primary site was noted in 8 patients (20%). Recurrence was noted from the center of gross tumor volume (GTV) in 4 of the 12 HPCs and 2 of the 14 OPCs, while recurrence from the planning target volume (PTV) margin was noted in 2 of the 15 NPCs. Both of the NPCs had T4 tumors, and recurrence or persistent tumor was noted at the posterior edge of the clivus or at the anterior wall of the sphenoid sinus. Residual or recurrence of neck lymph nodes was noted in 5 patients (12%), including 2 patients with neck nodes recurrence after salvage surgery for recurrent primary tumors. In 2 patients with NPCs, PTV delineation for the neck nodes was insufficient, and recurrences were noted at the posterior chain nodes and at the nodes near the parotid gland. As both the primary site and neck nodes recurrence were noted in one patient, the PTV marginal recurrence was noted in three (7%) NPCs. In three of the four marginal recurrences, keen review of the pretreatment MRI of the patients showed the involved nodes or the extension of the primary tumor at the edge of the PTV.

Conclusions: In some patients with head and neck cancer, local–regional recurrences after IMRT can be related to the insufficient delineation for the PTV. Keen evaluation of the pretreatment MRI is essential for the treatment planning of IMRT.

1023 POSTER

Phase II study of capecitabine (X) plus reirradiation in patients (pts) with recurrent squamous cell carcinoma of the head and neck (SCCHN)

<u>G. Kornek<sup>1</sup></u>, M. Burian<sup>2</sup>, L. Vormittag<sup>1</sup>, D. Radonjic<sup>3</sup>, E. Selzer<sup>3</sup>. <sup>1</sup> University of Vienna, Internal Medicine, Vienna, Austria; <sup>2</sup> University of Vienna, ENT, Vienna, Austria; <sup>3</sup> University of Vienna, Radiotherapy, Vienna, Austria

Background: X has shown substantial activity in SCCHN [Pivot et al. 2004] and is replacing 5-FU as the backbone of chemotherapy in a wide range of

solid tumours, particularly colorectal cancer, in both the metastatic and adjuvant setting. In addition, X has a unique survival benefit in breast cancer. There is substantial data to support using X in chemoradiation [Dunst et al. 2004]. This study evaluated the therapeutic impact and safety profile of reirradiation with concurrent oral X in SCCHN.

Materials and methods: Pts with locally advanced unresectable SCCHN who had prior radiation therapy (with or without chemotherapy) received X 450 mg/m² twice daily, 7 days a week. Radiation therapy was delivered as a standard fractionated regimen (2 Gy/day, 5 fractions/week) up to a total dose of 50–70 Gy. The cumulative (life-time) dose within overlapping fields was not to exceed 120–130 Gy.

Results: Currently, 12 male pts have completed treatment. Baseline characteristics were as follows: median age 56 (42–72), ECOG PS 0/1 3/9. Six pts had previous chemoradiation followed by radical resection; 4 had surgery and postoperative radiotherapy; 2 had concomitant chemoradiation (accelerated and hyperfractionated). Six pts experienced locoregional recurrences, 4 developed secondary primaries, and 2 had a regional lymph node relapse. The median time since prior radiation therapy was 38 months. Efficacy (as evaluated by WHO criteria): 3 pts had a complete response (25%); stable disease was observed in 8 pts, and tumour progression immediately after therapy was observed in 1 pt. Median duration of local control and median survival will be presented at the meeting. The most frequently observed toxicity was mucositis, which was grade 3 in only 2 pts; grade 1 or 2 skin reaction was seen in 8 pts. There was no evidence of systemic toxicity or myelosuppression, probably due to the very low dose of X. Radiation therapy was completed without delay in all pts, and X was prematurely interrupted in 1 pt due to mucositis grade 3.

Conclusions: Our preliminary data suggest that X chemoradiation is a well-tolerated and effective treatment modality for previously irradiated pts with SCCHN. Based on available data, a higher dose of X should be feasible and potentially more effective. Earlier use of X in SCCHN could also be evaluated.

1024 POSTER Do smokers really have more fun?

K. Jensen, A. Bonde Jensen, C. Grau. Aarhus University Hospital, Oncology, Aarhus C, Denmark

Background: Smoking is an important etiologic factor in the development of head and neck cancer. Smoking during radiotherapy influences survival and smoking after therapy, influences the recurrence rate and rate of second cancers. Many patients believe that "smokers have more fun" and thus better quality of life.

**Materials and methods:** A cross sectional quality of life and morbidity study was performed using EORTC C30 and H&N35 as well as the DAHANCA morbidity scoring system. The patients were attending follow up after single modality treatment with either radical radiotherapy (N = 83) or surgery (N = 33) for cancer of the larynx (N = 44), pharynx (N = 34) or oral cavity (N = 38). No data on social- or economical status or comorbidity were accessible.

Results: Fifty-two of 114 patients, with available smoking information, were registered as self reported smokers at follow up. Smoking status was not significantly correlated with any of the tumour or patient related endpoints registered on the DAHANCA forms including age, sex, tumour site, stage, time since therapy or therapy. Nevertheless smokers invariable have the lowest function scores and the highest symptoms scores in both DAHANCA and EORTC QLQ except fibrosis and HN Weight gain. This difference was significant in 20 of the 33 QoL scales, but none of the morbidity scores. The difference was apparent in both general endpoints (physical function, cognitive function, fatigue, nausea/vomiting, dyspnoea, appetite loss, constipation, diarrhoea and financial problems) and organ specific endpoints (HN pain, swallowing, senses, social eating, dry mouth, coughed, nutritional supplement, feeding tube, and weight loss). When dividing the non-smokers (N = 62) in patients admitting to smoke during initial therapy (N = 48) and never smokers (N = 14) a "dose" dependent effect could be seen with smokers having more symptoms than former smokers that had more symptoms than never smokers. The difference was significant between present and never smokers in 10 items, between former and present smoker in 10 items and between former and never smokers in 2 items

Conclusion: Smokers had a significantly reduced score of many items of the EORTC C30 and H&N35 quality of life questionnaires. Smoking was not correlated with any of the disease, patients or treatment related factors registered. There was a clear tendency towards a dose effect with previous smokers constituting an intermediate group between present-smokers and never-smokers. These findings support the recommendation that head and neck cancer patients should quit smoking and indicate that there could be an immediate benefit to the patient's health related quality of life of smoking cessation since former smokers did better than current smokers.