RADIATION ONCOLOGY CASE



Adaptive replanning of IMRT for head and neck cancer: A case report of replanning in a middle-aged patient with squamous cell carcinoma of the tonsil

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CASE SUMMARY

A 39-year-old male presented with a right-sided neck mass and sore throat. The patient was placed on antibiotics initially with no resolution in symptoms. The neck mass continued to enlarge and the patient was noted to have erythema of the right tonsil. Fine-needle aspiration of the right neck mass demonstrated atypical cells, prompting an examination under anesthesia and right-sided simple tonsillectomy for further evaluation. Pathology from the excised tonsil demonstrated a moderate to poorly differentiated, human papillomavirus (HPV)-positive, p16-positive, squamous cell carcinoma. The patient was referred to radiation oncology for definitive treatment of tonsillar bed and neck disease.

IMAGING FINDINGS

Computed tomography (CT) imaging of the neck with contrast demonstrated a 3.0-cm \times 1.3-cm \times 2.3-cm right level IIA, cystic nodal mass, and

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DIAGNOSIS

HPV-positive squamous cell carcinoma of the right tonsil

TREATMENT SUMMARY

A 7-field, step-and-shoot, intensity-modulated radiation therapy (IMRT) technique was utilized, treating to a dose of 72 Gy in 40 fractions at 1.8 Gy per fraction over 8 treatment weeks with concurrent cisplatin and 5-FU chemotherapy given in-house on weeks 1 and 4 of IMRT. Daily imageguided radiation therapy (IGRT) with cone-beam CT imaging (CBCT) ensured setup accuracy. Treatment planning margins included a 3-mm expansion from gross tumor volume (GTV) to clinical target volume (CTV-72), and a 3-mm additional expansion of CTV-72 to PTV-72 for both the initial plan and adaptive replan, making total expansion of 6 mm from GTV to PTV-72.

An adaptive radiation therapy (ART) planning CT scan in the treatment position occurred at fraction number 20 of 40, with implementation of the adapted IMRT plan starting on fraction 21. There was no interruption or delay in the patient's course of radiation during the ART replanning process.

During ART planning, CT imaging demonstrated significant anatomical and geometric changes compared to the initial pretreatment CT simulation images. Overlay of initial treatment contours and dose onto the mid-treatment CT using a rigid registration algorithm (MIM[®] Software) demonstrated significant anatomical shift in organs at risk (OAR) with associated changes in predicted delivered dose (Figure 1). Automated deformable image registration software using mutual information (MIM[®] Software) was used to aid in recontouring the



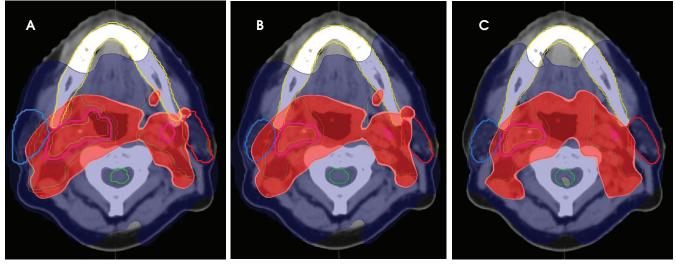
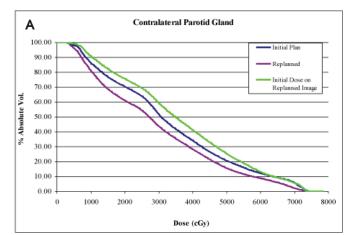
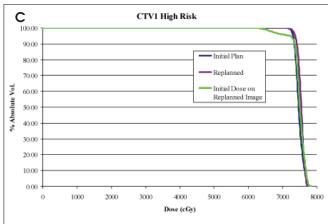


FIGURE 1. All figures contain the mid-treatment CT image obtained following fraction 20/40. (A) Initial ROI contours and dosimetry overlay the mid-treatment CT image. (B) Initial dosimetry overlies the newly re-contoured ROIs on the mid-treatment CT image. (C) Replanned ROI contours and dosimetry overlay the mid-treatment CT image.





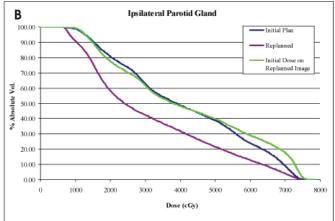


FIGURE 2. The dose-volume histograms (DVH) from the initial plan, replan, and transferred initial plan dose onto replanned image and contours are shown in (A) contralateral parotid gland, (B) ipsilateral parotid gland, and (C) CTV1 high-risk graphs.

OARs, CTV, and planning-target volumes (PTV) onto the mid-treatment CT scan images (Figure 1). Final analysis and comparison of the original IMRT and new, ART IMRT plans demonstrated a significant improvement in delivered dose to the OARs, CTVs, and PTVs (Figure 1).

DISCUSSION

Innovative techniques in radiation oncology have greatly reduced patient toxicity while maintaining, or in some cases improving, outcomes such as local-regional control and overall

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survival. IMRT has been shown to reduce overall toxicity compared to 3-dimensional radiation therapy (3DRT) in head and neck cancer.¹ Additional refinement of IMRT resulted from adaptive replanning to further reduce toxicity related to anatomical change and increased dosing to OARs. Anatomical change during treatment can result in dosimetric variations to OARs and target volumes that can be greatly improved with adaptive replanning.² Anatomical changes have been correlated with pretreatment nodal disease >4 cm, with the greatest change occurring in the first half of treatment.³ Our patient was determined to be at risk for significant anatomical change based upon the extent of nodal disease and predicted therapeutic response to chemoradiation. He was scheduled upfront, prior to the start of IMRT, for an ART, mid-treatment CT scan. New ROI contours were generated by MIM software using automated deformable image registration of the initial region of interest (ROI) contours onto the midtreatment CT followed by physician review and editing prior to ART planning.⁴ Significant anatomical change affecting delivered dose to OARs

was observed. Overall, ART planning improved the patient's therapeutic ratio by significantly reducing the maximum and mean doses to a number of OARs (Figure 2), preventing under dosing to $\mathrm{CTV}_{\mathrm{high}\,\mathrm{dose}}$ (Figure 2), and by delivering dose more conformally to CTV and PTV. ART planning has been shown to have beneficial effects on reducing chronic radiation-induced toxicity, while maintaining comparable local-regional control and survival outcomes.5 Our patient continues to be disease free 2 years post-ART planning with only mild, chronic radiation toxicity.

CONCLUSION

Our patient presented with HPVpositive, p16-positive, squamous cell carcinoma of the right tonsil with significant bulky nodal disease, placing him at risk for considerable anatomical and geometric changes during a course of definitive IMRT with concurrent cisplatin, 5-FU chemotherapy. If not accounted for, anatomical changes can result in overdosing of OARs and under-dosing of CTV or PTV target volumes, which may result in worse outcomes. The patient underwent ART planning at mid-treatment to account for the associated anatomical change. ART planning based upon midtreatment CT imaging improved this patient's therapeutic ratio by reducing delivered dose to OARs while ensuring conformal dose coverage of CTV and PTV target volumes. The patient continues to be disease free 2 years post-ART planning with minimal long-term toxicity.

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