

AbstractID: 11456 Title: A Dosimetric Analysis of IMRT Used to Replace and Mimic HDR Brachytherapy in the Treatment of Gynecologic Malignancy

Purpose: To evaluate the ability of intensity modulated radiation therapy (IMRT) to mimic the dose distribution of high-dose rate (HDR) intracavitary brachytherapy in the treatment of gynecologic malignancy, including regions of high dose in the target volume while limiting doses to organs at risk (OAR).

Methods: The HDR treatment planning CT scans of 4 patients with gynecologic malignancy were used in creating 3 different types of IMRT treatment plans. HDR dose levels were contoured and exported as volumes to the IMRT treatment planning system. The IMRT_{300%} and IMRT_{100%} plans were designed to mimic V_{300%(HDR)} and V_{100%(HDR)}, respectively. No attempt was made to spare the OAR. Results of IMRT_{300%} and IMRT_{100%} were used to determine the limitations of IMRT in order to create IMRT_{Tx}: a compromised plan, designed to replicate the V_{100%(HDR)} and the highest isodose volume possible, while sufficiently sparing the OARs.

Results: The IMRT_{300%} and IMRT_{100%} plans showed an increase in V_{50%} by a factor of 10 and 2, respectively. The IMRT_{Tx} plan mimicked V_{200%(HDR)} to within a mean of 16.4% while concurrently replicating V_{100%(HDR)} to within a mean of 79.7%. An associated

mean $\frac{V_{50\%(IMRT)}}{V_{50\%(HDR)}}$ of 510% and a mean D_{2cc} increase of 61.7% and 102.7% for the bladder and rectum were realized. Moreover, for

the patients who received supplementary conventional external-beam radiation therapy (CXRT), nodal treatment was integrated with IMRT_{Tx} in order to replace CXRT. The nodal mean D_{90(IMRT)} was within 3.9% of D_{90(CXRT)}. For all patients, IMRT_{Tx} OAR doses were below the regions of interest (ROI) constraints used clinically.

Conclusion: The present study suggests that IMRT may be capable of mimicking HDR brachytherapy in the treatment of gynecologic malignancy. As V_{200%(HDR)} and V_{100%(HDR)} are mimicked, increased V_{50%} and OAR doses must be accepted. Compliance to clinical ROI constraints can be achieved.