

and the maximum total dose for all fractions to any point at 2 cm from PTV (D2cm) divided to Dpr.

### Results

Depending on the volume of PTV, 5 groups are defined and the average results with standard deviation (SD) and maximum parameter values are presented in Table1.

Table 1

V <sub>PTV</sub> (cc)	CI	SD	CI <sub>max</sub>	NHI	SD	NHI <sub>max</sub>	IDMHT (%)	IDMHT <sub>max</sub> (%)	R <sub>95%</sub>	SD	R <sub>95%max</sub>	D <sub>low</sub> /D <sub>pr</sub> (%)	D <sub>low</sub> /D <sub>pr</sub> max (%)
<1	1.43	0.17	1.60	1.28	0.04	1.33	10.00%	14.69%	7.97	2.01	11.22	26.11%	38.57%
1 to 5	1.31	0.25	1.40	1.37	0.05	1.41	4.29%	5.28%	5.36	1.30	7.37	27.52%	32.88%
5 to 15	1.25	0.12	1.37	1.36	0.12	1.53	4.01%	9.60%	4.76	1.28	5.87	43.69%	50.00%
15 to 50	1.15	0.09	1.23	1.38	0.02	1.42	2.01%	4.09%	3.98	0.32	4.44	61.00%	67.43%
>50	1.16	0.07	1.26	1.29	0.14	1.40	1.00%	2.80%	3.51	0.12	3.65	61.42%	66.20%

The NHI shows a low sensitiveness to the PTV volume, and thus remains approximately within the same limits in all cases. The CI shows a clear dependence on the PTV volume and varies from 1.16 for volumes >50cc to 1.43 for volumes <1cc. Outside the PTV, the IDMHT does not show a clear dependency on volume, but it can be postulated that it should not exceed 10% for the PTV volumes >1cc. The gradient index outside the PTV decreases with increasing volume and on the basis of the results obtained, it is easy to calculate the radius of the shell to control the gradient outside the PTV for each particular case. The normalized maximum dose to any point at 2 cm from PTV increased with PTV volume from 26% for volume <1cc up to 61% and these result can also be used in prescribing subsequent stereotactic treatments.

### Conclusion

Data base was obtained and evaluated, based on the worksheet with parameters describing the isodose distribution in stereotactic treatments. The created protocol is used to prescribe the dose in and out PTV of each new patient. This results to an increasing in optimization parameters, but it facilitates to save time and makes treatment planning evidence based.

### EP-1930 Mixed beam radiotherapy for sternum and lung treatments

S. Mueller<sup>1</sup>, T. Risse<sup>1</sup>, M.K. Fix<sup>1</sup>, S. Tessarini<sup>1</sup>, F. Mueller<sup>1</sup>, K. Zaugg<sup>1</sup>, M.F.M. Stampanoni<sup>2</sup>, P. Manser<sup>1</sup>

<sup>1</sup>Division of Medical Radiation Physics and Department of Radiation Oncology, Inselspital Bern University Hospital and University of Bern, Bern, Switzerland

<sup>2</sup>Institute for Biomedical Engineering, ETH Zürich and PSI, Villigen, Switzerland

### Purpose or Objective

Current sternum and lung treatments using VMAT suffer from a large low dose bath delivered to the lungs, heart and other normal tissue. Combining photon and electron beams for mixed beam radiotherapy (MBRT) has the potential to reduce the dose delivered to normal tissue, because of the well-defined range of the electron beams without degrading the dose homogeneity in the target. This work presents an inverse treatment planning technique for MBRT and tests the given hypothesis by plan comparisons.

### Material and Methods

An inverse treatment planning technique for photon MLC based step & shoot MBRT is developed including a novel hybrid column generation and simulated annealing direct aperture optimization (DAO) algorithm. The hybrid DAO starts with an empty aperture pool and iteratively adds photon and electron apertures using the column generation algorithm. After each aperture addition, all apertures in the pool undergo a quasi-Newton weight optimization followed by a simulated annealing based simultaneous shape and weight optimization and a second quasi-Newton weight optimization. Thus, the optimizer has full freedom about the number, shapes and weights of photon and electron apertures and simultaneously optimizes them. After optimization, the deliverable dose

distribution of the apertures to be delivered with the photon MLC is calculated using Monte Carlo. MBRT plans with 50 apertures are generated for two sternum and a lung case with prescribed doses of 10, 30 and 50 Gy. Their deliverable dose distribution are compared to those of two arc VMAT plans in terms of planning target volume (PTV) dose homogeneity HI = V95% - V107%, mean dose to the lungs and the heart, D2% to the spinal cord and the low dose bath expressed as V10% of normal tissue.

### Results

Averaged over all three cases, the PTV dose homogeneity is 3% higher, mean dose to the lungs 23% lower, mean dose to the heart 11% lower, D2% to the spinal cord 36% lower and V10% of normal tissue 31% lower for MBRT plans compared to VMAT plans. The electron contribution defined as the integral dose in the PTV summed over all electron apertures is 27%, 27% and 43% for the MBRT plans determined for the first and the second sternum and the lung case, respectively.

### Conclusion

The MBRT plans outperformed the VMAT plans in all dosimetric aspects from PTV dose homogeneity, organs at risk sparing to the extension of the low dose bath. By utilizing electron apertures, the hybrid DAO is able to gain advantage over state-of-the-art photon only VMAT plans. This work was supported by Varian Medical Systems.

### EP-1931 Suitability of dynamic trajectory mixed beam radiotherapy for head and neck and brain treatments

S. Mueller<sup>1</sup>, P. Manser<sup>1</sup>, W. Volken<sup>1</sup>, D. Frei<sup>1</sup>, D.M. Aebbersold<sup>1</sup>, M.F.M. Stampanoni<sup>2</sup>, M.K. Fix<sup>1</sup>

<sup>1</sup>Division of Medical Radiation Physics and Department of Radiation Oncology, Inselspital Bern University Hospital and University of Bern, Bern, Switzerland

<sup>2</sup>Institute for Biomedical Engineering, ETH Zürich and PSI, Villigen, Switzerland

### Purpose or Objective

To demonstrate the clinical suitability of dynamic trajectory mixed beam radiotherapy (DT-MBRT) for head and neck as well as brain treatments.

### Material and Methods

A mixed photon-electron treatment technique is developed with the aim to exploit all major degrees of freedom of a conventional linear accelerator, namely the different particle types, intensity- and energy modulations and dynamic gantry, couch and collimator rotations. This is achieved by using dynamic trajectories (DTs) photon and step & shoot modulated electron beams collimated both using the photon MLC. The treatment planning process consists of several steps. Firstly, the couch and collimator rotations associated to the gantry rotation of the DTs are determined by minimizing the overlaps of the organs and risk (OARs) with the planning target volume (PTV) and by minimizing the area between a conformal MLC opening and the PTV, respectively. Afterwards, photon apertures along the DTs and electron apertures are simultaneously optimized using a simulated annealing based direct aperture optimization. Finally, the deliverable dose distribution of the electron apertures is calculated and based on this, the photon DTs are re-optimized using a finer control point resolution. DT-MBRT plans with two photon DTs, differing only by a 90° collimator rotation, and 16 electron apertures are generated for two head and neck and a brain case with prescribed doses of 66, 40 and 60 Gy and compared to VMAT plans with 5, 3 and 2 arcs, respectively. The deliverable dose distributions of the plans are compared in terms of PTV dose homogeneity HI = V95% - V107%, mean dose to the parallel OARs, D2% to the serial OARs and the low dose bath expressed as V10% of normal tissue.

## Results

Averaged over all three cases, the mean dose to the parallel OARs is 28% lower, D2% to the serial OARs is 28% lower and V10% to normal tissue is 14% lower for DT-MBRT plans compared to VMAT plans. For every case, the PTV dose homogeneity and coverage is similar for the DT-MBRT and the VMAT plan. The electron contribution defined as the integral dose in the PTV summed over all electron apertures is 42%, 32% and 40% for the DT-MBRT plans determined for the first and second head and neck and the brain case, respectively.

## Conclusion

Head and neck and brain treatments could remarkably benefit from DT-MBRT because of the large freedom for couch rotations and the targets which are at least partly superficial. Moreover, using DT-MBRT is not connected to large investments as it only exploits the degrees of freedom already provided by a conventional treatment unit. This work was supported by Varian Medical Systems.

## EP-1932 Assessment of Specific versus Combined Model Library in Knowledge Based Planning for Prostate Cancer

N. Dogan<sup>1</sup>, M. Duffy<sup>1</sup>, G. Simpson<sup>1</sup>, M. Abramowitz<sup>1</sup>, A. Pollack<sup>1</sup>, B. Bossart<sup>1</sup>

<sup>1</sup>University of Miami- Sylvester Comprehensive Cancer Center, Department of Radiation Oncology, Miami-Florida, USA

## Purpose or Objective

There may be large variations in the quality of the intensity modulated radiotherapy (IMRT) plans due to variations in experience and skills of the planners which may limit the desired critical structure sparing and target coverage. Recently, many investigations have demonstrated that the knowledge based planning (KBP) has a great potential to improve the quality and consistency of the treatment planning via KBP which utilizes a library of previously treated patient treatment plans. The main objective of this study was to assess the quality of the plans generated using a specific versus combined purpose model library for prostate cancer planning.

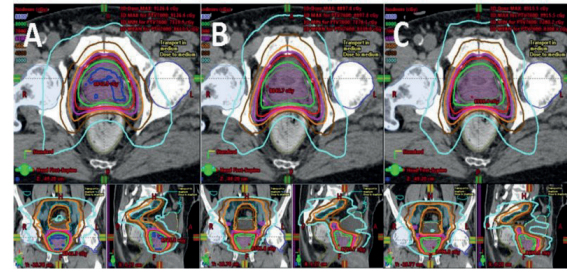
## Material and Methods

Ninety-seven prostate cancer patients were included in this retrospective study. First, three different KBP libraries were created using Eclipse RapidPlan software to benchmark KBP performance against clinical prostate IMRT plans. The original model libraries consisted of patients treated to the (a) prostate alone (P\_KBP, 66 patients), (b) prostate and pelvic lymph nodes (PPLN\_KBP, 31 patients), and (c) a model library combining the patients in model libraries (a) and (b) (P\_PPLN\_KBP, 97 patients). The number of dosimetric outliers in each library was identified and re-planned. Then, the refined P\_KBP, PPLN\_KBP and P\_PPLN\_KBP libraries which include replanned plans were created. Both original and refined three model libraries were validated on an independent set of ten patients treated to the prostate alone and ten patients treated to the prostate plus pelvic lymph nodes. All plans were normalized such that 96% of the prostate planning target volume (PTV) received 100% of the planned dose. All P\_KBP, PPLN\_KBP and P\_PPLN\_KBP based plans were compared against each other and clinical plans using the dose-volume constraints for targets and critical structures.

## Results

For both P\_KBP and PPLN\_KBP validation plans, no statistically significant differences ( $P > 0.05$ ) were found between plans generated by P\_KBP, PPLN\_KBP and P\_PPLN\_KBP libraries, with some critical structures being spared slightly better for one or the other model library, but no consistency as to which model library was better for any particular plan. The differences between plans

generated using original versus refined libraries were also negligible. However, there were 23% and 29% reduction in Dmax for left femur and right femur respectively using both PPLN\_KBP and P\_PPLN\_KBP libraries as compared to the manual clinical plans by an expert planner.



**Figure 1:** Treatment plans for one of the prostate plus pelvic lymph node patients included in the validation set. (A) Clinical plan by an expert planner, (B) PPLN\_KBP model library plan and (C) combined P\_KBP plus PPLN\_KBP model library plan. It is clear that there is very little difference between plans created using PPLN\_KBP library and P\_KBP + PPLN\_KBP library and the one created by the expert planner. Note that while the P\_KBP validation cases were replanned using both (b) and (c) model libraries, the PPLN\_KBP validation cases were replanned using both (b) and (c) model libraries (both original and refined libraries).

## Conclusion

This study demonstrated that no significant differences were observed between specific versus combined KBP model libraries in prostate planning. This may allow for fewer plans to be needed to create a model library. Refining model libraries did not further improve plans. Further studies are needed to evaluate benefits of combined model libraries for planning of complex sites such as head and neck.

## EP-1933 Half field VMAT for MLC leakage reduction and dosimetric impact in whole pelvis radiotherapy

H. Jang<sup>1</sup>, J.Y. Park<sup>2</sup>, M.H. Kim<sup>3</sup>, M. Chun<sup>3</sup>, O.K. Noh<sup>3</sup>, H.J. Park<sup>3</sup>, Y.T. Oh<sup>3</sup>

<sup>1</sup>Dongguk University School of Medicine, Radiation Oncology, Gyeongju, Korea Republic of

<sup>2</sup>University of Florida, Radiation Oncology, Florida, USA

<sup>3</sup>Ajou University School of Medicine, Radiation Oncology, Suwon, Korea Republic of

## Purpose or Objective

Recently, intensity modulated radiotherapy (IMRT) and volumetric modulated arc therapy (VMAT) techniques have been widely applied in patients with large irradiation field, such as whole pelvis radiotherapy (WPRT). However, if the irradiation field is large, multileaf collimator (MLC) leakage and non-blocking phenomenon are possible to be occurred by the limitation of MLC movement. We tried to minimize these problems by using half-field VMAT (HF-VMAT) planning technique.

## Material and Methods

We compared HF-VMAT plan with full-field VMAT (FF-VMAT) and modified full-field VMAT (MFF-VMAT) plan. Ten patients, who received whole pelvis radiotherapy with inguinal field, were included in present study. Cervical, anal, and vaginal cancer patients were 4, 4, and 2, respectively. The prescribed dose was 50 Gy (25 x 2 Gy). The normal organ dosimetric parameters for small bowel, bladder, rectosigmoid and femur head were compared according to radiotherapy planning technique. Normal tissue complication probability, conformity number (CN), and homogeneity index (HI) were also evaluated. In addition, we applied a modulation index (MI) value to support the superiority of the dose distribution by evaluating the MLC movement, gantry rotation, and dose rate.

## Results

Mean small bowel dose of HF-VMAT plan was significantly lower than FF-VMAT plan (29.6 vs 32.9,  $p < 0.05$ ), and V30 and V40 to small bowel were also significantly lower (V30: 46.4 vs 21.4, V40: 21.4 vs 28.7,  $p < 0.05$ ). Mean bladder dose of HF-VMAT plan was significantly lower than FF-VMAT and MFF-VMAT plan (33.6 vs 40.4 vs 37.2,  $p < 0.05$ ), and V30 to bladder were also significantly lower (62.6 vs 89.2 vs 86.2,  $p < 0.05$ ). There was no statistically significant differences in rectosigmoid. HF-VMAT showed