Dosimetric evaluation of dose calculation uncertainties for MR-only treatments of pelvic MRgRT

I. Coric¹, K. Shrestha³, T. Roque³, N. Paragios^{3,4}, D. Zips², D. Thorwarth¹, M. Nachbar¹

¹ Section for Biomedical Physics. Department of Radiation Oncology. University Hospital and Medical Faculty. Eberhard Karls University Tübingen, Tübingen. Germany

²Department of Radiation Oncology. University Hospital and Medical Faculty. Eberhard Karls University Tübingen. Tübingen. Germany

³ TheraPanacea. Paris. France

⁴CentraleSupelec. University of Paris-Saclay. Gif-sur-Yvette. France

Purpose:

MR-guided radiotherapy (MRgRT) allows plan adaptation on the MRI of the day. However, the current workflow for the 1.5 T MR-Linac (Elekta AB, Stockholm, Sweden) employs a structure specific bulk electron density (ED) overwrite derived from a planning CT for the calculation of dose distribution. This workflow does not only introduce uncertainties due to assignment of mean EDs, but still requires a planning CT. In this work, we investigated the uncertainty of the current patient specific (PSCT) dose calculation in contrast to the correct calculation on a CT and compare to MR-only workflows using population based bulk ED (PBCT) and artificial generative adversarial neural networks (GANs) intelligence-based generated pseudo CTs (AICT).

Methods:

Ten primary prostate cancer patients treated on the MR-Linac were chosen, based on best visual congruence between the planning CT and daily MRI. Treatment plans (20x3Gy) were optimized on the planning CT with the clinical TPS (Monaco 5.4). The CT dose distribution was then compared to the different synthetic CT approaches. (1) For PSCT, mean ED for femur, pelvis, sacrum, rectum, bladder, and patient were assigned based on mean CT densities. (2) Population-based mean EDs were derived based on 50 recent patient datasets and assigned to the structures for the PBCT approach. (3) A pseudo-CT AI-model was generated using end-to-end ensembled self-supervised GANs endowed with cycle consistency on a dataset of 42 patients with each one planning CTs and in average 5 T2w-MRIs. This model was used to create AICTs of the T2w-MRIs for dose calculation. For the analysis the planning CT was registered to the MRI, structures rigidly propagated, and the treatment plan recalculated on the correlating isocenter. For an evaluation differences in DVH-parameters were analyzed and dose distributions compared with global gamma criteria.

Results:

All three approaches reproduced for all patients the dose distribution based on a gamma criterion of 3mm/3% (40% threshold) with a pass rate greater than 98%. On a 2mm/2% criterion (40% threshold) a mean gamma pass rate for PSCT, PBCT and AICT of 96.53%, 96.32% and 96.9% was present (Fig. 1). The same is observed in terms of DVH-parameters (cf. table 1). Whilst the PTV/CTV parameters were better reproduced by the bulk density approach, differences for OARs were smaller for the AICT

Conclusion:

We presented a detailed analysis of uncertainties of pCT concepts for pelvic MRgRT. The current workflow depicts dose uncertainties based on bulk density approaches. Both a PBCT and AICT approaches, which bypass the need for a planning CT, might be considered clinically acceptable whilst reducing imaging dose and registration issues. Especially for air influenced OARs the AI-based pseudo-CT generation approach shows a better performance by more accurately predicting the different density gradients.

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Figure 1: Exemplary visualization of the dose distribution on the left for (A) the CT, (B) PSCT, (C) PBCT and (D) AICT with their corresponding gamma comparison (2mm/2%) to the CT. Dose was scaled to the evaluated 40% dose threshold.

Structure	DVH- parameter	Median Difference CT minus PSCT	Median Difference CT minus PBCT	Median Difference CT minus AICT
ΡΤΥ	D _{98%} [Gy]	-0.27 [-0.980.04]	-0.29 [-1.06 – 0.06]	0.52 [0.2 – 0.79]
	D _{2%} [Gy]	-0.11 [-0.54 – 0.07]	-0.14 [-0.57 – 0.25]	0.54 [0.36 – 1.18]
	D _{median} [Gy]	-0.27 [-0.74 – 0.03]	-0.23 [-0.79 – 0.27]	0.49 [0.32 – 1.08]
сти	D _{98%} [Gy]	-0.28 [-0.98 – 0.08]	-0.36 [-1 – 0.22]	0.46 [0.16 – 0.82]
	D _{2%} [Gy]	-0.14 [-054 – 0.12]	-0.13 [-0.87 – 0.25]	0.54 [0.3 – 1.2]
	D _{median} [Gy]	-0.3 [-0.86 – 0.02]	-0.25 [-0.96 – 0.18]	0.43 [0.29 – 1.2]
Rectum	V _{24,4Gy} [%]	-0.22 [-4.16 – 1.84]	-0.48 [-4.49 – 1.21]	0.34 [-2.7 – 1.90]
	V _{40,5Gy} [%]	-0.66 [-4.02 – 0.78]	-1.10 [-4.29 – 0.3]	-0.1 [-2.85 –1.37]
	V _{56,8Gy} [%]	-0.82 [-3.02 – 0.72]	-1.35 [-2.66 – -0.14]	0.18 [-1.13 – 2.71]
Bladder	V _{48,7Gy} [%]	1.18 [0.37 – 1.99]	1.35 [0.65 – 1.96	0.78 [-0.23 – 2.3]
	V _{56,76Gy} [%]	0.84 [0.05 – 1.41]	0.9 [0.33 – 1.87]	0.4 [-0.03 – 2.85]
PenileBulb	V _{40,5Gy} [%]	-4 [-16.9 – 2.56]	-3.89 [-17.24 – 0.71]	-0.14 [-5.94 – 13.82]

Table 1 DVH evaluation of the differences in dose distribution. Shown is the absolute median [range] difference against the CT based calculation for the evaluated approaches.