

Abstract

Implementation and Validation of a Software for Peak Skin Dose Calculation for a Fluoroscopy Equipment

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Purpose: An exceeding peak skin dose (PSD) in fluoroscopy procedures is associated with deterministic effects in skin, hence its assessment is required by Italian and European law for each patient. Many commercial software perform the PSD calculus and their validation is usually performed with specific gafchromic films, which are not always available. In a recent publication the American Association of Physicists in Medicine recommended the open source software Pyskindose for the calculation of skin dose maps. In this study we aimed to implement Pyskindose for a new fluoroscopy system and to estimate the associated error for future validation of dose tracking software.

Methods and materials: The configuration required the geometric and dosimetric characterization of the system, including table displacements, beam angles, table plus pad attenuation and half value layer at different voltages. Some customizations were also introduced, including the dose calculation for Cone Beam acquisitions and the construction of computational phantoms based on patients' Computed Tomography (CT) volumes from previous radiological exams. The error associated to PSD was estimated by considering the uncertainty of the source distance, HVL, table and pad attenuation and field homogeneity. In absence of suitable gafchromic films, the ones usually employed in radiotherapy were calibrated and irradiated with 4 simple exposure sequences as an additional test for previous assessments. The software was applied on 16 performed procedures and the PSD when using custom and standard phantoms were compared to the total Kerma at reference point.

Results: The estimated error of Pyskindose, to be considered when using the CT custom phantoms, was of 9%, which is comparable to the documented uncertainty associated to gafchromic films for fluoroscopy. This result could not be reproduced by the used gafchromic films, whose uncertainty was of 15%. However, the experimental maps reflected what was previously found with RDSR evaluation, i.e. differences inferior to 0.5cm in table displacements and beam collimation. The highest average dose difference in the considered regions of interest was of 19%. The maximum

observed discrepancy for the use of the standard phantom instead of the custom one was of 20%, observed in patients with large body size and angled procedures; while the total Kerma at reference point differed from PSD of at most 35% for procedures performed at a nearer distance from the source.

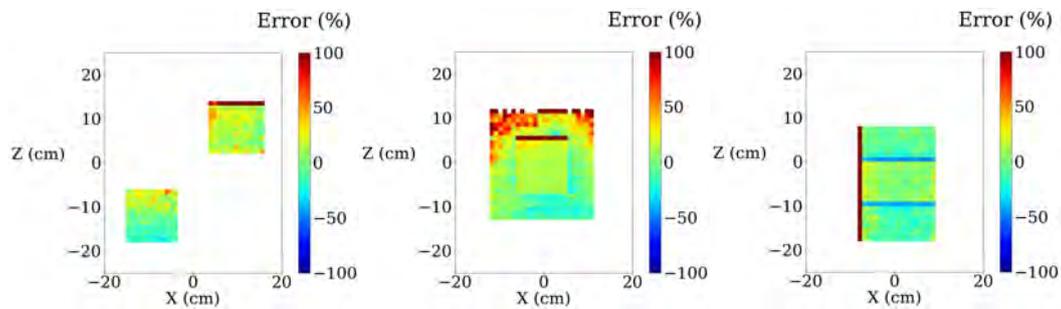


Figure 1: Difference maps between gafchromic films and Pyskindose software.

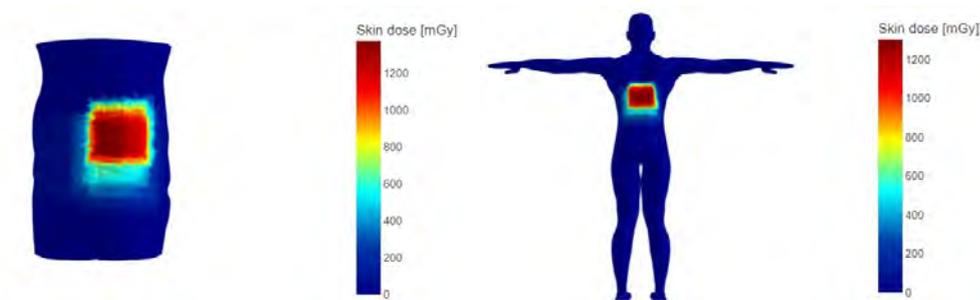


Figure 2: Skin dose maps using CT phantom (on the left) and the standard Pyskindose phantom (on the right).

Conclusion: Pyskindose was deemed accurate enough to perform the validation of new dose tracking software and to be employed in critical cases to increase the quality of the evaluation.

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