

# Commissioning an Anthropomorphic Spine and Lung Phantom for Remote Dose Verification of Institutions Participating in RTOG 0631

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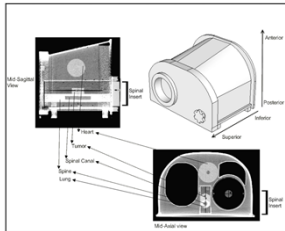
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## Introduction

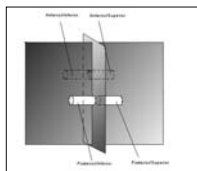
The RPC developed a new phantom to ensure comparable and consistent radiation administration in spinal radiosurgery clinical trials, particularly the Radiation Therapy Oncology Group protocol 0631. This study assessed the phantom's dosimetric and anatomic utility. The 'spine phantom' is a water filled thorax with anatomy encountered in spinal radiosurgery: target volume, vertebral column, spinal canal, esophagus, heart, and lungs. The dose to the target volume was measured with axial and sagittal planes of radiochromic film and thermoluminescent dosimeters (TLD). Four irradiations were administered: a four field box plan, a seven field conformal plan, a seven field IMRT plan, and a nine field IMRT plan. In each plan, at least 95% of the target volume received 8 Gy. For each irradiation the planned and administered dose distributions were registered via pinpricks, and compared using point dose measurements, isodose distributions, and gamma analyses. This gamma analysis test, along with the complementary assessments of the isodose distributions and point dose measurements, were used to determine if the spine phantom is a useful tool for the remote assessment of an institution's treatment planning and dose delivery regimen.

## Materials & Methods



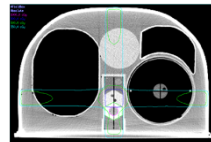
An illustration of the spinal phantom with mid-axial and mid-sagittal views from CT imaging. The anatomical features are labeled.

This image is an overview of the spine phantom design, and shows the relative location of the anatomical features within the phantom. Radiochromic film inserts bisect the spinal insert in the sagittal and axial aspects. Four TLD capsules about the axial and sagittal radiochromic film inserts within the target volume. The following image shows the relative location of the four TLD to the radiochromic film inserts.

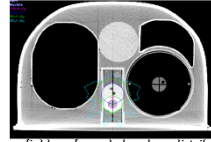


An illustration of the sagittal and axial film slices; each TLD capsule is located within the PTV, and their locations relative to the target volume center are labeled.

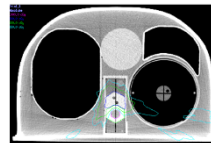
Four different treatment plans were designed in Philips Pinnacle 7.6 and administered to the spine/lung phantom: a four field box, a seven field conformal plan, a seven field IMRT plan, and a nine beam IMRT plan. 8 Gy was prescribed to 95% of the tumor volume in each administration. The following images show the relative dose distributions in each treatment plan; the dark blue contour is the 8 Gy prescription line.



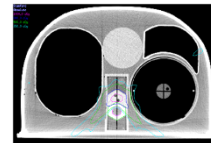
Four field box plan dose distribution



Seven field conformal plan dose distribution

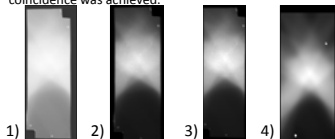


Seven field IMRT plan dose distribution



Nine field IMRT plan dose distribution

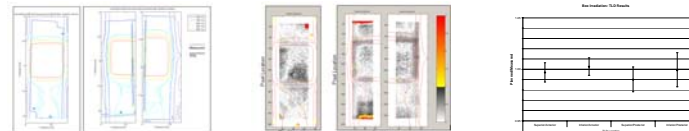
Each treatment plan was administered to the spine phantom in three trials. For each treatment administration, it was necessary to localize the physical isocenter to the radiation field isocenter with a high degree of accuracy. This was accomplished using cross patterns irradiated onto radiochromic film in the spine insert. The linear accelerator's collimator was set to a 1 mm 'slit', and the radiochromic film was irradiated. From the position of the cross pattern on the film, a shift of the treatment isocenter into coincidence with the radiation isocenter was calculated. The shift was applied to the treatment table, and this process was iterated until full coincidence was achieved.



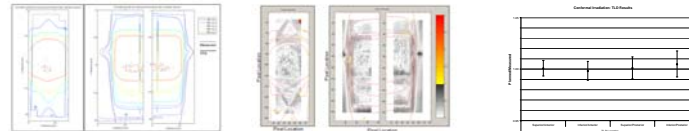
Each radiochromic film underwent the following data processing procedure: A Raw OD map was acquired through scanning (1). The OD data was then processed by the dose response curve of the film batch and smoothed with a median filter (2). The resultant dose distribution was downsampled with a bilinear interpolation, corrected to the TLD dose measurement (3), and rotated to match the pin-prick locations of the distribution from the planned dose plane (4).

## Results

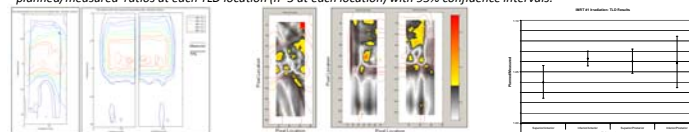
Previous RPC studies established a 95% pixel passing rate as the baseline for acceptable agreement between planned and measured dose at the 5%/3mm criteria. The following figures show a gamma analysis for one trial in the axial and sagittal planes for each treatment plan. Accompanying the gamma analyses are isodose distributions and point dose comparisons. The isodose distributions complement the gamma analyses, allowing for a qualitative assessment of the agreement between the planned and measured dose distributions. For the point dose comparisons, the ratio of the planned dose (from the TPS) to the measured TLD dose was calculated for each TLD location, and the 95% confidence intervals were calculated and plotted at each location. During the administration of the seven field IMRT plan, an error in localization was made; the phantom was shifted approximately 2.5 mm from the correct treatment position. The results from the seven field IMRT irradiations are included for illustrative purposes.



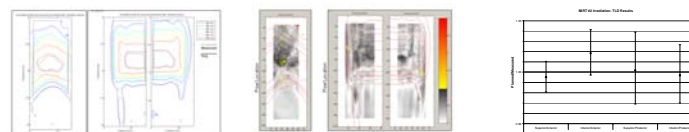
The isodose distribution and gamma analysis map from the first trial of the four field box irradiations, and the planned/measured ratios at each TLD location (n=3 at each location) with 95% confidence intervals.



The isodose distribution and gamma analysis map from the first trial of the seven field conformal irradiations, and the planned/measured ratios at each TLD location (n=3 at each location) with 95% confidence intervals.

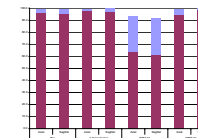


The isodose distribution and gamma analysis map from the first trial of the seven field IMRT irradiations, and the planned/measured ratios at each TLD location (n=3 at each location) with 95% confidence intervals.



The isodose distribution and gamma analysis map from the first trial of the nine field IMRT irradiations, and the planned/measured ratios at each TLD location (n=3 at each location) with 95% confidence intervals.

All trials of the four field box, seven field conformal, and nine field IMRT plans exceeded 95% pixel agreement in the axial and sagittal dose planes in gamma analysis at the 5%/3mm criteria. The seven field IMRT irradiations failed at these criteria, but with a known localization error of 2.5 mm. The ratio of planned to measured dose at the TLD locations for the four field box, seven field conformal, and nine field IMRT plans are not significantly different from unity at the 5% confidence level. The isodose distributions further illustrate the close agreement of the planned and measured dose distributions. Again, the seven field IMRT plan is the exception, with a notable shift on the isodose distribution. The adjacent chart shows the mean pixel agreement (average of the trials) for each plan in each dose plane. A tighter gamma criteria was also used in this chart: 3%/2 mm. For the four field box, seven field conformal, and nine beam IMRT, the pixel passing rate still exceeds 95%, while the passing rate for the seven field IMRT plan dropped off precipitously. This tighter criteria was explored to demonstrate that a higher standard may be feasible.



The mean percentage of pixels passing the gamma criteria for each dose plane of each irradiation; the 5% / 3mm criteria and a tighter criteria, 3% / 2mm, are shown. The mean was taken of all 3 trials from each dose plane.

## Conclusion

The spine phantom is to be used to test institution's ability to scan, plan, and administer a stereotactic radiosurgical treatment. The dosimetric utility of the spine phantom was tested using a variety of irradiation plans, from unmodulated beams to clinically applicable IMRT plans. This project assessed the dosimetric agreement between treatment planning and the TLD/radiochromic film system implemented in the phantom. This project demonstrated the dosimetric utility of the spine phantom for unmodulated and IMRT treatment plans at radiological dose levels. This project confirmed that the phantom is useful for assessing institutions participating in spinal radiosurgery protocols. The spine phantom provided a useful model for planning intensity modulated radiosurgery for spinal tumors.

## References

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 Low, D. A., W. B. Harris, S. Matic and J. A. Purdy (1998). "A technique for the quantitative evaluation of dose distributions." *Med Phys* 25(5): 656-61.  
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