

Development of test cases for comparisons of model-based dose calculations in low-energy brachytherapy



EM Elechart, F Ballester², L Beaulieu³, H Morrison⁴, A Poher³, MJ Rivard⁶, R Sloboda⁶, J Vijande², RM Thomson¹ (1) Carleton University, Ottawa, ON, CA, (2) University of Valancia (UV-IFIC-IRIMED), Burjassot, ES, (3) CHU de Quebec - Universite Laval, Quebec, QC, CA, (4) Tom Baker Cancer Centre, Calgary, AB, CA, (5) Rhode Island Hospital / Brown University, Providence, RI, USA, (6) Cross Cancer Institute, Edmonton, AB, CA

AIM

To outline a systematic framework for developing low-energy brachytherapy test cases to assist in comparing and benchmarking model-based dose calculation algorithms (MBDCAs) and implement it for the case of eye plaque brachytherapy

INTRODUCTION

 Brachytherapy dose distributions are calculated by summing the contributions from single seeds in a large water phantom (TG-43 formalism)

 TG-186 recommends adoption of MBDCAs which can account for patient geometry and tissue composition, source and applicator materials, radiation scatter, and more

 Low-energy brachytherapy is particularly sensitive to these factors due to the dominance of photoelectric interactions and their sensitivity to material cross-sections

METHODS

- Dose calculations were performed using four Monte Carlo (MC) codes for an eye plaque brachytherapy scenario
- Test cases were developed following the framework outlined in Fig. 1, progressing from simple to complex
- Local percentage dose difference was used to compare results:

$$\%\Delta D_{local} = \frac{D(r) - D_{ref}(r)}{D_{ref}(r)} \times 100\%$$
 (1)

where D(r) is the dose to the voxel at r from the MC being compared, and $D_{rel}(r)$ is the reference dose in the same voxel, here taken to be that calculated using egs_brachy



FIG 1. Framework for development of test cases

EYE PLAQUE TEST CASES

- 1) Single seed in water
- 2) Superposed seeds in water positioned as they would be in the plaque (TG-43)
- 3) Same as 2 but with interseed attenuation effects included
- 4) Seeds in water with full plaque and insert
- 5) Full plaque and insert in realistic eye phantom

RESULTS & DISCUSSION

- Local % dose difference was within ~2.5%
- Agreement was best for Test Case 1 and disagreement increased as the eye plaque and heterogeneous eye media were introduced
- Differences seen in simpler test cases can be used to explain differences seen in more complex ones – in Fig. 2, the single seed dose distribution (Test Case 1) can be used to explain the dose distributions in the more complex test cases
- Progression of test cases allows for troubleshooting and disentangling of user errors from fundamental differences between MBDCAs
- Although this work focused on ¹²⁵I eye plaque brachytherapy, the framework can readily be extended to other treatment sources and sites

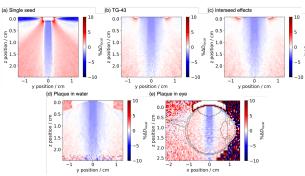


FIG 2. Local percent dose differences $(\% \Delta D_{local})$ as a function of position in the yz-plane for Penelope relative to egs_brachy for the five test cases outlined in this work

CONCLUSIONS

- Test cases for eye plaque brachytherapy were established and four MC codes were compared
- The framework established here provides a starting point for comparisons and can be extended to other applications where published benchmarked data are lacking

REFERENCES

- [1] Nath R et al. Dosimetry of interstitial brachytherapy sources: recommendations of the AAPM Radiation Therapy
- Committee Task Group No. 43. American Association of Physicists in Medicine. Med Phys. Feb 1995;22(2):209-34 [2] Rivard MJ et al. Update of AAPM Task Group No. 43 Report: A revised AAPM protocol for brachytherapy dose
- calculations. Med Phys. Mar 2004;31(3):633-74
- [3] Beaulieu L et al. Report of the Task Group 186 on model-based dose calculation methods in brachytherapy beyond the TG-43 formalism: Current status and recommendations for clinical implementation. Med Phys. 2012;39(10);6208-6236.

ACKNOWLEDGEMENTS

The authors acknowledge support from the Natural Sciences and Engineering Research Council of Canada (NSERC) [funding reference number 06267-2016], Canada Research Chairs (CRC) program, an Early Researcher Award from the Ministry of Research and Innovation of Ontario, the Queen Elizabeth II Graduate Scholarships in Science and Technology, the Ontario Graduate Scholarship, the Kiwanis Club of Ottawa Medical Foundation and Dr. Kanta Marwah Scholarship in Medical Physics, and the Carleton University Research Office.

CONTACT INFORMATION liz.fletcher@carleton.ca