Personalized Dosimetry in Radioembolization: A Paradigm Shift in Interventional Oncology



WEXNER MEDICAL CENTER

Mustaqueem Pallumeera, BS,¹ Marcus Hong, BS,¹ and Mina S Makary, MD²

The Ohio State University College of Medicine, Columbus, OH; ² Department of Radiology, The Ohio State University College of Medicine, Columbus, OH.

Purpose

- Radioembolization with yttrium-90 (Y-90) microspheres is a cornerstone therapy for hepatocellular carcinoma (HCC).
- Historically, treatment planning was standardized, ignoring patient- and tumor-specific variability.
- Recent advances in personalized dosimetry aim to optimize therapeutic efficacy while minimizing toxicity.
- This abstract highlights the clinical value of tailored dose planning and its implications for outcome improvement in interventional oncology.

Materials and Methods

• The primary methodology employed in this educational exhibit involves a comprehensive evidence- and case-based PubMed literature review of published outcomes to date.

Results

- Several prospective studies have demonstrated that tailoring the Y-90 dose based on tumor volume, location, and absorbed radiation can significantly improve outcomes.
- The DOSISPHERE-01 trial compared standard dosimetry (uniform dose of 120 ± 20 Gy) versus personalized dosimetry targeting ≥205 Gy to the tumor.
- Imaging techniques like pre-treatment 99mTc-MAA SPECT/CT and post-treatment Y-90 bremsstrahlung or PET/CT were integrated to assess dose distribution and treatment efficacy.^{2,3,5}
- In DOSISPHERE-01, the personalized dosimetry arm showed a significantly higher median overall survival (26.6 vs 10.7 months; HR, 0.42; P = .0096) and better tumor response rates.²
- Similarly, a secondary analysis of the SARAH trial demonstrated that an absorbed tumor dose ≥100 Gy was associated with longer median survival (14.1 vs 6.1 months; HR, 0.38; P < .001).
- When imaging correlation and tumor targeting were optimized, survival extended to 24.9 months.^{3,5}
- The SIRveNIB trial also supported selective internal radiation therapy (SIRT) as a viable alternative to systemic therapies like sorafenib in Asian populations.4
- These studies collectively emphasize that personalized dosimetry can turn a marginal treatment into a transformative therapy.

Conclusion

- Personalized dosimetry represents a significant evolution in radioembolization, underscoring interventional oncology's shift toward precision medicine.
- Integrating patient-specific imaging, predictive modeling, and dosimetric calibration enables maximization of tumor dose while sparing healthy tissue.^{2,3,5,6}
- To gain wider clinical adoption, interventional oncology must continue to produce prospective, multicenter evidence demonstrating meaningful improvements in survival and quality of life.^{2,3,4}
- With innovation in imaging, AI integration, and tumor biology analysis, personalized dosimetry is poised to become standard practice in locoregional cancer therapy.⁷⁻⁹

References

Garin E, Tselikas L, Guiu B, et al. Personalised versus standard selective internal radiation therapy in patients with locally advanced hepatocellular carcinoma (DOSISPHERE-01): a randomised, multicentre, open-label phase 2 trial. Lancet Gastroenterol Hepatol. 2021;6(1):17-29. doi:10.1016/S2468-1253(20)30290-9

Vilgrain V, Pereira H, Assenat E, et al. Efficacy and safety of selective internal radiotherapy with yttrium-90 resin microspheres compared with sorafenib in locally advanced and inoperable hepatocellular carcinoma (SARAH): an open-label randomised controlled phase 3 trial. Lancet Oncol. 2017;18(12):1624-1636. doi:10.1016/S1470-2045(17)30683-6 Hermann AL, Dieudonne A, Ronot M, et al. Relationship of tumor radiation—absorbed dose to survival and response in hepatocellular carcinoma treated with transarterial radioembolization with yttrium-90 in the SARAH trial. Radiology. 2020:296(3):673-684. doi:10.1148/radiol.2020191606

Chow PKH, Gandhi M, Tan SB, et al. SIRveNIB: selective internal radiation therapy versus sorafenib in Asia-Pacific patients with hepatocellular carcinoma. J Clin Oncol. 2018;36(19):1913-1921. doi:10.1200/JCO.2017.76.0892 Geschwind JF. Interventional oncology: fast forward to the future. Endovasc Today. 2021;20(10):90-93. Qu S, Worlikar T, Felsted A, et al. Non-thermal histotripsy tumor ablation promotes abscopal immune responses that enhance cancer immunotherapy. J Immunother Cancer. 2020;8(1):e000200. doi:10.1136/jitc-2019-000200 Zhong BY, Ni CF, Ji JS, et al. Nomogram and artificial neural network for prognostic performance on the albumin-bilirubin grade for hepatocellular carcinoma undergoing transarterial chemoembolization. J Vasc Interv Radiol. 2019;30(3):330-338. doi:10.1016/j.jvir.2018.08.026

Johnson PJ, Berhane S, Kagebayashi C, et al. Assessment of liver function in patients with hepatocellular carcinoma: a new evidence-based approach—the ALBI grade. J Clin Oncol. 2015;33(6):550-558. doi:10.1200/JCO.2014.57.9151 Abajian A, Murali N, Savic LJ, et al. Predicting treatment response to intraarterial therapies for hepatocellular carcinoma with the use of supervised machine learning—an artificial intelligence concept. J Vasc Interv Radiol. 2018;29(6):850-857. doi:10.1016/j.jvir.2018.01.769

Contact

mustaqueem.pallumeera@osumc.edu