

Accuracy and Safety of Scout Dose Y90 for Personalized Selective Internal Radiation Therapy Planning

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Background

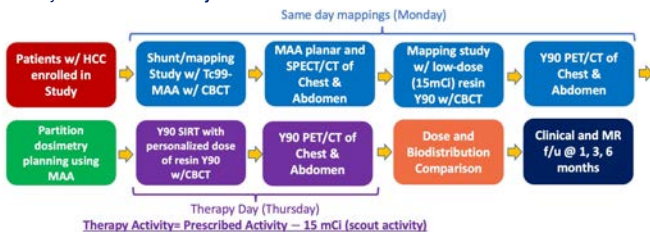
Technetium-99m macroaggregated albumin (Tc99m MAA) is an imperfect/flawed surrogate for yttrium-90 (Y90) biodistribution in the lungs and liver. Tc99m MAA can overestimate lung shunt fraction (LSF) and tumor-to-normal ratio (TNR) of activity compared to that of Y90.¹⁻⁵ This can result in inaccurate treatment planning for Y90 SIRT which result in suboptimal treatment response and treatment related toxicity.

Objectives

To evaluate the accuracy and safety of scout/low dose Y90 resin microspheres vs. Tc99m MAA in predicting the therapeutic dose Y90 biodistribution for personalized Selective Internal Radiation Therapy (SIRT) planning for hepatocellular carcinoma (HCC)

Methods

Prospective single-arm clinical trial (NCT04172714) of N=30 patients with HCC undergoing Y90 SIRT. Each patient underwent mapping angiography and MAA SPECT/CT followed by same-day mapping using 15 mCi Y90 resin microspheres and Y90 PET/CT. During MAA mapping, cone beam CT (CBCT) was used to detect and treat potential non-target embolization pre-MAA administration. Partition model was used for treatment planning using MAA data with the goal of >200 Gy to the tumors. Prescribed Y90 activity minus previously administered scout activity was administered 3 days post mappings. Same microcatheter type and administration position were used for all stages of the study (including CBCT) in each patient. Sureplan® (MIM Software, Cleveland, OH) was used for biodistribution analysis. Please see study flow chart below.



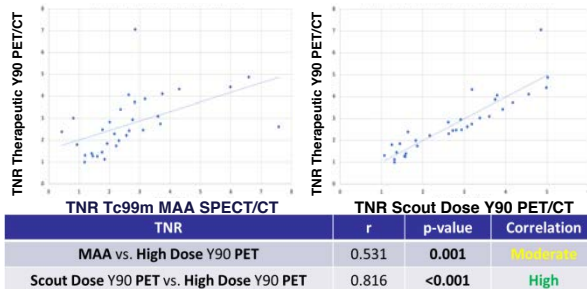
Mean LSF and TNR were compared between the MAA SPECT/CT, scout dose Y90 PET/CT, and therapeutic dose Y90 PET/CT (paired t-test). Linear correlation of LSF and TNR between Tc99m MAA and therapeutic dose Y90, as well as scout dose Y90, and therapeutic dose Y90 values were compared (Pearson's Correlation). The safety of scout dose resin microsphere was evaluated by determining non-target embolization seen on scout dose Y90 PET which was not detected and treated during mapping CBCT and discrepancy in extrahepatic activity between MAA SPECT/CT and scout dose Y90 PET/CT. Embolic effect of the scout dose was evaluated using degree of enhancement in the tumor and non-tumoral liver pre and post Y90 scout dose.

Results

N=30 patients, 33 treated tumors were treated (mean tumor volume: 44.9 cc). There was no difference for mean LSF and TNR between surrogates.

	MAA SPECT/CT	Scout Dose Y90 PET/CT	Therapeutic Dose Y90 PET/CT	p-values
Mean LSF	5.7%	6.4%	5.2%	p>0.05
Mean TNR	2.67	2.75	2.71	p>0.05

Greater linear correlation of TNR between scout dose and therapeutic dose Y90 (r=0.816), compared to MAA and therapeutic dose Y90 (r=0.531). Graphs below depict linear correlation between Y90 scout dose and MAA with therapeutic Y90 respectively.

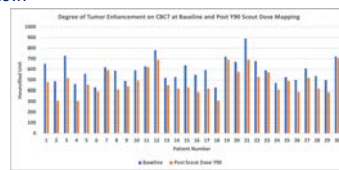


Greater linear correlation of LSF between scout dose and therapeutic dose Y90 (r=0.562), compared to MAA and therapeutic dose Y90 (r=0.394).

LSF	r	p-value	Correlation
MAA vs. High Dose Y90 PET	0.394	0.031	Low
Scout Dose Y90 PET vs. High Dose Y90 PET	0.562	0.001	Moderate

Two right gastric arteries were embolized prophylactically to prevent non-target embolization based CBCT findings during mapping. No extra-hepatic non-target embolization was seen. Specifically, there was no discrepancy between extrahepatic activity seen on MAA SPECT/CT and scout dose Y90 PET/CT. CBCT from microcatheter location for administration of scout dose Y90 predicted its distribution 100% of the time.

The degree of tumor enhancement was mildly decreased post scout dose Y90 (mean Hounsfield Unit (HU) pre scout: 589 vs. post scout: 483; p<0.001). The degree of enhancement for non-tumoral liver in the perfused area of CBCT remained unchanged between baseline (mean HU: 243) and post scout (mean HU: 233; p=0.464). Individual tumor enhancement pre and post Y90 scout dose depicted below.



Conclusions

- Scout dose Y90 is a safe and accurate surrogate for SIRT planning.
- Scout dose Y90 is more accurate than MAA for prospective therapy planning.
- 15 mCi 3-day pre-calibrated resin microsphere was safe and effective with no gross embolic effect or non-target embolization.
- Similar results should be confirmed with glass-based Y90 microspheres.

References

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