

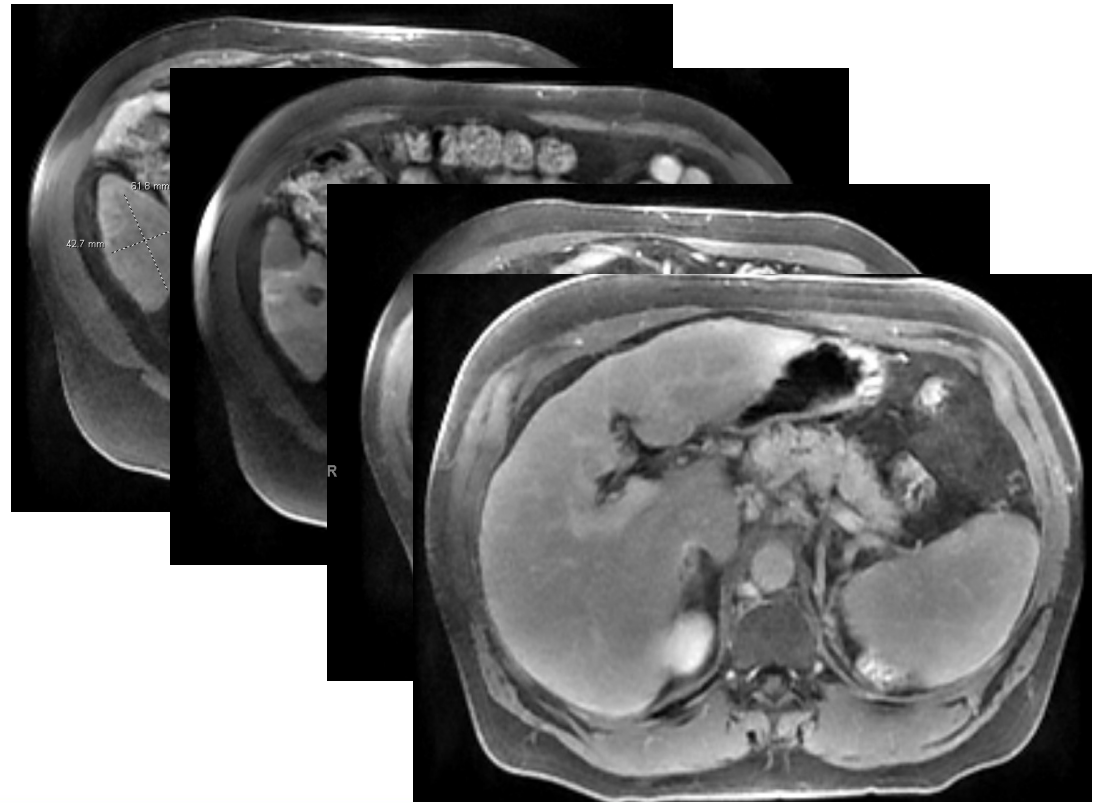
Outlining Traditional Treatment Pathway and Time to Treatment for Y90 Radioembolization

Venkatesh (Kavi) Krishnasamy, MD FCIRSE

Associate Professor of Radiology
Director of Interventional Oncology
Director of Interventional Research
University of Alabama Birmingham

Let's Start with a Case

- Imaging: Day zero
- Clinic 1 month
- 62 yo Male
- MASLD
- ALBI 2
- CP A5
- BCLC C
- ECOG 1
- AFP 5700

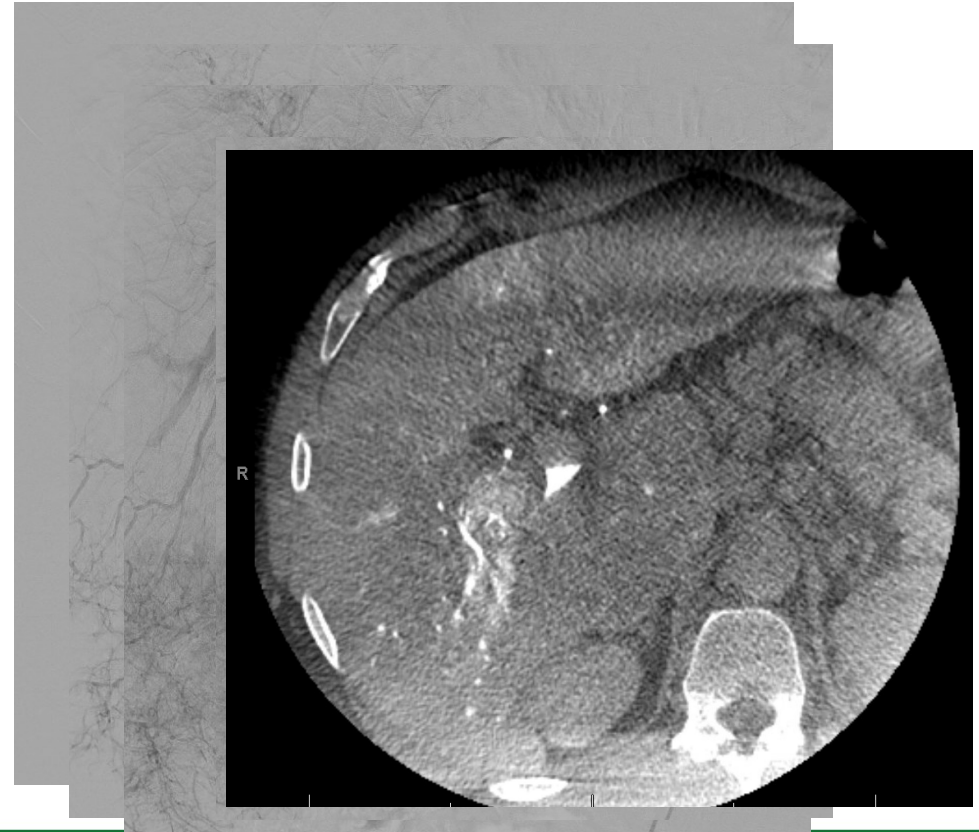


Dosing

- Mapping 1 month after clinic

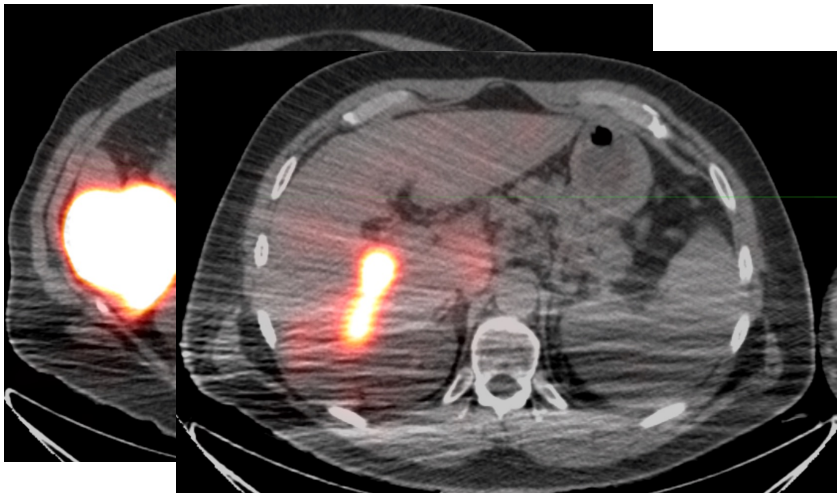
Lung Shunt and Volumes		
Lung Shunt =	3.12	%
Lungs Total Counts =	504.85	kCNTS
Liver Total Counts =	15694.3	kCNTS
Lungs Volume =	2756	cc
Liver Volume =	2356.75	cc

- Treatment 6 wks after clinic
 - First wk Tues
 - Segment 6 350cc 8gbq
 - 600gy 11gy lung
 - Segment 6 100cc 3gbq
 - 800gy 4.5gy lung
 - PVT 100cc 3gbq
 - 800gy 4.5gy lung

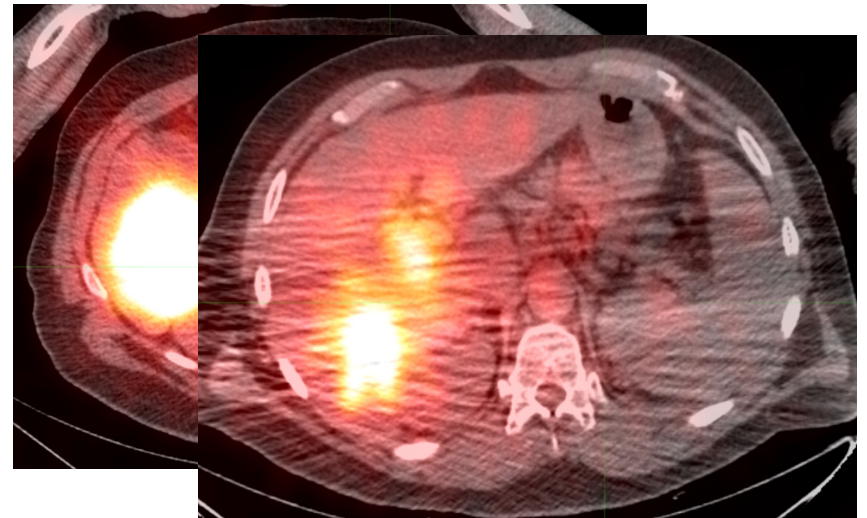


SPECT/CT

- Post MAA

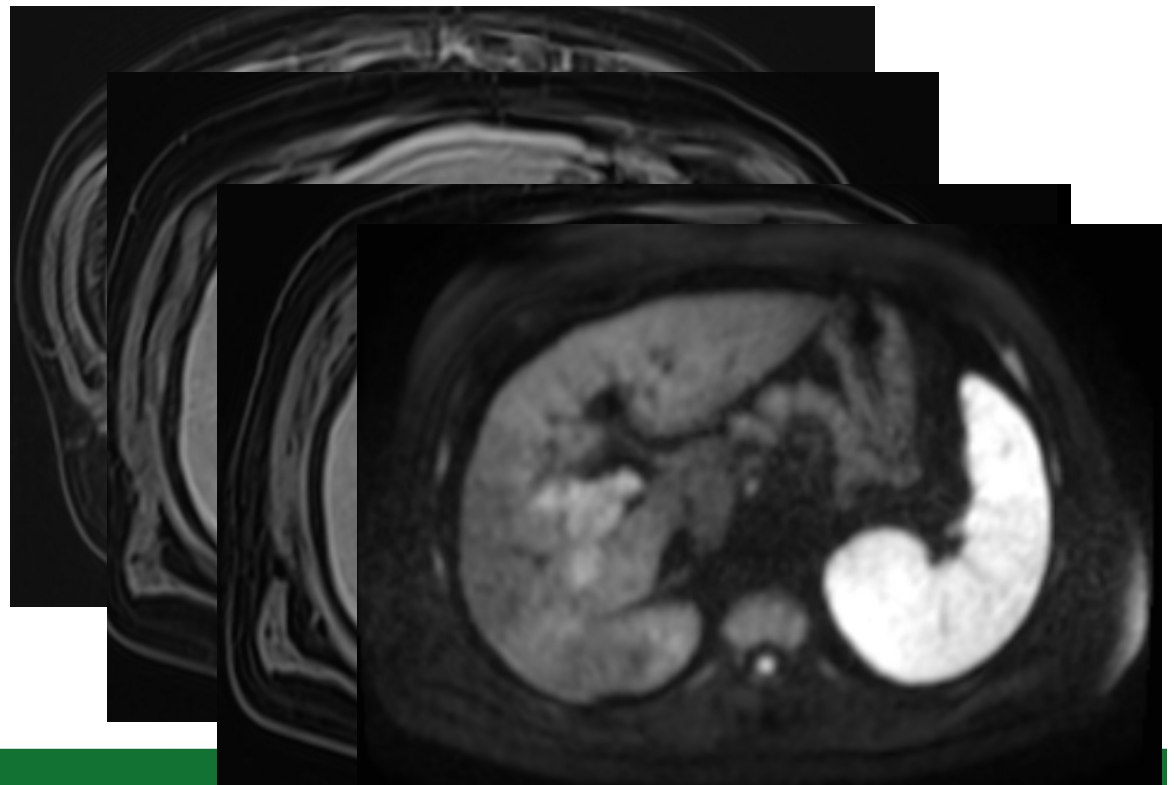


- Post Y90



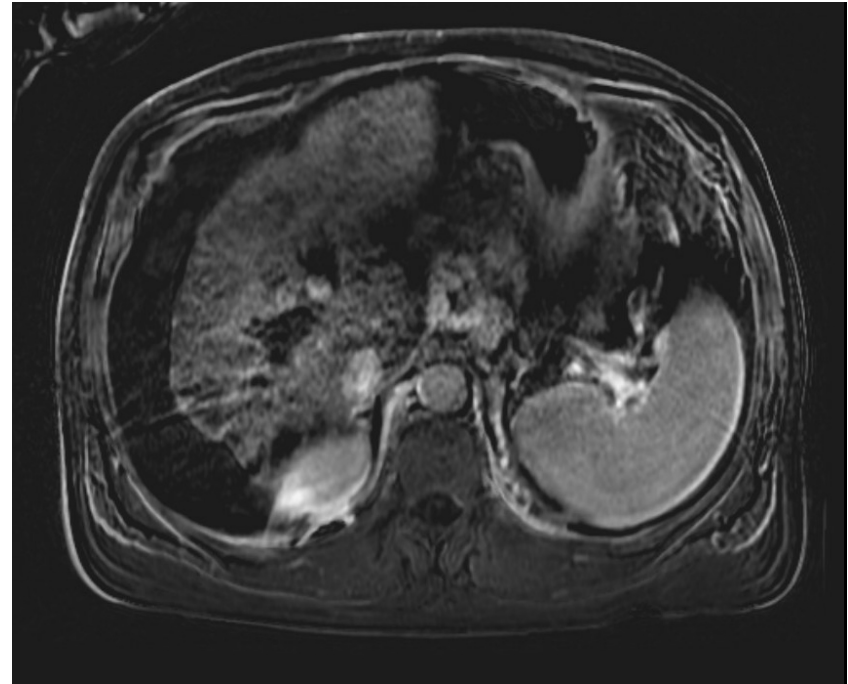
Initial Follow-Up

- 7 wks post treatment
- AFP 13,600
- CP B9
- ALBI 3



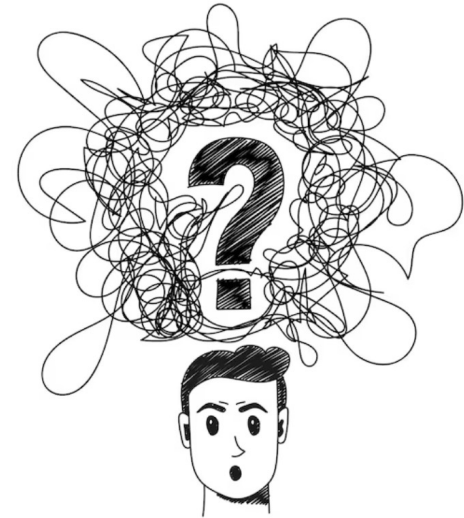
Long-Term Follow-Up

- 11 months after treatment
- AFP 350
- CP B9
- ALBI 3
- Lenvima[®] (lenvatinib)
- Recurrent paracentesis



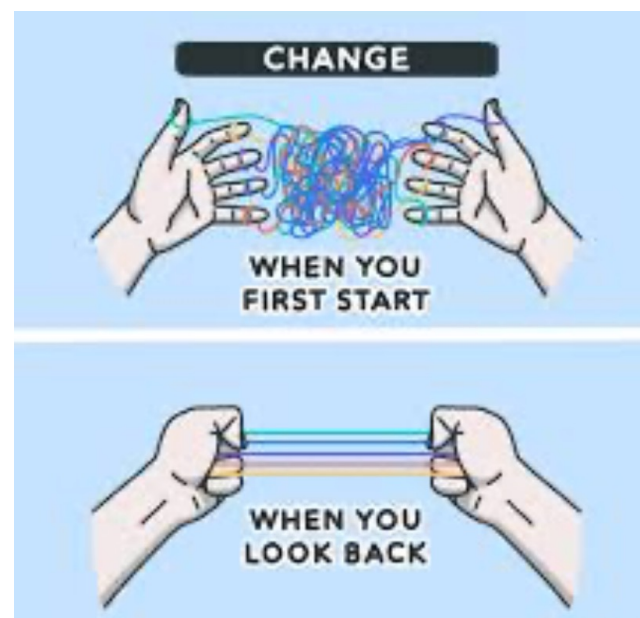
Thoughts?

- Bad biology
- Dosimetry
 - Treatment day
 - In combo with IO
- But.....
- **Treatment plans need to be individualized to the patient**

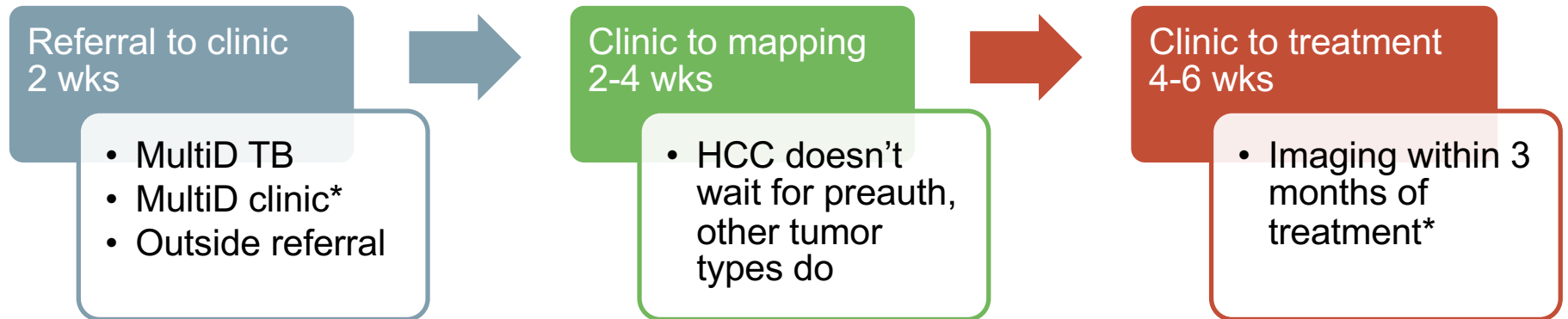


Traditional Workflow

- What was traditional is not contemporary!!!
- Not absolute
 - Treatment timeline
- But definitely for
 - Mapping
 - Dosimetry
 - Treatment
 - Follow-up



Pretreatment Workflow

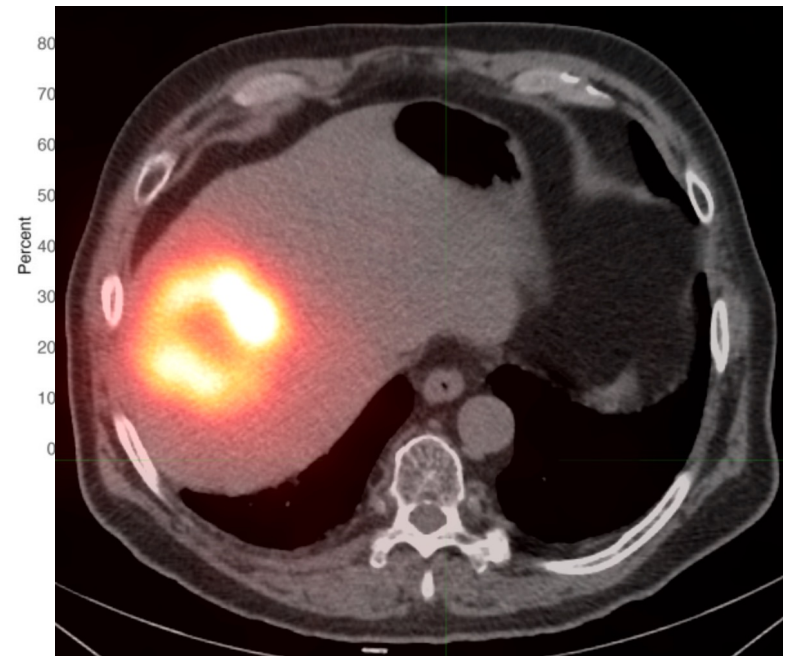


- MDTB or MTC is just the beginning of communication and collaboration
- Labs at clinic, mapping, and treatment

Weber, et al. *EJNMMI*. 2022. Mourad, et al. *Cancers*. 2024. Dahan, et al. *HC*. 2023.

Mapping Workflow

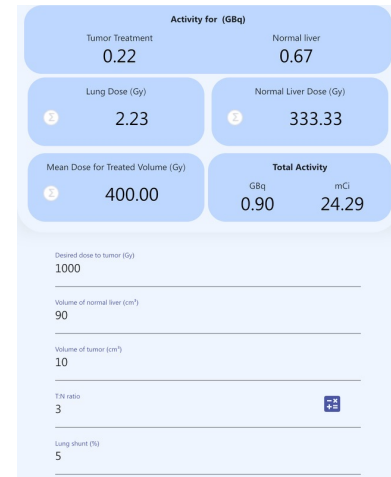
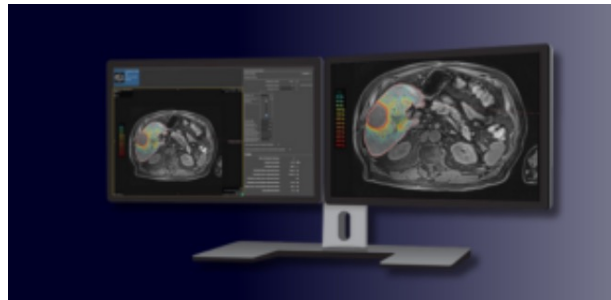
- Treatment intent
 - Curative
 - Bridging
 - Downstaging
 - Palliative
- Mapping
 - Nonselective CBCT
 - Multiple selective CBCTs
 - Every delivery spot
 - Drop the contrast rate and volume as needed
 - Volume accuracy
 - Nonselective or selective MAA



Salem, et al. *EJNMMI*. 2022.

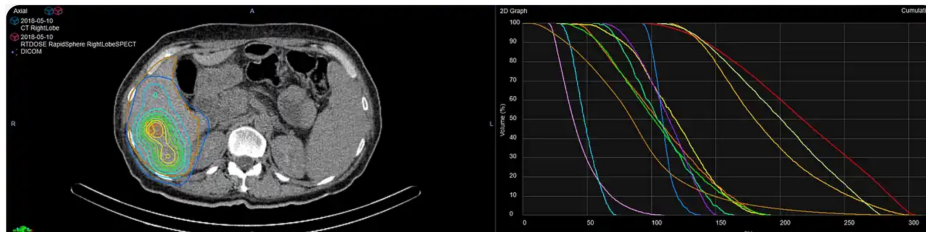
Dosimetry Software

Simplicit⁹⁰Y™
Personalised dosimetry



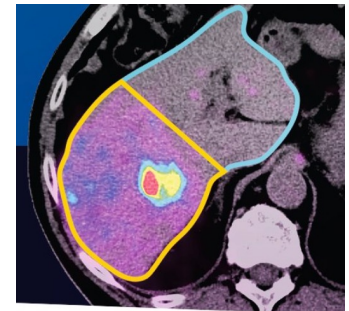
RapidSphere™

Image-guided voxel dosimetry for Y90 selective internal radiation therapy



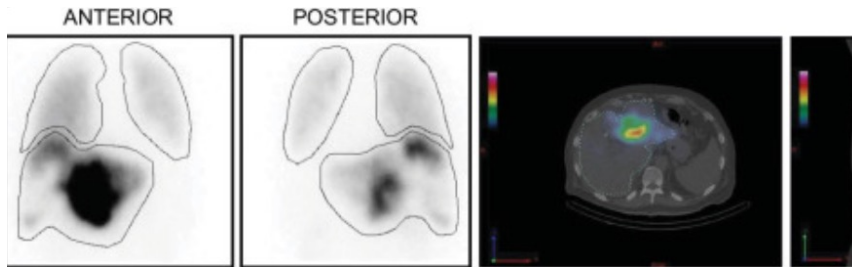
SurePlan™ LiverY90

Advanced dosimetry software for treatment of patients undergoing Y90 radioembolization



MAA Dosimetry

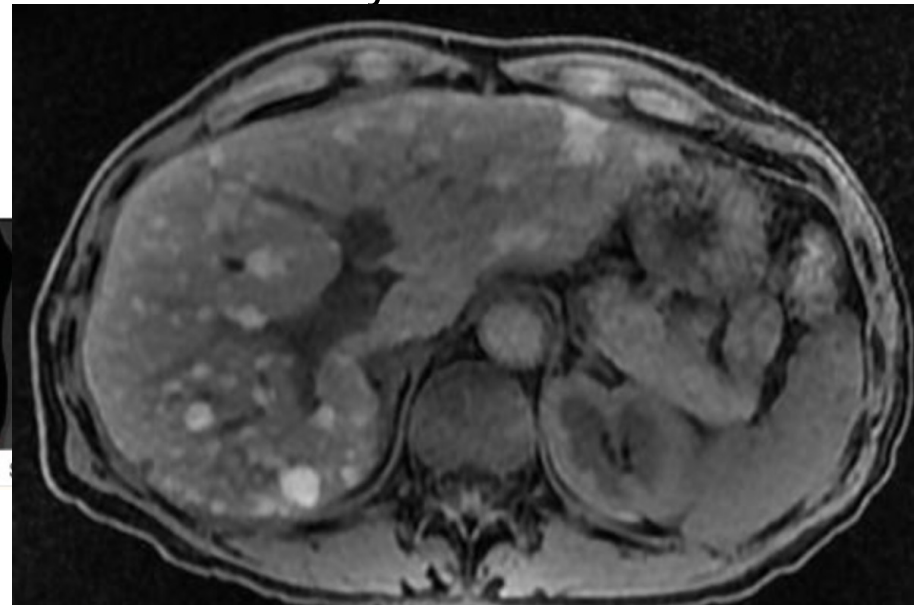
- Software – MIM
 - SPECT CT lung shunt
 - Especially over 10%
 - Whole liver volume



Planar Imaging LSF=24.2%

- Pertused volumes
 - Philips workstation

- Dosimetry



Georgiou, et al. *EJNMMI*. 2021. Elsayed, et al. *CVIR*. 2021.

MAA Dosimetry

- Partition dosimetry
 - > MIRD
 - > BSA
- Be cognizant
 - MAA to Y90 disparity

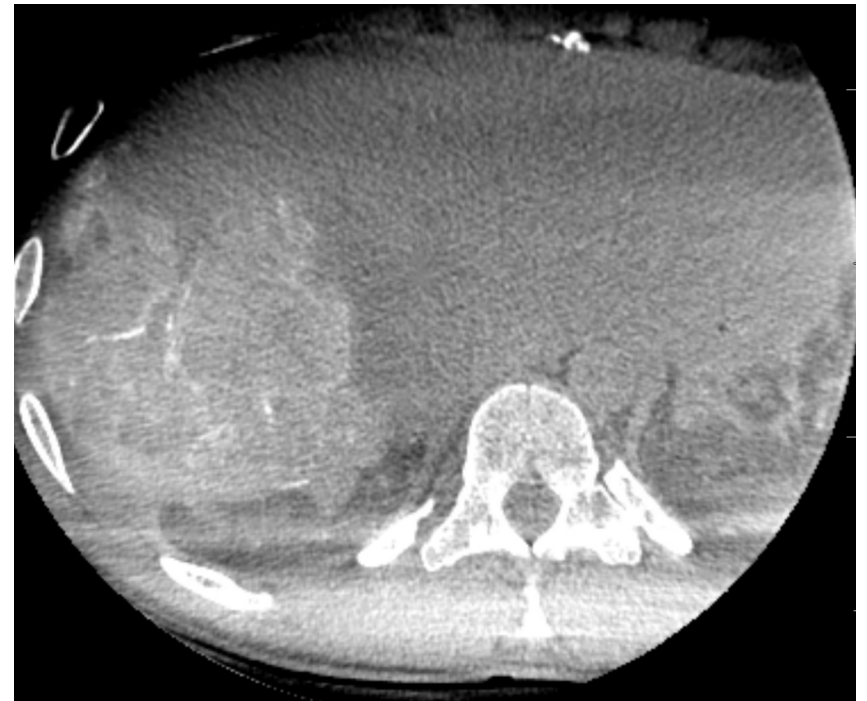
Target Region Statistics		
	l	Unit
Liver_Perfused_Target_Suffix =		
Target Region Volume =	799.67	cc
Percent of Liver Outside Target Region =	29.81	%
Percent of Liver Inside Target Region =	70.19	%
Liver Volume =	1091.18	cc
Tumor Involvement =	34.7	%
All Tumors (within Target Region) Volume =	277.52	cc
Total Tumor TNR =	3.92	:1
Tumor 1 (within Target Region) Volume =	173	cc
TNR Tumor 1 =	4.75	:1
Tumor 2 (within Target Region) Volume =	7.44	cc
TNR Tumor 2 =	1.06	:1
Tumor 3 (within Target Region) Volume =	97.08	cc
TNR Tumor 3 =	2.67	:1

Target Region Statistics		
	l	Unit
Liver_Perfused_Target_Suffix =		
Target Region Volume =	792.14	cc
Percent of Liver Outside Target Region =	33.25	%
Percent of Liver Inside Target Region =	66.75	%
Liver Volume =	1101.37	cc
Tumor Involvement =	42.35	%
All Tumors (within Target Region) Volume =	335.45	cc
Total Tumor TNR =	1.82	:1
Tumor 1 (within Target Region) Volume =	208.74	cc
TNR Tumor 1 =	2.12	:1
Tumor 2 (within Target Region) Volume =	8.8	cc
TNR Tumor 2 =	1.22	:1
Tumor 3 (within Target Region) Volume =	117.91	cc
TNR Tumor 3 =	1.33	:1

Kolligs, et al. *JHEP*. 2022. Schaefer, et al. *CCC*. 2022. Garin, et al. *Lancet*. 2021. Kappadath, et al. *JNM*. 2022. Villalobos, et al. *JVIR*. 2021. Wondergem, et al. *JNM*. 2013. Thomas, et al. *MP*. 2020. Alsultan, et al. *JNM*. 2021.

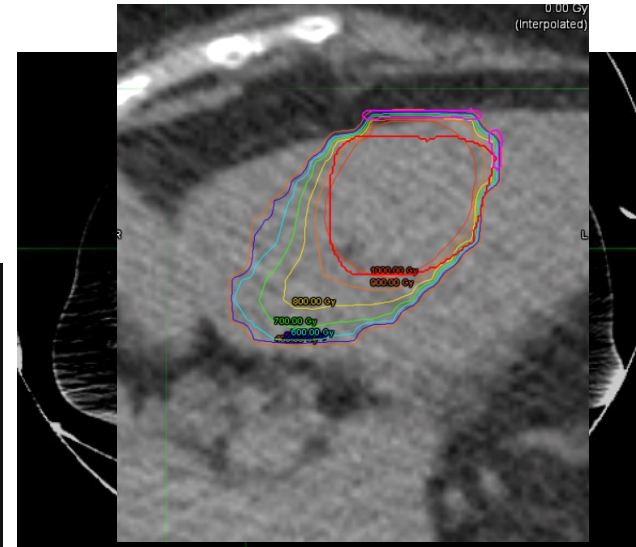
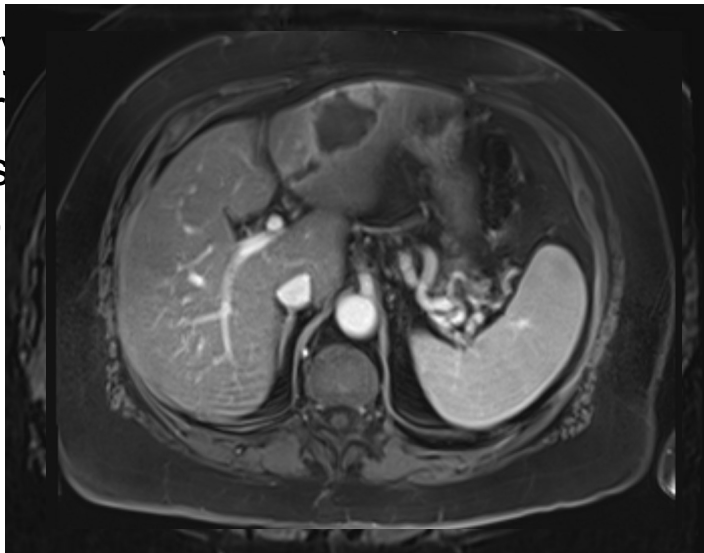
Delivery Workflow

- Replicate mapping
- Any change in plan
 - Repeat CBCT
- Something doesn't smell right
 - Repeat CBCT
- Deliver doses
 - Load catheter with contrast for glass
 - Push dose with 70/30 for resin



Y90 Dosimetry

- Post dosimetry
 - Tumor coverage
 - +/- voxel based
 - V400, D95



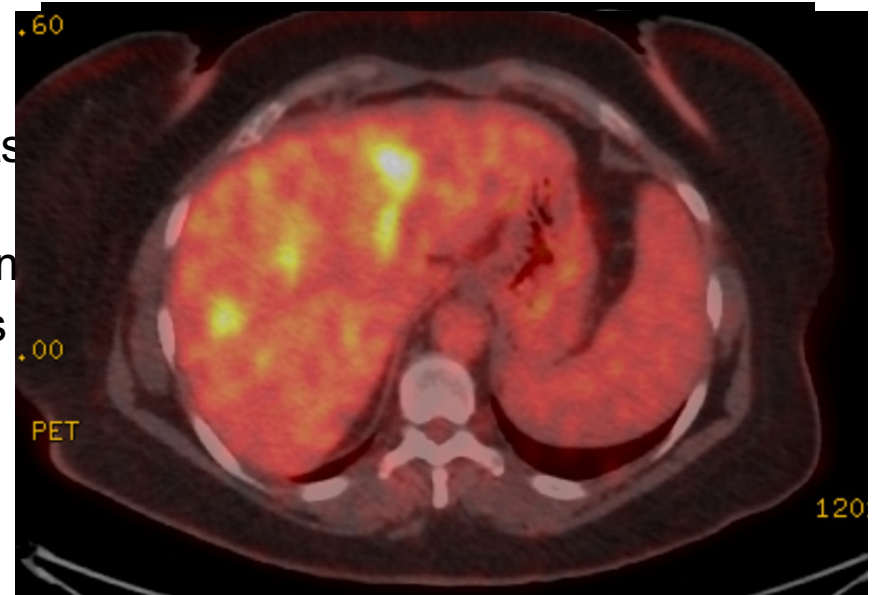
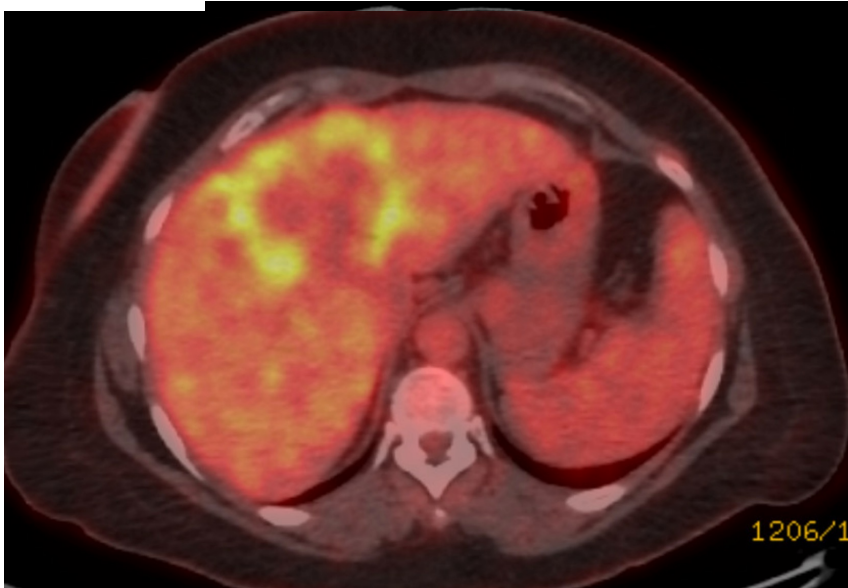
400.00	90.83	99.08
500.00	83.83	98.58
600.00	71.44	97.84
700.00	56.04	96.49
800.00	41.16	90.2

Mean dose perfused 770 Gy
 Mean dose tumor 1056 Gy

Sandow, et al. *JVIR*. 2024. Pianka, et al. *EJNMMI*. 2024.

Follow-Up Workflow

- 4-6 wks



- Must follow your patients
- Keep the referring physician in the loop

Stocker. *ER*. 2024. Weber, et al. *EJNMMI*. 2022. Prachanronarong, et al. *SIR*. 2021.

Summary

- Time to treatment is important
- Every patient is not the same
- Optimize your workflow and efficiency
 - Traditional treatment paradigm
 - Order/map/treat
 - Same day map/treat
 - Single session
 - Repeat treatment



Thank You

vkpk27@gmail.com
(330) 701-6765



SYMPOSIUM
ON CLINICAL
INTERVENTIONAL
ONCOLOGY

Understanding Resin FLEXDose Delivery Program, Order–Map–Treat

Ripal T. Gandhi, MD, FSIR, FSVM

Miami Cardiac & Vascular Institute

Miami Cancer Institute

Clinical Professor, FIU Herbert Wertheim College of Medicine



cio

Objectives

Order-Map-Treat Strategy

FlexDose Select– What is It?

FlexDosing – How I Do It

Resin Treatment Dose Thresholds

Sphere Count

Conclusions

Order–Map–Treat Strategy

Order-Map-Treat Strategy

Customized Dosing

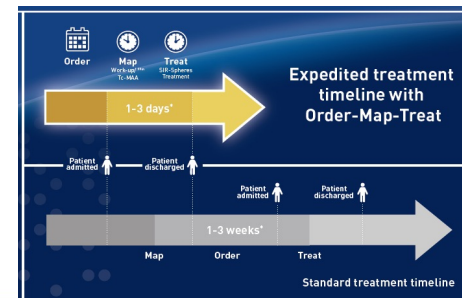
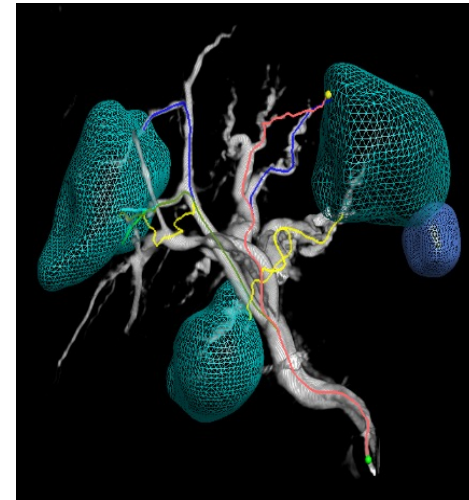
- Activity can be modified, tailored on day of therapy, and personalized to individual patient-specific needs
- Same-day activity draw

Order-Map-Treat Strategy

- Order dose in advance
- Have options of higher activity/fewer spheres and less activity/greater spheres with pre-cal doses (FLEXdose Delivery Programme)
- Same-day or same-week therapy allow for expedited treatment

Time-Sensitive Clinical Situations

- Allow for flexibility to modify treatment planning



Order-Map-Treat Strategy:

Fast Track from Referral to Treatment

For the Patient	For the Referring MD	For the Treating MD
<ul style="list-style-type: none">• Shorter hospital stay (Same-Day Map & Treat)• Reduced delays to treatment• Fewer trips to hospital (Same-Day Map & Treat)	<ul style="list-style-type: none">• Improved treatment efficiencies• Reduced interruptions between treatments• Supports continuum of care	<ul style="list-style-type: none">• Flexibility for interventional radiologists and nuclear medicine• Improved efficiencies for IR suites• Personalized dose adjustments up to time of treatment

Same Day/Same Week Y90

Resin Microspheres Can Easily Be Fractionated on Day of Procedure

- Choose appropriate FlexDose Select that you desire and order prior to mapping
- Dose can be fractionated as needed following mapping procedure and can be modified on the day of treatment as necessary

Glass Microspheres Come In Pre-Calibrated, Sealed Vials

- Dose vials must be ordered in advance
- Must do planning for anticipated treatment volume to order proper dose vials
- Cannot be fractionated

Can Perform Dosimetry After the Mapping

- Does not require much pre-procedural planning other than ordering dose
- If doing lobar Y90 or radiation segmentectomy, can calculate tumor volume and estimate liver volumes before the mapping

FlexDose Select Delivery Program

cio

FLEXDose Delivery Program

Flexibility to Deliver the Same Activity with More or Fewer Spheres

Changes:



27.3M microspheres/vial



Higher specific activity
option (5DPC)

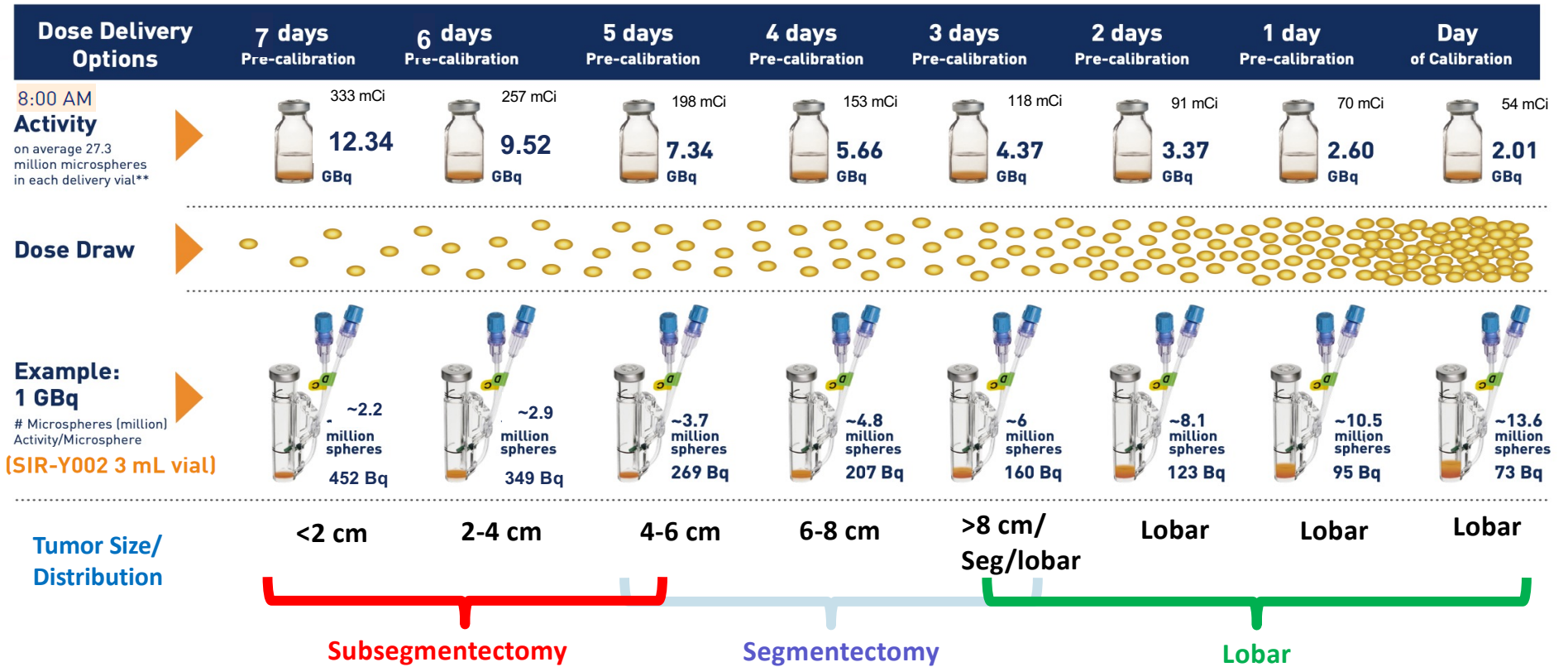
6DPC
7DPC

Will not change:

- Availability of other days of calibration
- Specific activity (per microsphere) of other vial options
- Approved indication

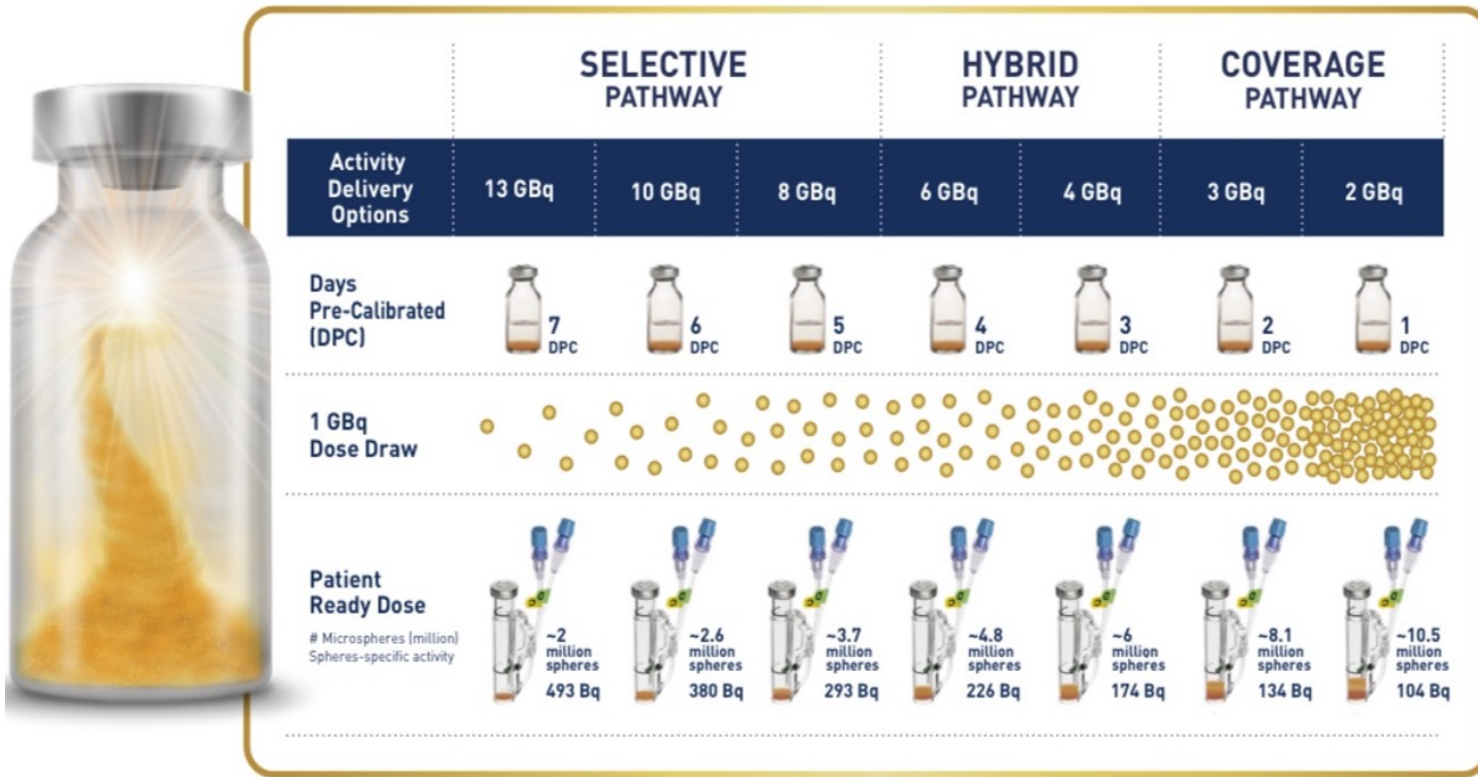
FLEXDose Delivery Program

Flexibility to Deliver the Same Activity with More or Fewer Spheres



FLEXDose Delivery Program

Flexibility to Deliver the Same Activity with More or Fewer Spheres

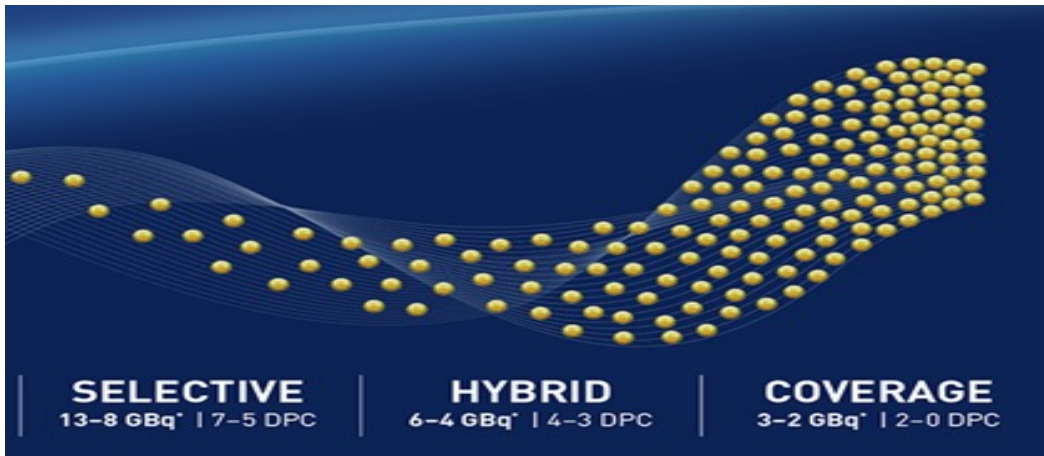


Resin Y90	Glass Y90
7-day precal	Week 2 Monday theraspheres
6-day precal	Week 2 Tuesday theraspheres



FLEXDose Delivery Program

Flexibility to Deliver the Same Activity with More or Fewer Spheres



For lobar or multifocal bilobar disease:

- 2-day precalibration
- 1-day precalibration (if very large tumor or large liver)

For RAD SEG:

- 3-day to 7-day precalibration
- Varies depending on tumor size (see table at left)

Tumor Size (Rad Seg)	Precalibration FLEXDose
<4 cm	5-7 day precalibration
4-8 cm	3-4 day precalibration

Benefits of FLEXDose SELECT Pathways

Treating Physician	For Patients
Flexibility and convenience	Customized treatment
Control – control of dosing and administration	Precise activity delivered
Ease of use	Optimal tumor coverage
Adaptability	

Dose Summary Review

Partition Model (aka 2 Compartment Model)

What is
Tumoricidal
Dose?

- Resin > 100-120 Gray
(ideally > 250 gy if unilobar)

What is
threshold
normal liver
dose?

- 40 Gray (70 Gy if unilobar)
- Less than 30 Gy if compromised
liver function, heavily pre-treated

What is
threshold
lung dose?

- 30 Gy single administration
- 50 Gy lifetime

European Journal of Nuclear Medicine and Molecular Imaging
<https://doi.org/10.1007/s00259-020-05163-5>

GUIDELINES



International recommendations for personalised selective internal radiation therapy of primary and metastatic liver diseases with yttrium-90 resin microspheres

Hugo Levillain¹ • Oreste Bagni² • Christophe M. Deroose³ • Arnaud Dieudonné⁴ • Silvano Gnesin⁵ • Oliver S. Grosser⁶ • S. Cheenu Kappadath⁷ • Andrew Kennedy⁸ • Nima Kokabi⁹ • David M. Liu¹⁰ • David C. Madoff¹¹ • Armeen Mahvash¹² • Antonio Martinez de la Cuesta¹³ • David C. E. Ng¹⁴ • Philipp M. Paprottka¹⁵ • Cinzia Pettinato¹⁶ • Macarena Rodríguez-Fraile¹³ • Riad Salem¹⁷ • Bruno Sangro¹³ • Lidia Strigari¹⁸ • Daniel Y. Sze¹⁹ • Berlinda J. de Wit van der veen²⁰ • Patrick Flamen¹

Received: 11 September 2020 / Accepted: 8 December 2020
© The Author(s) 2021

Chiesa A, et al. *QU Mol Im* 2012. Lau, et al. *Int J Radiat Oncol Biol Phys.* 2012;82:401. Levillain H, et al. *Eur J Nucl Med Mol Imaging* (online 1/21)

Dose Summary

Yttrium-90 Radioembolization Dosimetry: Dose Considerations, Optimization, and Tips

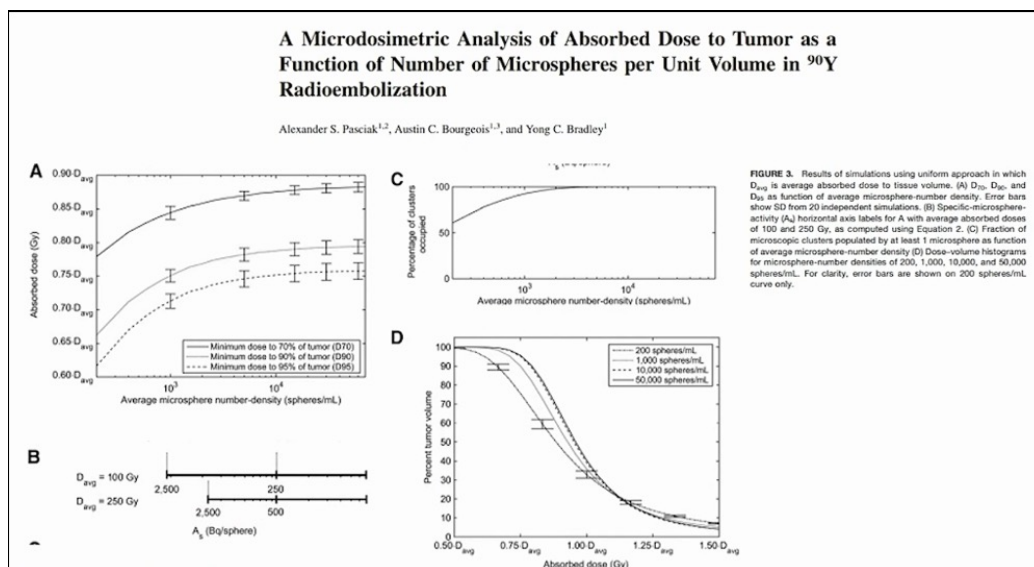
Alexander Villalobos, MD¹ Johannes L. du Pisanie, MD¹ Ripal T. Gandhi, MD, FSIR, FSVM¹
Nima Kokabi, MD, FRCPC¹

Table 4 Yttrium-90 radioembolization quick reference table—table references common clinical scenarios where yttrium-90 radioembolization can be used

Clinical scenario	Treatment intent	Ideal liver function (Child-Pugh)	Primary recommended dosimetry model	Dose goal ^a	
				Glass	Resin
Radiation segmentectomy (Y90-RS)	Curative	A–B7 >B7 ^b	Single compartment (MIRD model)	>400 Gy to angiosome	>250 Gy ^c to angiosome >350-400 Gy to angiosome → >435 Gy to tumor
Radiation lobectomy (Y90-RL)	Potentially curative; contralateral hypertrophy; “test of time”	A	Multicompartment (partition model)	Non-tumor: >88 Gy Tumor: >205 Gy	Non-tumor dose: >70 Gy Tumor: >250 Gy ^c ideally ➢ 180 Gy MIRD to lobe (3-day flex dose) ➢ >400 Gy to Segment and 180 Gy lobar (3-day)
Multifocal unilobar	Palliative	A–B7	Multicompartment (partition model)	Non-tumor w/ CP A: 100–120 Gy ^c Non-tumor w/ CP B: <70 Gy ^c Tumor: >205 Gy (ideally >250 Gy)	Non-tumor: 40–70 Gy ^c Tumor: >250 Gy ideally
Multifocal bilobar	Palliative	A	Multicompartment (partition model)	Non-Tumor: 40-70 Gy Tumor: >205 Gy (Ideally >250 Gy)	Non-tumor: 30–40 Gy Tumor: >100 Gy ^c
HCC with microvascular invasion and/or portal vein thrombus	Palliative	A ^b Ideally, must have good Tc-99m MAA uptake in MVI ^c	Multicompartment (partition model)	If unilobar MVI/PVT: R90-RL vs. Y90-unilobar approach ^c If bilobar MVI/PVT: Y90-bilobar approach ^c	If unilobar MVI/PVT: R90-RL vs. Y90-unilobar approach ^c If bilobar MVI/PVT: Y90-bilobar approach ^c

What about Sphere Count?

How Many Spheres Do You Need?



- Compared to 50k spheres/ml, decrease in D70 were statistically significant below 20k spheres/ml
- Differences in microsphere number density may have an effect on the microscopic tumor absorbed dose inhomogeneity, which may explain differences in treatment planning between glass and resin y90 devices

Pasciak AS, et al. *J Nucl Med.* 2016.

How Many Spheres Do You Need for Curative Intent



Journal of Vascular and Interventional Radiology

Available online 21 March 2023

In Press, Journal Pre-proof [What's this?](#)



Clinical Study

Yttrium-90 Radiation Segmentectomy of Hepatocellular Carcinoma: A Comparative Study of Effectiveness, Safety, and Dosimetry of Glass vs. Resin-based Microspheres

Alexander Villalobos MD¹, Linzi Arndt BS, MBA¹, Bernard Cheng MD¹, Howard Dabbous MD¹, Mohammed Loya MD¹, Bill Majdalany MD¹, Zachary Bercu MD¹, Nima Kokabi MD FRCPC¹

- 3-day pre-cal Resin
- Mean: **40,000** microspheres/cc tumor → **CR**

European Journal of Nuclear Medicine and Molecular Imaging
<https://doi.org/10.1007/s00259-025-07471-0>

ORIGINAL ARTICLE



Histopathologic outcomes of hepatocellular carcinoma treated with transarterial radioembolization with yttrium-90 resin microspheres

Ammar Sarwar¹, Imad Nasser², Jeffrey L. Weinstein¹, Mahmoud Odeh¹, Hafsa Babar³, Diana Dinh¹, Michael Curry⁴, Andrea Bullock⁵, Devin Eckhoff⁶, Martin Dib⁶, John A. Parker⁷, Muneeb Ahmed¹

- 3-day pre-cal Resin
- Median: **78,000** microspheres/cc tumor → **CPN**

cio

Miami Cancer Institute
BAPTIST HEALTH SOUTH FLORIDA

Miami Cardiac & Vascular Institute
BAPTIST HEALTH SOUTH FLORIDA

How Many Spheres Do You Need?

Methods

- Rad seg with Resin with post Y90 PET CT
- Median tumor size 3.4 cm HCC

Results

- Tumors 86% CR, 100% ORR at 6 months
- No differences between CR and incomplete responders in tumor size, volume, dose, or specific activity
- Incomplete responders had a lower D50 than CR (162 vs 273 gray), lower particle density (**36,310 vs 66,980 particles/cm³**), and higher underdosed tumor volumes
- **Note:** Montazeri SA, et al. showed specific activity >327 Bq/sphere was predictor of CR; however, in this study, 156 vs 158 Bq/sphere for CR vs incomplete response

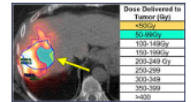
Note: D50 is minimum absorbed dose to 50% of volume

Vo NH, et al. *JVIR*. 2025.



Prediction of Recurrence of Hepatocellular Carcinoma Following Radiation Segmentectomy with Resin Microspheres Based on Underdosed Tumor Volume on Yttrium-90 Positron Emission Tomography/CT Dosimetry

Nhi H. Vo, MD, Muhammad Saad Malik, MD, Muhammad Mohid Tahir, MD, J. Anthony Parker, MD, PhD, Leo L. Tsai, MD, PhD, Diana Dinh, MD, Mamix Lam, MD, PhD, Jeffrey Weinstein, MD, Muneeb Ahmed, MD, and Ammar Sarwar, MD



ABSTRACT

Purpose: To evaluate the use of immediate post-⁹⁰Y-radiation segmentectomy (RS) positron emission tomography (PET)/computed tomography (CT) and dosimetry software in predicting incomplete response in early-stage hepatocellular carcinoma (HCC).

Materials and Methods: Patients with early-stage HCC undergoing resin ⁹⁰Y-RS (July 2014 to December 2022) prescribed using the single-compartment Medical Internal Radiation Dose (MIRD) model with post-⁹⁰Y transarterial radioembolization (TARE) PET/CT were reviewed. Forty patients (42 HCCs; 69 years [interquartile range (IQR), 63–77]; male, 75%), with a median tumor size of 3.4 cm (IQR, 2.2–4.6) met criteria. Dose-volume histograms were generated using dosimetry software (MIM SurePlan LiverY90, v7.2.3). Modified Response Evaluation Criteria in Solid Tumours (mRECIST) treatment responses were correlated with quantitative volumetric analyses of tumor absorbed dose (TAD) and underdosed tumor volume, defined as volume of tumor receiving less than 100 Gy (V₀₋₁₀₀). Additional subgroup dosimetry analysis was performed between tumors with complete response (CR) and incomplete response.

Results: Tumors treated showed 86% CR and 100% objective response rate at 6-month follow-up. Subgroup analysis between CRs and incomplete responders showed no difference in tumor size, volume, prescribed tissue dose, or prescribed, delivered, or specific activity. Incomplete responders had a lower D₅₀ than CRs (162 vs 273 Gy, *P* = .040), lower particle density (36,310 vs 66,980 particles/cm³, *P* = .040), and higher absolute (10 vs 0.7 mL; *P* = .011) and proportion (33% vs 5%; *P* = .001) of underdosed tumor volumes (V₀₋₁₀₀). An underdosed tumor volume (V₀₋₁₀₀) of ≥28% had an increased likelihood of incomplete response (odds ratio, 75; *P* = .001).

Conclusions: ⁹⁰Y-TARE PET/CT can identify tumors incompletely treated with an ablative ⁹⁰Y-RS dose and guide decision for closer follow-up or early retreatment.

Conclusions

FLEXDose

- Customized dosing depending on clinical situation
- For larger tumors/livers → day of calibration, 1-day or 2-day precalibration → I use 2-day precalibration
- For rad seg → 3-day to 7-day precalibration FLEXDose

Order Map Treat

- Allows for faster treatment
- Flexibility and ability to personalize dose to day of Y90

Tumoricidal Dose

- Rad Seg → MIRD → > 350-400 Gy to segment
- Lobar Therapies → Partition → at least 100 Gy to tumor and limit normal liver dose to <30-40 Gy
- Radiation Lobectomy: >70 Gy to background liver vs 180 Gy MIRD



Thank You



cio

 **Miami Cancer
Institute**
BAPTIST HEALTH SOUTH FLORIDA

 **Miami Cardiac & Vascular Institute**
BAPTIST HEALTH SOUTH FLORIDA

Understanding Ordering Surplus Glass Doses for Expedited Treatment

Osman Ahmed, MD
Professor of Radiology
University of Utah

Y-90 Glass Microspheres Expedited Delivery

Flexibility of **expedited delivery** services meets the needs of evolving physician schedules and emergent clinical situations.



Order for treatments
within 48 hrs



Personalized dose delivered
for patient treatment

Surplus ordering

Normal workflow – Just in time manufacturing

- Map to Treat 7-10 days
- Dose made for specific patient
- Dose arrives 24-48 hrs ahead of procedure

Surplus ordering is an “abnormal” workflow but can be done

Typically utilized for out-of-town patients/last minute deliveries

- Map to Treat in 48 hrs

Surplus Ordering

Orders placed by 12pm EST can ship same day for overnight delivery by 10:30am (local time)

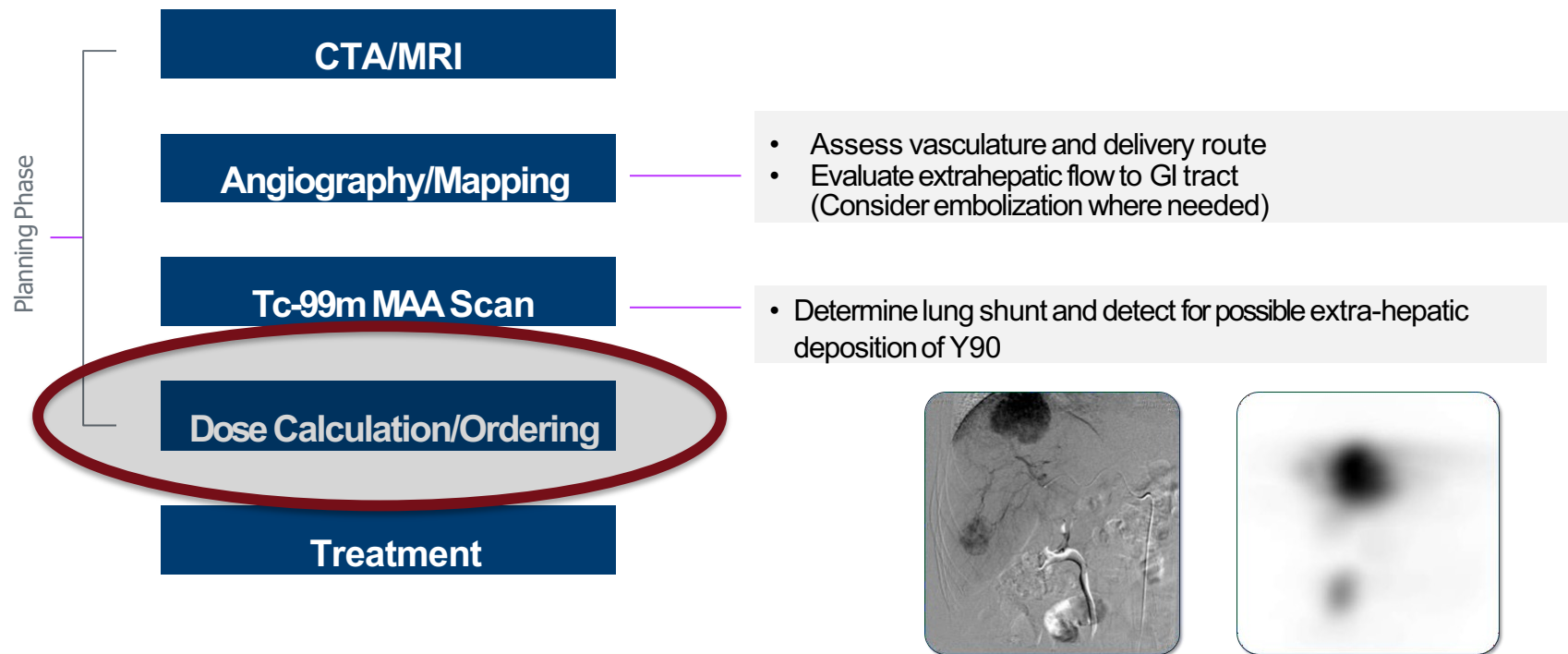
Surplus stock is standard doses (3,5,7,10,15, 20 GBq)

- Surplus stock kept at 15% based on historical usage

Requested dose not guaranteed

Work directly with rep to ensure smooth process

Treatment Pathway



Dose Options

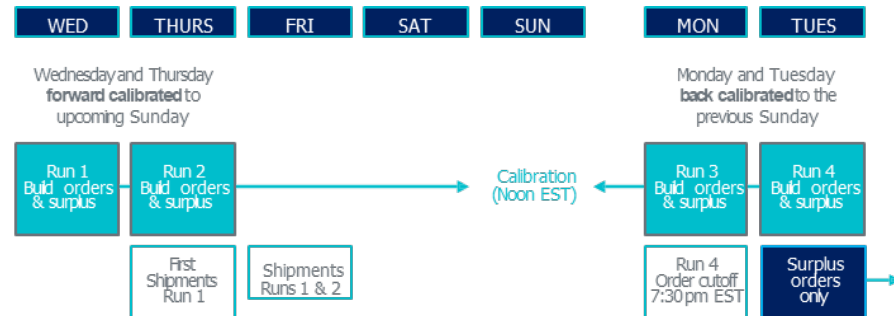
Standard Doses (GBq): 3, 5, 7, 10, 15 & 20

- Between 3 and 20 GBq in 0.5-GBq increments are available (approximately 400K spheres per GBq activity at calibration)

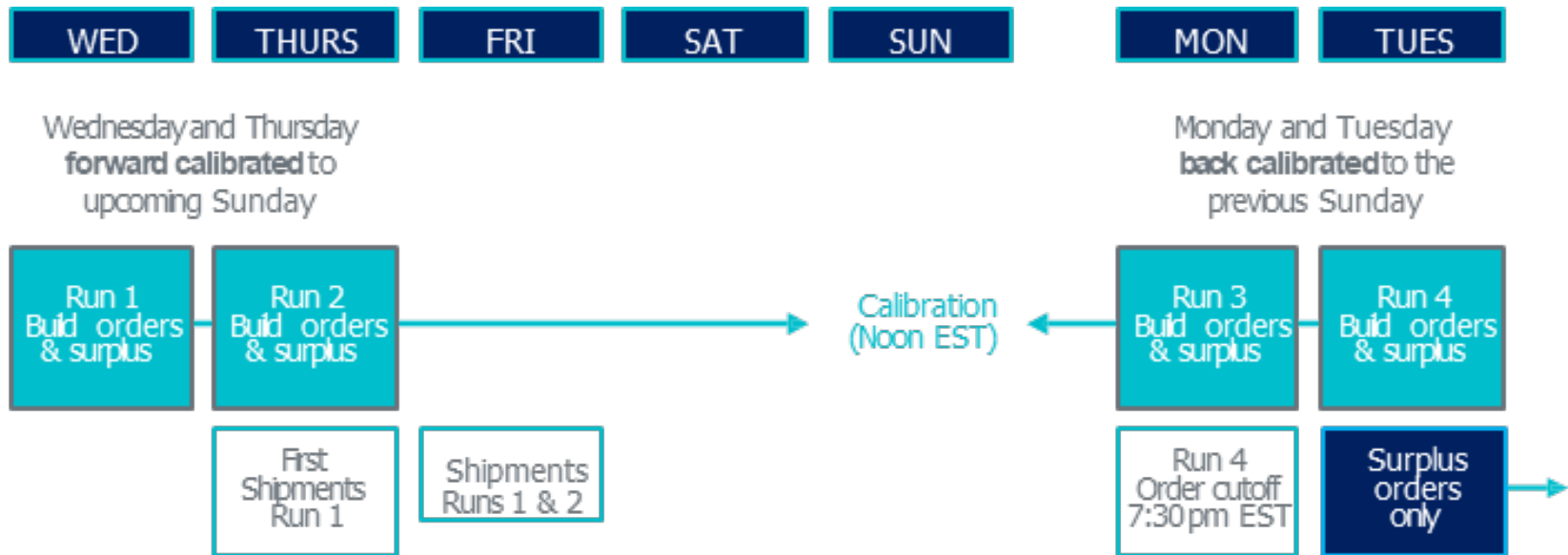
Manufacturing runs are referenced to a single calibration date (Sundays at Noon EST)

Preferably 7 days should be allowed between placing the order and treatment date (allows for shipment)

Glass dose vials arrive 1-2 days in advance of treatment date



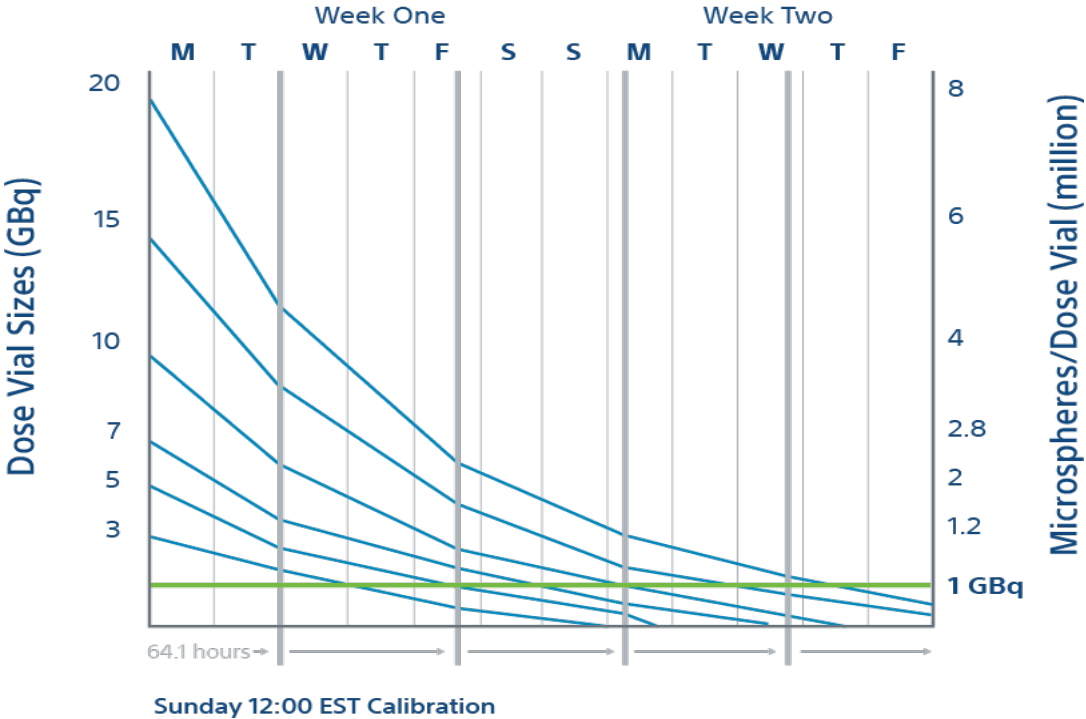
Dose Options



Dosing Flexibility for Personalized Treatment

Activity ($\pm 10\%$ GBq)	Mass of Microspheres (mg)	No. of Microspheres (millions)
3	27	1.2
5	45	2.0
7	63	2.8
10	90	4.0
15	135	6.0
20	180	8.0

Dosing Flexibility for Personalized Treatment



Conclusion

Typically use surplus orders for emergent situations or out-of-town patients (ie, to accommodate them)

- Will arrive 1-2 days prior to administration

Work directly with rep for surplus dosing

Can only order “standard” doses

Utility of Single Session Y90 Treatment

Nima Kokabi, MD, FRCPC

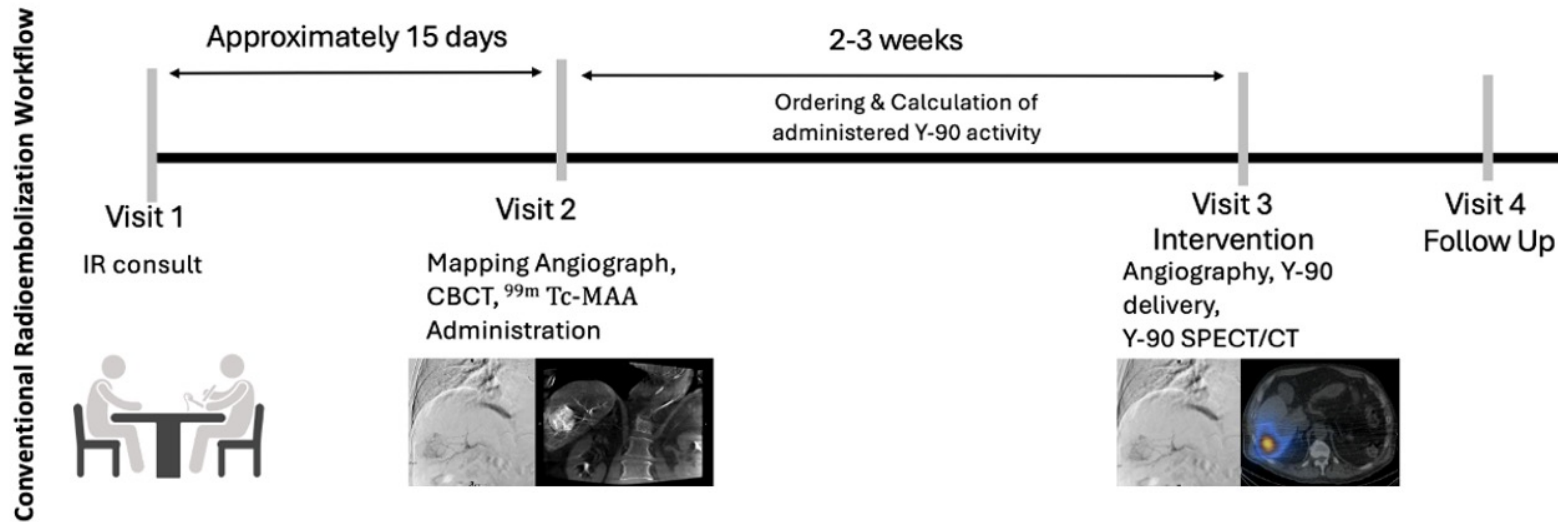
Associate Professor of Radiology, Hepatology, Medical Oncology

Vice Chair of Clinical Research

Director of Interventional Oncology

UNC School of Medicine

Conventional Workflow



SYMPOSIUM
ON CLINICAL
INTERVENTIONAL
ONCOLOGY

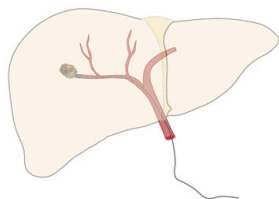


SYMPOSIUM
ON CLINICAL
INTERVENTIONAL
ONCOLOGY

- Rising cost of healthcare
- Limited resources
- Wait times for cancer care



Do we need to calculate LSF for small HCC?



Solitary HCC ≤ 5 cm **OR** ≤ 3 Tumors ≤ 3 cm



No prior TIPS placement



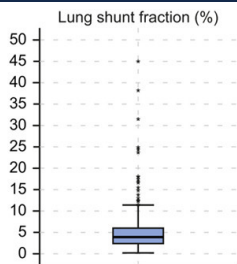
Planned selective TARE



Low lung shunt fraction



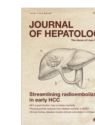
- Safe selective TARE without MAA study
- No radiation risk to the lungs



RESEARCH ARTICLE · Volume 72, Issue 6, P1351-1358, June 2020 [Download Full Issue](#)

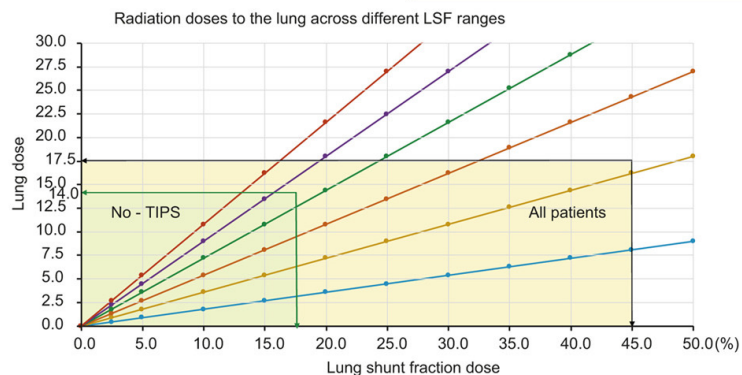
Streamlining radioembolization in UNOS T1/T2 hepatocellular carcinoma by eliminating lung shunt estimation

Ahmed Gabr¹ · Srirajkumar Ranganathan¹ · Samdeep K. Mauli¹ · Ahsun Riaz¹ · Vanessa L. Gates¹ · Laura Kulik² · Daniel Ganger² · Haripriya Maddur² · Christopher Moore² · Elias Hohlstos¹ · Nitin Katariya³ · Juan Carlos Caicedo³ · Aparna Kalyan⁴ · Robert J. Lewandowski^{1,3} · Riad Salem^{2,3,4} [Show less](#)



Retrospective
N=448 OPTN Stage T1/T2 HCC
Glass-based microsphere
Conventional 2-session mapping and treatment approach

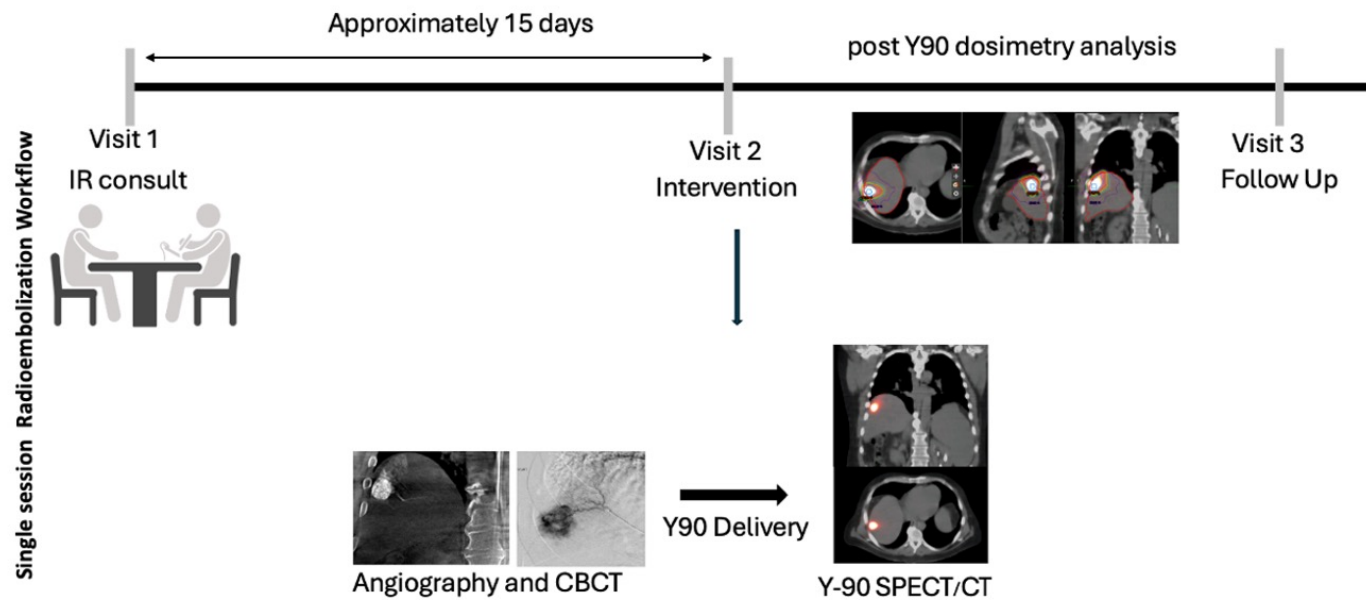
Median LSF: 3.9% (IQR 2.4–6%)
No LSF > 16%
No Lung dose > 14 Gy





SYMPOSIUM
ON CLINICAL
INTERVENTIONAL
ONCOLOGY

Single-Session Workflow



SYMPOSIUM
ON CLINICAL
INTERVENTIONAL
ONCOLOGY

Patient Selection

Patient Selection Criteria

- OPTN Stage I/II HCC (≤ 5 cm, ≤ 3 nodules)
- No macrovascular invasion, no TIPS
- Child–Pugh A–B7, bilirubin < 2 , ECOG < 2
- No arterial enhancement of venous structures on MRI/CT or intraprocedural CT



Glass-Based Microsphere

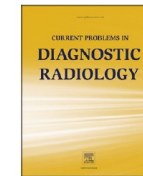
Current Problems in Diagnostic Radiology 54 (2025) 308–312



Contents lists available at [ScienceDirect](#)

Current Problems in Diagnostic Radiology

journal homepage: www.cpdjournal.com



Learning from Experience

Expediting care for hepatocellular carcinoma ≤ 3 cm by streamlining radiation segmentectomy: A quality improvement project

Cynthia De la Garza-Ramos^a, Steven Bussone^b, LaRissa L. Adams^c, Maeghan D. Barber^b, Gregory T. Frey^a, Andrew R. Lewis^a, Ricardo Paz-Fumagalli^a, Beau B. Toskich^{a,*}

^a Division of Interventional Radiology, Mayo Clinic Florida, Jacksonville, FL, USA

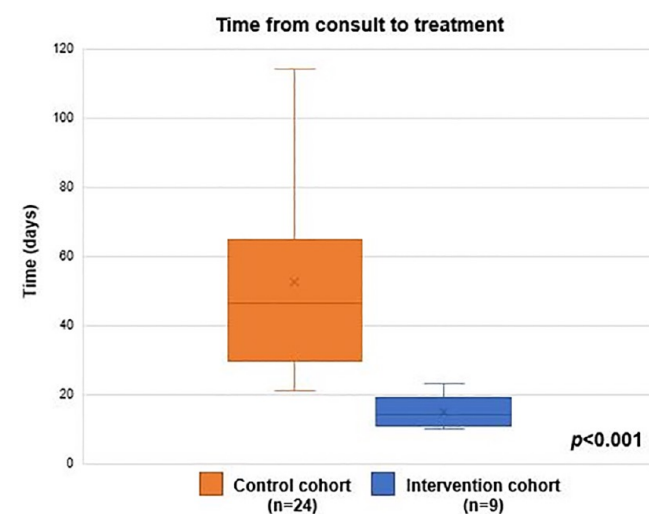
^b Department of Radiology, Mayo Clinic Florida, Jacksonville, FL, USA

^c Systems Quality Office, Mayo Clinic Florida, Jacksonville, FL, USA



SYMPOSIUM
ON CLINICAL
INTERVENTIONAL
ONCOLOGY

- Treatment N=9 patients (prospective)
- Control N=24 patients (retrospective)
- **Time to treatment**
 - 14 days (IQR 12–15) vs 47 days (IQR 31–64)
- **LSF**
 - 2.7 Gy (IQR 2.1, 3.6) vs 2.2Gy (IQR 1.0, 4.6)
- **Fluoroscopy time**
 - 8 min (IQR 14, 19) vs 28 min (IQR 24, 43)



Glass-Based Microsphere




Cardiovasc Intervent Radiol (2024) 47:1239–1245
<https://doi.org/10.1007/s00270-024-03799-6>



CLINICAL INVESTIGATION

INTERVENTIONAL ONCOLOGY

Single-Session Ablative Transarterial Radioembolization for Patients with Hepatocellular Carcinoma to Streamline Care: An Initial Experience

Zachary T. Berman¹  · Kurt Pianka¹ · Yousuf Qaseem¹ · Jonas Redmond¹ ·
Jeet Minocha¹



Results





- Retrospective N=15
 - 16 consecutive procedures
- Technical success: **88%**
 - **2 patients couldn't undergo single session b/c of intraprocedural findings**
- 25% required ≥ 2 vials for complete coverage
- **Median procedure time: 86 min**
 - **~\$7,000 saved per patient**
- **No radiation pneumonitis**
- Complete response: **94%**

Resin-Based Microsphere



Article

Initial Experience with Single-Session Resin-Based Transarterial Radioembolization Mapping and Treatment of Small Hepatocellular Carcinomas

Michael Mohnasky ^{1,*}, Sandra Gad ^{1,2}, Marco Fanous ¹, Johannes L. Du Pisanie ³, Marija Ivanovic ³, David M. Mauro ³, Hyeon Yu ³, Alex Villalobos ³, Andrew M. Moon ^{4,5}, Hanna K. Sanoff ⁶, Jingquan Jia ⁶ and Nima Kokabi ³



SYMPOSIUM
ON CLINICAL
INTERVENTIONAL
ONCOLOGY



Results

- **Retrospective**
- **Treatment N=10 pts (Resin-based microspheres)**
- **Control N=60 pts (Glass-based microspheres)**

Results

- No radiation pneumonitis
- **Complete response 90%**
- **No Grade 3+ AE**
- **Median to time Rx from IR clinic: 26.5 days vs 61 days ($p < 0.001$)**
- **Procedure time: 142 vs 151 min (NS)**
- **Fluoroscopy time: 25 vs 23 min (NS)**

Cost Savings

Table 4. Cost comparison of mapping and Y90 treatment procedures.

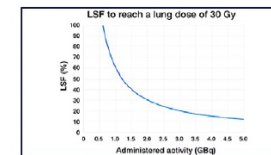
	Hospital Outpatient (OPPS) (USD)	Physician Services (MFFS) (USD)	Total Cost (USD)
Mapping Total Cost	25,400.86	1231.25	26,632.11
75,726: Angiography, visceral, RS&I	5405.70	90.57	
37,242: Arterial emb or occ, RS&I; arterial other than hem or tumor (includes catheter placement, 3D post-scan, ^{99m} Tc-MAA dose, radiopharmaceutical quantification measurement)	17,956.72	1042.99	
74,170: CT, abdomen; w and w/o contrast	178.02	63.40	
78,801: Planar imaging of multiple areas	401.83	32.35	
78,832: SPECT/CT imaging of multiple areas	1458.59	92.51	
Y90 Treatment Total Cost	36,685.53	1283.67	37,969.20
75,726: Angiography, visceral, RS&I	5405.70	90.57	
C2616: Brachytherapy source (yttrium-90 non-stranded–Medicare)	17,412.53	-	
37,243: Vascular emb or occ, inclusive of all RS&I, intraprocedural road mapping, and imaging guidance necessary to complete intervention; for tumors	11,340.57	858.64	
77,370: Special Medical Radiation Physics Consultation	132.77	-	
77,470: Special Treatment Procedure	578.47	104.48	
77,300: Basic Dosimetry Calculation	132.77	32.02	
79,445: Radiopharmaceutical therapy, intra-arterial particulate admin (1 doctor model (IR/AU))	224.13	105.45	
78,832: SPECT/CT imaging of multiple areas	1458.59	92.51	

↓41% Cost

Resin and Glass-Based Microsphere

CLINICAL STUDY

Streamlining Radioembolization without Lung Shunt Estimation versus Regular Radioembolization in Patients with Hepatocellular Carcinoma within the Milan Criteria



Hyo-Cheol Kim, MD, Minseok Suh, MD, Jin Chul Paeng, MD, Jong Hyuk Lee, MD, Myungsu Lee, MD, Jin Wook Chung, MD, and Jin Woo Choi, MD



SYMPOSIUM
ON CLINICAL
INTERVENTIONAL
ONCOLOGY

Results

- Retrospective, single-center, cohort study
- N = 100 consecutive patients
 - **Conventional** N=38
 - **Single session** N=62
- Microspheres: glass (n=84), resin (n=16)
- **Serious AEs \geq Grade 3**
 - **Conventional:** 7.9%
 - **Single session:** 3.2%

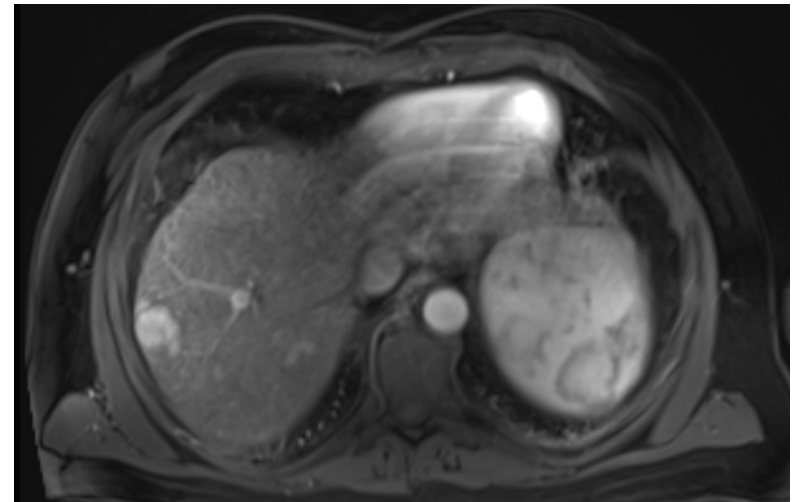
Results

- **Glass Microspheres**
- CR rates:
 - Conventional: 96.9%
 - **Single session: 90.4%**
- No significant difference

- **Resin Microspheres**
- Small numbers (n=16)
- CR rates:
 - Conventional: 66.7%
 - **Single session: 90%**
- No significant differences

Case Example

- 68-yr-old Male with history of HCV
- Bili 0.5, ECOG 1, Child-Pugh A5
- LR 5
- 2.4 cm HCC
- 28 days from consult to treatment



Dosimetry

- Angiosome volumetry by CBCT
- 5-day pre-cal Resin
- **Target >300 Gy**
- 2 options:
 - Call nuc med to draw the dose after volumetry → wait ~30 minutes for the box(es) to arrive
 - Have 3 boxes pre-drawn and ready in the room



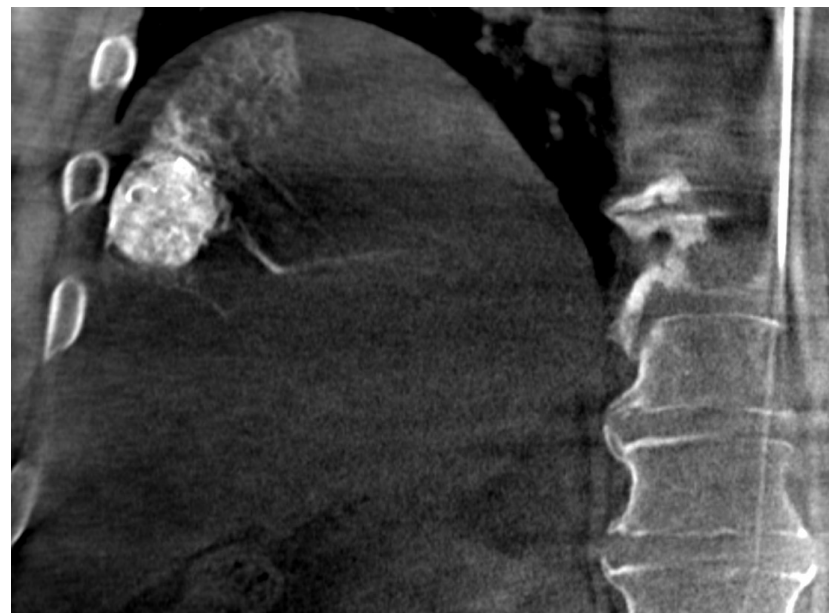
27 mci (150 cc)



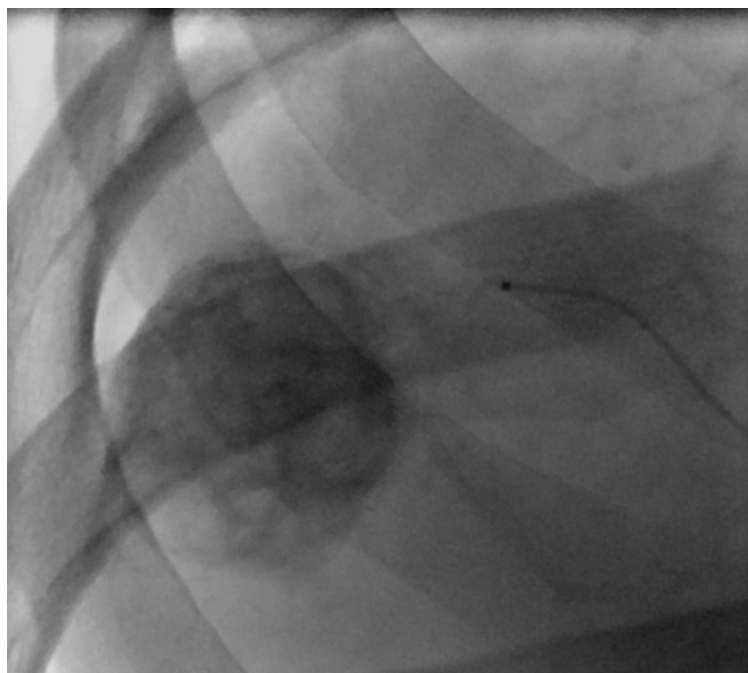
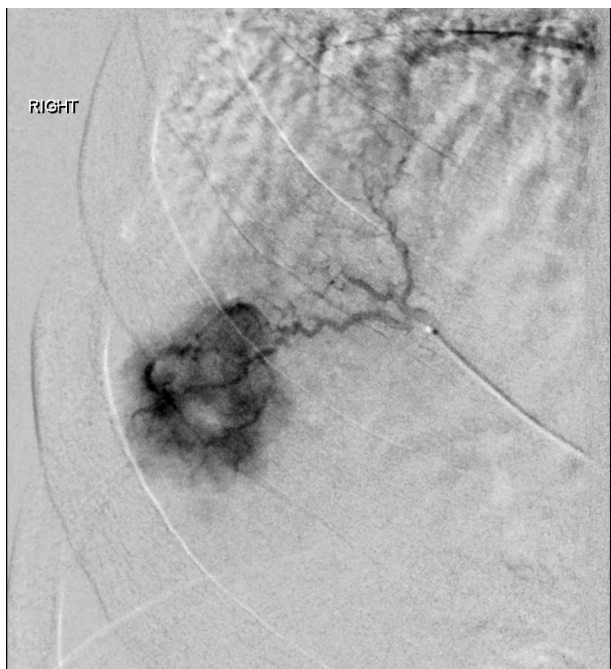
36 mci (200 cc)



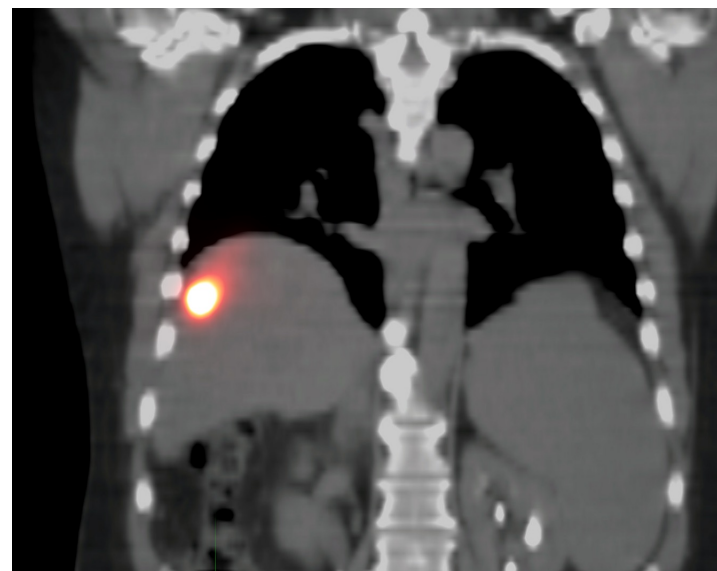
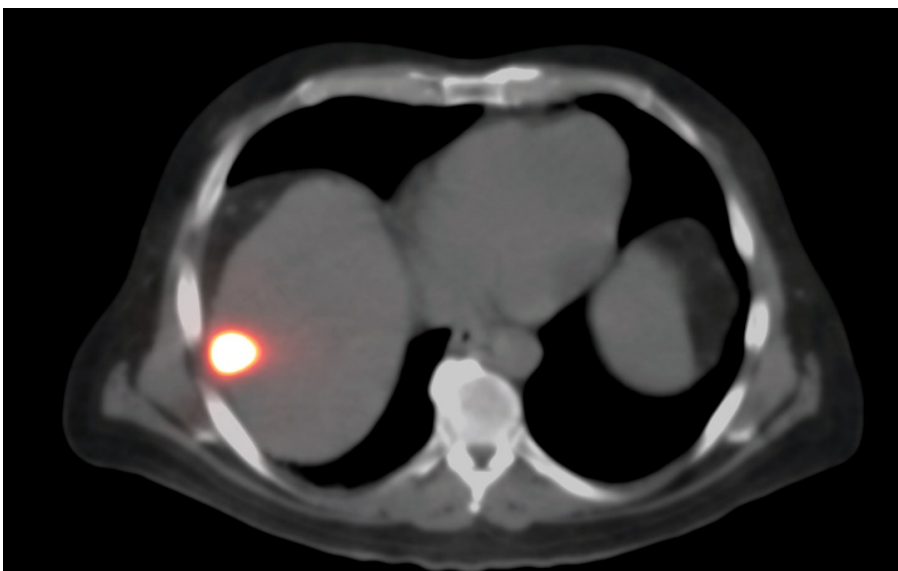
45 mci (250 cc)



SYMPOSIUM
ON CLINICAL
INTERVENTIONAL
ONCOLOGY



SYMPOSIUM
ON CLINICAL
INTERVENTIONAL
ONCOLOGY



SYMPOSIUM
ON CLINICAL
INTERVENTIONAL
ONCOLOGY

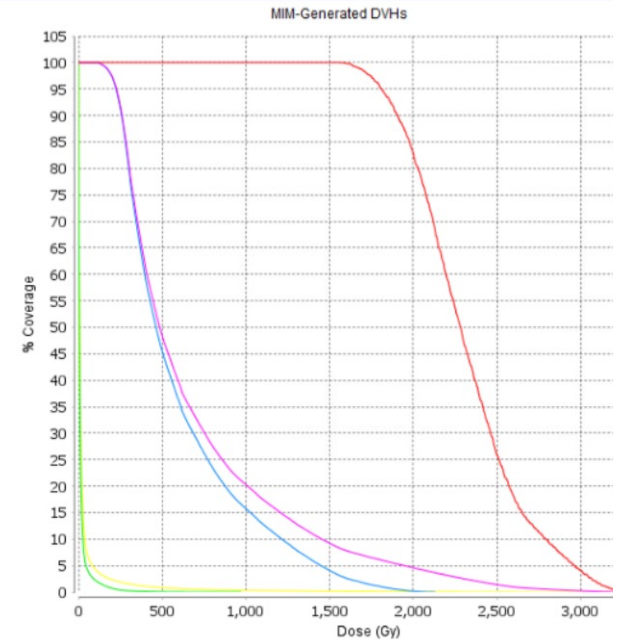
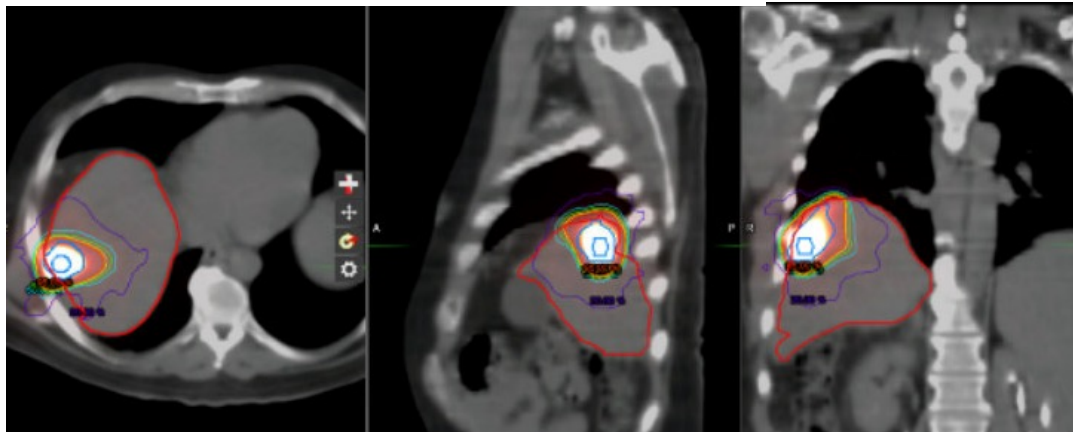
POST Y90 DOSIMETRY ANALYSIS

Intent: Curative
 Dosimetry Method: MIRD (ONE AND DONE TRIAL)
 Calibration Utilized: 5-day Pre-Cal

Total Liver Volume: 2,015 mL
 Targeted Angiosome Volume: 33 mL (1.6% of Liver)
 Targeted Tumor Volume: 2 mL

Targeted Angiosome Dose: >300 (ONE AND DONE TRIAL PROTOCOL)
 Actually Delivered Activity: 33 mCi
 Actually Achieved Angiosome Dose: 695 Gy
Actually Achieved Tumor Dose: 2,313 Gy

LSF: 5.78%



Dose ID	Dose Details				
D1	LDM with Known Activity Dose (Restricted to Liver)				
Contour	Line Style	Max Dose	Min Dose	Mean Dose	SD
	D1	D1	D1	D1	D1
Liver + Extrahepatic Activity	—	3303.70	0.00	26.86	113.48
Liver_Perfused_Target_I	—	3303.70	114.52	694.67	543.14
Non-Tumor	—	2127.50	114.52	603.18	385.93
Normal Liver	—	968.42	0.00	13.77	33.06
Tumor 1	—	3303.70	1567.17	2313.12	333.81

Glass vs Resin

Devices

- **Glass-Based Microspheres**
 - Deliver higher specific activity per particle with fewer particles
 - Pre-ordered vials
- **Resin-Based Microspheres**
 - Higher particle numbers but with lower specific activity per particle
 - Flexible dosing using mother vial aliquots
- Comparable outcomes

FUTURE DIRECTIONS



Streamlined Resin Y90 Radiation Segmentectomy for Small HCC (One&Done)

ClinicalTrials.gov ID ⓘ NCT06618300

Sponsor ⓘ UNC Lineberger Comprehensive Cancer Center

Information provided by ⓘ UNC Lineberger Comprehensive Cancer Center (Responsible Party)

Last Update Posted ⓘ 2024-11-07



Conclusion

- **Single-session Y90 is safe, effective, and cost-saving**
- **Promising for small HCCs in selected patients**
- **Potential to transform access in liver cancer care**