



Annual Users' Conference

**ADVANCING RADIATION SAFETY.
EMPOWERING PROGRESS.**

M Resort, Henderson, NV | June 15 – June 18



MIRION
TECHNOLOGIES

*Advancing Radiation Safety.
Empowering Progress.*

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From Measurement to Meaning

Building Accurate, Standardized Personalized RPTs

Carlos Uribe, PhD, MCCPM

Molecular Imaging and Therapy Physics Lead, BC Cancer

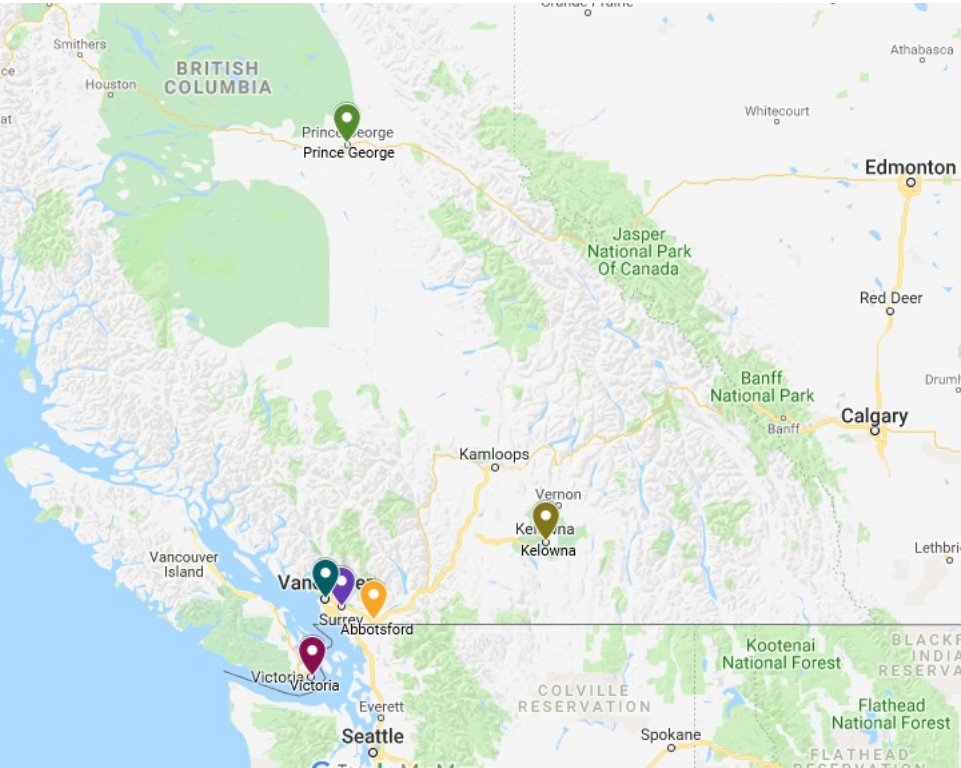
Mirion Connect | Annual Users' Conference 2026

Henderson, NV

Where I come from...



BC Cancer



Research Locations

Genomics



Deeley Research Centre
- Victoria

Drug Delivery



BC Cancer Research Centre
- Vancouver

Stem Cells

Medical Imaging
Radiation Therapy

Carlos Uribe, PhD, MCCPM

Immunology

Molecular Imaging and Therapy



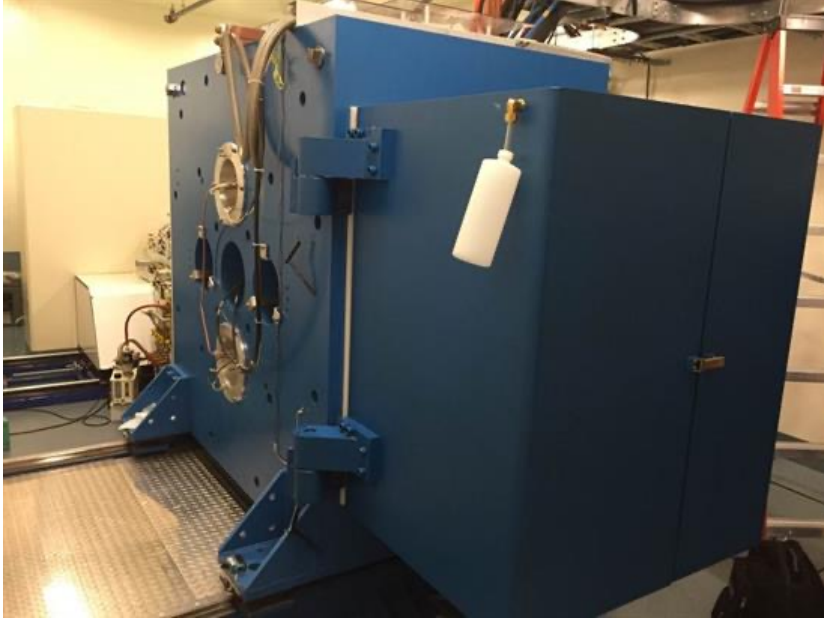
■ Multidisciplinary team

- NM Physicians
 - NM Physicists
 - NM Technologists
 - Radiochemists
 - Biologists
 - Computer Scientists
-
- Graduate Students
 - Postdoctoral Fellows

Molecular Imaging and Therapy



Molecular Imaging and Therapy



TR19 cyclotron
(ACSI, Richmond, BC)



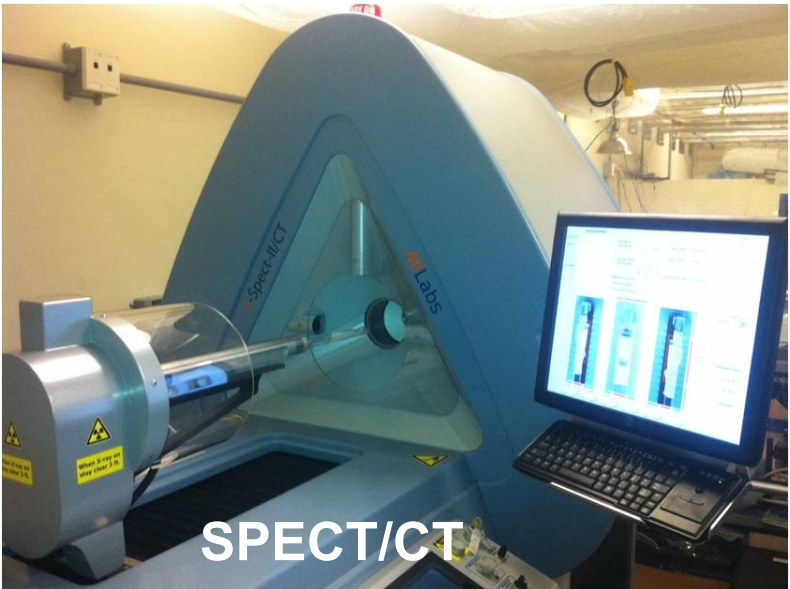
Two PET/CT Scanners

- GE DMI 5 Ring
- Siemens Vision Quadra

Small Animal Imaging Resource



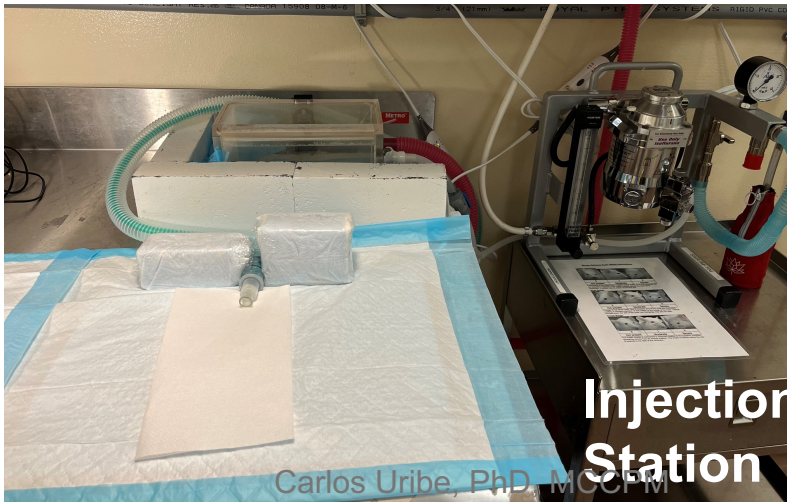
PET/CT



SPECT/CT



Clean

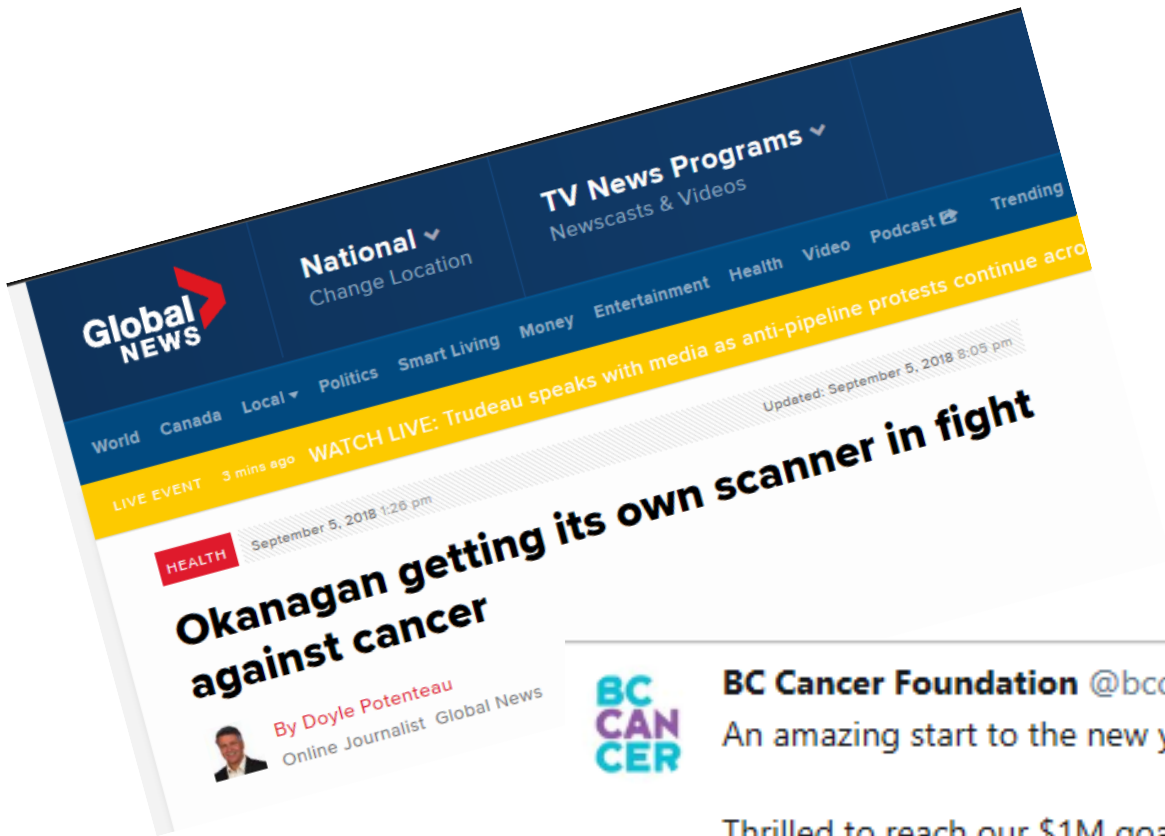


Injection Station



Necropsy Station

The Theranostics Expansion



The Theranostics Expansion

CBC

MENU

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British Columbia

Justin Trudeau announces funding to build nuclear medicine hub in Vancouver



PM says initiative will help Canada stay a leader in medical isotope research

The Canadian Press · Posted: Nov 01, 2018 5:51 PM PT | Last Updated: November 1, 2018



JUSTIN TRUDEAU, PRIME MINISTER OF CANADA

NEWSPHOTOS & VIDEOS

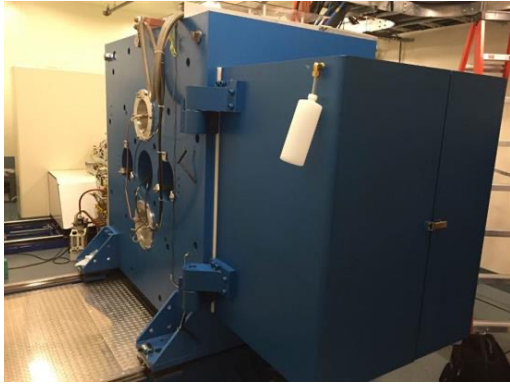
New home for nuclear medicine at the University of British Columbia

News » New home for nuclear medicine at the University of British Columbia



IAMI

Molecular Imaging and Therapy



3 Radiopharmacies
3 Cyclotrons

Two Nine PET/CT Scanners

- GE DMI 5 Ring
- Siemens Vision Quadra
- ...

Molecular Imaging and Therapy

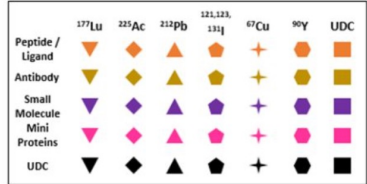


3 Radiopharmacies
3 Cyclotrons

Two Nine PET/CT Scanners

- GE DMI 5 Ring
- Siemens Vision Quadra
- ...

14



MIRION
Connect
AMERICAS

26

Annual Users' Conference

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Empowering Progress.
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Molecular Imaging and Therapy

SNMMI/ACNM Procedure Standard for Post-Treatment Imaging of ^{177}Lu -based Radiopharmaceuticals

Carlos Uribe^{1,2,3}, Amir Iravani⁴, Bital Savir-Baruch⁵, Heather Jacene⁶, Stephen A. Graves^{7,8,9}, Yuni K Dewaraja¹⁰, Courtney Lawhn Heath¹¹, Thomas A. Hope¹²

Research Article | Procedure Standard

Summary: SNMMI/ACNM Procedure Standard for Posttreatment Imaging of ^{177}Lu -Based Radiopharmaceuticals

Carlos Uribe, Amir Iravani, Bital Savir-Baruch, Heather Jacene, Stephen A. Graves, Yuni K. Dewaraja, Courtney Lawhn Heath and Thomas A. Hope
Journal of Nuclear Medicine October 2025, 66 (10) 1528-1537; DOI: <https://doi.org/10.2967/jnumed.125.270979>

Molecular Imaging and Therapy



Carlos Uribe, PhD, MCCPM

BC Cancer & My Role

BC CANCER

- *A provincial cancer agency serving all of British Columbia*
- *Molecular Imaging and Therapy program: a provincial program translating new radiopharmaceuticals from discovery to first-in-human*
- *Province-wide clinical nuclear medicine physics*

MY ROLE

- *Leader, Clinical Nuclear Medicine Physics (Provincial Operations)*
- *Co-lead, Qurit research program*
- *Member, MIRD Committee and SNMMI Dosimetry Task Force*
- *Led the SNMMI 177Lu Dosimetry Challenge*
- *Chair of the SNMMI Clinical Trial Network Scanner Validation committee*

The RPT problem...



Same administered activity

7.4 GBq
(200 mCi)



Tumor load

Very Low

Low

Moderate

High

Very High

Same administered activity

7.4 GBq
(200 mCi)



Tumor load

Very Low

Low

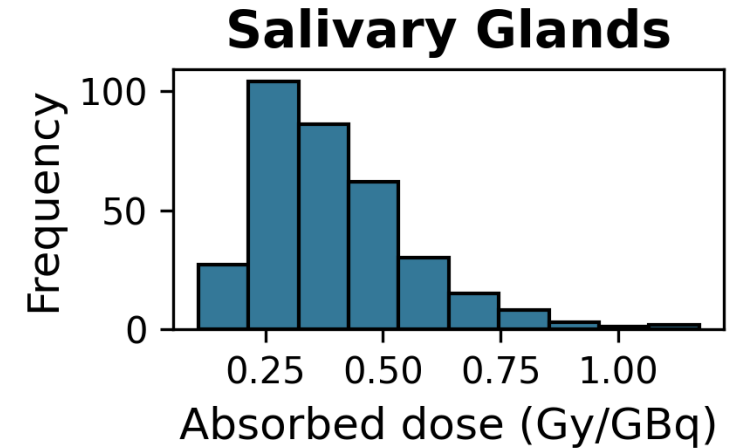
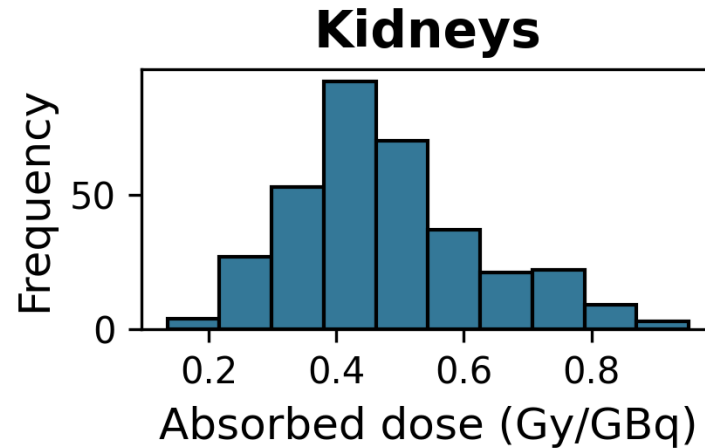
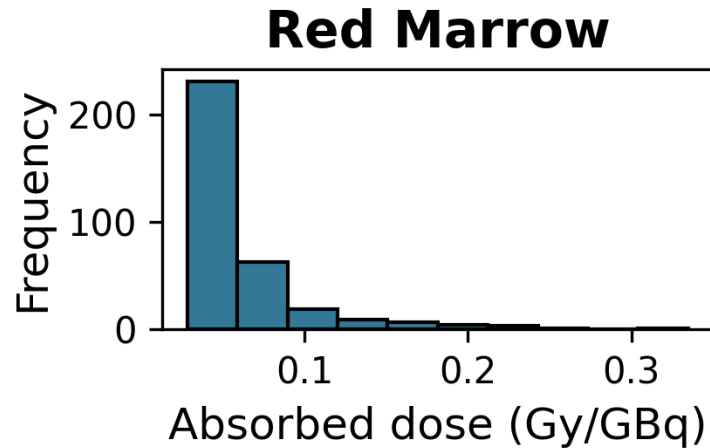
Moderate

High

Very High

But uptake, retention, clearance, tumor burden, and organ risk are patient-specific.

Same administered activity...different absorbed dose



But uptake, retention, clearance, tumor burden, and organ risk are patient-specific.

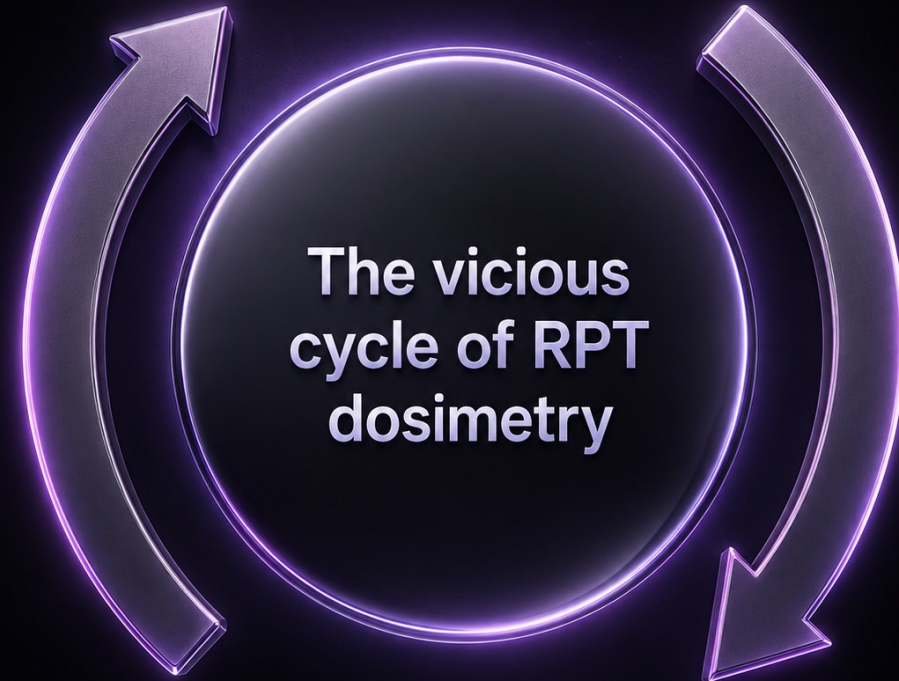


Rethinking Dosimetry: A European Perspective

Johannes Tran-Gia¹, Francesco Cicone², Michel Koole³, Francesco Giammarile^{4,5}, Jonathan Gear⁶, Emmanuel Deshayes^{7,8}, Pablo Minguez Gabiña^{9,10}, Marta Cremonesi¹¹, Jonathan Wadsley¹², Peter Bernhardt¹³, Manuel Bardiès^{7,8}, Silvano Gnesin¹⁴, Mattias Sandström¹⁵, Ulrike Garske-Román^{16,17}, Mona-Elisabeth R. Revheim^{18,19}, Frederik A. Verburg²⁰, Mark Konijnenberg²⁰, Bernd Joachim Krause²¹, Michael Lassmann¹, and Caroline Stokke^{22,23}



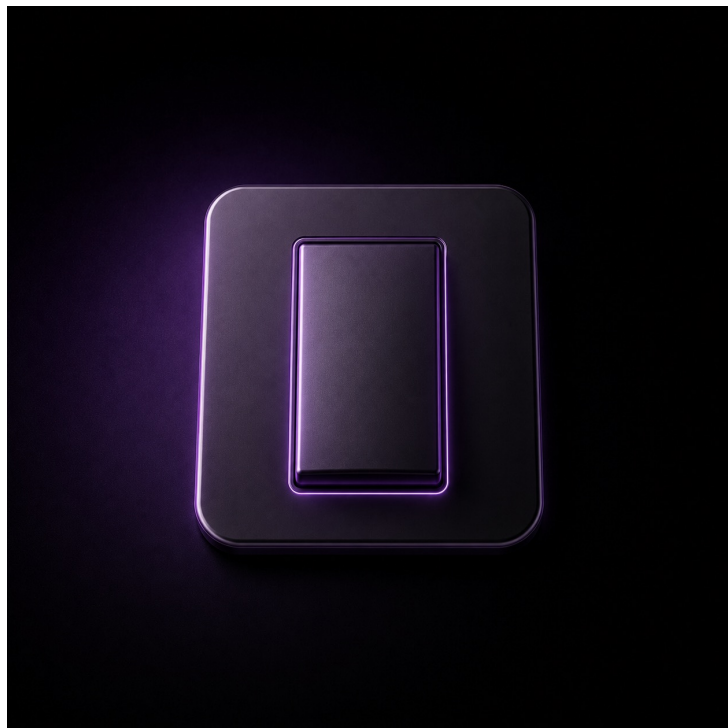
No routine dosimetry
due to lack of evidence



No evidence due to
lack of routine use

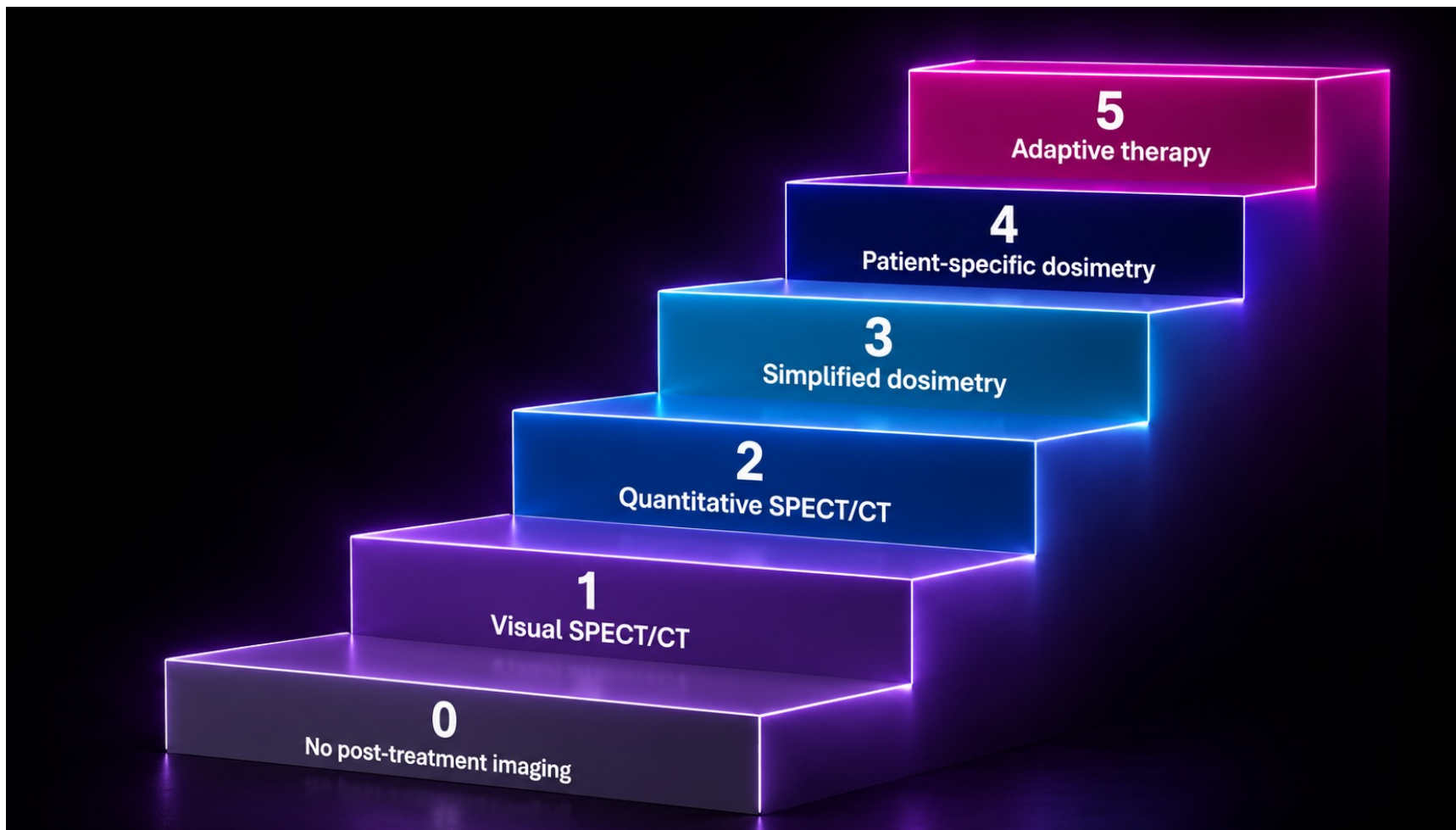
Dosimetry is a staircase, not an on/off switch

Start where the clinical workflow can support it. Build toward adaptive therapy.

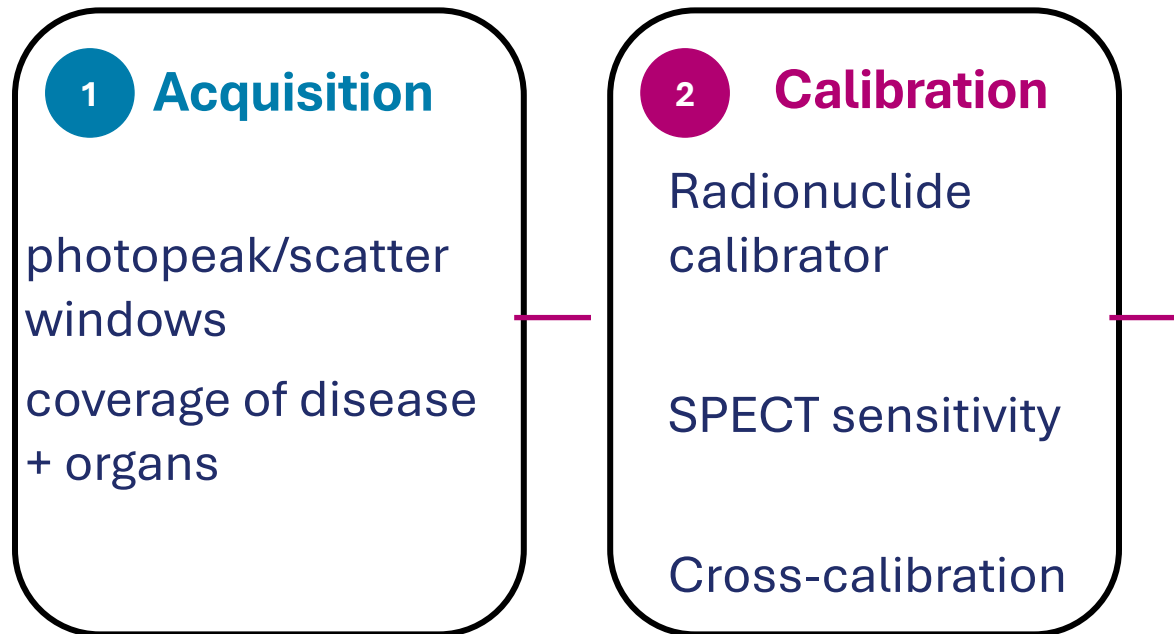


Dosimetry is a staircase, not an on/off switch

Start where the clinical workflow can support it. Build toward adaptive therapy.



Quantitative SPECT/CT is the foundation for dosimetry



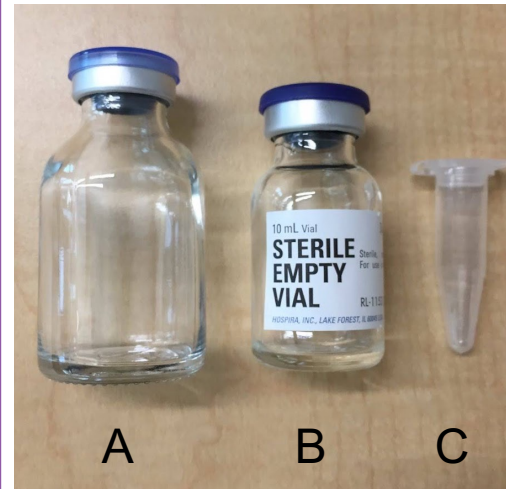
Radionuclide calibrator

Necessary to:

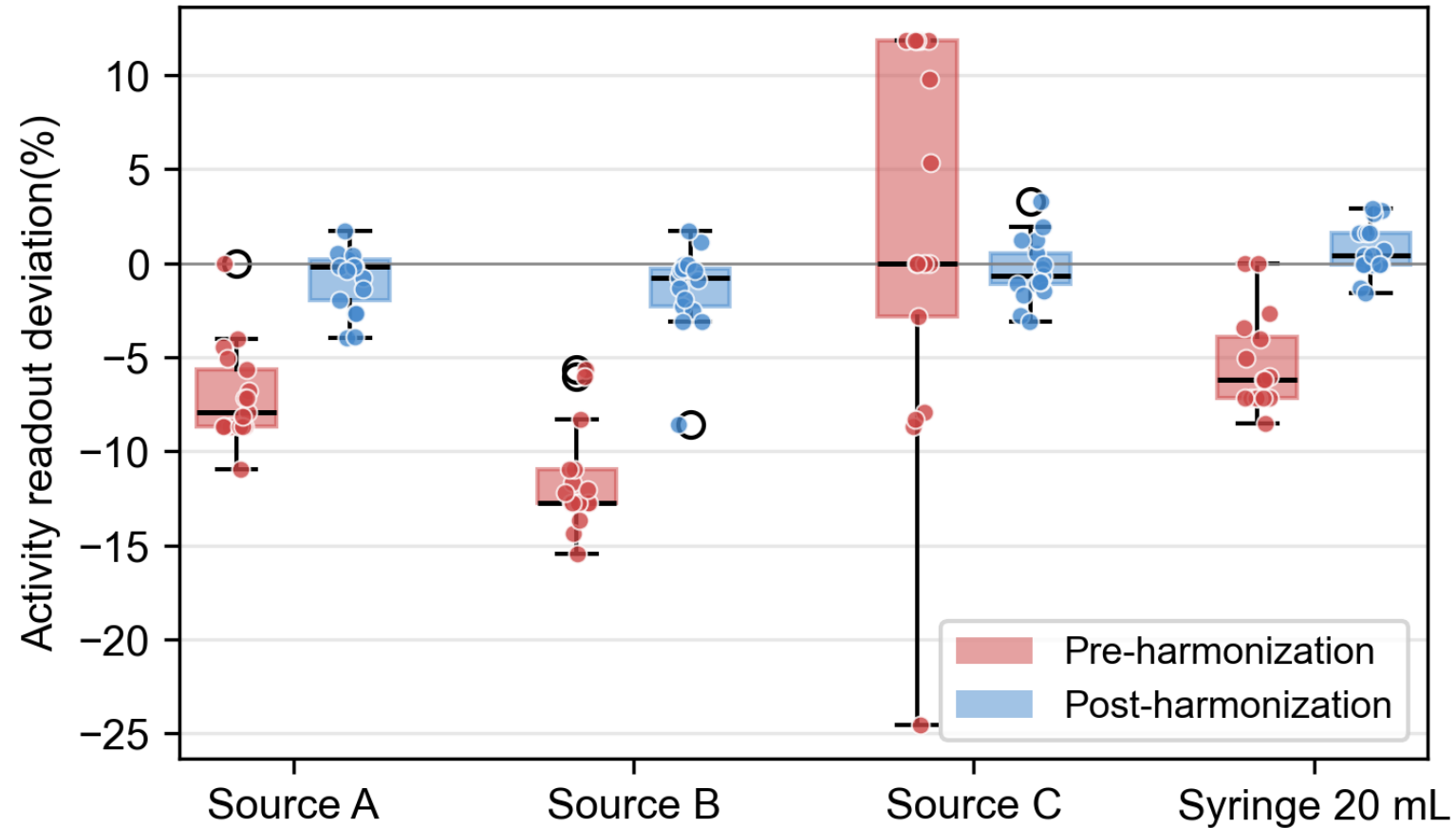
- Accurately measure how much activity is injected to the patient
- Cross calibrate with SPECT scanners to generate quantitative images.



Different source geometries



Radionuclide calibrator



SPECIAL CONTRIBUTIONS

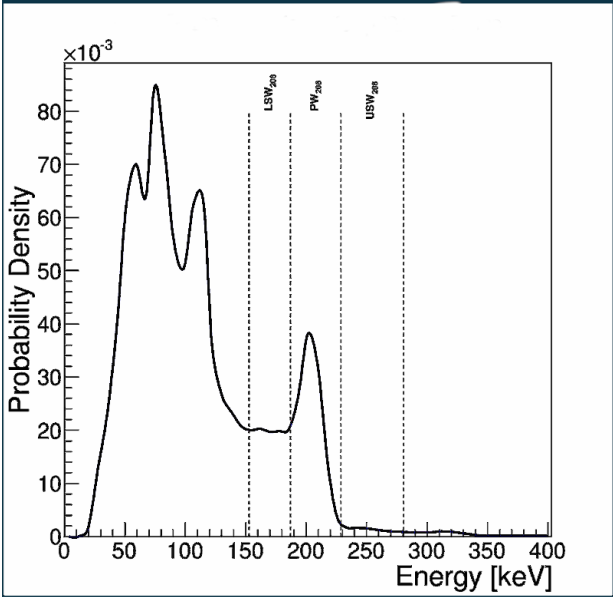
MIRD Pamphlet No. 26: Joint EANM/MIRD Guidelines for Quantitative ¹⁷⁷Lu SPECT Applied for Dosimetry of Radiopharmaceutical Therapy

Michael Ljungberg¹, Anna Celler², Mark W. Konijnenberg³, Keith F. Eckerman⁴, Yuni K. Dewaraja⁵, and Katarina Sjögreen-Gleisner¹

In collaboration with the SNMMI MIRD Committee: Wesley E. Bolch, A. Bertrand Brill, Frederic Fahey, Darrell R. Fisher, Robert Hobbs, Roger W. Howell, Ruby F. Meredith, George Sgouros, and Pat Zanzonico, and the EANM Dosimetry Committee: Klaus Bacher, Carlo Chiesa, Glenn Flux, Michael Lassmann, Lidia Strigari, and Stephan Walrand.

¹Department of Medical Radiation Physics, Lund University, Lund, Sweden; ²Radiology Department, Medical Imaging Research Group, University of British Columbia, Vancouver, Canada; ³Department of Nuclear Medicine, Erasmus University Medical Center, Rotterdam, Holland; ⁴Easterly Scientific, Knoxville, Tennessee; and ⁵Department of Radiology, University of Michigan Medical School, Ann Arbor, Michigan

Photopeak window	208 @20% [187.2 – 228.8] keV
Low scatter window	10% [166.4 – 187.2] keV
Upper scatter window	10% [228.8 – 249.6] keV



Quantitative SPECT



PET/CT Scanner Validation



A successful scanner validation will ensure that your study patients as well as your clinical patients are being scanned on a qualitatively and quantitatively accurate PET/CT. Contact ctnadmin@snmmi.org for more info.

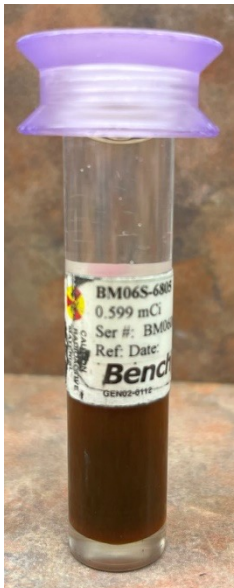
CTN works collaboratively with SNMMI Councils and Chapters as well as with external groups to move the entire field forward. In working with other imaging associations such as RSNA, government agencies such as the FDA and industry leaders, the CTN envisions having a key role in advancing the use of radiopharmaceuticals and optimizing the use of molecular imaging in clinical trials and for dissemination into clinical practice.

SPECT/CT Scanner Calibration

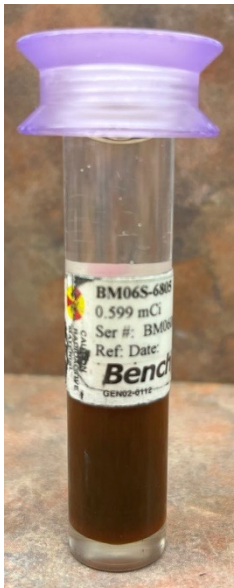


A successful radionuclide-specific SPECT/CT scanner calibration will allow conversion of the SPECT measured counts/voxel into a true radionuclide concentration measurement (Bq/ml) for accurate dosimetry measurements. Contact ctnadmin@snmmi.org for more info.

But there's a problem in the measurement chain



But there's a problem in the measurement chain

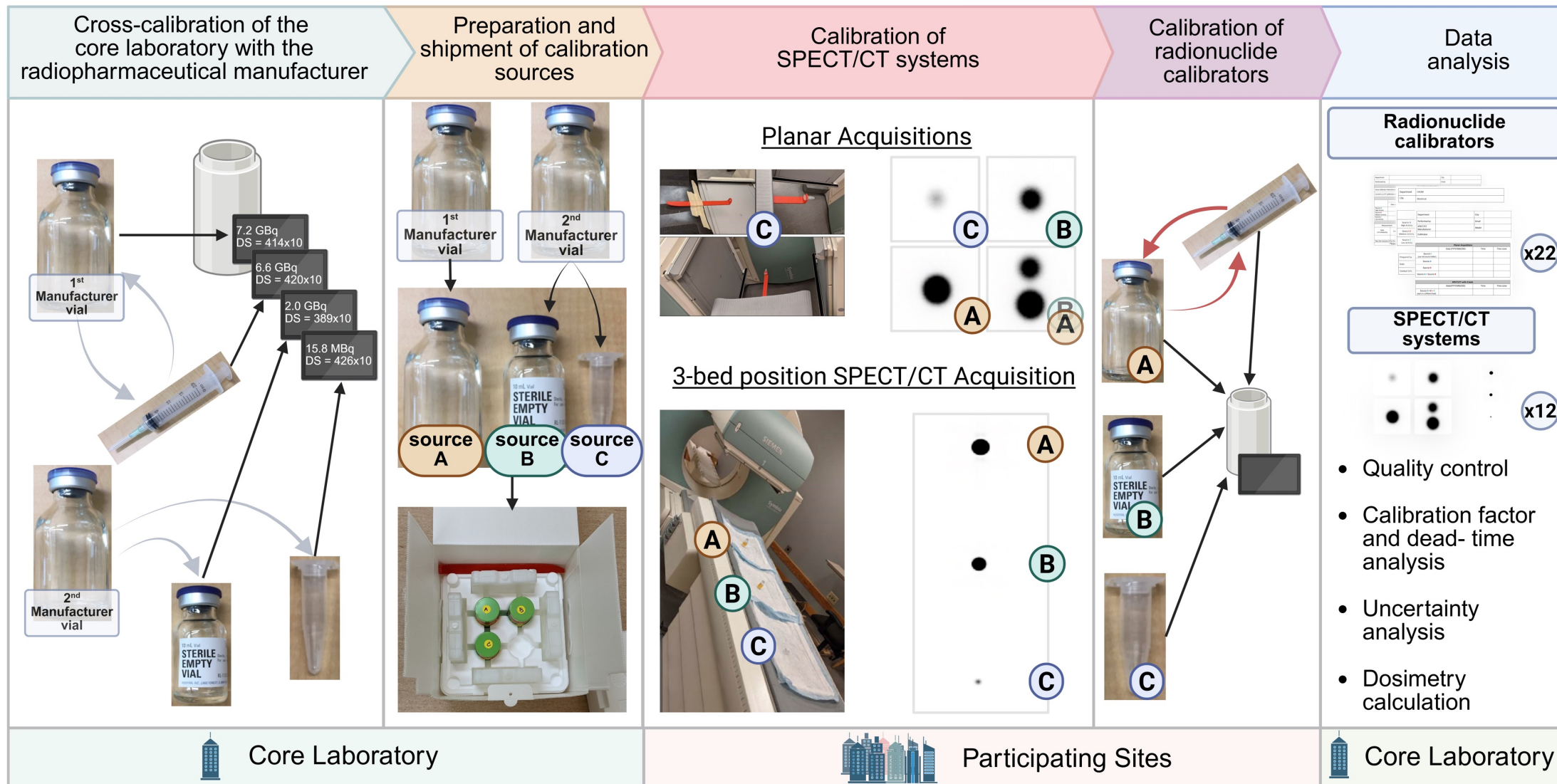


Lu-177
I-131
Ac-225
In-111
Pb-203
Pb-212

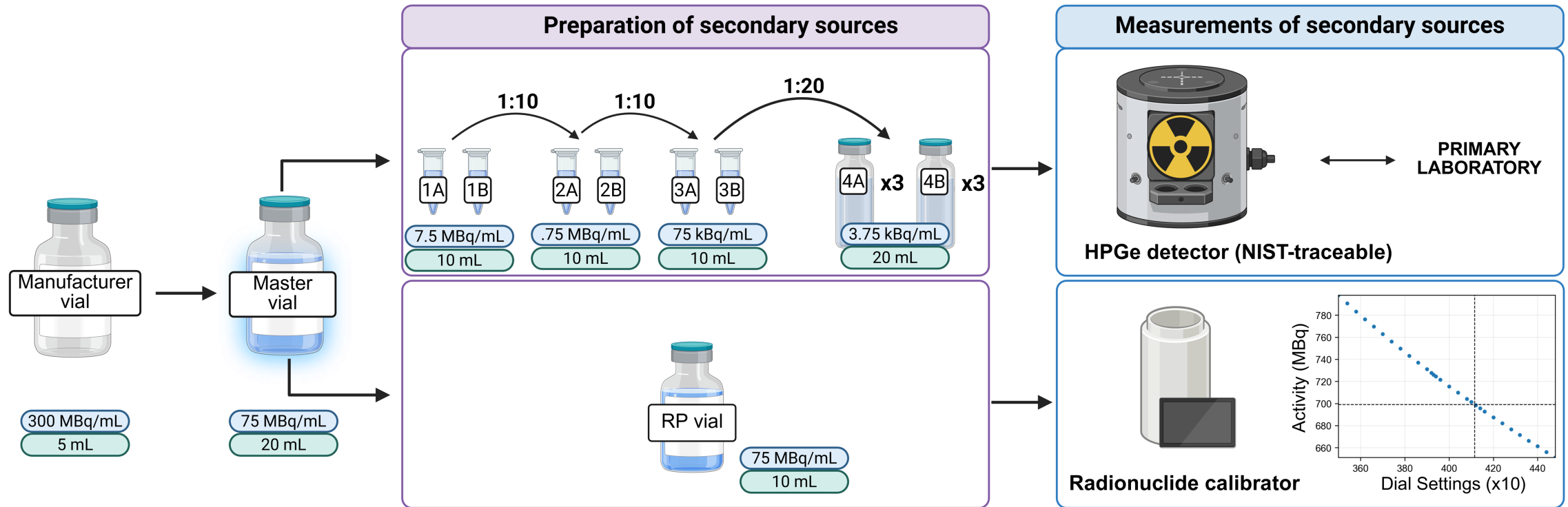
?

...

But there's a problem in the measurement chain

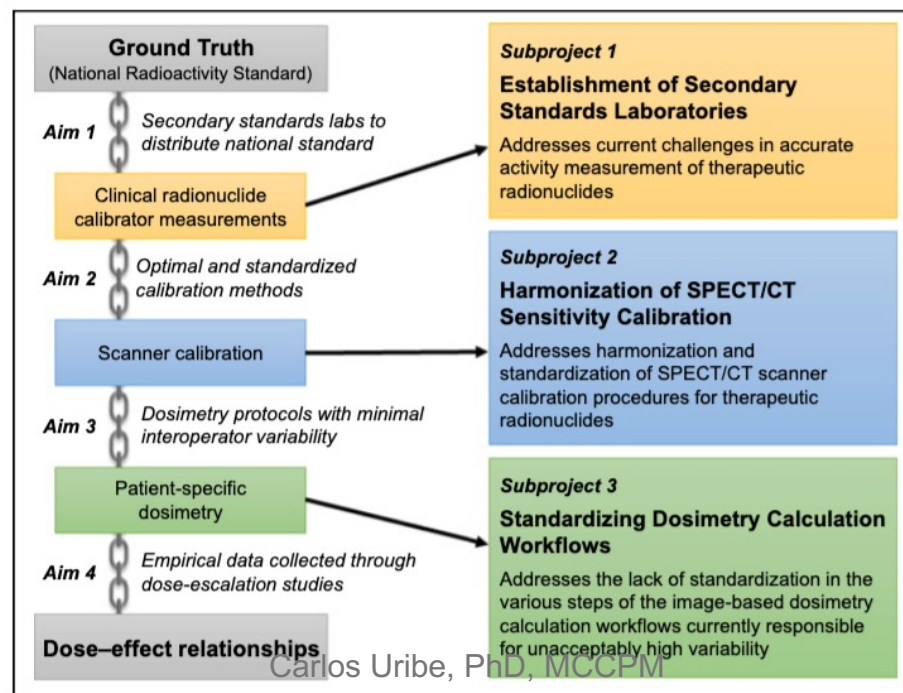


But there's a problem in the measurement chain



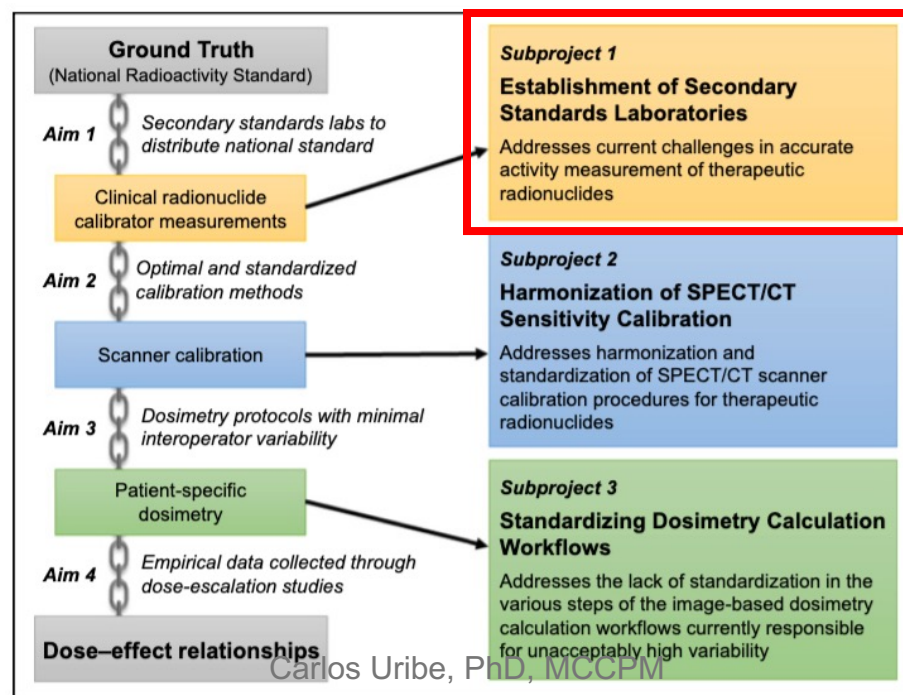
Pulling Together: A 5-Year Plan to Improve Theranostic Outcomes by Improving the Accuracy of Dosimetry—An FNHI Joint Academic, Clinical, and Industrial Collaboration

Dale L. Bailey¹, John C. Dickson², Suzanne E. Lapi³, Carlos Uribe⁴, Clarita Saldarriaga Vargas⁵, Price Jackson⁶, Caroline Stokke⁷, Julia Brosch-Lenz⁸, Althea Lang⁹, Stephen A. Graves¹⁰, and John J. Sunderland¹⁰

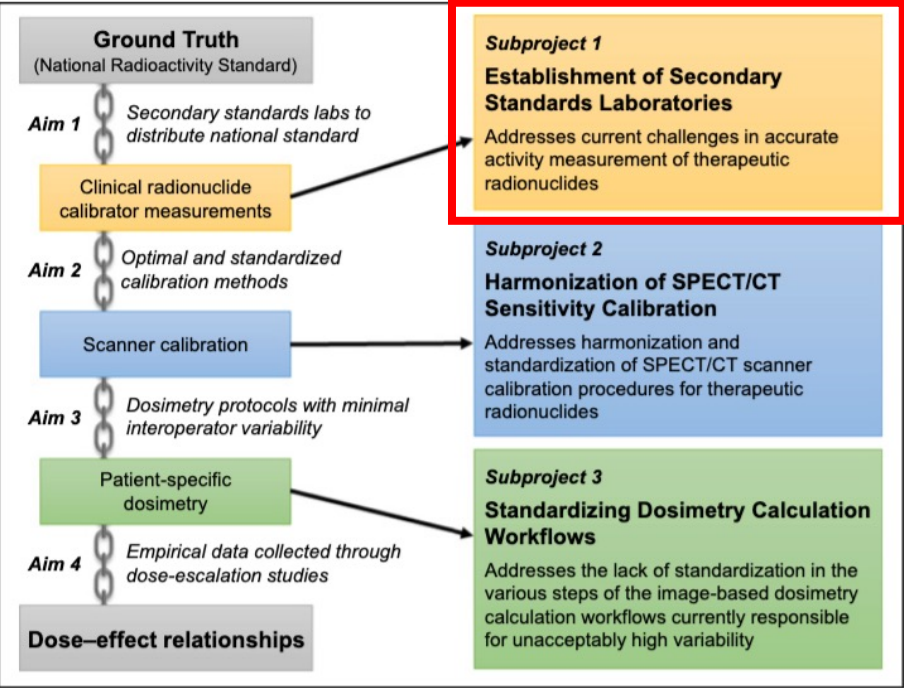


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HPGe spectrometers



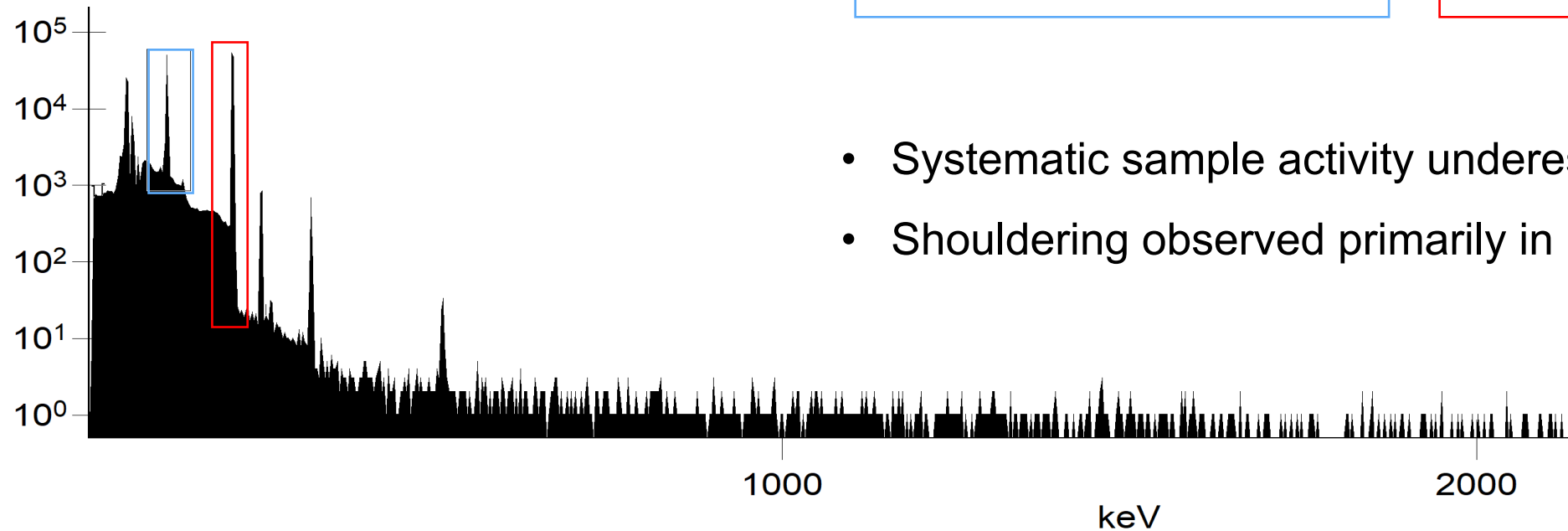
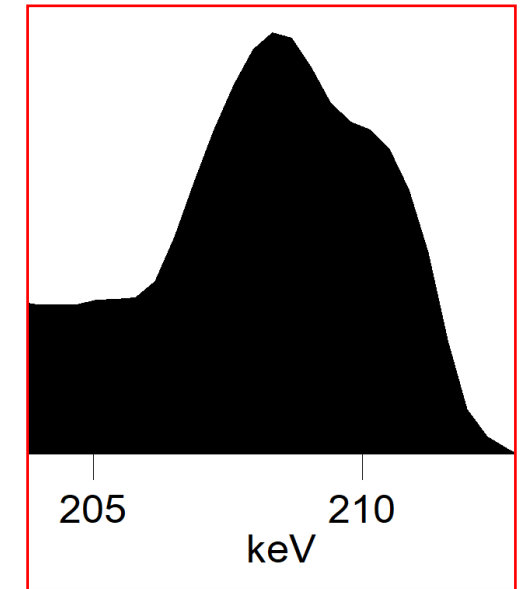
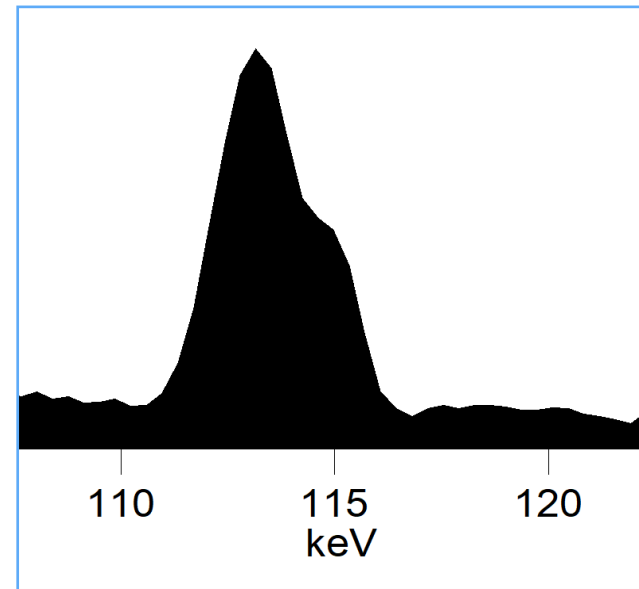
HPGe spectrometers



HPGe PDIB Test #1...

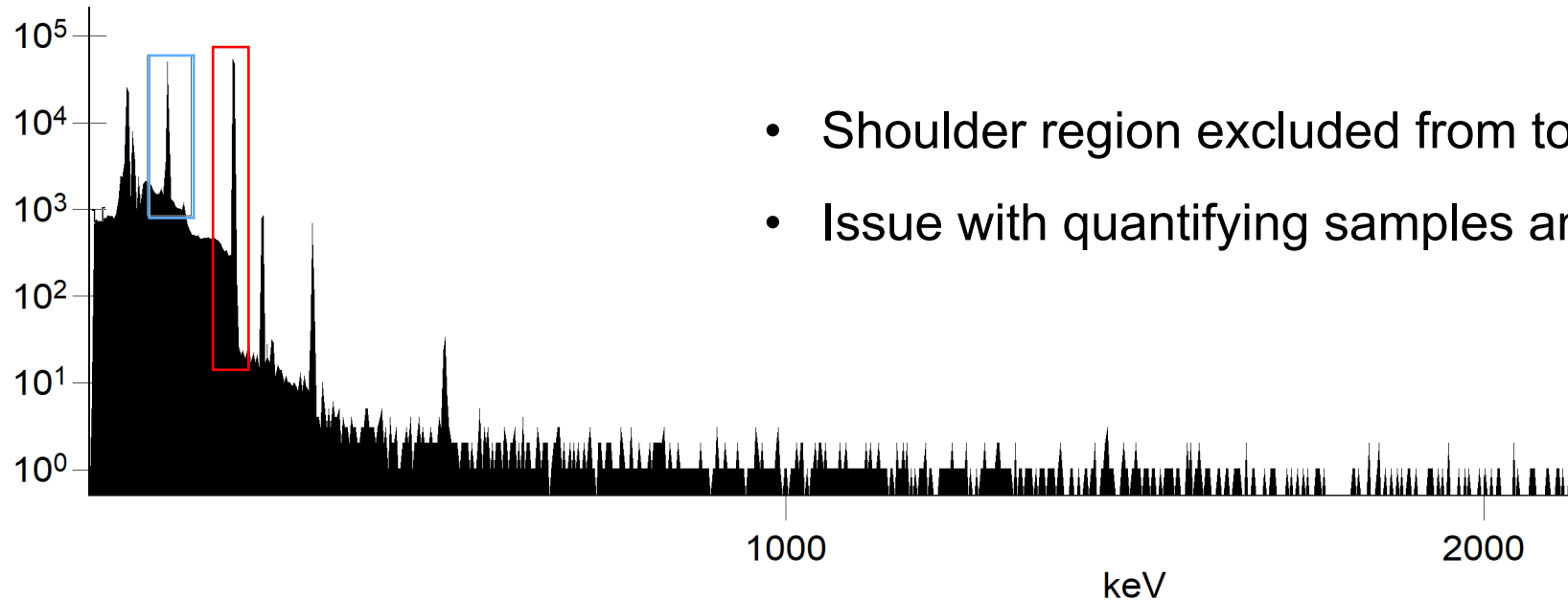
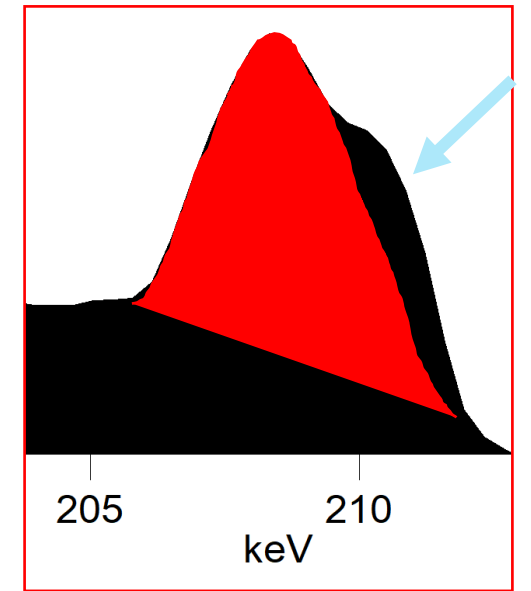
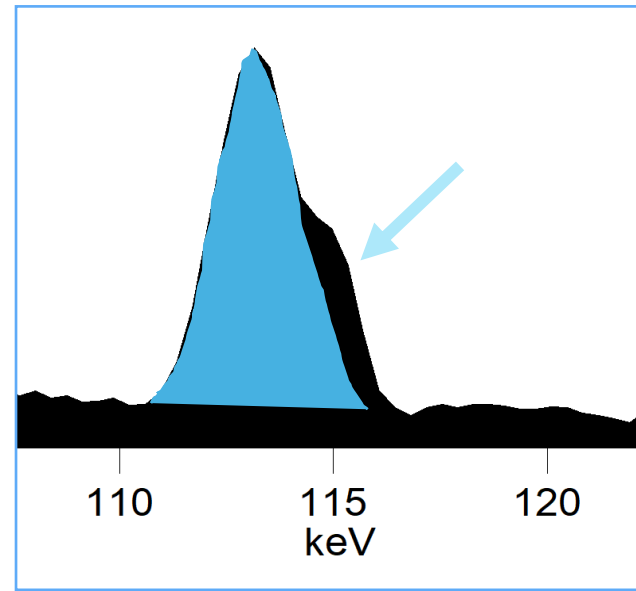


We Detected an Issue



- Systematic sample activity underestimation
- Shouldering observed primarily in low energy peaks

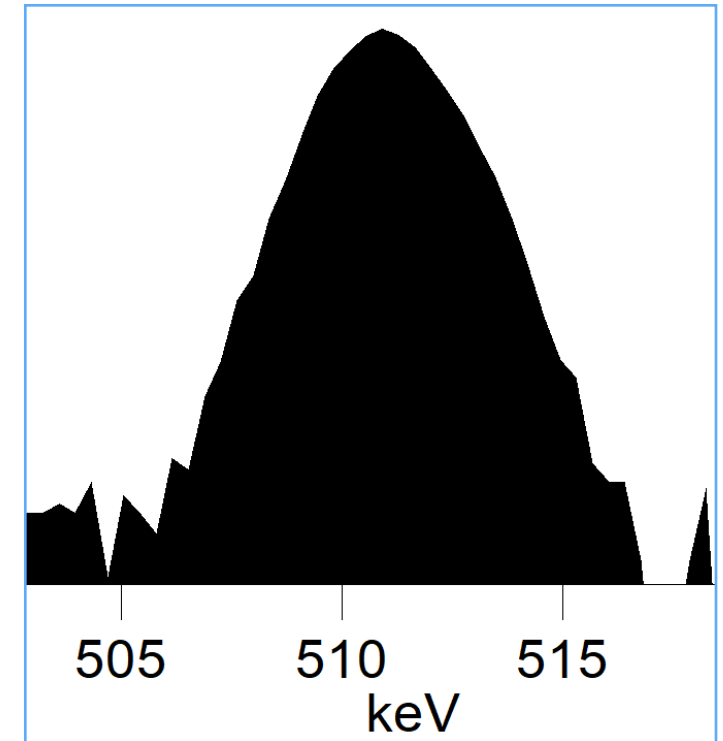
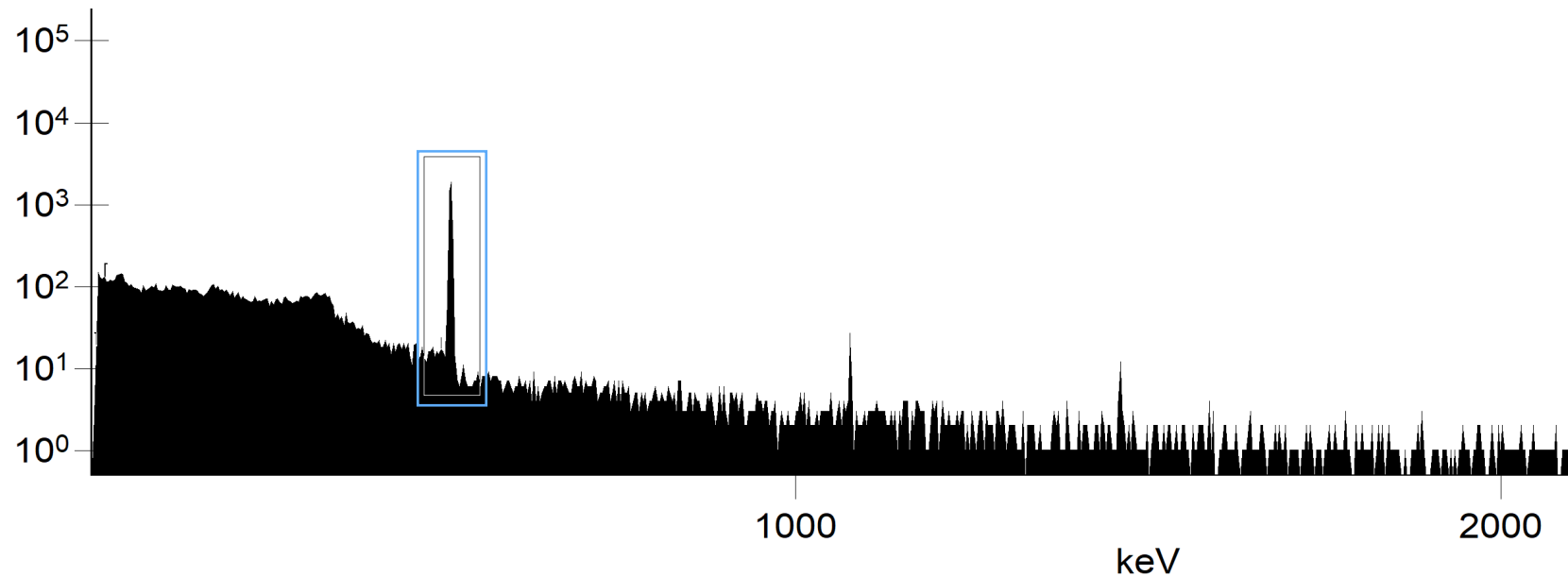
We Detected an Issue



- Shoulder region excluded from total peak counts
- Issue with quantifying samples and conducting efficiency calibration

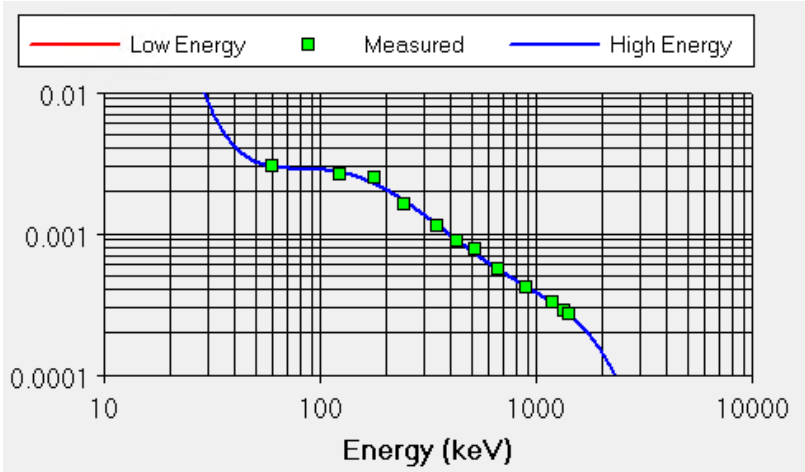
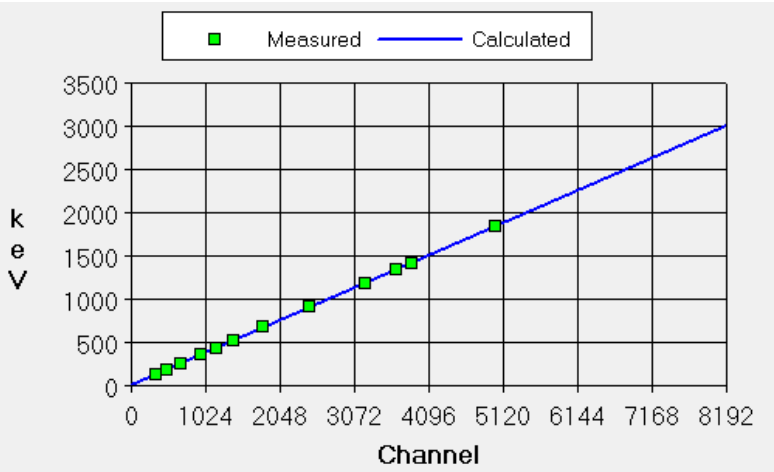
We Detected an Issue

- Detector primarily used for ^{18}F samples prior to 2025
- Shoulder region not visible in high FWHM peaks

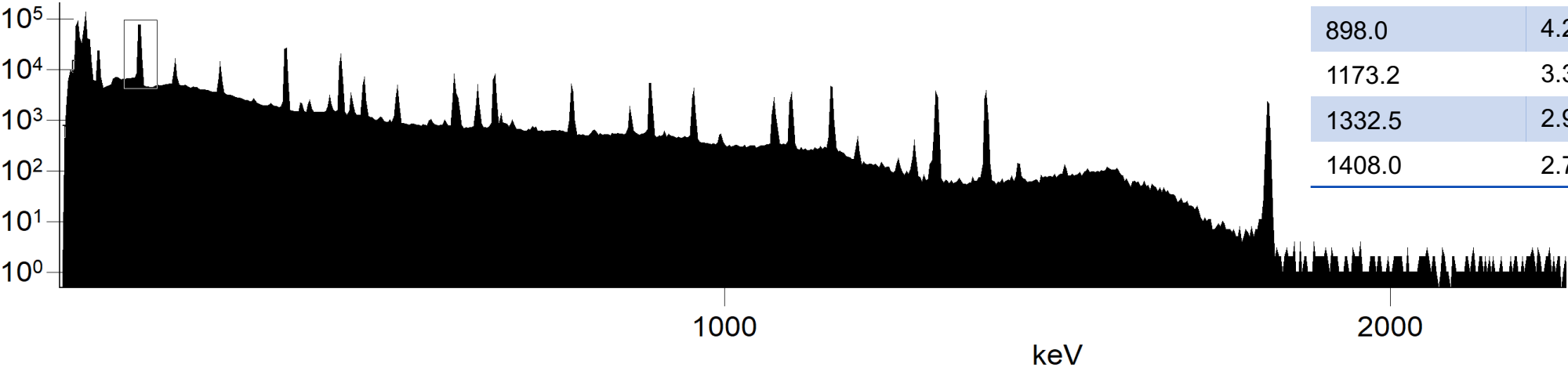


We Detected an Issue

Energy & efficiency calibration with shouldering observed in mixed standard

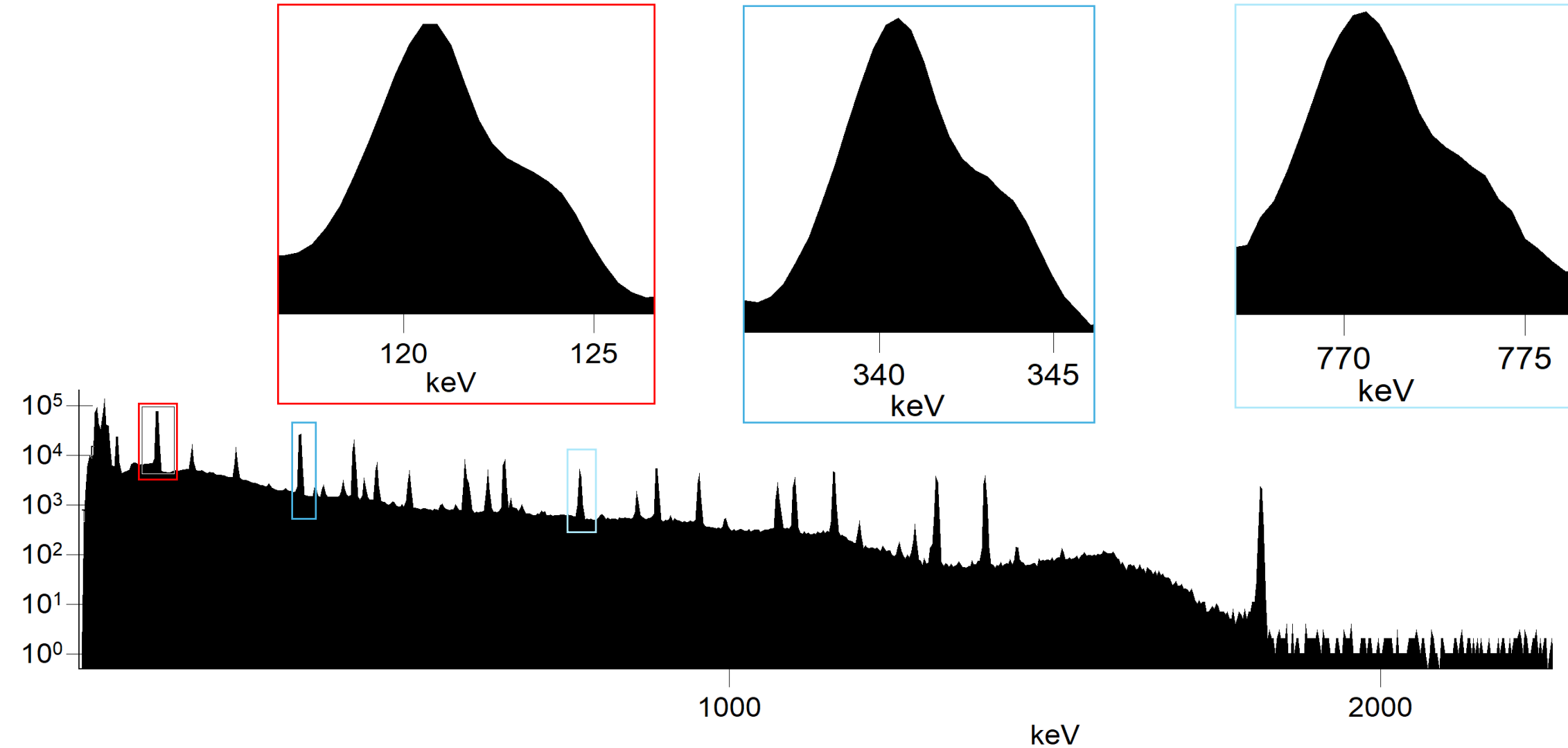


Peak Energy (keV)	Calculated Efficiency	Deviation from Measured Efficiency (%)
59.5	2.96E-3	-0.22
121.8	2.72E-3	2.87
176.3	2.26E-3	-8.35
244.7	1.70E-3	5.51
344.3	1.16E-3	1.97
427.9	9.0E-4	1.06
514.0	7.3E-4	-4.62
661.7	5.6E-4	0.46
898.0	4.2E-4	1.19
1173.2	3.3E-4	-0.19
1332.5	2.9E-4	1.50
1408.0	2.7E-4	-1.40

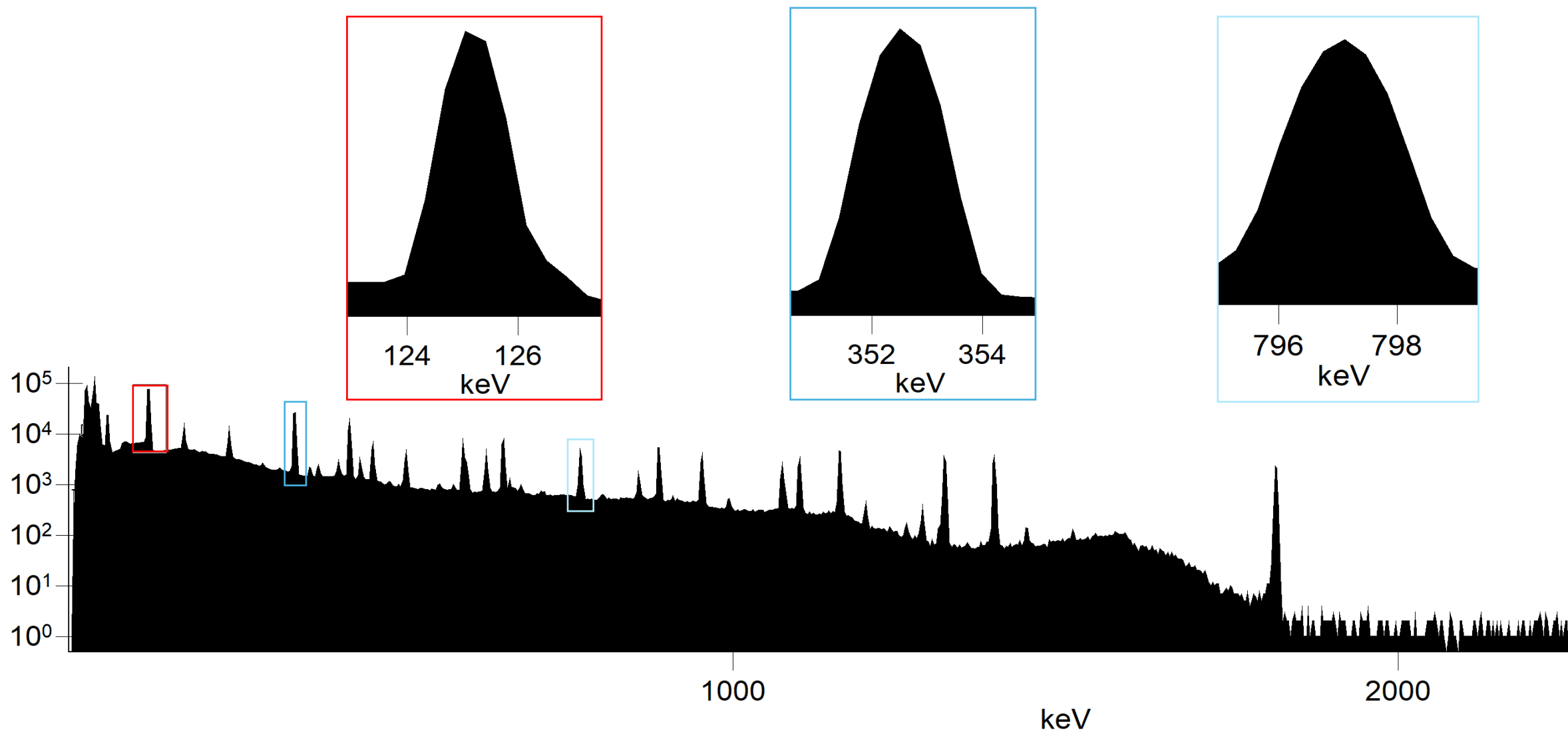


Mirion Support

1) Altered Flat Top & Rise Time → No change in shouldering

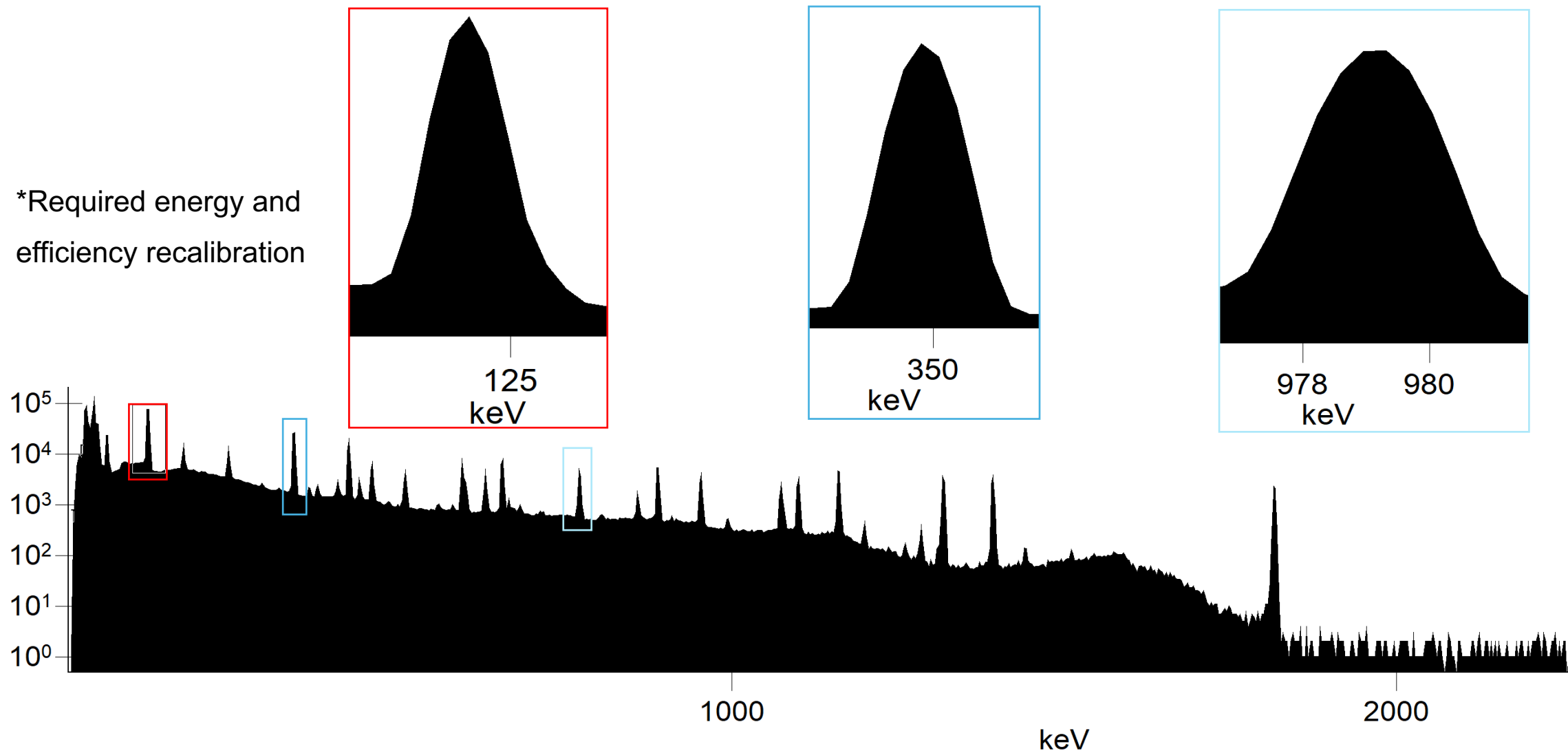


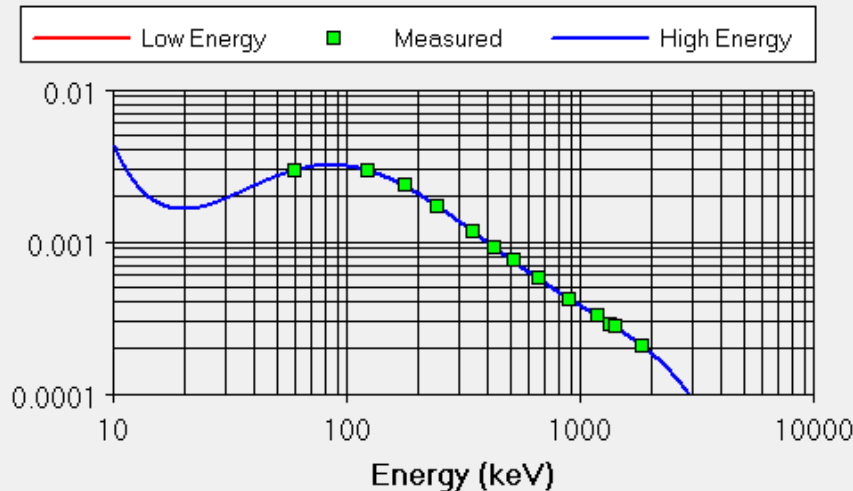
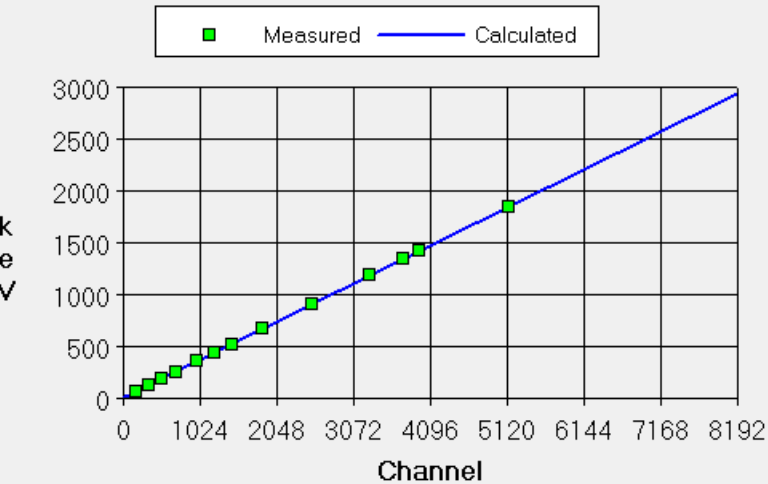
2) Thermal cycled detector → Shouldering removed, some high-energy tailing remaining



E&Z mixed standard, 10mL ISO glass vial, dead time: 2.05%

*Required energy and efficiency recalibration





Peak Energy (keV)	Calculated Efficiency	Deviation from Measured Efficiency (%)
59.5	2.95E-3	-0.08
121.8	2.93E-3	0.89
176.3	2.31E-3	-1.51
244.7	1.70E-3	-0.04
344.3	1.17E-3	1.18
427.9	9.1E-4	0.80
514.0	7.4E-4	-0.61
661.7	5.7E-4	-1.08
898.0	4.2E-4	0.13
1173.2	3.3E-4	-0.04
1332.5	2.9E-4	0.58
1408.0	2.7E-4	0.05
1836.1	2.1E-4	-0.21

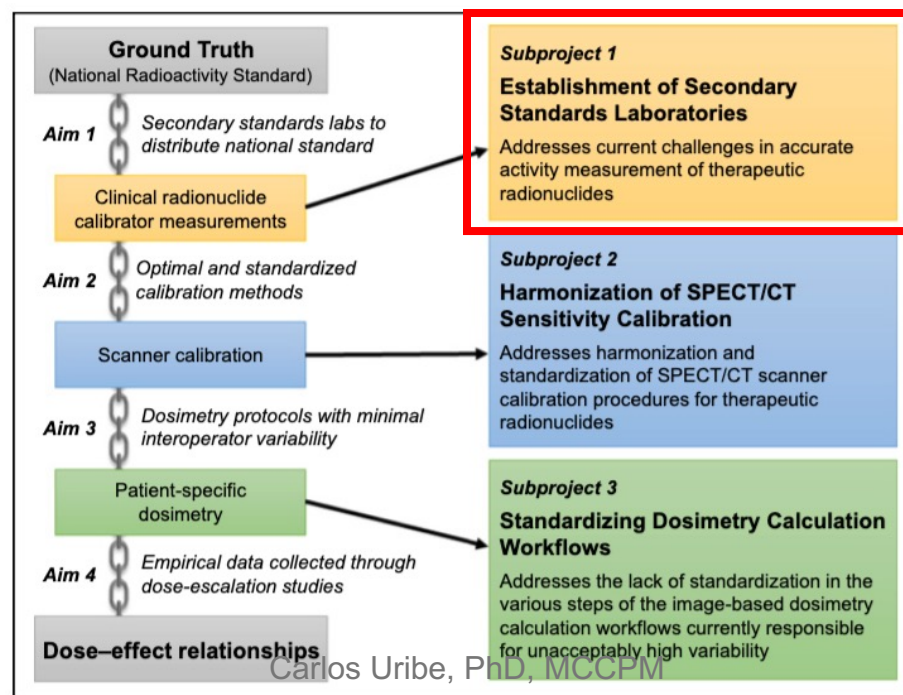
Deviation from national standards labs:

- Previously: **-5.0%**
- Now: **+1.1%**

Thank you to Henrik Persson, Steve Mettler, and the whole Mirion team for all the help

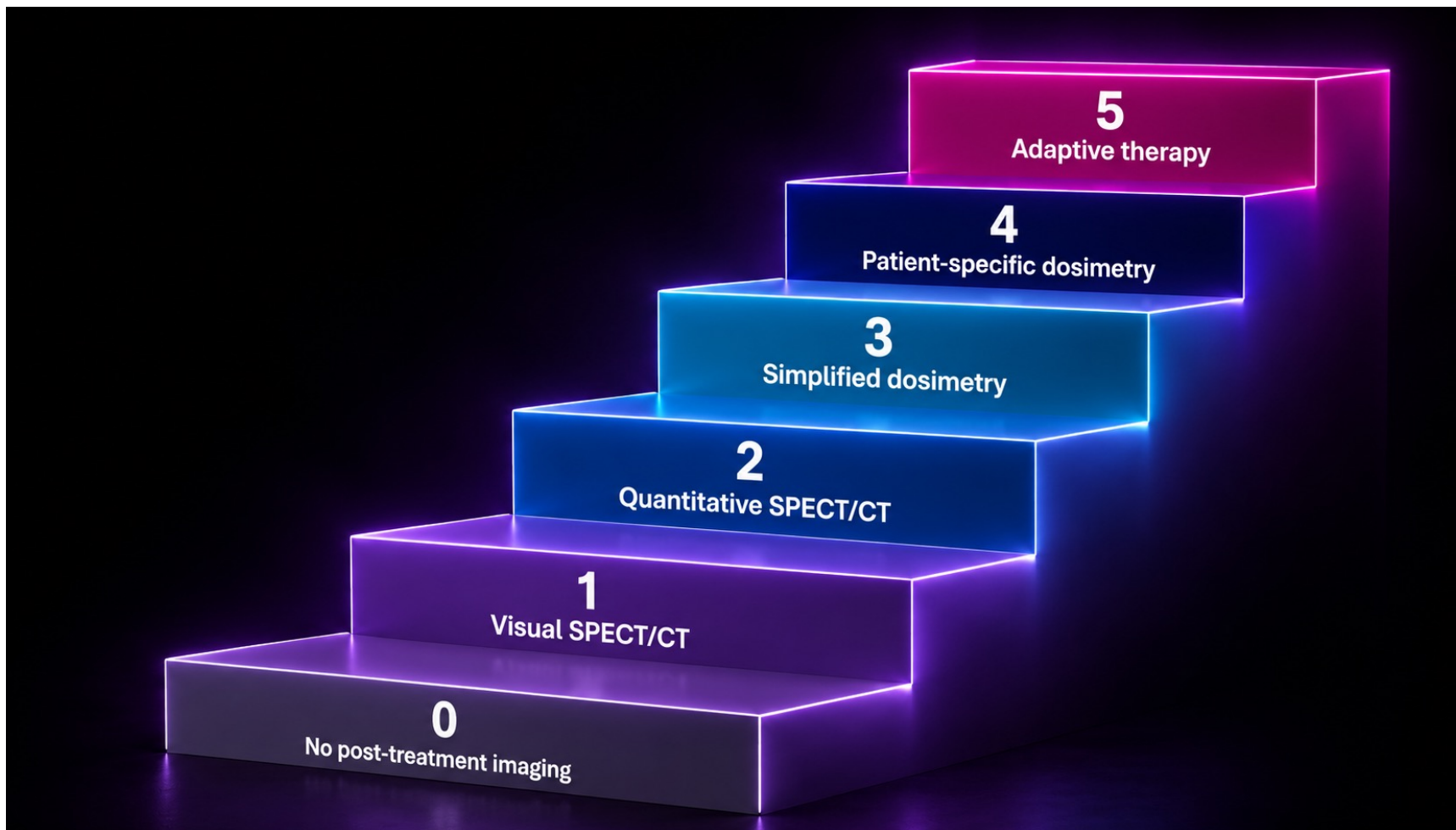
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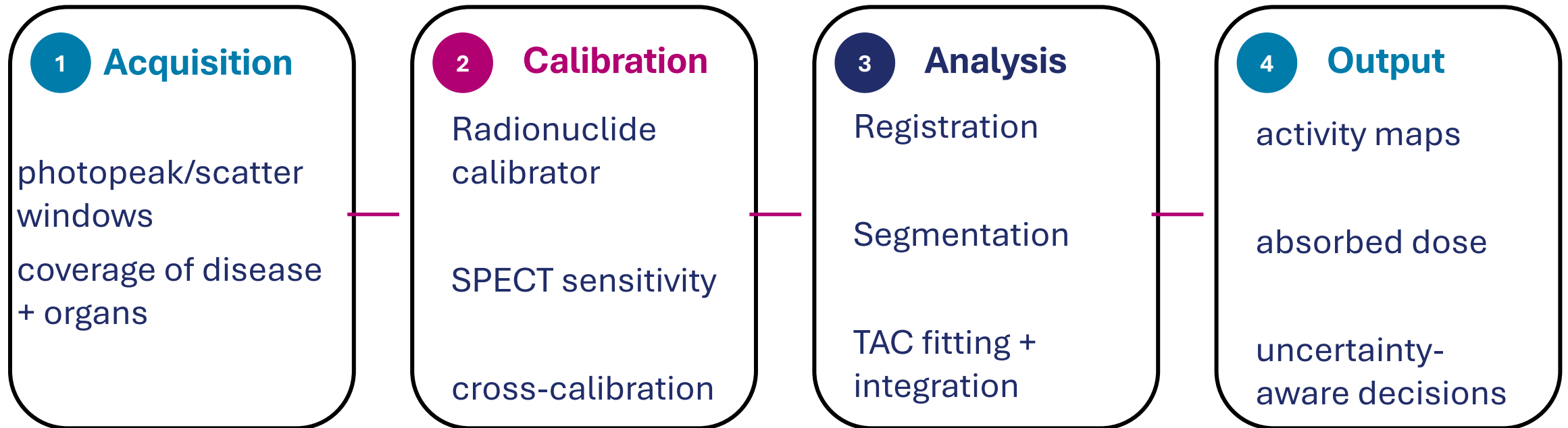


Dosimetry is a staircase, not an on/off switch

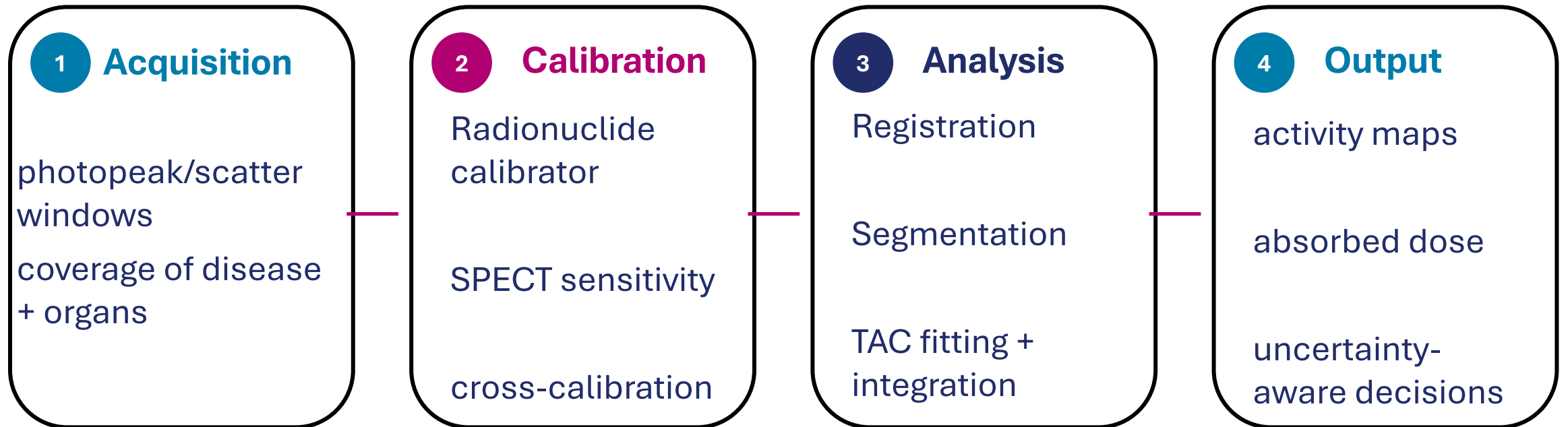
Start where the clinical workflow can support it. Build toward adaptive therapy.



Quantitative SPECT/CT is the foundation for dosimetry



Quantitative SPECT/CT is the foundation for dosimetry

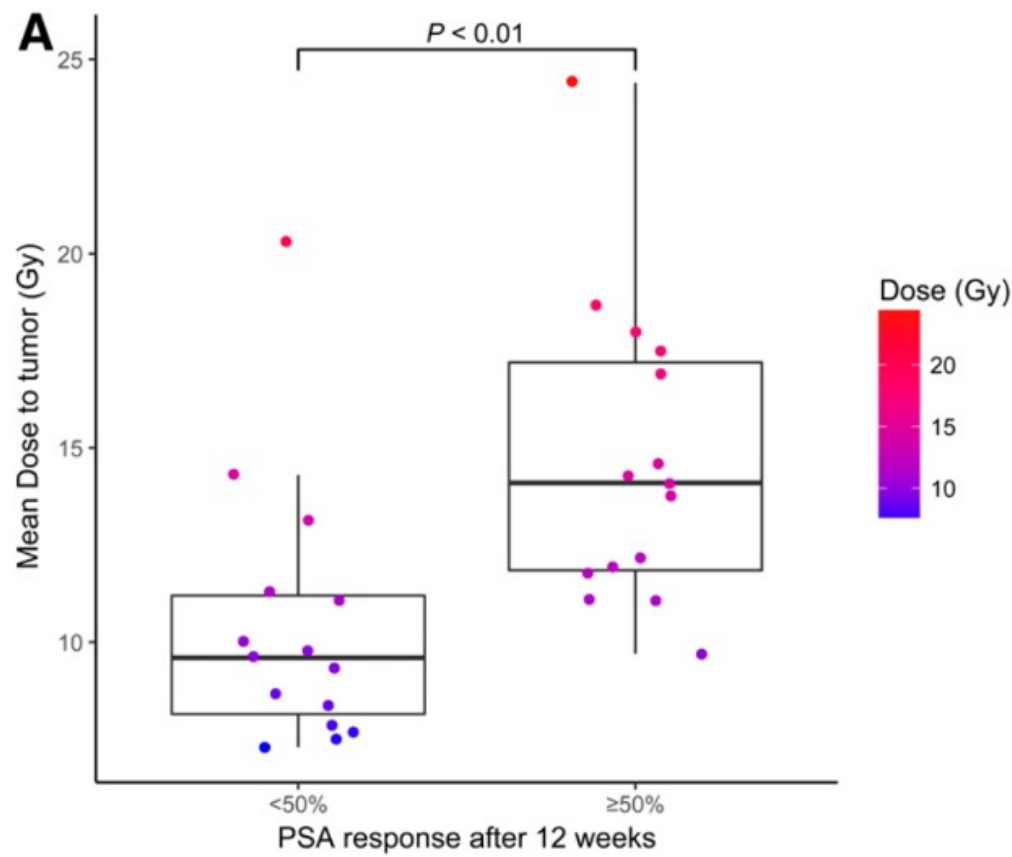


Without quantitative imaging, dosimetry is not a clinical measurement, it is just an assumption.



Carlos Uribe, PhD, MCCPM

Tumor absorbed dose is already linked to response in PSMA RPT



PSA ≥50% decline

Median tumor dose: 14.1 Gy

PSA <50% decline

Median tumor dose: 9.6 Gy

Low-dose group

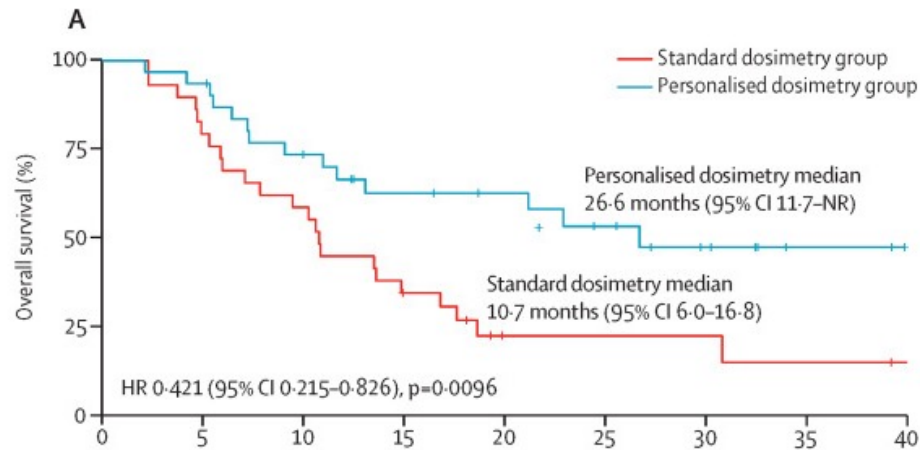
Only 1/11 patients below 10 Gy
achieved PSA ≥50% response

Violet et al. J Nucl Med. 2019;60:517–523

Personalized dosimetry has changed outcomes in RPTs

DOSISPHERE-01: proof of principle

Not the same disease or agent as ^{177}Lu RPT, but a powerful randomized example that absorbed-dose planning can change clinical outcomes.



Garin E, Tselikas L, Guiu B et al. The Lancet Gastroenterology & Hepatology, 2020; 6, 17-29

Response

↑ objective response

Survival

↑ overall survival

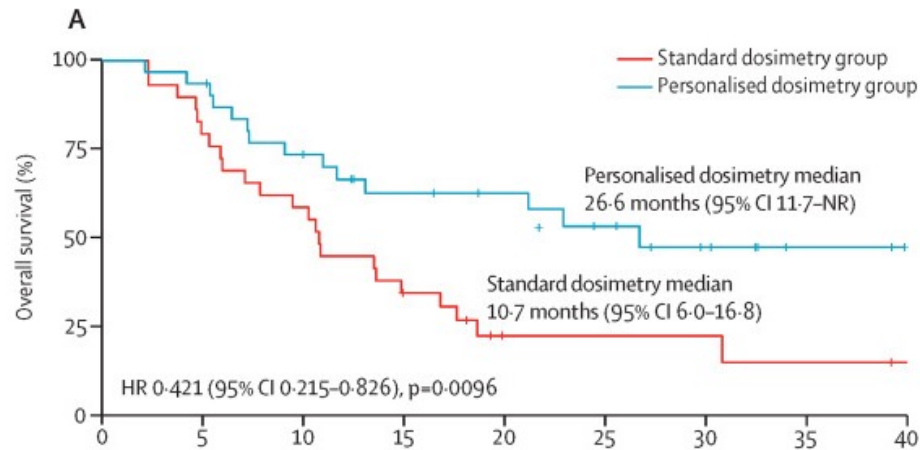
Downstaging

↑ surgical downstaging

Personalized dosimetry has changed outcomes in RPTs

DOSISPHERE-01: proof of principle

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Response

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Survival

↑ overall survival

Downstaging

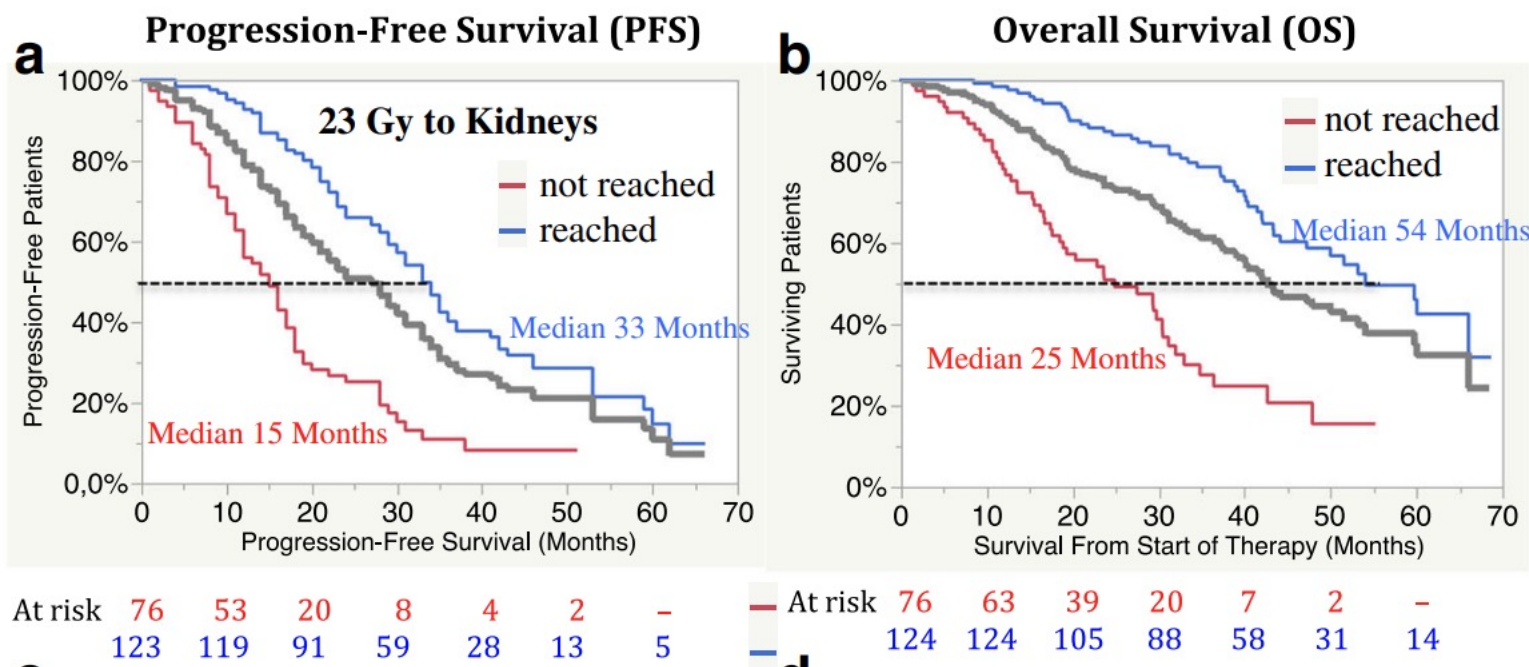
↑ surgical downstaging

The clinical question shifts from “what activity do we give?” to “what dose are we trying to deliver?”

Personalized dosimetry has changed outcomes in RPTs

Prospective observational study of ¹⁷⁷Lu-DOTA-octreotate therapy in 200 patients with advanced metastasized neuroendocrine tumours (NETs): feasibility and impact of a dosimetry-guided study protocol on outcome and toxicity

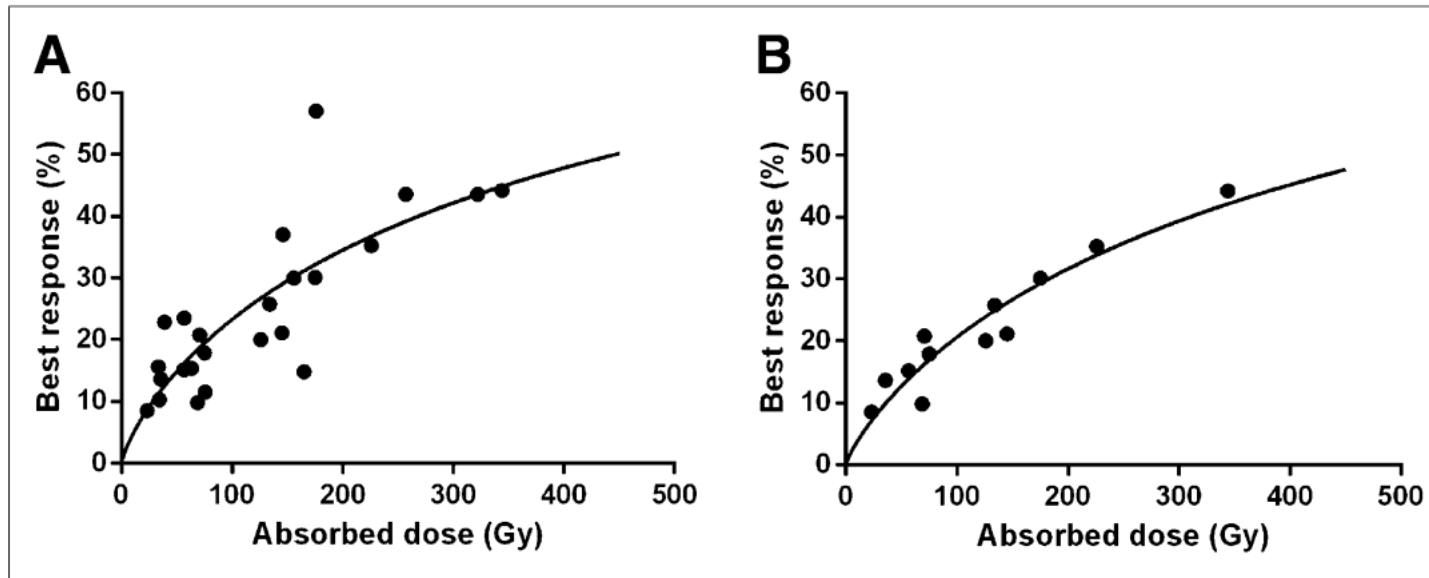
Ulrike Garske-Román^{1,2} • Mattias Sandström³ • Katarzyna Fröss Baron⁴ • Lars Lundin² • Per Hellman² • Staffan Welin⁴ • Silvia Johansson⁴ • Tanweera Khan⁴ • Hans Lundqvist³ • Barbro Eriksson⁴ • Anders Sundin² • Dan Granberg⁴



Tumors respond better with higher absorbed doses

Dose Response of Pancreatic Neuroendocrine Tumors Treated with Peptide Receptor Radionuclide Therapy Using ^{177}Lu -DOTATATE

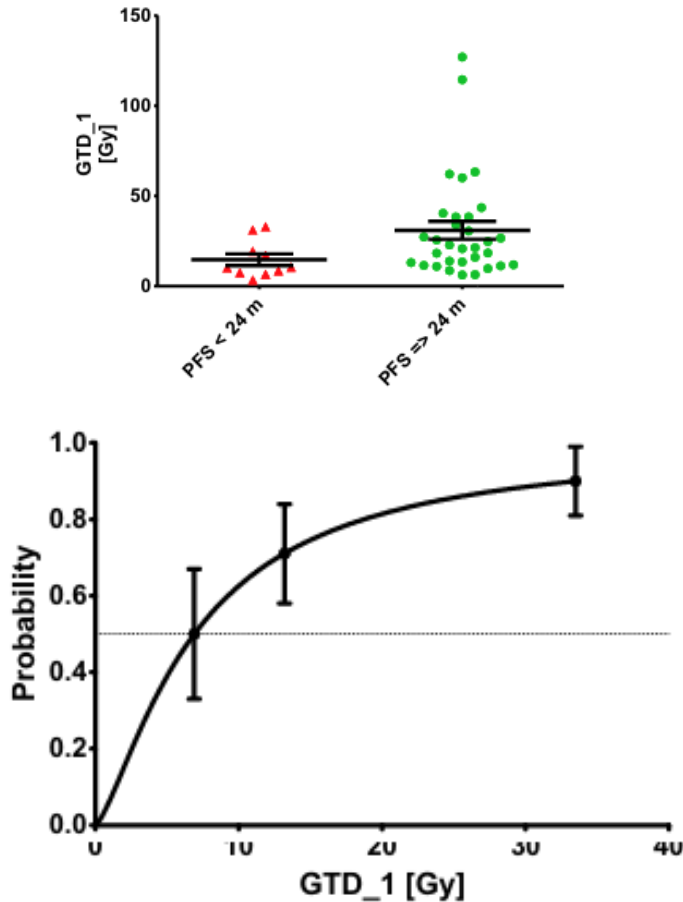
Ezgi Ilan^{1,2}, Mattias Sandström^{1,2}, Cecilia Wassberg^{1,3}, Anders Sundin^{1,3}, Ulrike Garske-Román^{1,3}, Barbro Eriksson⁴, Dan Granberg⁴, and Mark Lubberink^{1,2}



Tumors respond better with higher absorbed doses

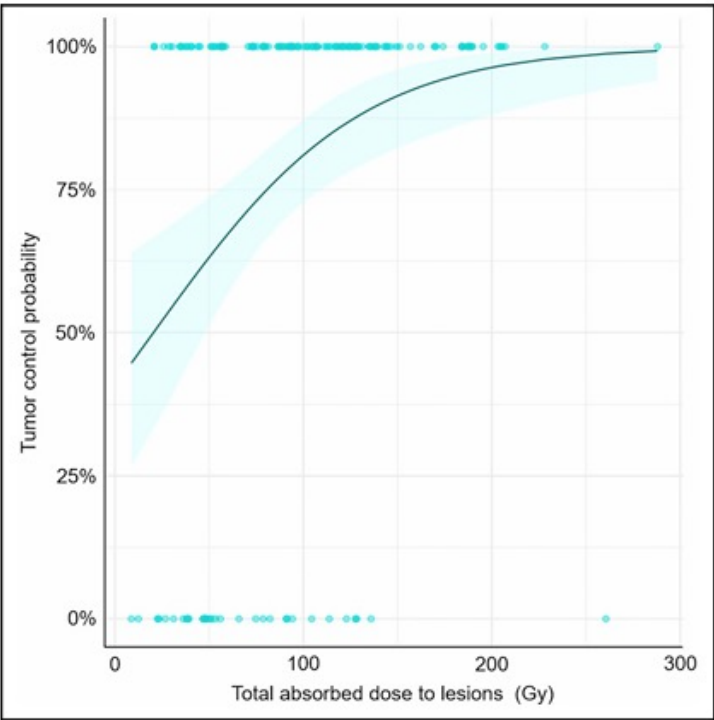
The LUTADOSE trial: tumour dosimetry after the first administration predicts progression free survival in gastro-entero-pancreatic neuroendocrine tumours (GEP NETs) patients treated with [¹⁷⁷Lu]Lu-DOTATATE

Marco Maccauro¹ · Mariarosaria Cuomo^{1,2} · Matteo Bauckneht^{3,4} · Matteo Bagnalasta¹ · Stefania Mazzaglia¹ · Federica Scalorbi¹ · Giovanni Argiroffi¹ · Margarita Kirienko¹ · Alice Lorenzoni¹ · Gianluca Aliberti¹ · Sara Pusceddu⁵ · Calareso Giuseppina⁶ · Garanzini Enrico Matteo⁶ · Ettore Seregni¹ · Carlo Chiesa¹



Absorbed Dose–Response Relationship in Patients with Gastroenteropancreatic Neuroendocrine Tumors Treated with [¹⁷⁷Lu]Lu-DOTATATE: One Step Closer to Personalized Medicine

Kévin Hebert^{*1} · Lore Santoro^{*1,2} · Maeva Monnier³ · Florence Castan³ · Ikrame Berkane¹ · Eric Assénat⁴ · Cyril Fersing^{1–5} · Pauline Gélibert⁶ · Jean-Pierre Pouget² · Manuel Bardiès^{1,2} · Pierre-Olivier Kotzki^{1,2} · and Emmanuel Deshayes^{1,2}



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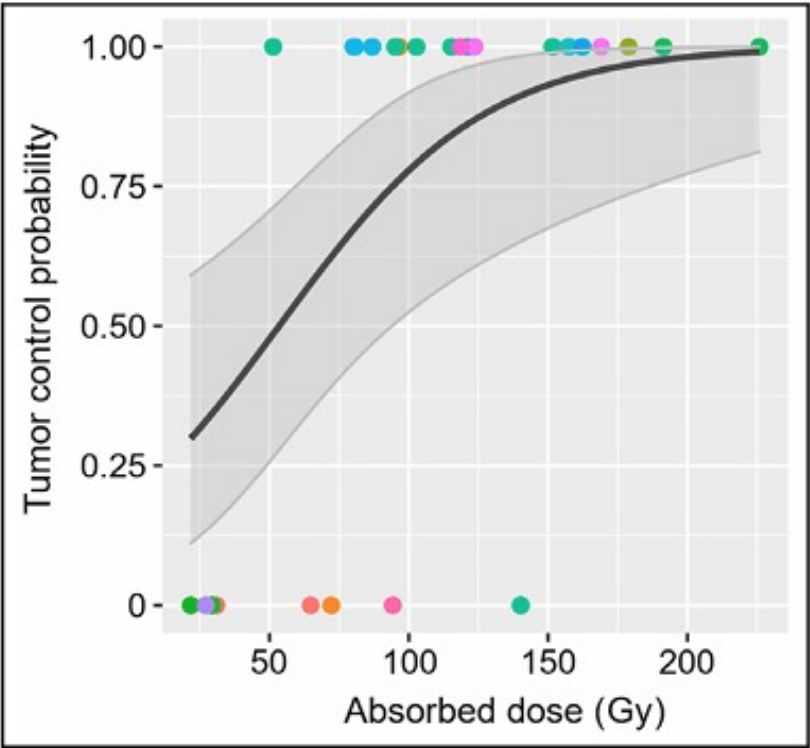
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Carlos Uribe, PhD, MCCPM

Tumors respond better with higher absorbed doses

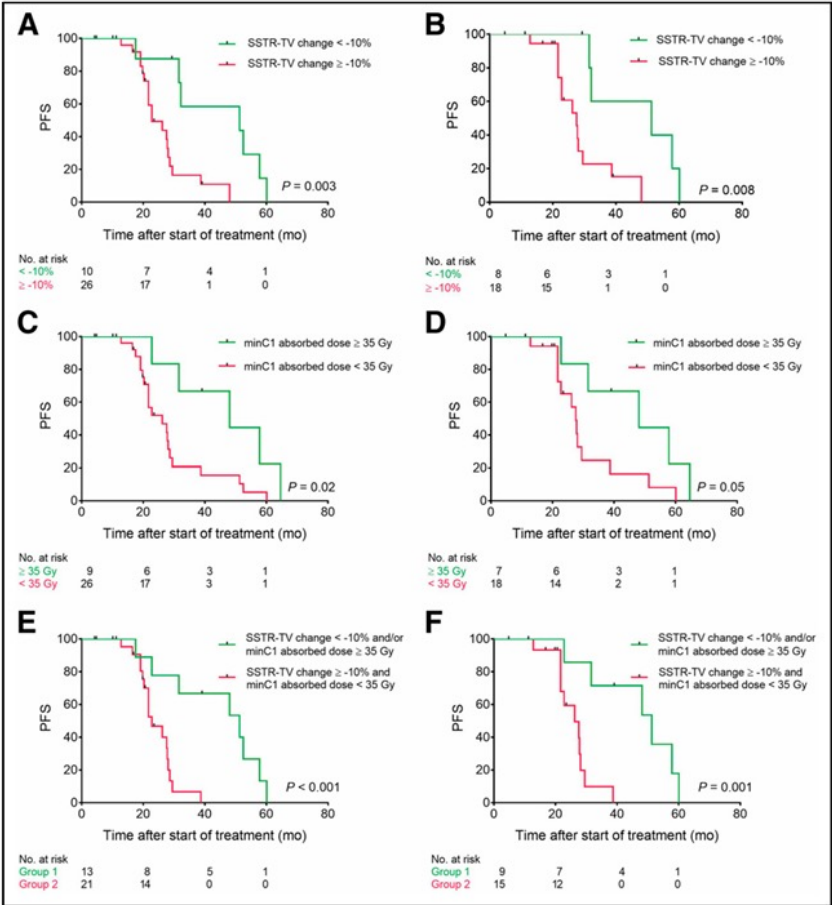
Relationship Between Absorbed Dose and Response in Neuroendocrine Tumors Treated with [¹⁷⁷Lu]Lu-DOTATATE

Carl Fredrik Warfvinge^{1,2}, Johan Gustafsson³, Daniel Roth³, Jan Tennvall², Johanna Svensson⁴, Peter Bernhardt^{5,6}, Anna Åkesson⁷, Elinore Wieslander¹, Anna Sundlöv², and Katarina Sjögreen Gleisner³



Prediction of ¹⁷⁷Lu-DOTATATE PRRT Outcome Using Multimodality Imaging in Patients with Gastroenteropancreatic Neuroendocrine Tumors: Results from a Prospective Phase II LUMEN Study

Magdalena Mileva¹, Gwennaëlle Marin², Hugo Levillain², Carlos Artigas¹, Camille Van Bogaert³, Clémentine Marin², Rachele Danieli², Amelie Deleporte⁴, Simona Picchia⁵, Konstantinos Stathopoulos⁵, Christiane Jungels⁴, Bruno Vanderlinden², Marianne Paesmans⁶, Lieveke Ameye⁶, Gabriela Critchi¹, Loubna Taraji-Schiltz¹, Chloe Velghe⁶, Zéna Wimana^{1,7}, Maria Bali⁵, Alain Hendlitz⁴, Patrick Flamen¹, and Ioannis Karfis¹



Dosimetry is about safety and opportunity

Safety

Identify patients with higher organ absorbed dose

Support risk-aware cycle decisions

Track cumulative exposure

Opportunity

Avoid systematic underdosing

Escalate when safe

Support retreatment and combinations

Learning

Build dose–response datasets

Refine organ limits

Improve trial design

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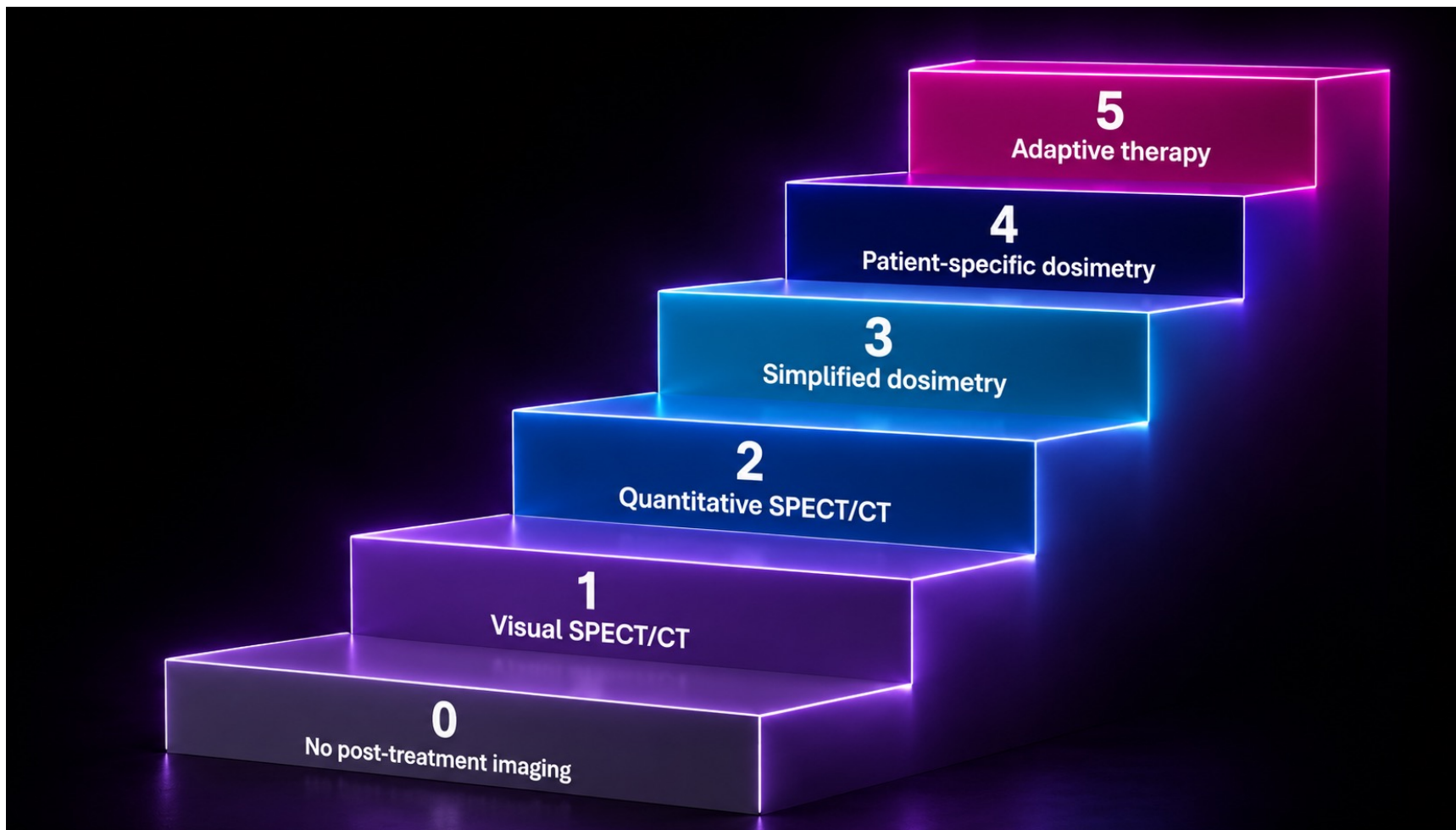
Improve trial design

A fixed activity can be simple and scalable, but it cannot tell us whether the patient was underdosed or near a limit.



Dosimetry is a staircase, not an on/off switch

Start where the clinical workflow can support it. Build toward adaptive therapy.



A practical clinical dosimetry workflow can start simple

Feasibility of Single-Time-Point Dosimetry for Radiopharmaceutical Therapies

Xinchi Hou¹, Julia Brosch², Carlos Uribe^{3,4}, Alessandro Desy^{5,6}, Guido Böning², Jean-Mathieu Beauregard^{5,6}, Anna Celler³, Arman Rahmim^{1,4,7}

Affiliations + expand

PMID: 33127625 PMCID: PMC8882881 DOI: 10.2967/jnumed.120.254656  

[Free PMC article](#)

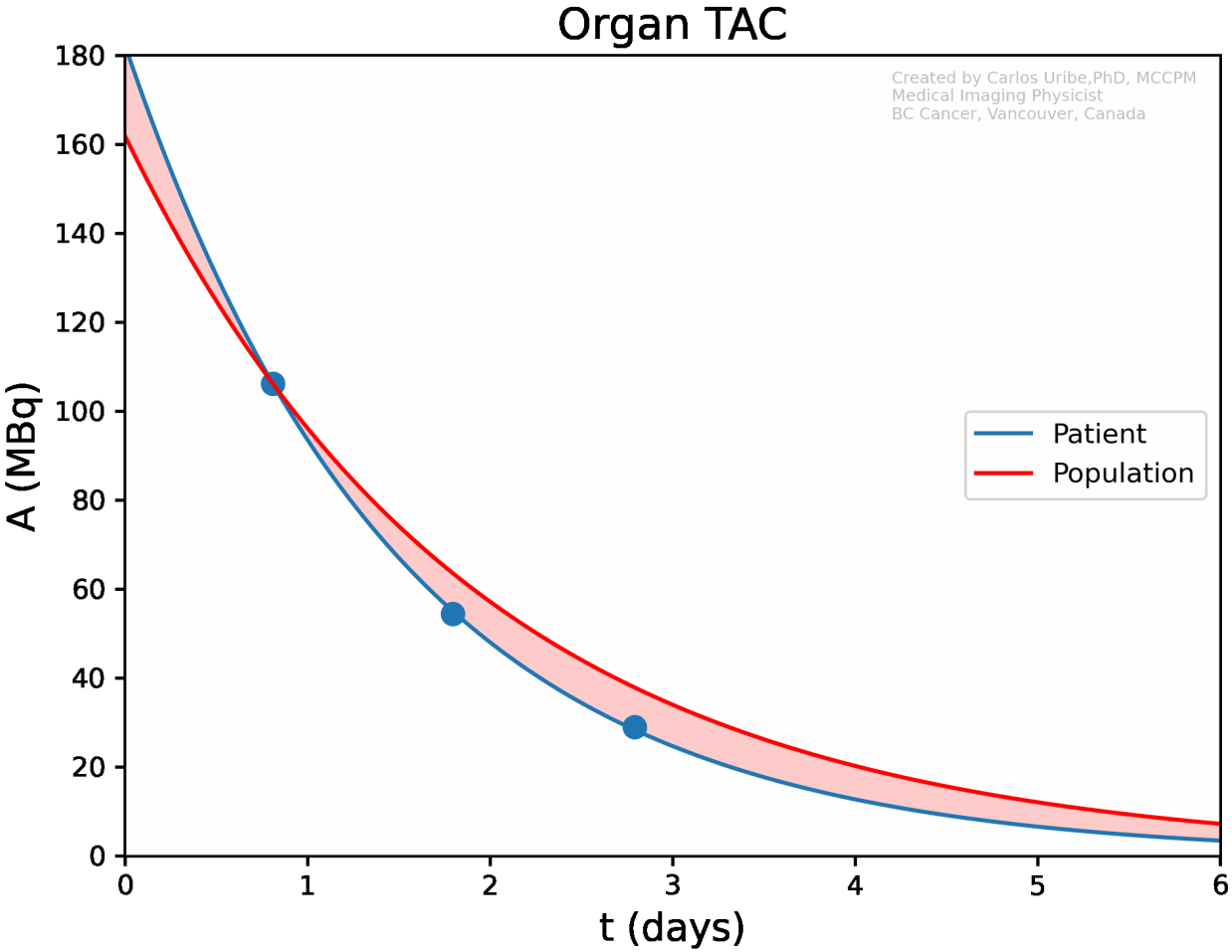
Toward Single-Time-Point Image-Based Dosimetry of ¹⁷⁷Lu-PSMA-617 Therapy

Julia Brosch-Lenz^{1,2}, Astrid Delker³, Friederike Völter³, Lena M Unterrainer³, Lena Kaiser³, Peter Bartenstein³, Sibylle Ziegler³, Arman Rahmim^{2,4,5}, Carlos Uribe^{4,6}, Guido Böning³

Affiliations + expand

PMID: 36657980 PMCID: PMC10152120 DOI: 10.2967/jnumed.122.264594  

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A practical clinical dosimetry workflow can start simple

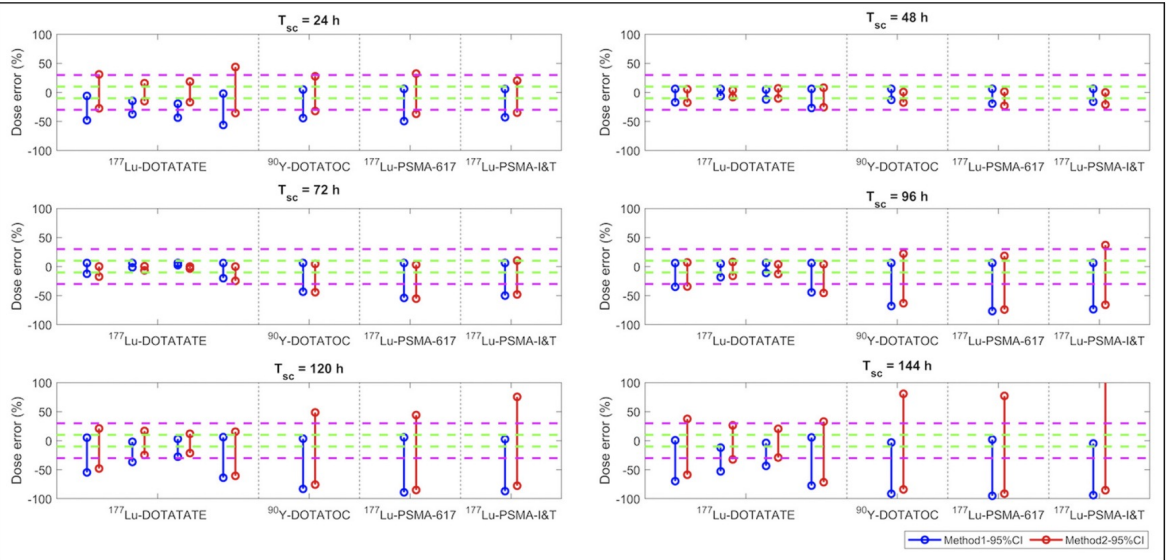
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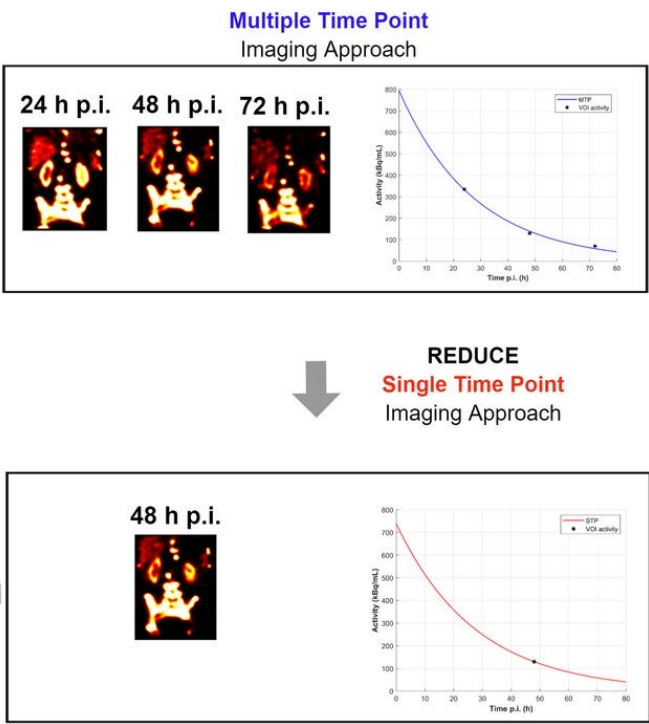
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Affiliations + expand

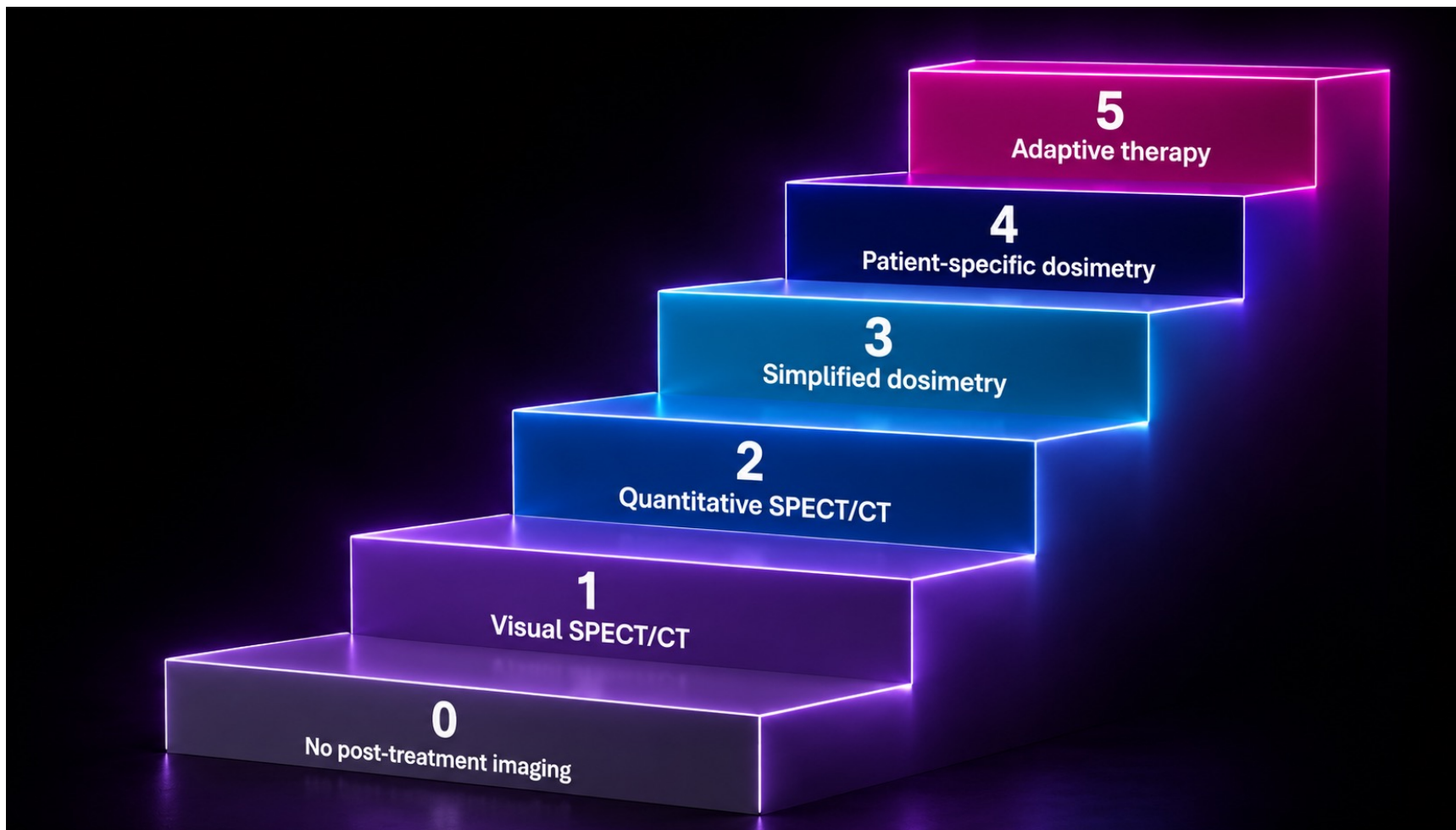
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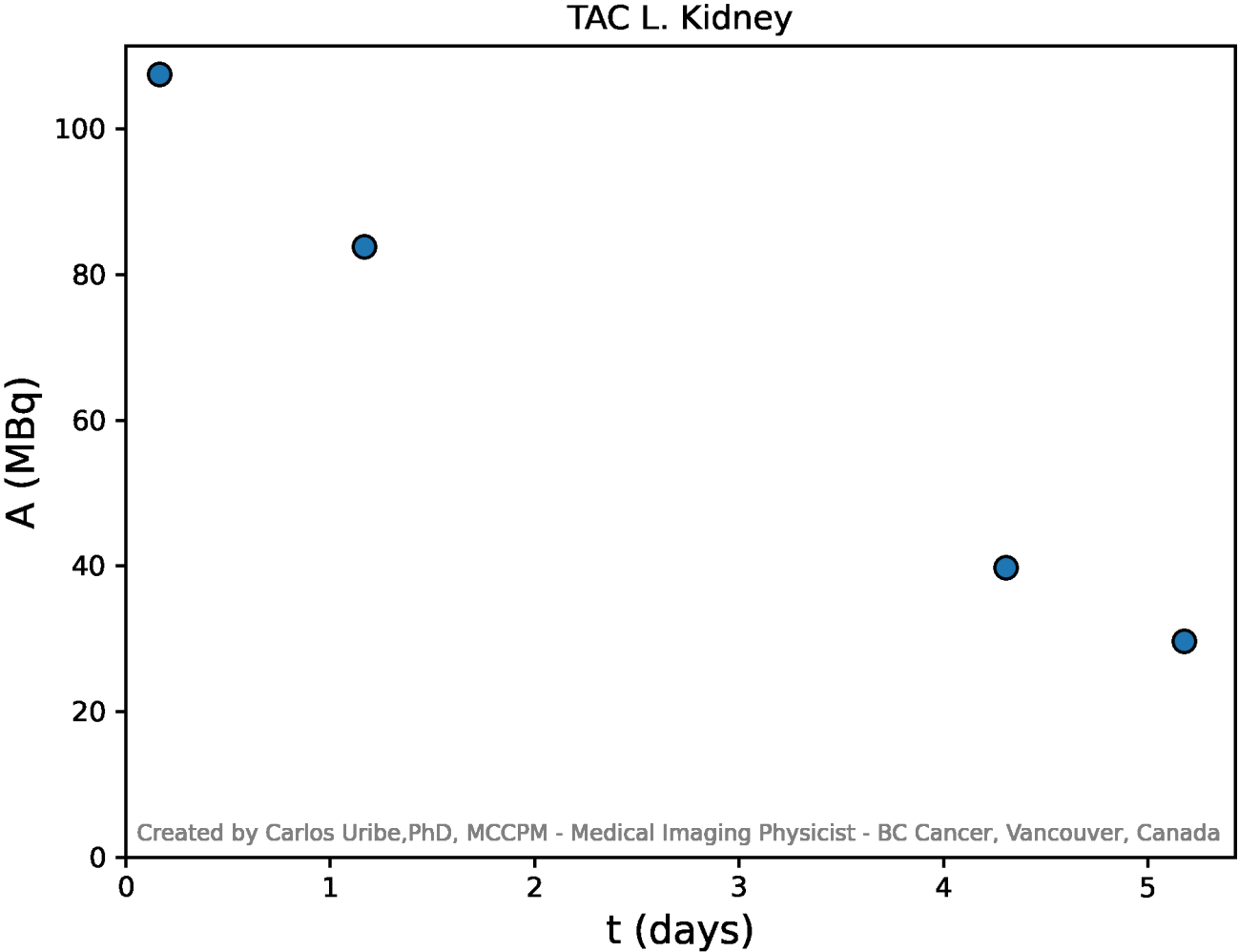
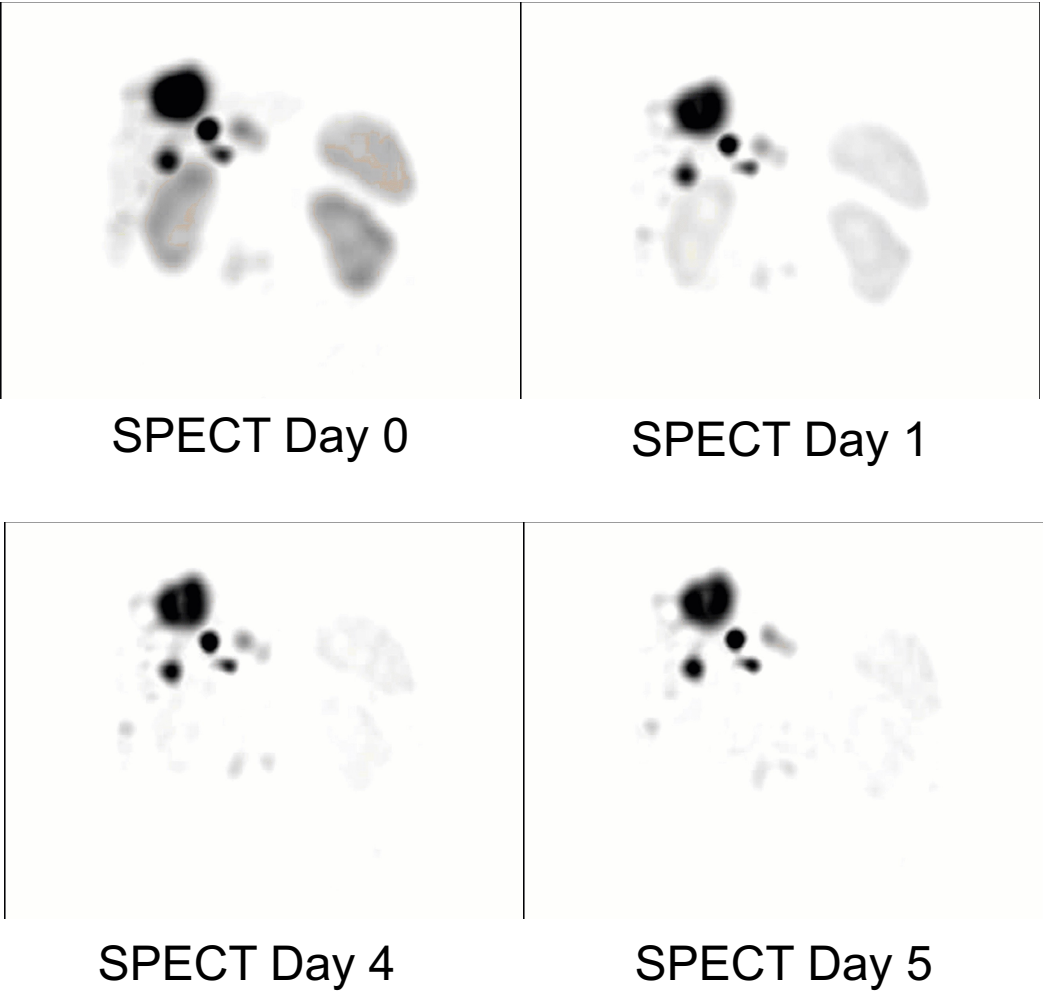


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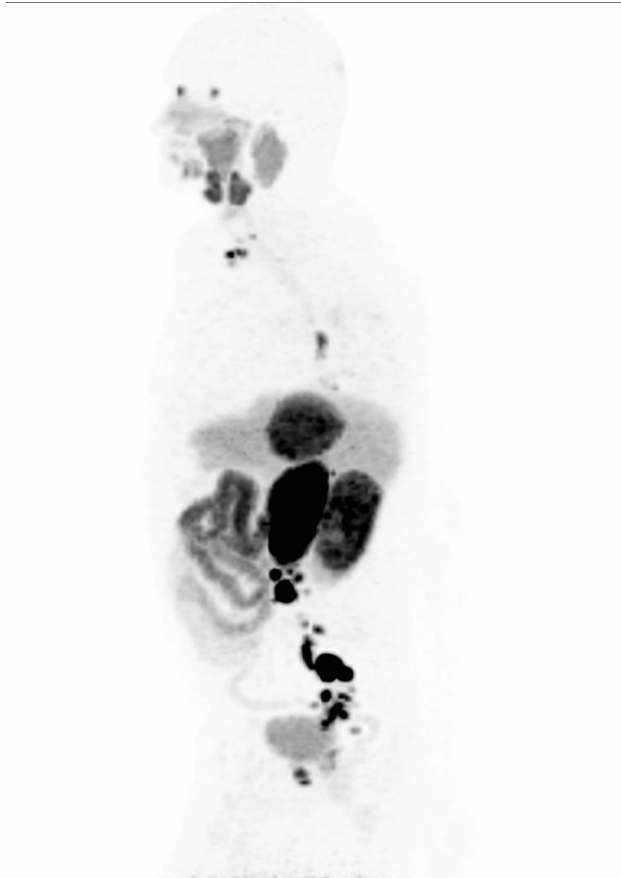


Patient specific

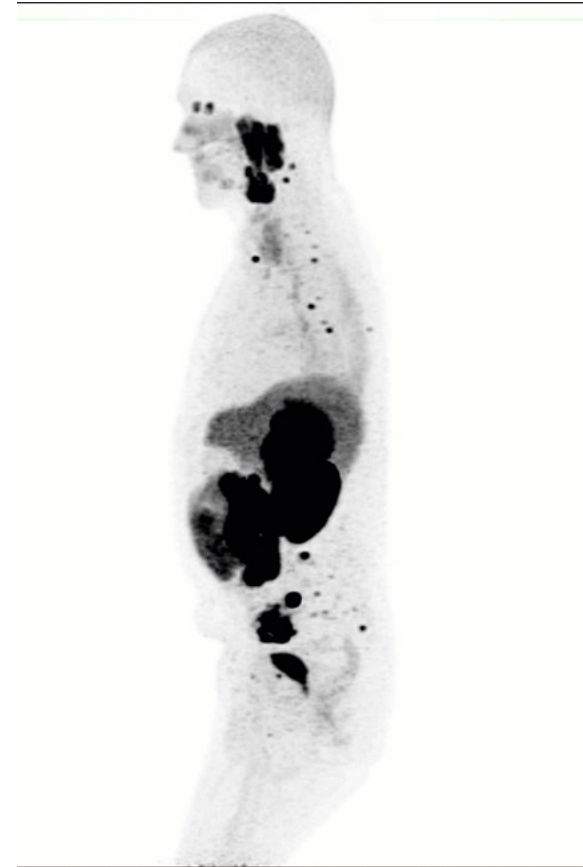


Patient specific

Mr. Ray Dioactive



Mr. Tom O'Grady



Patient specific

Mr. Ray Dioactive

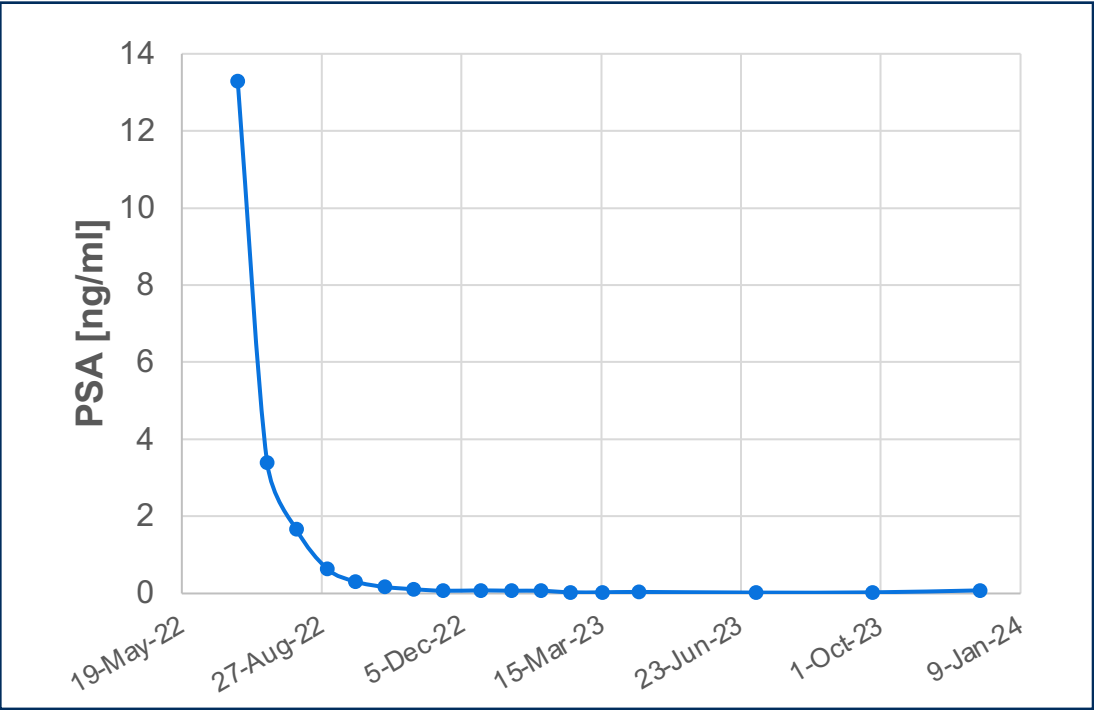
Injection		Tumors	
Cycle	Activity [MBq]	Cycle	Absorbed Dose [Gy]
1	7340	1	28.8
2	7700	2	14.2
3	7516	3	5.3
4	7908	4	7.5
5	7312	5	1.3
6	7432	6	0.7
		Total:	57.8

Mr. Tom O’Graphy

Injection		Tumors	
Cycle	Activity [MBq]	Cycle	Absorbed Dose [Gy]
1	8096	1	1.2
2	7357	2	2.2
3	8077	3	1.9
4	7795	4	1.4
5	7730	5	1.0
6	7384	6	1.3
		Total:	9

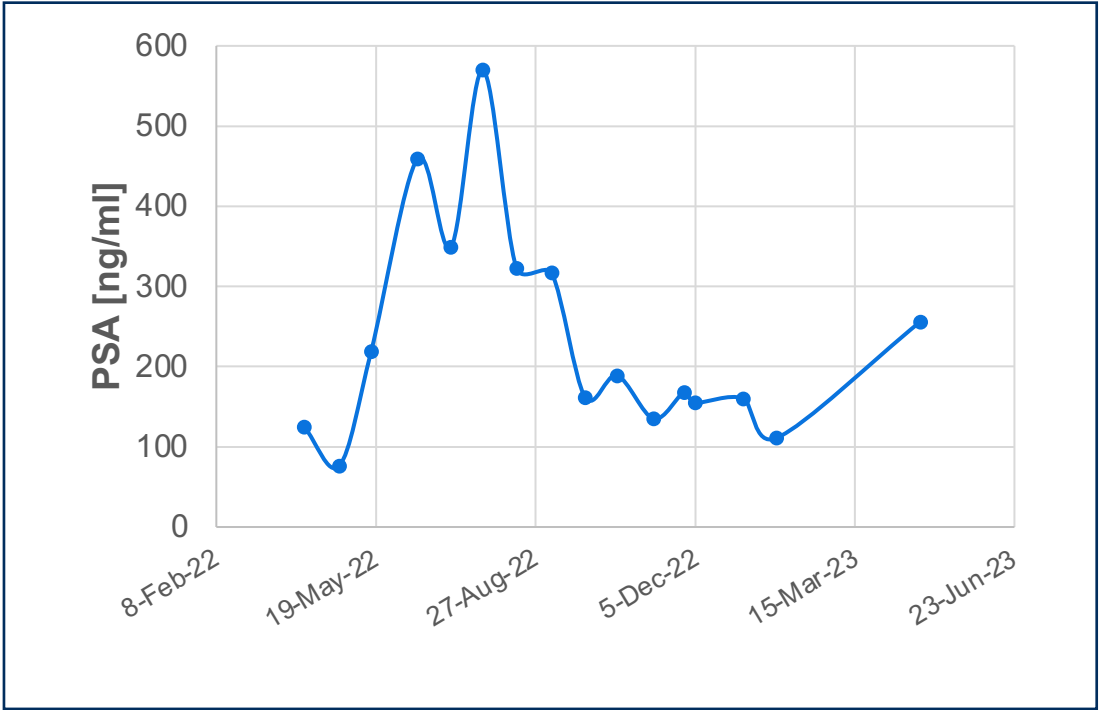
Patient specific

Mr. Ray Dioactive



Total: 57.8

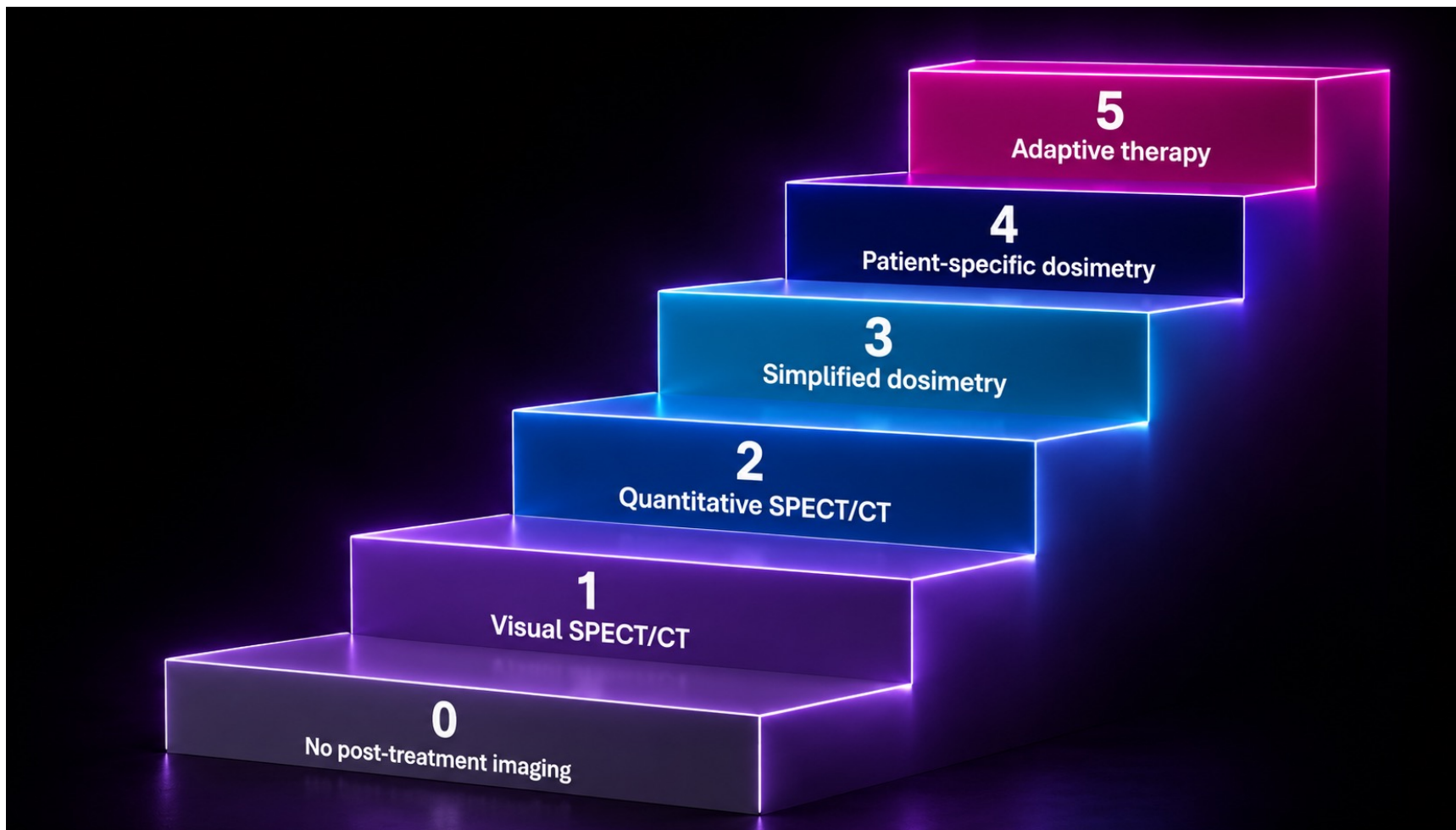
Mr. Tom O’Graphy



Total: 9

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¹⁷⁷Lu-HTK03170 Trial

Journal of Nuclear Medicine, published on April 30, 2025 as doi:10.2967/jnumed.124.269064

Lead Investigators:

- Don Wilson, MD
- Maryam Soleimani, MD
- Carlos Uribe, PhD, MCCPM

A Phase I/II Study of [¹⁷⁷Lu]Lu-HTK03170 with Personalized Dosimetry in Patients with Metastatic Castration-Resistant Prostate Cancer: Clinical Trial Protocol

Sara Harsini¹, Maryam Soleimani², Don Wilson¹, Carlos Uribe^{1,3}, Heather Sapruncoff¹, Ingrid Bloise¹, and Francois Benard^{1,3}

¹Department of Molecular Imaging and Therapy, BC Cancer, Vancouver, British Columbia, Canada; ²Department of Medical Oncology, BC Cancer, Vancouver, British Columbia, Canada; and ³Department of Radiology, University of British Columbia, Vancouver, British Columbia, Canada

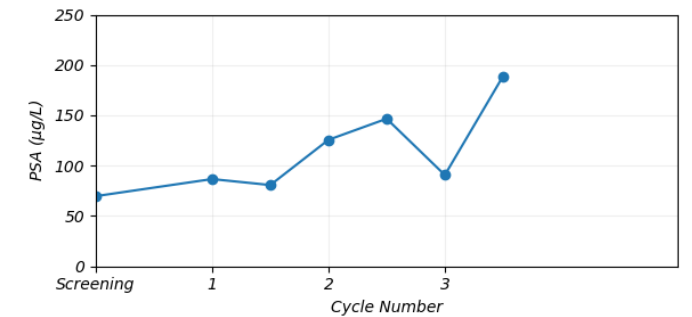
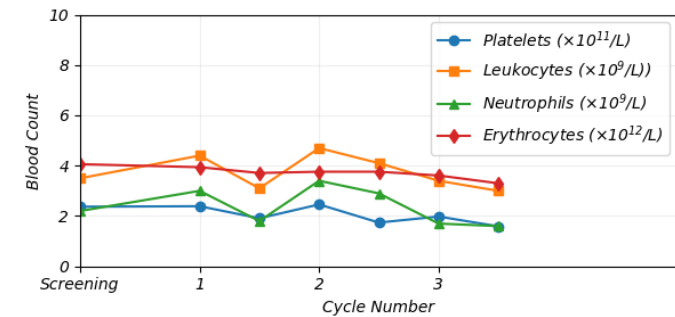
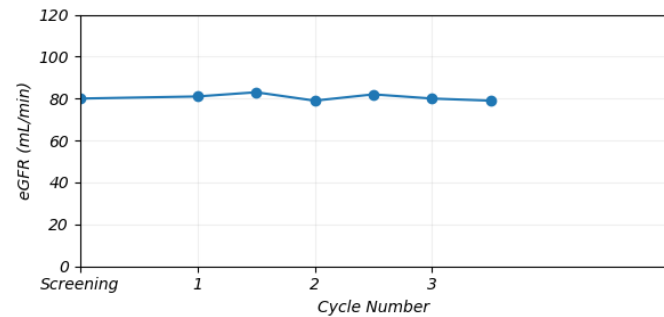
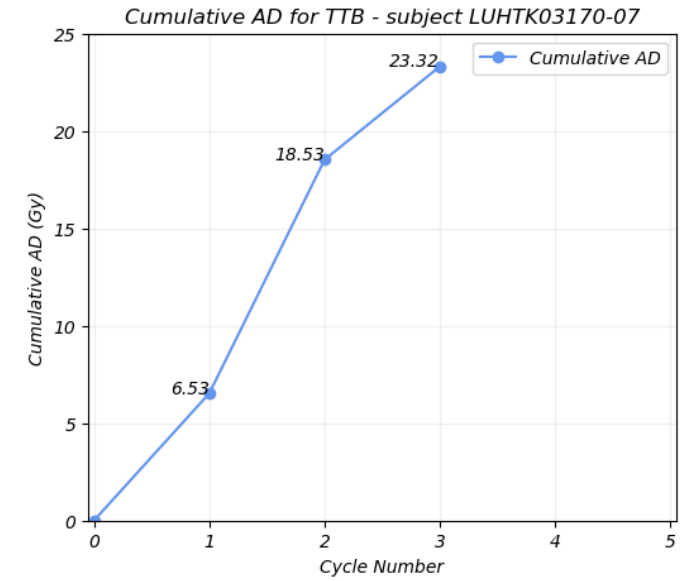
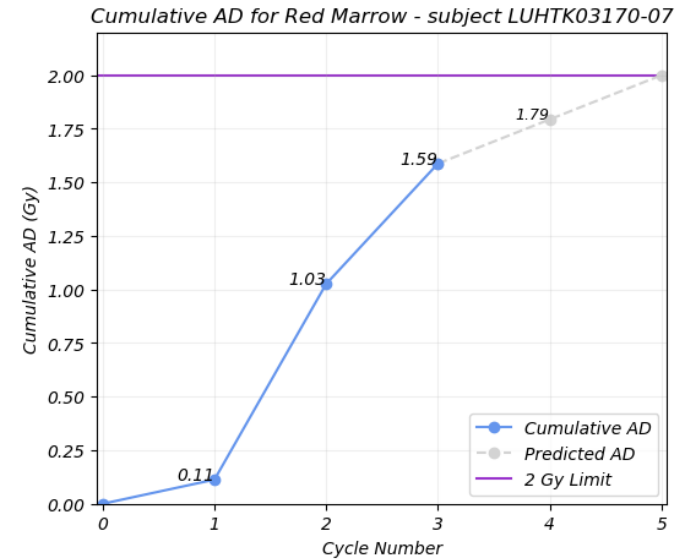
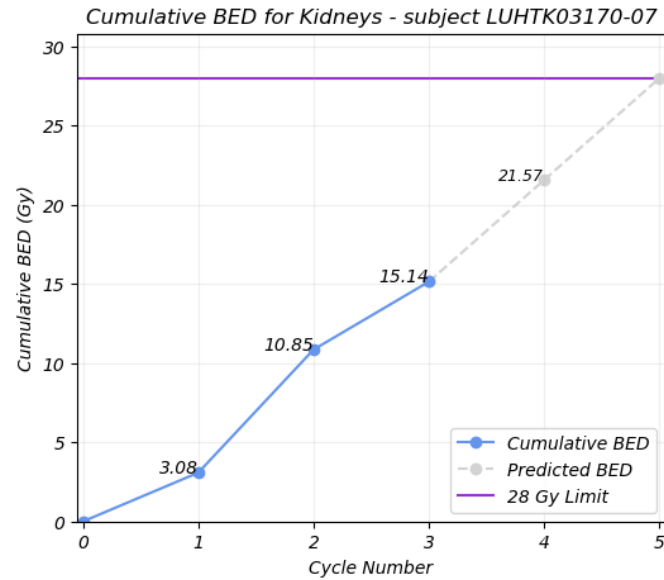
C1: 1.64 GBq, C2: 3.27 GBq , C3: 1.76 GBq)

Pre-treatment

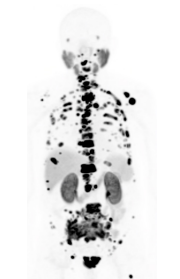
Cycle 1

Cycle 2

Cycle 3



C1: 1.64 GBq, C2: 3.27 GBq , C3: 1.76 GBq)



Pre-treatment



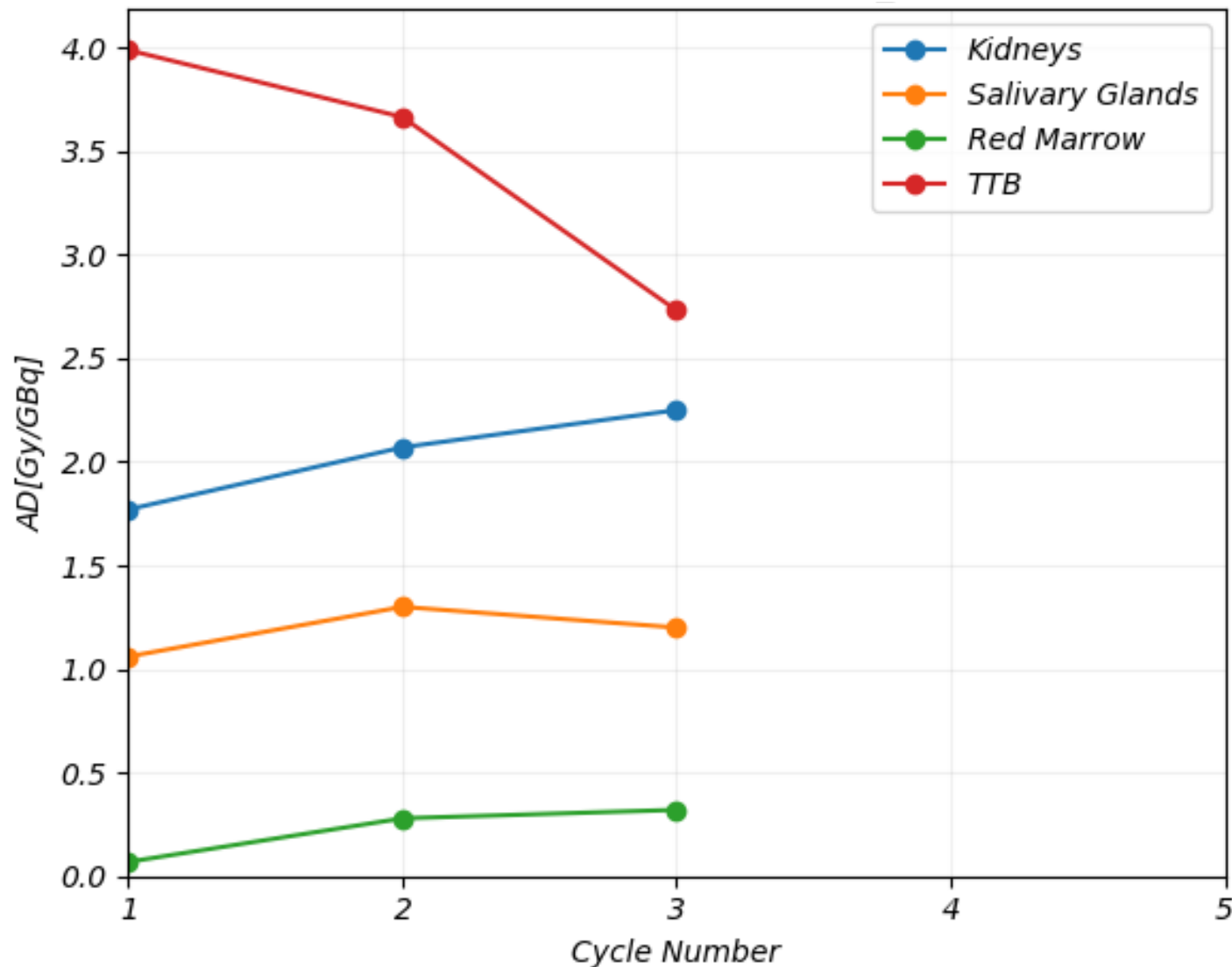
Cycle 1



Cycle 2



Cycle 3



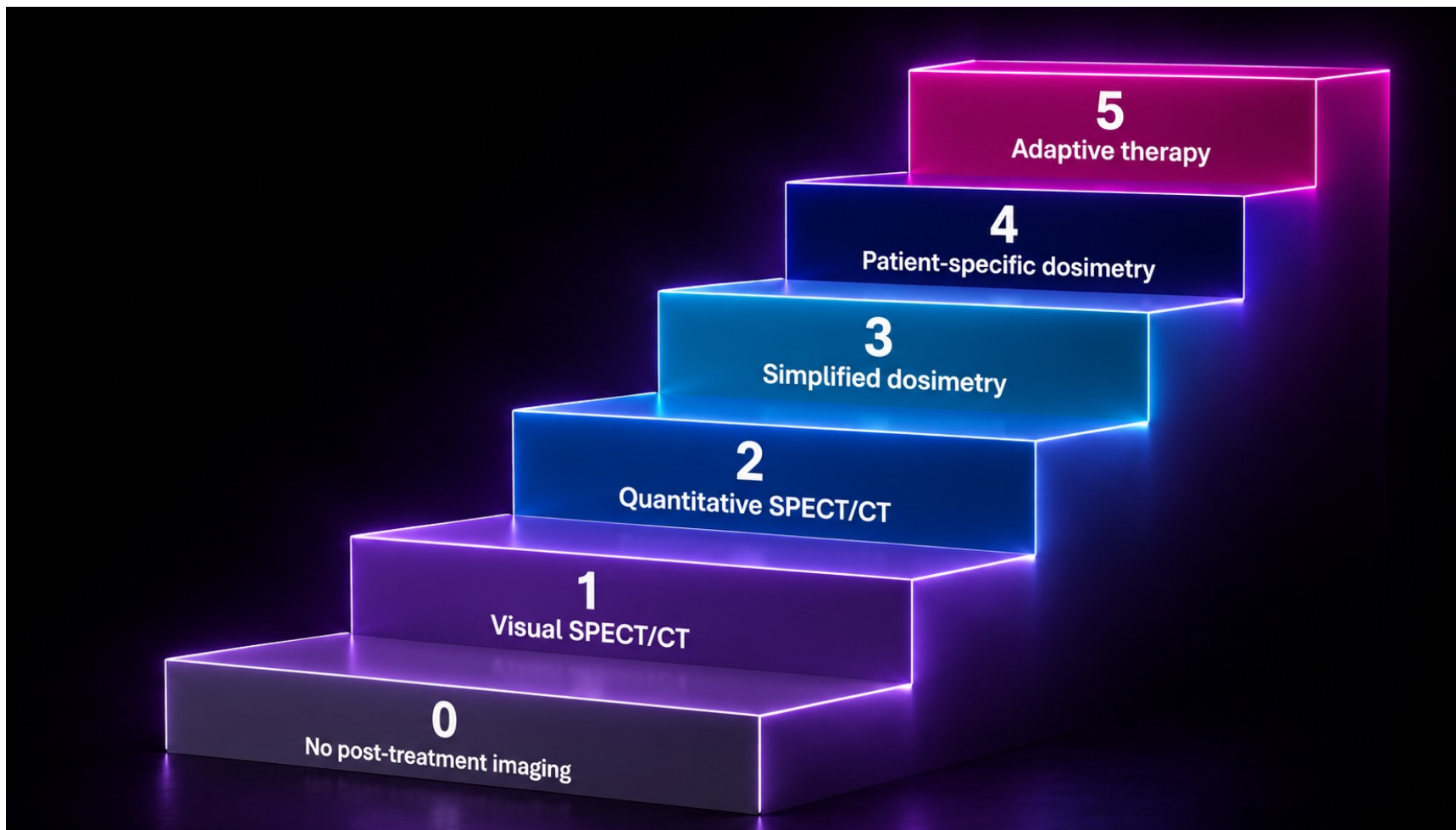
Carlos Uribe, PhD, MCCPM

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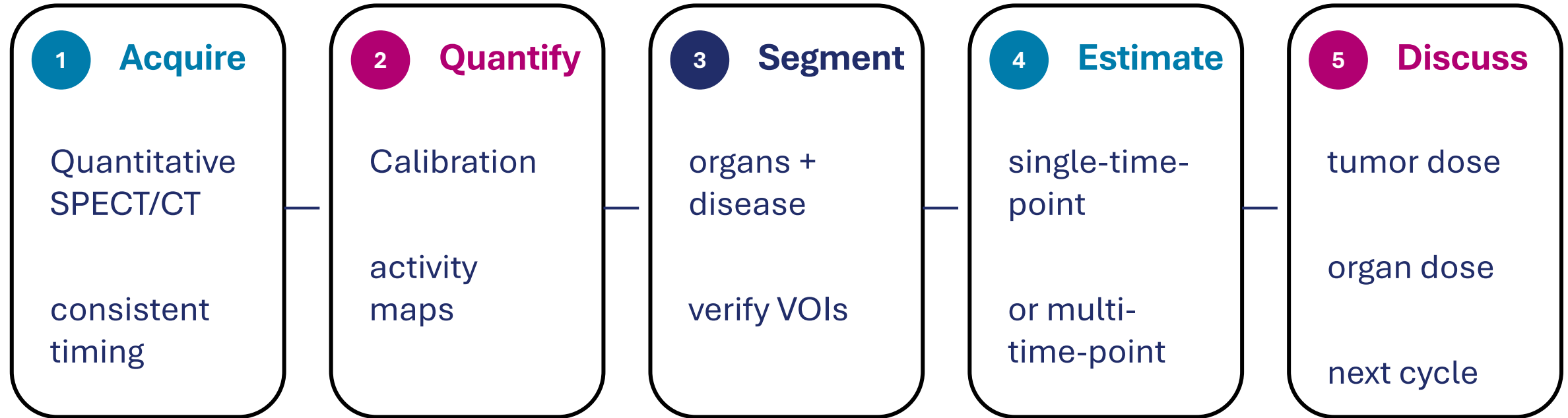
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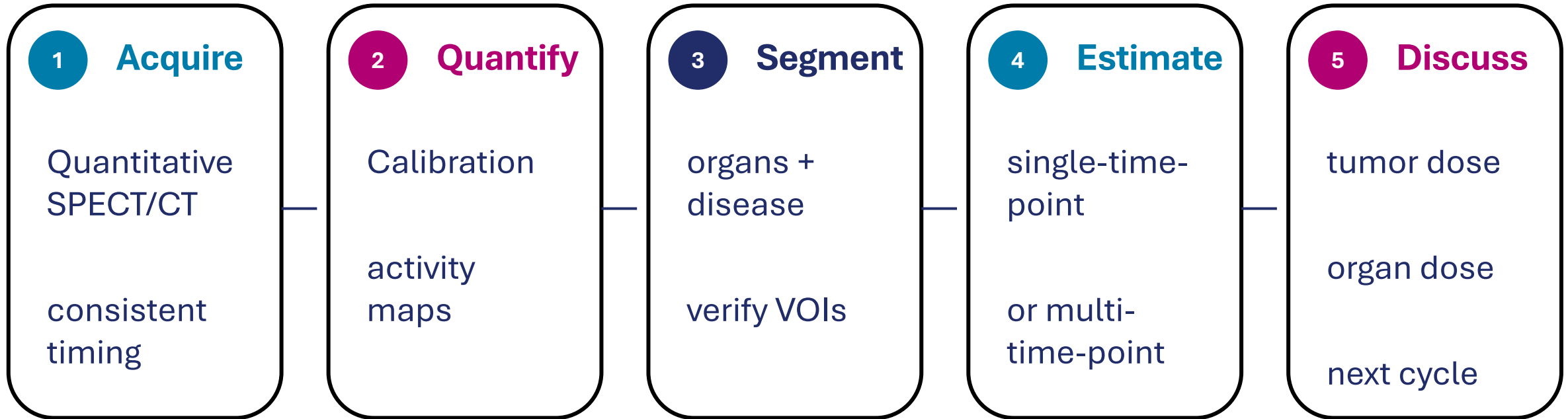
Start where the clinical workflow can support it. Build toward adaptive therapy.



A practical clinical dosimetry workflow can start simple



A practical clinical dosimetry workflow can start simple



The barrier is not that dosimetry is impossible. The barrier is deciding what level is appropriate for the decision.

Where should dosimetry be prioritized first?

- **Clinical trials and absorbed dose optimization studies**
- **Patients with renal, marrow, or salivary-gland risk**
- **Combination therapy with EBRT, chemotherapy, immunotherapy**
- **Retreatment or extended-cycle decisions**
- **Absorbed dose escalation and de-escalation strategies**
- **Alpha-emitter therapies and new agents**

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Dosimetry does not need to be universal tomorrow to be clinically valuable today.

The debate should not be “dosimetry or not dosimetry”

The better question is:

**What dosimetry is needed
for this decision?**

Routine imaging

visual + quantitative trends

Higher-risk patients

organ dose and cumulative dose

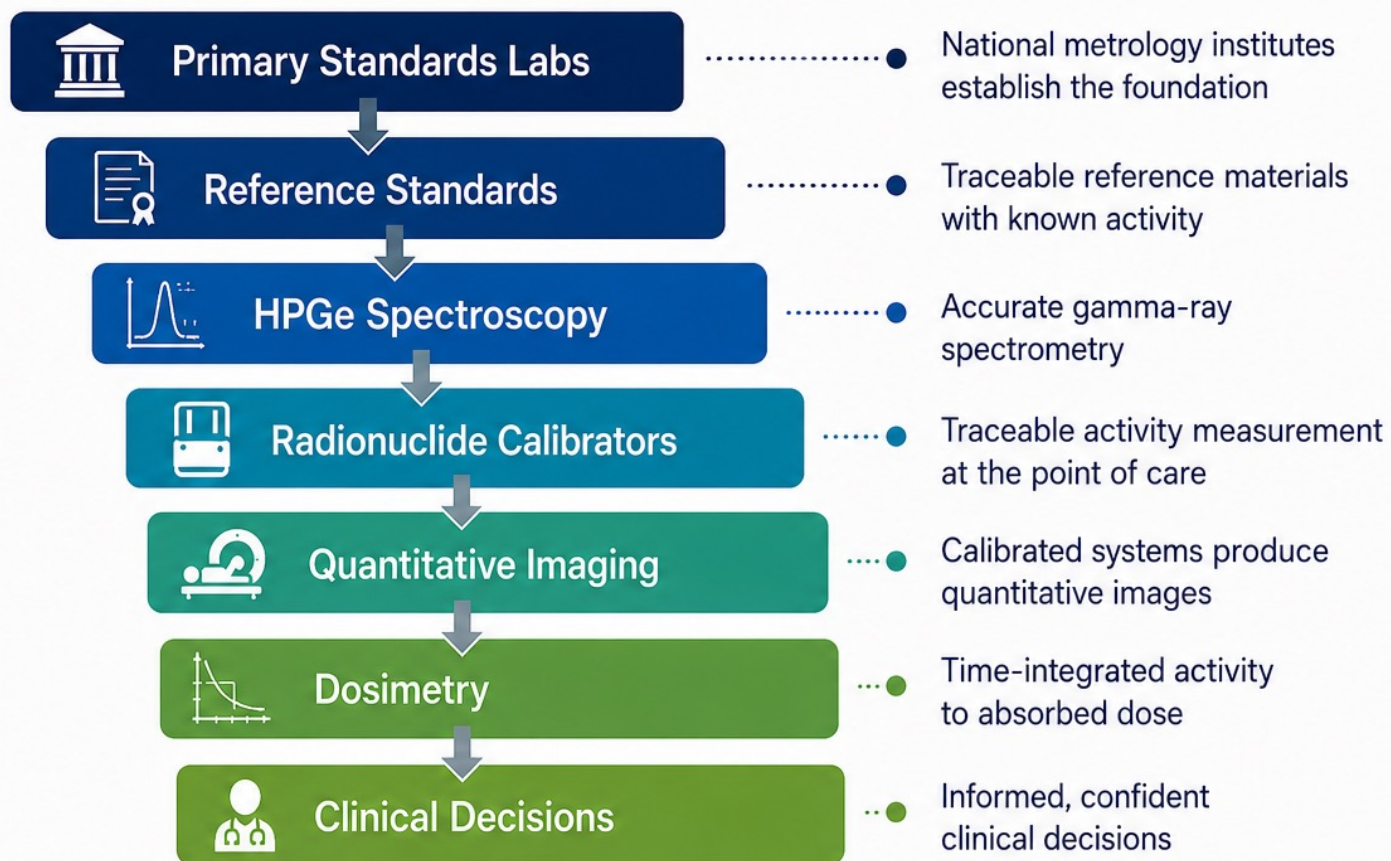
Trials / adaptation

patient-specific absorbed-dose
planning

Take-home messages

- **Post-treatment imaging is how we stop being blind.**
- **Quantitative SPECT is how we stop being purely descriptive.**
- **Dosimetry is how we start adapting treatment.**
- **Dosimetry is the quantitative language that lets our field learn from every treatment**
- **The field does not need perfection to begin learning.**
- **If the mechanism is radiation, and therapy is imageable, absorbed dose should become part of the language of theranostics**

None of this works without trustworthy measurements



Every absorbed dose estimate is only as reliable as the **measurement chain** before it.

Thank you

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