

5th International Workshop on PET in Lymphoma Metabolic Tumour Volume (MTV) & Total Lesion Glycolysis (TLG)

Menton (France), Palais de l'Europe,
September 19-20, 2014



Methodological Aspects

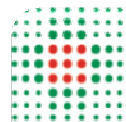
Annibale Versari

Nuclear Medicine

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**SERVIZIO SANITARIO REGIONALE
EMILIA-ROMAGNA**
Azienda Ospedaliera di Reggio Emilia

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Literature data

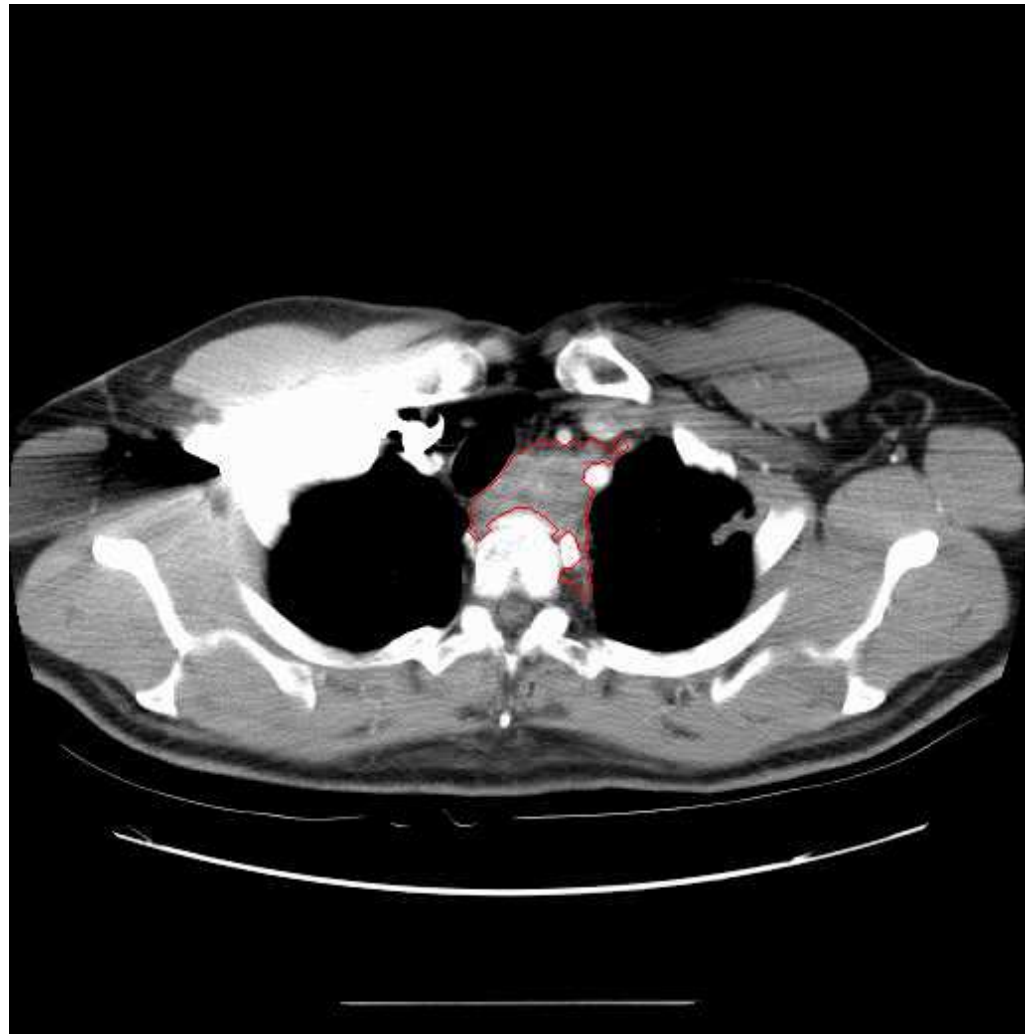
The Tumor Burden is an important prognostic tool in Lymphoma ...

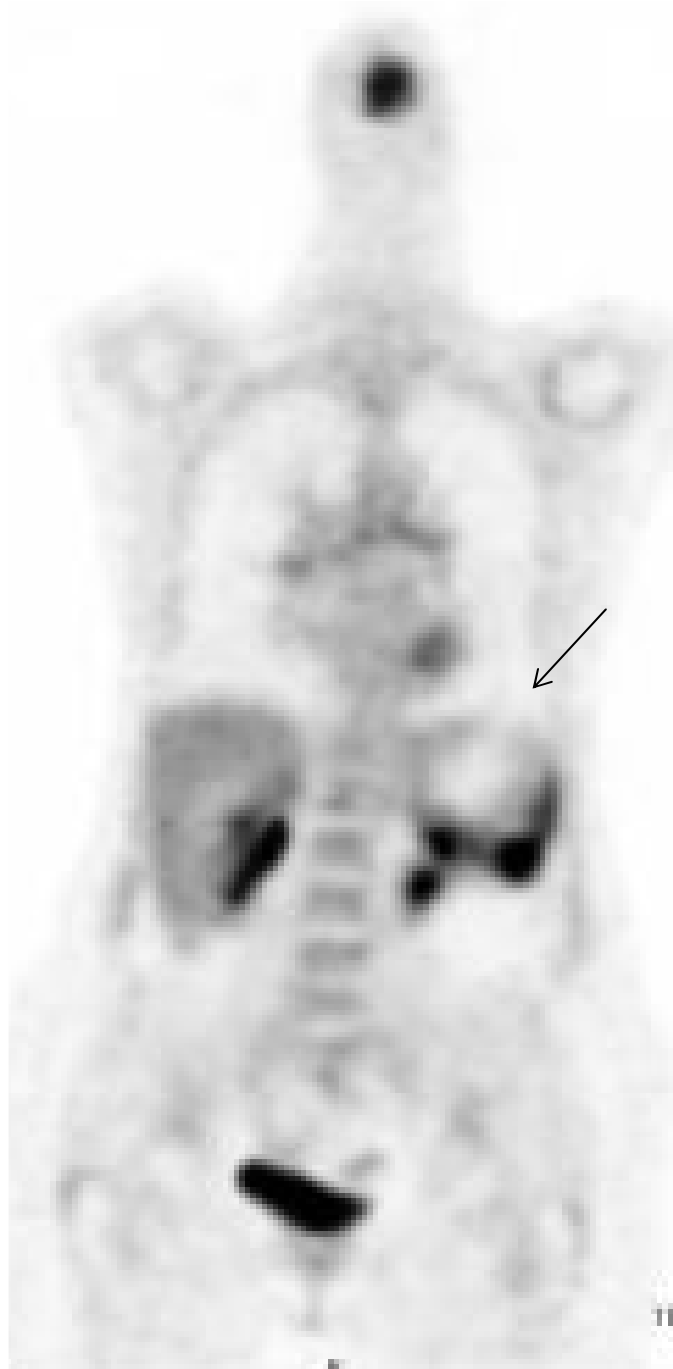
Volume CT

Morphologic
information

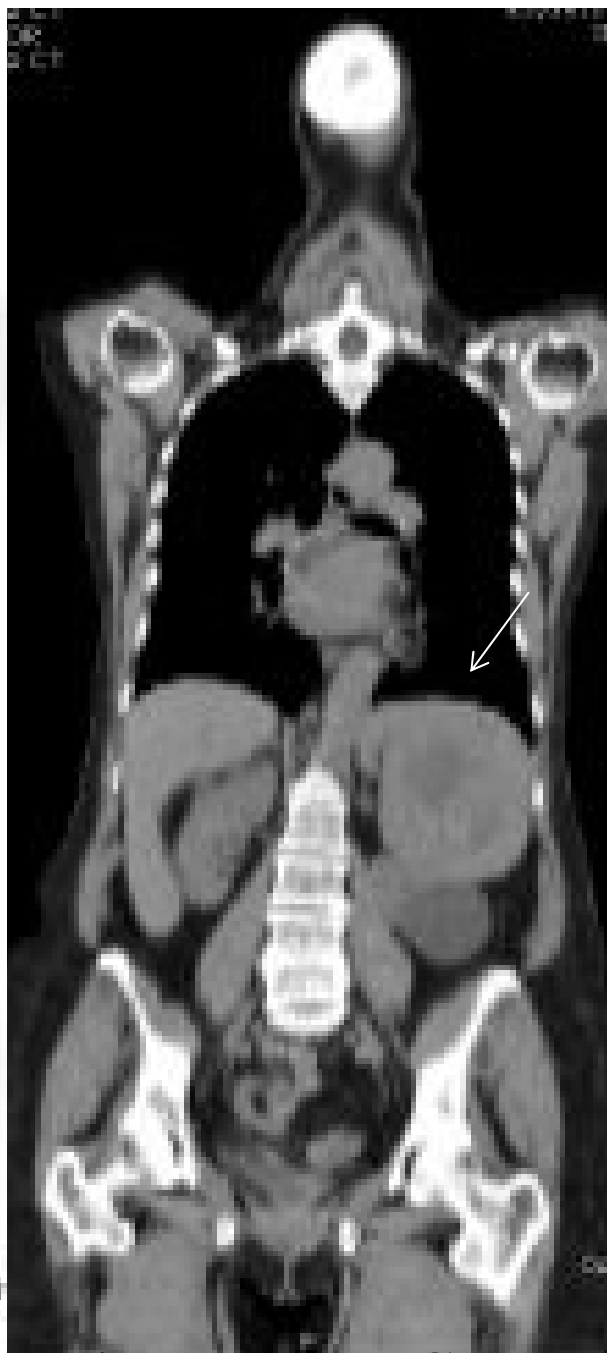
Contour definition

...but the definition on
CT images is complex
and time consuming

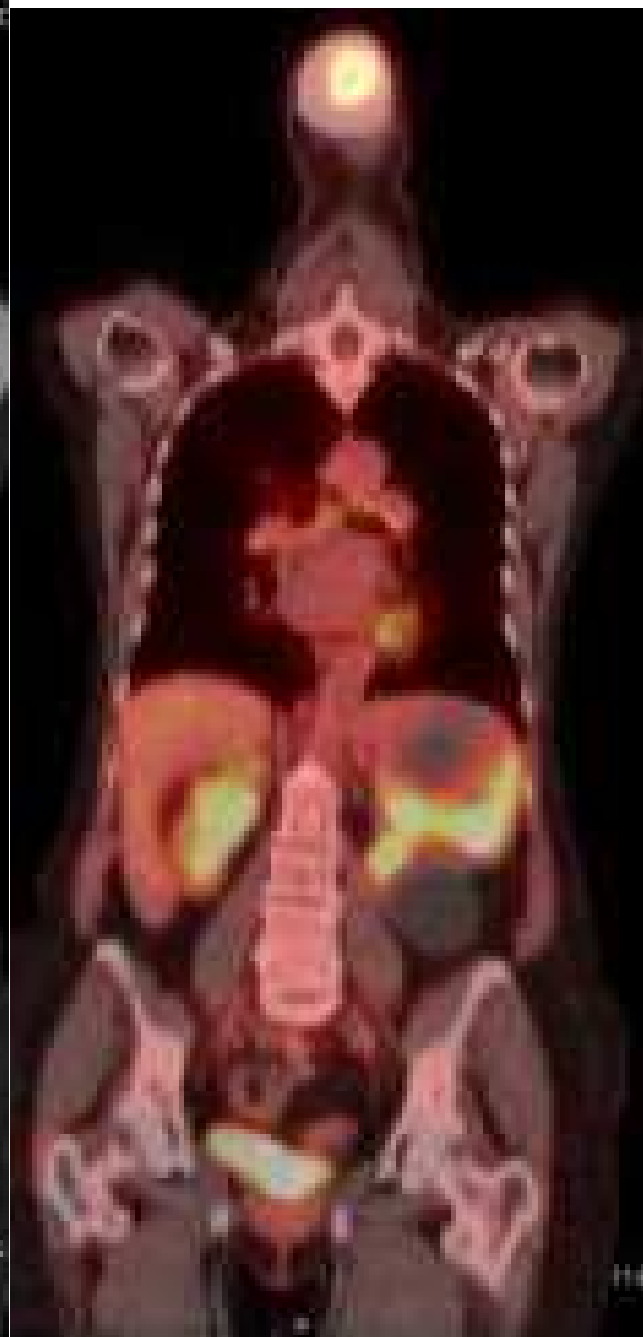




PET



Tumor Volume



CT

FDG PET

➤ Metabolic information (SUV)

but

➤ The metabolic tumor volume definition needs some rules

BTV definition: which method?

Several strategies in using PET for target volume definition in radiotherapy treatment planning are being investigated:

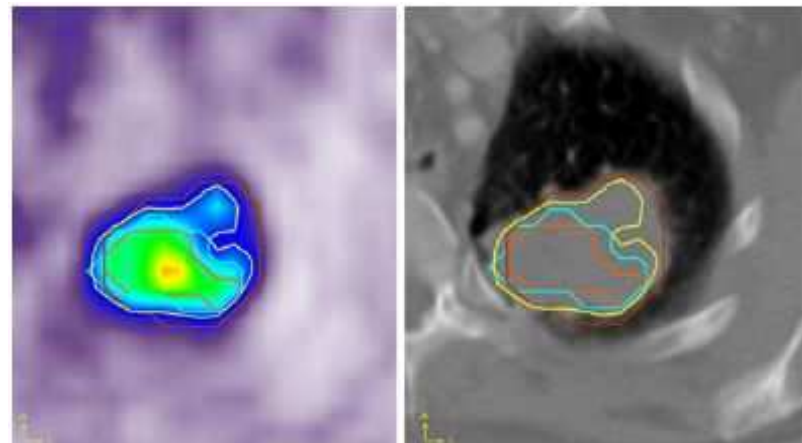
- **visual contouring**
- **Fixed or relative threshold** (SUV=2.5, 30-40-50 % of peak activity...)
- **adaptive threshold** (dependence on Signal/Bkg ratio and lesion size, dependence on reconstruction algorithm...)
- **gradient based** (adaptive region growing)
- **statistical techniques** (fuzzy locally adaptive bayesian, Markov models, k-means clustering...)

Reproducibility

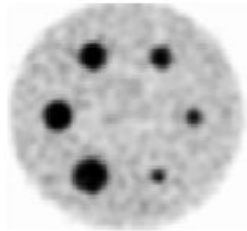
Robustness

Accuracy

Automation



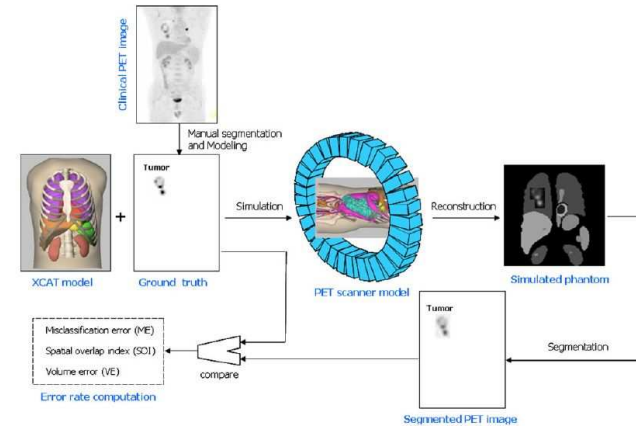
Validation = “fidelity to the truth”



Spherical and
homogeneous objects

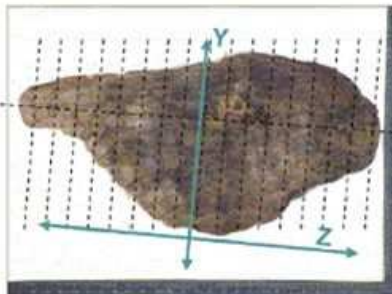


Anthropomorphic
phantoms



Simulated data

Ground truth



Histopathological
data

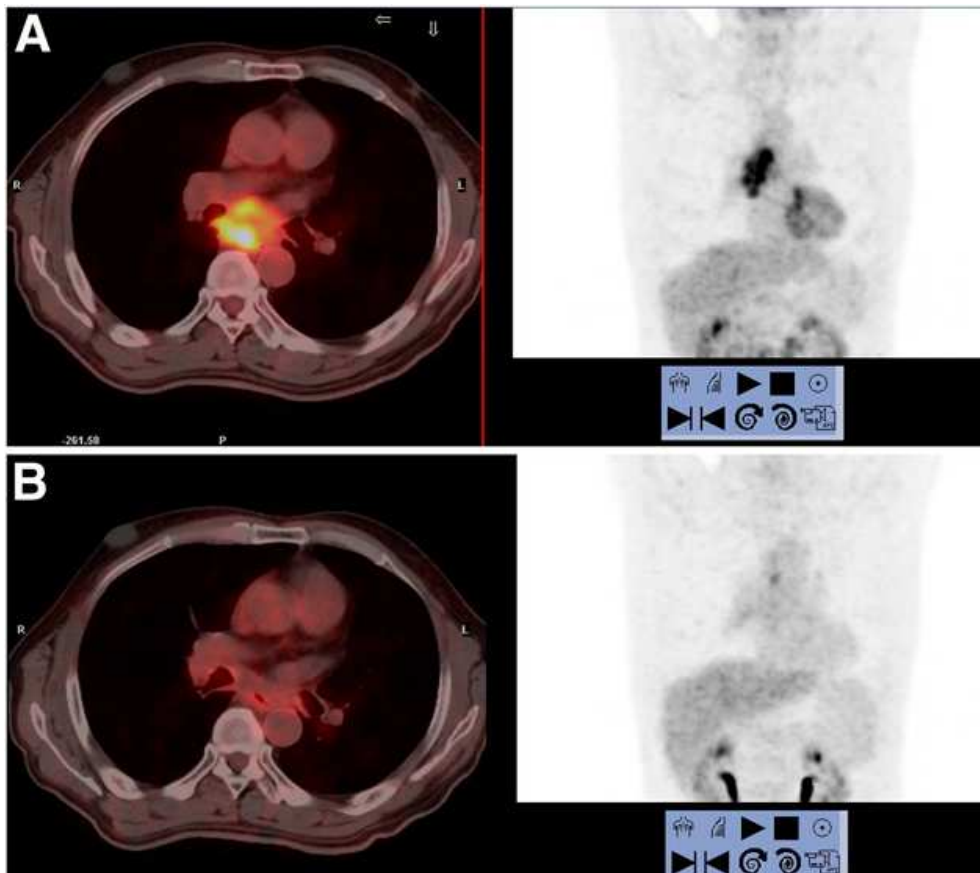
BUT the histopathological data evaluation is particularly complex, given that:

- PET study must be performed before surgical resection of the tumor
- the effects of sample shrinkage in the different conditions (in vivo and and in vitro) must be considered.

The volumetric analysis of the the surgical specimen must be done through an accurately and reliably procedure.

Value of Metabolic Tumor Volume on Repeated ^{18}F -FDG PET/CT for Early Prediction of Survival in Locally Advanced Non-Small Cell Lung Cancer Treated with Concurrent Chemoradiotherapy

Wei Huang^{*1}, Min Fan^{*1}, Bo Liu², Zheng Fu³, Tao Zhou¹, Zicheng Zhang¹, Heyi Gong¹, and Baosheng Li¹



... and many other tumors

- Esophagus
- Head&neck
- Ovarii
-

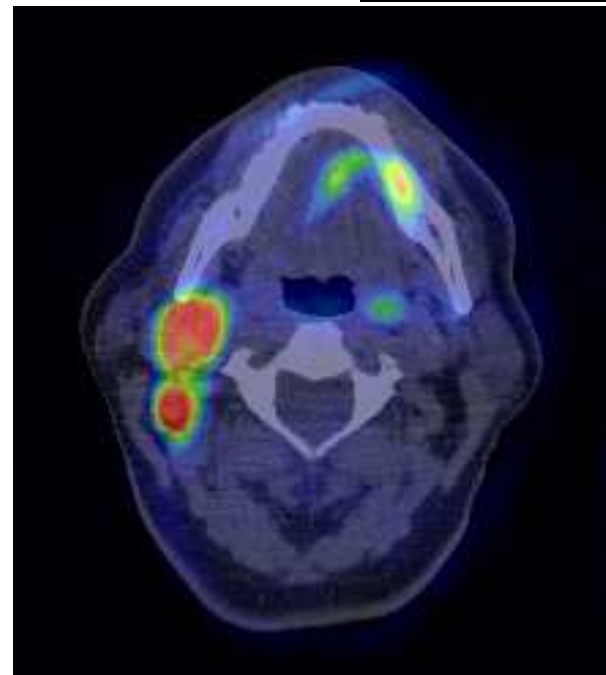
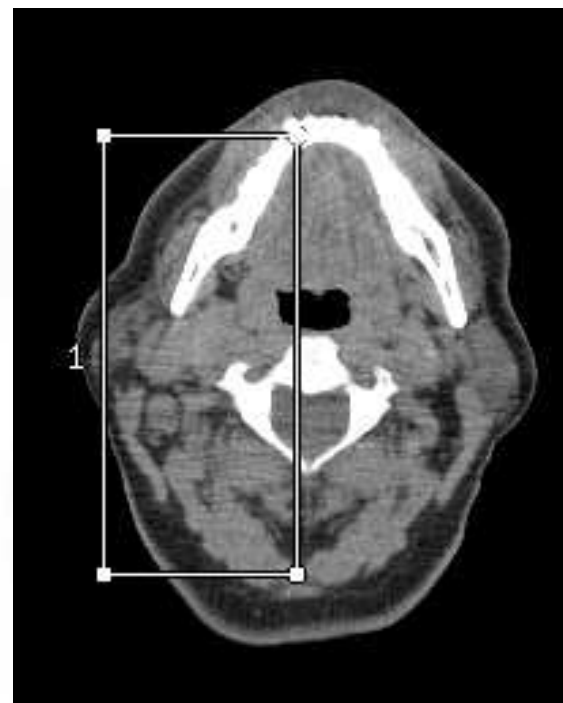
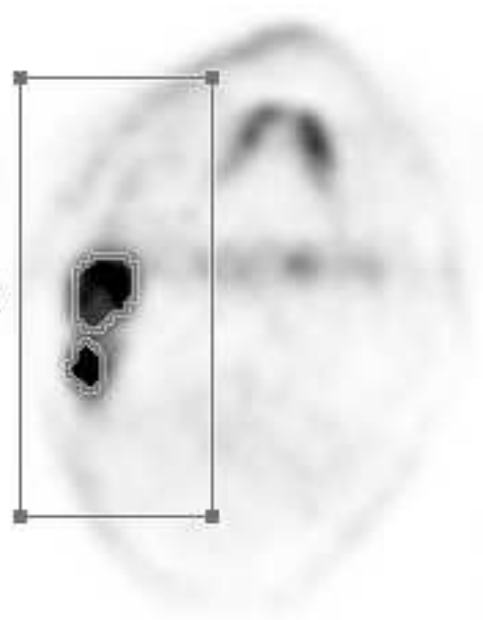
.... but Lymphoma has particular characteristics

- Heterogeneous disease

Many different

- sites
- volumes
- uptakes (SUV)
- backgrounds

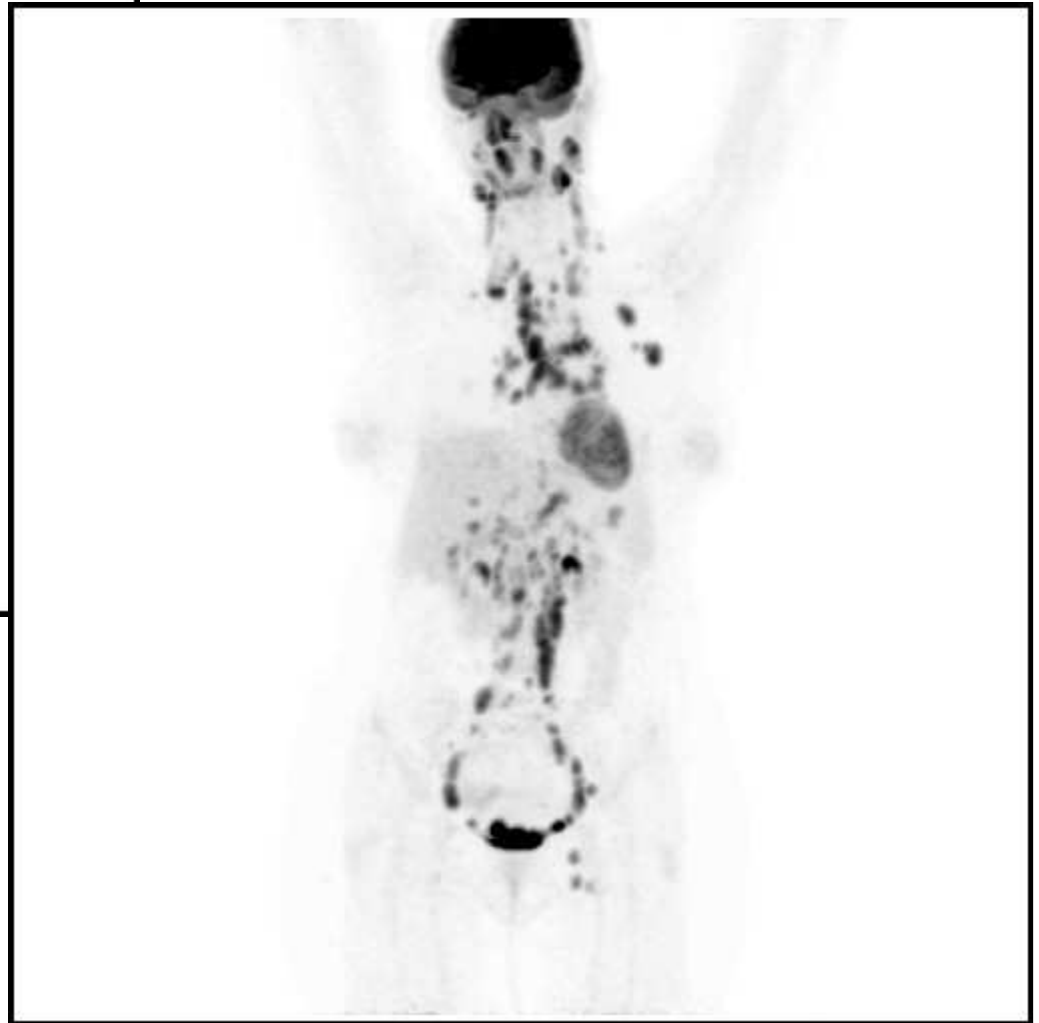
- Bulky – non-Bulky





**Hodgkin
Lymphoma**

**Non-Hodgkin
Lymphoma**



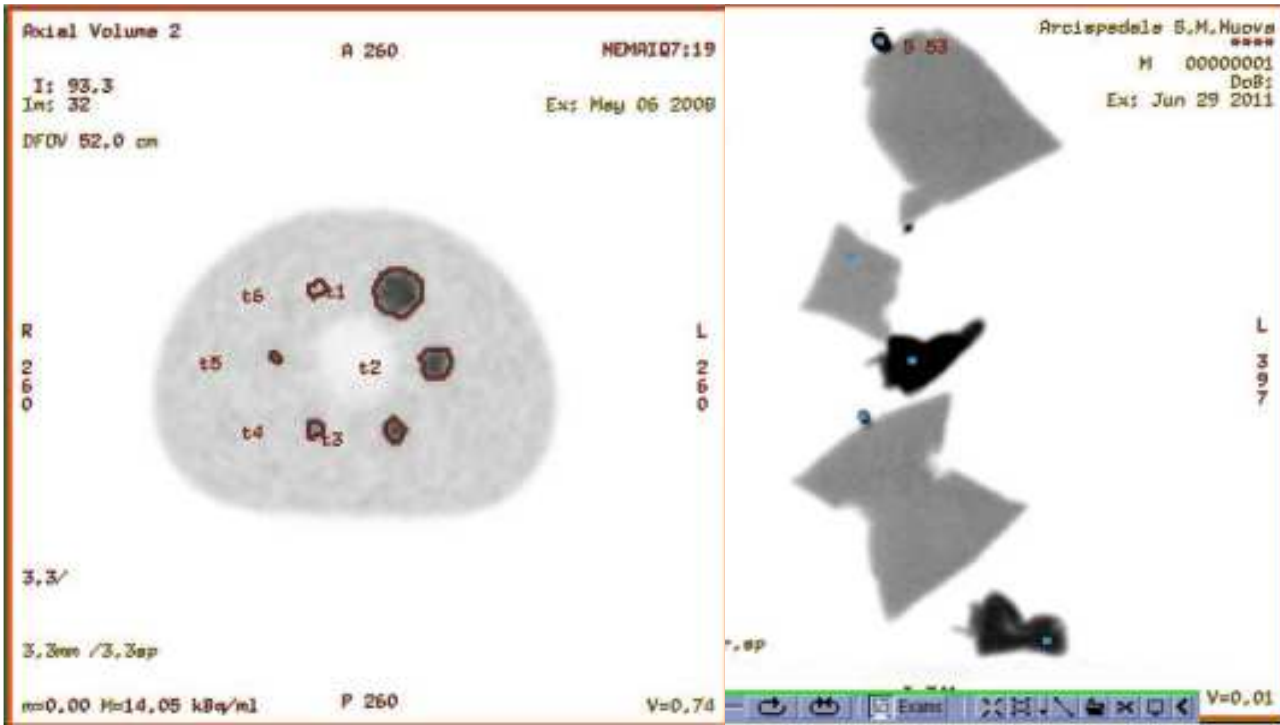
Metabolic tumour volumes measured at staging in lymphoma: methodological evaluation on phantom experiments and patients

Michel Meignan • Myriam Sasanelli • René Olivier Casasnovas •
Stefano Luminari • Federica Fioroni • Chiara Coriani • Helene Masset •
Emmanuel Itti • Paolo G. Gobbi • Francesco Merli • Annibale Versari

Comparison of 2 semiautomatic segmentation softwares in 2 centers

•**PET VCAR** - Volume Computer Assisted Reading – GE Healthcare
Nuclear Medicine Dept – Reggio Emilia (Italy)

•**KEOSYS software**
Nuclear Medicine Dept – Créteil (France)

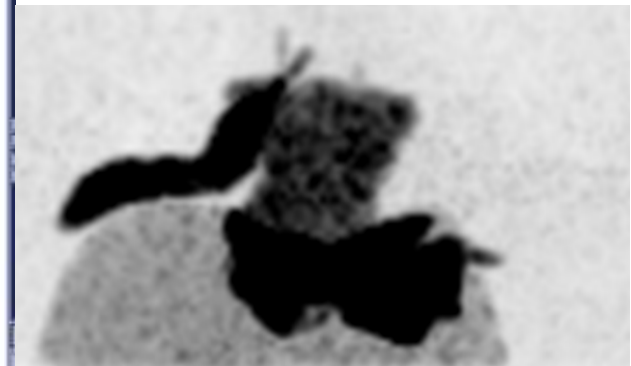
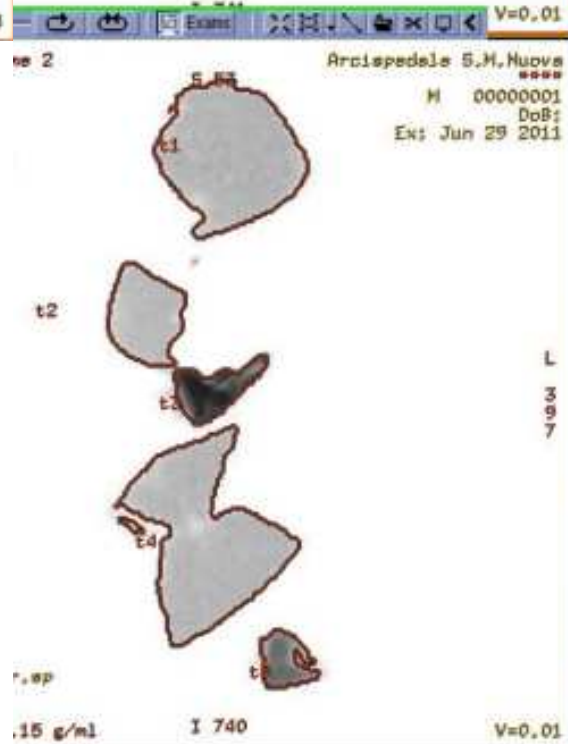


Nema phantom



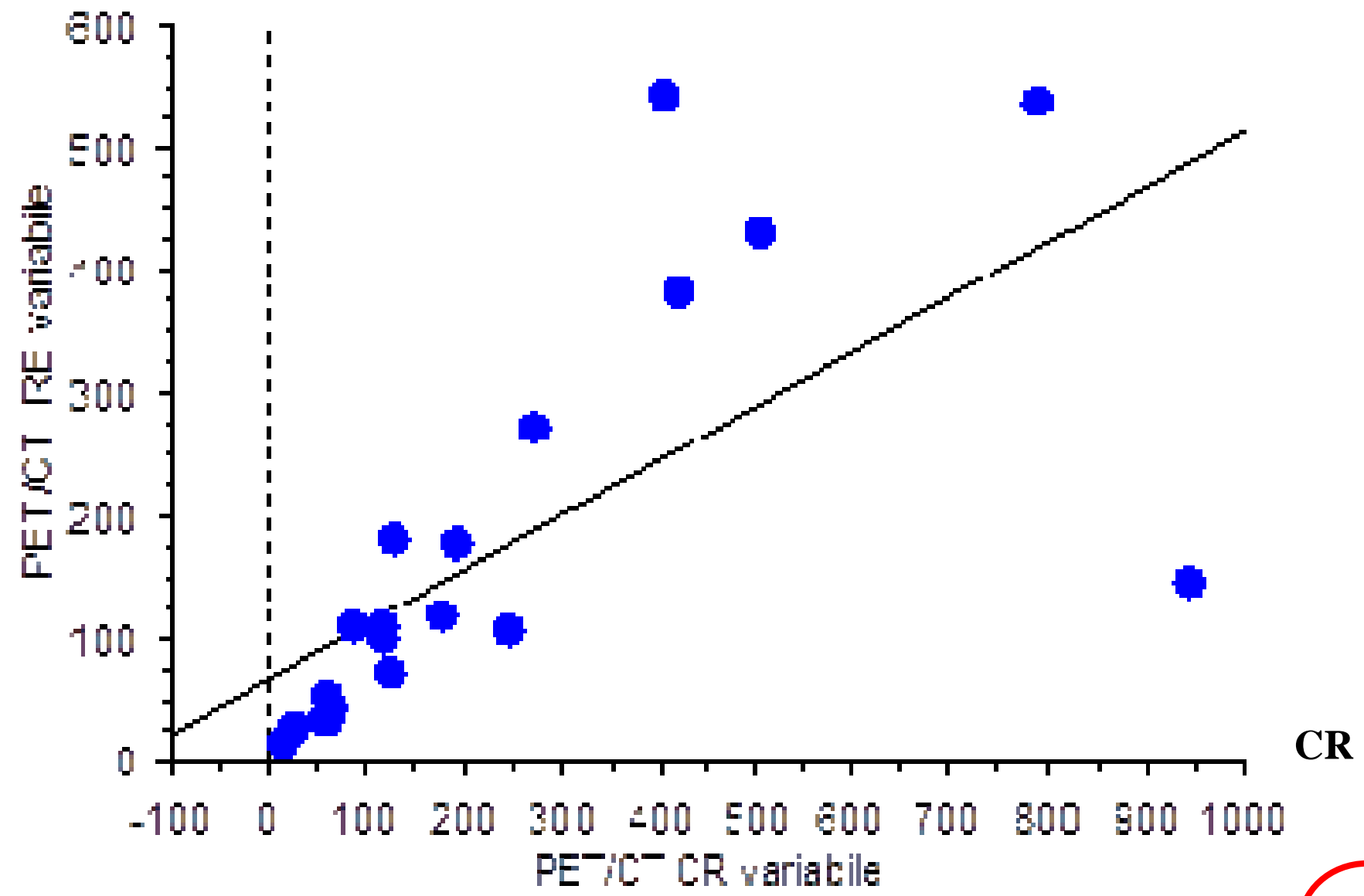
Anthropomorphic Phantom

Irregular phantoms

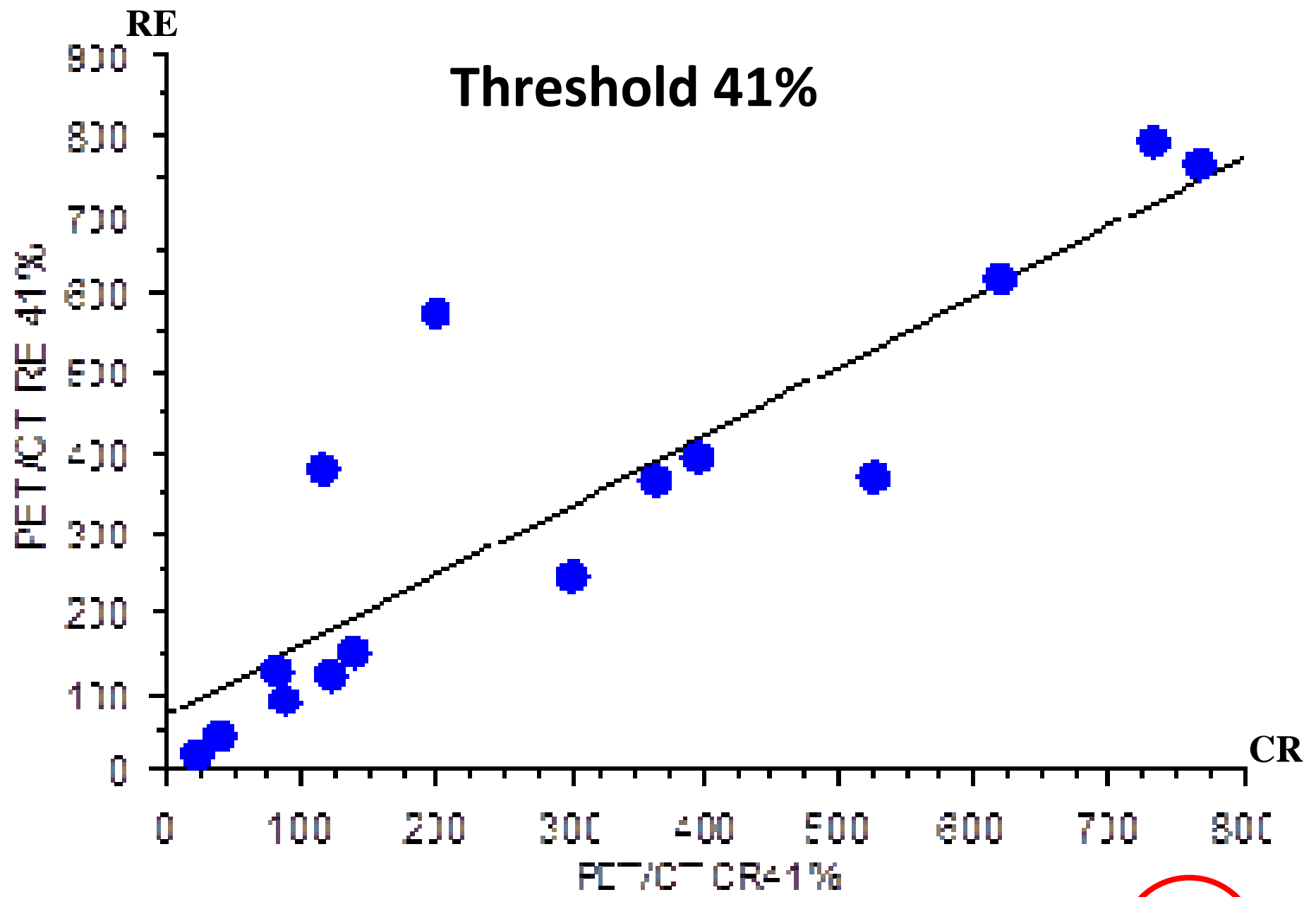


RE

Variable Threshold



PET/CT RE variable = 67.11 + .447 * PET/CT CR variable: R² = .46



PET/CT RE 41% = 71.461 - .88 * PET/CT CR 41%; R² = .77

FDG PET and PET/CT: EANM procedure guidelines for tumour PET imaging: version 1.0

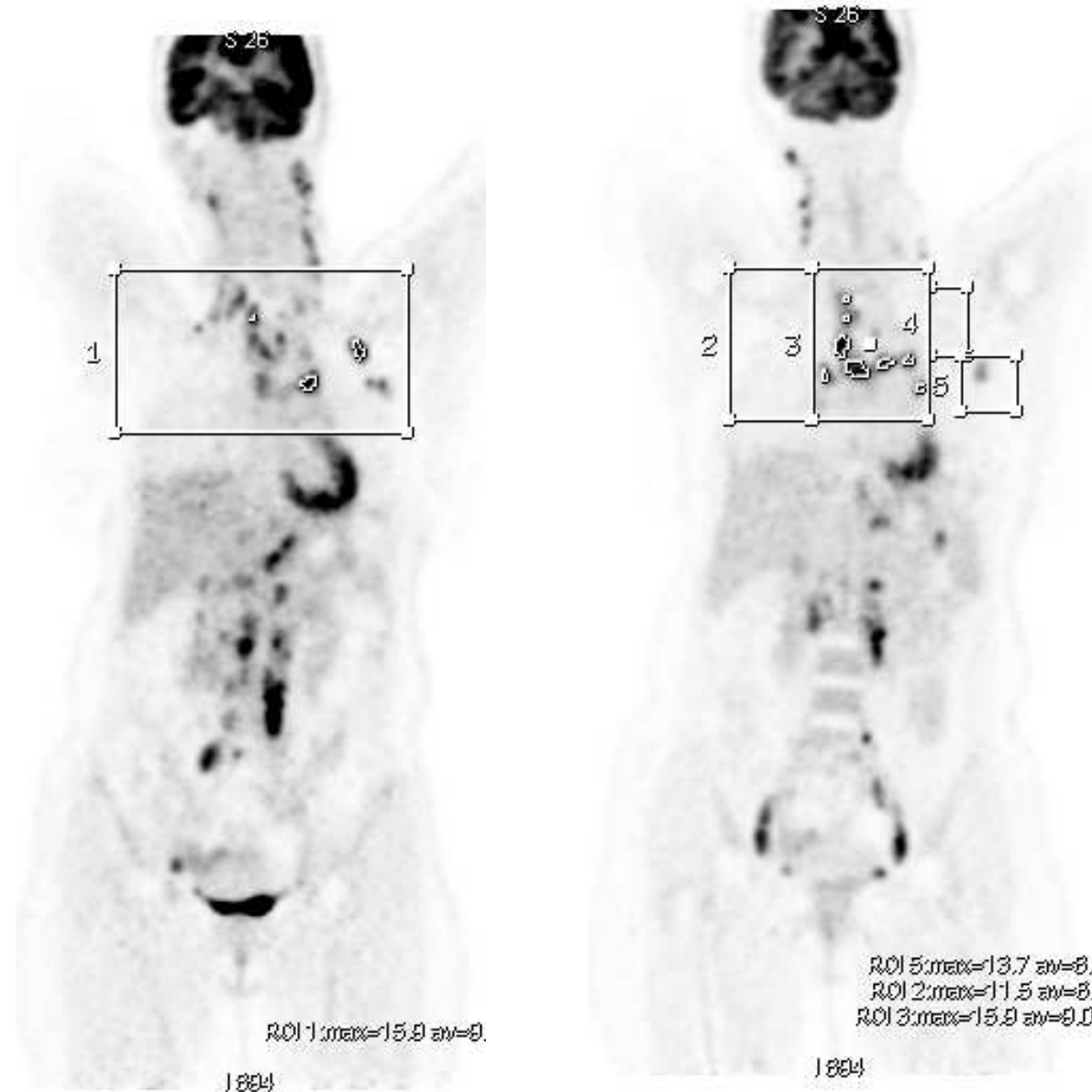
Ronald Boellaard • Mike J. O'Doherty • Wolfgang A. Weber • Felix M. Mottaghy • Markus N. Lonsdale • Sigrid G. Stroobants • Wim J. G. Oyen • Joerg Kotzerke • Otto S. Hoekstra • Jan Pruim • Paul K. Marsden • Klaus Tatsch • Corneline J. Hoekstra • Eric P. Visser • Bertjan Arends • Fred J. Verzijlbergen • Josee M. Zijlstra • Emile F. I. Comans • Adriaan A. Lammertsma • Anne M. Paans • Antoon T. Willemsen • Thomas Beyer • Andreas Bockisch • Cornelia Schaefer-Prokop • Dominique Delbeke • Richard P. Baum • Arturo Chiti • Bernd J. Krause

The isocontour described as A41 (41% of max pixel value)

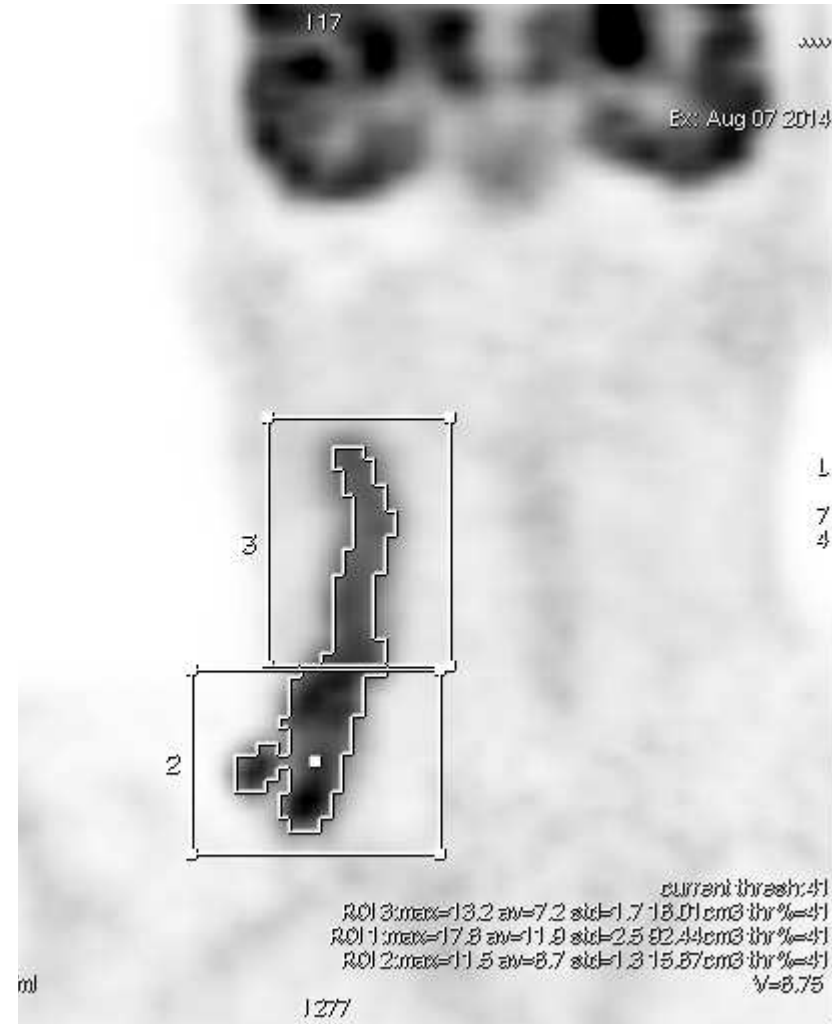
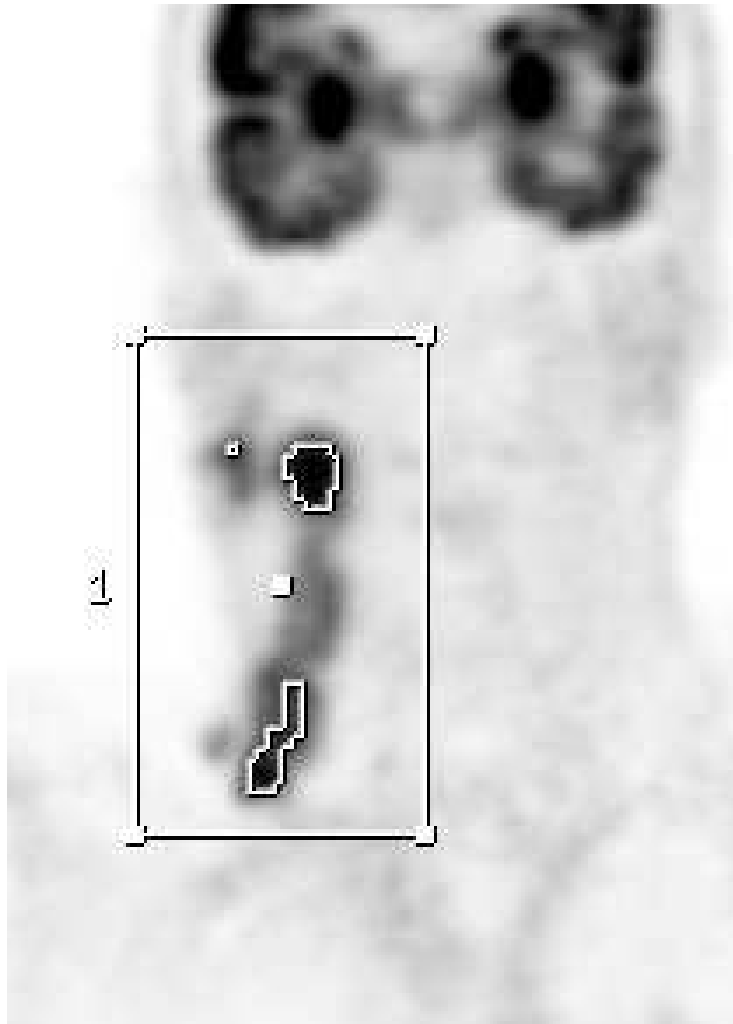
- **generally corresponds best with the actual dimensions of the tumour,**
- but only**
- **for higher tumour-to-background values and homogenous backgrounds**

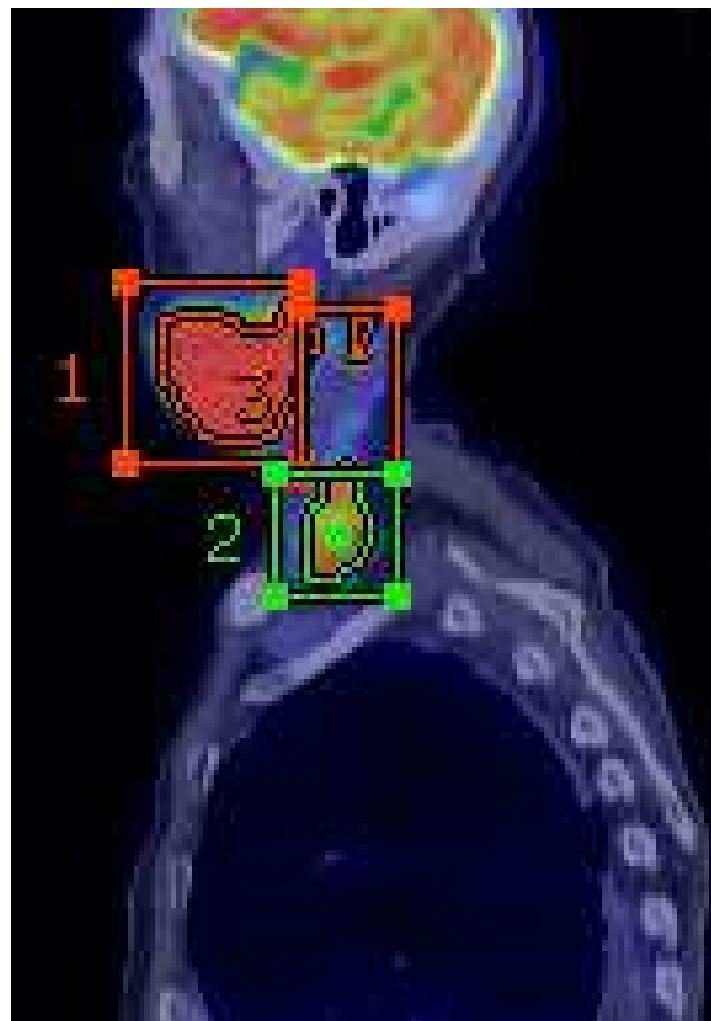
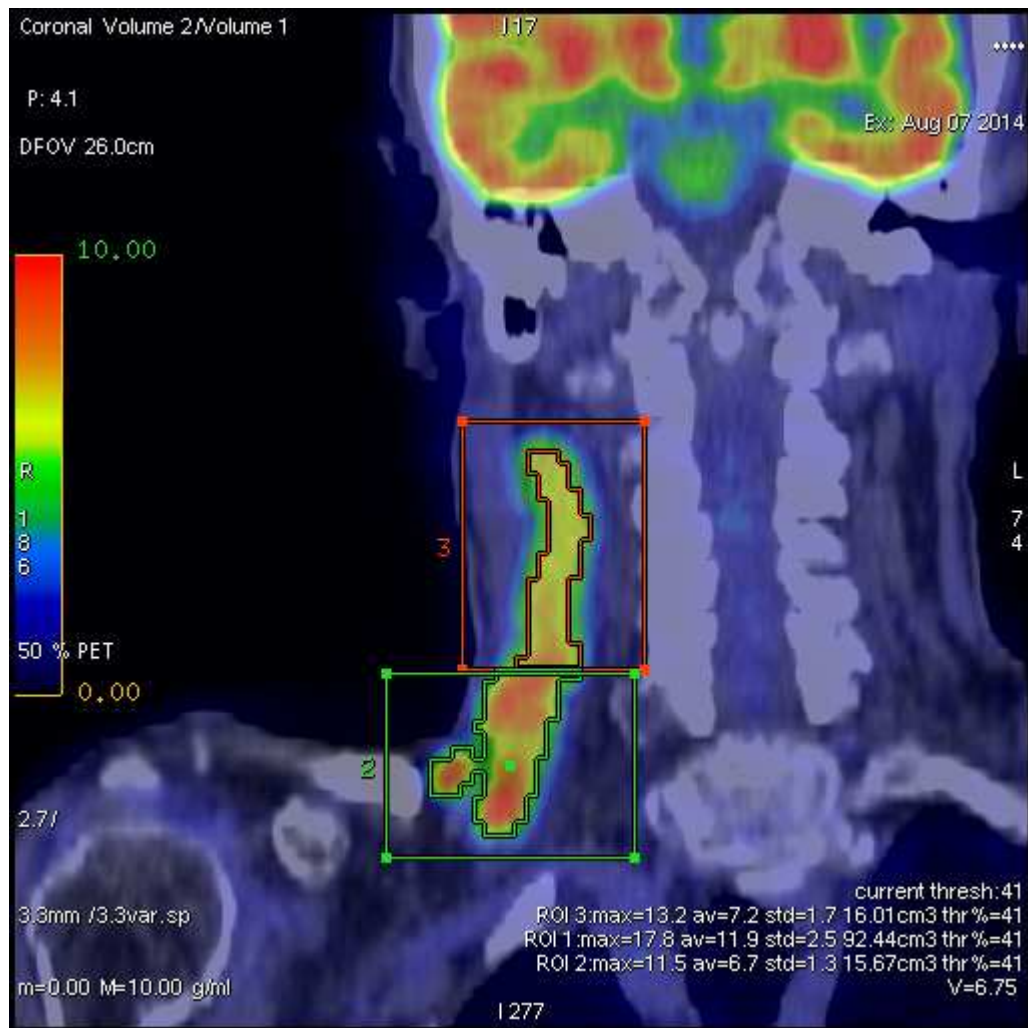
To determine TMTV0, volumes of interest (VOI) are identified by drawing spheres, cylinders or cubes around each focus of 18F-FDG uptake on pretreatment 18F-FDG PET/CT scans.

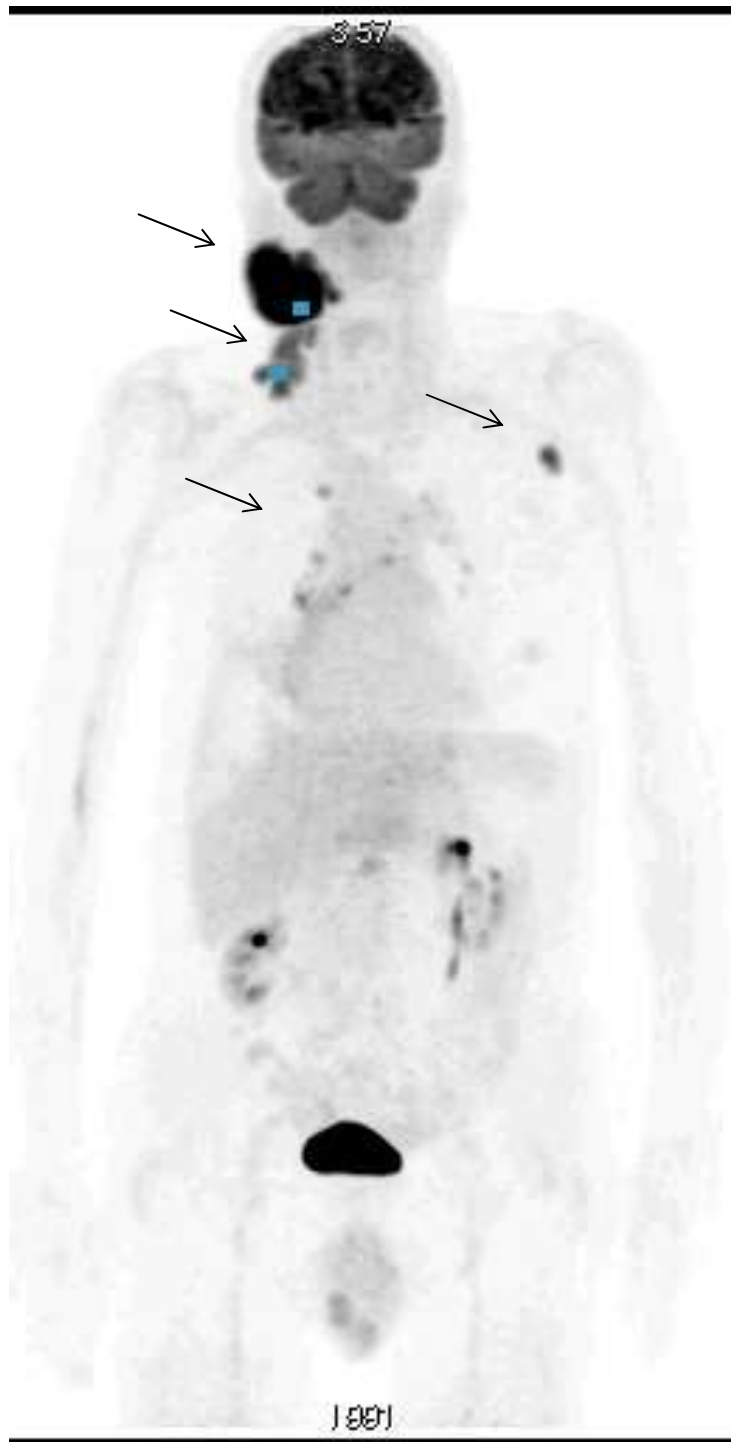
In each VOI, voxels with **41 %** or more of the SUVmax of the lesion are incorporated to define the volume of each individual lesion



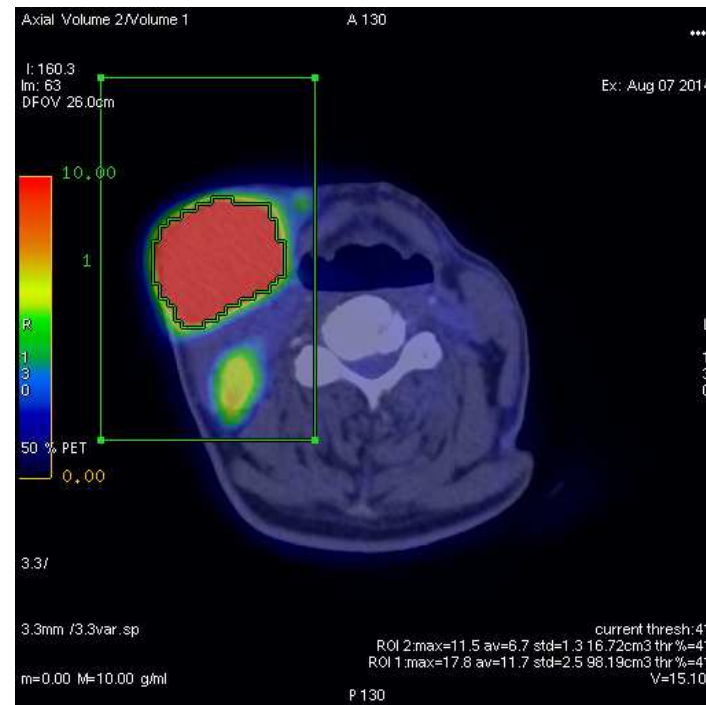
If the **uptake is heterogeneous**, high local uptake needs to be isolated in separate VOI to avoid underestimation of the tumour volume.

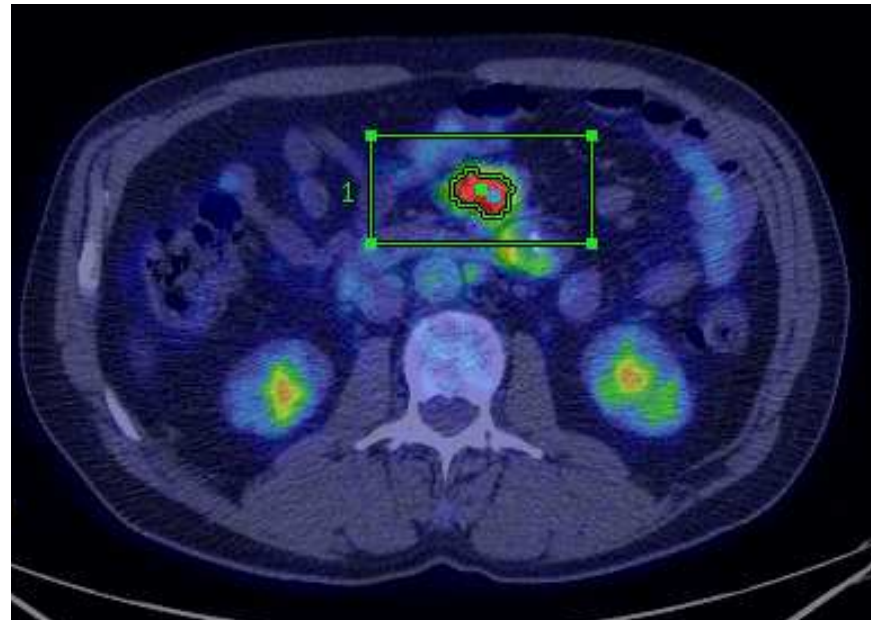
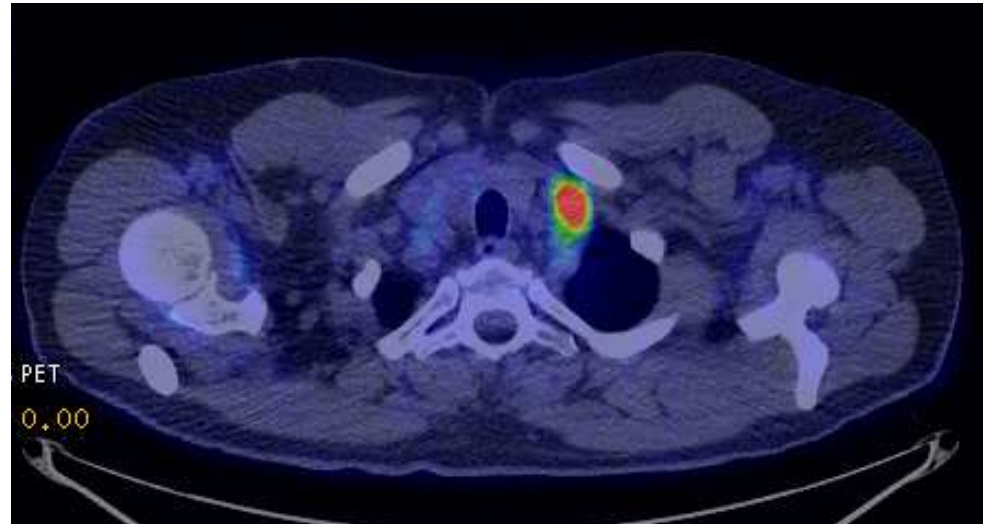
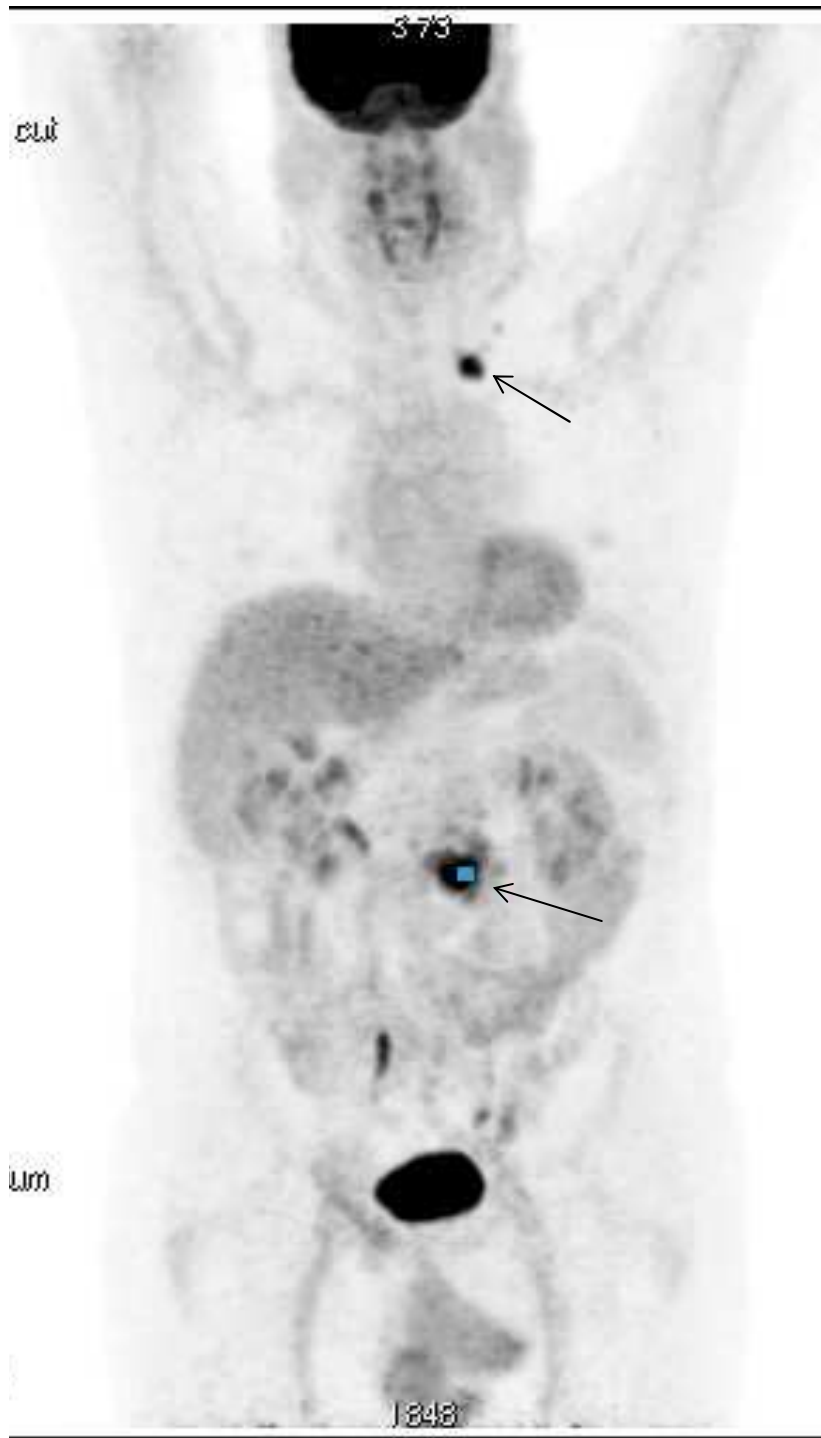






Different uptake





Different background

Problems

- **Extranodal involvement**
- **Bone marrow**
- **Spleen involvement**

Baseline metabolic tumour volume is an independent prognostic factor in Hodgkin lymphoma

Salim Kanoun • Cédric Rossi • Alina Berriolo-Riedinger • Inna Dygai-Cochet • Alexandre Cochet • Olivier Humbert • Michel Toubreau • Emmanuelle Ferrant • François Brunotte • René-Olivier Casasnovas

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Problems

Extranodal involvement

was considered in the volume calculation according to the following rules:

- the **liver, lung and bone marrow** were considered involved only if there was
- focal uptake**, and the volume of each individual hypermetabolic lesion was
- computed in a **separate VOI**

Eur J Nucl Med Mol Imaging (2014) 41:1735–1743
DOI 10.1007/s00259-014-2783-x

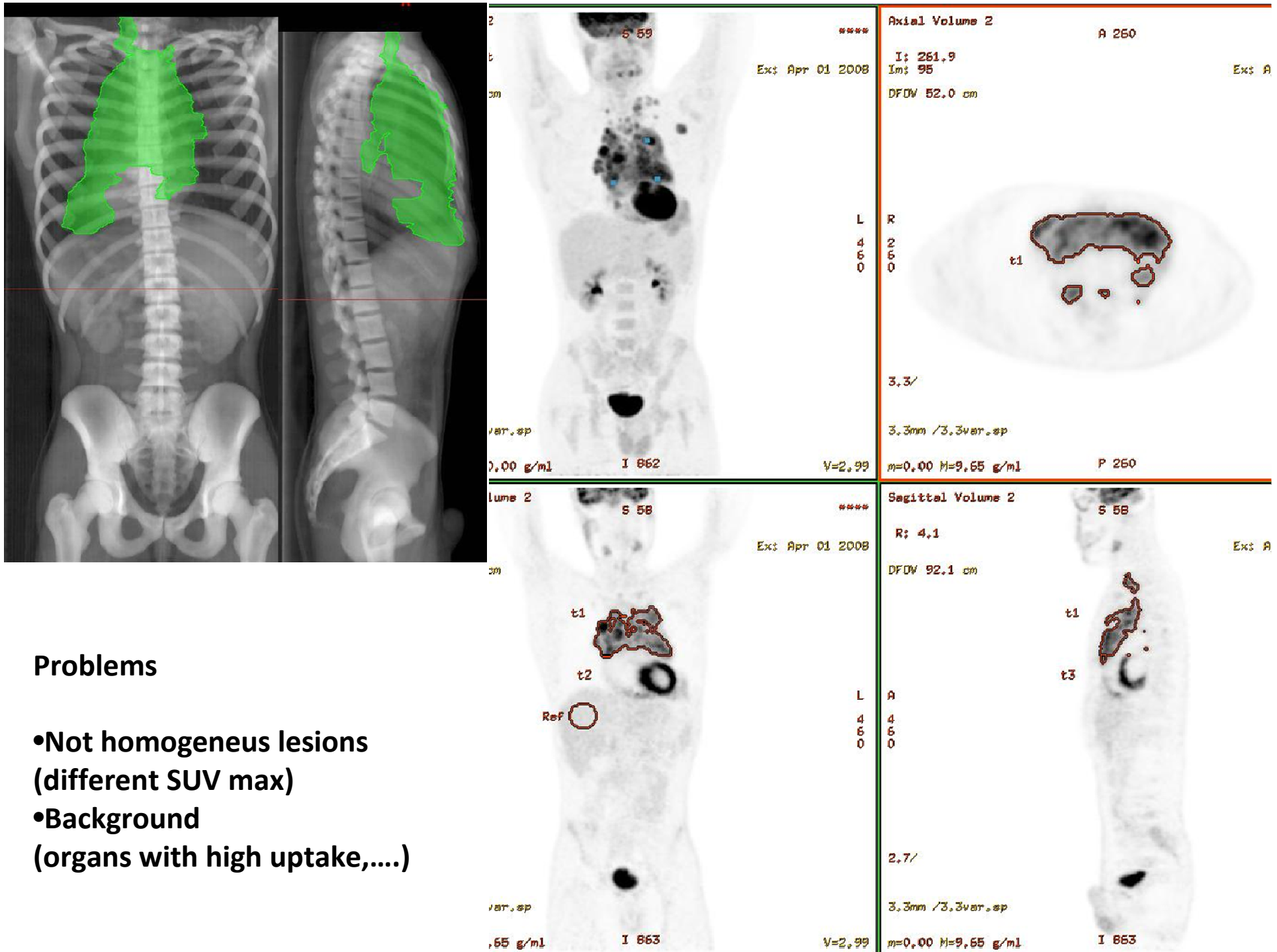
ORIGINAL ARTICLE

Baseline metabolic tumour volume is an independent prognostic factor in Hodgkin lymphoma

Salim Kanoun • Cédric Rossi • Alina Berriolo-Riedinger • Inna Dygai-Cochet • Alexandre Cochet • Olivier Humbert • Michel Toubeau • Emmanuelle Ferrant • François Brunotte • René-Olivier Casasnovas

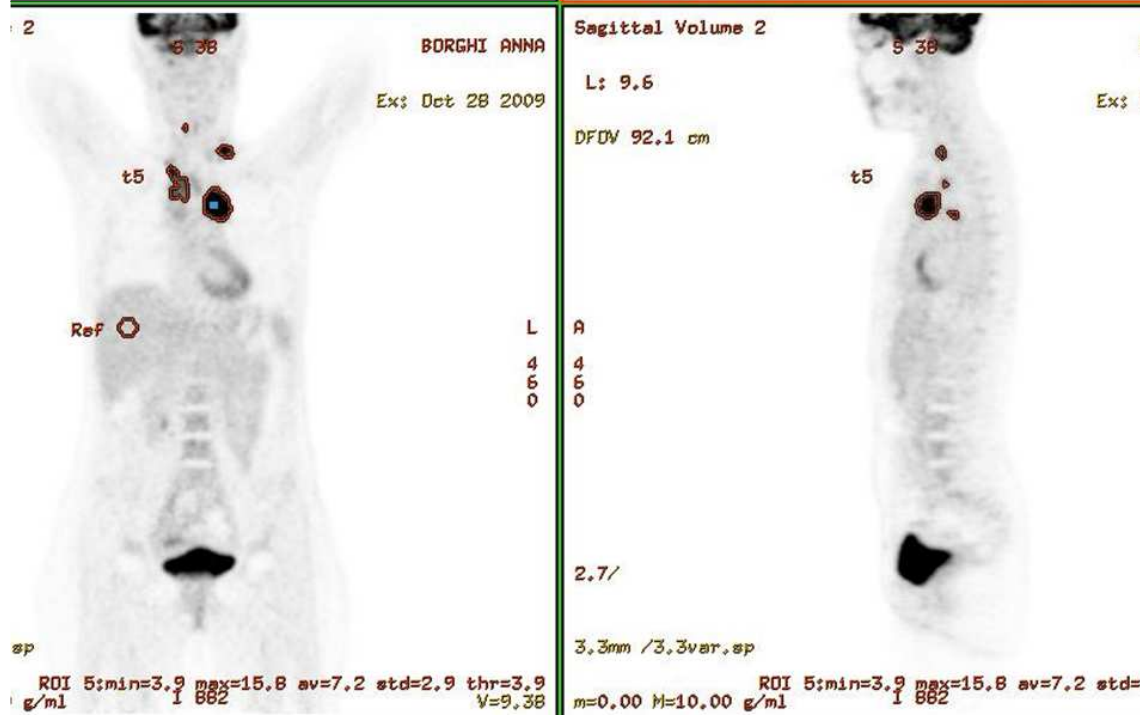
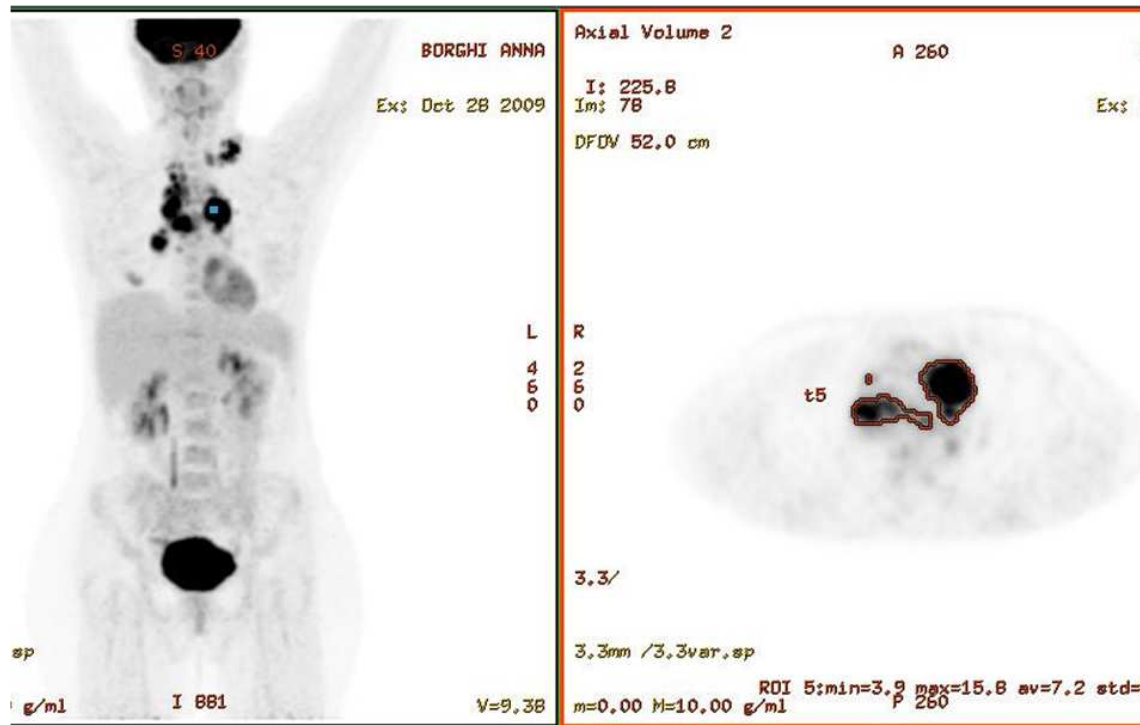
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- **Homogeneous bone marrow** uptake was not included in the tumour volume
- **Spleen involvement** was considered if there was **focal uptake or diffuse uptake higher than 150 % of the liver background**



Problems

- Not homogeneous lesions (different SUV max)
- Background (organs with high uptake,...)



All the individual lesion volumes are added together to give the **total metabolic tumour volume at baseline (TMTV0)**.

Total Lesion Glycolysis (TLG)

Calculation

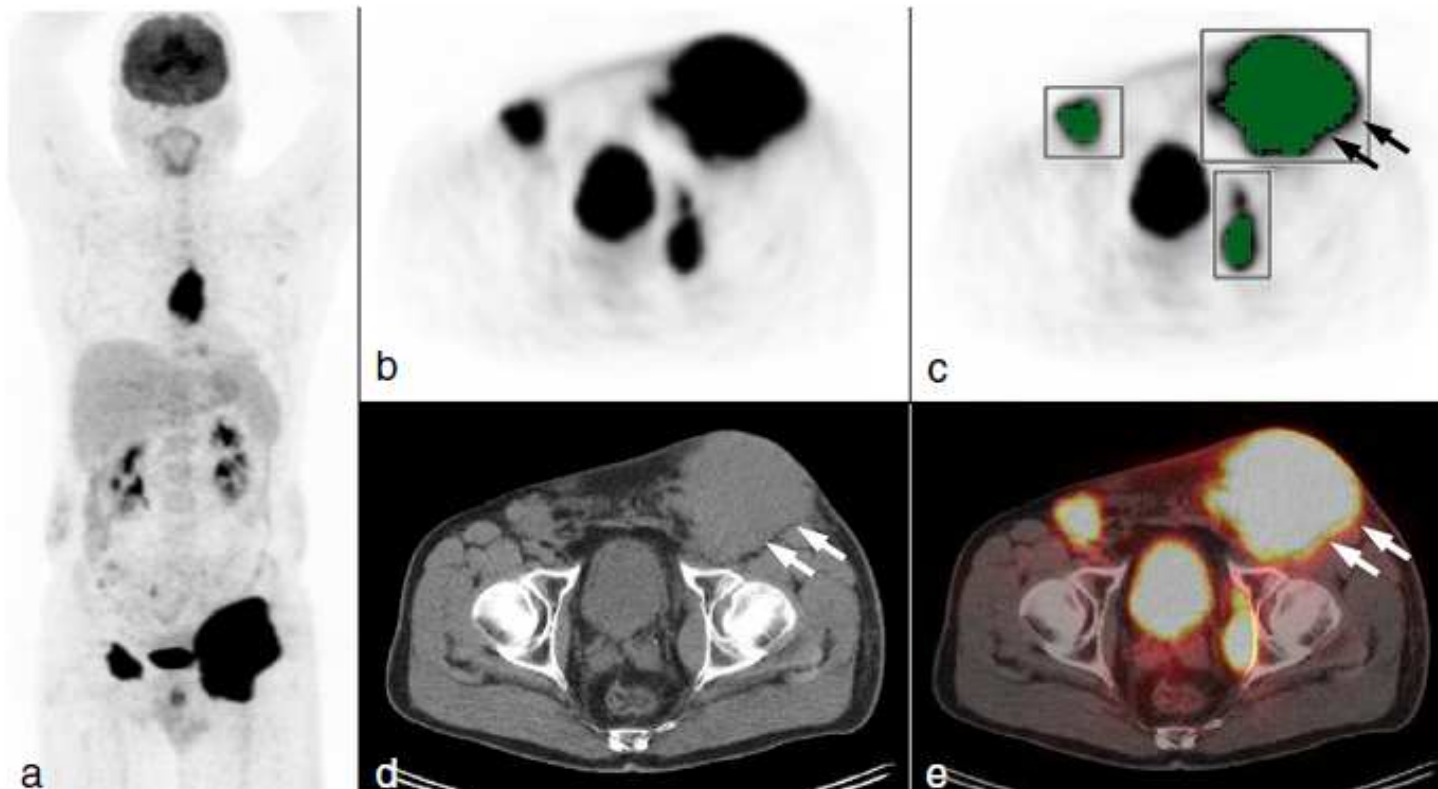
mean SUV

x

metabolic tumor volume

Total Lesion Glycolysis in Positron Emission Tomography Is a Better Predictor of Outcome Than the International Prognostic Index for Patients With Diffuse Large B Cell Lymphoma

Tae Min Kim, MD, PhD¹; Jin Chul Paeng, MD, PhD²; In Kook Chun, MD²; Bhumsuk Keam, MD¹; Yoon Kyung Jeon, MD, PhD³; Se-Hoon Lee, MD, PhD¹; Dong-Wan Kim, MD, PhD¹; Dong Soo Lee, MD, PhD²; Chul Woo Kim, MD, PhD³; June-Key Chung, MD, PhD²; Il Han Kim, MD, PhD⁴; and Dae Seog Heo, MD, PhD¹



Total Lesion Glycolysis (TLG)

mean SUV : advantages and problems of SUV

x

metabolic tumor volume

Conclusions

Metabolic Tumor Volume (MTV)

- Many software available
- Clear rules for definition are needed
- 41% threshold recommended for a better reproducibility

Total Lesion Glycolysis (TLG)

- Interesting
- Problems of SUV

Thanks for your attention

Calatrava Bridge – Reggio Emilia (Italy)

