

# Healthcare Challenges for CLASP Imaging Technology

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# Identified Challenges in Imaging

- Dose reduction in ionizing imaging modalities. Large patient imaging.
- Effect of patient movement, respiratory/cardiac, reducing resolution and accuracy of data.
- Sensitivity of detectors.
- Computer Aided Diagnosis/Pattern Recognition.
- Improved Image Reconstruction; Automatic Image Analysis.
- Image alignment from multiple modalities.

# Challenges Linked to 'Big Data'

- Data mining from large imaging archives – can we use this data to develop computer aided diagnosis which is machine/protocol specific.
- Data migration through system replacement. How do we manage technological advances and the use of normal ranges/results.
- Can we link imaging findings to biochemistry/pathology data.

# Potential Clinical Issues

- Ionizing v Non-ionizing modalities
- Function v Structure
- Multimodality
- Technology developments
- Image Guided Therapy –  
Radiotherapy/IMRT/IGRT
- Image Guided Surgery/Robotic Surgery
- Image Optimization/System Modelling

# An Example from PET/CT and Radiotherapy.

- How do we correct for respiratory/cardiac motion across different modalities and therapy regimes
- (Also applies to SPECT and SPECT/CT technologies – May be less of an issue in PET/MR if we can acquire simultaneously)
- (Will apply to any imaging conducted sequentially in different timescales)
- Motion always a degrading factor in any imaging.

# What are we trying to achieve with PET in Radiotherapy Planning?

- Identify and treat an anatomical structure/volume with a particular biological function with high sensitivity and specificity (better than current gold standards).
- Identify and quantify an anatomical sub structure volume with a particular (quantifiable) biological function to inform a treatment plan. (IMRT)

# What are we trying to achieve with PET in Radiotherapy Planning?

- A 'Biological Target Volume (BTV)' .
- The definition of a planned target volume (PTV) to effect a better cure rate with reduced normal tissue damage/complications.

**Mah et al, The impact of FDG PET on target and critical organs in CT-based treatment planning of patients with poorly defined NSCLC: a prospective study,  
Int. J. Radiat. Onc. Biol. Phys., 52, 339-350, 2002**

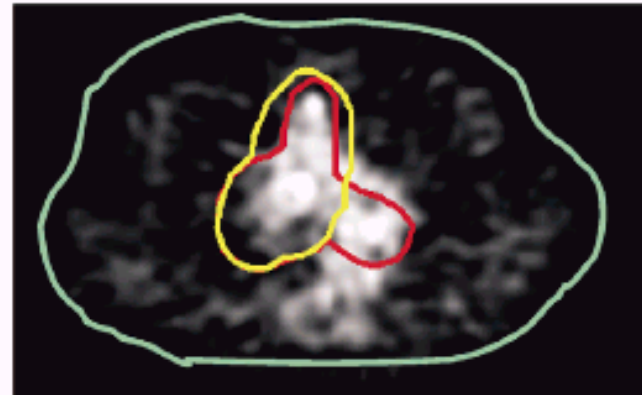
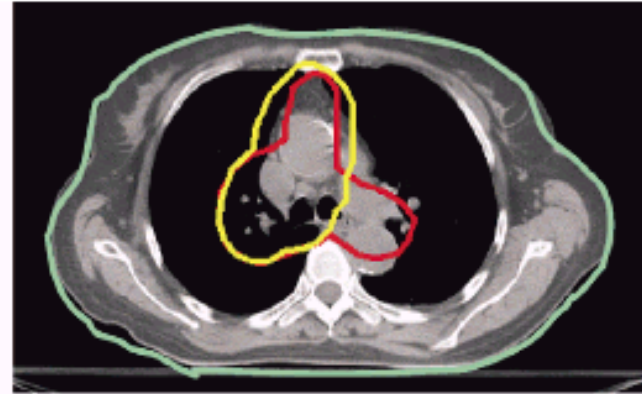
- 30 patients referred for radical treatment
- 3 radiation oncologists independently identified separate GTV's on the CT and fused PET-CT images
- With inclusion of PET information 7 of 30 patients changed to palliative intent (detection of disease spread)
- Addition of PET information led to change in size (reduction 24%-70%; increases 30%-76%) , shape and location of PTV
- Integration of FDG PET with CT lowers observer variation and provides more consistent definition of GTV



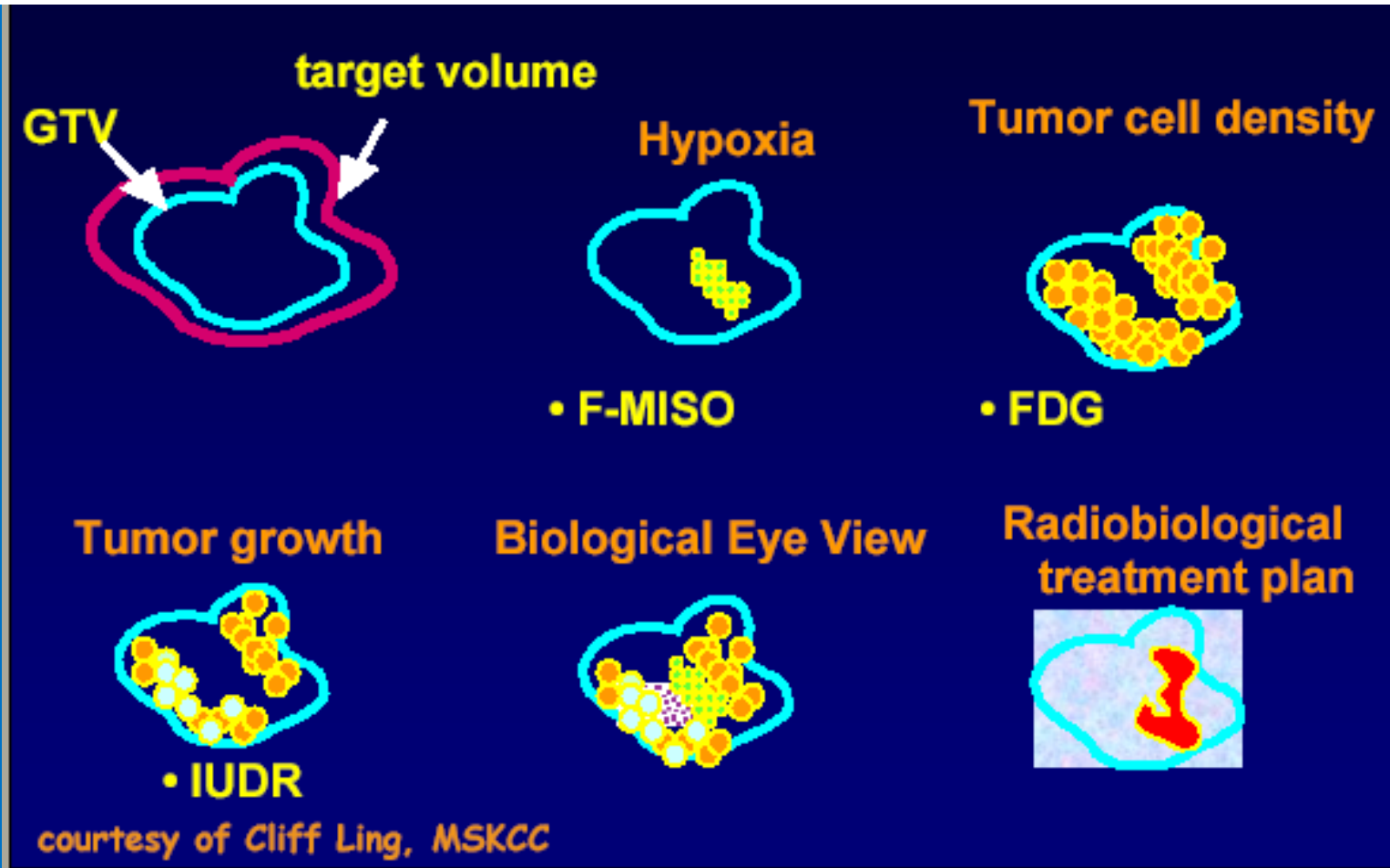
## Erdi et al, Radiotherapy treatment planning for patients with NSCLC using PET, Radiotherapy and Oncology, 62, 51-60, 2002

- 11 patients: referred for RT
- GE Advance PET Scanner
- Registered transmission PET with CT
- PET led to change in PTV for all patients - 7 increase, 4 decrease
- PET should be part of treatment planning, but outcome analysis needed to evaluate cost effectiveness and overall utility

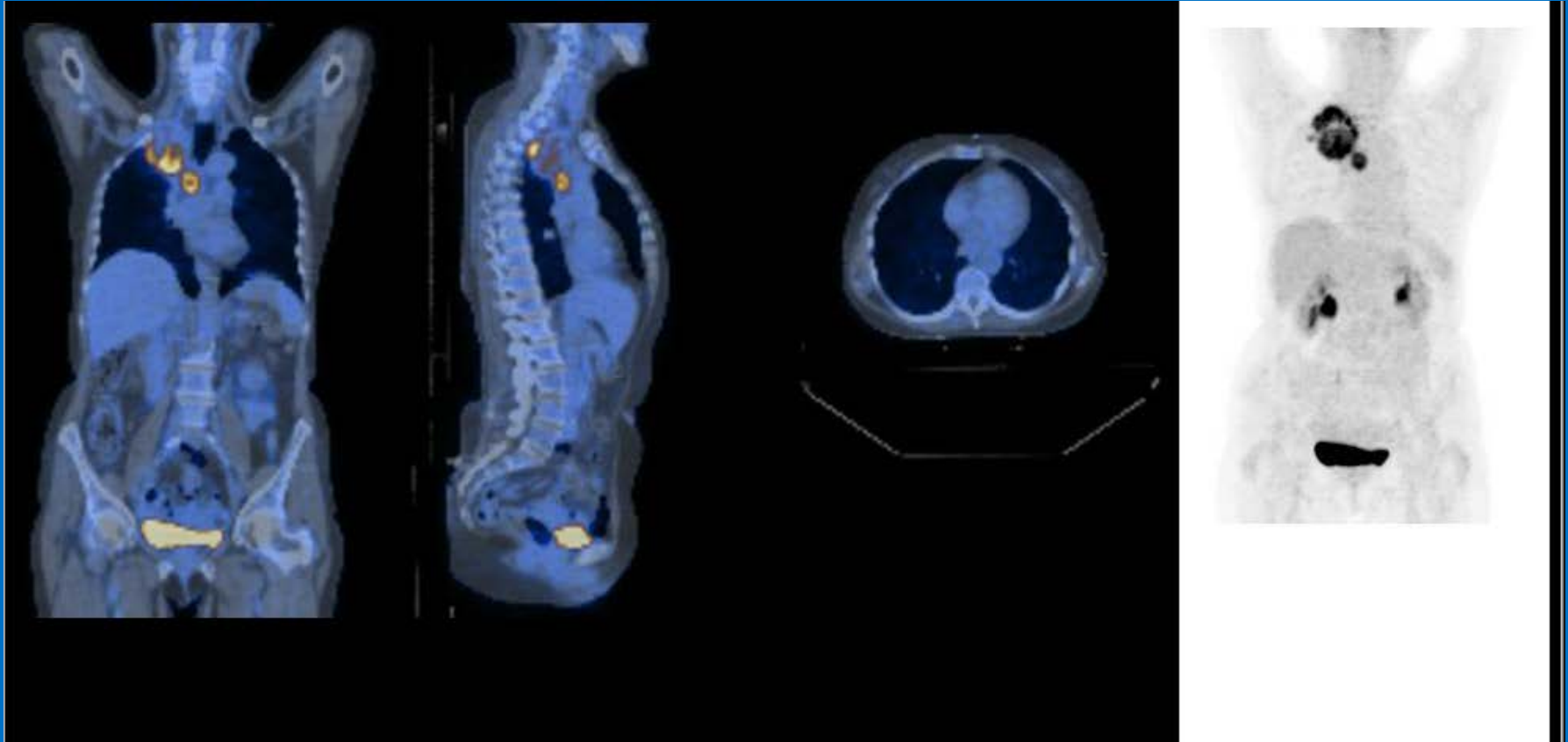
# Planning using PET/CT – Erdi et al



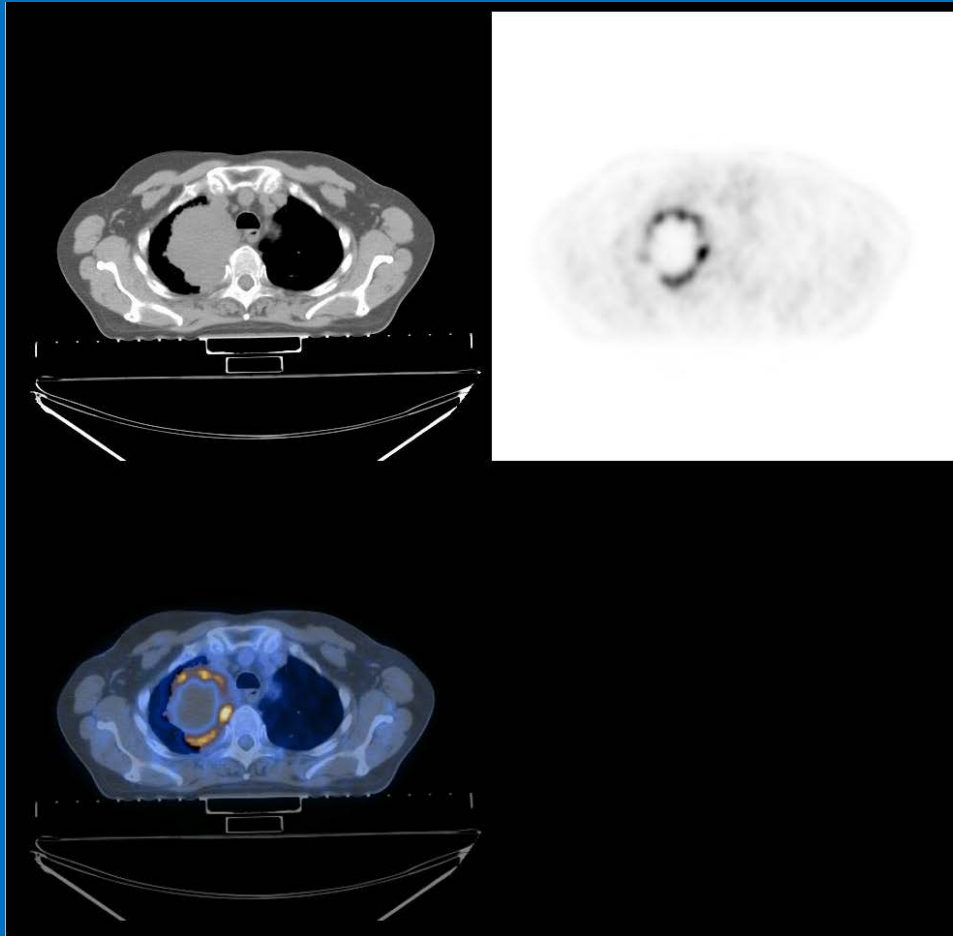
# A Biological Treatment Plan



# PET-CT images from radiotherapy planning patient



# PET/CT Planning Dataset



# PET/CT Integration - Issues

- Misalignment of transmission and emission data primarily due to differential motion either respiratory or gross movement.
- CT generated artefacts from heavy metal objects, IV and Oral contrast.
- Emission acquisition slow and averages patient physiological motion.

# Respiratory motion

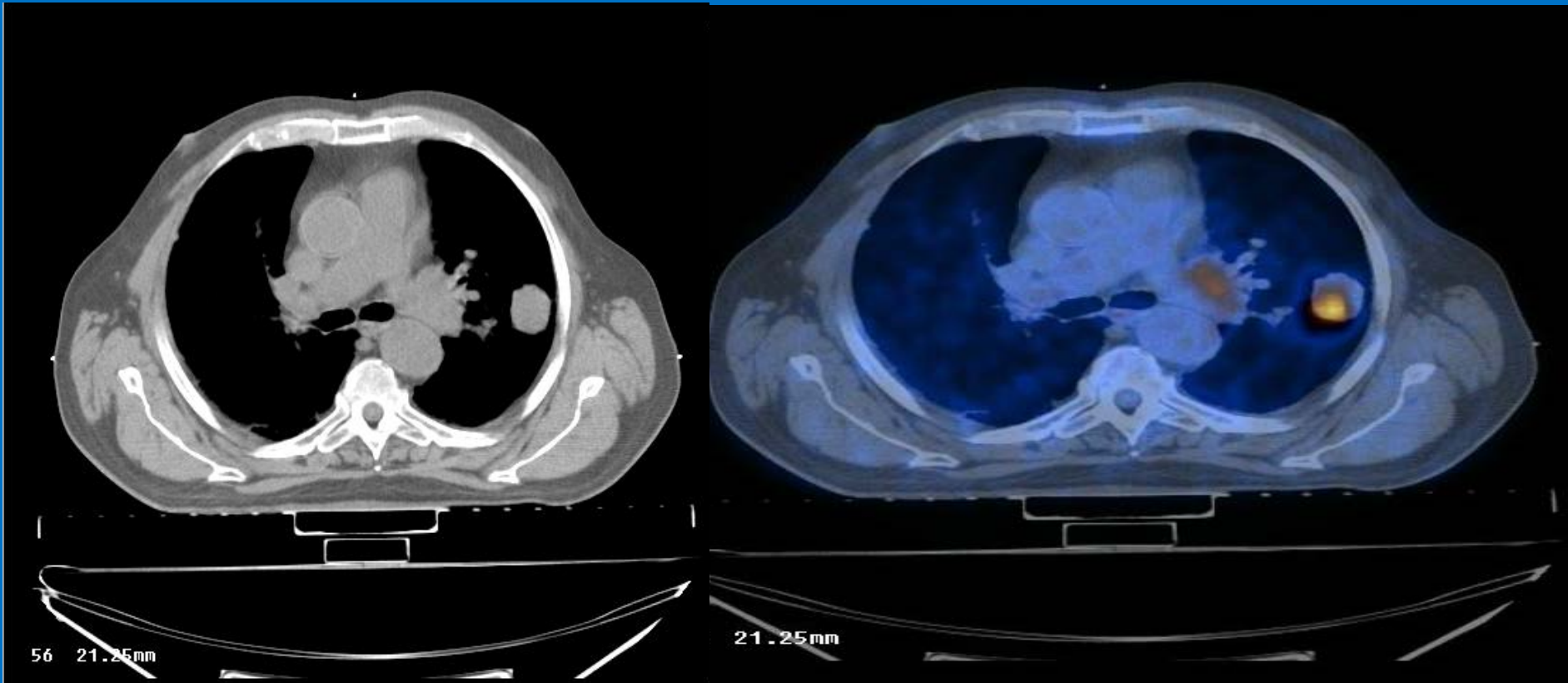


# Effect of respiration in PET- CT

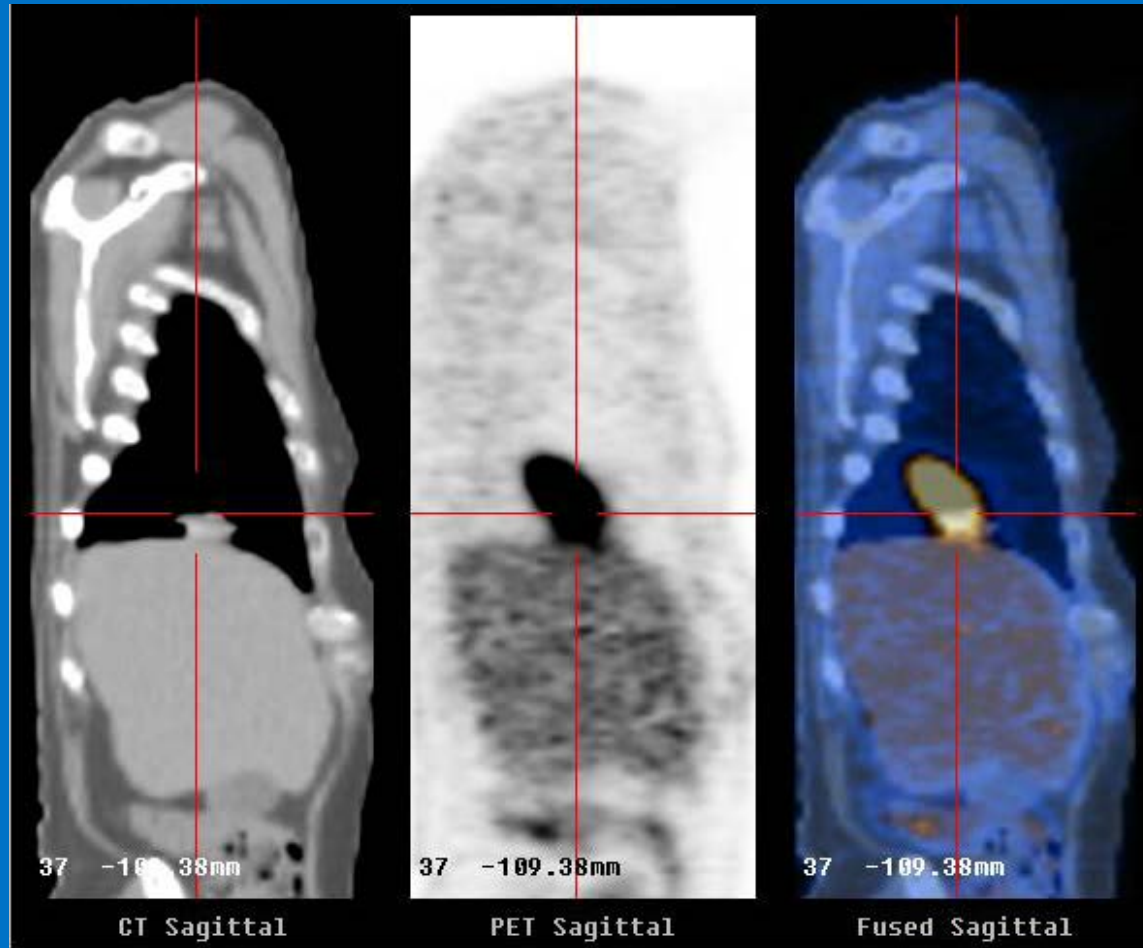
- Blurs the PET images
  - Reduces image quality
  - Reduces calculated FDG uptake
  - Reduces contrast in the images
  - Affects size of determined lesion
- Causes mis-alignment with CT
  - position
  - size/shape
- Incorrect Attenuation Correction
  - Unknown impact on uptake values.



# Mis-registration of PET and CT due to breathing?



# Example of breathing artifact in PET-CT data



# Example of breathing artifact in PET-CT data



Attenuation correction using a respiration average transmission map with radioactive rod sources

Attenuation correction using a CT map acquired at end inspiration

Attenuation correction using a CT map acquired at end expiration or under free breathing

# How to deal with respiratory motion?

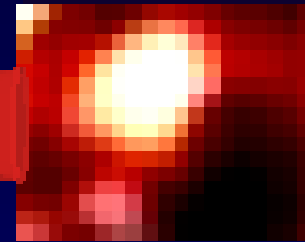
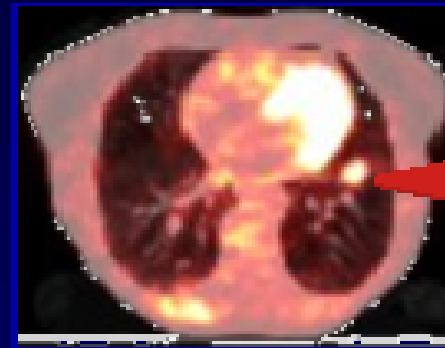
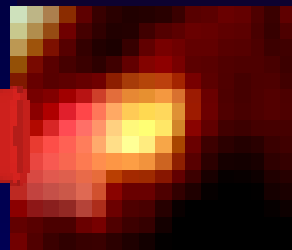
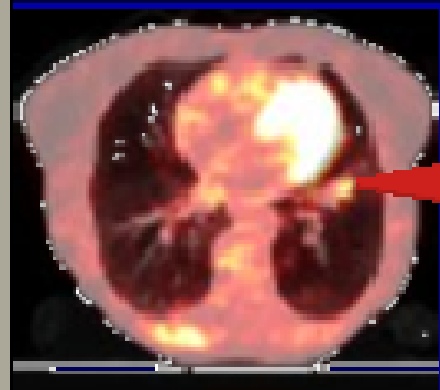
- Assume PET volume includes breathing motion (CTV + IM)
- CT breathing protocols to improve registration with PET e.g. max expiration (inspiration) breath hold
- Respiratory gated PET/CT (and RT)
- Respiratory-correlated Dynamic PET

# Effect of Gating

- Reduction in lesion volume
- Improves quantification (increases SUV)
- Improves registration with CT



# Advantages of 4D PET - Improved SUV



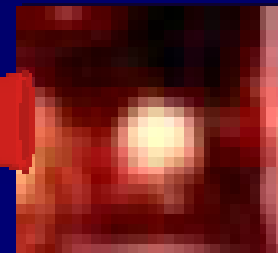
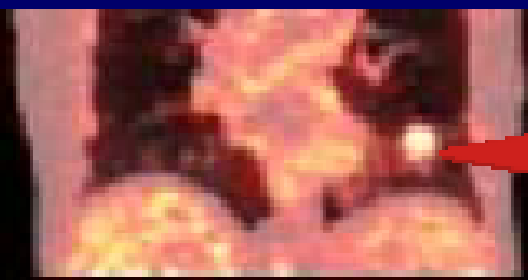
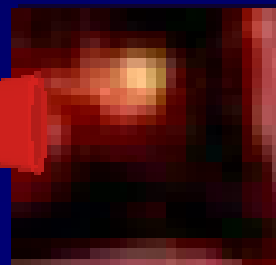
$SUV_{max} = 3.3$



34 %



$SUV_{max} = 4.4$



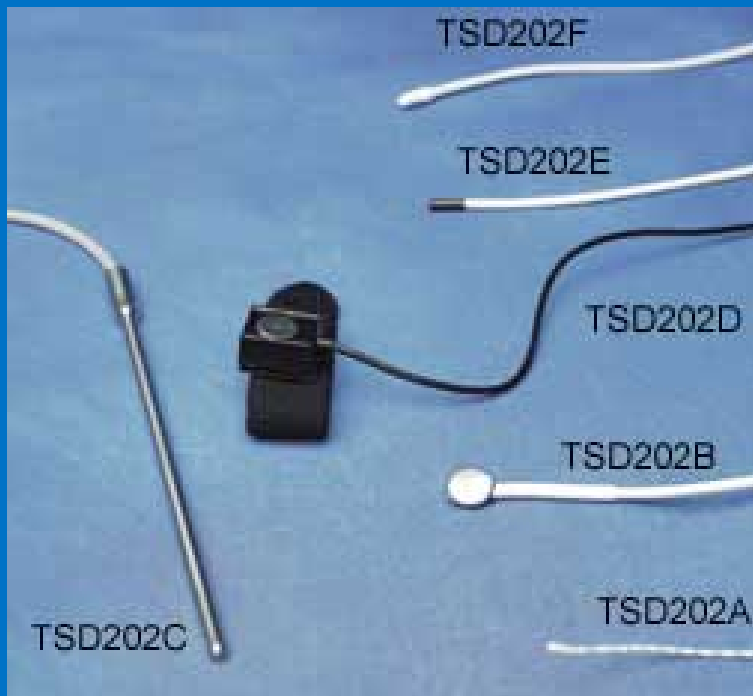
4D PET with Clinical CT

4D PET with 4D CT

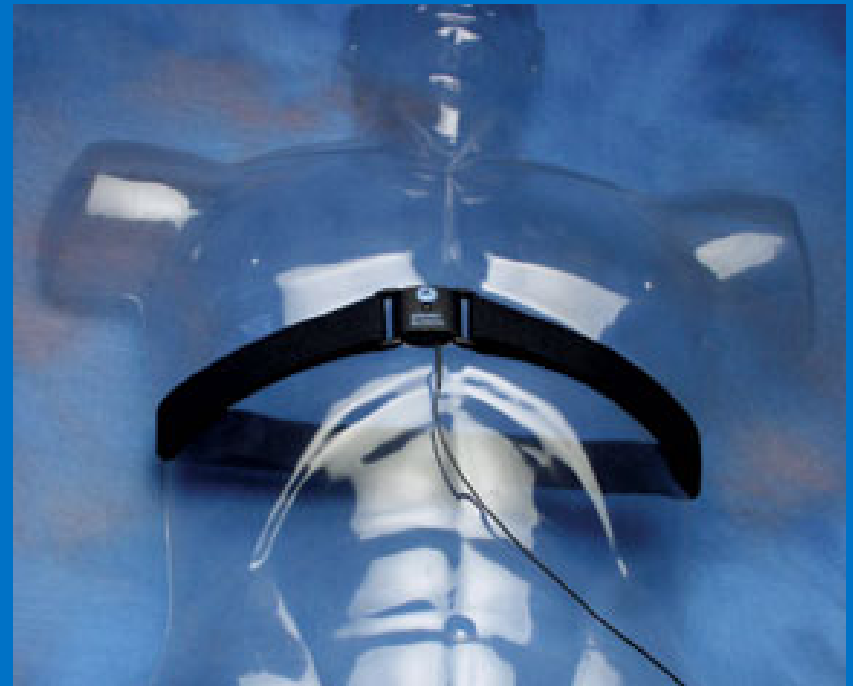
# Respiratory Gating

- Breathing synchronisation should be based upon periods of equal lung volume.
- Segmentation based on equal time slices or phase slicing will be inappropriate.
- Physiological signals providing data on lung volume required.
- Intrinsic gating from dynamic/list mode data.
- Synchronization of data streams required.

# Options for respiratory monitoring



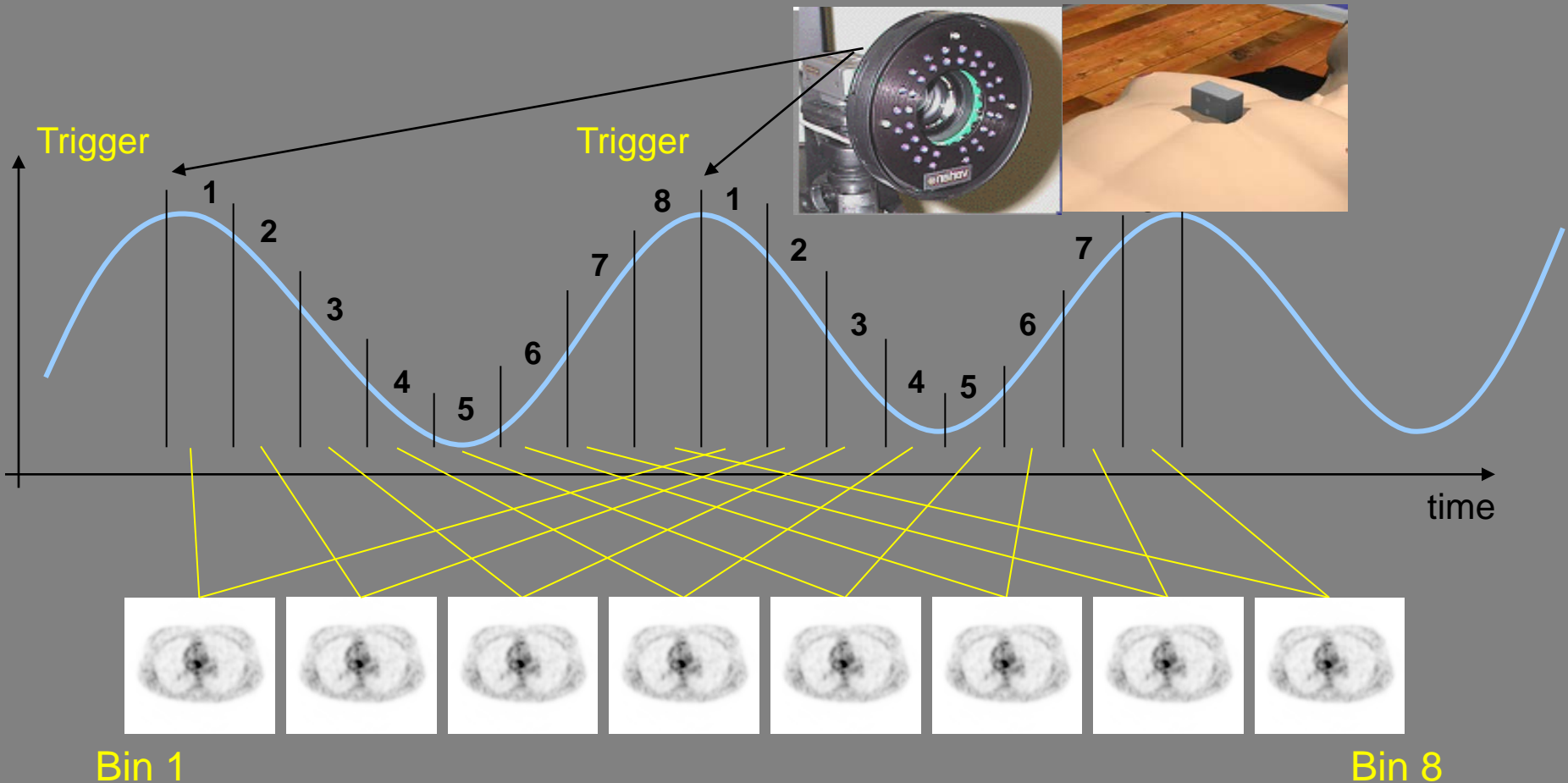
Thermistor



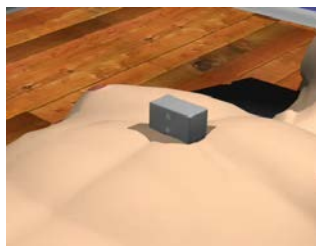
Respiratory belt



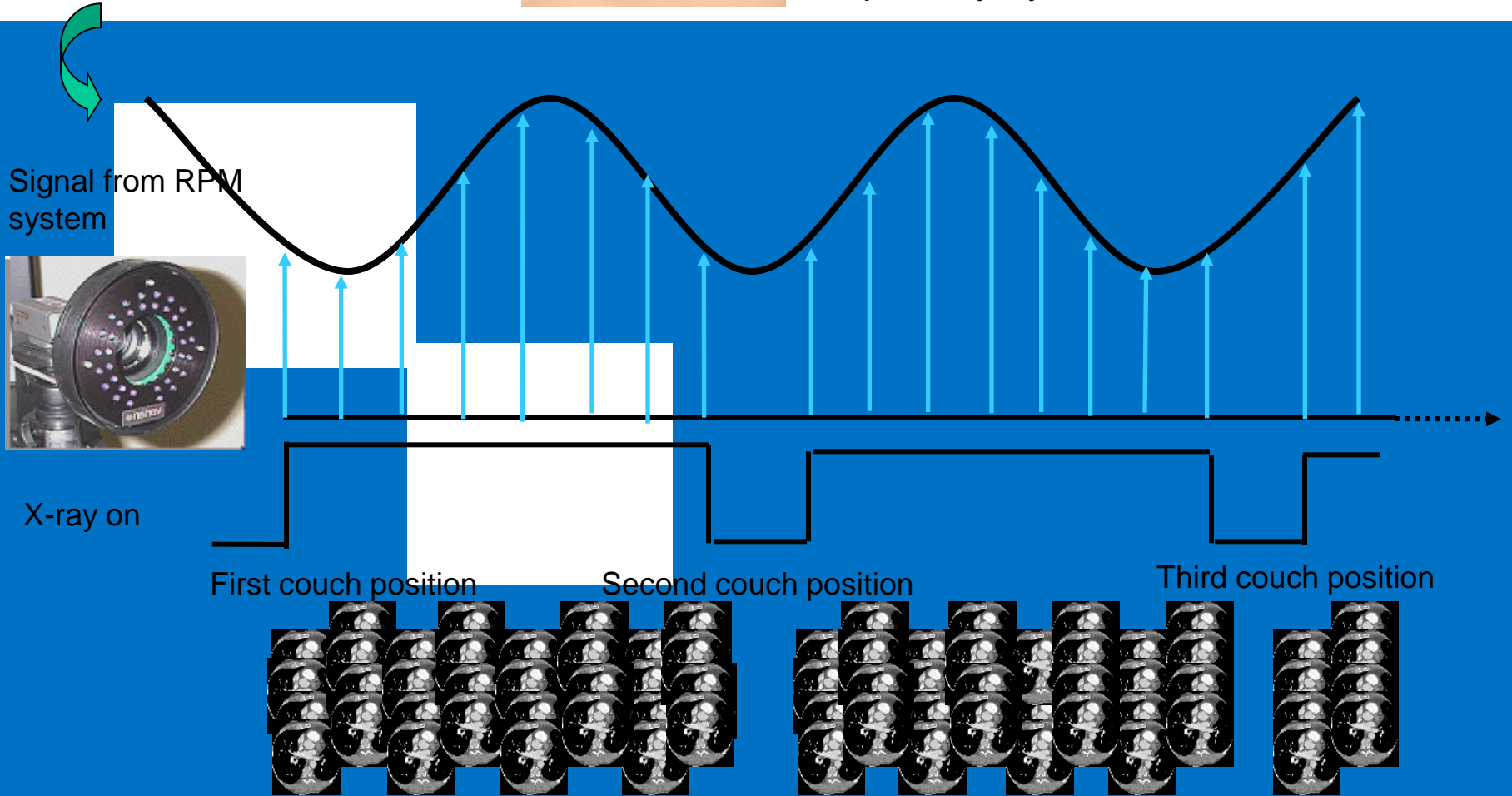
# Infra-red reflectors



# 4D PET-CT



- Respiratory tracking with Varian RPM optical monitor
- CT images acquired over complete respiratory cycle



# Using Anatomical Information from CT to correct PET

Gated CT taken over one respiratory cycle



Find transformations to ref CT frame

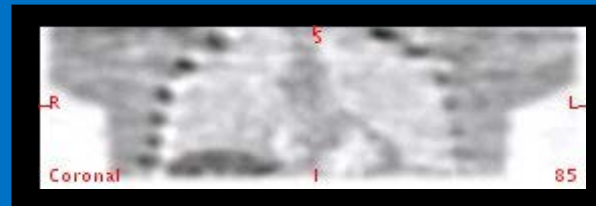


Gated PET over one respiratory cycle : low statistical quality



Apply transformations from CT-CT registration

Corrected PET



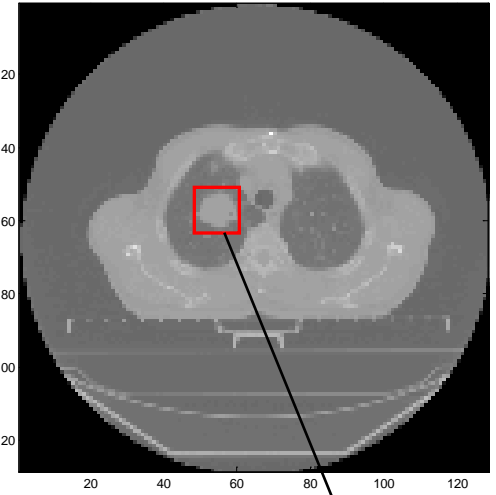
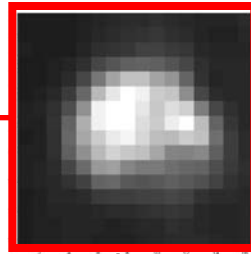
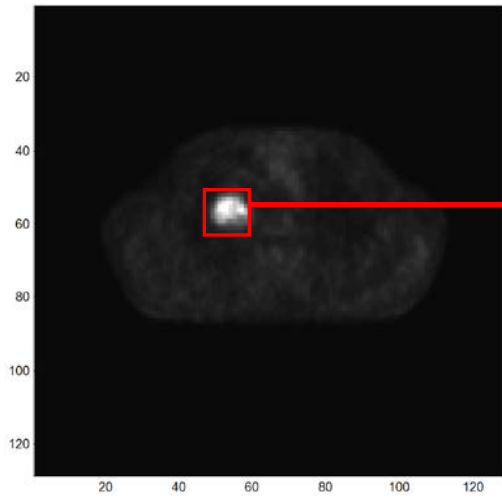
# Technology Issues

- Respiration correction during acquisition a compromise.
- How do we manage motion during delivery of therapy?
- How do we manage changes in anatomy/function during therapy treatment?
- Multiple use of ionizing radiation increases dose and cancer risk. Higher sensitivity/different technology required.
- How do we optimize dose/image quality?

# Tumour Delineation.

- Identification of tumour volumes if defined from PET still require validation. The use of a fixed threshold value is inadequate for all lesion volumes, shapes and target to background ratios.
- Signal to noise ratios very poor for respiratory gated studies.
- Assumptions of uniform uptake in tumour invalid.

ROIs center on lesion : 16x16



SEGMENTATION ALGORITHMS

T50

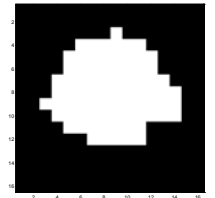
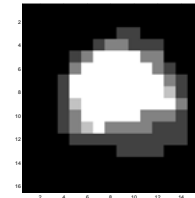
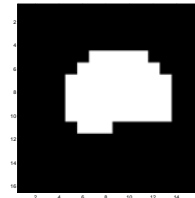
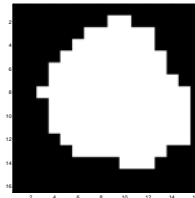
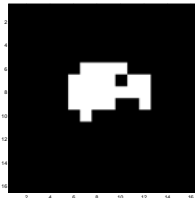
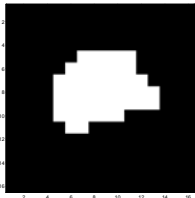
T75

T<sub>Std</sub>

FCM

FHMC

Manual



Vol. = 3536

Vol. = 1807 mm<sup>3</sup>

Vol. = 9114 mm<sup>3</sup>

Vol. = 4007 mm<sup>3</sup>

Vol = 6522 mm<sup>3</sup>

Vol. = 6207 mm<sup>3</sup>

C<sub>avg</sub> = 24843

C<sub>avg</sub> = 28438

C<sub>avg</sub> = 14897

C<sub>avg</sub> = 23714

C<sub>avg</sub> = 18572

T50 = Threshold 50% of max pixel value, T75 = Threshold 75% of max pixel value.

T<sub>Std</sub> = Threshold 3 x Std dev. Measured on background ROI. FCM = Fuzzy C-Means clustering

FHMC = Fuzzy Hidden Markov Chains, 2 hard classes and 2 fuzzy levels

C<sub>avg</sub> = average count density

# Challenges

- Routine motion correction methodology for whole body PET imaging.
- Accurate and automatic uptake and volume quantification of uptake distributions.
- The utilization of high resolution anatomical data to restore low resolution PET distributions.

# Opportunities

- To ensure that radiotherapy treatments are delivered and optimized to provide the maximum cure rate whilst minimizing damage to normal tissues.
- To integrate the advances in imaging technology with advances in radiotherapy delivery methods.
- To ensure that new methodologies are evaluated through clinical trials