
Quantitative Imaging: CT and MRI

Sources and Mitigation of CT Quantitative Imaging Errors

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Disclosures

- UCLA Department of Radiological Sciences has a Master Research Agreement with Siemens Healthineers



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Quantitative Imaging

- What does it take to make Imaging Quantitative?
- Go from making an Image
- To
- Making a Measurement



3

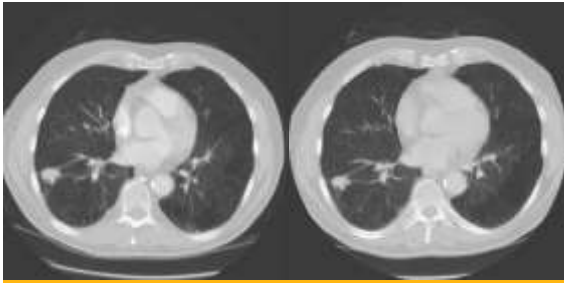
Example: How Big Is This Lesion?



What size metric should we use? Currently use one or two linear measurements



Example: Did Lesion Change in Size?



Quantitative Imaging in CT

- CT is inherently Quantitative (isn't it?)
- Each voxel reports a CT number
- And it even has units (HU)
- Which are defined internationally

• CT number =
$$\left(\frac{\mu_{\text{tissue}} - \mu_{\text{water}}}{\mu_{\text{water}}} \right) * 1000$$

- Water ($\mu = \mu_{\text{water}}$) ---> 0 HU
- Air ($\mu \sim 0$) ---> -1000 HU



CT to Measure Change

- Change in Size
- Change in Density
- Change in Texture
- Change in Function (Perfusion, etc.)

- Can we measure these Changes Reliably?
 - Good enough to aid Dx?
 - Or Assess Treatment Efficacy?



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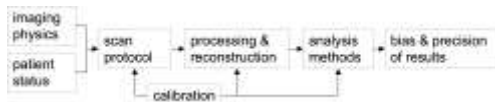
CT to Measure Change

- Can we do this in a robust fashion
 - Across scanners
 - Across centers
 - Across patients (with similar condition/disease)



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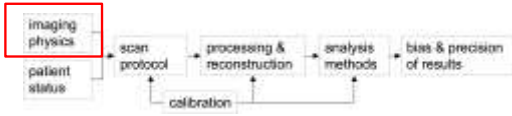
System Description – Sources of Variability





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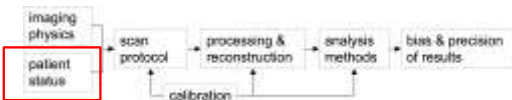
System Description – Sources of Variability



CT Imaging Physics Considerations

- Scanner Design
 - Geometry e.g. Number of Detector Rows
- Scanner Operation
 - kV, mAs, pitch
- Capabilities:
 - Dose reduction technologies
 - Advanced (Iterative) reconstruction
 - Dual Energy
 - Other

System Description – Sources of Variability



Patient Considerations

- Health Status of Individual patient
 - Ability to breathhold if required
 - Ability to use oral or IV contrast
 - Ability to perform study without motion
- Abnormalities and Concomitant Disease
 - Inflammation which may mask progression
 - Patient Health Status during trial



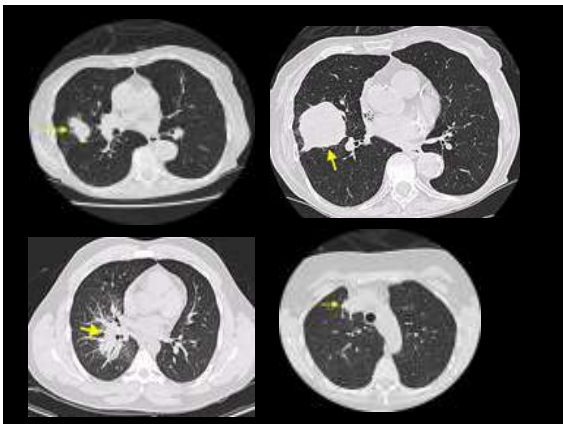
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Tumor Related Considerations

- Complexity of Tumor
 - Shape (Spherical or Complex) can make determining boundaries "difficult" (i.e. not reproducible)
 - Location
 - Physiology (contrast uptake, washout)



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System Description – Sources of Variability



Scan Protocol Considerations

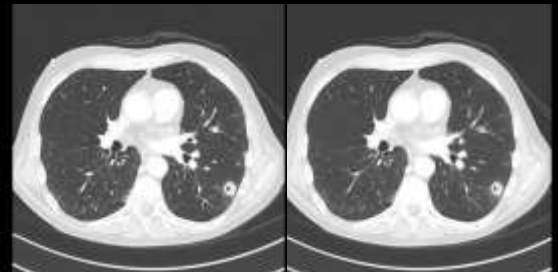
- Reconstructed image thickness
- Reconstructed image interval
- Reconstruction filter

- Resolution and Noise

Example Lung Lesion

1 mm B45

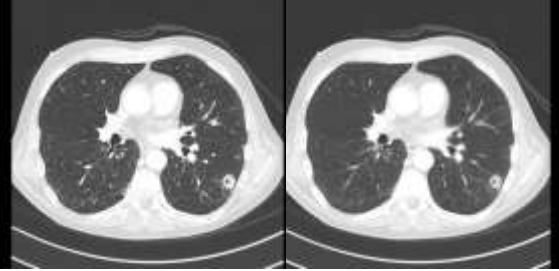
3 mm B45



Example Lung Lesion

1 mm B45

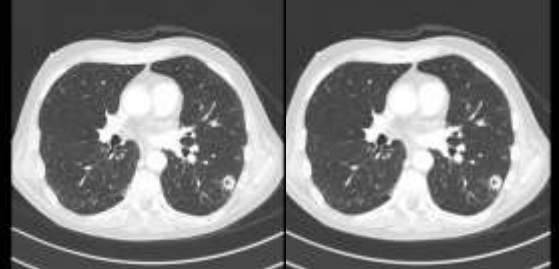
5 mm B45



Example Lung Lesion

1 mm B45

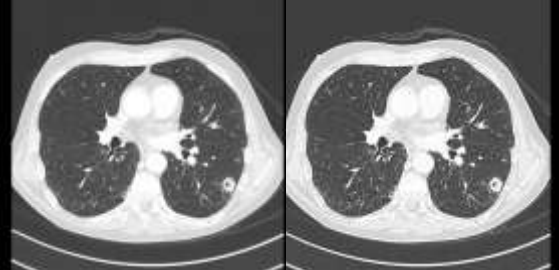
1 mm B10



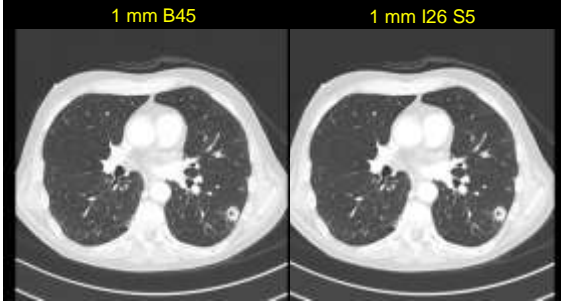
Example Lung Lesion

1 mm B45

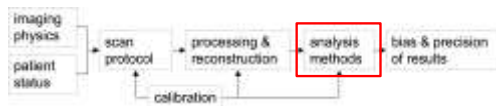
1 mm B80



Example Lung Lesion



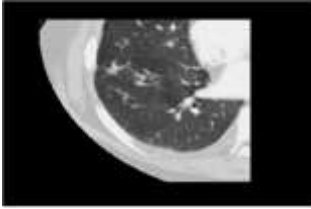
System Description – Sources of Variability



Analysis Method

- Fully Automated
- Some human intervention
 - Radiologist measuring diameter
 - Contouring boundary
- Measurement itself
 - Diameter
 - Volume
 - Mass/density
- Registration method if change is measured

Original Image



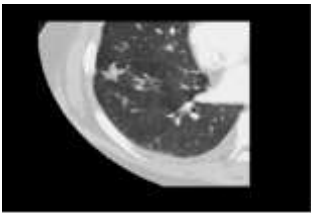
From Meyer et al, Acad Radiol. 2006



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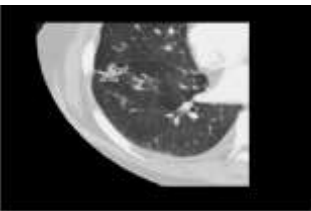
Contour 1



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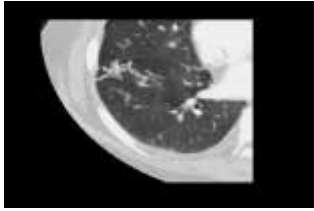
Contour 2



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Contour 3



Possible Mitigation Strategies

- Limiting the acceptable conditions under which measurements are made
 - Only allow measurements under specified conditions for each of the elements in the measurement chain;
 - Imaging Physics conditions
 - Patient conditions (disease, health status, nodule size/shape)
 - Acquisition/Reconstruction Conditions
 - Analysis (Segmentation, Feature Calculation) Conditions

Some Attempts at Standardization

- National Lung Screening Trial (NLST)
 - No quantitative measure as an endpoint
 - Positive Screen meant there was a nodule present or not on CT or CXR
- ACRIN 6678
- COPD/Gene

- These have informed the RSNA QIBA Protocols
<https://www.rsna.org/QIBA-Profiles-and-Protocols/>

From Cagnon et al Academic Radiology, 2006

Table 1
Acquisition of CT Technique Comparison Chart: Scanner Specific Techniques Monitored by MIST Protocol

Parameter	GE QX 4 × 160/4.5 sec	GE L5 Pro 4 × 160/3.5 sec	GE 160 4 × 160/3.5 sec	GE L5 V 16 × 160/3.5 sec	GE VCT Pro 16 × 160/3.5 sec
KT	100	100	100	100	100
Scatter Reduction Time	8.9 sec	10.9 sec	6.5 sec	9.1 sec	8.3 sec
mAs (Regular pattern Large patient reduced)	80-90	80-90	80-90	80-90	80-90
mAs (Regular Large)	80-90	80-90	80-90	80-90	80-90
Scanner effective mAs ¹ (Reg Log)	37.7-39	36.5-38	36.6-38.2	39.1-40.3	37.3-38
Detector Collimation (mm)–T	2.5 mm	2.5 mm	2.5 mm	2.5 mm	2.5 mm
Number of active detectors–R	8	8	8	16	16
Detector Collimation–R, L, T	8 × 2.5 mm	8 × 2.5 mm	8 × 2.5 mm	16 × 2.5 mm	16 × 2.5 mm
MIST (Thin/Thick)	1.0/1.0	1.0/1.0	1.0/1.0	1.0/1.0	1.0/1.0
Table acceleration (cm/min/sec)	6.0 mm	6.0 mm	12.0 mm	11.0 mm	10.0 mm
Pitch (cross-table) from collimation–R/L	1.0	1.0	1.25	1.0	0.98
Table Speed (mm/sec)	18.75 mm/sec	18.75 mm/sec	22.5 mm/sec	18.75 mm/sec	18.75 mm/sec
Scan Time (min:sec)	32 sec	15 sec	18 sec	7.5 sec	6.5 sec
Minimal Reconstructed Slice Width	2.5 mm	2.5 mm	2.5 mm	2.5 mm	2.5 mm
Reconstruction Interval ²	2.5 mm	2.5 mm	2.5 mm	2.5 mm	2.5 mm
Reconstruction Algorithm ³	FC	FC	FC	FC	FC
# Images/Scan (at 100 mm interval)	390	390	390	390	390
C-FLU (Scan to mAs) (Regular Large)	0.9-1.0 mAs	0.9-1.0 mAs	0.9-1.0 mAs	0.9-1.0 mAs	0.9-1.0 mAs

Parameter	Philips 4 × 160/3.5 sec 4 × 2.5	Philips 4 × 160/3.5 sec 4 × 1	Philips 4 × 160/3.5 sec 16 × 7.5	Siemens 16 × 160/3.5 sec 4 × 160/3.5 sec	Siemens 16 × 160/3.5 sec 16 × 160/3.5 sec
KT	100	100	100	100	100
Scatter Reduction Time	10.9 sec	8.9 sec	10.9 sec	8.9 sec	8.9 sec
mAs (Regular pattern Large patient reduced)	70-75	80-85	70-75	80-85	80-85
mAs (Regular Large)	70-75	80-85	70-75	80-85	80-85
Scanner effective mAs ¹ (reconstruction) (Reg Log)	35-38	36-40	35-38	36-39	36-39

Possible Mitigation Strategies

- Post-Acquisition/Reconstruction steps
 - Image DeNoising
 - Image Normalization
 - Regardless of the original Acquisition/Reconstruction Parameters, transform the images to a standard set of conditions (thickness, spatial resolution, contrast resolution, noise, etc.)
 - Deep Learning (!!!!!)

Possible Mitigation Strategies

- These strategies may vary based on task
- Mitigation used to measure change in volume MAY NOT BE same used measure change in texture.

Summary

- Converting an Imaging Modality to a Measurement System is not easy
- LOTS of sources of variability (including patient)
- Mitigation methods being explored
- Sources of Variability AND Mitigation methods may be task dependent.

Summary

- As we listen to presentations today, let's consider:
- What is the task? (Or is a specific task identified?)
- What are sources of variability?
- What is the mitigation approach?
- How do these affect QI metric?
 - Repeatability?
 - Reproducibility?
- (NOTE: Each presentation does NOT have to solve ALL of these problems at once! ☺)

Thank You

Terminology

From a previous QIBA Annual Meeting
“Ten Things to Remember from the QIBA Metrology
Workshop” by Nancy Obuchowski, PhD
“1. Do not use *repeatability* and *reproducibility*
interchangeably”



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Repeatability

- This is the *within-subject* variability
- It is the agreement between measurements made within a short period of time (test-retest) holding variables constant
 - Example: Coffee Break Experiments
- Includes variability due to scanner adjustment, image noise, subject positioning



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Reproducibility

- The observations are performed on the same subject (usually) over a short period of time, **but the location, operator and/or measuring system differs**



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