



Correlating Radiomics Information with Clinical Outcomes for Lung SBRT

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Radiation Oncology

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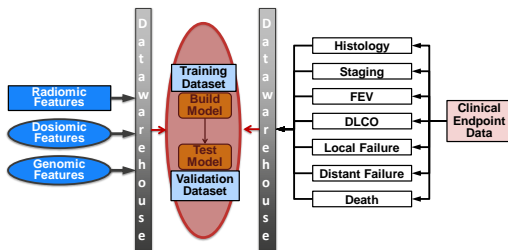
Physics

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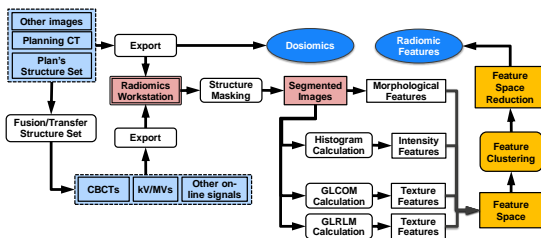
Objectives

- Correlation study between patient specific radiomics data and clinical outcomes extracted from lung SBRT
- Techniques for extracting large amount radiomics data in CT and CBCT images used for lung SBRT
- Effective data modeling for treatment assessment and prediction in lung SBRT

Applications to Lung SBRT



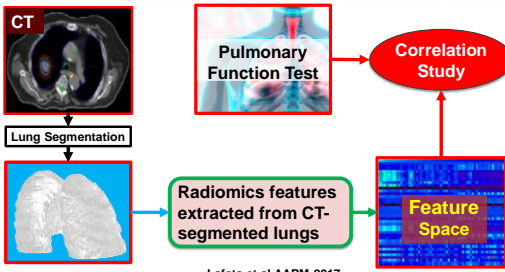
Example – Image/Data Flow



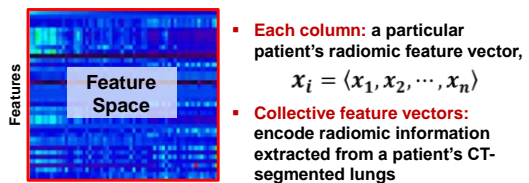
Sample Radiomic Features

Total 39 features	Autocorrelation	Intensity	Energy	Morphology	Lung Volume
	Cluster Prominence		Entropy		Lung Surface Area
	Cluster Shade	Kurtosis	4	2	
	Cluster Tendency	Skewness			
Fine Texture	Contrast	Coarse Texture	Short Run Emphasis	11	
	Correlation		Long Run Emphasis		
	Differential Entropy		Gray Level Non-uniformity		
	Dissimilarity		Run Length Non-uniformity		
	2D Energy		Run Percentage		
	2D Entropy		Run Percentage		
	Homogeneity 1		Low Gray Level Run Emphasis		
	Homogeneity 2		High Gray Level Run Emphasis		
	Information Measure of Correlation 1		Short Run Low Gray Level Emphasis		
	Information Measure of Correlation 2		Short Run High Gray Level Emphasis		
	Inverse Difference Moment Normalized		Long Run Low Gray Level Emphasis		
	Inverse Variance	Long Run High Gray Level Emphasis			
	Maximum Probability				
	Sum Average				
	Sum Entropy				
	Sum Variance				
	Variance				

CT Radiomic Features and Pulmonary Function



Build A Pulmonary Radiomics Feature Space



Feature Analysis: Multivariate analysis using a novel clustering algorithm based on Quantum Langevin Clustering (QLC)
Lafata et al AAPM 2017

Feature Clustering: Quantum Langevin Clustering

Development of Quantum Langevin Clustering (QLC): 3 steps

Step 1: Map radiomics feature vectors, x_i , to a function space, $\psi(x)$, by associating each feature with a Gaussian of width, σ , and constructing their sum (mapping from Euclidean space into Hilbert space):

$$x_i = \langle x_1, x_2, \dots, x_n \rangle \rightarrow \psi(x) = \sum_{i=1}^m e^{-\frac{1}{2\sigma^2}(x-x_i)^2}$$

- ✓ $\psi(x)$ represents probability density function of data
- ✓ Interpreted as a **quantum mechanical wavefunction**

Lafata et al AAPM 2017

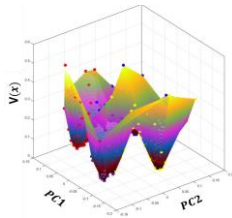
Feature Clustering: Quantum Langevin Clustering

Step 2: Inversely search for corresponding Potential Function, $V(x)$

$$\psi(x) = \sum_{i=1}^m e^{-\frac{1}{2\sigma^2}(x-x_i)^2}$$

$$V(x) = \frac{\sigma^2}{2} \nabla^2 \psi(x) + E$$

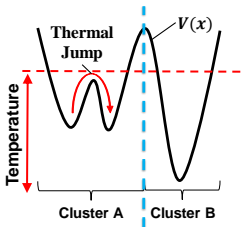
- ✓ Satisfies the **Schrodinger Equation** with solution $\psi(x)$



Lafata et al AAPM 2017

Feature Clustering: Quantum Langevin Clustering

Step 3: Propagate feature vectors through $V(x)$ via Langevin dynamics



$$dx = p dt$$

$$dp = \underbrace{-\nabla V(x) dt}_{\text{Global Force Term}} - \underbrace{\gamma p dt}_{\text{Damping Term}} + \underbrace{\sqrt{2\gamma k_B T} dB}_{\text{Brownian Motion}}$$

- ✓ **Global Force:** Potential gradient acts globally to push vectors downhill
- ✓ **Brownian Motion:** Acts locally to help vectors thermally jump local barriers into locations otherwise forbidden, controlled by **Temperature T**

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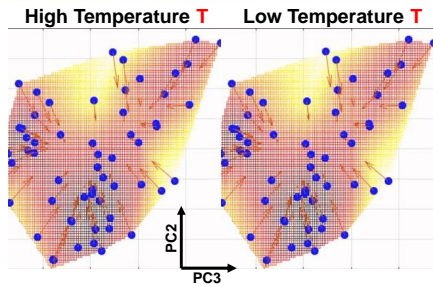
Correlation Study between QLC and Lung Function

Correlation study using QLC: Map a relationship between the radiomic feature space and pulmonary function test (PFT)



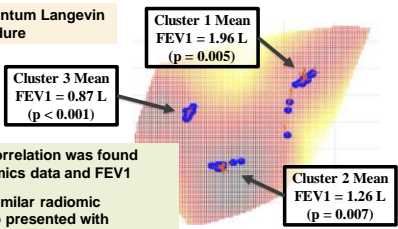
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Demonstration of Clustering Analysis



Results using QLC for Clustering Analysis

Final state of Quantum Langevin Clustering procedure

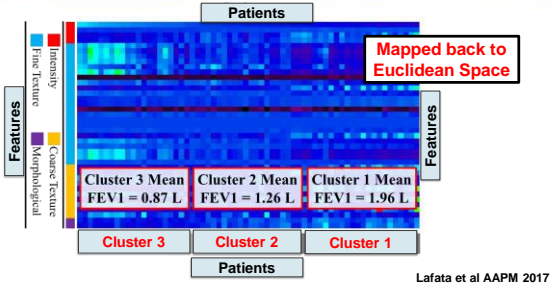


✓ A significant correlation was found between radiomics data and FEV1

✓ Patients with similar radiomic signatures also presented with comparable spirometry measurements

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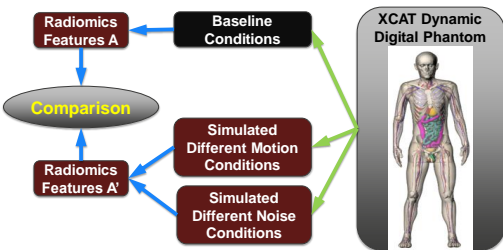
Results: Clustering Analysis



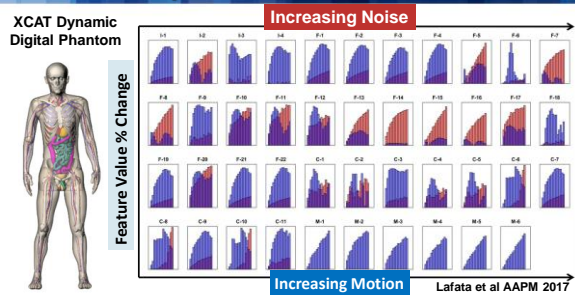
Sensitivity Analysis of Radiomics

- Effect of imaging noise
- Effect of motion
- Effect of HU inhomogeneity

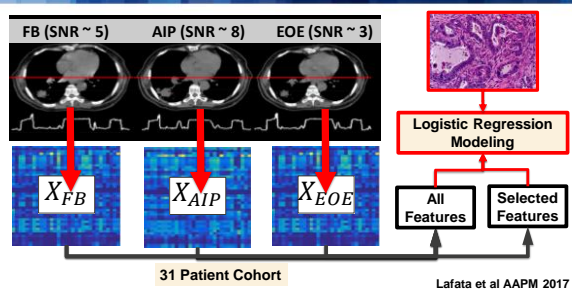
Simulation Study: Effect of Noise Motion



Simulation Study: Effect of Noise and Motion

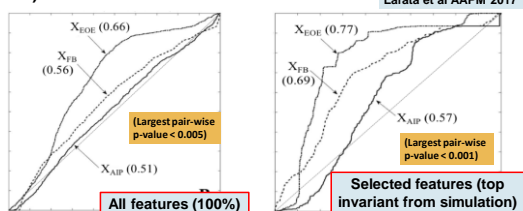


Histology Classification of NSCLC



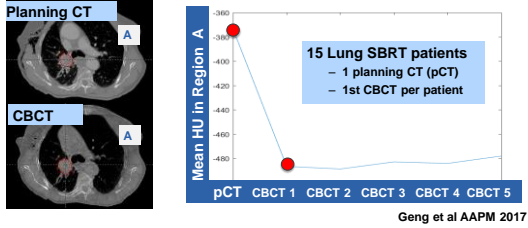
Results: Feature-Specific Model Response

- Effect on histology classification:
- 1) image noise and motion – FB, EOE, AIP
 - 2) feature selection

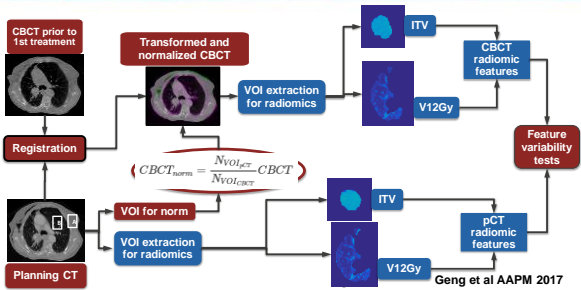


Effect of Inhomogeneous HU between CT/CBCT

- Radiomics feature variations on CBCT and planning CT
- Harmonization between planning CT (pCT) and on-board CBCT images

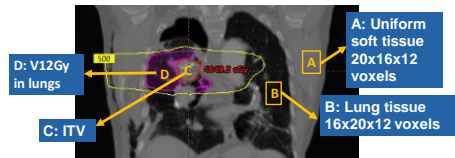


Data Harmonization of CT and CBCT: Workflow



Volumes of Interest (VOIs) Selection for Normalization

VOIs at the same locations on pCT and registered CBCT (VOI A & B: outside of 5 Gy isodose line → tissues in VOIs unchanged by radiation)



Wilcoxon signed rank test ($p < .05$) and Bonferroni correction (DOF=55) were used to test each pair of feature and to identify number of variable features before and after each harmonization

Feature Variability Reduction on ITV

# of variable features before harmonization	Features	VOI: Soft tissue (A)	VOI: Lung tissue (B)	VOI: Average of A & B	VOI: ITV
1	Gray level histogram	0%	0%	0%	-100%
4	Gray level co-occurrence	25%	-200%	50%	50%
7	Gray level run length	85.7%	-14.2%	50%	42.9%
8	Gray level size zone	37.5%	-12.5%	12.5%	37.5%
0	Neighborhood gray tone difference	0%	-100%	0%	-100%

For radiomics calculation in ITV, harmonization with VOI in soft tissue or average of 2 is more effective in feature variability reduction

Feature Variability Reduction on Lung V12Gy

Features	# of variable features before harmonization	VOI: Soft tissue (A)	VOI: Lung tissue (B)	Average of A & B	VOI: V12Gy
Gray level histogram	1	100%	0%	100%	100%
Gray level co-occurrence	0	0%	-1400%	0%	0%
Gray level run length	2	0%	-100%	50%	100%
Gray level size zone	1	0%	-200%	0%	0%
Neighborhood gray tone difference	0	0%	-200%	0%	-200%

For radiomics calculation in V12 Gy in lungs, harmonization with average of 2 VOIs provided the highest efficiency of feature variability reduction.

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Summary

- Radiomics features are useful for treatment assessment
- Radiomics features are affected by motion, noise and data variability
- Quantification of radiomics features are important for their clinical applications



Thank you for your attention

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- Q1 Which of following is not a radiomics feature?
- a. Grey-level co-occurrence matrix correlation
 - b. Grey-level co-occurrence matrix energy
 - c. Grey-level co-occurrence matrix homogeneity
 - d. Morphology
 - e. Mean dose

Answer: e

Ref: Sarah A. Mattonen et al, Detection of Local Cancer Recurrence After Stereotactic Ablative Radiation Therapy for Lung Cancer: Physician Performance Versus Radiomic Assessment. Int J Radiation Oncol Biol Phys, Vol. 94, No. 5, pp. 1121-1128, 2016
<http://dx.doi.org/10.1016/j.ijrobp.2015.12.369>
