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Radiomic hypotesis

Molecular heterogeneity of cancer lesions is cause of different clinical outcome.

Such heterogeneity can be captured, *in vivo*, on the entire lesion volume, by high-throughput quantitative radiomics descriptors from 3D image of cancer lesion.

Different expression level of a signature of radiomic features are able to predict different prognosis or treatment response of patients with similar cancer diagnosis (statistical analysis and predictive models).

Radiology

Radiomics: Images Are More than Pictures, They Are Data¹

Robert J. Gillies, PhD Paul E. Kinahan, PhD Hedvig Hricak, MD, PhD, Dr(hc)

In the past decade, the field of medical image analysis has grown exponentially, with an increased number of pattern recognition tools and an increase in data set sizes. These advances have facilitated the development of processes for high-throughput extraction of quantitative features that result in the conversion of images into mineable data and the subsequent analysis of these data for decision support; this practice is termed radiomics. This is in contrast to the traditional practice of treating medical images as pictures intended solely for visual interpretation. Radiomic data contain first-, second-, and higher-order statistics. These data are combined with other patient data and are mined

Radiomics: a new approach for the study of cancer



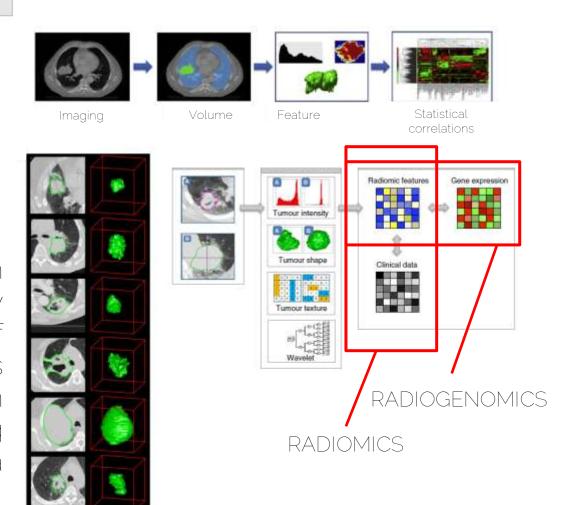
Published in final edited form as: Eur J Can et 2012 March 1 45 4): 441–446. doi:10.1016/j.ejca.2011.11.036.

Radiomics: Extracting more information from medical images using advanced feature analysis

Philippe Lambin^{a,*,c,f}, Emmanuel Rios-Velazquez^{a,c}, Ralph Leijenaar^{a,c}, Sara Carvalho^{a,c}, Ruud G.P.M. van Stiphout^{a,e}, Patrick Granton^{a,e}, Catharina M.L. Zegers^{a,e}, Robert Gillies^{b,e}, Ronald Boellard^{c,c}, André Dekker^{a,c}, and Hugo J.W.L. Aerts^{a,d,c}

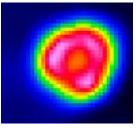
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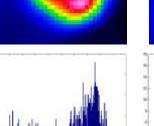
Comprehensive quantification of disease phenotypes by applying a large number of quantitative image features representing lesion heterogeneity and correlating with omics and clinical data

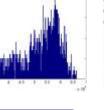


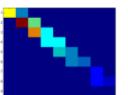
Texture and shape features

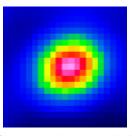
Feature	Description		Examples
Texture- First order	Grey level frequency distribution from histogram Analysis	Global	Minimum, mean and maximum intensity
Texture-First order Texture-Second order Texture-Third order	Histogram Anatysis		Standard deviation
			Skewness
			Kurtosis
			Percentile values
			Range of intensities
Texture-	Standard deviation Skewness Kurtosis Percentile values Range of intensities From spatial grey level dependence matrices Local (SGLDM) or co-occurrence matrices They express how often a pixel of intensity i finds itself within a certain relationship to another pixel of intensity j Percentile values Range of intensities Entropy Energy Contrast Homogeneity Dissimilarity Uniformity Correlation Percentile values Range of intensities Entropy Energy Contrast Homogeneity Dissimilarity Uniformity Correlation Percentile values Range of intensity Energy Romander Homogeneity Dissimilarity Uniformity Correlation Romander From neighbourhood grey-tone difference matrices (NGTDMs) From voxel alignment matrices Regional Run-length and emphasi Run-length variability From grey level size zone matrices Regional They reflect regional intensity variations or the distribution of homogeneous regions	Entropy	
Second		Local	Energy
order	They express how often a pixel of intensity i		Contrast
Texture- Second order	finds itself within a certain relationship to		Homogeneity
	another pixet of intensity j		Dissimilarity
			Uniformity
			Correlation
Taytura-		Local	Coarseness
			Contrast
			Busyness
			Complexity
	From spatial grey level dependence matrices Local (SGLDM) or co-occurrence matrices Entropy They express how often a pixel of intensity i finds itself within a certain relationship to another pixel of intensity j From neighbourhood grey-tone difference matrices (NGTDMs) From voxel alignment matrices From grey level size zone matrices They reflect regional intensity variations or the distribution of homogeneous regions Range of intensity interpret. Entropy Energy Contrast Homogeneit Homogeneit Coarseness Contrast Busyness Complexity Run-length a Run-length a Size-zone value of intensity variations or the distribution of homogeneous regions Spericity Compactness Eccentricity	Run-length and emphasis	
From v	Trom voxet dag. more madrood	rtogioriat	Run-length variability
	From arev level size zone matrices	Regional	Zone emphasis
	- ·	J	Size-zone variability
Shape and			Spericity
			Compactness
			Eccentricity
			Surface Area
			Sperical Disproportion
			Surface to Volume ratio
			Solidity

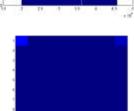


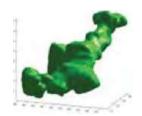


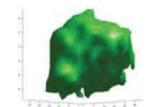


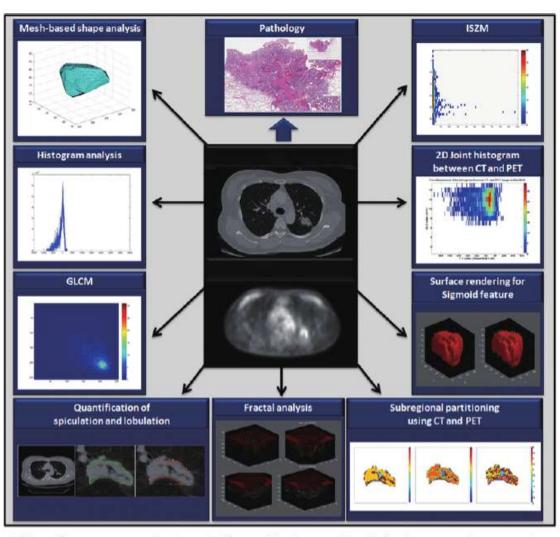












https://www.researchgate.net/figure/Various-radiomic-features-such-as-mesh-based-shape-histogram-gray-level-co-occurrence_fig3_315902486

Morphological features

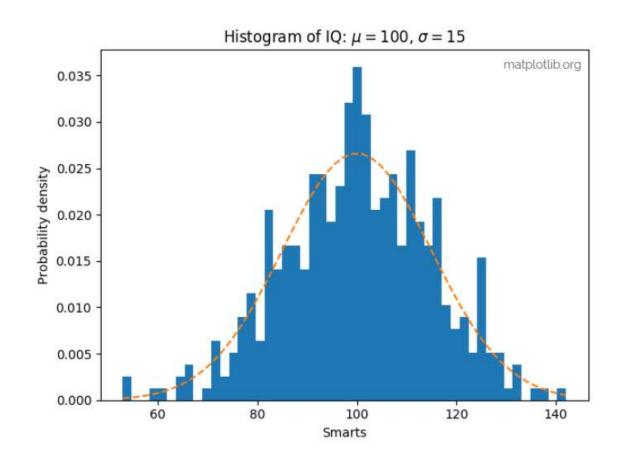
- 1. Metabolic Target Volume (MTV)
- 2. Surface
- 3. Spherical disproportion (ratio between measured surface of the lesion and surface of an equivalentsphere in terms of volume)
- 4. Sphericity
- 5. Surface-to-volume ratio

Normal	Cancer	
000		Large, variably shaped nuclei
090		Many dividing cells;
927		Disorganized arrangement
		Variation in size and shape
		Loss of normal features

http://sphweb.bumc.bu.edu/otlt/MPH-Modules/PH/PH709_Cancer/PH709_Cancer7.html

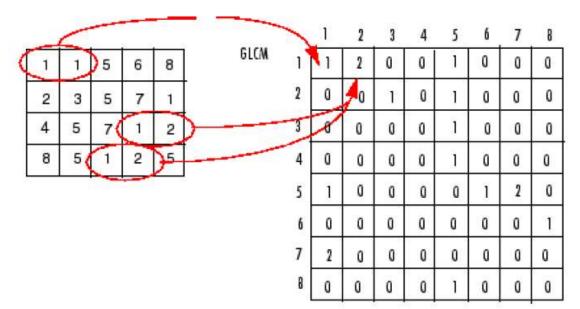
Histogram-based features

- 1. Maximum
- 2. Minimum
- 3. Mean
- 4. Median
- 5. Mean Absolute Deviation (MAD)
- 6. Root Mean Square (RMS)
- 7. Energy
- 8. Entropy
- 9. Kurtosis
- 10. Skewness
- 11. Standard Deviation
- 12. Uniformity
- 13. Variance



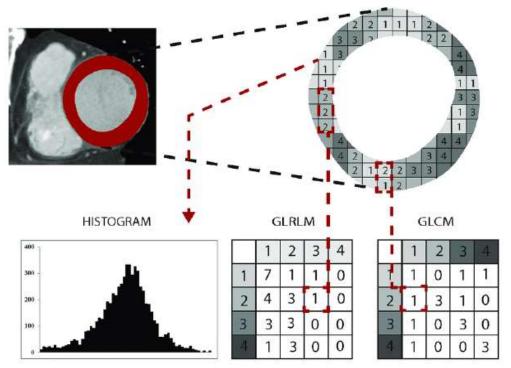
Texture descriptors Gray-Level Co-occurrence Matrix (GLCM)*

- 1. Energy
- 2. Contrast
- 3. Entropy
- 4. Homogeneity
- 5. Correlation
- 6. Sum Average
- 7. Variance
- 8. Dissimilarity
- 9. Auto Correlation
- * A Gray Level Co-occurrence Matrix (GLCM) quantifies the number of times the combination of levels X and Y occur in two pixels in the image that are separated by a distance of D pixels along angle A.



mathworks.com

Texture descriptors Gray-Level Run Length Matrix (GLRLM)*



https://www.researchgate.net/figure/Principles-of-generating-texture-analysis-features-Principles-of-generating-the_fig2_320821651

*A Gray Level Run Length Matrix (GLRLM) quantifies gray level runs, which are defined as the length in number of pixels, of consecutive pixels that have the same gray level value. It describes # runs with gray level G and length L that occur in the image along angle A.

Texture descriptors Gray-Level Size Zone Matrix (GLSZM)

1	2	3	4
1	3	4	4
3	2	2	2
4	1	4	1

Level	Size zone, s		
g	1	2	3
1	2	1	0
2	1	0	1
3	0	0	1
4	2	0	1

http://thibault.biz/Research/ThibaultMatrices/GLSZM/GLSZM.html

* A gray level zone is defined as a the number of connected voxels that share the same gray level intensity.

Contrary to GLCM and GLRLM, the GLSZM is rotation independent, with only one matrix calculated for all directions in the ROI

|Radiomi<u>cs</u>|

Texture descriptors Neighbouring Gray Tone Difference Matrix (NGTDM)*

* A Neighbouring Gray Tone Difference Matrix quantifies the difference between a gray value and the average gray value of its neighbours within distance D

PREPROCESSING	FEATURE EXTRACTION	FEATURE SELECTION	ON	TRAINING AND INTERNAL TESTING	
EXTRACT THE FOL	LOWING FEATURE FAMILIE	S			
MORPHOLOGICAL	FEATURES	TEXTURE-E	BASED	FEATURES	
✓ Morphology (25 f	features 3D or 10 features 2D)	✓ Grey-lev	el co-c	occurrence matrix (25 features)	
INTENSITY-BASED	FEATURES	✓ Grey-lev	el run l	length matrix (16 features)	
✓ Intensity-based s	statistic (18 features)	✓ Grey-lev	el size	zone matrix (15 features)	
✓ Intensity histogra	am features (16 features)	✓ Neighbo	ourhood	d grey tone difference matrix (5 features)	
		✓ Neighbo	ouring g	grey level dependence matrix (16 features	5)
APPLY THE FOLLO	WING TRANSFORMATION F	FILTERS FOR FILTER	-BASE	D FEATURES (111 features for each filte	er)
Wavelet - not rec	commended for 2D modalities	it will greatly increase	the re	quired time (minutes instead of seconds	for each image)
Square					
Squareroot					
Logarithm					
Exponential					
Gradient					
Laplacian of gaus	ssian (LoG)				
Local binary patte	erns (LBP) - the use of this filte	er will greatly increase	the req	quired time (minutes instead of seconds fo	or each image)

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Exploring feature-based approaches in PET images for predicting cancer treatment outcomes

El Naqa, Ph.D.³, P. Grigsby, M.D.³, A. Apte, M.Sc³, E. Kidd, M.D.³, E. Donnelly, M.D.³, D. Khullar, M.Sc³, S. Chaudhari, B.Sc³, D. Yang, Ph.D.³, M. Schmitt, B.Sc⁵, Richard Laforest, Ph.D.⁵, W. Thorstad, M.D.³, and J. O. Deasy, Ph.D.⁵

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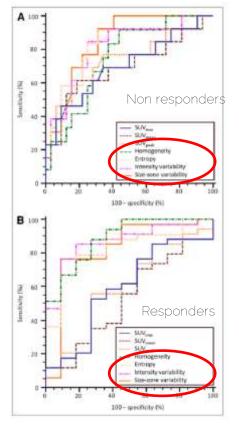


Table 2

Association between different extracted features and overall survival in a cohort of 9 head and neck patients measured by Spearman's rank correlation (rs) and the area under the ROC curve (AUC).

Variable		Spearman (13)	AUC
Tumor volum	W.	0,6928	9.8750
	Maximum	9.3464	0.7000
	Minimum	-0.2642	0.6000
	Mean	0.1752	0.6500
	Standard deviation	0.3464	0.6750
	110	0.1732	0.7000
	190	0.0	0.5000
	$I_{10.90}$	0.2598	0.6750
IVH Intensity-volume metrics	V ₁₀	-0.1732	0.5750
	V _{AB}	-0.7794	0.9500
VH Intensity-volume metrics	V ₁₀₋₉₀	0.0866	0.5000
	Energy	0.0866	0.5000
T	Contrast	-0.5196	0.8000
Texture-based features	Local homogeneity	0.5196	0.8250
	Entropy	-0.1732	0.5250
	Eccentricity	0.2598	0.6500
	Euler Number	0.6766	0.8500
Shape-based features	Solidin	-0.6088	0.850
	Extent	-0.6062	0.8500

J Nucl Med 2011; 52:369-378

Intratumor Heterogeneity Characterized by Textural Features on Baseline ¹⁸F-FDG PET Images Predicts Response to Concomitant Radiochemotherapy in Esophageal Cancer

Florent Tixier¹, Catherine Cheze Le Rest^{1,2}, Mathieu Hatt¹, Nidal Albarghach^{1,3}, Olivier Pradier^{1,3}, Jean-Philippe Metges^{3,4}, Laurent Corcos⁴, and Dimitris Visvikis¹

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A study in which we hope not to be cited...



RESEARCH ARTICLE

False Discovery Rates in PET and CT Studies with Texture Features: A Systematic Review

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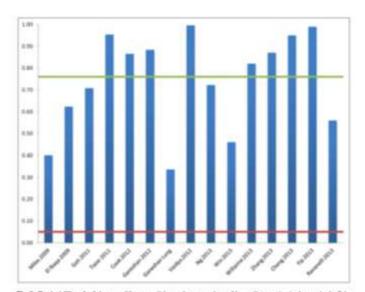


Fig 2. Probability of a false positive result based on number of hypotheses tested per study (blue columns) for all study categories. 5% type-I error probability = red line, average type-I error probability (76%) over all studies = green line (Note—additional infletion of the type-I error probability due to the use of the optimum out-off approach is not included here).

doi:10.1371/journal.pone.0124165.g002

Key methodological issues

- Repeatability, the closeness of the agreement between the results of successive radiomic measurements under the same conditions of measurement
- Riproducibility, the closensess of the agreement between the results of radiomic measurement under similar conditions of measurements
- Significance, the ability of radiomic in effectively characterizing cancer lesion heterogeneity

Stability