

ARTIFICIAL INTELLIGENCE IN LUNG CANCER: USING RADIOMICS TO PREDICT TUMOR RECURRENCE

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Non-Small Cell Lung Cancer

- Survival rates remain quite poor despite advances in diagnosis and treatment.
- ► 5-year survival rates:
 - > Stage I: 55-75%
 - > Stage II: 40-50%
 - > Stage III: 5-35%
 - > Stage IV: <5%



Lang-Lazdunski, L. et al. Eur Resp Soc. (2013) 382-404. SEER Stat Fact Sheets: Lung and Bronchus Cancer.

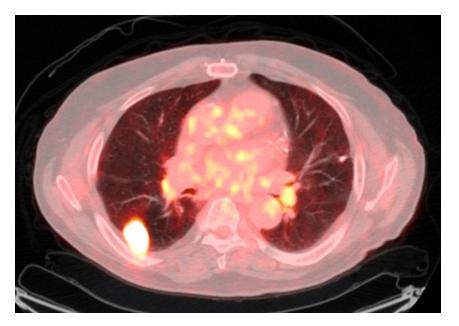


The Role of Imaging

Computed Tomography (CT)



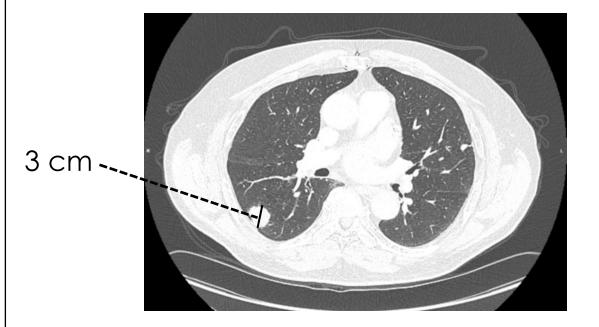
Positron Emission Tomography (PET)



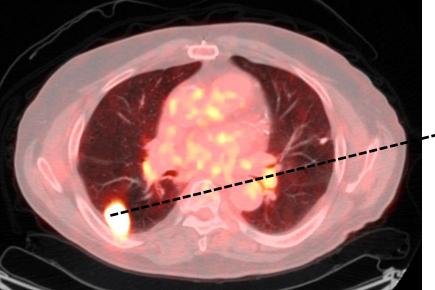


The Role of Imaging

Computed Tomography (CT)



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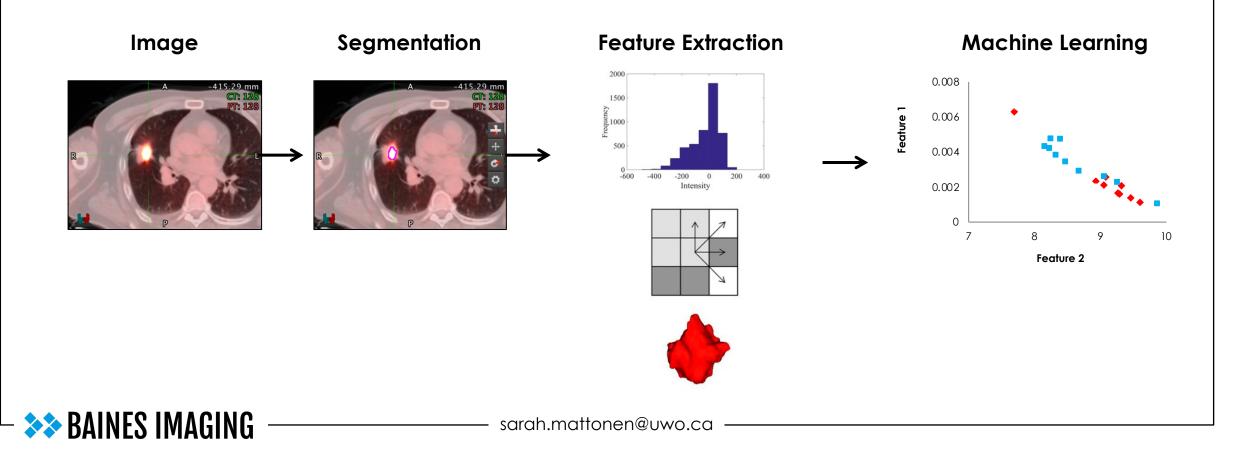


SUVmax = 9



Radiomics

 Radiomics aims to extract more complex quantitative information (e.g., texture) from standard medical images.



Positron Emission Tomography (PET)

 SUVmax has been shown to predict a higher risk of recurrence or death in NSCLC.

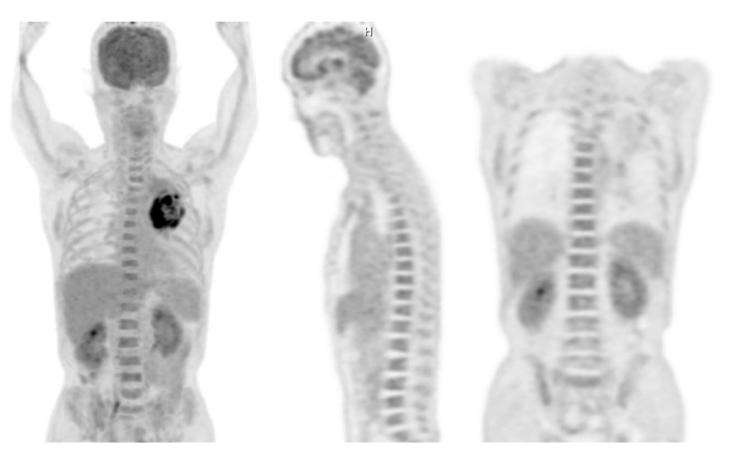


Travis WD, et al. J of Thor Onc. (2011) 6:244-285. Kadota K, et al. J of Thor Onc. (2015) 10:806-814. Lee et al., European Radiology (2017) 27:1912–1921.



Positron Emission Tomography (PET)

- Tumour invasion from the main tumour mass.
- Dissemination of disease throughout the body.



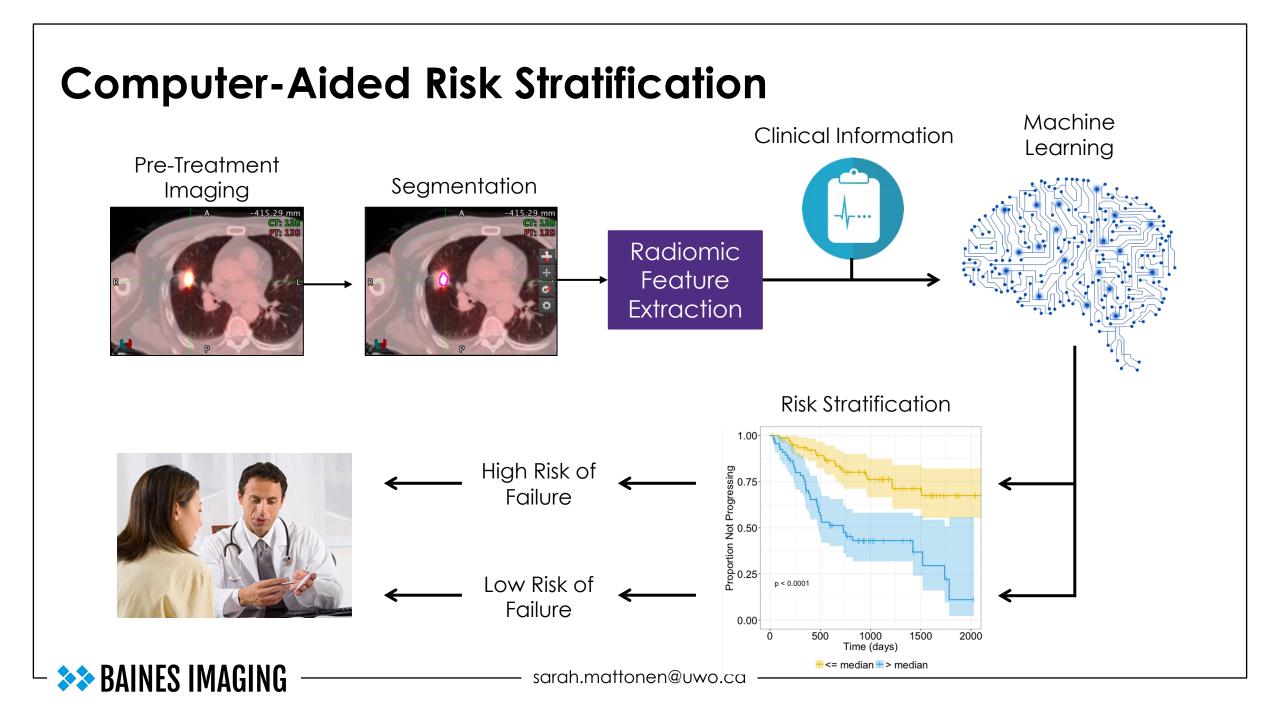
Travis WD, et al. J of Thor Onc. (2011) 6:244-285. Kadota K, et al. J of Thor Onc. (2015) 10:806-814. Lee et al., European Radiology (2017) 27:1912–1921.



Objective

To develop a software system integrating PET imaging and non-imaging biomarkers to improve lung cancer prognosis and risk stratification.





Materials

- ► Training Cohort (n = 145):
 - > Selected from two local medical centers.
 - > All patients had pre-operative PET/CT performed prior to surgery.
 - Feature selection and model development

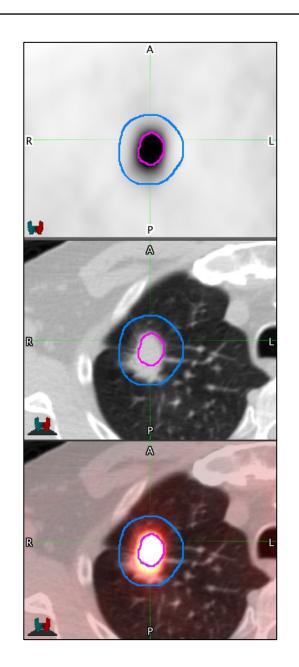
► Testing Cohort (n = 146):

- > Selected from three local medical centers.
- Underwent PET/CT imaging prior to definitive treatment as part of observational biomarker study.
 - Model evaluation

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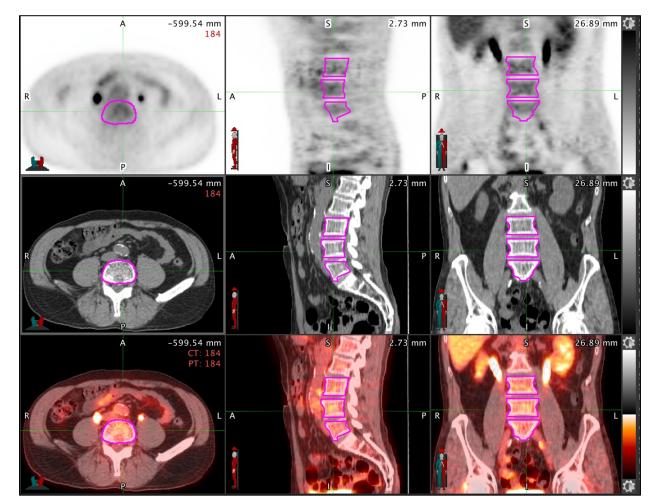
Segmentation: Tumour

- The metabolic tumour volume (MTV) was segmented on the PET image.
- A 3-dimensional penumbra region was also generated surrounding the MTV to sample surrounding uptake.
- ► Three regions were evaluated:
 - > MTV only
 - > Penumbra only (excluding the MTV)
 - > MTV plus penumbra





Methods: Bone Marrow Segmentation

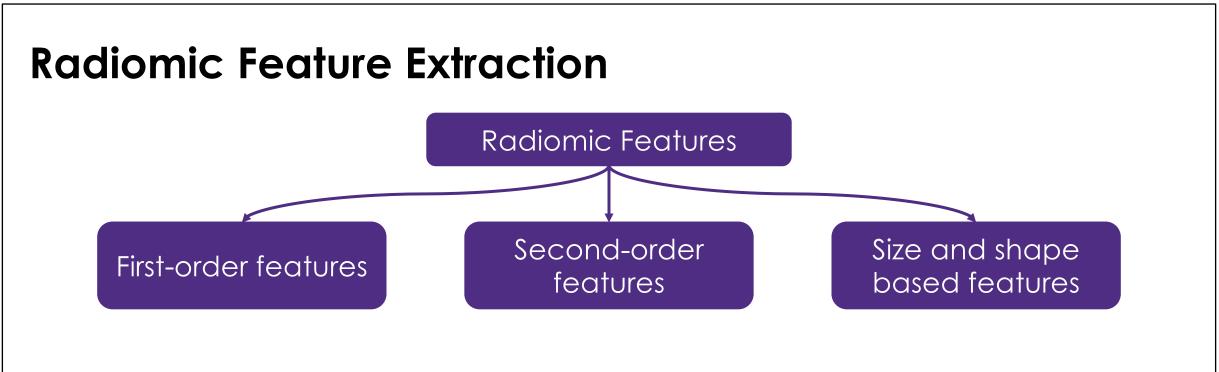


MIM (MIM Software Inc., Cleveland, OH).



sarah.mattonen@uwo.ca

Mattonen, et al. Radiology: 293(2), 451-459, 2019.



A total of 668 radiomics features were extracted from the volumes of interest.

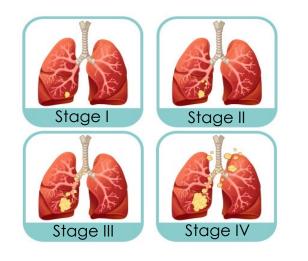
Echegaray, S., et al. (2017) J Digit Imaging, 1-12.

GitHub: riipl/3d_qifp



Methods: Model Training

► Top predictive features were selected using randomizations of 4-fold cross-validation of LASSO Cox regression.



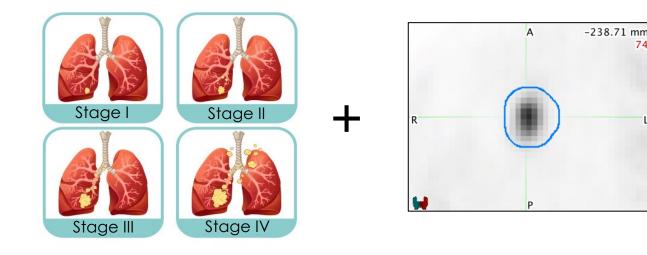
Clinical

- **>>** BAINES IMAGING

Methods: Model Training

Top predictive features were selected using randomizations of 4-fold cross-validation of LASSO Cox regression.

74



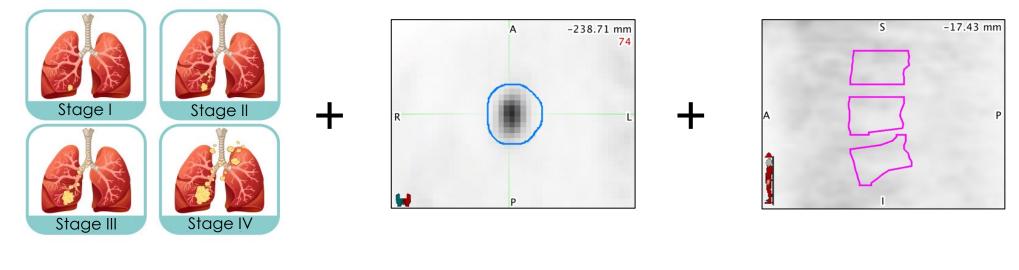
Tumour Plus Penumbra



Clinical

Methods: Model Training

► Top predictive features were selected using randomizations of 4-fold cross-validation of LASSO Cox regression.



Clinical

Tumour Plus Penumbra

Bone Marrow



Results: Multivariate Model Clinical + Tumour + Bone Marrow

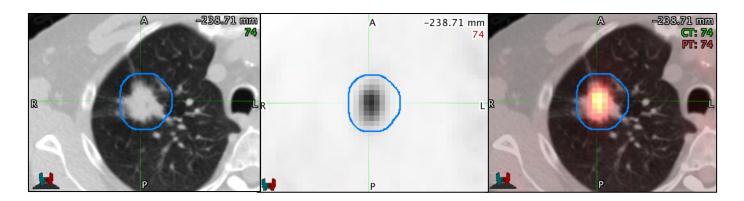
Feature Type	Feature	HR [95% CI]	p-value
Clinical	Stage	1.98 [1.45-2.70]	p<0.001*
Blood	WBC (1000/uL)	0.99 [0.88-1.11]	p=0.81
	Hemoglobin (g/dL)	0.99 [0.82-1.20]	p=0.93
	Platelets (1000/uL)	1.00 [1.00-1.01]	p=0.93
Tumor	MTV Plus Penumbra GLCM Energy (MAD)	0.69 [0.40-1.19]	p=0.18
	Penumbra GLCM Entropy (IQR)	1.35 [0.97-1.86]	p=0.07
	Penumbra GLCM Cluster Shade (Max)	1.17 [0.84-1.63]	p=0.36
Bone Marrow	GLCM Sum Mean (Skewness)	0.52 [0.32-0.84]	p=0.008*
	GLCM Cluster Tendency (Skewness)	1.62 [1.02-2.59]	p=0.04*



Results: Risk Stratification Testing Cohort Training Cohort (B) (A) 1.00 1.00 0.75 0.75-Disease Free Survival Disease Free Survival 0.25 0.25 p<0.001 p<0.001 0.00 0.00 1500 2000 1000 500 1000 2500 500 1500 2000 0 0 Time (days) Time (days) + <= median 🕂 > median <= median 🕂 > median +Mattonen, et al. Radiology, 2019. **SAINES IMAGING**

Qualitative Results

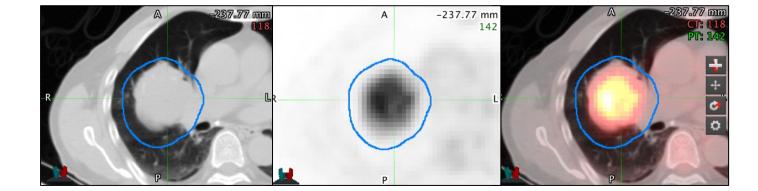
(A)



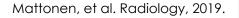
 $SUV_{max} = 10.3$ Stage I High-Risk Radiomics Recurrence



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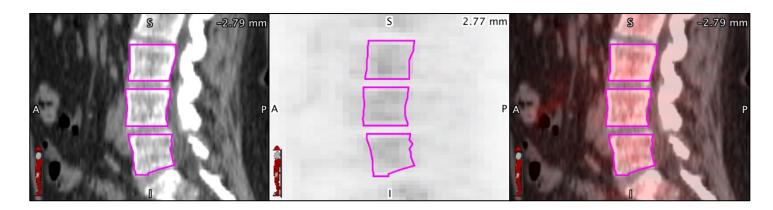


SUV_{max} = 10.1 Stage II Low-Risk Radiomics **No Recurrence**



Qualitative Results

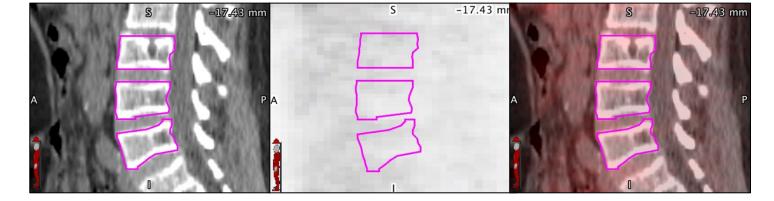
(A)



Stage I High-Risk Radiomics **Recurrence**

(B)

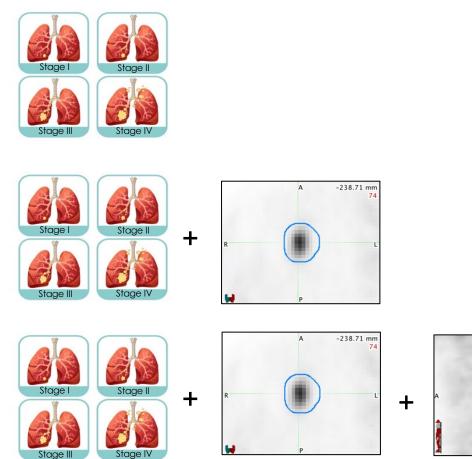
Solution BAINES IMAGING



Stage III Low-Risk Radiomics **No Recurrence**

Mattonen, et al. Radiology, 2019.

Results: Summary



Concordance = 0.69 [0.60-0.77]

Concordance = 0.75 [0.67-0.82]

Concordance = 0.78 [0.70-0.85]



-17.43 mm

CAN ADDING CT FEATURES IMPROVE PERFORMANCE?

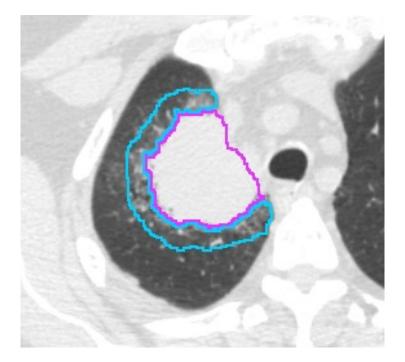
Jaryd Christie, CAMPEP PhD Candidate





Tumor and Peri-tumoral CT Segmentation

MATLAB based-GUI for semi-automatic tumor segmentation on CT

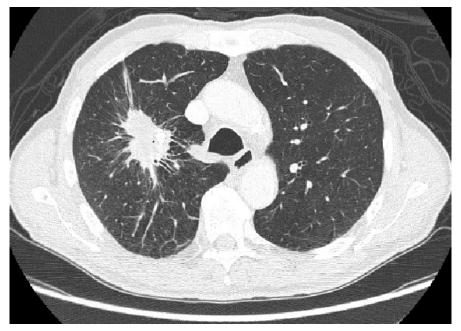


Christie, Jaryd R., et al. Vol. 12036. SPIE, 2022. <u>https://github.com/baines-imaging-mattonen-lab/CT-Lung-Tumour-Segmentation</u>

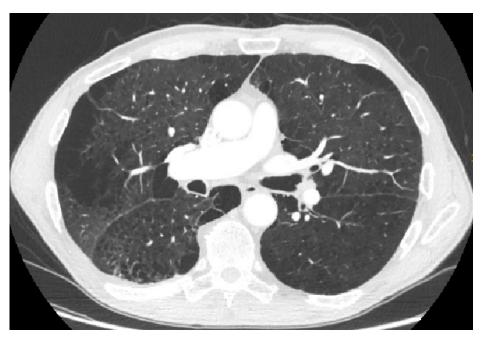


Qualitative Features

- ► Tumour features that describe the location and geometry
- ► Features which characterize the lung tissue, bronchi, and lumen



Spiculated



Severe Emphysema



Results: Feature Selection

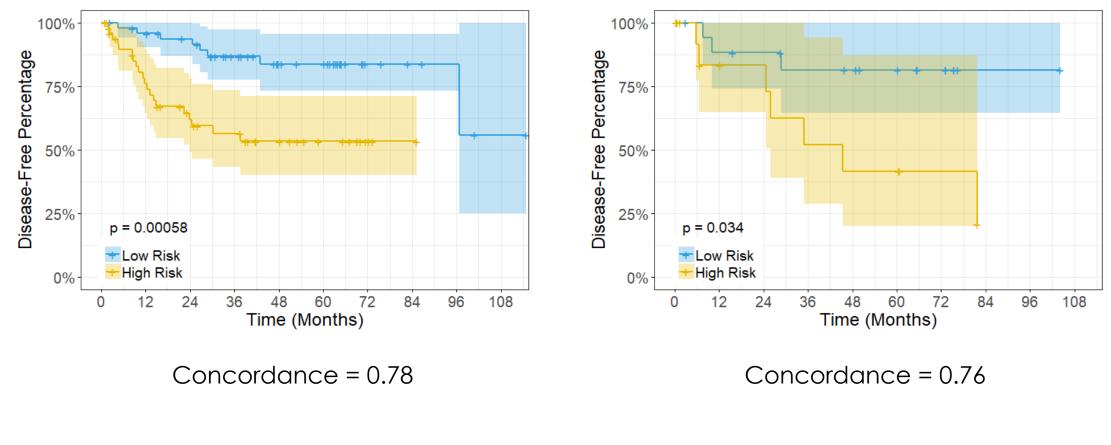
- Seven selected features:
 - > One clinical feature
 - Stage
 - > Six radiomic features (3 texture, 3 first-order)
 - Three CT (2 Tumour, 1 Peritumoural)
 - Three PET (2 Peritumoural, 1 Bone Marrow)

Results: Model Evaluation

- Training: Stage vs Radiomics + Stage
 - > **Concordance:** 0.67 [95% CI: 0.58 0.76] vs 0.78 [95% CI: 0.70-0.86]
 - > p < 0.005
- Testing: Stage vs Radiomics + Stage
 - > **Concordance:** 0.60 [95% CI: 0.48 0.74] vs 0.76 [95% CI: 0.59-0.87]
 - > **p = 0.008**
- Radiomics model significantly stratified patients into high- and lowrisk of recurrence

Results: Risk Stratification

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Testing, n=34

Training, n=101

Conclusions

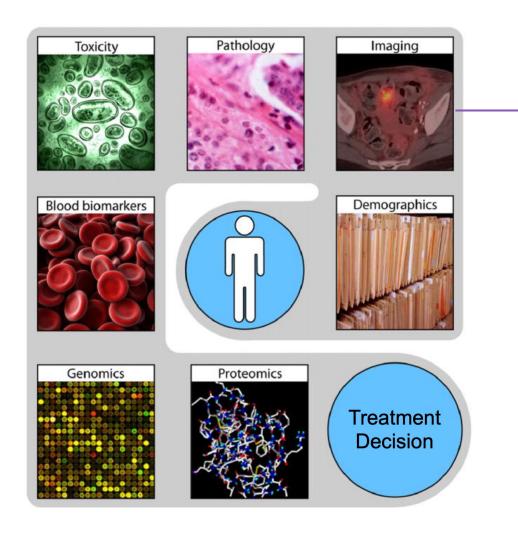
- These radiomics based tools have the potential to identify NSCLC patients at a higher risk of recurrence and may add clinical utility for risk stratification.
- This assist physicians in distinguishing patients who may benefit from adjuvant or more aggressive personalized treatment options.

Next Steps

- Collaborations for external validation of models.
- Implementation of standardized radiomics features and opensource software



Translational Cancer Imaging



Computer-Aided Decision Support



Improve Patient Outcomes





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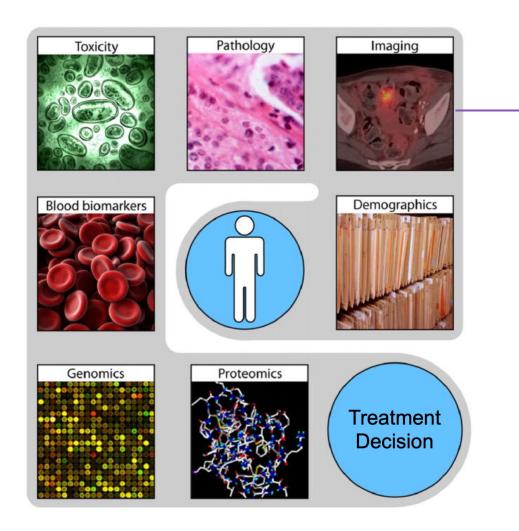
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Translational Cancer Imaging



Computer-Aided Decision Support



Improve Patient Outcomes



