## Cancer Data Science

Thursday, July 24th, 2025



### **Background: Breast Cancer**

- Very prevalent cancer
- High burden of disease
- Patients can experience recurrence or death
- Many variables can impact outcomes

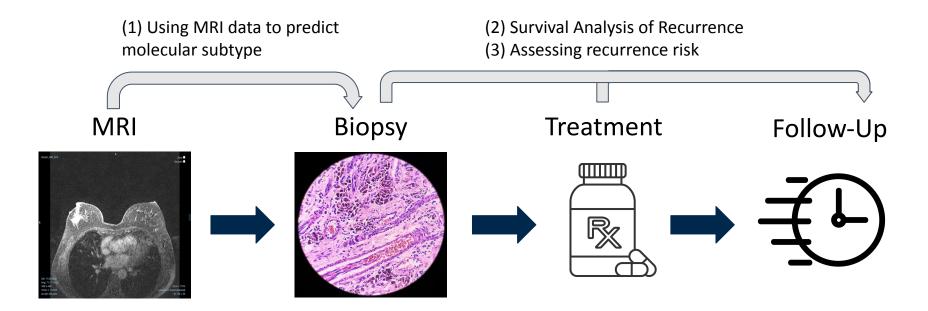


24,240 will **die** from Breast Cancer in 2025

Source: <a href="https://www.nationalbreastcancer.org/breast-cancer-facts/">https://www.nationalbreastcancer.org/breast-cancer-facts/</a>



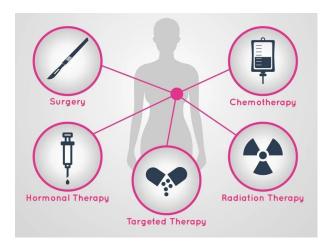
#### **The Breast Cancer Care Process**



### **Nuances and Acknowledgements**

 Our dataset and projects represent only a small snapshot of the journey patients go through.

- Treatment is not a "one size fits all"
  - Neoadjuvant treatments: before surgery
  - Adjuvant treatments: after surgery
  - Many factors go into deciding treatment such as cancer type, clinician expertise, and patient preferences.



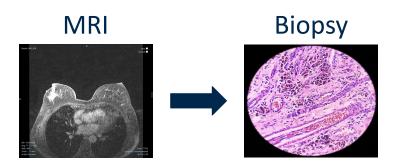
Source: <a href="https://nhcancerclinics.com/cancer-types/breast-cancer/">https://nhcancerclinics.com/cancer-types/breast-cancer/</a>

## Background: Dynamic contrast-enhanced MRIs of breast cancer patients

- The Cancer Imaging Archive: Duke Breast Cancer MRI
- Dataset: "single-institutional, retrospective collection of 922 biopsy-confirmed invasive breast cancer patients, over a decade"
- Combination of 96 clinical and 529 imaging features

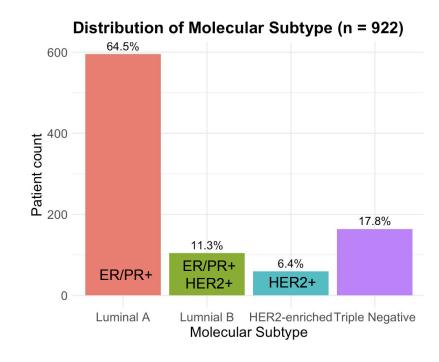
# Using radiomics features to predict molecular subtype of breast cancer

Albert Kang, Lucy Malmud, Ritish Natesan, Kate Podrebarac



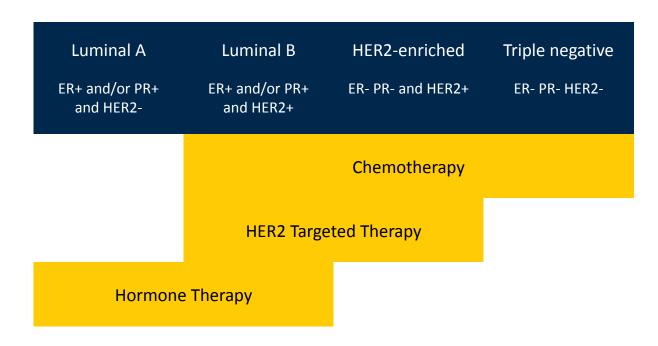
### **Background: Molecular Subtype**

- Combination of Estrogen Receptor (ER), Progesterone Receptor (PR), and HER2 status
- Provides deeper insights into tumor pathology and progression than from histology
- Crucial for accurate diagnosis and precision treatment



## **Background: Clinical Importance**

- Decrease need for invasive biopsy procedures
- Decrease cost of care
- Faster treatment decisions





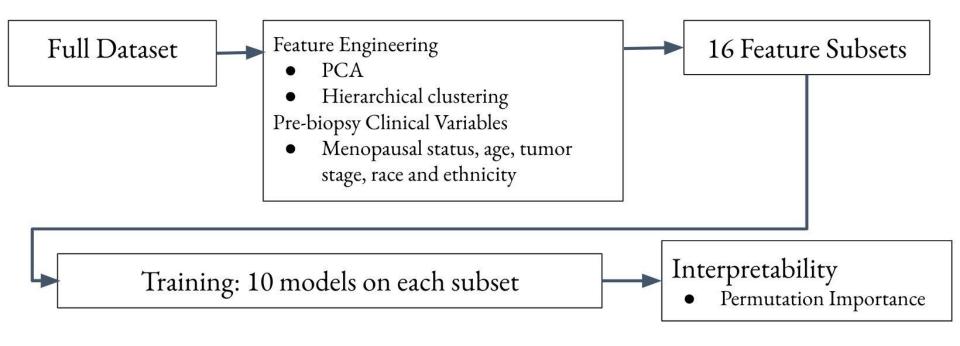
## **Background: Reference Paper + Dataset**

- Radiomics = the field of study concerned with extracting large amounts of quantitative data from images
  - 529 radiomics features partitioned into 10 feature groups
- In a 2018 study<sup>1</sup> by Saha and colleagues, used random forest model to predict molecular subtype
- Can we do better?

Target	Reference AUC
ER	0.649
PR	0.622
HER2	0.500

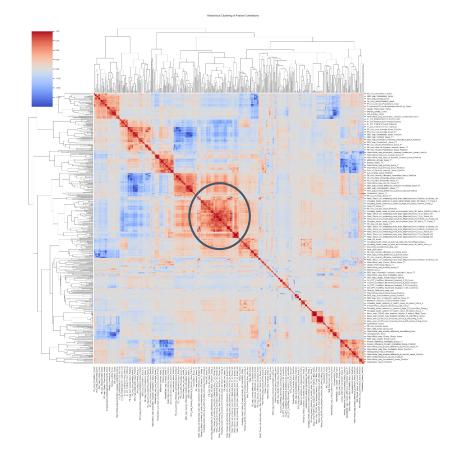
<sup>&</sup>lt;sup>1</sup>A Machine Learning Approach to Radiogenomics of Breast Cancer: a study of 922 subjects and 529 DCF-MRI features

#### **Our Process**



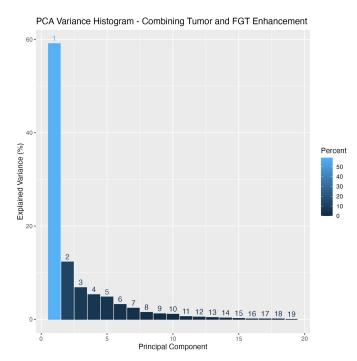
## Feature Engineering: Hierarchical Clustering

- Correlation-based hierarchical clustering
- "Slice" the dendrogram at a level corresponding to correlation > .90
- Select one feature from each of these groups
- 529  $\rightarrow$  251 covariates



## **Feature Engineering:**Principal Component Analysis (PCA)

- Transforms high dimensional data into a new coordinate system
- First principal component explains the most variation in the data
- Successive components explain less variation
- Each covariate has one loading factor per component
  - Represents how much that covariate contributes to the component



## **Training:** Models

## **Generalized Linear Models:**

- Logistic Regression
- LASSO
- Elastic Net

#### **Clustering:**

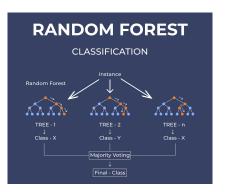
- K-Means
- Gaussian Mixture Model

#### **Machine Learning Models:**

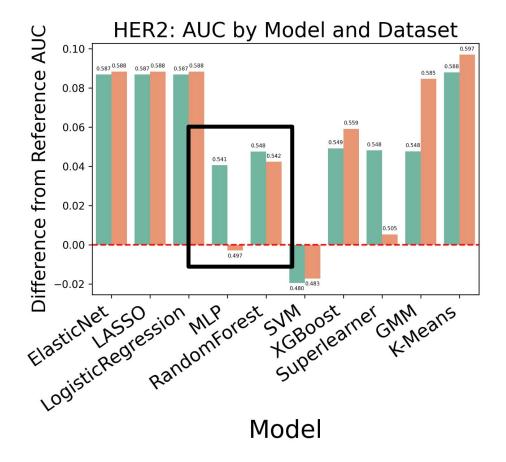
- Support Vector Machine (SVM)
- Multilayer Perceptron (MLP)

#### **Ensemble Methods:**

- Random Forest
- XGBoost
- Superlearner
  - Metalearner: Gradient Boosting Classifier
  - Base Learners: Random Forest, Decision Tree, Logistic Regression, Gradient Boosting Classifier



Source: https://diagramskiniatinguiccnf.z21.web.core.windows.net/random-forest-model-diagram.html

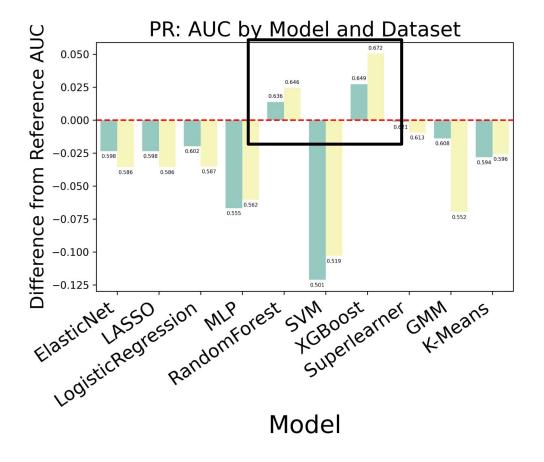


#### **Feature Subsets**

Top Three Principal
Components Per
Feature Group
Raw Data Values
from Top Three Covariates
in PC1

PCA scores yield better results than raw data values.

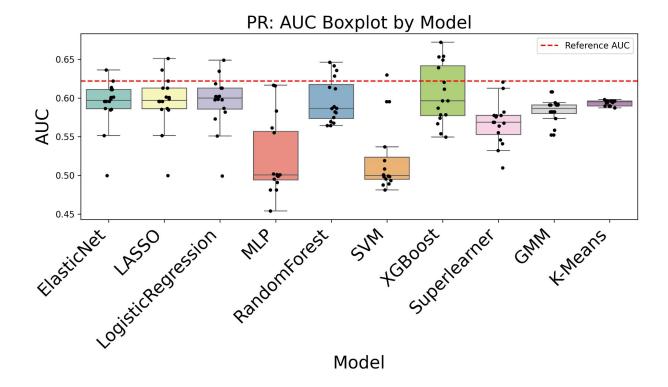
Improvement in performance by capturing more information in the same number of predictors.



**Feature Subsets** 

Unengineered Data
Unengineered Data
+ Pre-Biopsy Clinical

Adding pre-biopsy clinical features improves performance as compared to the original imaging data in some cases.



- SVM and MLP consistently perform poorly
- Linear models
   (ElasticNet, LASSO,
   LogisticRegression)
   comparable
   performance
- XGBoost,
   RandomForest
   consistently best

#### **Feature Subsets**

						PR: A	UC H	eatm	ap b	у Мос	del ar	nd Da	taset					_				
	ElasticNet	0.60	0.59	0.59	0.61	0.61	0.60	0.60	0.64	0.62	0.50	0.55	0.60	0.58	0.60	0.61	0.60		- (	0.66	1: All Imaging Data	9: Enough PCs to Explain 90%
	GMM	0.61	0.55	0.61	0.55	0.59	0.59	0.58	0.58	0.59	0.58	0.59	0.58	0.59	0.56	0.57	0.59				2: All Imaging + Pre-Biopsy Clinical	Variance per Feature Group
	S		0.55	0.01	1 0.55 0.59 0.59 0.58 0.58 0.59 0.58 0.59 0.58 0.59 0.58 0.59 0.56 0.57 0.59	- (	0.64	3: Hierarchical Clustering	10: Enough PCs t													
	K-Means	0.59	0.60		0.60	0.59	0.59	0.60	0.60	0.59	0.60	0.59	0.60	0.59	0.60	0.59	0.60			0.62	4: Hierarchical Clustering + Clinical	Variance per Feature Group + Clinical
	LASSO	0.60	0.59	0.59	0.61	0.61	0.60	0.60	0.64	0.62	0.50	0.55	0.60	0.58	0.60	0.65	0.60		ľ	0.02	5: Top PC\ Per Feature Group	11: Raw Data Val
del 1	LASSO LogisticRegression	0.60	0.59	0.58	0.61	0.61	0.60	0.60	0.63	0.62	0.50	0.55	0.60	0.57	0.60	0.65	0.60		- (	0.60 On	6: Top Three PCs Per Feature Group	Covaraiates in P 12: Raw Data Val from Top
Mod	MLP	0.56	0.56	0.50	0.62	0.62	0.50	0.50	0.58	0.50	0.50	0.50	0.50	0.49	0.48	0.45	0.48		- 0.58 ₹ - 0.56 - 0.54	0.58₹	7: Top Three PCs s Per Feature Group + Clinical	Covaraiate in PC1 + Clinical
	RandomForest	0.64	0.65	0.64	0.58	0.59	0.61	0.59	0.61	0.63	0.57	0.57	0.58	0.59	0.56	0.57	0.56			0.56	8: Top PC Per Feature Group + Clinical	13: Raw Data Val from Top Three C in PC1
	SVM	- 0.50	0.52	0.54	0.60	0.60	0.50	0.48	0.63	0.49	0.50	0.51	0.49	0.49	0.50	0.50	0.50			0.54	+ Cillical	14: Raw Data Val from Top Three C in PC1 + Clinical
	Superlearner	0.62	0.61	0.58	0.58	0.58	0.57	0.57	0.56	0.56	0.55	0.57	0.58	0.58	0.54	0.53	0.51	- 0.52	0.52		15: Raw Data Val	
	XGBoost	0.65	0.67	0.65	0.60	0.60	0.62	0.57	0.61	0.64	0.55	0.55	0.59	0.57	0.58	0.65	0.58			0.32		Explain 90% of Variance per Feature Group
		1	Ź	3	4	5	6	7 Fee	8 ature	9 Sub	10 set	11	12	13	14	15	16		- (	0.50		16: Raw Data Val from PCs to Explain 90% of Variance per Feature Group

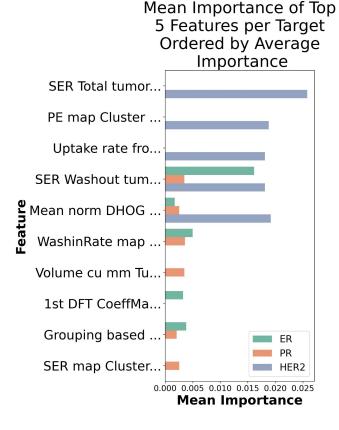
56	1: All Imaging Data	9: Enough PCs to Explain 90%
	2: All Imaging + Pre-Biopsy Clinical	Variance per Feature Group
54	3: Hierarchical Clustering	10: Enough PCs to Explain 90% of
52	4: Hierarchical Clustering + Clinical	Variance per Feature Group + Clinical
, ,	5: Top PC\ Per Feature Group	11: Raw Data Values from Top
50	6: Top Three	Covaraiates in PC1
2	PCs Per Feature Group	12: Raw Data Values from Top
58₹	7: Top Three PCs s Per Feature Group	Covaraiate in PC1 + Clinical
66	+ Clinical 8: Top PC Per	13: Raw Data Values from Top Three Covariates in PC1
00	Feature Group + Clinical	
54		14: Raw Data Values from Top Three Covariates in PC1 + Clinical
52		15: Raw Data Values from Enough PCs to Explain 90% of Variance per Feature Group
50		16: Raw Data Values from PCs to Explain 90% of

+ Clinical

### Results

Target	Best Model	Best Dataset	AUC*	Reference AUC		
ER	XGBoost	Hierarchical Clustering + Clinical	0.661	0.649		
PR	XGBoost	Unengineered + Clinical	0.672	0.622		
HER2	Random Forest	Raw Data Values from PCs to Explain 90% of Variance per Feature Group + Clinical	0.655	0.500		

Final model and dataset combination with highest \*mean AUC across 5-fold cross-validation.



### **Feature Importance**

- Permutation importance measures change in AUC when values of each covariate are permuted
- Insight into what covariates are most important in determining outcome

#### **Important Features**

Mean norm DHOG... (3)

Quantification of texture

SER Washout tumor... (3) + WashinRate map... (2)

Speed of contrast agent movement

#### Conclusion

#### Summary:

- Performing dimensionality reduction was able to increase prediction performance in some settings
- Adding pre-biopsy clinical predictors increased performance
- Training multiple models helped us find the best model for each target

#### Limitations:

- Imbalanced molecular subtype outcome
- Dataset is restricted to one hospital

#### Future:

Using raw MRI data for SVM and MLP

Source: https://www.svchs.com/blog/southwest-virginia-community-health-systems-recognizes-the-month-of-october-as-breast-cancer-awareness-month/



## Thank You!