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### Performance Analysis of Low and High-Grade Breast Tumors Using DCE MR Images and LASSO Feature Selection

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### Introduction

**Breast Cancer Facts - In** 2020, there were 2.3 million women diagnosed with breast cancer. (*WHO 2020*).

**Causes of Breast cancer**-Genetic, Environment, Early menstruation, and Late menopause.

**Breast Anatomy-** Parts-Ducts, Lobes, Lobules, Lymph node.

**DCE MRI for breast imaging** produces highresolution images for women at high risk of breast cancer and is also effective for evaluating dense female breasts

• Detect microscopic lesions in a (potentially) large volume of tissue. High temporal resolution while preserving high spatial resolution.



# Introduction- Grade Facts

- Grade Prognostic factor and aggressive potential
- The 3 factors for one of the scoring systems are (the Nottingham Histologic Score system)
  - amount of gland formation
  - nuclear features
  - mitotic activity



# Motivation

- Low grades (Grade I and Grade II) are less aggressive and show an avascular nature with less proliferation of tumors.
- High Grade is a more aggressive, highly intense, highly vascularized, and heterogenous large mass where necrotic, and apoptotic processes take place in the tumor.
- Needle biopsy may be a misinterpretation of the actual grade due to tumor heterogeneity.
- It is essential to ascertain suitable machine learning methods for differentiating low and high-grade breast tumors.

# Aim & Objectives

**Aim:** To analyze Radiomics-based low and high-grade DCE-MR breast tumor classification with a collection of classifiers using LASSO feature selection

### **Objectives:**

- Analysis of clinicopathological characteristics
- Feature selection by LASSO model
- classification of high-grade and low-grade tumors by using a collection of classifiers

# Materials & Methods

#### Dataset description

- A total of 638 patients included in our study where 431(67.55%) were low-grade and 207 (32.44%).
- A total of 529 features named tumor enhancement, shape, enhancement of tissues surrounding, texture, and shape were extracted from the segmented tumor

#### Feature Selection

• LASSO regression analysis techniques are frequently employed in feature selection and binary classification.

$$L_{lasso}(\hat{\beta}) = \sum_{i=1}^{n} \left( y_i - x_i' \hat{\beta} \right)^2 + \alpha \sum_{j=1}^{m} \left| \hat{\beta}_j \right|$$

• Pairwise Pearson Correlation Coefficient Matrix (PCCM) identified high-correlated feature pairs

# Materials & Methods

#### Classifiers

- Logistic regression (LR), k-nearest Neighbors (KNN), Linear discriminant analysis (LDA), Gaussian Naïve Bayes (GNB), Linear Support Vector Machines (LSVM), and Random Forest (RF) were implemented for the classification of Low and High grade
- The performance of different classification models was analyzed by using evaluation matrices such as Accuracy, Sensitivity, Area Under the receiver operating characteristic Curve (AUC), specificity, F1-score, Precision, Positive Predictive Value (PPV), and Negative Predictive Value (NPV).



Representative set of breast DCE MR Images of two different Highgrade patients acquired in the axial plane (a) one can appreciate highintensity tumor and (b) one cannot appreciate high-intensity tumor



Representative set of breast DCE MR Images of two different Low-grade patients acquired in the axial plane (a) one can appreciate moderate-intensity tumor and (b) one cannot appreciate moderate-intensity tumor.

### Clinicopathologic Characteristics

	Low grade	High grade	p-value		
No of subjects	431(67.55%)	207(32.44 %)			
Age(Mean±SD)	54.69±10.86	49.90 ± 11.61	0.6921		0.575 Average across the folds
Estrogen receptors status			<.00001	0.050 -	alpha: CV estimate
Positive	376(87.23%)	105(50.72%)			0.550 -
Negative	55(12.76%)	102(49.27%)		2 0.025 -	
Progesteron			<.00001		ē 0.525 -
receptor status					2 a saa
Positive	338(78%)	75(36.23%)			ā 0.500 -
Negative	93(21.57%)	132(63.76%)		-0.025 -	<b>D</b>
HER2 status			0.00239		⊆ 0.475
Positive	62(13.38%)	50(24.15%)			<u> </u>
Negative	369(85.61%)	157 (75.84%)		-0.075 -	0.450 -
Response status			<.00001	<u>ت</u>	and the second
PCR	9(2.08%)	37 (17.87%)		-0.100	0.425 -
Non-PCR	83(19.25%)	59 (28.50%)			
Not Available	332(77.03%)	103 (49.75%)		····· · · · · · · · · · · · · · · · ·	········
Others	7(1.62%)	9 (4.34%)		$10^{-2}$ $10^{-1}$ $10^{0}$ $10^{1}$ $10^{2}$	10 <sup>4</sup> 10 <sup>5</sup> 10 <sup>6</sup>
Menopausal Status			0.02629	alpha	alphas
Premenopausal	179(41.53%)	109(52.65%)		LASS	SO Analysis
Postmenopausal	241(55.91%)	95(45.89%)			
Not Available	11(2.55%)	3(1.44%)			
Bilateral status			0.00451		
Bilateral	25(5.80%)	2(0.96%)			
Non -Bilateral	406(94.66%)	205(99.03%)			



**Performance Analysis of Different Classifiers for Categorizing Low and High-Grade** 

Classifiers	Accuracy (%)	AUC	Sensitivity (%)	F1-score	Specificity (%)	Precision	NPV
LD	74.6	0.78	91.53	0.82	39.68	0.75	0.69
LR	75.6	0.76	92.30	0.83	41.26	0.77	0.72
GNB	73.6	0.74	90.76	0.82	38.09	0.75	0.67
L-SVM	77.9	0.79	96.15	0.86	39.52	0.82	0.58
C-KNN	73.6	0.70	91.53	0.82	36.50	0.74	0.67
RF	74.4	0.71	91.36	0.84	30.18	0.76	0.57

#### LASSO Selected Features

**Selected Features** 

Inf\_mea\_of\_corr2\_Tumor'

Grouping\_based\_proportion\_of\_tumor\_voxels\_3D\_tumor\_Group\_1

Mean\_norm\_DLBP\_max\_timepoint\_binsize\_256\_with\_filling\_Tumor

SER\_Total\_tumor\_vol\_cu\_mm WashinRate\_map\_information\_measure\_correlation2\_tumor WashinRate\_map\_inverse\_difference\_normalized\_tumor WashinRate\_map\_skewness\_tumor PE\_map\_information\_measure\_correlation2\_tissue\_PostCon



#### AUC for L-SVM Classifier

# Conclusions

- An experiment was conducted to classify breast tumor grades using different classifiers.
- LASSO feature selection method with optimal hyperparameter selection has selected 8 optimal features for the evaluation process.
- The clinical and histopathological characteristics tabulation revealed highly significant differences between the clinical parameters and tumor grades.
- For the feature's multi-collinearity identification, a Pearson Correlation Heat Map has been generated.
- Lastly, the collection of classifiers was involved in tumor grade classification.

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