



MICHIGAN INSTITUTE
FOR DATA & AI IN SOCIETY
UNIVERSITY OF MICHIGAN



BINARY CLASSIFICATION TO PREDICT TUMOR TYPE (BENIGN OR MALIGNANT)

BENIGN BY DESIGN 

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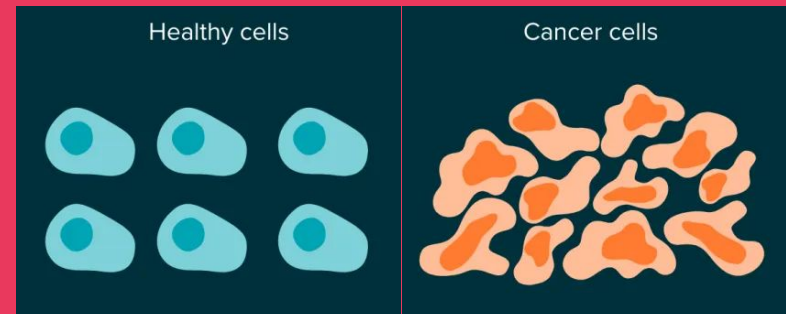
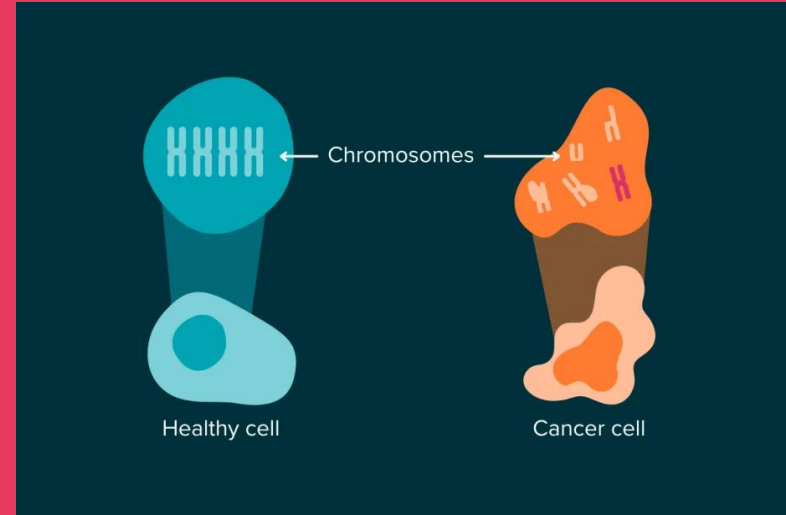
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INTRODUCTION

What could go wrong?

- The contents of the nucleus are very different for cancer cells
- Looking at cells, it can be determined if they are
 - M (Malignant): cancerous
 - B (Benign): non-cancerous



WHAT ARE WE MEASURING?

- The dataset features describing cell nuclei characteristics.
 - Each instance represents a single breast mass sample.
- Categories:
 - Radius, Texture, Perimeter, Area, Smoothness, Compactness, Concavity, Concave points, Symmetry, and Fractal dimension
- Three measurements/features each:
 - Mean, Standard Error, and Worst

(Kaggle.com)

SIMPLE MODELS ARE A SCIENTISTS BEST FRIEND

Reliability

Accountability

Interpretability

Sustainability

Executability

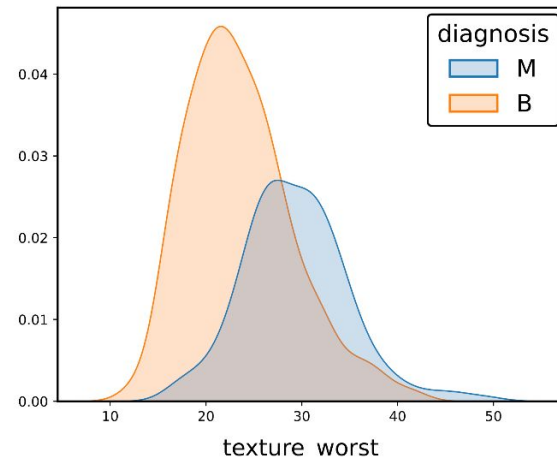
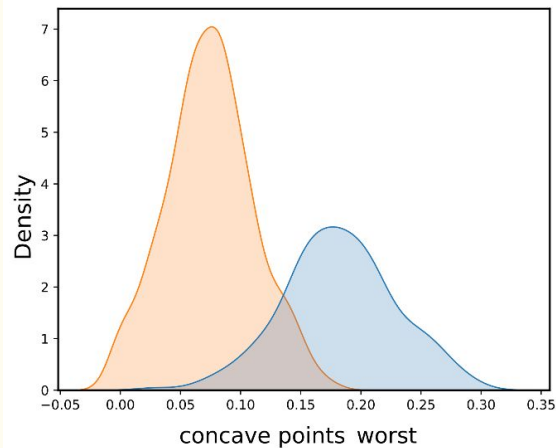
(Forbes 2021)

RESEARCH QUESTION

- **How many features are necessary to accurately predict a patient's breast cancer diagnosis using simple ML models?**
- **Measurements of success:**
 - High accuracy & Low false positive rate

Data Analysis

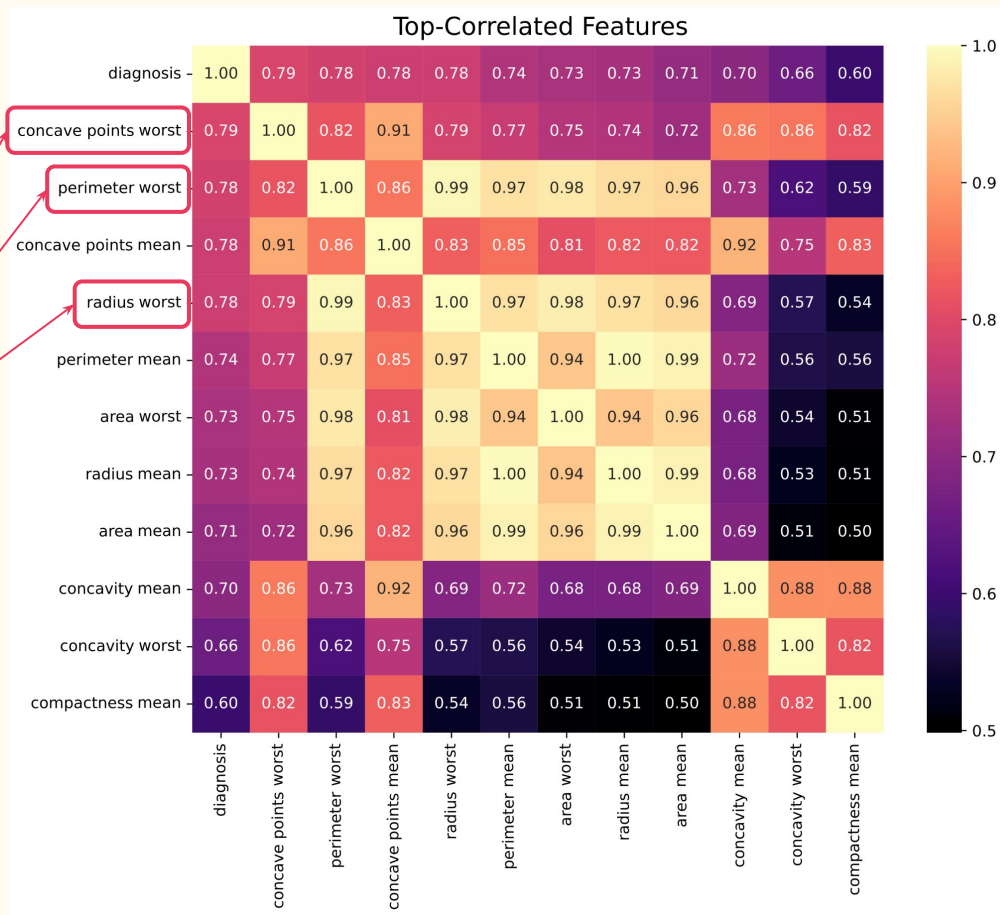
Bimodal Distribution



Data Analysis

Top-Correlated Features

Top 3 Features

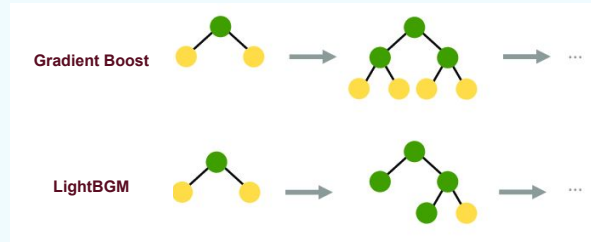
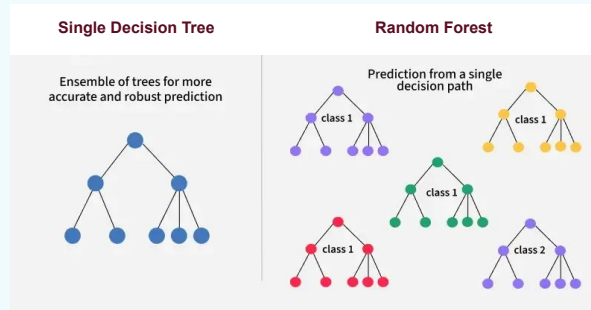
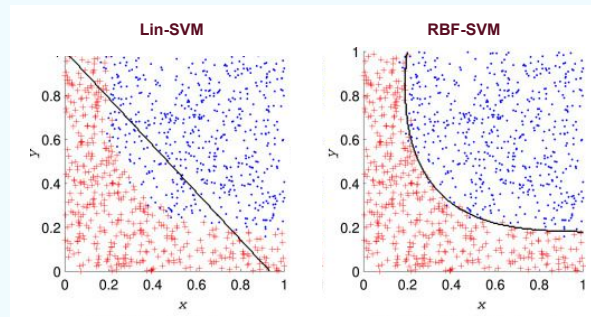
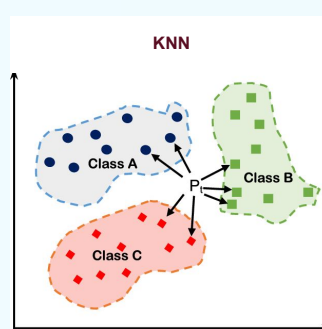
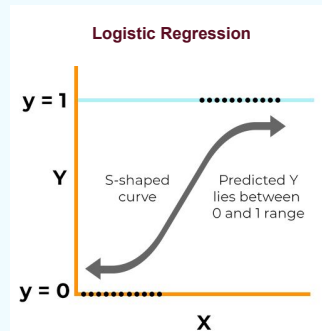




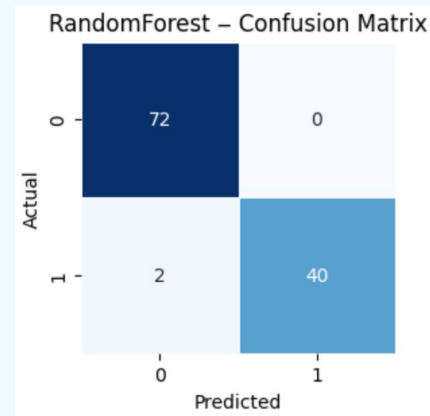
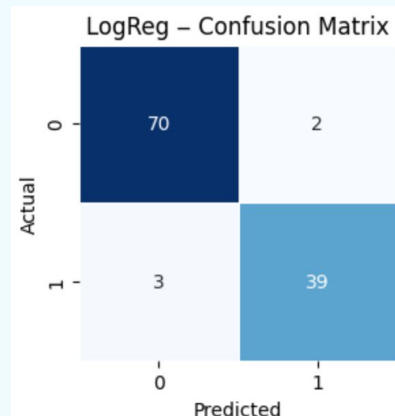
PREDICTING THE DIAGNOSIS

Setup

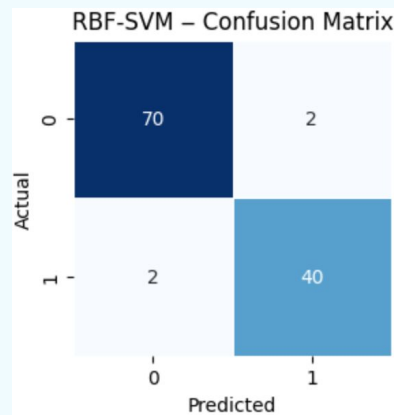
- Split data set 80/20
- Tested 8 different ML models
 - Logistic Regression, SVM (Lin & RBF), KNN, Decision Tree, Random Forest, Gradient Boost, LightGBM
- Tested use of different features (of the 30 possible)



Preliminary Results



Models	Accuracy	ROC-AUC
LogReg	0.956	0.994
Random Forest	0.982	0.996
RBF-SVM	0.965	0.995



Representative Model

Models	ROC-AUC			Accuracy		
	3	10	All	3	10	All
LogReg	0.994	0.996	0.994	0.947	0.965	0.956
Random Forest	0.984	0.994	0.996	0.939	0.965	0.982
RBF-SVM	0.995	0.994	0.995	0.956	0.974	0.965

3 Parameters

	Model	ROC-AUC	Accuracy	TP	FP	FN	TN
0	LinSVM	0.995	0.947	41	5	1	67
1	RBF-SVM	0.995	0.956	39	2	3	70
2	LogReg	0.994	0.947	41	5	1	67
3	KNN	0.993	0.956	39	2	3	70
4	LightGBM	0.988	0.921	40	7	2	65
5	RandomForest	0.984	0.939	39	4	3	68
6	GradBoost	0.982	0.921	39	6	3	66
7	DecisionTree	0.854	0.860	35	9	7	63

10 Parameters

	Model	ROC-AUC	Accuracy	TP	FP	FN	TN
0	LinSVM	0.996	0.965	40	2	2	70
1	LogReg	0.996	0.965	40	2	2	70
2	RBF-SVM	0.994	0.974	40	1	2	71
3	RandomForest	0.994	0.965	40	2	2	70
4	LightGBM	0.993	0.965	40	2	2	70
5	GradBoost	0.991	0.947	40	4	2	68
6	KNN	0.982	0.982	40	0	2	72
7	DecisionTree	0.941	0.939	40	5	2	67

30 Parameters

	Model	ROC-AUC	Accuracy	TP	FP	FN	TN
0	LinSVM	0.997	0.965	40	2	2	70
1	RandomForest	0.996	0.982	40	0	2	72
2	RBF-SVM	0.995	0.965	40	2	2	70
3	LogReg	0.994	0.956	39	2	3	70
4	LightGBM	0.993	0.974	39	0	3	72
5	GradBoost	0.992	0.939	40	5	2	67
6	KNN	0.981	0.974	39	0	3	72
7	DecisionTree	0.930	0.930	39	5	3	67

CONCLUSION

What can we conclude? Future directions?

1

Simple models

- Easy-to-understand algorithms
- Choose the ones most linked to the result
- Avoid using similar features

2

You don't need 30 features to predict a diagnosis

- With only 3 of top features achieves >95% fidelity
- Using one feature from each category is approx. same as using all

3

Future improvements

1. Can combine (stack or blend) models
2. Find larger dataset with more diverse data (not interdependent measurements)



**“KEEP IT
SIMPLE”**