

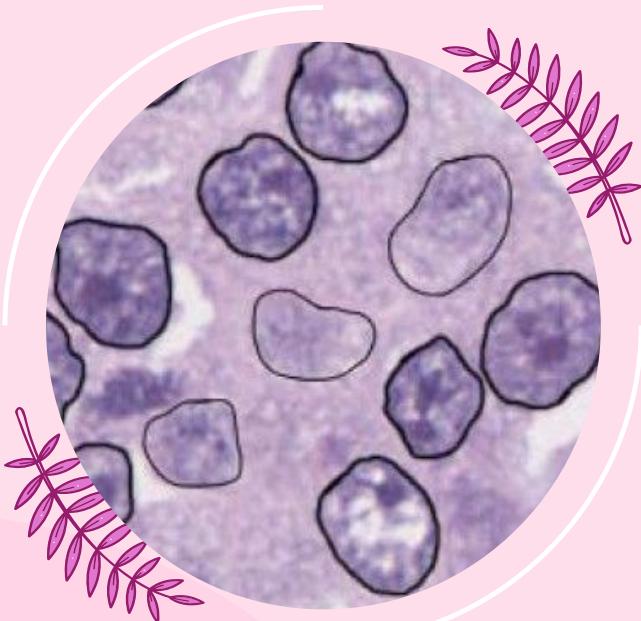
Breast Cancer Wisconsin Data Analysis

CellSight Diagnostics



The Diagnostic Challenge

- Our client is a radiology network exploring computer-aided diagnosis (CAD) systems for breast cancer screening
- We had received cell nuclei imaging data from fine needle aspirate (FNA) samples.
- Our team worked to identify which cellular characteristics best distinguish malignant from benign tumors to create a machine learning model that accurately predicts breast cancer



Why Our Mission Matters

- Breast cancer is the 2nd most common cancer and 2nd leading cause of cancer death among women in the U.S
- It is the leading cause of cancer death for black and hispanic women
- 13% of U.S women will develop invasive breast cancer in their lifetime
- Early detection at the localized stage could yield a >99% 5-year survival rate



Dataset Description

Wisconsin Diagnostic Breast Cancer Dataset (WDBC)

- **Source:** UCI ML Repository / Kaggle
- **File:** data.csv
- **Rows / Columns:** 569 rows, 33 columns

Target Variable:

- **Diagnosis (Categorical)**
 - M = Malignant
 - B = Benign
- **Purpose:**
 - Binary classification to predict if a tumor is malignant or benign.

Features:

- 30 numeric tumor measurements:
Radius, texture, perimeter, area, smoothness, compactness, concavity, concave points, symmetry , fractal dimension.
- Each of the 10 attributes is recorded in all three groups, giving 30 total numeric features
- Measurement Groups: **Mean, Standard Error, Worst values**
- One empty column: **Unnamed: 32**

Data Dictionary

Column	Description	Feature Type	Valid Values/Range	Notes/Issues
id	Unique patient/sample identifier	Identifier	8670-911320502 integers	Variable length, inconsistent formatting
diagnosis	Tumor classification	Binary Categorical	M (malignant), B(benign)	Target variable
radius_mean	Mean distance from center to nucleus perimeter	Continuous	6.98 - 28.11	Larger = more likely malignant
texture_mean	Mean gray-scale variation in nucleus	Continuous	9.71-39.28	Higher = more irregular surface
perimeter_mean	Mean nucleus perimeter	Continuous	43.79 - 188.50	Correlated with radius
area_mean	Mean area of cell nucleus	Continuous	143.50 - 2501.00	Malignant cells typically larger
smoothness_mean	Mean variation in radius lengths	Continuous	0.05 - 0.16	Lower values indicate smoother borders, higher values indicate irregular borders
compactness_mean	Mean nucleus compactness	Continuous	0.02 - 0.35	0 = perfect circle; higher = more irregular
concavity_mean	Mean severity of concave contour portions	Continuous	0.00 - 0.43	Higher = more indentations
concave points_mean	Mean number of concave contour points	Continuous	0.00 - 0.20	Malignant tumors have more concave points
symmetry_mean	Mean nucleus symmetry	Continuous	0.11 - 0.30	Lower = more symmetric = likely benign
fractal_dimension_mean	Mean boundary complexity	Continuous	0.05 - 0.10	Higher = more irregular border

Analysis of 10-Point Inspection

Core Findings

- **Dimensions:** 569 biopsy samples across 30 clinical features
- **Data Integrity:** 100% complete features, no missing values or duplicates
- **Target Balance:** Split is 62.7% Benign vs. 37.3% Malignant – clinically representative and sufficient for model training

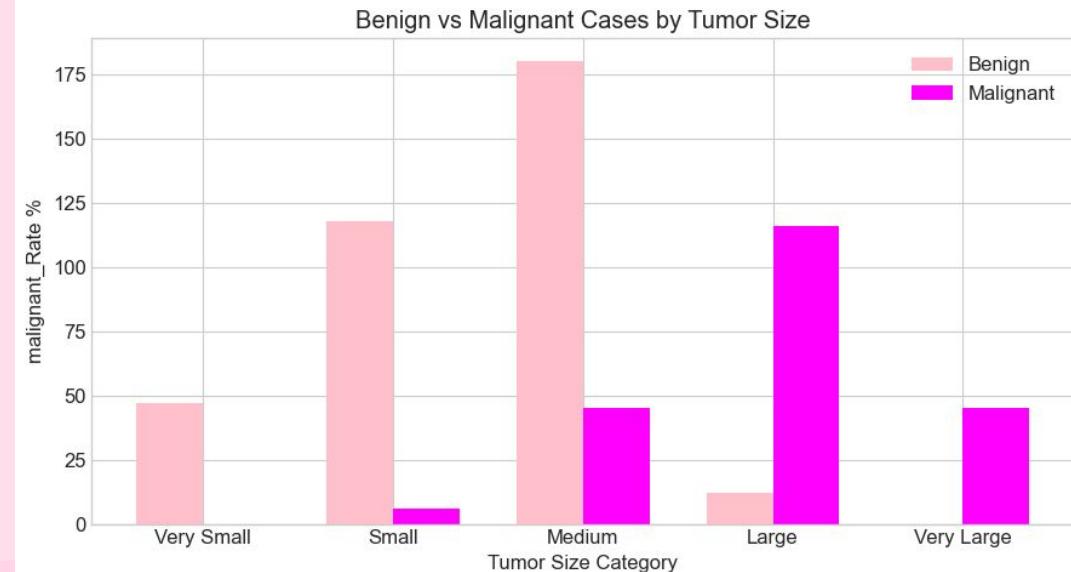
Data Quality Surprises

- **Empty Feature:** 'Unnamed: 32' column with 100% null values (likely CSV trailing comma error)
- **Class Distribution:** Malignant class is more frequent (~37%) than general population averages (~20%), providing a stronger signal for analysis

Overall: The dataset is high-quality, verified for internal logic, and required minimal cleaning

Tumor Size Categories

Tumor Size	Benign	Malignant	Total	Malignant Rate (%)
Very Small	47	0	47	0
Small	118	6	124	4.8
Medium	180	45	225	20
Large	12	116	128	90.6
Very Large	0	45	45	100



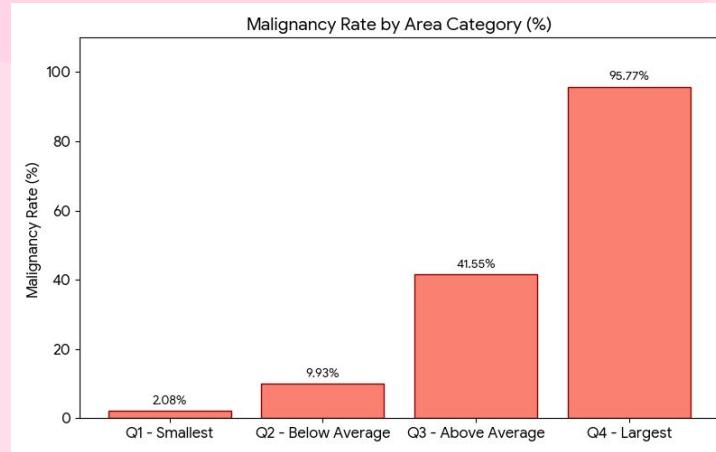
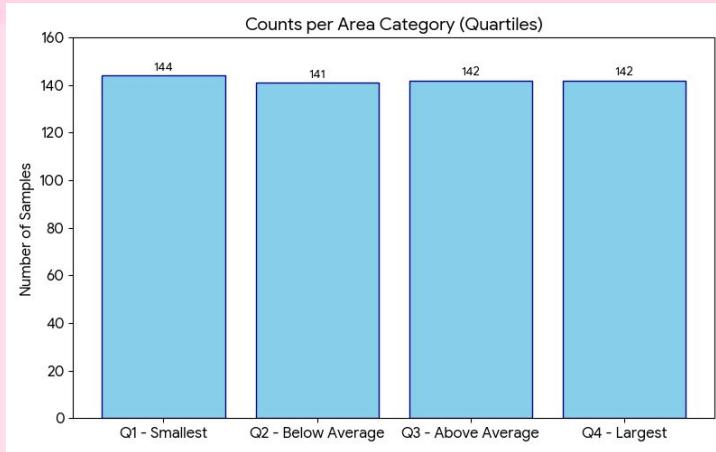
Malignant Tumors

- Correlation with size: **0.82**
- P-value: **0.089**

Benign Tumors

- Correlation with size: **-0.60**
- P-value: **0.28**

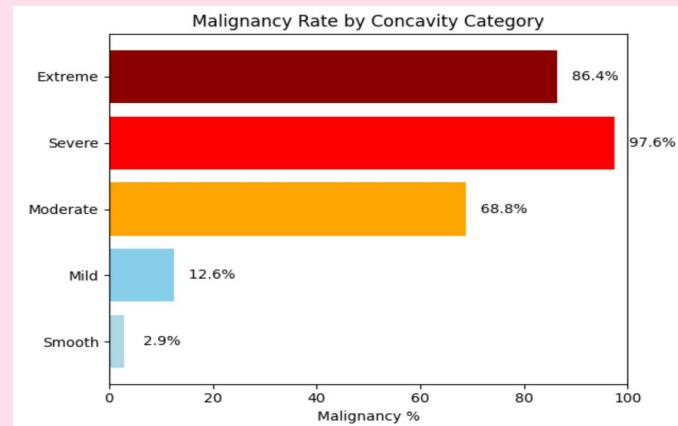
Tumor Area Categories



- **Quartiles for area_mean:** 25th Percentile (Q1) - 420.3, 50th Percentile (Median) - 551.1, and 75th Percentile (Q3) - 782.7
- **Distribution nearly equal** across the four categories
- Dramatic increase in malignancy rate as tumor area increases
 - **Strong positive correlation** between tumor size and malignancy

Cell Irregularity Categories

Category <i>(Intensity of cell indentations)</i>	Concavity Mean
Smooth	< 0.035
Mild	< 0.085
Moderate	< 0.155
Severe	< 0.255
Extreme	0.255 +



Smooth: 172 cases (2.91% Malignant)

Mild: 167 cases (12.57% Malignant)

Moderate: 125 cases (68.80% Malignant)

Severe: 83 cases (97.59% Malignant)

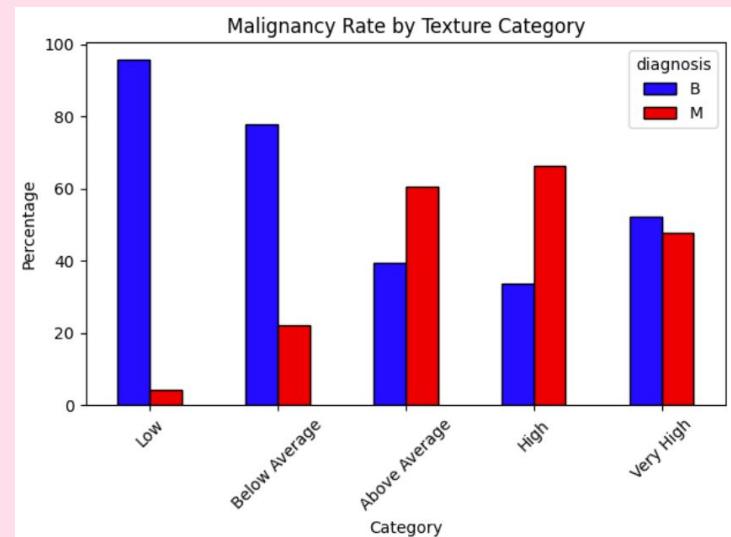
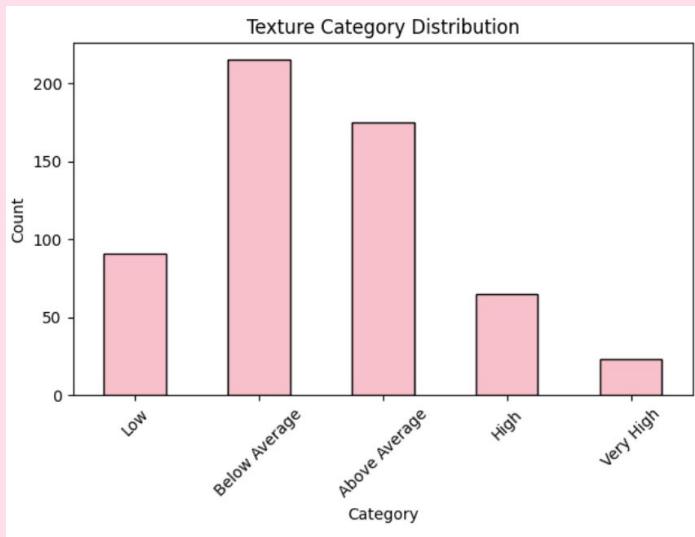
Extreme: 22 cases (86.36% Malignant)

Total malignant: 212 (37.26%)

Total benign: 357 (62.74%)

Total cases: 569

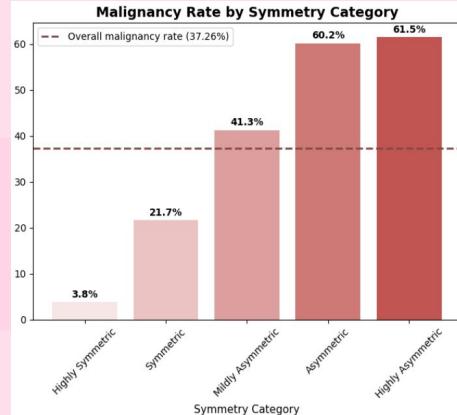
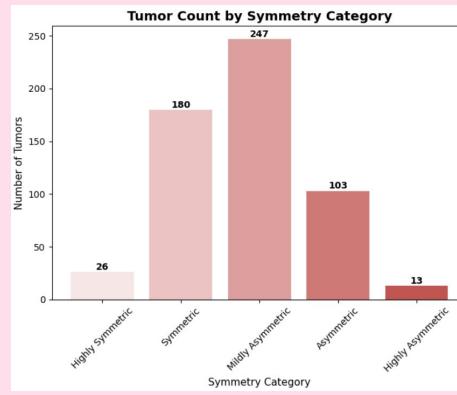
Texture Variability Categories



- Texture categories were made based on mean (19.29) and standard deviation (4.30)
- Malignant tumors had a higher average texture_mean (21.60) than benign tumors (17.91)
- Positive relationship between higher texture variability and malignancy

Symmetry-Based Categories

- Asymmetric Cells = Higher Malignancy Risk:
 - Cell division, chromosomal instability, unequal cell division, gene mutations all affect nuclear shape
- Most tumors (43.4%) fall in Mildly Asymmetric
- Malignancy rates rise as asymmetry increases
 - 3.8% -> 61.5%
- Symmetry is a useful diagnostic feature but cannot be used alone





Observations

- Malignancy Rate of Each Feature:
 - Very large tumor size = 100%
 - Most malignant tumors were in the very or large category
 - Large area category = 95.77%
 - Very high texture variability = 47.8%
 - Extreme cell irregularity = 86.4%
 - Texture and irregularity weren't perfectly proportional with malignancy because largest categories had lower malignancy rates than the 2nd largest categories
 - Highly asymmetric = 61.5% malignancy rate
 - Symmetry can't be used alone because most tumors were classified as mildly asymmetric

Recommendations & Conclusion

Key Findings:

- Tumor **size** and **area** are the most accurate and strongest predictors of malignancy.
 - Tumors in the **very large** category were **always** malignant
 - Tumors in the **large** category were malignant in more than 95% of cases

What features should the CAD system prioritize? What are the limitations?

- CAD system should prioritize tumor size and area
 - The sample data indicated that those features were the best indicators of tumor malignancy
- This dataset is limited to non-temporal data
 - More data on the progression of tumors may be helpful in predicting the malignancy of cells



Thank You!

ANY
QUESTIONS?