Classification of Dermatological Diseases

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Agenda

- → About the research
- → Objective and Business Gains
- → Dataset Description
- Dataset Preprocessing
- → Statistical Insights
- → Model Building
- → Challenges
- → Summary

About The Research

- 6 types of diseases to be classified
- → 12 clinical attributes and 22 histopathological attributes considered
- Clinical trials physical traits and symptoms
- Histopathological attributes structures at cellular level

Problem Statement

- Classification of six types of diseases based upon the records of patients
- → Misclassification of diseases leads to various complications on both sides; the patients as well as the doctors

Dataset Description

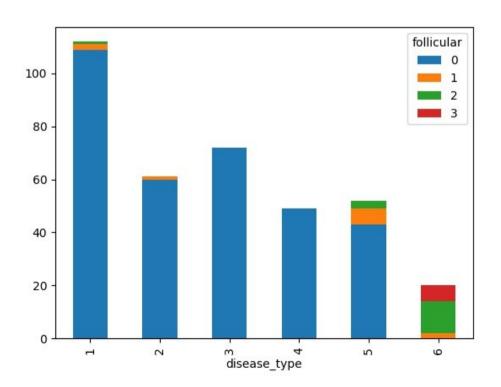
- → 366 records
- → Multi class Classification Problem 6 levels
- → 33 features are categorical
- → Age feature is numerical
- → 32 categorical features have levels as:
 - → 0: Not Present in patient
 - → 1,2: Intermediate levels
 - → 3: Maximum Level of symptoms reached
- → Family History is either 0 or 1

Dataset Preprocessing

- → No class imbalance
- All numerical features converted back to categorical
- → Age feature converted back to numerical feature
- → 8 missing values imputed via mean imputation, KNN imputation and using missForest package
- → The missForest package gave the best results for imputation

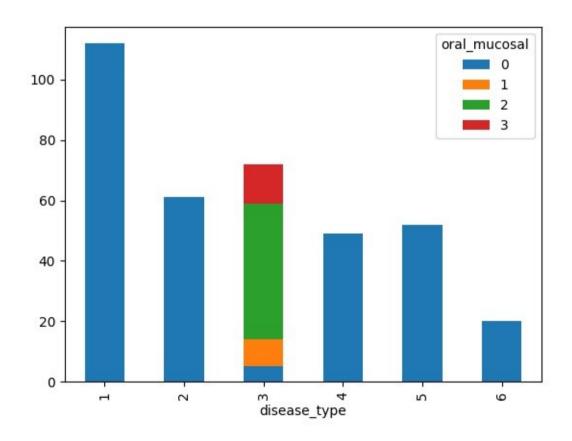
Statistical Insights: Clinical

- → Clinical trials :
 - → Follicular Papules main cause for Pityriasis Rubra Pilaris



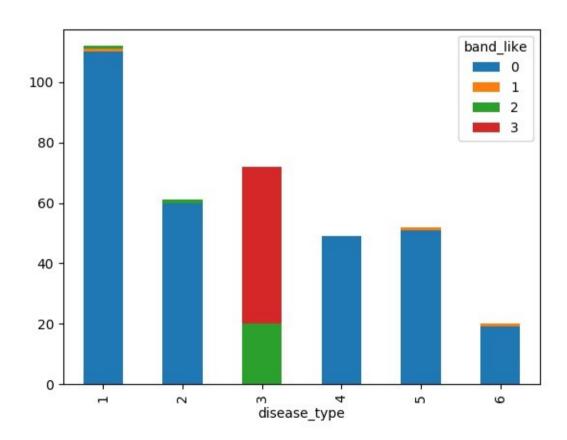
Clinical Trials

→ Oral Mucosal a huge cause for Lichen Planus



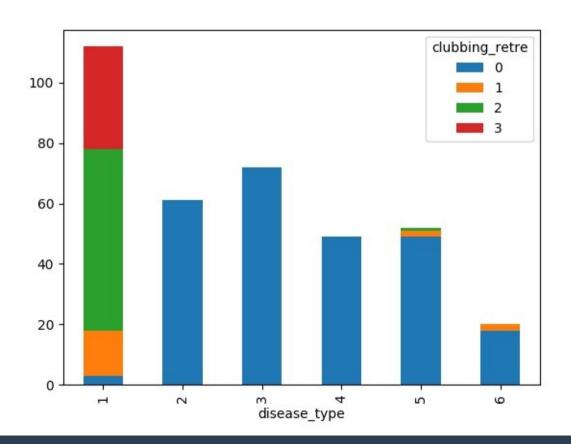
Histopathological Features

Band like structure of cell is mainly found in patients of lichen planus



Histopathological Features

 Clubbing of the retre ridges mainly found in patients suffering from psoriasis



Histopathological Features

- → Fibrosis of the papillary dermis mainly found in Chronic Dermatitis
- → Focal Hypergranulosis mainly in Lichen Planus
- → Horn Plug like structure found in Pityriasis Rubra Pilaris
- Melanin Inconsistencies, munro microabcess, perifollicular parakeratosis, Saw-Tooth like structures: All show the same behaviour

Insights: Summary

- Clinical traits generally present in all 6 types of diseases
- Histopathological features are more disease-specific
- → The structures and inconsistencies at cellular level more effective at classifying the diseases

Model Building

- → Data split into train, validation and test in ratio of 60:20:20
- → Models built:
 - → Multinomial Logistic
 - → C 5.0
 - **→** CART
 - → Bagging CART
 - → Random Forest
 - → Gradient Boost

Multinomial Logistic Model

	Accuracy	Precision	Recall	F1-Score
Class 1	94.23	100	88.46	93.87
Class 2	86.74	85.71	75	80
Class 3	100	100	100	100
Class 4	98.41	84.61	100	91.66
Class 5	99.20	91.66	100	95.65
Class 6	99.27	83.33	100	90.90

C 5.0 Model

	Accuracy	Precision	Recall	F1-Score
Class 1	98.07	100	96.15	98.03
Class 2	93.75	100	87.50	93.33
Class 3	99.18	92.85	100	96.29
Class 4	100	100	100	100
Class 5	100	100	100	100
Class 6	99.27	83.33	100	90.90

CART Model

	Accuracy	Precision	Recall	F1-Score
Class 1	96.15	100	92.30	96
Class 2	91.47	70	87.50	77.78
Class 3	100	100	100	100
Class 4	95.45	100	90.90	95.23
Class 5	100	100	100	100
Class 6	99.27	83.33	100	90.90

Bagging CART Model

	Accuracy	Precision	Recall	F1-Score
Class 1	100	100	100	100
Class 2	98.48	80	100	88.89
Class 3	100	100	100	100
Class 4	90.90	100	81.81	90
Class 5	100	100	100	100
Class 6	100	100	100	100

Random Forest

	Accuracy	Precision	Recall	F1-Score
Class 1	100	100	100	100
Class 2	99.18	92.85	100	96.29
Class 3	100	100	100	100
Class 4	92.85	100	85.71	92.30
Class 5	100	100	100	100
Class 6	100	100	100	100

Gradient Boosting

	Accuracy	Precision	Recall	F1-Score
Class 1	100	100	100	100
Class 2	98.36	86.66	100	92.85
Class 3	100	100	100	100
Class 4	85.71	100	71.42	83.33
Class 5	100	100	100	100
Class 6	100	100	100	100

Best Model

- → All the models give a good result on all the error metrics
- → However, random forest gives the best results

Challenges

- → Method of imputation : Only 8 missing values
 - → Introduced missing values at specific intervals in random columns
 - → Compared imputed values with original data
- → Choice of error metric

Summary

- → Objective
- → Insight
- → Model Selection
- → Challenges

Thank You

For Codes, please refer to www.github.com/AchintKumar/Internship