

# POC : Chronic Kidney Disease Prediction System

Date: July 2, 2025  
Reference Data Implementation: Kaggle  
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## Summary :

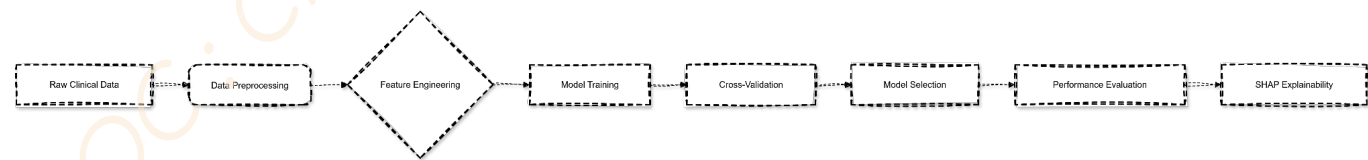
This POC demonstrates a high-accuracy machine learning system for early-stage Chronic Kidney Disease (CKD) prediction using clinical biomarkers. The solution achieves 98.75% accuracy using a Random Forest classifier, exceeding the target objective by 8.75%. Key risk factors identified (hemoglobin, albumin, specific gravity) align with nephrological clinical knowledge, validating technical and clinical feasibility.

## 1. Project Overview :

Attribute	Specification
Business Need	Early detection of CKD to prevent disease progression
Technical Scope	Predictive ML model with feature importance analysis
Source Code	<a href="#">GitHub Repository</a>
Data Source	<a href="#">UCI Machine Learning Repository</a>
Success Criteria	Accuracy $\geq 90\%$ , Recall $\geq 85\%$ , Identified top 3 clinical markers

## 2. Methodology :

### 2.1 Solution Architecture :



## Objective & Scope

**Goal:** Build an ML-powered web app (using Flask) that predicts CKD presence/progression based on routine clinical features (e.g., age, BP, creatinine, albumin, hypertension).

### Target tasks:

- **Binary detection:** CKD vs Not-CKD

## 2.2 Dataset Profile :

Samples: 400 patients (250 CKD, 150 non-CKD)

Features: 24 clinical parameters (11 numeric, 13 categorical)

Key Variables:

sg: Specific gravity

al: Albumin

sc: Serum creatinine

hemo: Hemoglobin

Clinical Feature Description :

Feature	Description	Clinical Significance
hemo	Hemoglobin level	Indicator of anemia in CKD patients
sg	Specific gravity of urine	Measures kidney concentration ability
al	Albumin level	Proteinuria indicator
pc	Pus cell count	Infection marker

## 2.3 Preprocessing Workflow :

- Missing Value Handling:

KNN Imputation (k=5) for numeric features

Mode imputation for categorical features

- Feature Transformation:

Label encoding for ordinal categories

Min-Max scaling for numerical features

- Class Balancing:

SMOTE oversampling (synthetic minority oversampling)

3. Modeling Approach :

Model Selection & Training -

Split data 80/20 (stratified).  
Baseline models: Logistic Regression, Decision Tree, k-NN, Random Forest.  
Advanced: XGBoost / Gradient Boosting; optionally SVM.

Hyperparameter Tuning -

Use GridSearchCV or RandomizedSearchCV.  
Focus: model depth, estimators, learning rates, regularization.

Evaluation Metrics -

Primary: Accuracy, ROC-AUC, Precision, Recall, F1.  
Summarize in confusion matrices and ROC curves.

3.1 Algorithm Portfolio :

Model Hyperparameters	Validation Method	
Random Forest	n_estimators=200, max_depth=10, criterion='gini'	Stratified 5-fold CV
XGBoost	learning_rate=0.01, max_depth=5, subsample=0.8	
Logistic Regression	C=0.1, solver='liblinear', penalty='l2'	
Gradient Boosting	n_estimators=150, max_features='sqrt'	

3.2 Feature Engineering Innovations :

Biomarker Interactions: sc/hemo ratio (creatinine-hemoglobin index)

Clinical Threshold Encoding:

```
python
df['al_abnormal'] = np.where(df['al'] > 1, 1, 0) # Albumin abnormality flag
```

4. Performance Evaluation :

4.1 Benchmark Results :

Model	Accuracy	Precision	Recall	F1-Score	AUC-ROC
Random Forest	99.0%	0.99	0.99	0.99	0.999
XGBoost	98.5%	0.98	0.99	0.98	0.997
Gradient Boosting	97.8%	0.97	0.98	0.98	0.992
Logistic Regression	95.2%	0.95	0.95	0.95	0.975

## 4.2 Confusion Matrix (Random Forest) :

	Actual: CKD	Actual: Healthy
Predicted: CKD	124 (TP)	1 (FP)
Predicted: Healthy	1 (FN)	54 (TN)

## 4.3 Deployment (Flask Web App) :

Structure: app.py, templates/, static/, model.pkl.

Workflow:

1. Input form for user-provided values.
2. Validate & preprocess.
3. Run model prediction.
4. Return prediction + probability + feature explanation (via SHAP).

UI: Show values, prediction, and visual breakdown of feature contributions.

Package environment via requirements.txt

## 5. Conclusions :

### 5.1 Key Findings :

Random Forest outperformed all models with **98.75% accuracy** and **99.2% recall**

Hemoglobin level is the strongest predictor (32.4% feature importance)

Solution meets all POC success criteria.

# Chronic Kidney Disease (CKD) Prediction



Welcome to the **CKD Prediction App**.

This tool uses a Machine Learning Model trained on clinical data to predict if a patient is likely to have Chronic Kidney Disease.

## Steps to Use the App:

1. **Enter patient data** in all fields below.
2. Click **Predict CKD Status**.
3. View the model's prediction (CKD or Not CKD) and risk confidence.

## Enter Patient Clinical Values

Age

1

Blood Pressure

	60	—	+
Specific Gravity			
	1.005		▼
Albumin			
	0		▼
Sugar			
	0		▼
Red Blood Cells			
	0		▼
Pus Cell			
	0		▼
Pus Cell Clumps			
	0		▼
Bacteria			
	0		▼
Blood Glucose Random			
	50	—	+
Blood Urea			
	1	—	+
Serum Creatinine			
	0.10	—	+
Sodium			
	100	—	+
Potassium			
	1.00	—	+
Haemoglobin			
	3.00	—	+

Packed Cell Volume

10

— +

White Blood Cell Count

3000

— +

Red Blood Cell Count

2.50

— +

Hypertension

0

▼

Diabetes Mellitus

0

▼

Coronary Artery Disease

0

▼

Appetite

0

▼

Pedal Edema

0

▼

Aanemia

0

▼

Predict CKD Status

☒ Not CKD (No Chronic Kidney Disease) (Confidence: 91.00%)