

# FINAL PROJECT - HEART FAILURE PREDICTION

## About this Dataset

Cardiovascular diseases (CVDs) are the number 1 cause of death globally, taking an estimated 17.9 million lives each year, which accounts for 31% of all deaths worldwide. Heart failure is a common event caused by CVDs and this dataset contains 12 features that can be used to predict mortality by heart failure.

Most cardiovascular diseases can be prevented by addressing behavioural risk factors such as tobacco use, unhealthy diet and obesity, physical inactivity and harmful use of alcohol using population-wide strategies.

People with cardiovascular disease or who are at high cardiovascular risk (due to the presence of one or more risk factors such as hypertension, diabetes, hyperlipidaemia or already established disease) need early detection and management wherein a machine learning model can be of great help.

## Feature Descriptions

Age : age [years]

anaemia : Decrease of red blood cells or hemoglobin (boolean)

creatinine\_phosphokinase : Level of the CPK enzyme in the blood (mcg/L)

diabetes : If the patient has diabetes (boolean)

ejection\_fraction : Percentage of blood leaving the heart at each contraction (percentage)

high\_blood\_pressure : If the patient has hypertension (boolean)

platelets : Platelets in the blood (kiloplatelets/mL)

serum\_creatinine : Level of serum creatinine in the blood (mg/dL)

serum\_sodium : Level of serum sodium in the blood (mEq/L)

sex : Woman or man (binary)

smoking : If the patient smokes or not (boolean)

time : Follow-up period (days)

DEATH\_EVENT : If the patient deceased during the follow-up period (boolean). This will be our Dependent Variable.

## Data Retrieving

```
%config Completer.use_jedi = False
```

```
import numpy as np
import pandas as pd
import matplotlib.pyplot as plt
import seaborn as sns
from collections import Counter
from scipy import stats
import random
```

```
data =
pd.read_csv('Downloads/heart_failure_clinical_records_dataset.csv')
data.head()
```

	age	anaemia	creatinine_phosphokinase	diabetes
0	75.0	0	582	0
20				
1	55.0	0	7861	0
38				
2	65.0	0	146	0
20				
3	50.0	1	111	0
20				
4	65.0	1	160	1
20				

	high_blood_pressure	platelets	serum_creatinine	serum_sodium	sex
0	1	265000.00	1.9	130	1
1	0	263358.03	1.1	136	1
2	0	162000.00	1.3	129	1
3	0	210000.00	1.9	137	1
4	0	327000.00	2.7	116	0

	smoking	time	DEATH_EVENT
0	0	4	1
1	0	6	1
2	1	7	1
3	0	7	1
4	0	8	1

## EDA & Data Cleaning

```
data.describe()
```

	age	anaemia	creatinine_phosphokinase	diabetes	\
count	299.000000	299.000000	299.000000	299.000000	
mean	60.833893	0.431438	581.839465	0.418060	
std	11.894809	0.496107	970.287881	0.494067	
min	40.000000	0.000000	23.000000	0.000000	
25%	51.000000	0.000000	116.500000	0.000000	
50%	60.000000	0.000000	250.000000	0.000000	
75%	70.000000	1.000000	582.000000	1.000000	
max	95.000000	1.000000	7861.000000	1.000000	

	ejection_fraction	high_blood_pressure	platelets	\
count	299.000000	299.000000	299.000000	
mean	38.083612	0.351171	263358.029264	
std	11.834841	0.478136	97804.236869	
min	14.000000	0.000000	25100.000000	
25%	30.000000	0.000000	212500.000000	
50%	38.000000	0.000000	262000.000000	
75%	45.000000	1.000000	303500.000000	
max	80.000000	1.000000	850000.000000	

	serum_creatinine	serum_sodium	sex	smoking
time \				
count	299.000000	299.000000	299.000000	299.000000
299.000000				
mean	1.39388	136.625418	0.648829	0.32107
130.260870				
std	1.03451	4.412477	0.478136	0.46767
77.614208				
min	0.50000	113.000000	0.000000	0.00000
4.000000				
25%	0.90000	134.000000	0.000000	0.00000
73.000000				
50%	1.10000	137.000000	1.000000	0.00000
115.000000				
75%	1.40000	140.000000	1.000000	1.00000
203.000000				
max	9.40000	148.000000	1.000000	1.00000
285.000000				

	DEATH_EVENT
count	299.00000
mean	0.32107
std	0.46767
min	0.00000
25%	0.00000
50%	0.00000
75%	1.00000
max	1.00000

data.isnull().sum()

```

age                                0
anaemia                           0
creatinine_phosphokinase          0
diabetes                          0
ejection_fraction                 0
high_blood_pressure               0
platelets                         0
serum_creatinine                  0
serum_sodium                      0
sex                               0
smoking                           0
time                              0
DEATH_EVENT                       0
dtype: int64

```

*Note: as shown above there are no null values in our dataset!*

```
data.shape
```

```
(299, 13)
```

```
data.info()
```

```
<class 'pandas.core.frame.DataFrame'>
```

```
RangeIndex: 299 entries, 0 to 298
```

```
Data columns (total 13 columns):
```

#	Column	Non-Null Count	Dtype
0	age	299 non-null	float64
1	anaemia	299 non-null	int64
2	creatinine_phosphokinase	299 non-null	int64
3	diabetes	299 non-null	int64
4	ejection_fraction	299 non-null	int64
5	high_blood_pressure	299 non-null	int64
6	platelets	299 non-null	float64
7	serum_creatinine	299 non-null	float64
8	serum_sodium	299 non-null	int64
9	sex	299 non-null	int64
10	smoking	299 non-null	int64
11	time	299 non-null	int64
12	DEATH_EVENT	299 non-null	int64

```
dtypes: float64(3), int64(10)
```

```
memory usage: 30.5 KB
```

```
data.columns.to_list()
```

```

['age',
 'anaemia',
 'creatinine_phosphokinase',
 'diabetes',
 'ejection_fraction',
 'high_blood_pressure',

```

```
'platelets',
'serum_creatinine',
'serum_sodium',
'sex',
'smoking',
'time',
'DEATH_EVENT']
```

```
data.nunique()
```

```
age                47
anaemia            2
creatinine_phosphokinase  208
diabetes           2
ejection_fraction  17
high_blood_pressure  2
platelets          176
serum_creatinine    40
serum_sodium        27
sex                 2
smoking             2
time               148
DEATH_EVENT         2
dtype: int64
```

```
#Dividing features into numerical and categorical
```

```
col = list(data.columns)
categorical_features = []
numerical_features = []
for i in col:
    if len(data[i].unique()) > 2:
        numerical_features.append(i)
    else:
        categorical_features.append(i)
```

```
print('Categorical Features :',*categorical_features)
print('Numerical Features :',*numerical_features)
```

```
Categorical Features : anaemia diabetes high_blood_pressure sex
smoking DEATH_EVENT
Numerical Features : age creatinine_phosphokinase ejection_fraction
platelets serum_creatinine serum_sodium time
```

```
data['age'] = data['age'].astype(int)
data['platelets'] = data['platelets'].astype(int)
df = data.copy(deep = True)
df
```

```
   age  anaemia  creatinine_phosphokinase  diabetes
ejection_fraction \
0      75         0                    582         0
```

20					
1	55	0		7861	0
38					
2	65	0		146	0
20					
3	50	1		111	0
20					
4	65	1		160	1
20					
..	...	...		...	...
..					
294	62	0		61	1
38					
295	55	0		1820	0
38					
296	45	0		2060	1
60					
297	45	0		2413	0
38					
298	50	0		196	0
45					

	high_blood_pressure	platelets	serum_creatinine	serum_sodium
sex \				
0	1	265000	1.9	130
1				
1	0	263358	1.1	136
1				
2	0	162000	1.3	129
1				
3	0	210000	1.9	137
1				
4	0	327000	2.7	116
0				
..	...	...	...	...
..				
294	1	155000	1.1	143
1				
295	0	270000	1.2	139
0				
296	0	742000	0.8	138
0				
297	0	140000	1.4	140
1				
298	0	395000	1.6	136
1				

	smoking	time	DEATH_EVENT
0	0	4	1
1	0	6	1

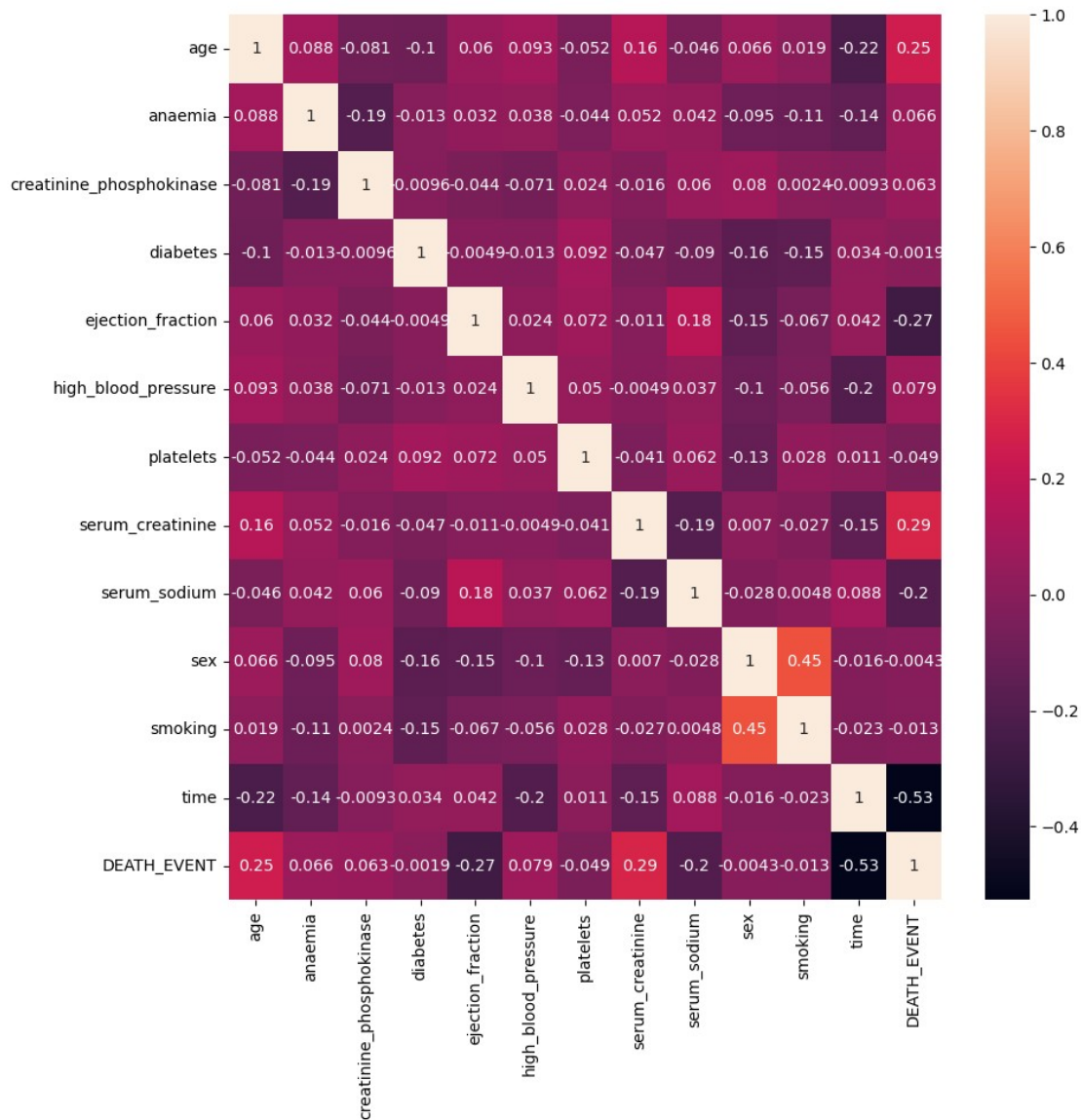
2	1	7	1
3	0	7	1
4	0	8	1
..	...	...	...
294	1	270	0
295	0	271	0
296	0	278	0
297	1	280	0
298	1	285	0

[299 rows x 13 columns]

```

correlation = df.corr()
plt.figure(figsize=(10, 10))
sns.heatmap(correlation, annot=True)
plt.show()

```



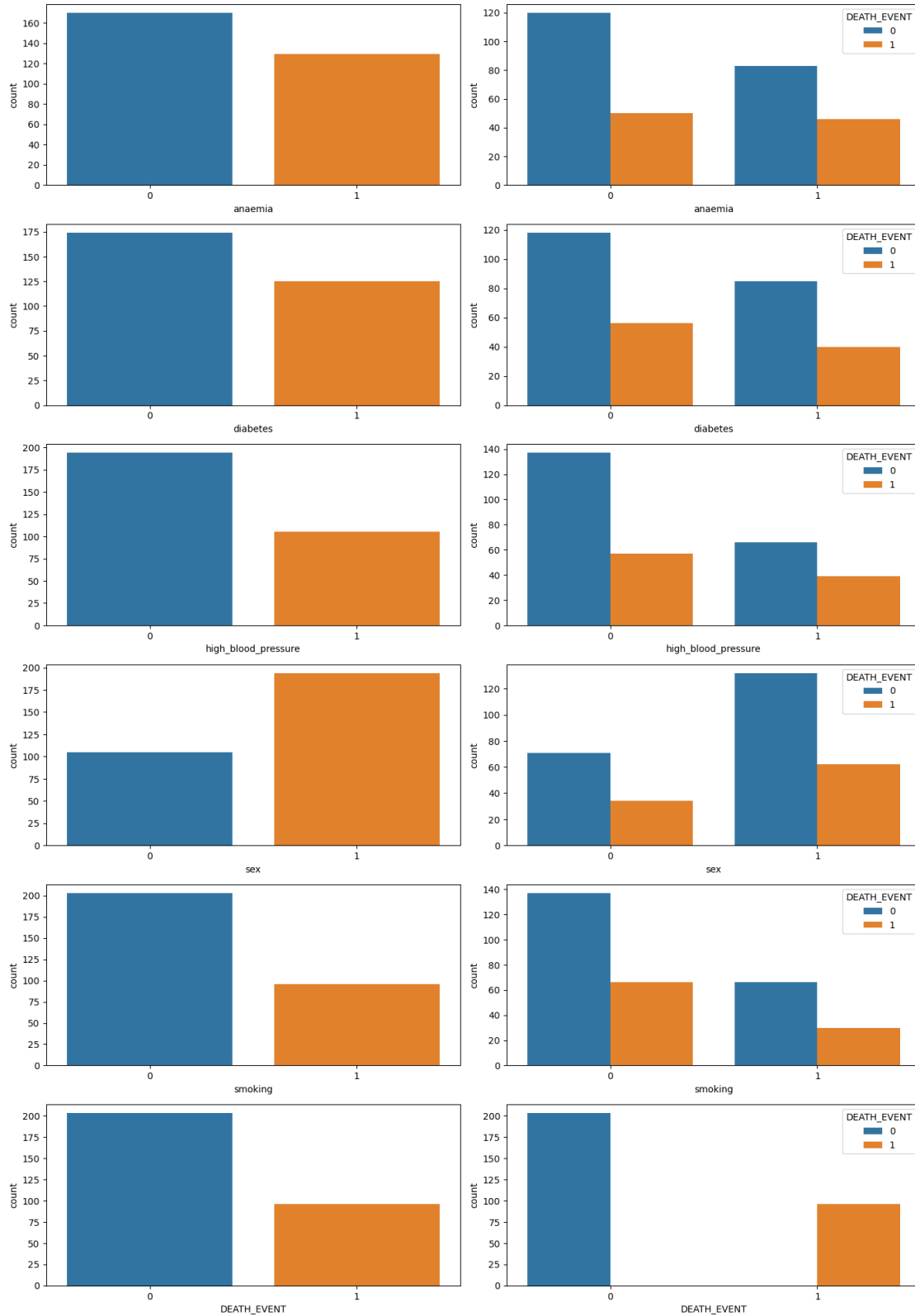
## Observations

There is nothing to conclude from categorical features correlation matrix. From the correlation matrix for numerical features, time is inversely correlated to death. Thus patients with less follow up time are prone to heart failure. Based on EDA, features such as anaemia, diabetes, age, sex, smoking are less contributing.

```
fig, ax = plt.subplots(len(categorical_features), 2, figsize=(14,20))
```

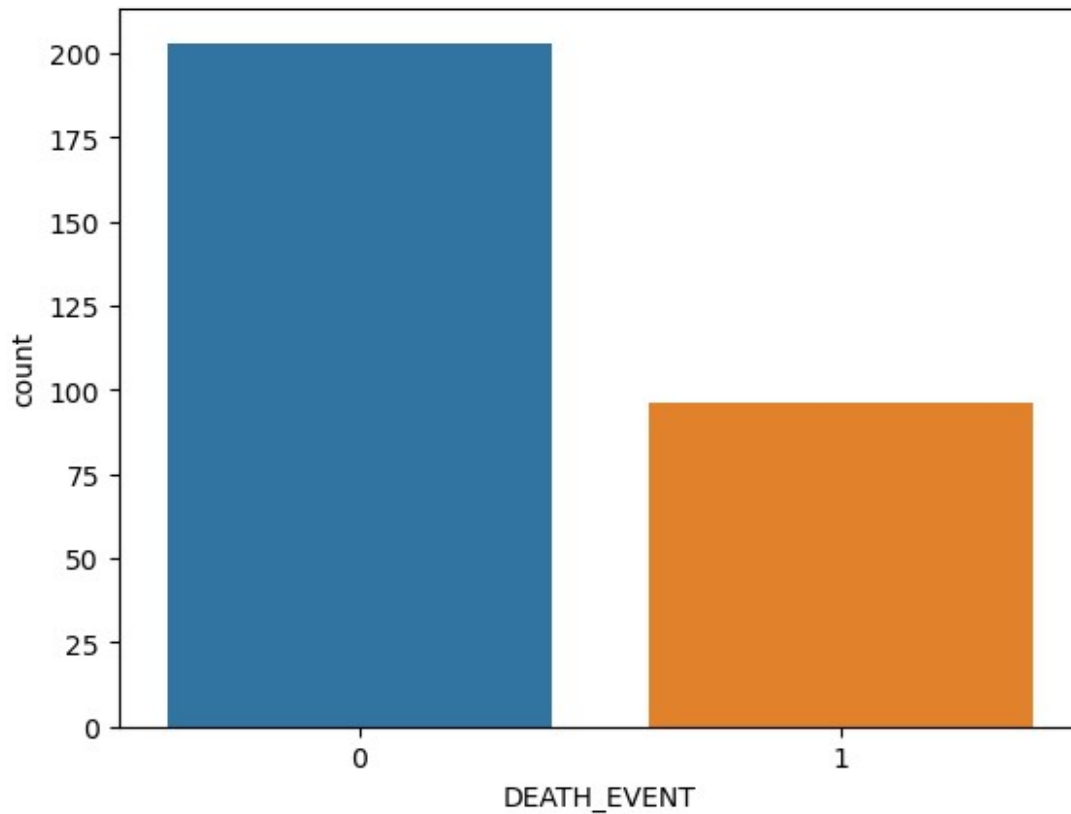
```
for i in range(len(categorical_features)):
    sns.countplot(ax=ax[i, 0], x=categorical_features[i], data=df)
    sns.countplot(ax=ax[i, 1], x=categorical_features[i],
hue='DEATH_EVENT', data=df)
fig.tight_layout(pad=1)
plt.show()
```





```
sns.countplot(x='DEATH_EVENT', data=df)
```

```
<AxesSubplot:xlabel='DEATH_EVENT', ylabel='count'>
```

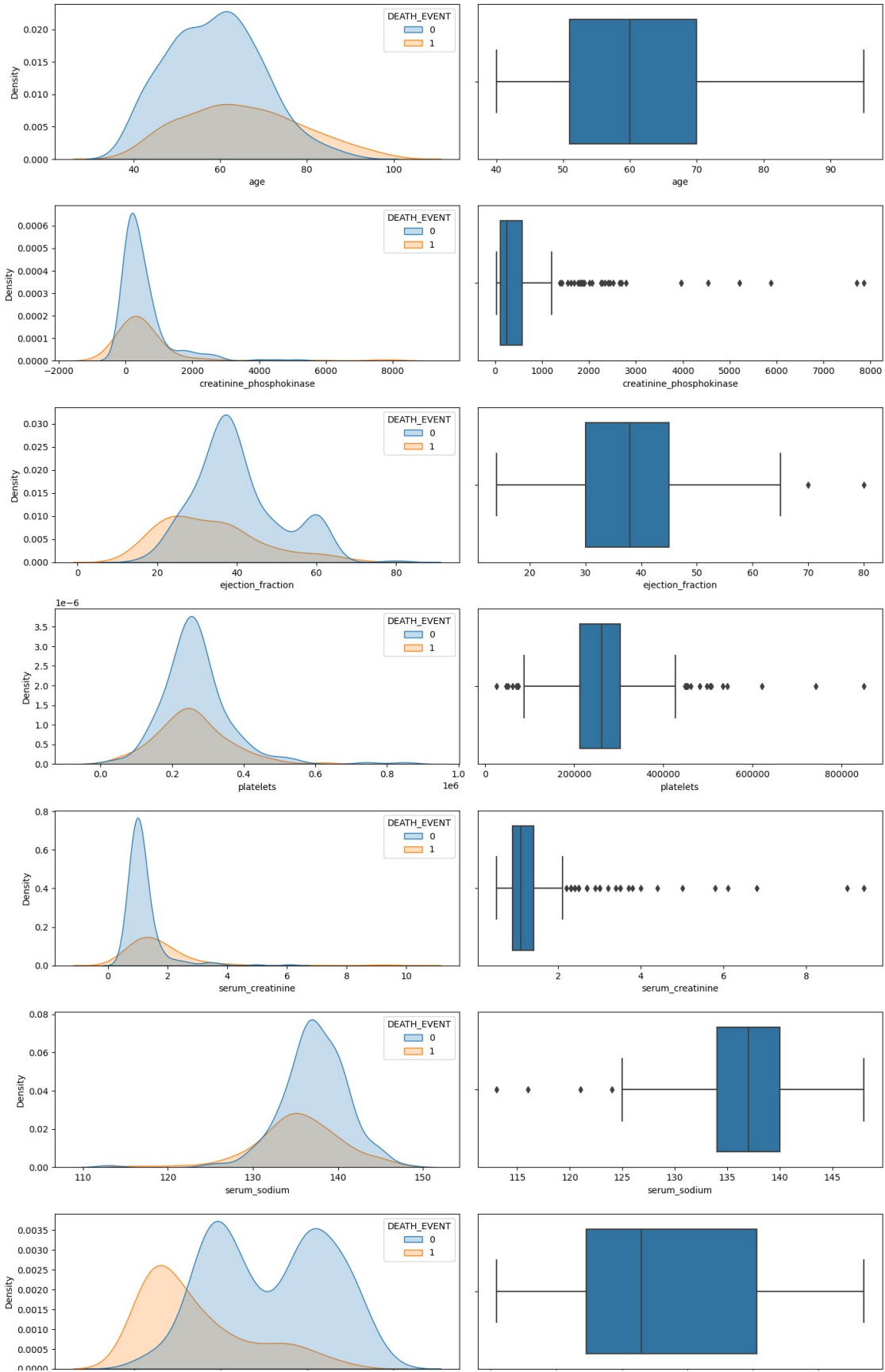


### Observations

There is an imbalance with the target variable, so we can apply cross validation technique with over sampling method compared to under sampling as the data size is small.

```
fig, ax = plt.subplots(len(numerical_features), 2, figsize=(14,22))

for i in range(len(numerical_features)):
    sns.kdeplot(ax=ax[i, 0], x=numerical_features[i],
hue='DEATH_EVENT', data=df, fill = True)
    sns.boxplot(ax=ax[i, 1], x=numerical_features[i], data=df)
fig.tight_layout(pad=1)
plt.show()
```



## Observations

- No missing values present in the data.
- From the correlation matrix for continuous features, time is inversely correlated to death. Thus patients with less follow up time are prone to heart failure.
- Smoking and Sex features are slightly correlated.
- Based on EDA, features such as anaemia, diabetes, age, sex, smoking are less contributing.
- There is an imbalance with the target variable, so we can apply cross validation technique with over sampling method compared to under sampling as the data size is small.
- creatinine\_phosphokinase, serum\_creatinine and serum\_sodium are highly skewed.
- From KDE Plots and boxplots, we can find that there are outliers in the data.
- creatinine\_phosphokinase, serum\_creatinine contains many outliers and can be treated using IQR Formula.

## Hypothesis Testing

### Hypothesis 1

**alpha =0.05**

**Null Hypothesis (H01):** There is no relation between smoking and sex.

**Alternate Hypothesis (Ha1):** There is a relationship between the means of smoking and sex of patient as it is a observation of EDA.

```
t_score, p_val = stats.ttest_ind(df['smoking'], df['sex'])
print(t_score, p_val)
```

```
-8.47376579265773 1.8637689471019785e-16
```

```
dof = 25 + 25 - 2 # degree of freedom
t_dist = stats.t(dof)
print(2 * t_dist.cdf(t_score))
```

```
4.2756596819011004e-11
```

```
if p_val < 0.05:    # alpha value is 0.05 or 5%
    print(" we are rejecting null hypothesis")
else:
    print("we are accepting null hypothesis")
```

```
we are rejecting null hypothesis
```

*Result: As p-value is less than 0.05, null hypothesis 1 is rejected. So there is a relationship between the means of smoking and sex of patient as it is a observation of EDA.*

## Hypothesis 2

**Null Hypothesis (H02):** The mean age of the sample set is not equal from that of complete data.

**Alternate Hypothesis (Ha2):** The mean age of the sample set is almost equal to complete data.

```
sample_age_set = np.random.choice(list(df['age']), 20)

_, p_val = stats.ttest_1samp(sample_age_set, df['age'].mean())
print(p_val)

0.282528137071198

if p_val < 0.05:
    print(" we are rejecting null hypothesis")
else:
    print("we are accepting null hypothesis")

we are accepting null hypothesis
```

*Result: As p-value is more than 0.05, null hypothesis 2 is accepted. So the mean age of the sample set is not equal from that of complete data although its close enough.*

```
print('Mean of complete data: ', np.mean(data['age']))
print('Mean of sample data: ', np.mean(sample_age_set))

Mean of complete data: 60.82943143812709
Mean of sample data: 63.8
```

## Hypothesis 3

**Null Hypothesis (H03):** There is a some connection between high BP and risk of heart failure as high BP stress on the heart functioning thus might affecting the patients predictions.

**Alternate Hypothesis (Ha3):** There is no relation between high BP and heart failure.

```
_, p_val = stats.ttest_ind(df['DEATH_EVENT'],
df['high_blood_pressure'])
print(p_val)

0.43675818272737343

if p_val < 0.05:
    print(" we are rejecting null hypothesis")
else:
    print("we are accepting null hypothesis")
```

we are accepting null hypothesis

*Result: As p-value is greater than 0.05, null hypothesis 3 is accepted. So there is a some connection between high BP and risk of heart failure as high BP stress on the heart functioning thus might affecting the patients predictions.*

## Future Scope

Will try out different classifiers and selecting one with highest recall score. Recall represents the False Negative values which is very crucial in medical diagnosis. As the dataset is imbalanced, need to tackle the major issue. Alongwith that data rows are less, so will opt for oversampling or SMOTE or weighted class method. Feature selection is also crucial as some features are proven to be more prominent based on EDA, thus contributing more rather than selecting and processing all.

## Conclusion

In this way, we have analysed the heart failure patient dataset, understanding the aspects of it. One more thing to observe is the data consists of time and death event variables. These variables are more important with respect to the survival model analysis. The same data can be used for survival analysis of the patients thus predicting the risk score and survival probability of the patient over the course of time.