Time to heart failure survival analysis

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Introduction

Heart failure is a chronic condition characterized by the heart's inability to pump an adequate amount of blood to meet the body's demands. It can occur when the heart muscle becomes weakened or damaged, resulting in symptoms such as shortness of breath, fatigue, and fluid retention. Various factors, including coronary disease, diabetes, and obesity, can contribute to the development of heart failure. In this study, our goal is to evaluate the significance of different parameters on the survival of patients with heart failure. We analyze the occurrence of patient deaths as the event of interest.

Data description

The individuals of the data were patients admitted to Institute of Cardiology and Allied hospital Faisalabad-Pakistan during April-December (2015). From the 299 patients of the dataset, 105 are women and are 194 men. They are between 40 and 95 years. All have left ventricular systolic dysfunction, belonging to New York Heart Association (NYHA) class III and IV. Follow up time was between 4 to 285 days. — maybe we remove the rest of this section Class III means patients have marked limitations of physical activity. They are comfortable at rest but experience symptoms with less than ordinary physical activity. Class IV means patients are unable to carry out any physical activity without discomfort. They may have symptoms even at rest and are often bedridden.

The database has 13 features, including Age, Anemia, High Blood Pressure, Creatinine phosphokinase, Diabetes, Ejection Fraction, Sex, Platelets, Serum Creatinine, Serum Sodium, Smoking, Time, and Death Event. Out of these features, 5 are Boolean variables, namely Anemia, High Blood Pressure, Diabetes, Sex, and Smoking. We added two new features to the dataset: "over60" and "EF_levels." The "over60" feature categorizes individuals as either over 60 years old or not, based on their age. The "EF_levels" feature categorizes individuals into different groups based on their Ejection Fraction. These new features allow us to create Kaplan-Meier survival curves and analyze the data based on these specific characteristics.

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The presence of time and death event in this dataset makes it suited for survival analysis. The unit of time in the dataset is measured in days. Since not all patients experienced the event of interest (death), the dataset contains right-censored data.

Univariate analysis: group comparison

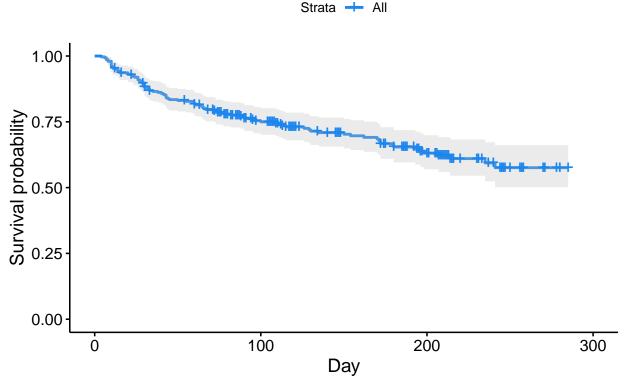
Overall Kaplan-Meyer estimator

We estimate the survival probability with the Kaplan - meier estimator.

```
## Call: survfit(formula = Surv(time, DEATH_EVENT) ~ 1, data = data)
##
## n events median 0.95LCL 0.95UCL
## [1,] 299 96 NA NA NA
```

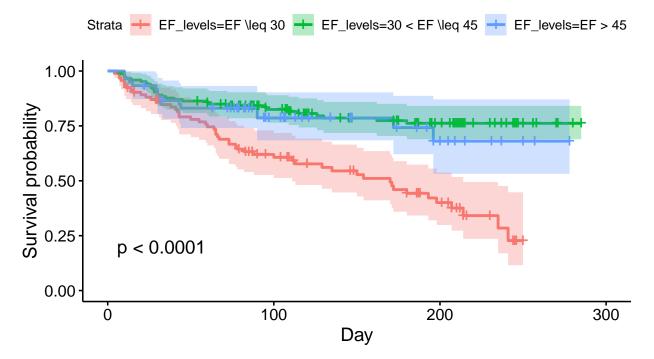
96 (32%) patients died due to the Cardiovascular Heart Disease (CHD). The median, 0.95LCL and 0.95UCl are NA because too many data are right censored. We need to go deeper in the analysis.

Kaplan-Meier Curve for Heart Falure Survival



As the EF level and the high tension are the directly heart related covariates, we use them for the Kaplan Meier survival estimate

Kaplan-Meier Curve per EF levels for Heart Failure Surviva



With a p-value less than 0.0001, the EF levels are indeed statistically significant to the death for patient with heart failure as shown by the plot.

We apply the same method to the others covariates. The p-value for continuous covariates are obtained with Cox proportional hazards model. The results are summerized in the following tab:

								ser.	
covariate	Sex	Smoking	Diabetes	Aenemia	Age	creat.phos platelets		creat	ser.sodium
p-value	0.95	0.96	0.84	0.099	8.36e-07	0.26	0.4585	1.18e-07	0.00052

Since the p-value is less than 0.05 we reject the null hypothesis for age, serum creatinine and serum sodium, this means these covariates have a statistically significant impact in survival.

Cox Proportional Hazards Model

Univariate Cox regression

We will be examining the significance of each covariate using the Cox regression model. The results will be presented in a table, which includes the covariates, their beta coefficients, hazard ratios, lower confidence intervals, upper confidence intervals, and p-values.

```
columns_to_remove <- c("time", "DEATH_EVENT", "EF_levels")
column_names <- column_names [!column_names %in% columns_to_remove]

#indices_to_delete <- c(5,12, 13,14,15)
covariates <- column_names #[-indices_to_delete]
#covariates <- append(covariates, 'EF_levels')
covariates</pre>
```

```
##
    [1] "age"
                                    "anaemia"
    [3] "creatinine phosphokinase" "diabetes"
##
##
   [5] "ejection_fraction"
                                    "high_blood_pressure"
   [7] "platelets"
                                    "serum creatinine"
  [9] "serum_sodium"
                                    "sex"
##
## [11] "smoking"
                                    "bad_platelet"
## [13] "over60"
```

print(updated_df)

```
##
                                   beta
                                          HR lower_ci
                                                       upper_ci p.value
                                                 1.00
                                                            1.10 8.4e-07
## age
                                4.2e-02 1.00
## anaemia
                                3.4e-01 1.40
                                                 0.94
                                                            2.10 1.0e-01
## creatinine_phosphokinase
                                1.1e-04 1.00
                                                 1.00
                                                           1.00 2.6e-01
## diabetes
                              -4.2e-02 0.96
                                                 0.64
                                                           1.40 8.4e-01
## ejection_fraction
                              -4.6e-02 0.95
                                                 0.93
                                                           0.98 1.7e-05
## high_blood_pressure
                               4.4e-01 1.50
                                                 1.00
                                                           2.30 3.7e-02
## platelets
                              -7.8e-07 1.00
                                                 1.00
                                                           1.00 4.7e-01
## serum creatinine
                               2.9e-01 1.30
                                                 1.20
                                                           1.50 1.2e-07
## serum sodium
                              -6.8e-02 0.93
                                                 0.90
                                                           0.97 5.2e-04
                                1.4e-02 1.00
                                                 0.67
                                                           1.50 9.5e-01
## sex
```

```
## smoking
                              -9.6e-03 0.99
                                                 0.64
                                                           1.50 9.7e-01
## bad_platelet
                               2.2e-01 1.20
                                                 0.73
                                                           2.10 4.2e-01
## over60
                              -4.5e-01 0.64
                                                 0.43
                                                           0.95 2.7e-02
## 30 <= EF vs 30 < EF <= 45 -1.2e+00 0.31
                                                 0.20
                                                           0.49 4.4e-07
## 30 <= EF vs EF > 45
                              -9.0e-01 0.41
                                                 0.23
                                                           0.74 3.0e-03
```

For table above we say that for all these covariates, including anaemia, creatinine_phosphokinase, diabetes, platelets, sex, and smoking, have p-values greater than the chosen significance level (e.g., 0.05). This suggests that these covariates are statistically insignificant, indicating that there is no strong evidence of a significant association between these variables and the hazard rate.

Ejection Fraction (EF) appears to be a significant factor as it shows statistical significance for both comparisons: $30 \le EF \le 30 \le EF \le 45$ and $30 \le EF \le 45$, with p-values below 0.05. For both EF level comparisons, the negative beta coefficients indicate a negative association between EF levels and the hazard rate. The hazard ratios of 0.31 and 0.41 suggest lower hazard rates for the specified EF levels compared to the baseline group.

High Blood Pressure is another significant factor with p-value = 0.037. These covariates have The positive beta coefficient of 0.44 suggests a positive association High Blood Pressure and risk of death.

Multivariate Cox regression analysis

additive effect of age and anaemia.

p-value = 0.054 There is no difference in survival time with respect to age and anaemia

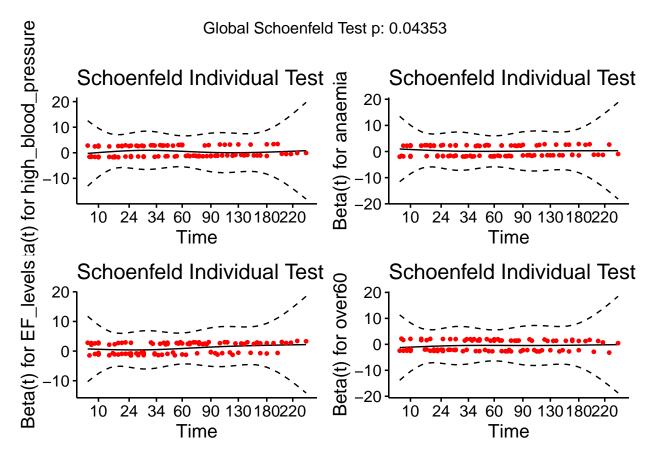
additive effect of high_blood_pressure and anaemia.

```
## p-value = 0.06584754 There is no additive effect of hbp and anaemia
```

effect of EF + hbp + age

```
##
                                     coef exp(coef)
                                                         se(coef)
                                                                      Pr(>|z|)
                             0.0738706418 1.0766675 1.548158e-02 1.828421e-06
## age
## creatinine_phosphokinase 0.0002350038 1.0002350 9.982031e-05 1.855922e-02
                            -0.0633925801 0.9385749 2.626050e-02 1.577889e-02
## ejection_fraction
## high_blood_pressure1
                             0.4751724707 1.6082916 2.214657e-01 3.190646e-02
## serum_creatinine
                             0.2864581650 1.3317025 7.482003e-02 1.288590e-04
## serum_sodium
                            -0.0442830531 0.9566831 2.228763e-02 4.693464e-02
## over60>60
                             0.7409593957 2.0979473 3.589062e-01 3.897099e-02
```

additive effect of high_blood_pressure and anaemia and EF_levels and age over 60.



Patients presenting high_blood_pressure, EF_levels30 < EF <= 45, EF_levelsEF > 45, age > 60, serum_creatinine, serum_sodium are more likely to experience heart failure and die. Moreover, presenting a high_blood_pressure increases the risk of heart failure by a hazard rate of 1.53, holding other covariates fixed, while, EF_levels30 < EF <= 45, EF_levelsEF > 45, age > 60, serum_creatinine and serum_sodium increases the risk of dying from heart failure with hazard rate magnitude greater or almost equal to 1.

Results

In this application, we were to analyze time to heart failure in this cohort, the event being death. We have found that, there is no median time to death. Our results show that there is an effect of high blood pressure, EF and ageing, but surprisingly no effect of Smocking, nor diabetes or anaemia. There is a group difference in survival time in EF and high blood pressure. Half of subject having EF less than 30, are expected to die after almost 175 days. An additive model with covariates, high_blood_pressure1, EF_levels30 < EF <= 45, EF_levelsEF > 45, age > 60, serum_creatinine and serum_sodium shows that, patients presenting a profile with theses covariates are more likely to experience heart failure and die. Also, There is a noticeable effect of ageing (age > 60).

Discussion

We aim at analyzing time to heart failure. We have found that, there is an effect of high blood pressure, EF, serum_creatinine, serum_sodium and ageing, but surprisingly no effect of Smocking, nor diabetes or anaemia. (why: to document)

Conclusions

Following our results, we can conclude that ageing, Ejection fraction, serum_creatinine, serum_sodium and high blood pressure influence time to heart failure and the occurence of CHD as well as ageing. Overall, above half of these patients died early of heart failure. Though there was a high risk of death in this cohort..

References

Ahmad T, Munir A, Bhatti SH, Aftab M, Raza MA (2017) Survival analysis of heart failure patients: A case study. PLoS ONE 12(7): e0181001. https://doi.org/10.1371/journal.pone.0181001