

1 **Title: Risk Factor for Postoperative Pneumonia after Coronary Artery Bypass Grafting**

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42 **Risk Factor for Postoperative Pneumonia after Coronary Artery Bypass Grafting**

43 **ABSTRACT**

44 **Background:** Postoperative pneumonia (POP) is a common infectious complication of
45 coronary artery bypass grafting (CABG), leading to significant morbidity, mortality, and
46 increased healthcare costs. This study found that the prevalence of POP was nearly double that
47 reported in previous studies, underscoring the urgent need to identify specific risk factors.
48 These findings emphasize the importance of local data in refining preventive strategies and
49 improving clinical outcomes in CABG patients.

50 **Material and Methods:** This is a retrospective cohort study. The subjects comprised patients
51 who underwent CABG procedures at a single institution between June 2020 and June 2024. A
52 logistic regression analysis model for evaluating the risk of POP was established.

53 **Results:** This study observed a POP rate of 41.7%, significantly exceeding the 2–24% range
54 reported in previous studies. Key risk factors included elevated creatinine levels, eGFR <60
55 ml/min/1.73 m², and low early postoperative albumin. POP strongly correlated with prolonged
56 hospitalization, with an odds ratio of 13.043 (95% CI: 6.130–27.751, p<0.001), underscoring
57 its substantial impact on patient outcomes.

58 **Conclusions:** The present study delineates renal impairment and hypoalbuminemia
59 postoperative as pivotal risk factors for POP following CABG. It emphasizes the importance
60 of tailored interventions, structured institutional practices, and continuous research to enhance
61 preventive strategies and patient outcomes.

62 **Keywords:** coronary artery bypass grafting; postoperative pneumonia; risk factor

63 1. Introduction

64 Coronary revascularization may be conducted through two principal approaches: surgical
65 coronary artery bypass grafting (CABG) and percutaneous coronary intervention (PCI). CABG
66 is generally considered a higher-risk procedure in comparison to PCI and is frequently linked
67 to a more extended recovery period. Nevertheless, in the long term, CABG is correlated with
68 a reduced recurrence rate of coronary artery disease.^{1,2} Despite advancements in CABG,
69 significant risks persist due to complications during and after the procedure, raising morbidity
70 and mortality rates. To reduce these risks, comprehensive strategies for prevention, monitoring,
71 evaluation, and prompt management of complications are essential. Understanding patient risk
72 factors, healthcare performance, and postoperative conditions is crucial to effectively prevent
73 and manage post-CABG complications. Addressing these factors is key to improving patient
74 outcomes and reducing complication rates procedure.³

75 Postoperative infections are common complications, occurring in up to 16% of cases.
76 They can delay healing, prolong hospital stays, and increase mortality. Various strategies exist
77 to reduce these infections, starting with preoperative screening and extending to postoperative
78 care ICU.⁴ Postoperative pneumonia constitutes a significant infectious complication
79 subsequent to cardiac surgery, resulting in substantial increases in morbidity, mortality, and
80 healthcare expenditures. Postoperative pneumonia incidence varies between 2.1% and 21.6%.
81 The demographic profile of cardiac surgery patients has evolved significantly. Despite
82 advancements in surgical techniques and anesthesia, the aging population with multiple
83 comorbidities and rising antibiotic-resistant pathogens increases the number of high-risk
84 patients for complications POP.⁶

85 The majority of investigations concerning risk factors associated with POP have taken
86 place in developed regions, including the United States and Europe, while data from Southeast
87 Asia remain limited. Variations in patient demographics and medical histories have the
88 potential to affect these risk factors, thereby underscoring the necessity for region-specific
89 research. This study endeavors to fill this gap by identifying crucial risk factors for POP
90 following CABG and aims to offer valuable insights that could enhance clinical outcomes in
91 Southeast Asian populations.

93 **2. Materials and Methods**

94 This study utilized an analytical observational approach with a retrospective cohort
95 design. The target population was CAD patients after CABG-only surgery. The accessible
96 population in this study were CAD patients after CABG-only surgery registered in the “Cardiac
97 Surgery Registry” at Dr. Hasan Sadikin General Hospital, Bandung, with the ethical number
98 DP.04.03/D.XIV.6.5/324/2024.

99 The research focused on patients with heart disease who underwent CABG procedures
100 at Dr. Hasan Sadikin General Hospital in Bandung. The study included all CABG patients from
101 June 2020 to June 2024. Patients with the following conditions were excluded from this study:
102 acquired pneumonia within two weeks before surgery, death or discharge within 48 hours after
103 surgery, and combination surgery with another procedure besides CABG. Patients who had the
104 CABG procedure more than once were also excluded

105 Clinical data were collected from the hospital's records system. Preoperative variables
106 included demographics (sex, age, height, weight, BMI, smoking history), comorbidities
107 (hypertension, diabetes, COPD, peripheral arterial disease, renal disease), left ventricular
108 ejection fraction, and laboratory values. Intraoperative variables included CPB time.
109 Postoperative variables covered mechanical ventilation duration, early albumin levels, hospital
110 stay length, and mortality.

111 Postoperative pneumonia was defined by new or progressive pulmonary infiltrates on
112 chest radiographs and at least two of the following: fever over 38 °C without known cause,
113 leukocytosis $>12 \times 10^9/L$, leukopenia $<4 \times 10^9/L$, and purulent secretions. Semiquantitative
114 cultures were employed to ascertain the microbiological etiology of pneumonia from sputum
115 through initial microscopic examination alongside quantitative bacterial cultures. Hypertension
116 is defined as a blood pressure of 140/90 mmHg or higher, a prior diagnosis, or use of
117 antihypertensive medication. Diabetes mellitus is characterized by fasting glucose of 126 mg/dl
118 or more, random glucose of 200 mg/dl or higher, a past diagnosis, or the use of diabetes
119 medication. A history of smoking includes both current and past daily smoking habits. Chronic
120 Obstructive Pulmonary Disease (COPD) is defined by an FEV1/FVC ratio of ≤ 0.7 from
121 spirometry. Renal insufficiency is indicated by serum creatinine >1.24 mg/dL or a prior
122 diagnosis. The creatinine level and estimated glomerular filtration rate (eGFR) were calculated
123 one day prior to surgery. Early postoperative albumin refers to the serum albumin levels

124 measured within the first 24 hours following surgery. The area under the curve in the receiver
125 operating characteristic plot is utilized as a cutoff point for analysis.

126 Statistical analysis was performed using SPSS (IBM SPSS Statistics, version 26).
127 Patients with and without POP were compared by univariate analysis using the Chi Square test
128 for categorical variables and the Fisher exact test for discrete variables. Variables with a p value
129 < 0.25 on univariate analysis were entered into a multivariate logistic regression analysis to
130 identify the independent risk factors. Significance was considered at a p-value <0.05.

131

132 **3. Results**

133 From June 2020 to June 2024, 256 CABG procedures were performed. Fifty patients
134 were excluded from the analysis: thirty patients underwent combination cardiac surgery besides
135 CABG (valvular surgery, closure defects intracardiac, repair septal rupture), sixteen patients
136 died during the first 48 hours, and four patients got pneumonia before surgery. The remaining
137 206 patients underwent isolated CABG procedures instituted by the cohort. POP was
138 diagnosed in the 86 patients (41.7%). Table 1 shows the baseline characteristics of patients.

139 The most common microorganism isolated in this study was *Klebsiella pneumoniae*
140 (16%), followed by *Staphylococcus epidermidis*, *Pseudomonas aeruginosa*, and
141 *Staphylococcus haemolyticus* (9%). Polymicrobial POP was detected in 14% of cases.
142 However, in 54% of cases, no microorganism growth was found in culture examination. The
143 prophylactic antibiotic administered to patients is ceftriaxone 2 grams. According to the
144 antibiotic resistance data gathered from our hospital, *Klebsiella pneumoniae* has been identified
145 as the second most prevalent microorganism isolated from sputum cultures, following
146 *Acinetobacter baumannii*, in the intensive care unit. This organism demonstrates the highest
147 level of sensitivity to amikacin (>64%). Sensitivity to ceftriaxone has been documented to vary
148 between 15% and 30%. *Pseudomonas aeruginosa* ranks as the third most prevalent

149 microorganism isolated from sputum cultures within our hospital, exhibiting the highest
150 sensitivity level recorded towards amikacin (>64%). Currently, there is no available data
151 concerning pseudomonas sensitivity to the aforementioned antibiotic ceftriaxone.

152 No statistically significant differences were observed in univariate analysis between POP
153 and non-POP patients, except for estimated GFR <60 (32.6% in the POP group vs. 18.3% in
154 the non-POP group; p=0.019), early postoperative albumin (2.73 ± 0.49 in the POP group vs.
155 3.07 ± 0.58 in the non-POP group; p<0.001), acute clinical presentation preoperatively (29.1%
156 in the POP group vs. 12.5% in the non-POP group; p=0.003), cardiopulmonary bypass duration
157 (26.7% in the POP group vs. 10.8% in the non-POP group; p=0.003), and hospital length of
158 stay (84.9% in the POP group vs. 30% in the non-POP group; p<0.001).

159 Univariate analysis of risk factors for POP is presented in Table 2. Some variables had a
160 p-value of less than 0.25: sex, COPD, creatinine, eGFR < 60 ml/min/1.73 m², early
161 postoperative albumin, NLR value, RDW-CV value, preoperative clinical presentation, CPB
162 time, duration of mechanical ventilation, and hospital LOS. Chronic obstructive pulmonary
163 disease has a p-value of less than 0.25, but it has missing data of more than 5%; therefore, to
164 prevent substantial bias that may weaken the integrity of multivariate analysis, this variable
165 cannot be processed further into multivariate analysis.

166 Multivariate analysis identified three independent risk factors for POP (Table 3):
167 creatinine level (odds ratio [OR], 3.88; 95% CI, 1.04–14.43), eGFR < 60 ml/min/1.73 m² (odds
168 ratio [OR], 6.93; 95% CI, 1.87–25.73), and early postoperative albumin < 2.93 g/dL (OR, 3.22;
169 95% CI, 1.54–6.7). POP occurrence has a strong association with prolonged hospital stays
170 (odds ratio [OR], 13.04; 95% CI 6.13-27.75).

171

172 **4. Discussion**

173 Predictive models for postoperative pneumonia (POP) after cardiac surgery exist, but
174 data on CABG-only procedures are limited, especially in Southeast Asia. CABG is the most
175 common cardiac surgery globally, making it vital to explore unique risk factors for POP in
176 these cases. Independent risk factors for POP differ across studies due to variations in
177 population characteristics and diagnostic definitions. This study offers insights into POP risk
178 factors for CABG-only procedures, addressing a regional gap and enhancing global
179 understanding of this complication. Identified patient characteristics align with previous
180 research, underscoring the need for tailored risk assessment and management approaches
181 prevention.

182 Kilic et al. developed a validated 33-point risk score for POP from 6,222 patients
183 (2005–2012), noting a 4.5% incidence. Their model identified six key perioperative predictors:
184 advanced age, chronic lung disease, peripheral vascular disease, prolonged cardiopulmonary
185 bypass (CPB) time, intraoperative blood transfusion, and use of an intra-aortic balloon pump
186 period.⁸ Wang et al. reported a higher incidence of POP (9.96%) among 5,323 patients and
187 developed a 32-point risk score incorporating 13 independent risk factors, including age > 60
188 years, hypertension, diabetes mellitus, smoking, COPD, BMI > 24 kg/m², renal insufficiency,
189 NYHA class III-IV, preoperative anemia, hypoalbuminemia, CPB time > 120 minutes, and
190 blood transfusion.⁹ Allou et al. proposed a scoring system from a study of 5,582 patients with
191 a 3.1% incidence of POP. Their model identified four key risk factors: advanced age, COPD,
192 low preoperative LVEF, and the interaction of RBC transfusion with prolonged CPB
193 duration.¹⁰ These studies underscore the multifactorial nature of POP risk and the importance
194 of identifying patient-specific and procedure-specific factors to guide preventive strategies.

195 This study identified three independent risk factors for postoperative pneumonia (POP)
196 following CABG: elevated creatinine levels, eGFR <60 ml/min/1.73 m², and early
197 postoperative albumin <2.93 g/dL. The observed POP rate of 41.7% was significantly higher

198 than previously reported rates (2–24%).^{11–15} Gram-negative bacteria were the predominant
199 pathogens, with *Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, *Staphylococcus*
200 *epidermidis*, and *Staphylococcus haemolyticus* as the most frequently isolated species. These
201 findings align with prior studies in which *K. pneumoniae* and *P. aeruginosa* were common
202 causes of POP after cardiac surgery.^{9,10,16} Mortality among patients with POP was significantly
203 higher compared to that of those without pneumonia, underscoring the severe impact of POP
204 on outcomes. While the overall mortality rate was consistent with previous studies, these
205 findings highlight the critical need for effective prevention and management of POP to reduce
206 associated mortality risks.¹¹

207 In the present study, both elevated creatinine levels and reduced eGFR were identified
208 as independent risk factors for POP after cardiac surgery. Elevated creatinine levels were
209 significantly associated with POP, with an odds ratio (OR) of 3.883 (95% CI: 1.045–14.434, p
210 = 0.043) in the final model. Similarly, reduced eGFR <60 ml/min/1.73m² showed an even
211 stronger association, with an OR of 6.935 (95% CI: 1.869–25.734, p = 0.004). Impaired renal
212 function, indicated by these markers, likely contributes to systemic inflammation, immune
213 dysfunction, and fluid imbalances, increasing susceptibility to pulmonary infections. Research
214 by Wang et al. and Kilic et al. highlighted similar associations, with creatinine and eGFR
215 serving as robust predictors of postoperative complications, particularly infections.^{8,9}
216 Estimated GFR <60 ml/min/1.73 m² was observed in 24.3% of patients, predisposing them to
217 infection due to systemic inflammation, fluid imbalance, and impaired immune function.
218 Studies have consistently shown that renal dysfunction is a strong predictor of postoperative
219 complications, including pneumonia.^{9,14,17} Improving renal function preoperatively may reduce
220 complications and enhance prognosis in renal dysfunction patients.

221 In the early postoperative period, albumin levels at or below 2.93 g/dL were observed
222 in 57.8% of patients, thereby indicating nutritional deficiencies. Hypoalbuminemia undermines

223 wound healing, reduces immune responses, and increases infection susceptibility. Malnutrition
224 adversely affects surgical outcomes, leading to higher complication rates, longer recovery, and
225 extended hospitalization. Albumin, the main human protein, is a key nutritional status marker
226 and predicts surgical outcomes. Its quick decline post-surgery indicates surgical stress and
227 metabolic response. Hubner et al. showed that postoperative serum albumin reductions
228 correlate with surgical stress and predict more complicated recovery course.¹⁸ Perioperative
229 nutritional support, including immune-modulating formulas, has been shown to reduce
230 infectious complications and hospital stays after major surgeries.¹⁹

231 The mean age of patients with POP in this study was 58.29 ± 7.633 years, which is
232 lower than the mean age reported in previous studies (68 ± 13 years).¹⁰ Older patients face
233 higher complication risks due to weakened immune function, reduced pulmonary reserve, and
234 age-related comorbidities. However, the lower POP rate in earlier studies suggests other
235 influential factors. In this research, younger patients might have had greater comorbidity
236 burdens, indicated by renal impairment, smoking history, and hypoalbuminemia—strong risk
237 factors for POP. These conditions likely outweigh the benefits of youth. In contrast, older
238 populations in prior studies might have had fewer risk factors or benefited from improved
239 perioperative care, resulting in lower complication rates age.

240 In this study, 87.2% of patients with POP were male compared to 65% in Allou et al.'s
241 study. The higher proportion of male patients with POP compared to females suggests that
242 gender may play a role in the increased incidence of pneumonia following surgery. Male
243 patients also often have higher rates of smoking, which can impair lung function and increase
244 vulnerability to respiratory infections.²¹ In this study, 75.6% of patients with POP had a
245 smoking history compared with 20% in the previous study. This behavior is strongly linked to
246 increased risks of postoperative complications, including pneumonia.

247 This study found a lower occurrence of COPD among POP patients compared to Allou's
248 study; however, missing data exceeding 5% rendered it inappropriate for multivariate analysis.
249 This limitation restricts the ability to fully evaluate the impact of COPD on POP rates within
250 this study. However, the univariate analysis indicated a COPD prevalence of 13.3% among
251 POP patients, with an OR of 3.846 (95% CI: 0.736–20.111, $p = 0.139$). Although not
252 statistically significant, these findings suggest that COPD remains a relevant factor influencing
253 POP risk. However, the inability to analyze COPD data in the multivariate model underscores
254 the challenge of accurately assessing its independent contribution to POP in this cohort.

255 In this study, neither the neutrophil-to-lymphocyte ratio (NLR) nor the red cell
256 distribution width (RDW) significantly influenced postoperative pneumonia (POP). Univariate
257 analysis revealed median NLR values of 2.02 in POP patients and 2.42 in non-POP patients (p
258 = 0.126). Similarly, the median RDW-CV was 13.20 in POP patients and 13.10 in non-POP
259 patients ($p = 0.244$). These findings indicate that NLR and RDW failed to differentiate between
260 POP and non-POP patients in this cohort. NLR and RDW are nonspecific markers of systemic
261 inflammation, influenced by factors like infections, comorbidities, and perioperative stress.
262 The lack of significant differences in comorbidities between POP and non-POP groups limits
263 variability in baseline inflammation, reducing NLR and RDW's predictive power.

264 Left Ventricular Ejection Fraction of less than 40% is a widely recognized indicator
265 of heart failure with reduced ejection fraction (HFrEF), which is significantly correlated with
266 postoperative complications, particularly pulmonary infections. Research conducted by
267 Hosseini et al. revealed an odds ratio (OR) of 2.95 (95% CI: 1.2–7.6) for pneumonia among
268 patients exhibiting LVEF below 40%, thereby underscoring its predictive value strength.²² In
269 a similar vein, Pieri et al. identified LVEF <40% as a significant predictor of complications
270 following cardiac procedures surgery.²³ In this study, however, LVEF was not significantly
271 correlated with POP occurrence. The median LVEF was comparable between POP and non-

272 POP groups: 49% (38–59.25%) vs. 53% (38–60%) ($p = 0.699$). The prevalence of LVEF $\leq 40\%$
273 was nearly identical (30.2% vs. 30.8%, OR = 0.972, $p = 0.926$). Unlike prior studies with higher
274 proportions of patients with severe heart failure, such as Allou et al., this cohort had a moderate
275 proportion of reduced LVEF cases (30.6%), with most patients presenting LVEF $>40\%$
276 (69.4%). This discrepancy highlights cohort variations, where fewer reduced heart failure cases
277 may have weakened the statistical power to confirm LVEF $<40\%$ as an independent risk factor
278 for POP. More research with higher reduced heart failure proportions is needed to clarify its
279 predictive value.

280 This study found no significant correlation between mechanical ventilation duration
281 and POP occurrence, unlike previous studies that identified prolonged ventilation as a major
282 predictor, particularly for VAP. The lack of significance might be due to shorter ventilation in
283 this cohort or improvements in respiratory therapy and early extubation. Larger studies with
284 clinically relevant ventilation thresholds are needed to clarify this impact on POP.

285 Patients who developed POP had a substantially higher likelihood of requiring
286 extended hospitalization. Specifically, 84.9% of POP patients had a hospital stay longer than 6
287 days, compared to only 30.0% of non-POP patients. The odds ratio (OR) for extended LOS
288 among POP patients in the final analysis was 13.043 (95% CI: 6.130–27.751, $p < 0.001$),
289 indicating a strong association between the occurrence of POP and longer hospital stays.

290

291 **5. Study Limitations**

292 This study has several limitations. The lack of data on COPD over 5% prevented its
293 inclusion in the multivariate analysis and may have affected the assessment of its link to POP.
294 Additionally, inadequate preoperative albumin data hindered analysis of delta albumin. The
295 single-center design and small sample size limit the generalizability of the findings.
296 Unmeasured confounding factors, such as infection control practices, intraoperative

297 transfusions, provider performance, and postoperative care quality, were not addressed and
298 might have influenced POP incidence. Future research should further explore these variables
299 using multi-center designs to validate and enhance these findings.

300

301 **6. Conclusion**

302 This study identifies renal impairment and hypoalbuminemia as key risk factors for POP
303 after isolated CABG, providing insights for perioperative risk stratification. It emphasizes the
304 importance of institutional practices and patient-specific factors, highlighting the need for
305 tailored interventions to optimize outcomes. Additionally, it lays a groundwork for future
306 research to explore unmeasured variables and improve preventive strategies, thus enhancing
307 care for CABG patients.

308

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385 TABLE LEGENDS

386 Table 1. Baseline Characteristics of Patients

387 Table 2. Univariate analysis of risk factors for POP

388 Table 3. Multivariate analysis of risk factors of POP

389
 390 **Table 1. Baseline Characteristics of Patients**

Variable	N=206
Age	
Mean±Std	57.76±8.485
Age Category	
>=60 y.o	92(44.7%)
<60 y.o	114(55.3%)
Sex	

Male	171(83.0%)
Female	35(17.0%)
BMI	
Median	24.70
IQR (Q1-Q3)	4.34(22.86-27.20)
BMI Category	
>22.9	151(73.3%)
<22.9	55(26.7%)
Risk Factors	
DM	56(27.2%)
Hypertension	123(59.7%)
Dyslipidemia	75(36.4%)
Smoking	147(71.4%)
COPD (n=97)	8(8.2%)
PAD	89(43.2%)
LVEF	
Median	50.00
Range (min-max)	22.00(38.00-60.00)
LVEF Category	
<=40%	63(30.6%)
>40%	143(69.4%)
Hemoglobin	
Mean±Std	13.80±1.799
WBC	
Median	7925.00
IQR (Q1-Q3)	2227.50(6932.50-9160.00)
Platelet	
Median	260000.00
IQR (Q1-Q3)	83500.00(220500.00-304000.00)
Creatinin	
Median	1.10
IQR (Q1-Q3)	0.40(0.93-1.33)
eGFR	
Mean±Std	73.51±23.103
eGFR Category	
<60	50(24.3%)
>60	156(75.7%)
Early Postoperative Albumin	

Mean±Std	2.93±0.568
Early Postoperative Albumin	
≤2.93 g/dL	119(57.8%)
>2.93 g/dL	87(42.2%)
NLR	
Median	2.15
IQR (Q1-Q3)	1.52(1.67-3.18)
RDW-CV	
Median	13.15
IQR (Q1-Q3)	1.20(12.70-13.90)
RDW-SD	
Median	41.50
IQR (Q1-Q3)	3.93(39.88-43.80)
Clinical presentation preoperative	
ACS	40(19.4%)
CCS	166(80.6%)
Pump CABG	
On pump	77(37.4%)
Off pump	129(62.6%)
CPB Time	
Median	0.00
IQR (Q1-Q3)	81.25(0.00-81.25)
CPB Time Category	
>92 minutes	36(17.5%)
<92 minutes	170(82.5%)
Mechanical Ventilaton Duration	
Median	0.00
IQR (Q1-Q3)	1.00(0.00-1.00)
Hospital LOS	
Median	7.00
IQR (Q1-Q3)	4.00(5.00-9.00)
Complication	
Dengan POP	86(41.7%)
Tanpa POP	120(58.3%)
Mortality	
Yes	10(4.9%)
No	196(95.1%)

Descriptions: SD: Standard Deviation; BMI: Body-Mass Index; CABG: Coronary Artery Bypass Grafting; COPD: Chronic Obstructive Pulmonary Disease; eGFR: estimated Glomerular Filtration Rate; LOS: Length of Stay; LVEF: Left Ventricle Ejection Fraction; PAD: Peripheral Artery Disease; POP: Postoperative Pneumonia; WBC: White Blood Cell

391 **Table 2. Univariate analysis of risk factors for POP**

Variable	Complication		OR CI 95%	Nilai P
	With POP N=86	Without POP N=120		
Age				0.446
Mean±Std	58.29±7.633	57.38±9.059		
Age Category			1.049	0.866
≥60 y.o	39(45.3%)	53(44.2%)	(0.601-1.831)	
<60 y.o	47(54.7%)	67(55.8%)		
Sex			1.705	0.174
Male	75(87.2%)	96(80.0%)	(0.785-3.700)	
Female	11(12.8%)	24(20.0%)		
BMI				0.630
Median	25.37	24.38		
IQR (Q1-Q3)	4.60(22.74-27.34)	4.09(22.89-26.99)		
BMI Category			0.996	0.990
>22.9	63(73.3%)	88(73.3%)	(0.533-1.862)	
<22.9	23(26.7%)	32(26.7%)		
Risk factors				
DM	21(24.4%)	35(29.2%)	0.785	0.450
			(0.418-1.473)	
Hypertension	52(60.5%)	71(59.2%)	1.056	0.851
			(0.600-1.857)	
Dyslipidemia	35(40.7%)	40(33.3%)	1.373	0.279
			(0.773-2.436)	
Smoking	65(75.6%)	82(68.3%)	1.434	0.256
			(0.768-2.678)	
COPD (n=97)	6(13.3%)	2(3.8%)	3.846	0.139
			(0.736-20.111)	
Peripheral Artery Disease	37(43.0%)	52(43.3%)	0.987	0.965
			(0.565-1.727)	
LVEF				0.699
Median	49.00	53.00		
Range (min-max)	21.25(38.00-59.25)	22.00(38.00-60.00)		

LVEF			0.972	0.926
Category			(0.533-1.774)	
<=40%	26(30.2%)	37(30.8%)		
>40%	60(69.8%)	83(69.2%)		
Hemoglobin				0.662
Mean±Std	13.74±1.810	13.85±1.798		
WBC				0.831
Median	7950.00	7805.00		
IQR (Q1-Q3)	2317.50(6812.50-9130.00)	2242.50(6945.00-9187.50)		
Platelet				0.578
Median	263000.00	256500.00		
IQR (Q1-Q3)	76500.00(220500.00-297000.00)	97750.00(219000.00-316750.00)		
Creatinin				0.186
Median	1.15	1.10		
IQR (Q1-Q3)	0.52(0.95-1.47)	0.34(0.92-1.26)		
eGFR				0.281
Mean±Std	71.45±22.681	74.98±23.383		
eGFR			2.150	0.019
Category			(1.127-4.103)	
<60	28(32.6%)	22(18.3%)		
>60	58(67.4%)	98(81.7%)		
Early Postoperative Albumin				<0.001
Mean±Std	2.73±0.488	3.07±0.580		
Early Postoperative Albumin			3.438	<0.001
			(1.881-6.283)	
<=2.93 g/dL	64(74.4%)	55(45.8%)		
>2.93 g/dL	22(25.6%)	65(54.2%)		
NLR				0.126
Median	2.02	2.42		
IQR (Q1-Q3)	1.46(1.62-3.08)	1.48(1.71-3.19)		
RDW-CV				0.244
Median	13.20	13.10		
IQR (Q1-Q3)	1.30(12.70-14.00)	0.88(12.70-13.58)		
RDW-SD				0.569

Median	41.55	41.50		
IQR (Q1-Q3)	4.95(39.05-44.00)	3.60(40.00-43.60)		
Clinical Presentation preoperative			2.869 (1.405-5.857)	0.003
ACS	25(29.1%)	15(12.5%)		
CCS	61(70.9%)	105(87.5%)		
Pump CABG			1.387 (0.784-2.455)	0.260
On pump	36(41.9%)	41(34.2%)		
Off pump	50(58.1%)	79(65.8%)		
CPB Time				0.077
Mean±Std	42.79±53.303	29.37±42.961		
Median	0.00	0.00		
IQR (Q1-Q3)	94.75(0.00-94.75)	77.75(0.00-77.75)		
CPB Time category			3.005 (1.422-6.348)	0.003*
>92 minutes	23(26.7%)	13(10.8%)		
<92 minutes	63(73.3%)	107(89.2%)		
Mechanical Ventilaton Duration				0.083
Median	0.00	0.00		
IQR (Q1-Q3)	1.00(0.00-1.00)	1.00(0.00-1.00)		
Hospital LOS				<0.001
Median	8.00	5.50		
IQR (Q1-Q3)	6.00(7.00-13.00)	2.00(5.00-7.00)		
Hospital LOS Category			13.103 (6.458-26.584)	<0.001
>6 days	73(84.9%)	36(30.0%)		
<6 days	13(15.1%)	84(70.0%)		
Mortality			2.175 (0.595-7.955)	0.326
Yes	6(7.0%)	4(3.3%)		
No	80(93.0%)	116(96.7%)		

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393 **Table 3. Multivariate analysis of risk factors of POP**

	B	S.E.	Wald	Nilai P	OR	CI 95%	
						Lower	Upper

INITIAL MODEL	Sex	0.805	0.575	1.962	0.161	2.238	0.725	6.905	
	Creatinin	1.429	0.712	4.024	0.045	4.176	1.033	16.873	
	eGFR Category	2.131	0.722	8.720	0.003	8.427	2.048	34.682	
	Early Postoperative Albumin Category	1.111	0.388	8.191	0.004	3.037	1.419	6.501	
	NLR	0.176	0.114	2.386	0.122	1.192	0.954	1.491	
	RDW-CV	0.007	0.151	0.002	0.961	1.007	0.749	1.355	
	Clinical Presentation Preoperative	0.247	0.465	0.283	0.595	1.280	0.515	3.185	
	CPB Time Category	0.527	0.485	1.182	0.277	1.694	0.655	4.384	
	Mechanical Ventilation Duration	0.018	0.144	0.016	0.899	1.019	0.767	1.352	
	Hospital LOS Category	2.427	0.412	34.669	<0.00 1	11.320	5.047	25.388	
	FINAL MODEL	Creatinin	1.357	0.670	4.101	0.043	3.883	1.045	14.434
		eGFR Category	1.937	0.669	8.379	0.004	6.935	1.869	25.734
		Early Postoperative Albumin Category	1.168	0.375	9.713	0.002	3.215	1.542	6.700
		Hospital LOS Category	2.568	0.385	44.450	< 0.00 1	13.043	6.130	27.751

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Pre