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Characteristics of D-Dimer Levels in Patients Confirmed Positive for COVID-19 Aged 45 – 70 Years at UKI Hospital for the Period December 2020 – September 2021

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Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

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ABSTRACT

COVID-19 is a disease caused by the SARS-CoV-2 virus. COVID-19 is divided into several clinical symptoms based on a person's experiences. In severe clinical signs of COVID-19, D-Dimer will be significantly higher. This study aims to determine D-Dimer levels in COVID-19 patients aged 45 – 70 and uses a descriptive-analytic method with a retrospective cross-sectional design. The number of samples was 198 patients whose D-Dimer levels were checked and 127 patients who received

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anticoagulant prophylaxis. The research samples were all medical records of patients suffering from COVID-19 aged 45 - 70 years and who had checked D-Dimer levels at RSU UKI from December 2020 to September 2021 found that 145 patients (73.2%) had normal D-Dimer levels, 53 patients (26.8%) had increased D-Dimer levels. The distribution of women were 100 patients (50.5%), and men were 98 patients (49.5%), the highest range was aged 54 - 62 years with 74 patients (37.4%), clinical symptoms were being experienced by 122 patients (61.6%), and comorbid hypertension was the most common disease with 64 patients (32.3%). A significant relationship was found between D-Dimer levels and comorbidities ($p=0.04$). Anticoagulant prophylaxis was given to 127 patients, with mild symptoms experienced by five patients (3.6%), moderate symptoms experienced by 73 patients (57.5%), and severe symptoms experienced by 49 patients (38.6%). However, a significant relationship was found ($p=0.66$) between administering anticoagulant prophylaxis and D-Dimer levels in patients with mild, moderate, or severe symptoms.

Keywords: D-dimer; age; covid-19; anticoagulants.

1. INTRODUCTION

Currently, people around the world are experiencing difficulties due to epidemics that arise and spread rapidly. This incident began at the end of 2019 when China reported a mysterious pneumonia infection to WHO [1]. Were 7,736 cases recorded in China on January 1, 2020, and they have spread worldwide [2]. The rapid development of the virus, called Coronavirus Disease 2019 (COVID-19) by WHO, is triggered by transmission from humans to other humans through droplets that come out when coughing or sneezing [3].

When someone is infected with COVID-19, that person will show symptoms of fever and at least signs of respiratory illness, such as coughing and shortness of breath, and require hospitalization. Diagnosis is made based on the results in NAAT, the recommended method is a Real-Time Reverse-Transcription Polymerase Chain Reaction [4]. In addition to NAAT, hematological examination is also essential for disease progression. In hematological analysis, the tests that can be carried out are Prothombin Time, Activated Partial Thromboplastin Time, and D-Dimer [5].

In patients with severe clinical symptoms, D-Dimer will be significantly higher. Increases in D-Dimer levels occur with age and during pregnancy. D-dimer levels themselves are related to the severity of the disease. Currently, D-dimer examination is used as a prognostic biomarker in COVID-19 patients [6]. Diseases such as diabetes, cancer, and stroke, which are common in elderly patients, can also trigger D-Dimer levels in COVID-19 patients [7]. Several studies suggest that COVID-19 makes patients more susceptible to arterial and venous

thrombosis. Measuring D-dimer levels and coagulation parameters in the early stages of the disease can help control and manage COVID-19 [8].

In COVID-19 patients with moderate, severe, and critical symptoms, administering anticoagulants is highly recommended as prophylaxis and treating macrothrombi [9]. Apart from that, anticoagulants can also be given to COVID-19 patients with mild symptoms after evaluating the patient's D-Dimer levels.

Based on this, researchers are interested in researching the profile of D-Dimer levels in patients confirmed positive for COVID-19 by RT-PCR examination at 45 - 70 years old at the Indonesian Christian University General Hospital. The problem in this research is: What is the profile of D-Dimer levels in patients confirmed positive for COVID-19 by RT-PCR examination aged 45 - 70 years at the UKI General Hospital from December 2020 - September 2021? The research aims to determine D-Dimer levels in patients confirmed positive for COVID-19.

Coronavirus-Coronavirus is a single positive strand RNA virus encapsulated and not segmented [10]. Coronavirus has four main protein structures, namely, protein N (nucleocapsid), glycoprotein M (membrane), glycoprotein spike S (spike), and protein E (sheath) [11]. Coronaviruses belong to the order Nidovirales and the family Coronaviridae. There are four genera in the coronaviridae family, namely alphacoronavirus, betacoronavirus, gammacoronavirus, and deltacoronavirus [12].

Coronavirus has been around since the 1930s and is known to occur in animals. Extraordinary events occurred in 2002 and 2012 caused by

betacoronavirus, namely SARS, which first appeared in China, and MERS, which first appeared in Saudi Arabia [13]. At the end of 2019, a new coronavirus was detected and classified by WHO as novel coronavirus 2019 (nCoV-2019), which CREVSG later changed to Severe Acute Respiratory Syndrome Coronavirus 2. SARS-CoV-2 belongs to the betacoronavirus genus and the disease it causes has been named COVID-19 by WHO [14].

Coronavirus Disease 2019 is caused by a new coronavirus first identified in Wuhan, China. COVID-19 was not initially identified as a coronavirus infection but rather a sudden occurrence of pneumonia of unknown cause. After carrying out a genome sequencing examination, it was found that the cause was SARS-CoV-2 [13]. The first COVID-19 cluster is known to be associated with the Huanan Seafood Market in Wuhan, China. The SARS-CoV-2 virus is zoonotic, where the pathogen transfers from animals, in this case bats sold at the Huanan Seafood Market, to humans [15]. Like SARS and MERS, the symptoms that appear in COVID-19 patients are similar to pneumonia caused by viruses, namely fever, dry cough, difficulty breathing, and several additional symptoms such as headache, vomiting, weakness, and diarrhea [16]. COVID-19 has spread throughout the world and more than 219,000,000 people in the world are positive for COVID-19 [13].

SARS-CoV-2 is a single-stranded RNA virus measuring 29.9 kb with an envelope containing a nucleocapsid. 10,12 Coronaviruses are divided into four major classifications, namely, α , β , γ , and δ . SARS-CoV, MERS-CoV, and SARS-CoV-2 are classified as betacoronaviruses, which can only infect mammals [17]. Coronavirus has four primary proteins, namely, protein N (nucleocapsid), glycoprotein M (membrane), glycoprotein spike S (spike), and protein E (sheath), which have different functions [11].

The spike protein is a transmembrane trimetric glycoprotein that protrudes from the virus's surface. Two functional subunits form the spike; the S1 subunit is responsible for binding to host cell receptors, and the S2 subunit is responsible for binding to viruses and cell membranes. ACE2 in the lower respiratory tract has been identified as a functional receptor for the SARS-CoV and MERS-CoV spike glycoproteins. Structural and functional analysis revealed that the genetic sequence, almost the same as SARS-CoV,

means that the SARS-CoV-2 spike protein can also bind to ACE2 to infect humans. The SARS-CoV-2 spike binds to the human ACE2 receptor with approximately ten to twenty times higher affinity than the SARS-CoV spike, allowing the virus to spread more easily from person to person. Once it enters the alveolar epithelium, SARS-CoV-2 replicates rapidly, eliciting a strong immune response and causing a cytokine storm and lung tissue damage. In addition, in COVID-19 patients, the total number of T cells, CD4+ T cells, and CD8+ T cells decreases and causes a decrease in immune function, which will worsen the patient's condition [13;17].

The SARS-CoV-2 virus has a high level of pathogenicity and transmissibility. SARS-CoV-2 can be transmitted from humans to other humans through direct contact with sufferers, via droplets, and indirectly through objects or air contamination. [18] Droplets released by sufferers through coughing, sneezing, singing, or talking can be inhaled or stick to the mucous membranes of the eyes, nose, or mouth and cause a person to be infected with the SARS-CoV-2 virus [13,18,19]. Contamination of objects by droplets can also transmit COVID-19.

The incubation period for the SARS-CoV-2 virus is one to fourteen days. In general, COVID-19 is detected on the third to seventh day [20]. According to Hidayani WR (2020), there are several risk factors associated with COVID-19, including:

A person experiences symptoms when sick with COVID-19, ranging from asymptomatic or no signs, mild and moderate symptoms, to clinical conditions requiring mechanical ventilation and support in the ICU [21]. Mild symptoms are often defined as signs of acute upper respiratory tract infection without complications and may be accompanied by fever, fatigue, cough (with or without sputum), anorexia, malaise, sore throat, nasal congestion, or headache. COVID-19 patients with severe pneumonia are characterized by fever and one of the following symptoms: (1) respiratory rate of more than thirty breaths per minute, (2) severe shortness of breath, or (3) oxygen saturation of 93% without oxygen support [2].

Low lymphocyte counts and low monocyte, basophil, and eosinophil results are found in severe clinical cases. In addition, proinflammatory mediators (TNF- α , IL1, IL6, and IL 8) increased, but helper T cells, suppressor T

cells, and regulatory T cells decreased [22]. X-rays of COVID-19 patients often show multiple ground glass shadows and infiltration shadows in both lungs. In severe cases, lung consolidation may occur. Apart from the respiratory organs, SARS-CoV-2 can cause disorders of the cardiovascular and gastrointestinal systems and acute kidney failure [18].

The normal mechanism carried out by the body in the process of stopping bleeding at the site of injury is called hemostasis [23]. The hemostasis stage or process can be divided into three main steps, namely (1) vascular vasoconstriction, (2) formation of platelet plugs (primary hemostasis), and (3) blood clotting (secondary hemostasis). The hemostasis process is balanced by blood clotting control mechanisms and the fibrinolysis process (tertiary hemostasis) [24].

In the blood coagulation (clotting) process, important substances can influence clotting, such as procoagulants, which function as substances that make clotting easier, and anticoagulants, which function as clotting inhibitors. When blood vessels are damaged, procoagulants are more active so that wounds can close [23]. The coagulation theory commonly used today, proposed by Mac Farlane, Davie, and Ratnoff, is a cascade or waterfall theory to describe the coagulation process [24].

The cascade theory explains that each coagulation factor is a proenzyme converted into an active enzyme, where some of these factors act as co-factors and some act as coagulant factors. The nomenclature of coagulation factors is a nomenclature that uses Roman numerals in the order in which they are found. For writing, fibrinogen (factor I), Prothrombin (factor II), tissue thromboplastin (factor III), Ca ions (factor IV), prekallikrein (PK), and HMWK are usually not written in Roman numerals [23,24]. Factor V, Factor VII, HMWK, and thromboplastin are co-factors for other factors. Factor XII, Factor XI, Factor [24]. In the cascade theory, the blood clotting process is divided into two pathways: the intrinsic pathway, where all components are present in the blood, and the intrinsic pathway, which involves subendothelial membranes and tissue factors [24,25].

Contact factors (factors XII, Exposure to FXII (Hageman Factor) on foreign surfaces converts XII to FXIIa. FXIIa converts prekallikrein to kallikrein, which in turn increases FXII activation. Simultaneously, kallikrein activates FVII to FVIIa in the extrinsic pathway, activates plasminogen to plasmin in the fibrinolytic system, and converts

Kininogen to Kinin, which plays a role in the inflammatory response. FXIIa, with the help of the HMWK co-factor, divides FXI into FXIa, and then FIXa divides FIX into IXa. FIXa, FXa, and thrombin cleave FVIII to form FVIIIa. Finally, FIXa, FVIIIa, calcium ions, and negatively charged phospholipids form a three-molecule complex called tenase. Tenase then changes FX to FXa. FXa binds to the co-factor FVa to form a prothrombinase complex. This complex converts the proenzyme prothrombin into the enzyme thrombin. Thrombin converts fibrinogen into fibrin monomers, rapidly polymerizing to form fibrin clots [24,25].

The injured vascular endothelium is the origin of the extrinsic pathway. FVII comes into contact with extravascular tissue factor and forms a tissue factor-FVIIa complex where FVII activation is carried out by tissue factor in a non-proteolytic manner. The binding between tissue factors FVIIa forms an enzyme complex that can convert FX to FXa. FXa formed in the extrinsic pathway functions almost the same as tenase in the intrinsic pathway, namely binding to the FV co-factor and ultimately converting fibrinogen into fibrin. The cascade will continue along a common pathway with FXa, formed from intrinsic and extrinsic pathways [25].

FX activation begins a common pathway through intrinsic, extrinsic, or both pathways. FXa is assisted by FV, calcium, and phospholipid ions to convert prothrombin into thrombin. The main function of thrombin is to catalyze the proteolysis of plasma-dissolved fibrinogen into soluble fibrin monomers. Monomeric fibrin then polymerizes into polymer fibrin, which will defend blood cells. In addition, thrombin can convert FXIII to FXIIIa and mediate cross-linking of fibrin polymers to form slightly soluble and stable fibrin. Thrombin can also catalyze the formation of co-factors FVa and FVIIIa, leading to increased coagulation [25]. The cascade coagulation theory is illustrated in Fig. 1.

D-Dimer- The D-Dimer test determines whether there are blood clots in the body. The D-Dimer test is generally used as an additional test to rule out the diagnosis of DVT and pulmonary embolism and confirm the diagnosis of DIC. [26] D-Dimer itself is the final product of the fibrinolytic process in the form of fibrin degradation. [26] Fibrinolysis is a physiological mechanism that destroys fibrin deposits with a fibrinolytic system that continuously works through the coagulation system to ensure

smooth blood flow to body tissues or peripheral organs [23,24]. The fibrinolytic system is activated by the presence of a plasminogen activator, which breaks down plasminogen into plasmin. Active plasmin is divided into free plasmin, which is neutralized by antiplasmin, and fibrin-bound plasmin [24]. Fibrin-bound plasmin impairs fibrin formation by inducing FDP. Initially, an X piece will be formed. Piece X will be divided into pieces Y, D, and E [24,27]. Two pieces of D and one piece of E will bond and form a D-Dimer [27,28].

In testing D-Dimer concentration, the principle used is monoclonal antibodies. [26] There are several testing methods to check D-Dimer levels, namely ELISA, Latex Agglutination, Whole Blood Agglutination, immunoturbidimetry, and immunofiltration. The gold standard in D-Dimer testing is testing using the ELISA method [27].

Standardization of normal D-Dimer values does not exist to date. However, the D-Dimer levels commonly used range from 0 – 300 ng/mL. Increased D-dimer levels occur with age and during pregnancy [29]. In addition, increased D-Dimer levels also occur in CAP and COPD patients [29;30]. The severity of the disease is related to D-Dimer levels; currently, D-Dimer examination is used as a prognostic biomarker in COVID-19 patients. [30] In the early stages of COVID-19 disease, D-Dimer and fibrinogen levels increase, which leads to a poor prognosis. D-Dimer greater than 1 µg/mL is a risk factor for death in COVID-19 patients [30].

Diseases such as diabetes, cancer, and stroke can also increase D-Dimer levels in COVID-19 patients. Several studies show that COVID-19 makes patients more susceptible to arterial and venous thrombosis. Measuring D-Dimer levels and coagulation parameters in the early stages of the disease can be useful in controlling and managing COVID-19 [31].

Anticoagulants are substances used to prevent blood clotting by inhibiting certain blood clotting factors or inhibiting the formation of thrombin, which is needed to convert fibrinogen into fibrin [32]. In severe COVID-19 symptoms, a significant increase in D-Dimer levels was found, which could occur due to massive inflammation and impaired blood coagulation. Elevated D-Dimer levels can be a marker of thromboembolic events that occur in COVID-19 patients. Giving anticoagulants to COVID-19 patients can be a solution in preventing the formation of

microvascular and macrovascular thrombi which can cause vital organ dysfunction. The International Society of Thrombosis and Haemostasis recommends administering prophylactic doses of LMWH to all COVID-19 patients, except those with contraindications and high bleeding activity. In moderate, severe, and critical COVID-19 patients, administering LMWH or UFH anticoagulants is highly recommended, both useful as prophylaxis and in treating macrothrombi [33]. Apart from that, anticoagulants can also be given to COVID-19 patients with mild symptoms after evaluating the patient's D-Dimer levels [34].

2. METHODS

This research type is analytical descriptive research with data collected retrospectively cross-sectionally from secondary data obtained from the Medical Records Archives Section of RSU UKI. The research was conducted in the medical records section of RSU UKI from December 2021 to January 2022. The population in this study were COVID-19 sufferers who were treated and had their D-Dimer levels checked. The research samples were COVID-19 patients who were treated and had their D-Dimer levels checked and recorded in the medical record at RSU UKI during the treatment period from December 2020 to September 2021. The sample size in the study was calculated using the Slovin formula: The sample size required in this research was 154 patients. This research used research instruments in the form of medical records obtained from the Medical Records Archives Section of RSU UKI from December 2020 to September 2021.

The data from the data collection process will be displayed in descriptive tables and analyzed using the SPSS application with the Chi-Square correlation method or Fisher test on computer software. The data processing process by the SPSS program consists of several steps: a) Coding, to translate the data collected during the research into symbols suitable for analysis; b) Data entry, data entry into the computer; c) Verification, visually checking the data that has been entered into the computer; d) Computer output, prints the results of the analysis carried out by the computer in the previous step.

3. RESULTS AND DISCUSSION

In this study, data collection was carried out from December 18, 2021 to January 15, 2022. The data collected was secondary in the form of

medical records of patients confirmed positive for COVID-19 who underwent inpatient treatment at RSU UKI from December 2020 to September 2021. Of 689 confirmed positive for COVID-19 patients, 198 research samples met the inclusion criteria. Of the 198 samples, 127 samples were given anticoagulant prophylaxis therapy. Data processing was carried out using the SPSS version 22 application. Research data is presented in the form of tables and diagrams. Of the 198 samples obtained in the study, there were 145 people (73.2%) with normal D-Dimer levels, namely <500 ng/dl, and 53 people (26.8%) with D-Dimer levels >500 ng/dl (increase).

3.1 Sample Characteristics Based on Age

The age group most exposed to COVID-19 at RSU UKI from December 2020 to September 2021 was 54 - 62, namely 74 people (37.4%), followed by 63 - 70 years old, 66 people (32.3%). and aged 45 - 53 years, as many as 60 people (30.3%).

3.2 Sample Characteristics Based on Gender

The research results on patients confirmed positive for COVID-19 aged 45 - 70 years found that 100 patients were female (50.5%) and 98 were male (49.5%).

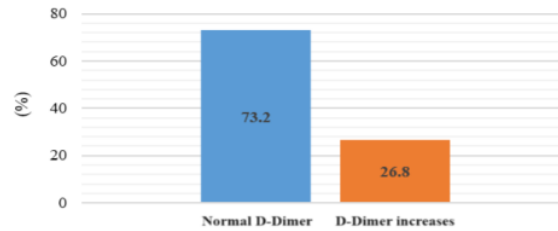


Fig. 1. Sample characteristics based on D-dimer levels

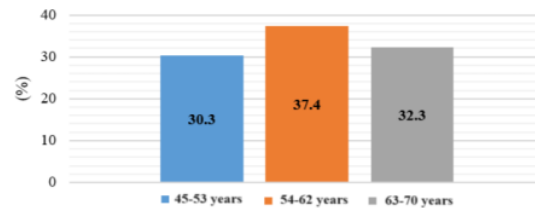


Fig. 2. Sample characteristics based on age

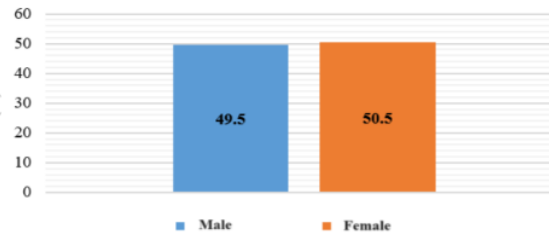


Fig. 3. Sample Characteristics Based on Gender

3.3 Sample Characteristics Based on Clinical Symptoms

The most clinical symptoms experienced by COVID-19 patients at RSU UKI were moderate clinical symptoms in as many as 122 people (61.6%), followed by severe clinical symptoms in as many as 60 people (30.3%) and mild clinical symptoms in as many as 16 people (8.1%).

3.4 Characteristics of Clinical Symptoms Based on D-Dimer Levels

Of the 122 people (61.6%) who experienced moderate symptoms of COVID-19, 96 people (48.5%) had normal D-Dimer levels, and 26 people (13.1%) had increased D-Dimer levels.

Furthermore, of the 60 people (30.3%) who experienced severe symptoms of COVID-19, 37 people (18.7%) had normal D-Dimer levels, and increased D-Dimer levels were experienced by 23 people (11.6%). Next, of the 16 people (8.1%) who experienced mild symptoms of COVID-19, 12 people (6.1%) had normal D-Dimer levels, and four people (2%) had increased D-Dimer levels.

3.5 Sample Characteristics Based on Concomitant Diseases

The most common comorbidity experienced by patients at RSU UKI was hypertension, 64 people (32.3%).

Table 1. Sample characteristics based on clinical symptoms

Clinical Symptoms	Total(n)	Percentage(%)
COVID-19 Mild Symptoms	16	8,1
COVID-19 Moderate Symptoms	122	61,6
COVID-19 Severe Symptoms	60	30,3
Total	198	100

Table 2. Characteristics of Clinical Symptoms Based on D-Dimer Levels

			D-Dimer results (ng/dl)		Total
			D-Dimer Normal	D-Dimer Increases	
Degree of COVID-19	COVID-19 Mild Symptoms	Total	12	4	16
		Percentage	6,1%	2%	8,1%
	Covid-19 Moderate Symptoms	Total	96	26	122
		Percentage	48,5%	13,1%	61,6%
	Covid-19 Severe Symptoms	Total	37	23	60
		Percentage	18,7%	11,6%	30,3%
Total		Total	145	53	198
		Percentage	73,2%	26,8%	100%

Table 3. Sample Characteristics Based on Concomitant Diseases

Concomitant Diseases	Total (n)	Percentage(%)
Hypertension	64	32,3
DM Type 2	20	10,1
Hypertension and Type 2 DM	20	10,1
Hypertension and heart problems	9	4,5
Hypertension and CKD	5	2,5
Type 2 DM and heart problems	3	1,5
Hypertension, Type 2 DM, and ACKD	3	1,5
Heart problems	3	1,5
DM Type 2 and ACKD	2	1
Hypertension, Type 2 DM, ACKD, and heart problems	2	1
Hypertension, Type 2 DM, and CKD	2	1
Hypertension, Type 2 DM, and heart problems	2	1
COPD	2	1
CA Nasopharynx	1	0,5

Concomitant Diseases	Total (n)	Percentage(%)
CKD	1	0,5
Type 2 DM, ACKD, and heart problems	1	0,5
Type 2 DM and COPD	1	0,5
Drug eruption	1	0,5
Hypertension and ACKD	1	0,5
Hypertension and mammary CA	1	0,5
Hypertension and Dementia	1	0,5
Hypertension, Type 2 DM and Bronchial Asthma	1	0,5
Hypertension, Type 2 DM and CA colon	1	0,5
Hypertension, Type 2 DM, CKD, and heart problems	1	0,5
Hypertension, Type 2 DM, CKD, and Stroke	1	0,5
Hypertension and Parkinson's	1	0,5
Hypertension and COPD	1	0,5
Hypertension and stroke	1	0,5
Hypertension and transaminitis	1	0,5
Schizophrenia	1	0,5
Pulmonary TB	1	0,5
Transaminitis	1	0,5
No disease	42	21,2
Total	198	100

3.6 Relationship between Concomitant Diseases and D-Dimer Levels in COVID-19 Patients

Of the 156 people (78.8%) who had comorbidities, 47 people (23.7%) had increased D-Dimer levels, and 109 people (55.1%) had normal D-Dimer levels. Furthermore, of the 42 people (21.2%) who did not have comorbidities, six people (3%) had increased D-Dimer, and 36 people (18.2%) had normal D-Dimer levels.

3.7 Sample Characteristics Based on Administration of Anticoagulant Prophylaxis

In this study, it was found that prophylactic anticoagulant therapy was given to 127 COVID-19 patients, and when an evaluation was carried out by checking D-Dimer levels, it was found that 81 people (64%) had normal D-Dimer levels, and 46 people (36%) had D-Dimer levels. -Dimer increases.

3.8 Correlation between Anticoagulant Prophylaxis and D-Dimer Levels in Patients with Mild, Moderate, and Severe Clinical Symptoms

From the data collected, five people (3.6%) had mild clinical symptoms and were given anticoagulant prophylaxis, of which three people (2.3%) had normal D-Dimer levels, and

two people (1.6%) had D-Dimer levels. -Dimer increases. Followed by 73 people (57.5%) with moderate clinical symptoms and given anticoagulant prophylaxis, of which 49 people (38.6%) had normal D-Dimer levels, and 24 people (18.9%) had increased D-Dimer levels. Then, there were 49 people (38.6%) with severe clinical symptoms and were given anticoagulant prophylaxis, of which 29 people (22.9%) had normal D-Dimer levels, and 20 people (15.7%) had increased D-Dimer levels.

3.9 Characteristics of D-Dimer Levels in Patients who have Concomitant Diseases and are Receiving Anticoagulant Prophylaxis

The data showed that 107 patients had comorbidities and were given anticoagulant prophylaxis. It was found that 66 patients (62%) had normal D-Dimer levels, and 41 patients (38%) had increased D-Dimer levels.

Coronavirus Disease 2019 is caused by the SARS-CoV-2 virus, first identified in Wuhan, China. When the SARS-CoV-2 virus enters the human body, the virus will bind to the ACE2 receptor, causing a cytokine storm and damage to lung tissue [13]. In patients with severe clinical symptoms, D-Dimer will be significantly higher. D-dimer levels themselves are related to the severity of the disease, and currently, D-dimer examination is used as a prognostic biomarker in COVID-19 patients [30].

Table 4. Relationship between Concomitant Diseases and D-Dimer Levels in COVID-19 Patients

			D-Dimer Results (ng/dl)		Total	p-value
			D-Dimer Normal	D-Dimer Increases		
Concomitant Diseases	Yes	Total	109	47	156	0.040
		Percentage	55,1%	23,7%	78,8%	
	None	Total	36	6	42	
		Percentage	18,2%	3%	21,2%	
Total		Total	145	53	198	
		Percentage	73,2%	26,8%	100%	

From statistical analysis, $p = 0.04$ was obtained, which means a relationship exists between comorbidities and D-Dimer levels in COVID-19 patients

Table 5. Sample Characteristics Based on Anticoagulant Prophylaxis

D-Dimer Results(ng/dl)	n	%
D-Dimer Normal	81	64
D-Dimer Increases	46	36
Total	127	100

Table 6. Relationship between anticoagulant prophylaxis and D-dimer levels in patients with mild, moderate, and severe clinical symptoms

		D-Dimer Results(ng/dl)				Total		p-value
		D-Dimer Normal		D-Dimer Increases				
		(n)	(%)	(n)	(%)	(n)	(%)	
Giving Anticoagulant Prophylaxis to Patients with Clinical Symptoms	Mild	3	2,3	2	1,6	5	3,9	0.660
	Moderate	49	38,6	24	18,9	73	57,5	
	Severe	29	22,9	20	15,7	49	38,6	
Total		81	63.8	46	36.2	127	100	

From statistical analysis, $p = 0.66$ was obtained, meaning no relationship exists between administering anticoagulant prophylaxis and D-Dimer levels in COVID-19 patients with mild, moderate, and severe symptoms

Table 7. Characteristics of D-Dimer Levels in Patients Who Have Concomitant Diseases and Are Receiving Anticoagulant Prophylaxis

D-Dimer Results(ng/dl)	n	%
D-Dimer Normal	66	62
D-Dimer Increases	41	38
Total	107	100

The UKI RSU Laboratory determines the normal level of D-Dimer at <500 ng/dL. Of the 198 samples obtained in the study, 145 COVID-19 patients (73.2%) did not experience increased D-Dimer levels. Meanwhile, 53 patients (26.8%) experienced increased D-Dimer levels of >500 ng/dL. It is in line with research conducted by Djajalaksana et al [34]. in Malang, which stated that there was an increase in D-Dimer levels above normal limits in patients confirmed positive for COVID-19. D-Dimer is found if there is fibrin

degradation and is a specific indicator for fibrinolysis. Here increasing D-Dimer levels can be related to the severity of the disease [34]. Elevated D-Dimer is also the most characteristic finding in patients with COVID-19 and coagulopathy.

The age group with the most positive cases of COVID-19 was in the 54 – 62 year age range, totaling 74 people (37.4%). It aligns with research conducted by Khaerunisa et al. in

Bekasi, where the age most exposed to COVID-19 was 46 – 59 [35] Apart from that, in research conducted by Elviani et al. [36] in Palembang, it was found that the age range of 56 - 69 years was the second highest. People of all ages can be infected with COVID-19. However, an elderly person has many risk factors for being infected with the disease. One of the risks experienced is the deterioration of body cells, which triggers a decrease in the function of the body's tissues and organs, making it easier for disease to enter due to a decrease in immunity.

Based on gender, most COVID-19 sufferers at RSU UKI from December 2020 to September 2021 were women, namely 100 people (50.5%) and 98 people (49.5%) were men. It is in line with research conducted by Widjaja et al [37]. In Bandung, where the results obtained were that women were more exposed to COVID-19 than men. However, based on the literature, men are more susceptible to being infected with COVID-19. It is because men are more often outside the home due to work factors and are thought to be more active smokers than women. Smoking can increase the risk of exposure to COVID-19 because there is an increase in ACE2 receptors, making it easier for the SARS-CoV-2 virus to enter the body. The lower incidence of COVID-19 cases in women can be attributed to protection from sex hormones, progesterone, and the X chromosome, which provide innate and adaptive immunity.

In this study, the degree of clinical symptoms of COVID-19 that most patients experienced was moderate clinical symptoms with 122 people (61.6%), of which 96 people (48.5%) had normal D-Dimer levels, and 26 people (13.1%) had increased D-Dimer levels. Of the 26 people with elevated D-Dimer levels, 22 were given anticoagulant prophylaxis and comorbidities, and two were given anticoagulant prophylaxis and did not have comorbidities. Followed by severe clinical symptoms in 60 people (30.3%), of which 37 people (18.7%) had normal D-Dimer levels, and 23 people (11.6%) had increased D-Dimer levels. Of the 23 people with elevated D-Dimer levels, 20 were given anticoagulant prophylaxis and had comorbidities. Then, the clinical symptoms were mild in 16 people (8.1%), of which 12 people (6.1%) had normal D-Dimer levels, and four people (2%) had increased D-Dimer levels. Two of the four people who experienced increased D-Dimer levels were given anticoagulant prophylaxis and had comorbidities. Statistical analysis of the

relationship between administration of anticoagulant prophylaxis and D-Dimer levels in mild, moderate, and severe levels of clinical symptoms showed $p = 0.66$, which means there is no relationship between administration of anticoagulant prophylaxis and D-Dimer levels in mild and moderate clinical symptoms, and heavy.

This research shows that the most common comorbidity experienced by patients is hypertension, with a total of 64 people (32.3%). This study's results align with research by Berger et al. [38] in New York; hypertension is the most common comorbidity experienced by COVID-19 patients. The incidence of hypertension and COVID-19 is related to endothelial dysfunction and RAAS dysregulation. The Renin-Angiotensin-Aldosterone System itself is a system that controls blood pressure through aldosterone, angiotensin II, and angiotensin 1-7 with the help of the enzymes ACE1 and ACE2. When the SARS-CoV-2 virus enters the human body, the virus will bind to the ACE2 receptor, causing a decrease in the ACE2 enzyme in the RAAS and ultimately causing endothelial dysfunction, disruption of blood pressure regulation processes, and cytokine storms.

Based on the relationship between comorbidities and D-Dimer levels, of the 156 people (78.8%) who had comorbidities, 47 people (23.7%) had increased D-Dimer levels, and 109 people (55.1%) had normal D-Dimer levels. Elevated D-Dimer levels continued to occur in 42 people with comorbidities despite being given prophylactic anticoagulant therapy. Furthermore, of the 42 people (21.2%) who did not have comorbidities, six people (3%) had increased D-Dimer, and 36 people (18.2%) had normal D-Dimer levels. The statistical analysis showed that $p = 0.04$, which means a relationship exists between comorbidities and D-Dimer levels in COVID-19 patients.

In 107 patients who had comorbidities and were given anticoagulant prophylaxis, 66 people (42.3%) had normal D-Dimer levels, and 41 people (26.3%) had increased D-Dimer levels. In the study, it was found that as many as 127 people were given anticoagulant prophylaxis. It aligns with research by Pratiwi and Adhityasmara [39] in Semarang, which stated that anticoagulant therapy is recommended in treating COVID-19 patients because it can reduce the risk of death. Research conducted by Nugroho et al [40]. in Semarang also found that after giving anticoagulant therapy to COVID-19

patients for several days, there was a significant reduction in D-Dimer levels. The latest management guidelines recommend administering anticoagulant therapy to all patients who have or have not been proven to have thrombosis [39]. It can prevent the risk of thromboembolism in patients and the risk of death [40].

4. CONCLUSION

Based on the results of research conducted at the Medical Records of the Indonesian Christian University General Hospital, the following conclusions were obtained: a) 145 people (73.2%) had normal D-Dimer levels for the period December 2020 – September 2021 D-Dimer increased by 53 people (26.8%); b) The highest proportion of COVID-19 patients based on age were patients aged 54 – 62 years, 74 people (37.4%) and the lowest proportion were patients aged 45 – 53 years, 60 people (30.3%); c) There were 100 female COVID-19 patients (50.5%) and 98 male (49.5%) and d) Based on clinical symptoms, there were 122 COVID-19 patients with moderate clinical symptoms (61.6%), with severe clinical symptoms as many as 60 people (30.3%) and mild clinical symptoms as many as 16 people (8.1%). a) The most common comorbidity experienced by COVID-19 patients from RSU UKI is hypertension with 64 people (32.3%); b) Of the 156 people (78.8%) who had comorbidities, 47 people (23.7%) had increased D-Dimer levels, and 109 people (55.1%) had normal D-Dimer levels. Furthermore, of the 42 people (21.2%) who did not have comorbidities, six people (3%) had increased D-Dimer, and 36 people (18.2%) had normal D-Dimer levels; c) Prophylactic anticoagulant therapy was given to 127 COVID-19 patients, and when an evaluation was carried out by checking D-Dimer levels it was found that 81 people (64%) had normal D-Dimer levels, and 46 people (36%) had increased D-Dimer levels; d) This research shows that there is no relationship between the administration of anticoagulant prophylaxis and D-Dimer levels in COVID-19 sufferers with mild, moderate and severe clinical symptoms ($p=0.66$); and e) Of the 107 patients who had comorbidities and were given anticoagulant prophylaxis. It was found that 66 patients (62%) had normal D-Dimer levels, and 41 patients (38%) had increased D-Dimer levels.

Based on the conclusions outlined, suggestions that can be taken into consideration and included are as follows: a) Hospitals are advised to

continue paying attention to counseling/ education to the public regarding preventing the spread of COVID-19; b) Hospital staff are advised to check D-Dimer levels on the first day the patient is treated after the patient is declared positive for COVID-19 via RT – PCR examination, and provide anticoagulants for prophylaxis or therapy; c) For further research, it is highly hoped that the filling in of medical records will be carried out more completely, such as always including the results of laboratory examination of D-Dimer levels so that further research can obtain complete information; and d) It is hoped that future researchers can expand this research to a higher and more specific level of evidence, as well as analyze factors that can prevent an increase in D-Dimer levels.

2 CONSENT AND ETHICAL APPROVAL

It is not applicable.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

1. Anwar A, Malik M, Raees V, Anwar A. Role of mass media and public health communications in the COVID-19 pandemic. *Cureus*. 2020 Sep 14;12(9).
2. Burki TK. Coronavirus in China. *The Lancet Respiratory Medicine*. 2020 Mar 1;8(3):238.
3. Li X, Geng M, Peng Y, Meng L, Lu S. Molecular immune pathogenesis and diagnosis of COVID-19. *Journal of pharmaceutical analysis*. 2020 Apr 1;10(2):102-8.
4. Ogawa M, Sato M, Yamakami A, Hayasaka Y, Saya Y, Une K, Iwai S, Tsuchiya S, Mikami Y, Fukazawa C, Uehara Y. Diagnostic test property of transcription-reverse transcription concerted reaction reagent TRCReady® SARS-CoV-2 i using nasopharyngeal swab samples. *Journal of Infection and Chemotherapy*. 2023 Jan 1;29(1):115-7.
5. Tomo S, Karli S, Dharmalingam K, Yadav D, Sharma P. The clinical laboratory: A key player in diagnosis and management of COVID-19. *EJIFCC*. 2020 Nov;31(4):326.
6. Wan J, Yang X, He W, Zhu Y, Zhu Y, Zeng H, Liu P, Xia L, Lu N. Serum D-dimer levels at admission for prediction of outcomes in acute pancreatitis. *BMC*

- gastroenterology. 2019 Dec;19(1): 1-7.
7. Rostami M, Mansouritorghabeh H. D-dimer level in COVID-19 infection: A systematic review. *Expert review of hematology*. 2020 Nov 1;13(11):1265-75.
8. Eljilany I, Elzouki AN. D-dimer, fibrinogen and IL-6 in COVID-19 patients with suspected venous thromboembolism: A narrative review. *Vascular Health and Risk Management*. 2020 Nov 13:455-62.
9. Poor HD. Pulmonary thrombosis and thromboembolism in COVID-19. *Chest*. 2021 Oct 1;160(4):1471-80.
10. Ezzikouri S, Nourilil J, Benjelloun S, Kohara M, Tsukiyama-Kohara K. Coronavirus disease 2019—Historical context, virology, pathogenesis, immunotherapy, and vaccine development. *Human Vaccines & Immunotherapeutics*. 2020 Dec 1;16(12):2992-3000.
11. Sari EY, Cahyarini AS. Sensitivity and specificity of serological test compared to real-time polymerase chain reaction (rt-pcr) in covid-19 patients.
12. Wang B, Liu Y, Ji CM, Yang YL, Liang QZ, Zhao P, Xu LD, Lei XM, Luo WT, Qin P, Zhou J. Porcine deltacoronavirus engages the transmissible gastroenteritis virus functional receptor porcine aminopeptidase N for infectious cellular entry. *Journal of Virology*. 2018 Jun 15;92(12):10-128.
13. Li H, Liu SM, Yu XH, Tang SL, Tang CK. Coronavirus disease 2019 (COVID-19): Current status and future perspectives. *International Journal of Antimicrobial Agents*. 2020 May 1;55(5):105951.
14. Bchetnia M, Girard C, Duchaine C, Laprise C. The outbreak of the novel severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2): A review of the current global status. *Journal of Infection and Public Health*. 2020 Nov 1;13(11):1601-10.
15. Dhama K, Khan S, Tiwari R, Sircar S, Bhat S, Malik YS, Singh KP, Chaicumpa W, Bonilla-Aldana DK, Rodriguez-Morales AJ. Coronavirus disease 2019—COVID-19. *Clinical microbiology reviews*. 2020 Sep 16;33(4):10-128.
16. Gralinski LE, Menachery VD. Return of the coronavirus: 2019-nCoV. *Viruses*. 2020 Jan 24;12(2):135.
17. Yuki K, Fujiogi M, Koutsogiannaki S. COVID-19 pathophysiology: A review. *Clinical Immunology*. 2020 Jun 1;215:108427.
18. Lotfi M, Hamblin MR, Rezaei N. COVID-19: Transmission, prevention, and potential therapeutic opportunities. *Clinica Chimica Acta*. 2020 Sep 1;508:254-66.
19. Ouassou H, Kharchoufa L, Bouhrim M, Daoudi NE, Imtara H, Bencheikh N, ELbouzidi A, Bnouham M. The pathogenesis of coronavirus disease 2019 (COVID-19): Evaluation and prevention. *Journal of Immunology Research*. 2020 Jul 10;2020.
20. Jiang X, Niu Y, Li X, Li L, Cai W, Chen Y, Liao B, Wang E. Is a 14-day quarantine period optimal for effectively controlling coronavirus disease 2019 (COVID-19)? *MedRxiv*. 2020 Mar 18:2020-03.
21. HIDAYANI WR. Pneumonia: Epidemiologi, faktor risiko pada balita.
22. Levani Y, Prastya AD, Mawaddatunnadila S. Coronavirus disease 2019 (COVID-19): Patogenesis, manifestasi klinis dan pilihan terapi. *Journal Kedokteran dan kesehatan*. 2021 Jan 27;17(1):44-57.
23. Durachim A, Astuti D. Bahan Ajar Teknologi Laboratorium Medik (TLM): Hemostasis. Kementrian Kesehatan RI, Jakarta. 2018.
24. Umar I, Sujud RW. Hemostasis dan disseminated intravascular coagulation (dic). *Journal of Anaesthesia and Pain*. 2020 Jun 28;1(2):19-32.
25. Kurniawan LB, Arif M. Hemostasis berlandaskan sel hidup (in Vivo). *Indonesian Journal of Clinical Pathology and Medical Laboratory*. 2013;19(3):204-10.
26. Rostami M, Mansouritorghabeh H. D-dimer level in COVID-19 infection: A systematic review. *Expert Review of Hematology*. 2020 Nov 1;13(11):1265-75.
27. Luhulima DE. D-dimer pada kehamilan.
28. Eljilany I, Elzouki AN. D-dimer, fibrinogen and IL-6 in COVID-19 patients with suspected venous thromboembolism: A narrative review. *Vascular Health and Risk Management*. 2020 Nov 13:455-62.
29. Poudel A, Poudel Y, Adhikari A, Aryal BB, Dangol D, Bajracharya T, Maharjan A, Gautam R. D-dimer as a biomarker for assessment of COVID-19 prognosis: D-dimer levels on admission and its role in predicting disease outcome in hospitalized patients with COVID-19. *Plos one*. 2021 Aug 26;16(8):e0256744.
30. Yao Y, Cao J, Wang Q, Shi Q, Liu K, Luo Z, Chen X, Chen S, Yu K, Huang Z, Hu B. D-dimer as a biomarker for disease

- severity and mortality in COVID-19 patients: A case control study. *Journal of Intensive Care*. 2020 Dec;8:1-1.
31. Rostami M, Mansouritorghabeh H. D-dimer level in COVID-19 infection: A systematic review. *Expert Review of Hematology*. 2020 Nov 1;13(11):1265-75.
 32. Depasse F, Binder NB, Mueller J, Wissel T, Schwes S, Germer M, Hermes B, Turecek PL. Thrombin generation assays are versatile tools in blood coagulation analysis: A review of technical features and applications from research to laboratory routine. *Journal of Thrombosis and Haemostasis*. 2021 Dec 1;19(12):2907-17.
 33. Chandra A, Chakraborty U, Ghosh S, Dasgupta S. Anticoagulation in COVID-19: Current concepts and controversies. *Postgraduate Medical Journal*. 2022 May;98 (1159): 395-402.
 34. Listyoko AS, Djajalaksana S, Sugiri YJ. Analisis faktor koagulasi: Korelasi fibrinogen dengan rendahnya derajat oksigenasi pada pasien covid-19: Faktor koagulasi pada COVID-19. *Medica hospitalia: Journal of Clinical Medicine*. 2021 Jul 15;8(2):172-8.
 35. Rumana NA, Khaerunnisa R, Sitoayu L, Yulia N. Risiko terinfeksi Covid 19 pada pasien dengan komorbid diabetes melitus dan hipertensi di rumah sakit mekar sari kota bekasi. *Journal of Hospital Management ISSN (Print):2615:8337*.
 36. Elviani R, Anwar C, Sitorus RJ. Gambaran usia pada kejadian Covid-19. *Jambi Medical Journal" Journal Kedokteran dan Kesehatan"*. 2021 May 1;9(2): 204-9.
 37. Widjaja JT, Kwee L, Giantara AK, Suabgiyo HA, Edwin C, Putri RL. Karakteristik pasien Covid-19 rawat inap di RS immanuel bandung, Indonesia. *Journal of Medicine and Health*. 2021 Aug 17;3(2).
 38. Berger JS, Kunichoff D, Adhikari S, Ahuja T, Amoroso N, Aphinyanaphongs Y, Cao M, Goldenberg R, Hindenburg A, Horowitz J, Parnia S. Prevalence and outcomes of D-dimer elevation in hospitalized patients with Covid-19. *Arteriosclerosis, Thrombosis and Vascular Biology*. 2020 Oct;40(10):2539-47.
 39. Pratiwi AD, Adhityasmara D. Gambaran penggunaan antikoagulan pada pasien Covid-19 di salah satu rumah sakit rujukan Covid-19 di kota Semarang. *Sebatik*. 2021 Dec 1;25(2):442-8.
 40. Nugroho TE, Mochamat M, Famila F. penggunaan heparin dosis tinggi pada pasien Covid-19 dengan ards dan hipertensi di unit perawatan intensif (ICU). *JAI (Journal Anestesiologi Indonesia)*. 2020;12(3):1-8.

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