

# Clinical Outcomes of Non-diabetic COVID-19 Patients with Different Blood Glucose Levels: A Nationwide Turkish Study (TurCoGlycemia)

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## Research Article

**Keywords:** COVID-19, SARS-CoV-2, hyperglycemia, hospitalization, ICU admission, mechanical ventilation, mortality

**Posted Date:** May 10th, 2021

**DOI:** <https://doi.org/10.21203/rs.3.rs-497846/v1>

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# Abstract

## Purpose

New coronavirus disease 2019 (COVID-19) has a worse prognosis in patients with diabetes. However, there are insufficient data about the effect of hyperglycemia on COVID-19 prognosis in non-diabetic patients. This study aimed to investigate the relationship between random blood glucose levels measured at the time of diagnosis and prognosis of COVID-19 disease in non-diabetic patients.

## Methods

A nationwide retrospective cohort of non-diabetic patients with confirmed COVID-19 infection from 11 March to 30 May 2020 in the Turkish Ministry of Health database was investigated. The patients were stratified into three groups according to blood glucose levels which were <100 mg/dL in group-1, in the range of 100-139 mg/dl in group-2, and the range of 140-199 mg/dl in group-3. Clinical characteristics and outcomes were compared among the groups. The primary outcome was mortality.

## Results

A total of 12,817 non-diabetic patients (median age [IQR]: 44 [25] years, females: 50.9%) were included. Patients in group-2 (5%) and group-3 (14%) had higher mortality rates than patients in group-1 (2.1%). The rates of hospitalization, hospital stays longer than 8 days, intensive care unit (ICU) admission, ICU stay more than 6 days, and mechanical ventilation were also significantly higher in group-3 patients. Likewise, glucose levels in the range of 140-199 mg/dL were an independent associate of mortality and composite of ICU admission and/or mechanical ventilation.

## Conclusion

Hyperglycemia at the time of COVID-19 diagnosis is associated with poor prognosis in non-diabetic patients. Clinicians should be more careful in the treatment of non-diabetic COVID-19 patients with hyperglycemia.

## Introduction

New coronavirus disease 2019 (COVID-19) caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) has been one of the most important epidemics of the century threatening human health worldwide. Due to the COVID-19, more than 130 million patients have been infected, and more than 2.8 million deaths have occurred around the world [1, 2].

Mortality and morbidity rates of COVID-19 disease increase significantly in certain population groups such as males, older adults, or patients with comorbidities. Type 2 diabetes mellitus (T2DM) is one of the most frequent comorbidities in COVID-19 patients with considerably increased mortality and morbidity rates [3, 4]. Moreover, recent data indicate that new-onset hyperglycemia is common among hospitalized

COVID-19 patients with no history of diabetes in the past. Besides, hyperglycemia on hospital admission is a significant prognostic factor for COVID-19 patients without diabetes mellitus [5-8]. Studies from the United States, Italy, Spain, and China have consistently shown an increased risk of mortality, intensive care unit (ICU) admission, and a need for mechanical ventilation in patients with on admission hyperglycemia but no previous diagnosis of diabetes mellitus [5-9].

On the other hand, not all COVID-19 patients require hospitalization at the time of diagnosis. While the studies that have shown the impact of hyperglycemia on COVID-19 outcomes were performed exclusively in hospitalized patients, very little information has been obtained from the less severe COVID-19 population who comprise a great proportion of the confirmed cases. Moreover, criteria to exclude patients with diabetes and define hyperglycemia were not consistent across different studies mentioned above. In this study, we aimed to investigate the association of blood glucose levels by the time of COVID-19 diagnosis and clinical outcomes including hospitalization, mortality, and ICU admission and/or a need for mechanical ventilation in COVID-19 patients without diabetes mellitus.

## Materials And Methods

### 2.1. Study design and participants

This nationwide study was conducted using the COVID-19 registry of the Turkish Ministry of Health National Electronic Database. To promote data sharing with the scientific society, demographic, clinical, and outcome data of all symptomatic confirmed cases of COVID-19 infection were made available to the author group under the supervision of the Ministry of Health, and several publications have been made so far.[10, 11] The study population included 149,671 adult patients with confirmed COVID-19 (PCR positive) diagnosis from 11 March through 30 May 2020. **Figure 1** illustrates the patient enrollment. Briefly, patients with type 1 diabetes mellitus (n=370), T2DM (n=33,478) or unclassified (n=715, see "Definitions" below), with blood glucose level above 200 mg/dl (n=187) and no glucose measurement (n=102,104) at the time of COVID-19 diagnosis were excluded from the analysis. The final study population comprised of 12,817 confirmed COVID-19 patients without previously known diabetes. The patients were stratified into three groups according to glucose level which was <100 mg/dL in group-1, in the range of 100-139 mg/dl in group-2, and in the range of 140-199 mg/dl in group-3. The design and procedures in the study are in accordance with the Declaration of Helsinki and the study protocol was approved by the COVID-19 Investigation Review Board under the General Directorate of Health Services Bioethics Committee (IRB no: 95741342-020:186404/28.10.2020).

### 2.2. Data collection

Sociodemographic characteristics (gender, age, body mass index (BMI), smoking, and education), comorbid diseases, and information on medications were recorded. Blood glucose, low-density lipoprotein cholesterol (LDL-cholesterol), high-density lipoprotein cholesterol (HDL-cholesterol), total cholesterol, and triglycerides, creatinine, alanine and aspartate aminotransferases (ALT and AST), C-reactive protein (CRP), lymphocyte count, procalcitonin, lactate dehydrogenase (LDH), ferritin, fibrinogen,

and D-dimer were obtained from the national database. Blood glucose levels measured within 24 hours of COVID-19 diagnosis were recorded regardless of fasting state. Chest computerized tomography (CT) results were available in the national database and recorded as positive (bilateral distribution of patchy shadows or ground-glass opacity) or negative for COVID-19.

### **2.3. Definitions**

Smoking was defined as currently smoking at the time of the COVID-19 diagnosis. Hypertension, dyslipidemia, chronic obstructive pulmonary disease (COPD), asthma, heart failure, cardiovascular disease (CVD) (combined coronary artery, peripheral artery, and cerebrovascular diseases) were identified using the International Classification of Diseases System-10 (ICD-10) codes. BMI was calculated as the ratio of weight to the square of heights ( $\text{kg}/\text{m}^2$ ). Obesity was defined as  $\text{BMI} \geq 30 \text{ kg}/\text{m}^2$ . Chronic kidney disease was defined as a decreased estimated glomerular filtration rate (eGFR:  $<60 \text{ mL}/\text{min}/1.73 \text{ m}^2$ ) based on the CKD-EPI equation [12]. Renin-angiotensin system (RAS) blocker use was defined as receiving any angiotensin-converting enzyme inhibitors or angiotensin receptor blockers with/without their combination forms. Patients who did not have an ICD-10 code for type 1 diabetes or T2DM in twelve months past or patients who had an HbA1c value below 6.5% along with the absence of any antihyperglycemic medication were defined as non-diabetic. Patients with missing data were excluded.

### **2.4. Study outcomes**

The primary outcome was mortality. The secondary outcomes were the rates of hospitalization and ICU admission and/or mechanical ventilation.

### **2.5. Statistical analyses**

Numerical data were expressed as median (interquartile range-IQR) and categorical variables as counts (n) and percentage (%). Normality of distribution was assessed using the Kolmogorov-Smirnov test. Differences between groups were assessed using the Chi-square test for categorical variables and the student's t-test or the Mann-Whitney U test, as appropriate. Univariate analyses were performed to evaluate the potential variables associated with mortality, hospitalization, and ICU admission, and/or mechanical ventilation and presented with an odds ratio (OR) and 95% confidence intervals (CI). Multivariable logistic regression analysis was used to study the independently associated risk factors of the three outcomes. Variables with significant univariate association with the outcomes and variables which could be potential predictors despite the lack of significant univariate association were included in a multivariate model. Hosmer-Lemeshow and the likelihood ratio tests were used to assess final model fitting. Statistical significance was defined as a two-sided p-value of  $\leq 0.05$ . All data were analyzed using Statistical Package for the Social Sciences (SPSS) for Windows 25.0 (SPSS Inc. 111 Chicago, IL).

## **Results**

### **3.1. Basic characteristics**

The median age of the study population (n=12,817) was 44 (25) years, and 50.9% was female. The demographic and clinical parameters of the three groups are displayed in **Table 1**. The percentage (number) of patients in group-1, 2, and 3 were 46.1% (n=5911), 47.3% (n=6058), and 6.6% (n=848), respectively. Median age, male sex, and mean BMI showed increases from group 1 to group 3. The current smoking rates in the groups were 21.2%, 19.1%, and 22.4%, respectively.

As shown in **Table 1**, most laboratory test results showed a significantly worse direction from group-1 to group-3. Notably, D-dimer, CRP, LDH, ferritin, and lymphopenia that were more commonly referred to COVID-19 severity, increased gradually across the groups. Only three laboratory variables, total cholesterol, LDL-cholesterol, and fibrinogen, were not different from normal glucose to high glucose group. Lung involvement on CT increased considerably from group-1 to group-3.

The prevalences of hypertension, dyslipidemia, obesity, asthma/COPD, heart failure, CVD, chronic kidney disease, and cancer were significantly different among the three groups, with more than double the increase in the frequencies of CVD, heart failure, chronic kidney disease, and medications in group 3 (**Table 1**). Similarly, the use of RAS blockers, statins, and acetylsalicylic acid (ASA) was more common in group-3.

### 3.2. Outcome analysis

There were significant differences in the primary and secondary outcomes across the groups. Mortality rate was found to be 14% in group-3, which was significantly higher than the mortality rate of 5% and 2.1% in group-2 and group-1, respectively. Mortality was also significantly higher in group-2 compared to group-1. Similar differences were also found in the rate of hospitalization, hospital stays longer than the median duration (8 days), ICU admission, ICU stay longer than the median duration (6 days), and mechanical ventilation (**Table 2**).

### 3.3. Independent associations of mortality and secondary outcomes

Multivariate analyses indicated that age (OR, 95% CI: 1.02, 1.01-1.02), lung involvement (OR, 95% CI: 3.12, 2.58-3.77), lymphopenia (OR, 95% CI: 1.26, 1.03-1.55), chronic kidney disease (OR, 95% CI: 1.62, 1.12-2.33), cancer (OR, 95% CI: 1.49, 1.02-2.18) and the use of ASA (OR, 95% CI: 1.68, 1.24-2.27) were associated with the risk of hospitalization (**Table 3**). Additionally, age (OR, 95% CI: 1.07, 1.06-1.09), male gender (OR, 95% CI: 2.68, 1.73-4.15), lung involvement (OR, 95% CI: 1.55, 1.04-2.32), glucose levels in the range of 140-199 mg/dl (OR, 95% CI: 2.66, 1.41-5.00), lymphopenia (OR, 95% CI: 1.1.87, 1.25-2.80), CRP (OR, 95% CI: 3.11, 1.69-5.73), hypertension (OR, 95% CI: 1.95, 1.00-3.79) and chronic kidney disease (OR, 95% CI: 2.16, 1.40-3.35) associated with mortality (**Table 2**). Also, age (OR, 95% CI: 1.04, 1.03-1.05), male gender (OR, 95% CI: 1.92, 1.46-2.53), lung involvement (OR, 95% CI: 1.63, 1.25-2.13), glucose level in the range of 140-199 mg/dl (OR, 95% CI: 2.26, 1.45-3.52), lymphopenia (OR, 95% CI: 2.03, 1.54-2.66), CRP (OR, 95% CI: 2.31, 1.67-3.20) and heart failure (OR, 95% CI: 1.94, 1.26-2.98) were significantly associated with the composite of ICU admission and/or mechanical ventilation (**Table 2**).

The Kaplan-Meier graph of 30-day mortality curves in the total sample is given in **Figure 2**. The rate of mortality in group 3 patients was significantly higher (Log-Rank test:  $p < 0.001$ ) than in other groups.

## Discussion

The present study showed that non-diabetic COVID-19 patients who had increased glucose levels at the time of diagnosis were at increased risk of mortality, longer hospitalization, ICU admission, and a need for mechanical ventilation than patients with lower blood glucose. On the other hand, glucose levels at the time of diagnosis was not independently associated with the occurrence of hospitalization. Additional variables such as age, male gender, lung involvement, lymphopenia, CRP, hypertension, and chronic kidney disease were associated with reduced survival in non-diabetic patients with COVID-19 disease.

Many studies reported that diabetes mellitus, especially T2DM, is one of the most frequent comorbidities in COVID-19 patients [10, 11, 13-15]. The findings also indicated that hyperglycemic state in patients is very important in the prognosis of COVID-19 disease. Furthermore, some recent studies reported that glycemia levels on hospital admission had a negative impact on the prognosis of COVID-19 disease in non-diabetic patients [5, 6, 8, 9, 16]. With one of the largest patient groups, the present study showed that hyperglycemia at the time of COVID-19 diagnosis in non-diabetic patients was an independent predictor of mortality and the composite endpoint of ICU admission and/or mechanical ventilation. Especially, glucose levels in the range of 140-199 mg/dl increased the mortality 2.7 fold and the rate of ICU admission and/or mechanical ventilation 2.3 fold, compared to normoglycemia. However, glucose levels in the range of 100-139 mg/dl had no effect on these outcomes. Our results are consistent with the findings of other studies which showed increased mortality due to hyperglycemia in non-diabetic patients with COVID-19 disease [5-9]. However, each of these studies used different criteria to identify the non-diabetic patients and also used the various thresholds to define the hyperglycemia when investigating the effect of glycemia levels. Therefore, there may be some methodological issues, especially in identifying non-diabetic patients and in defining hyperglycemia levels in these studies. For instance, in the study by Coppeli et al., the normoglycemia group was defined as the patients who had glucose levels  $< 140$  mg/dl [7]. It can be speculated that there may be the probability of having undiagnosed diabetes patients in the normoglycemia group. Also, in another recent study by Carrasco et al., hyperglycemia groups were defined as patients who had glycemia levels in the range of 140-180 mg/dl, and  $\geq 180$  mg/dl [9]. Similarly, Bode et al., defined hyperglycemia in patients without diabetes as two or more blood glucose levels  $> 180$  mg/dL occurred within any 24-hour period with an HbA1c  $< 6.5\%$  or no HbA1c testing performed during hospitalization [5]. In both studies, hyperglycemia in non-diabetic patients was significantly associated with increased mortality compared to patients with normoglycemia. However, diabetes mellitus diagnosis may be missed in a significant proportion of patients with glycemia levels above 180 mg/dl in both studies. In the current study, we used the glucose cut-off levels for normoglycemia as  $< 100$  mg/dl and did not include the glucose values above 200 mg/dl in order to eliminate false negativity regarding the diagnosis of diabetes. In addition, we excluded patients who use antidiabetic drugs, have HbA1c  $\geq 6.5\%$ , and have a diagnosis of diabetes with an ICD-10 diagnostic code.

Therefore, we think that we had a very homogeneous patient population in order to demonstrate the effect of hyperglycemia in non-diabetic patients.

It is not surprising that hyperglycemia was associated with mortality in non-diabetic patients with COVID-19 disease. It is well known that hyperglycemia affects mortality in critical illnesses and cardiovascular diseases [17]. Also, previous studies have reported that hyperglycemia was associated with mortality in SARS and the Middle East respiratory syndrome (MERS) [18, 19]. There is a bidirectional relationship between hyperglycemia and SARS-CoV-2 infection. SARS CoV-2 uses the angiotensin-converting enzyme 2 (ACE-2) as the receptor for entry into the cells. Hyperglycemia classically impairs the immune system response by affecting neutrophilic chemotaxis and phagocytosis, but in SARS-CoV-2 infection, it may also increase the expression of the ACE-2 receptors and cause a susceptibility [20, 21]. Since the ACE-2 receptors are present in other organs too that regulate glucose hemostasis, it is also possible for SARS CoV-2 to cause hyperglycemia affecting these organs [20]. Although the underlying mechanisms are not fully elucidated, our results support the negative effect of hyperglycemia on the prognosis of COVID-19 disease.

It has been also shown that diabetes is an important risk factor for hospitalization in COVID-19 disease [22], but the effect of hyperglycemia for hospitalization of non-diabetic patients is unknown. Unlike other studies, the present study is the first one using random glycemia levels at the time of COVID-19 diagnosis to search for the effect of hyperglycemia levels on the prognosis of COVID-19 disease. This method eliminates the increase in glucose levels due to inflammation that may occur during the course of COVID-19 disease or hyperglycemia that may develop secondary to drug use such as steroid therapy. Also, the use of glucose values at the time of diagnosis gave the opportunity to demonstrate the effect of hyperglycemia on hospitalization rates for the first time. Despite the striking effects of hyperglycemia on the clinical severity of COVID-19 disease, no effect was found on the hospitalization rates compared to normoglycemia levels. The possible reason for this result may be related to the hospitalization policy of the Ministry of Health, at the beginning of the pandemics. Patients, especially those with chronic diseases were hospitalized regardless of the symptom severity according to this policy.

There may be several limitations of the current study. First of all, the cross-sectional nature of the study may preclude the cause-effect relationship between clinical and laboratory parameters and outcomes. Also, there may be residual or unmeasured confounding factors. Second, although strict criteria have been used to identify non-diabetic patients, there is also the possibility of undiagnosed patients with diabetes in the groups. To eliminate the risk of misclassification of patients with diabetes mellitus into the study, we performed a targeted database search to identify any anti-hyperglycemic prescription in 12 months past. Additional to this limitation, we were not able to identify in the database whether glucose measurement was performed at a fasting state or random. Therefore, the present sample does not successfully represent the population with prediabetes. On the other hand, this may also be meaningful within the real-world conditions as most COVID-19 patients undergo blood sampling at any time of the day, though services may differ in other countries. Third, as the data obtained from the electronic data recording system of the Ministry of Health, some missing data could have a potential impact on the

results. However, we believe that large patient population and the strict inclusion and exclusion criteria are the important strength of the present study.

In conclusion, this study demonstrated that glycemia levels especially in the range of 140-199 mg/dL at the time of COVID-19 diagnosis significantly increased the rates of ICU admission and/or a need for mechanical ventilation and mortality in non-diabetic patients. However, hyperglycemia was not found to have an impact on hospitalization rates. In the light of these findings, clinicians should be more careful, especially in the treatment of non-diabetic COVID-19 patients who had hyperglycemia at the time of diagnosis.

## Declarations

**Funding:** This research did not receive any funding.

**Conflicts of interest:** On behalf of all authors, the corresponding author states that there is no conflict of interest.

**Availability of data and material:** Not applicable.

**Code availability:** Not applicable.

**Ethics approval:** This study was approved by the COVID-19 Investigation Review Board (IRB) under the General Directorate of Health Services Bioethics Committee designated by the Ministry of Health. (IRB no: 95741342-020/27112019)

**Consent to participate:** Not applicable.

**Consent for publication:** Not applicable.

**Acknowledgements:** We acknowledge all health care professionals with gratitude and respect who have devoted their lives to combating COVID-19 and have provided these data.

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## Tables

**Table 1.** Clinical and laboratory parameters of patients according to the different blood glucose levels.

	Group-1 Glucose <100 mg/dl (n=5911, 46.1%)	Group-2 Glucose 100- 139 mg/dl (n=6058, 47.3%)	Group-3 Glucose 140- 199 mg/dl (n=848, 6.6%)	Total available data (n=12,817)	p
Age, years, median (IQR)	35 (19) <sup>a,b</sup>	45 (22) <sup>c</sup>	55(20)	12817	<b>&lt; 0.001</b>
Gender, female, n (%)	3137 (53.1) <sup>a,b</sup>	3009 (49.7) <sup>c</sup>	383 (45.2)	12817	<b>&lt; 0.001</b>
Smoking (current smoker - n,%)	952 (21.2)	832 (19.1)	125 (22.4)	9417	<b>0.022</b>
BMI, kg/m <sup>2</sup> , median (IQR)	25.5 (6.1)	27.2 (6.1)	28.4 (5.6)	1483	<b>&lt;0.001</b>
Comorbid conditions					
Hypertension, n (%)	1950 (33.0)	2739 (45.2)	503 (59.3)	12817	<b>&lt;0.001</b>
Dyslipidemia, n (%)	601 (10.2)	849 (14.0)	178 (21.0)	12817	<b>&lt;0.001</b>
Obesity, n (%)	121 (17.5)	187 (27.0)	34 (34.3)	1482	<b>&lt;0.001</b>
Asthma/COPD, n (%)	1233 (20.9)	1561 (25.8)	271 (32.0)	12817	<b>&lt;0.001</b>
Heart failure, n (%)	179 (3.0)	266 (4.4)	60 (7.1)	12817	<b>&lt;0.001</b>
CVD, n (%)	868 (14.7)	1309 (21.6)	293 (34.6)	12817	<b>&lt;0.001</b>
Chronic kidney disease, n (%)	306 (5.6)	551 (9.7)	125 (15.7)	11947	<b>&lt;0.001</b>
Cancer, n (%)	309 (5.2)	384 (6.3)	88 (10.4)	12817	<b>&lt;0.001</b>
Laboratory values					
Chest CT on admission consistent with COVID-19, n (%)	1615 (29.2) <sup>a,b</sup>	2126 (36.5) <sup>c</sup>	354 (43.4)	12161	<b>&lt;0.001</b>
Glucose, mg/dL, median (IQR)	90 (10)	110 (16)	152 (32)	12769	<b>&lt;0.001</b>
Total cholesterol, mg/dL, median (IQR)	178 (58.7)	179.5 (66.1)	196 (68)	1141	0.151
Triglycerides, mg/dL, median (IQR)	105 (90.5)	126 (98.5)	136 (103)	1556	<b>0.003</b>
HDL-cholesterol,	48.7 (21.3)	45 (18)	47 (16.4)	1205	<b>0.026</b>

mg/dL, median (IQR)					
LDL-cholesterol, mg/dL, median (IQR)	106.5 (46.8)	111.5 (58)	118 (53.9)	1090	0.235
eGFR, ml/min/1.73 m <sup>2</sup> , median (IQR)	105.3 (53.1)	99.2 (48.6)	91.1 (54.1)	8563	<0.001
AST, >ULN, n (%)	311 (12.8)	526 (18.4)	104 (24.9)	5714	<0.001
ALT, >ULN, n (%)	392 (16.2)	545 (19.2)	101 (24.3)	5684	<0.001
D-dimer >ULN, n (%)	374 (33.2)	429 (38.4)	85 (51.5)	2406	<0.001
CRP, >ULN, n (%)	1760 (49.3)	2405 (61.8)	390 (68.3)	8031	<0.001
Procalcitonin, >ULN, n (%)	35 (6.3)	51 (9.8)	21 (22.6)	1168	<0.001
Lactate dehydrogenase, >ULN, n (%)	620 (26.1)	976 (38.2)	168 (46.4)	5286	<0.001
Ferritin, >100 ng/mL, n (%)	716 (42.8)	867 (52.4)	159 (67.4)	3565	<0.001
Fibrinogen, >ULN, n(%)	176 (64.7)	199 (69.3)	35 (61.4)	616	0.351
Lymphopenia, Lym # <1000, n (%)	698 (17.1)	961 (22.7)	164 (29.1)	8869	<0.001
Treatments					
RAS blocker, n (%)	960 (16.2)	1601 (26.4)	330 (38.9)	12817	<0.001
Statin, n (%)	236 (4.0)	378 (6.2)	108 (12.7)	12817	<0.001
Acetylsalicylic acid, n (%)	701 (11.9)	1011 (16.7)	209 (24.6)	12817	<0.001
<b>Post-hoc tests: <sup>a</sup>p&lt; 0.05 group-1 vs. group-2, <sup>b</sup>p&lt; 0.05 group-1 vs. group-3, <sup>c</sup>p &lt;0.05 group-2 vs. group-3</b>					
<b>Abbreviations:</b> IQR, interquartile range; BMI, body mass index; CT, computerized tomography; HDL-cholesterol, high-density lipoprotein cholesterol; LDL-cholesterol, low-density lipoprotein cholesterol; eGFR, estimated glomerular filtration rate; ULN, upper limit of normal; AST, aspartate amino transferase; ALT, alanine amino transferase; CRP: C-reactive protein; Lym #, lymphocyte count; COPD, chronic obstructive pulmonary disease; CVD, cardiovascular disease; RAS, renin angiotensin aldosterone system.					

**Table 2.** Primary and secondary outcomes according to the different blood glucose levels

	Group-1 Glucose <100 mg/dl (n=5911, 46.1%)	Group-2 Glucose 100- 139 mg/dl (n=6058, 47.3%)	Group-3 Glucose 140- 199 mg/dl (n=848, 6.6%)	Total available data (n=12,817)	p
Hospitalization	3839 (64.9) <sup>a,b</sup>	4282 (70.7) <sup>c</sup>	641 (75.6)	12817	<b>&lt;0.001</b>
Hospital stay more than 8 days, n (%)	1739 (45.3) <sup>a,b</sup>	2118 (49.5) <sup>c</sup>	379 (59.1)	8762	<b>&lt;0.001</b>
ICU admission, n (%)	307 (8.0) <sup>a,b</sup>	568 (13.3) <sup>c</sup>	168 (26.4)	8748	<b>&lt;0.001</b>
ICU stay more than 6 days, n (%)	135 (44.0) <sup>b</sup>	288 (50.9) <sup>c</sup>	100 (59.5)	1041	<b>0.005</b>
Mechanical ventilation, n (%)	160 (52.1) <sup>a,b</sup>	342 (60.2) <sup>c</sup>	116 (69.0)	1043	<b>0.001</b>
Death, n (%)	123 (2.1) <sup>a,b</sup>	300 (5.0) <sup>c</sup>	119 (14.0)	12817	<b>&lt;0.001</b>
<b>Post-hoc tests: <sup>a</sup>p &lt;0.05 group-1 vs. group 2, <sup>b</sup>p &lt;0.05 group-1 vs. group-3, <sup>c</sup>p &lt;0.05 group-2 vs. group-3</b>					
<b>Abbreviations:</b> ICU, intensive care unit.					

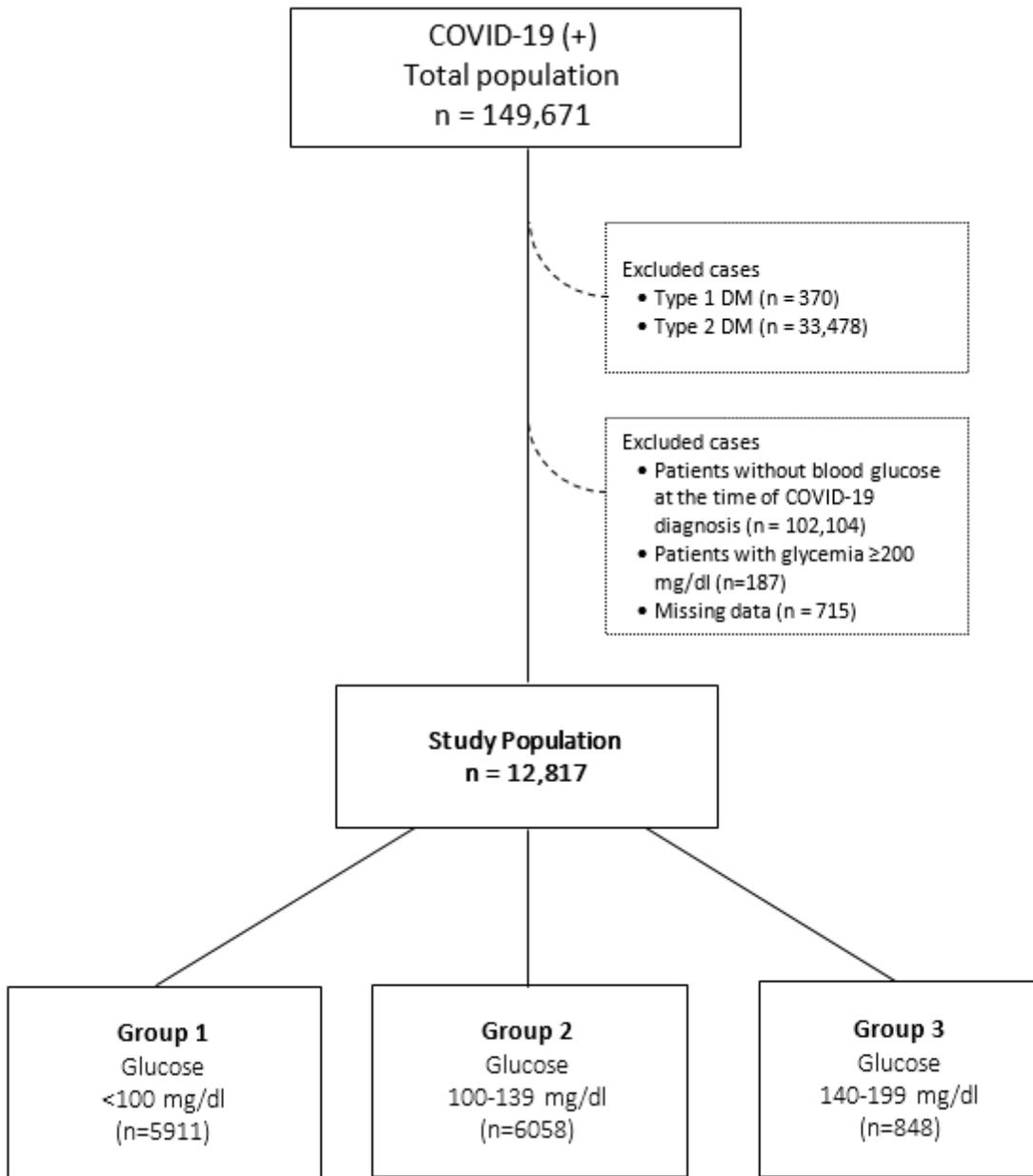
**Table 3.** Adjusted ORs and 95% CIs for different levels of blood glucose from multivariate logistic regression (dependent variables: hospitalization, ICU admission and/or mechanical ventilation, and mortality)

	Hospitalization		ICU Admission and/or Mechanical Ventilation		Mortality	
	OR (95% CI)	p	OR (95% CI)	p	OR (95% CI)	p
Glucose <100 mg/dl, ref.						
Glucose 100-139 mg/dl	-	-	1.24 (0.93-1.66)	0.145	1.44 (0.91-2.29)	0.119
Glucose 140-199 mg/dl	-	-	2.26 (1.45-3.52)	<b>&lt;0.001</b>	2.66 (1.41-5.00)	<b>0.002</b>
Age	1.02 (1.01-1.02)	<b>&lt;0.001</b>	1.04 (1.03-1.05)	<b>&lt;0.001</b>	1.07 (1.06-1.09)	<b>&lt;0.001</b>
Gender, male ref.	-	-	1.92 (1.46-2.53)	<b>&lt;0.001</b>	2.68 (1.73-4.15)	<b>&lt;0.001</b>
CT findings of COVID-19	3.12 (2.58-3.77)	<b>&lt;0.001</b>	1.63 (1.25-2.13)	<b>&lt;0.001</b>	1.55 (1.04-2.32)	<b>0.031</b>
Lymphopenia (Lym# < 1000/micL)	1.26 (1.03-1.55)	<b>0.027</b>	2.03 (1.54-2.66)	<b>&lt;0.001</b>	1.87 (1.25-2.80)	<b>0.002</b>
CRP	-	-	2.31 (1.67-3.20)	<b>&lt;0.001</b>	3.11 (1.69-5.73)	<b>&lt;0.001</b>
Hypertension	-	-	-	-	1.95 (1.00-3.79)	<b>0.049</b>
Chronic kidney disease	1.62 (1.12-2.33)	<b>0.010</b>	-	-	2.16 (1.40-3.35)	<b>0.001</b>
Heart failure	-	-	1.94 (1.26-2.98)	<b>0.003</b>	-	-
Cancer	1.49 (1.02-2.18)	<b>0.037</b>	-	-	-	-
Acetylsalicylic acid	1.68 (1.24-2.27)	<b>0.001</b>	-	-	-	-

**Abbreviations:** ICU, intensive care unit; OR, odds ratio; CI, confidence intervals; CT, computerized tomography; COVID-19, new coronavirus-19; CRP, C-reactive protein.

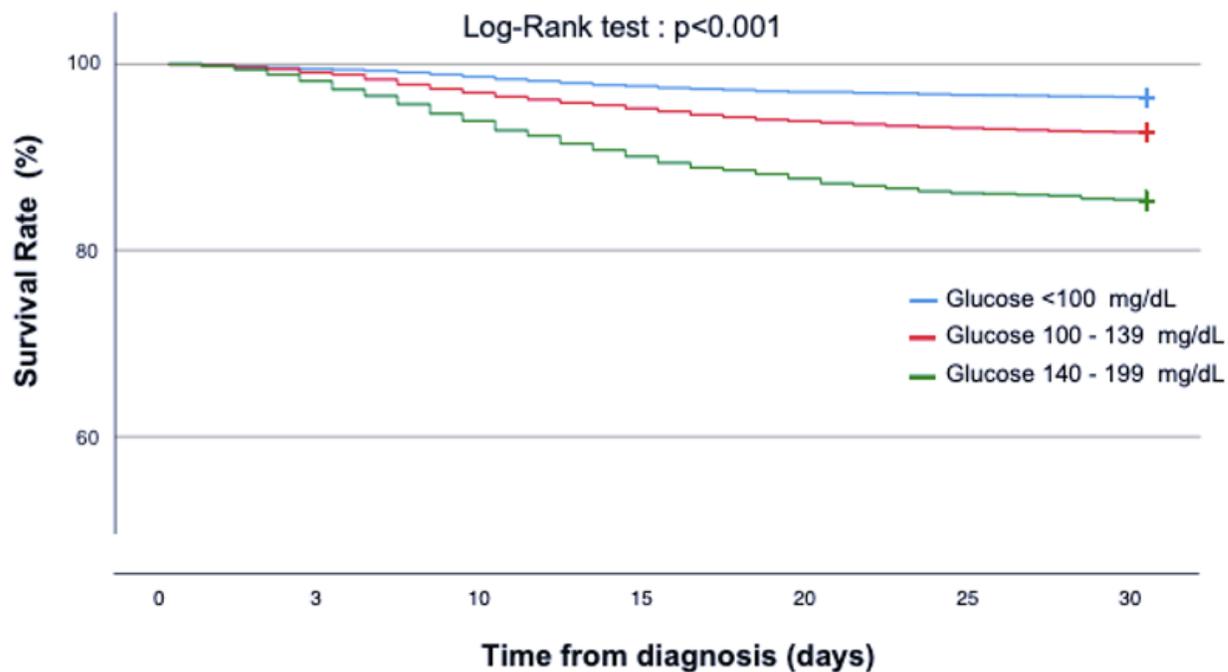
**Variables included in the multivariate analyses:** Glucose levels <100 mg/dl, Glucose levels between 100-140 mg/dl, Glucose levels between 140-200 mg/dl, Age, Gender, Pulmonary CT findings of COVID-19, Hypertension, Dyslipidemia, Obesity, Asthma/COPD, Heart failure, Cardiovascular disease, Chronic kidney disease, Cancer, Lymphopenia, RAS blockers ± combinations, Statins, Acetylsalicylic acid (only the significant associates were given in the table).

# Figures



**Figure 1**

Study inclusion flow chart



	Number at risk						
	0	3	10	15	20	25	30
Glucose <100 mg/dL	5025	4992	4946	4899	4877	4859	4846
Glucose 100 - 139 mg/dL	6419	6349	6197	6096	6018	5974	5952
Glucose 140 - 199 mg/dL	1892	1842	1759	1692	1650	1629	1615

**Figure 2**

Survival analyses of patients according to different blood glucose levels