

COVID-19 in Adults With Dementia: Clinical Features and Predictive Factors of Mortality. A Clinical Cohort Study on 125 Patients

Agathe Vrillon (✉ agathe.vrillon@aphp.fr)

APHP: Assistance Publique - Hopitaux de Paris <https://orcid.org/0000-0002-0761-7885>

Elsa Mhanna

Fondation Léopold Bellan, France

Clément Aveneau

AP-HP: Assistance Publique - Hopitaux de Paris

Manon Lebozec

AP-HP: Assistance Publique - Hopitaux de Paris

Lina Grosset

AP-HP: Assistance Publique - Hopitaux de Paris

Diane Nankam

Fondation Léopold Bellan, France

Fernanda Albuquerque

Fondation Léopold Bellan, France

Raphaëlle Razou Feroldi

Fondation Léopold Bellan, France

Barbara Maakaroun

Fondation Léopold Bellan, France

Iana Pissareva

Fondation Léopold Bellan, France

Dalenda Cherni Gherissi

Fondation Léopold Bellan, France

Julien Azuar

AP-HP: Assistance Publique - Hopitaux de Paris

Véronique François

AP-HP: Assistance Publique - Hopitaux de Paris

Claire Hourrègue

AP-HP: Assistance Publique - Hopitaux de Paris

Julien Dumurgier

AP-HP: Assistance Publique - Hopitaux de Paris

Lisette Volpe-Gillot

Fondation Léopold Bellan

Claire Paquet

AP-HP: Assistance Publique - Hopitaux de Paris

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Abstract

BACKGROUND

There is limited evidence on the characteristics and outcome of patients with dementia hospitalized for novel coronavirus infection (COVID-19).

METHOD

We conducted a prospective study in 2 gerontologic Covid Units in Paris, France, from March 14th 2020 to May 7th 2020. Patients with dementia hospitalized for confirmed COVID-19 infection were systematically enrolled. A binary logistic regression analysis was performed to identify factors associated with mortality at 21 days.

RESULTS

We included 125 patients. Median age was 86 (IQR 82-90); 59.4% were female. Most common causes of dementia were Alzheimer's disease, mixed dementia and vascular dementia. 67.2% had ≥ 2 comorbidities; 40.2% lived in a long-term care facility. The most common symptoms at COVID-19 onset were confusion and delirium (82.4%), asthenia (76.8%) and fever (72.8%) before polypnea (51.2%) and desaturation (50.4%). Falls were frequent at the initial phase of the disease (35.2%). The fatality rate at 21 days was 22.4%. Chronic kidney disease and CRP at admission were independent factors of death. Persisting confusion, mood and behavioral disorders were observed in survivors (19.2%).

CONCLUSION

COVID-19 in demented individuals is associated with severe outcome in SARS-CoV-2 infection and is characterized by specific clinical features and complications, with confusion and delirium at the forefront. COVID-19 testing should be considered in front of any significant change from baseline.

Introduction

On January 30th 2020, the World Health Organization (WHO) drew attention to a new coronavirus disease 2019 (COVID-19) declaring it a public health emergency of international concern. As of February the 4th 2021, there had been more than 100 millions of cases and 2 millions deaths worldwide. The clinical spectrum of COVID-19 infection appears broad, encompassing asymptomatic infections, mild upper respiratory tract illness and severe pneumonia with respiratory failure, systemic complications and multi-organ failure [1].

Severe COVID-19 affects elderly with chronic diseases, including cognitive decline, in high proportion compared to general adult population [2,3]. According to recent studies, dementia is a major risk factor for COVID-19 severity [4,5]. Concomitantly, the risk of exposure to the infection is more important in patients with dementia, highly exposed in the context of long-term care facilities, frequent

hospitalizations and intellectual decline. So far, specific clinical features and prognostic factors of COVID-19 in demented patients remain unclear [6,7].

In order to identify specific features and risk factor of death in demented people, we report a cohort study on 125 patients with dementia hospitalized for a confirmed COVID-19 infection.

Methods

Study design

This prospective cohort study systematically included patients over 65 with dementia hospitalized for COVID-19 infection in two centers : Leopold Bellan Hospital, Paris, France and Lariboisière Fernand Widal University Hospital Paris, France, between March 14th and May 7th 2020. The definite diagnosis of COVID-19 was determined through reverse-transcription polymerase chain reaction (RT-PCR) testing of pharyngeal swabs or chest CT. We collected demographics, medical history, clinical presentation, laboratory results, treatment, complications and outcome. The outcome was death at 21 days, to assess short-term case fatality rate. Dementia was defined according to the American Psychiatric Association's Diagnostic and Statistical Manual (DSM-5) as a significant cognitive decline from a previous level of performance in one or more cognitive domains interfering with independence in everyday activities [8]. Etiology of the dementia was collected when available. Comorbidities were evaluated using the Charlson Index [9]. Blood oxygen desaturation was defined by a saturation under 93 % or a loss of 3% or more. Acute kidney injury was diagnosed according to KDIGO definition [10]. Cardiac injury was diagnosed clinically or throughout, abnormalities observed on electrocardiography and serum level of troponin and Brain natriuretic Peptide (BNP). COVID-19 severity and Acute Respiratory Distress Syndrome were defined following the WHO recommendations. None of the patients included in this study were deemed candidates for the intensive care unit after staff discussions with the physician in charge, palliative care team and patients and caregivers.

Statistical analysis

Descriptive data are shown as median (interquartile interval) or percentage (number of subjects). Chi-square test or Fisher test was used to compare qualitative data between groups. Mann Whitney test and Kruskal Wallis were applied to analyze non-normally distributed data. A binary logistic regression analysis was performed to identify clinical and demographic characteristics associated with mortality. We included the variables that had differed in descriptive analysis between survivors and non survivors. Variables with missing data were excluded from this analysis. A two-tailed p-value < 0.05 was considered significant. Statistical analysis was performed using GraphPad Prism and SPSS 85 Statistics version 26.0.

Ethical statement: we obtained the required approval from the Commission Nationale Informatique et Liberté (CNIL) to collect anonymized data.

Results

We included a total of 125 patients. Patients demographic and clinical characteristics are summarized in **Table 1**. The median age was 86 (82-91) year-old (yo), and 41.6% of included subjects were male. All patients had a clinical diagnosis of dementia. 40 % of the patients had received a specific diagnosis: most frequent cause of dementia was Alzheimer's Disease (AD) (10.4%), vascular dementia (7.2%), multiple etiologies dementia (7.2%). Parkinson's disease and atypical parkinsonian syndromes, fronto-temporal dementia alcohol related cognitive impairment and psychosis accounted for other diagnoses. Patients frequently presented with associated mood disorder (31.2%) and psychotic symptoms (12%). A stroke history was noted in 32% of patients. A former history of fall was reported for 38.4% of patients. Sixty percent of patients received psychotropic treatment: antidepressants/mood regulators (35.2%), anxiolytic (30.4%) and neuroleptics (12.0%). Most patients were living at home (59.2%) while 40.8% were living in a long-term care facility.

Non neurological comorbidities were frequent (91.2%) with 67.2% of patients with 3 or more comorbidities. Median Charlson comorbidities index was 6 (5-7). Most frequent comorbidities were hypertension (80%), dyslipidemia (48.8%) and history of cardiovascular disease (40.8%).

Clinical presentation and biological findings

Infection was confirmed in 93.2% of patients by PCR test and in 6.8% by chest CT (**Table 2**). The most common initial symptoms were confusion and delirium observed in 82.4% of patients. It was notably the only symptom at onset and during the evolution of the disease for three patients in our cohort, thus COVID-19 infection was screened for considering the pandemic context. The most frequent associated general symptoms were asthenia (76.8%), fever (72.8%) and anorexia (56.8%). Most frequent respiratory symptoms were: polypnea (51.2%), desaturation (50.4%) and cough (49.6%). An initial fall was reported in 35.2%. No patient complained of loss of taste or smell.

The most common biochemical abnormality was lymphopenia (84.8%) with a median value of lymphocytes of 0.99 (0.73-1.29), increased CRP at admission and low albumin. Thrombopenia was observed in 16% of cases. Troponin was available for 22 patients and increased in 35.5% of patients.

Treatment and outcome

Treatment and outcome are reported in Table 3. Persisting behavioral disorders during hospitalization were observed in 19.2%. Other neurological complications included strokes (2.4%, 3 patients) and seizure (2.4%, 3 patients). Treatment mostly associated oxygen therapy (60%) and antibiotherapy (61.6%). A minority of patients received specific treatment for COVID-19 (hydroxychloroquine 16.8% and corticosteroids 8.0%). ARDS (Acute respiratory distress syndrome) occurred in 25.6 % of patients. Associated complications were bacterial surinfection (19.2%), acute kidney injury (16.8%), cardiac failure

(13.6%) and diabetic acidoketosis (2.4%). Treatment by neuroleptics or antidepressant was not associated to outcome.

Mortality at 21 days was 22.4%. The large majority of deaths (92.9%) were attributed to ARDS. Cardiac failure and osmolar coma were associated causes of death. Non-survivors were overrepresented in men (60.7 % versus 36.1% in survivors, $P=0.029$), presenting more frequent chronic kidney disease. Respiratory distress, lower lymphocytes count, high CRP and positive troponin were more frequent in non-survivors. No statistically significant difference was observed in the rate of death between patients with a diagnosed neurodegenerative disorder compared to the rest of the population ($P=1.000$) or in those living in long-term facilities ($P=0.663$).

Male sex, chronic kidney disease, desaturation, dyspnea as well as lymphocytes count, CRP, and serum creatinine at admission were included in a multivariate binary logistic regression model to identify associated factors of death (**Table 4**). Two variables were independently associated to death at 21 days. CRP at admission: (OR=1.013, $P=0.004$) and chronic kidney disease (OR=4.631, $P=0.025$).

Discussion

In this study, we aimed at describing specific clinical features and prognostic factors of mortality in demented patients. Confusion with or without general symptoms is the most frequent initial presentation and was not a predictive factor of death while history of chronic kidney disease and CRP level at admission were significantly associated with mortality.

Studies focusing on the neurological features of COVID-19 suggest that confusion occurred in 20–30% of hospitalized patients increasing to 60–70% for severe forms [11,12]. Older adults are more prone to experiencing confusion and delirium and dementia is the higher predisposing risk factor (OR from 2.3–4.7), before age, visual and hearing impairment, and polypharmacy [13,14]. Several series and cases reports highlight COVID-19 infection presenting as isolated and persistent confusion [15–17] and as a factor of negative outcome [18]. In our serie, confusion & delirium occurred independently of the severity of the COVID-19 infection and did not appear as a factor of negative outcome. The discrepancy between all these results could be explained by the high prevalence of confusion in our cohort that might have not allowed to study his weight on prognosis.

Overall, the prevalence of confusion and delirium was higher than the one reported in older adults admitted with non-COVID-19 pneumonia [19,20] and in patients admitted with severe acute respiratory syndrome (SARS) and Middle East respiratory syndrome (MERS) [21].

The pathophysiological explanations remain unclear. COVID-19 could have a direct neuronal toxicity through CNS invasion [22]. Cognitive symptoms could also indirectly be related to neuroinflammation, corresponding to "sickness behavior" to which demented subjects have been shown to present with increased vulnerability [23]. An increase of neurofilaments lights and Glial Fibrillary Acid Protein, respectively reflecting neuronal injury and glial activation has been observed in patients with moderate

and severe COVID-19 with or without dementia [24]. The neuronal and synaptic fragility in demented patients may be particularly prone to injury induced by COVID-19, either through direct infectious lesion or through indirect inflammatory mechanisms. Another frequently reported neurologic symptom in our series is falls, as already reported [5,25]. General risk of fall is high in demented older adults; gait impairment and falls are more prevalent in dementia than in normal aging and are related to the severity of cognitive impairment [26]. Rupture of homeostasis in the context of viral infection would account for their frequent occurrence [27].

Beside confusion, delirium and falls, a broad range of neurological complications, caused directly or indirectly by the virus, including infectious, para-infectious, and post-infectious encephalitis, stroke related to coagulopathy, and acute neuropathies have been reported [28].

Stroke and seizure were observed during disease evolution for respectively 3 of our patients, at a similar rate as the one observed in cases of a cohort of around 5000 subjects [29]. Case fatality rate at 21 days was 22.4%, in line with previous findings in a large cohort [2,5]. A study by Canavelli *et al* [30] evaluating the prevalence of dementia in a random sample of confirmed COVID-19 infected patients, found that patients with dementia accounted for 15.8% of overall COVID-19 related death. In a meta-analysis of the mortality of individuals with dementia was increased compared to not demented subjects (OR=5.17)[31]. More specifically, Matias-Guiu *et al* have shown that AD patients showed a higher risk of death in COVID-19 than patients with fronto-temporal dementia. One explanation could be that the APOE e4 genotype (the highest risk factor for AD) has been reported as a predictive factor of severe COVID-19 [19] and death [20]. In our series, we did not find any difference between patients with a diagnosed dementia (AD, vascular dementia, alcohol related cognitive impairment) and patients with unexplored cognitive impairment.

History of chronic kidney disease and high CRP at admission were independently associated with death. Our overall data are in accordance with previous reports in which male sex, multiple comorbidities, elevated CRP and low lymphocyte count were observed in the majority of COVID-19 deaths [3,32]. None of our patient was deemed suitable candidate for ICU or invasive ventilation. Patients received variable treatment, associating antibiotherapy, rarely antiviral treatment, high flow oxygen and corticosteroids or immunomodulatory drugs for a small number. Given the size of our cohort, we were not able to assess efficiency or draw any recommendation. Therapeutic essays have been ongoing regarding the potential beneficial effect of serotonin reuptake inhibitor (SSRI) through modulation inflammatory response during sepsis[33]; no difference in outcome was observed regarding treatment by antidepressant in our cohort.

All in all, in the actual context of a second wave of COVID-19, this work demonstrates that special attention should be given to demented patients who manifest confusion with acute behavioral changes and falls, with or without asthenia, fever and lymphopenia.

Limitations

This study has several limitations. Our studied population was drawn from geriatric hospitalized samples of demented patients in need of hospital care, limiting its generalization. Patients presenting with atypical symptoms and lacking respiratory symptoms or fever may not have been identified as COVID cases and included in our analysis. Due to low group numbers, we could not assess the difference in severity and mortality between the different etiologies of dementia. We focused on short-term mortality; long term follow-up could allow us to identify additional prognostic factors.

Conclusion

In our cohort of individuals with dementia, we have observed that COVID-19 was characterized by atypical presentation with prevalent nonspecific symptoms at the initial phase of the disease, namely persisting confusion and behavioral disorders as well as frequent falls. Early recognition of COVID-19 in demented adults should help provide early treatment and adequate care and isolation and COVID-19 testing should be considered in front of any significant change from baseline.

Declarations

Ethics approval and consent to participate

We obtained the required approval from the Commission Nationale Informatique et Liberté (CNIL) to collect anonymized data.

Consent for publication

Not applicable.

Availability of data and material

The full datasets used during the current study are available from the corresponding author on reasonable request.

Competing interests

The authors declare that they have no competing interests

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Author's Contributions

Concept & design: AV, EM, LVG, CP.

Data acquisition, analysis and interpretation of the data: all authors.

Statistical analysis: AV, EM, JD.

Drafting the manuscript: AV, EM, CP.

Writing-review & editing: all authors.

Supervision of the study: CP, LVG.

AV and EM have full access to all the data in the study and take full responsibility for the integrity of the data and the accuracy of the data analysis.

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Tables

Table 1- Characteristics of included patients

	Total (n=125)	Survivors (n=97)	Non- survivors (n=28)	<i>p</i>
Age	86 (82-91)	87 (82-91)	86 (82-93)	0.815
Male sex	41.6% (52)	36.1% (35)	60.7% (17)	0.029
Admission from				0.663
Home	59.2 (74)	57.7% (56)	64.3% (18)	
Long-term care facility	40.8% (51)	42.3% (41)	35.7% (10)	
Medical history				
Comorbidities	91.2% (114)	91.8% (89)	89.3% (25)	0.709
>2 Comorbidities	67.2% (84)	67.0% (65)	64.3% (18)	0.822
Charlson Comorbidity Index	6 (5-7)	6 (5-7)	6 (5-7)	0.292
Neurological status				
<i>Etiology of dementia</i>				
Unspecified cognitiv impairment	60.0% (75)	59.8% (58)	60.7%(17)	1.000
Diagnosed dementia	40.0 % (50)	40.2% (39)	39.3% (11)	1.000
Alzheimers Disease	10.4% (13)	11.3% (11)	7.1% (2)	0.731
Vascular Dementia	7.2% (9)	6.2% (6)	10.7% (3)	0.418
Other causes of dementia	7.2% (9)	5.2% (5)	14.3% (4)	0.113
Parkinson disease, parkinsonian syndromes	9.6% (12)	10.3% (10)	7.1% (2)	1.000
Psychosis	3.2% (4)	4.1% (4)	0.0% (0)	1.000
Alcohol related dementia	2.4% (3)	3.1% (3)	0.0% (0)	1.000
<i>Neurological symptoms& history</i>				
Mood disorder	31.2% (39)	33.0% (32)	25.0 % (7)	0.640
Psychotic symptoms	12.0% (15)	12.4% (12)	10.7% (3)	1.000
History of stroke	32.0% (40)	32.0% (31)	32.1% (9)	1.000
History of fall	38.4% (48)	36.1% (35)	46.4% (13)	0.373
Walk	70.4% (88)	70.1% (68)	71.4% (20)	1.00
Psychotrop Treatment	60.0% (75)	60.8% (59)	57.1% (16)	0.826
Anxiolytic	30.4% (38)	32.0% (31)	25.0 % (7)	0.642
Antidepressant, mood regulator	35.2% (44)	35.1% (34)	35.7% (10)	1.000
Neuroleptics	12.0 % (15)	11.3% (11)	14.3% (4)	0.742
Non neurological comorbidities				
Cardiovascular disease	40.8% (51)	38.1% (37)	50.0% (14)	0.383
Arterial hypertension	80.0% (100)	79.4% (77)	82.1 % (23)	1.00
Hypercholesterolemia	48.8% (61)	46.4% (45)	57.1% (16)	0.392
Atrial fibrillation	24.8% (31)	24.7% (24)	25.0 % (7)	1.00
Coronaropathy	16.0% (20)	14.4% (14)	21.4 % (6)	0.388
Cardiac insuffisancy	24.0% (30)	25.8% (25)	17.9% (5)	0.460
Thrombosis	9.6% (12)	11.3% (11)	3.6% (1)	0.296
COPD	8.8% (11)	6.2% (6)	17.9% (5)	0.068
Respiratory insuffisancy	3.2% (4)	3.1% (3)	3.6% (1)	1.000
Diabete	16.8% (21)	15.5% (15)	21.4 % (6)	0.566
Denutrition	33.6% (42)	37.1% (36)	21.4 % (6)	0.172
Malignancy	9.6% (12)	11.3% (11)	3.6% (1)	0.296
Chronic liver disease	1.6% (2)	2.1% (2)	0.0% (0)	1.000
Chronic renal disease	16.0% (20)	11.3% (11)	32.1% (9)	0.017

Data are median (IQI), %(n). P values >0.05 [**Bold**] indicates significant differences between survivors and non-survivors.

*Other causes of dementia: included mixed dementia (AD & vascular dementia, n=7), Fronto temporal dementia (n=1) and vascular dementia and alcohol-related dementia (n=1)

COPD: Chronic Obstructive Pulmonary Disease

Table 2- COVID clinical symptoms and laboratory results

Symptoms at admission	Total (125)	Survivors (n=97)	Non survivors (n=28)	P
Fever	72.8% (91)	71.1 % (69)	78.6% (22)	0.482
Cough	49.6% (62)	51.5 % (50)	42.9% (12)	0.521
Expectorations	21.6% (27)	18.6 % (18)	32.1% (9)	0.190
Polypnea	51.2% (64)	42.3% (41)	82.1% (23)	<0.0001
Desaturation	50.4% (63)	43.3% (42)	75.0% (21)	0.002
Minimum Saturation	90 (88-92)	90 (88-92)	90 (88-91)	0.167
Digestive symptoms	16.8% (21)	18.6% (18)	10.7% (3)	0.400
Asthenia	76.8 % (96)	72.2% (70)	92.9% (26)	0.023
Myalgia	15.2 % (19)	15.5% (15)	14.3% (4)	1.000
Anorexia	56.8% (71)	53.6% (52)	67.9% (19)	0.201
Headache	4.0% (5)	4.1% (4)	3.6% (1)	0.688
Confusion, delirium	82.4% (103)	81.4% (79)	85.7% (24)	0.780
Initial fall	35.2% (44)	33.0 % (32)	42.9% (12)	0.373
Laboratory findings				
Hemoglobin, g/dL	12.1 (10.9-13.08)	12.2 (10.9-13.3)	11.6 (10.3-12.7)	0.152
Lymphocytes count(mm3)	0.99 (0.73-1.29)	1.0 (0.8-1.3)	0.9 (0.6-1.2)	0.032
Lymphopenia, %	84.8% (106)	85.6% (83)	82.1% (23)	0.544
White blood cell count (mm3)	5.8 (3.9-7.9)	5.4 (3.9-7.1)	6.2 (4.1-8.7)	0.070
Platelets count (mm3)	207 (164.3-265)	210(170-261)	202(150-283)	0.995
Thrombopenia,%	16.0% (20)	13.4% (13)	25.0% (7)	0.146
Initial C-reactive protein, mg/dL	30.85 (8.75-84.75)	25.8(8.0-64.5)	76.0 (33.0-44.0)	<0.0001
Initial Creatinine, µmol/L	78 (64-103.8)	74.0 (63.5-94.0)	127.0 (70.0-159.0)	0.003
Aspartate aminotransferase, U/L	36 (24-54)	35 (24-53)	46 (32-61)	0.053
Alanine aminotransferase, U/L	18 (13-31)	17.00 (13.00-31.00)	23.50 (12.75-34.75)	0.438
Albumin, g/L	30 (26-34)	30.8(27-34)	28.5 (23-33)	0.088
Troponin positivity, % [#]	35.5 % (22)	30.4% (14)	50.0% (8)	0.032

Data are presented as median (IQR), %(n). P values >0.05 [**Bold**] indicates significant differences between survivors and non-survivors. [#]:Troponine was available for 62 subjects.

Table 3- Treatment, complication and outcome

	Total	Survivor (n=97)	Non-survivors(N=28)	P
Covid Severity*				0.001
Mild	37.6% (47)	48.5% (47)	0.0% (0)	
Moderate	27.2% (34)	35.0% (34)	0.0% (0)	
Severe	35.2% (44)	16.5% (16)	100% (28)	
Treatment				
Oxygenotherapy	60.0% (75)	50.5% (49)	92.9% (26)	<0.0001
Maximal flow (L/min)	3.0 (2.0-6.0)	3.0 (2.0-6.0)	5.0 (2.5 -15.0)	0.018
Antibiotherapy	61.6% (77)	57.7% (56)	75.0% (21)	0.074
Combination antibiotherapy	33.6% (42)	29.9% (29)	46.4% (13)	0.082
Betalactamine	54.4 % (68)	50.5% (49)	67.9% (19)	0.079
Macrolides	37.6% (47)	35.1% (34)	46.4% (13)	0.279
2nd line antibiotherapy	17.6% (22)	14.4% (14)	28.6% (8)	0.096
Steroids	8.0% (10)	8.2% (8)	7.1% (2)	1.000
Hydroxychloroquine	16.8% (21)	16.5% (16)	17.9% (5)	1.000
Neuroleptics	4.8% (6)	5.2% (5)	3.6% (1)	0.678
Complications				
ARDS	25.6% (32)	6.2% (6)	92.9% (26)	<0.0001
Cardiac injury	13.6% (17)	9.3% (9)	28.6% (8)	0.024
ACKI	16.8% (21)	12.4% (12)	32.1% (9)	0.021
Bacterial surinfection	14.4% (18)	11.3% (11)	25.0% (7)	0.122
Behavioral disorder	19.2% (24)	21.6% (21)	10.7% (3)	0.278
Diabetic acidocetosis	2.4% (3)	3.1% (3)	0.0% (0)	1.000
Seizure	2.4 % (3)	2.1% (2)	3.6% (1)	0.536
Stroke	2.4% (3)	2.1% (2)	3.6% (1)	1.000
Outcome	Total	Survivor	Non survivor	
Total mortality	22.4% (28)			
Delay admission-death		8.5 (3.7-16)		
Lengh of stay	16 (11-19.7)			
Discharge home before end of followup	40.2% (39)			

Data are median (IQR), %(n). P values >0.05 [**Bold**] indicates significant differences between survivors and non-survivors.

*COVID severity defined according to WHO guidelines

ACKI: Acute Kidney Injury; ARDS: Acute Respiratory Distress Syndrome

Table 4-Predictive factors of mortality at 21 days of diagnosis

	Exp(B)	95% C.I.for EXP(B)		Sig.
		Lower	Upper	
Male sexe	1.120	0.379	3.310	0.838
Chronic renal disease	4.631	1.215	17.649	0.025
Desaturation	1.879	0.418	8.452	0.411
Dyspnea	2.100	0.452	9.765	0.344
Lymphocytes	1.096	0.649	1.852	0.732
Creatinine	0.999	0.992	1.007	0.850
CRP	1.013	1.004	1.021	0.004

Multivariate binary logistic regression for identifying predictive factors of outcome. Troponine was excluded of the analysis due to missing data. P<0.05 indicates significance [**Bold**]

95%CI: 95% Confidence Interval. CRP: C-reactive Protein