

Pattern of SARS-CoV-2 infection among dependant elderly residents living in long-term care facilities in Marseille, France, March-June 2020

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Tran Duc Anh Ly, Didier Zanini, Vincent Laforge, Sylvie Arlotto, Stephanie Gentile, et al.. Pattern of SARS-CoV-2 infection among dependant elderly residents living in long-term care facilities in Marseille, France, March-June 2020. International Journal of Antimicrobial Agents, 2020, 56 (6), 10.1016/j.ijantimicag.2020.106219. hal-03149233

HAL Id: hal-03149233 https://amu.hal.science/hal-03149233

Submitted on 15 Dec 2022

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- 1 Pattern of SARS-CoV-2 infection among dependant elderly residents living in long-term
- 2 care facilities in Marseille, France, March-June 2020
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- **Running title:** dependant elderly residents and COVID-19
- 40 **Keywords**: COVID-19; SARS-CoV-2; elderly resident; long-term care facility; mass testing;
- 41 hydroxycloroquine.

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- 42 **Abstract (250/250 words)**
- 43 **Objectives.** We aimed to report the results of SARS-CoV-2 PCR-based screening campaigns
- 44 conducted on dependent elderly residents (compared to staffs members) in long-term care
- 45 facilities (LTCFs) in Marseille, France, and the follow-up of positive cases.
- 46 **Methods.** Data from 1,691 elderly residents and 1,000 members of staff were retrospectively
- 47 collected through interviewing the medical teams in 24 LTCFs and using the hospitals'
- 48 electronic health recording systems.
- **Results.** Elderly residents were predominantly female (64.8%) with a mean age of 83.0 years.
- 50 SARS-CoV-2 detection among residents (226, 13.4%) was significantly higher than among
- staff members (87, 8.7%), with p<0.001. Of the 226 infected residents, 37 (16.4%) were
- detected on a case-by-case basis due to their COVID-19 symptoms and 189 (83.6%) were
- detected through mass screening. Most (77.0%) had possible COVID-19 symptoms, including
- respiratory symptoms and signs (44.5%) and fever (46.5%) and 23.0% were asymptomatic. 116
- 55 (51.4%) patients received a course of oral hydroxychloroquine and azithromycin (HCQ-
- AZM) for ≥ 3 days, and 47 (20.8%) died. Through multivariate analysis, the death rate was
- positively associated with being male (30.7%, vs. 14.0%, OR=3.95, p=0.002), being older than
- 85 years (26.1%, vs. 15.6%, OR=2.43, p=0.041), and receiving oxygen therapy (39.0%, vs.
- 59 12.9%, OR=5.16, p<0.001) and negatively associated with being diagnosed through mass
- screening (16.9%, vs. 40.5%, OR=0.20, p=0.001) and receiving HCQ-AZM treatment ≥ 3 days
- 61 (15.5%, vs. 26.4%, OR=0.37, p=0.02).
- 62 **Conclusion.** High proportion of asymptomatic COVID-19 patients and independent factors
- 63 for mortality suggests that early diagnosis and treatment of COVID-19 patients in LTCFs may
- be effective in saving lives.

Introduction

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In France, as of 2 June 2020, 10,350 elderly residents living in long-term care facilities or medical-social establishments had died from COVID-19 (27.6% lethality rate) accounting for 55.6% of COVID-19 deaths in France [1]. Similar pictures have also been reported in many European countries [2] and worldwide [3]. The prevalence of chronic conditions such as cardiovascular diseases, hypertension and diabetes mellitus is high among elderly persons living in long-term care facilities; coronavirus disease (COVID-19) in this population may therefore have severe outcomes with a high mortality rate [2, 4, 5]. Other drivers of mortality among elderly persons living in long-term care facilities already include type of facility, the amount of persons visiting the facilities during the week previous to lockdown, staff ratios [3], and lagged infection in staff members [6]. The treatment of COVID-19 has been the subject of widespread controversy, in particular with regards to the use of hydroxychloroquine (HCQ) [7]. It appears to us that some of the elements of the controversy are the heterogeneity of protocols using HCQ, with doses ranging from 800 mg to 1200 mg per day, the duration of treatment, whether or not it is combined with azithromycin (AZM), and the stage of the disease at which patients were being treated. It can be considered that there is a purely viral phase of the disease with a more or less strong immune response, which can become predominant in what has been referred to as the cytokine storm, followed in a number of cases, by necrotic lesions, linked to pulmonary infarctions [8]. Furthermore, mortality depends very significantly on age and thus, in Europe, almost all deaths have been among persons over the age of 60, with more than 50% in persons over the age of 85 years [9]. Under these conditions, it is very difficult to carry out comparative studies addressing the effect of HCQ on COVID-19-associated deaths. Very few randomised studies have been conducted and their interpretations have also led to heated

89 debate. To assist the debate, we believe that it may be important to assess whether there is a

clear reduction in mortality in the most at-risk groups.

In Marseille, over a period of approximately two months, we were able to test and treat COVID-19 patients in long-term care facilities (Etablissement d'Hébergement pour Personnes Agées Dépendantes –EHPAD) with a combination of HCQ-AZM, as we have described it on several occasions [8, 10-12]. The objective of this study was to estimate the prevalence of SARS-CoV-2 carriage among elderly residents and among staff members working in 24 EHPAD, Marseille, France. We also aimed to estimate the lethality among elderly residents treated in these EHPADs and informally compare it to the lethality of patients in these EHPAD

who were not treated s and the general lethality of patients in EHPADs in France.

Methods

Setting, study design and population

SARS-CoV-2 cross-sectional mass screening campaigns were conducted among residents and staff members from 24 long-term care facilities (EHPADs) in Marseille, between 24 March and 2 June 2020. In some centres, screening campaigns were conducted following the diagnosis of confirmed COVID-19 cases in symptomatic patients who were sampled on a case-by-case strategy. In other centres, screening campaigns were conducted systematically. In all cases, screening campaigns were conducted following a request from the directors and medical staff of the long-term care facilities. Nasopharyngeal samples were processed for SARS-CoV-2 PCR testing at the Institut Hospitalo-Universitaire (IHU) Méditerranée Infection at Assistance Publique-Hôpitaux de Marseille (AP-HM), as previously described [13] or in private laboratories in Marseille, in some cases. Residents who tested positive were either i) treated at their long-term care facilities by local medical staff only, ii) treated at their long-term care facilities in coordination with the AP-HM Home Hospitalisation Unit (HHU), iii)

admitted to the IHU (day-care hospital or conventional units), or iv) transferred to AP-MH Intensive Care Units (ICU). For confirmed cases, information on demographics, chronic medical conditions, COVID-19 treatment and clinical data including fever, asthenia, anorexia and weight loss, respiratory symptoms and signs (cough, rhinorrhoea, dyspnoea, chest pain, acute respiratory distress syndrome) and death was collected retrospectively from interviews with the medical team of 24 long-term care facilities and the electronic health recording systems of the AP-HM.

Statistical methods

Statistical procedures were performed using STATA 11.1. We used Pearson's chi-square or Fisher's exact tests to compare differences between groups of patients where appropriate. A two-sided p-value of less than 0.05 was considered to be statistically significant. A separate logistic regression analysis was used to identify independent risk factors for SARS-CoV-2 death prevalence among all elderly residents testing positive for SARS-CoV-2. The results were presented by percentages and odds ratio (OR) with a 95% confidence interval (95%CI). The initial model included variables presenting a p-value <0.2. The stepwise regression procedure and likelihood-ratio tests were applied to determine the final model.

Results

Over the study period, 1,691 elderly residents and 1,000 staff members were tested (Table 1). For residents, the sex ratio (male to female) was 1:1.8 and the mean age (± standard derivation [SD]) was 83.0 (±10.6) years (ranging from 50 to 106 years). For staff members, the sex ratio was 1:3.5 and the mean age (± SD) was 40.8 (±12.8) years (ranging from 18 to 87 years). It should be noted that two religious staff members at one long-term care facility were aged 75 and 87, respectively.

Overall, 313 participants (of 2,691, 11.6%) were confirmed positive for SARS-CoV-2. The prevalence among residents (226 of 1,691, 13.4%) was significantly higher than among staff members (87 of 1,000, 8.7%), p<0.001). With regard to the housing facilities, at least one individual was positive in 11/24 (45.8%) centres with prevalence of SARS-CoV-2 detection ranging from 0% to 57.6% among residents and from 0% to 24.1% among staff members (Table 1). The lethality rate among residents was 20.8% while no deaths occurred among staff members (p<10⁻⁴).

Characteristics of 226 elderly residents testing positive for SARS-CoV-2 (Tables 2 and 3)

Of the 226 SARS-CoV-2-positive elderly residents, 37 were diagnosed on a case-by-case basis through selected sampling of patients with COVID-19 symptoms and 189 (83.4%) were detected through mass screening. Regarding co-morbidities, the most frequent chronic condition was hypertension (39.6%), followed by other cardiovascular diseases (37.1%), dementia (28.9%) and other mental disorders (23.6%). In terms of clinical findings, 77.0% had possible COVID-19 symptoms, including respiratory symptoms and signs (44.5%), and fever (46.5%); and 23.0% had no COVID-19 symptoms representing 24.8% (40/161) of individuals tested through mass screening (Table 2).

When it came to therapeutic management, 62 (27.4%) patients were managed within their long-term care facilities by local medical staff only, 117 (51.8%) were managed within their long-term care facility in collaboration with the HHU, 16 (7.1%) were admitted to IHU, and 31 (13.7%) were transferred to ICU. Overall, 116 (51.4%) patients received an oral HCQ (200 mg three times daily for ten days), and AZM (500 mg on day 1 followed by 250 mg daily for the next four days) for at least three days and were monitored as described in previous studies [10-12]. Of the 110 others (48.6%), one (0.4%) received a two-day course of HCQ-AZM, one (0.4%) received HCQ alone, 37 (16.4%) received AZM alone, and 71 (31.4) did not receive either drug. The prevalence of HCQ-AZM treatment for at least three days ranged from 0%-

87.5% according to the housing facilities. Other treatments are described in Table 2. A total of 161 179 patients survived (79.2%) and 47 (20.8%) died. 162 The baseline characteristics of 116 patients who received HCQ-AZM treatment for at least 163 three days compared with 110 patients who did not receive the treatment were largely similar 164 (Table 3). A higher proportion of patients with a history of stroke was observed in the treated 165 group (15.8%) compared with the untreated group (5.2%, p=0.04). 166 167 Table 4 shows the lethality rate among elderly residents with SARS-CoV-2 infection, according to demographics, chronic conditions, circumstance of diagnosis, type of medical management 168 of patients, use of HCO-AZM and housing facility effect according to prevalence of HCO-169 AZM treatment for at least three days in each housing facility. Under univariate analysis, death 170 from COVID-19 was significantly associated with male gender. In addition, patients who were 171 diagnosed on a case-by case basis due to their COVID-19 symptoms were more likely to die 172 (40.5%) than those diagnosed through systematic screening (16.9%). Finally, patients who 173 received oxygen treatment were more likely to die (39.0%) than those who did not receive such 174 175 a treatment (12.9%), by contract, patients who received HCQ-AZM treatment for at least three 176 days were less likely to die (15.5%) than those who did not receive such a treatment (26.4%). Through multivariate analysis, the death rate was positively associated with being male 177 178 (30.7% vs. 14.0%, OR=3.95 [1.65-9.44], p=0.002), being older than 85 years (26.1% vs. 15.6%,

OR=2.43 [1.04-5.69], p=0.041), and receiving oxygen therapy (OR=5.16 [2.26-11.76], p< 10^{-4})

and negatively associated with being diagnosed through mass screening (16.9%, vs. 40.5%,

OR=0.20 [0.08-0.53], p=0.001) and receiving HCQ-AZM treatment for at least three days

(OR=0.37 [0.17-0.86], p=0.02).

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Discussion

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In Marseille, the first case of COVID-19 in the general population was diagnosed on 3 March 2020 and the epidemic peaked during the first week of April and remained active until the end of the month. Our survey of long-term care facilities began when the entire French population was placed under a strict lockdown (17 March) and when the epidemic was active in Marseille. All long-term care facilities became confined environments with very strict restrictions being place upon visits. We found a 13.4% SARS-CoV-2 positivity rate among dependant elderly residents in Marseille that was significantly higher than the 5.4% positivity rate among all French dependant elderly residents according to a national survey (37,405 confirmed cases in an estimated 695,060 French dependant elderly residents, p<0.001, 2 June update) [1, 14]. We observed an overall 20.8% COVID-19 lethality rate among infected residents in Marseille that was significantly lower than that in all French long-term care facilities or medical-social establishments (27.7% lethality rate, p=0.026, 2 June update) [1]. The main drivers of mortality in Marseille residents were older age and male sex, as already reported in many studies [15]. In addition, systematic screening by PCR was identified as an independent protective factor against death from COVID-19. A symptom-based diagnostic strategy is less effective in long-term care facilities, most likely because elderly patients with comorbidities such as chronic respiratory or cardiovascular diseases may be unable to accurately report new symptoms suggestive of COVID-infection or may present with atypical symptoms that challenge medical staff [13, 17]. Furthermore, in our experience, more than 23% of SARS-CoV-2 infected residents had no symptoms at the time of sampling. A very high prevalence (around 80%) was observed in a cross-sectional study conducted on elderly residents living in 2074 Belgian long-term care facilities [18]. In this study, we show that there was a significant difference in lethality between patients treated with our standardised treatment and untreated patients, as already reported in study conducted among elderly patients living in a Spanish

public nursing home in the same period [14]. Treatment with HCQ alone was demonstrated to be associated with lower mortality in patients admitted with COVID-19 [20-23]. Another cohort study conducted among American patients with rheumatic conditions showed an association between long-term HCQ treatment and reduced COVID-19 lethality rate [24]. The potential mechanisms of HCQ in the decrease of mortality in COVID-19 might be its inhibitory effects upon the production of pro-inflammatory cytokines interleukin (IL)-1-β, TNF-α and IL-6, and chemokines (CCL2 and CCL3) involved in the recruitment of proinflammatory cells in the lung [25]. There are some limitations to our study. Our study population was not randomly and homogenously recruited. Data regarding demographics, chronic conditions and clinical status was not systematically documented. Frailty that has been shown to be a major risk factor for mortality in COVID-19 was not evaluated in this study due to its retrospective design [26]. The use of individual preventive measures was not documented. Nevertheless, we believe that even if there are biases, as in any comparative study including randomisation, these biases are relatively neutralised by the multifactorial study. Above all, we demonstrate that the mortality in patients treated in EHPADs in Marseille was half that of patients in nursing homes across France who, in most cases very likely did not receive specific treatment since its use is restricted to the hospital setting [27, 28]. We believe that focusing on the population with the highest mortality, to show a significant effect, is important and agree in this sense with several studies that have shown a reduction in mortality of 30 to 50% by HCQ-AZM in populations most at risk [29, 8].

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| 231 | Declarations |
|-----|---|
| 232 | Funding. This work was supported by the French Government under the "Investments for the |
| 233 | Future" programme managed by the National Agency for Research (ANR), Méditerranée |
| 234 | Infection 10-IAHU-03, and was also supported by Région Provence-Alpes-Côte d'Azur. This |
| 235 | work received financial support from the Fondation Méditerranée Infection. |
| 236 | Competing Interests. No potential conflict of interest relevant to this article was reported. |
| 237 | Ethical Approval: Ethical approvals were obtained from the Marseille Institutional Review |
| 238 | Board and Ethics Committee (N° 2020-028). |
| 239 | |
| 240 | Acknowledgements. The authors thank all healthcare workers, laboratory technicians and the |
| 241 | Marins Pompiers de Marseille for their efforts in the fight against COVID-19. Our thanks also |
| 242 | go to all directors and coordinating doctors of the 24 long-term care facilities for their support. |
| 243 | |
| 244 | Authors' contributions statement: |
| 245 | Writing – original draft: TD, PG |
| 246 | Writing – review & editing: TD, DZ, VL, SA, SG, HM, MF, DM, OQ, PM, AM, PL, GD, BL, |
| 247 | AM, AC, PA, PP, EC & DR |
| 248 | Conceptualization: PG |

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| 360 | Table 1. SARS-CoV-2 testing among residents and staff members at 24 long-term care |
|-----|--|
| 361 | facilities in Marseille, France, 27 March– 2 June 2020 |
| 362 | Table 2. Comorbidities, symptoms and signs, diagnostic and therapeutic management among 226 |
| 363 | elderly residents testing positive for SARS-CoV-2. |
| 364 | Table 3. Characteristics of group receiving HCQ-AZ for at least 3 days and untreated group. |
| 365 | Table 4. Associations between multiple factors and SARS-CoV-2 deaths among 226 infected |
| 366 | elderly residents (univariate and multivariate analysis) |
| 367 | |

Table 1. SARS-CoV-2 testing among residents and staff members at 24 long-term care facilities in Marseille, France, 27 March– 2 June 2020

| | | Resider | nts | No. (07.) | | Staff member | S No. (%) | | | То | tal |
|----------------------|---|------------|------------------|--|---------------|---------------------|--|--------------------------|--------------------------|------------|------------------|
| Characteristics | Date of mass testing | No. tested | No. (%) positive | No. (%) deaths among positive cases (lethality rate) | No. tested | No. (%) positive | deaths among positive cases (lethality rate) | p- value ¹ | p- value ² | No. tested | No. (%) positive |
| Total | | 1691 | 226 (13.4) | 47 (20.8) | 1000 | 87 (8.7) | 0 (0) | 4.10-4 | <10-4 | 2691 | 313 (11.6) |
| Centre (2691) | | | | | | | | | | | |
| Centre-01 | 01 April, 08 April, 19 April | 99 | 57 (57.6) | 17 (29.9) | 83 | 20 (24.1) | 0 (0) | 2.10-3 | 0.04 | 182 | 77 (42.3) |
| Centre-02 | 08 April, 19 April, 20 May | 112 | 50 (44.6) | 9 (18.0) | 71 | 17 (24.0) | 0 (0) | 7.10 ⁻³ | 0.053 | 183 | 67 (36.6) |
| Centre-03 | 20 April, 26 April, 04 May, 11 May, 18 May, 25 May, 02 June | 52 | 23 (44.2) | 2 (8.7) | 35 | 7 (20.0) | 0 (0) | 2.10 ⁻³ | N/A | 87 | 30 (34.5) |
| Centre -04 | 06 April, 21 April | 89 | 24 (27.0) | 8 (33.3) | 108 | 12 (11.1) | 0 (0) | 7.10 ⁻³ | 0.03 | 197 | 36 (18.3) |
| Centre -05 | 08 April, 29 April | 37 | 10 (27.1) | 3 (30.0) | 32 | 1 (3.1) | 0 (0) | 0.035 | N/A | 69 | 11 (16.0) |
| Centre -06 | 08 April, 17 April, 22 April | 230 | 45 (18.0) | 7 (15.6) | 180 | 15 (8.3) | 0 (0) | 2.10 ⁻³ | 0.18 | 410 | 60 (14.9) |
| Centre -07 | 02 Avril, 27 April, 25 May | 81 | 8 (9.9) | 0 (0) | 57 | 11 (19.3) | 0 (0) | 0.18 | N/A | 138 | 19 (13.8) |
| Centre -08 | 13 April, 06 May | 77 | 7 (9.1) | 1 (14.3) | 24 | 1 (4.2) | 0 (0) | 0.67 | N/A | 101 | 8 (7.9) |
| Centre -09 | 21 April | 54 | 0 (0) | N/A | 44 | 3 (6.8) | 0 (0) | 0.08 | N/A | 98 | 3 (3.1) |
| Centre -10 | 23 April | 46 | 1 (2.2) | 0 (0) | 12 | 0 (0) | N/A | N/A | N/A | 58 | 1 (1.7) |
| Centre -11 | 15 April | 118 | 1 (0.9) | 0 (0) | 60 | 0 (0) | N/A | N/A | N/A | 178 | 1 (0.6) |

| Centre -12 | 15 April | 66 | 0 (0) | N/A | 18 | 0 (0) | N/A | N/A | N/A | 84 | 0 (0) |
|--------------------|----------|-----------------|------------|-----------|------------|-------|-----|-----|-----|-------------|-------|
| Centre -13 | 28 April | 96 | 0 (0) | N/A | 39 | 0 (0) | N/A | N/A | N/A | 135 | 0 (0) |
| Centre -14 | 30 April | 45 | 0 (0) | N/A | 12 | 0 (0) | N/A | N/A | N/A | 57 | 0 (0) |
| Centre -15 | 17 April | 64 | 0 (0) | N/A | 27 | 0 (0) | N/A | N/A | N/A | 91 | 0 (0) |
| Centre -16 | 22 April | 48 | 0 (0) | N/A | 19 | 0 (0) | N/A | N/A | N/A | 67 | 0 (0) |
| Centre -17 | 25 April | 61 | 0 (0) | N/A | 29 | 0 (0) | N/A | N/A | N/A | 90 | 0 (0) |
| Centre -18 | 15 April | 52 | 0 (0) | N/A | 18 | 0 (0) | N/A | N/A | N/A | 70 | 0 (0) |
| Centre -19 | 27 April | 32 | 0 (0) | N/A | 24 | 0 (0) | N/A | N/A | N/A | 56 | 0 (0) |
| Centre -20 | 27 April | 29 | 0 (0) | N/A | 15 | 0 (0) | N/A | N/A | N/A | 44 | 0 (0) |
| Centre -21 | 24 April | 25 | 0 (0) | N/A | 11 | 0 (0) | N/A | N/A | N/A | 36 | 0 (0) |
| Centre -22 | 20 April | 53 | 0 (0) | N/A | 22 | 0 (0) | N/A | N/A | N/A | 75 | 0 (0) |
| Centre -23 | 14 April | 100 | 0 (0) | N/A | 52 | 0 (0) | N/A | N/A | N/A | 152 | 0 (0) |
| Centre -24 | 24 April | 25 | 0 (0) | N/A | 8 | 0 (0) | N/A | N/A | N/A | 33 | 0 (0) |
| Sex (2471) | | | | | | | | | | | |
| Female, n (%) | | 1,069 (64.8) | 135 (12.6) | 19 (14.1) | 646 (77.7) | | | | | 1705 (69.0) | |
| Male, n (%) | | 581 (35.2) | 91 (15.7) | 28 (30.8) | 185 (22.3) | | | | | 766(31.0) | |
| Age (years) (2556) | | | | | | | | | | | |
| Mean±SD | | 83.0±10.6 | 83.4±10.6 | 86.8±10.2 | 40.8±12.7 | | | | | 68.3±23.1 | |
| Range (min-max) | | 50-106 | 56-103 | 59-103 | 18-87 | | | | | 18-106 | |
| 18-34,n (%) | | 0 (0) | N/A | N/A | 326 (36.4) | | | | | 326 (12.8) | |
| 35-49, n (%) | | 0 (0) | N/A | N/A | 292 (32.6) | | | | | 292 (11.4) | |
| 50-59, n (%) | | 34 (2.1) | 3 (8.8) | 1 (33.3) | 236 (25.4) | | | | | 270 (10.6) | |
| 60-69, n (%) | | 189 (11.4) | 25 (13.2) | 3 (12.0) | 38 (4.2) | | | | | 227 (8.9) | |
| 70-79, n (%) | | 348 (21.0) | 46 (13.2) | 5 (10.9) | 1 (0.1) | | | | | 349 (13.7) | |
| 80-89, n (%) | | 552 (33.2) | 78 (14.1) | 16 (20.5) | 1 (0.1) | | | | | 553 (21.6) | |
| 90-99, n (%) | | 505 (30.3) | 67 (13.3) | 19 (28.4) | 0 (0) | | | | | 505 (19.8) | |
| >99, n (%) | | 34 (2.1) | 7 (20.6) | 3(42.9) | 0 (0) | | | | | 34 (1.3) | |
| | | | | | | | | | | | |

Abbreviation: N/A, not applicable;

¹Comparison of positive testing prevalence between resident group and staff member group.

²Comparison of lethality rates between infected resident group and infected staff member group.

³ Number of individuals for whom data was available.

Table 2. Comorbidities, symptoms and signs, diagnostic and therapeutic management among 226 elderly residents testing positive for SARS-CoV-2.

| Parameters | n (%) | |
|---|------------|--|
| Comorbidities (159) 1 | | |
| Hypertension | 63 (39.6) | |
| Cardiovascular diseases (other than hypertension) | 59 (37.1) | |
| Dementia | 46 (28.9) | |
| Mental disorder | 39 (23.6) | |
| Diabetes mellitus | 25 (15.7) | |
| Chronic lung diseases | 19 (12.0) | |
| Stroke | 17 (10.7) | |
| Cancer | 15 (9.4) | |
| Chronic neurological disorder | 12 (7.6) | |
| Obesity | 7 (4.4) | |
| Chronic kidney diseases | 7 (4.4) | |
| Asthma | 3 (1.9) | |
| Symptoms and signs (200) | | |
| Respiratory symptoms and signs | 89 (44.5) | |
| Fever | 93 (46.5) | |
| Asthenia, anorexia, weight loss | 21 (10.5) | |
| No COVID-19 symptoms | 46 (23.0) | |
| Circumstances of diagnosis (226) | | |
| Case-by-case testing in patients with COVID-19 symptoms | 37 (16.4) | |
| Mass testing | 189 (83.6) | |
| Medical management of patients (226) | | |
| Managed at long-term care facilities by local medical staff only | 62 (27.4) | |
| Managed at long-term care facilities in coordination with the HHU | 117 (51.8) | |

| Admitted to IHU | 16 (7.1) | |
|--|--------------|--|
| Transferred ICU | 31 (13.7) | |
| HCQ-AZM therapy (226) | | |
| At least a three-day course | 116 (51.4) | |
| two-day course | 1 (0.4) | |
| HCQ alone | 1 (0.4) | |
| AZM alone | 37 (16.4) | |
| No HCQ, no AZM | 71 (31.4) | |
| HCQ-AZM therapy at least a three-day course according to the housing | | |
| facilities (226) | | |
| Centre-07, n/N (%) | 7/8 (87.5) | |
| Centre-01, n/N (%) | 39/50 (78.0) | |
| Centre-02, n/N (%) | 43/57 (75.4) | |
| Centre -05, n/N (%) | 4/10 (40.0) | |
| Centre -06, n/N (%) | 14/45 (31.1) | |
| Centre -04, n/N (%) | 4/23 (17.3) | |
| Centre -03, n/N (%) | 4/24 (16.7) | |
| Centre -08, n/N (%) | 1/7 (14.3) | |
| Centre -10, n/N (%) | 0/1 (0) | |
| Centre -11, n/N (%) | 0/1 (0) | |
| Oxygen therapy (199) | 59 (29.7) | |
| Ceftriaxone or ertapenem therapy (199) | 63 (31.6) | |
| Low-molecular-weight heparin therapy (199) | 24 (12.1) | |

Abbreviation: HCQ, hydroxychloroquine; AZM, azithromycin; HHU, Home Hospitalisation Unit, Institut Hospitalo-Universitaire; ICU, Intensive Care Units.

¹ Number of individuals for whom data was available.

Table 3. Characteristics of group receiving HCQ-AZ for at least 3 days and untreated group.

| Charact | eristics | Patient receiving HCQ-AZ≥3 days N=116 | Patient not receiving HCQ-AZ≥3 days N=110 | p-value |
|-----------------------------|---------------|---|---|---------|
| Demographic factors (226) 1 | | | | |
| Gender | Female, n (%) | 70 (60.3) | 65 (59.1) | 0.85 |
| | Male, n (%) | 46 (39.7) | 45 (40.9) | |
| Age (years) ² | 50-85, n (%) | 66 (56.9) | 49 (44.5) | 0.07 |
| | >85, n (%) | 50 (43.1) | 61 (55.5) | |
| Chronic conditions (159) | | | | |
| Cardiovascular diseases | No, n (%) | 51 (62.2) | 49 (63.6) | 0.85 |
| | Yes, n (%) | 31 (37.8) | 28 (36.4) | |
| Hypertension | No, n (%) | 46 (56.1) | 50 (64.9) | 0.26 |
| | Yes, n (%) | 36 (43.9) | 27 (35.1) | |
| Dementia | No, n (%) | 61 (74.4) | 52 (67.5) | 0.34 |
| | Yes, n (%) | 21 (25.6) | 25 (32.5) | |
| Mental disorder | No, n (%) | 61 (74.4) | 59 (76.6) | 0.74 |
| | Yes, n (%) | 21 (25.6) | 18 (23.4) | |
| Diabetes mellitus | No, n (%) | 69 (84.1) | 65 (84.4) | 0.96 |
| | Yes, n (%) | 13 (15.9) | 12 (15.6) | |
| Chronic lung diseases | No, n (%) | 72 (87.8) | 68 (88.3) | 0.92 |
| | Yes, n (%) | 10 (12.2) | 9 (11.7) | |

| Stroke | No, n (%) | 69 (84.2) | 73 (94.8) | 0.04 |
|--|--|-----------|-----------|------|
| | Yes, n (%) | 13 (15.8) | 4 (5.2) | |
| Cancer | No, n (%) | 74 (90.2) | 70 (90.9) | 0.86 |
| | Yes, n (%) | 8 (9.8) | 7 (9.1) | |
| Chronic neurological disorder | No, n (%) | 76 (92.7) | 71 (92.2) | 0.9 |
| | Yes, n (%) | 6 (7.3) | 6 (7.8) | |
| Circumstances of diagnosis (226) | Case-by-case testing in patients with COVID-19 symptoms, n (%) | 23 (19.8) | 14 (12.7) | 0.1 |
| | Mass testing, n (%) | 93 (80.2) | 96 (87.3) | |
| Facility management of patients (226) ³ | In long-term care facilities only | 0 (0) | 62 (56.4) | N/A |
| | Other | 116 (100) | 48 (43.7) | |
| Oxygen therapy (199) | No, n (%) | 79 (68.1) | 61 (73.5) | 0.4 |
| | Yes, n (%) | 37 (31.9) | 22 (26.5) | |
| Ceftriaxone or ertapenem therapy (199) | No, n (%) | 81 (69.8) | 55 (66.3) | 0.59 |
| шстару (199) | Yes, n (%) | 35 (30.2) | 28 (33.7) | |
| Low-molecular-weight heparin therapy (199) | No, n (%) | 98 (84.5) | 77 (92.8) | 0.08 |
| шстару (199) | Yes, n (%) | 18 (15.5) | 6 (7.2) | |

Abbreviation: HCQ, hydroxychloroquine; AZM, azithromycin; NA, Not applicable.

¹ Number of individuals for whom data was available.

² Median of the variable is used for analysis.

³ Indication of HCQ-AZ treatment was compulsorily administrated in coordination with hospital.

Table 4. Associations between multiple factors and SARS-CoV-2 deaths among 226 infected elderly residents (univariate and multivariate analysis)

| Characteristics | | Deaths Survivors | Survivors | Univariate | e | Multivariate | | |
|--------------------------|------------------------|------------------|------------|------------------|---------|------------------|---------|--|
| | | N=47 | N=179 | OR [95%CI] | p-value | aOR [95%CI] | p-value | |
| Demographic facto | ors (226) ¹ | | | | | | | |
| Gender | Female, n (%) | 19 (14.0) | 116 (86.0) | Ref | | Ref | | |
| | Male, n (%) | 28 (30.7) | 63 (69.2) | 2.71 [1.40-5.24] | 0.003 | 3.95 [1.65-9.44] | 0.002 | |
| Age (years) ² | 50-85, n (%) | 18 (15.6) | 97 (84.4) | Ref | | Ref | | |
| | >85, n (%) | 29 (26.1) | 82 (73.9) | 1.90 [0.99-3.67] | 0.055 | 2.43 [1.04-5.69] | 0.041 | |
| Chronic conditions | (159) | | | | | | | |
| Cardiovascular | No, n (%) | 21 (21.0) | 79 (79.0) | Ref | | | | |
| diseases | Yes, n (%) | 12 (20.3) | 47 (79.7) | 0.98 [0.43-2.12] | 0.92 | | | |
| Hypertension | No, n (%) | 23 (24.0) | 73 (76.0) | Ref | | | | |
| | Yes, n (%) | 10 (15.9) | 53 (84.1) | 0.59 [0.26-1.36] | 0.22 | | | |
| Dementia | No, n (%) | 28 (24.8) | 85 (75.2) | Ref | | | | |
| | Yes, n (%) | 5 (10.9) | 41 (89.1) | 0.37[0.13-1.02] | 0.057 | | | |
| Mental disorder | No, n (%) | 25 (20.9) | 95 (79.1) | Ref | | | | |
| | Yes, n (%) | 8 (20.5) | 31 (79.5) | 0.98 [0.40-2.39] | 0.96 | | | |
| Diabetes mellitus | No, n (%) | 27 (20.2) | 107 (79.8) | Ref | | | | |
| | Yes, n (%) | 6 (24.0) | 19 (76.0) | 1.25 [0.45-3.43] | 0.66 | | | |
| Chronic lung | No, n (%) | 26 (18.6) | 114 (81.4) | Ref | | | | |

| diseases | Yes, n (%) | 7 (36.9) | 12 (63.1) | 2.55 [0.91-7.12] | 0.073 | | |
|--|--|-----------|------------|------------------|-------------------|-------------------|-------|
| Stroke | No, n (%) | 31 (21.8) | 11 (78.2) | Ref | | | |
| | Yes, n (%) | 2 (11.7) | 15 (88.3) | 0.47 [0.1-2.20] | 0.34 | | |
| Cancer | No, n (%) | 28 (19.4) | 116 (80.6) | Ref | | | |
| | Yes, n (%) | 5 (33.3) | 10 (66.7) | 2.07 [0.65-6.54] | 0.215 | | |
| Chronic neurological | No, n (%) | 30 (20.4) | 117 (79.6) | Ref | | | |
| disorder | Yes, n (%) | 3 (25.0) | 9 (75.0) | 1.30 [0.33-5.10] | 0.71 | | |
| Diagnostic and thera | apeutic management factors | 3 | | | | | |
| Circumstances of diagnosis (226) | Case-by-case testing in patients with COVID-19 symptoms, n (%) | 15 (40.5) | 22 (59.5) | Ref | | Ref | |
| | Mass testing, n (%) | 32 (16.9) | 157 (83.1) | 0.30 [0.14-0.64] | 0.002 | 0.20 [0.08-0.53] | 0.001 |
| Facility management of | In long-term care facilities only | 12 (19.4) | 50 (80.3) | Ref | | | |
| patients (226) | Other | 35 (21.3) | 129 (78.7) | 1.13 [0.54-2.35] | 0.74 | | |
| HCQ-AZM treatment for at least | No, n (%) | 29 (26.4) | 81 (73.6) | Ref | | Ref | |
| three days (226) | Yes, n (%) | 18 (15.5) | 98 (84.5) | 0.51 [0.26-0.99] | 0.047 | 0.37 [0.17-0.86] | 0.02 |
| Housing facility effect ³ (226) | >75% | 26 (22.6) | 89 (77.4) | Ref | | | |
| effect (226) | 25%-75% | 11 (20.0) | 44 (80.0) | 0.85 [0.38-1.89] | 0.7 | | |
| | <25% | 10 (17.9) | 46 (82.1) | 0.74 [0.33-1.67] | 0.48 | | |
| Oxygen therapy (199) | No, n (%) | 18 (12.9) | 122 (87.1) | Ref | | Ref | |
| | Yes, n (%) | 23 (39.0) | 36 (61.0) | 4.33 [2.1-8.89] | <10 ⁻⁴ | 5.16 [2.26-11.76] | <10-4 |
| Ceftriaxone or | No, n (%) | 26 (19.1) | 110 (80.9) | Ref | | | |

| ertapenem therapy (199) | Yes, n (%) | 15 (23.8) | 48 (76.2) | 1.32 [0.64-2.71] | 0.45 | |
|----------------------------------|------------|-----------|------------|------------------|------|--|
| Low-molecular- weight heparin | No, n (%) | 36 (20.6) | 139 (79.4) | Ref | | |
| therapy (199) | Yes, n (%) | 5 (20.8) | 19 (79.2) | 1.01 [0.35-2.90] | 0.97 | |

Abbreviation: Ref, Reference; NA, Not applicable; OR, Odds-ratio; aOR, adjusted Odds-ratio; HCQ, hydroxychloroquine; AZM, azithromycin

¹ Number of individuals for whom data was available.

² Median of the variable is used for analysis.

³ According to prevalence of HCQ-AZM treatment for at least three days among infected residents in each housing facility as seen in Table 2. Bold lines indicate the variables recruited in initial multivariate mode.