NEPHROLOGY - LETTER TO THE EDITOR



Prevalence of antibodies against SARS-CoV-2 in hemodialysis patients

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Editor,

Elderly patients with comorbidities, such as diabetes and hypertension, are more susceptible to severe COVID-19. Hemodialysis (HD) patients have suboptimal immunity [1, 2] in addition to the expanded risk of being in constant contact with medical facilities. Contingency strategies focus on identifying patients with specific symptoms, using PCR-based testing [3]. There is little information on the use of rapid serological tests to demonstrate the presence or absence of antibodies [4]. This strategy is readily accessible to the medical staff without formal laboratory training.

All patients in our HD unit were tested within the 10 days after the detection of 1 confirmed case by positive PCR test, on May 21st, 2020. Samples (n = 154) were obtained by pinprick using the COVID-19 IgG/IgM Rapid Test Cassette, (Healgen Scientific Limited Liability Company. Houston, USA), and these tests have sensitivity 90.11%, specificity 98.46%, and accuracy 95.31%.

The mean age was 54.3 years (19-87), 52.6% (n=81) were female, 61.7% attending twice a week, 32.9% thrice a week, and 2.7% once a week. The chronic kidney diseases (CKD) were: diabetic nephropathy (58.3%), hypertension (18.8%), glomerulopathies (4.7%), and neurogenic bladder (3.7%).

Out of the154 test that were performed, 18 patients were positive for COVID-19, 11 tested positive for IgG (61.1%%),

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3 for IgG and IgM (16.7%), and 4 had positive IgM only (22.2%) (Fig. 1), and 4 (22.2%) of these patients shared a HD session with the patient positive PCR. Only 33% of the antibody-positive patients presented at least 1 symptom with an average of 9.5 days before sample collection; predominantly cough, fever, dyspnea, and diarrhea, and 67% were pauci-symptomatic. Of the antibody-negative patients (n = 136), 81% (n = 110) were pauci-symptomatic and the remaining 19% (n = 26) referred at least 1 symptom in the last 5 days. Only the patient in the initial case had severe pneumonia, the rest of the patients with positive antibodies presented mild and moderate symptoms.

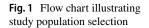
Antibody-positive patients had a mean age which was 52.6 years, 51% were female, the etiology of CKD diabetes 77.8%, 72.2% attended twice a week, and public transportations was used by 61.1%.

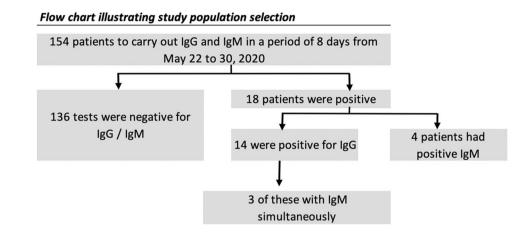
A previous study demonstrated the high transmissibility of SARS-CoV-2 by the rapid appearance of COVID-19 among 230 dialysis patients with a total of 37 HD patients and 4 members of the medical staff infected, post-performing tests after the first case [5]. In our center, 11 patients were identified with IgG antibodies, 3 with IgG and IgM, and 4 only with IgM. With this information, a stratification of patients could be made in every hemodialysis unit.

Of the 18 antibody-positive patients, 33% had mild symptoms compatible with COVID-19 compared to 19% of the patients with negative antibodies. The term pauci-symptomatic was used for patients with CKD in HD treatment, since this population frequently presents symptoms associated with volume overload, such as dyspnea and fatigue, for that many patients with CKD curse unnoticed SARS-CoV-2 infection.

There was no difference in symptoms between antibody positive vs negative patients, suggesting that mild symptoms are not an adequate screening tool for COVID-19 in this population. We acknowledge limitations; mainly, the lack of FDA approved antibody testing that at the time of the study was not available in our region.







In the present study, the prevalence of positive antibodies against SARS-CoV-2 in HD patients was 27.36%, 12 weeks after the first case was identified in the region. Determining the presence of antibodies could help to rapid diagnostic, attention, and severity classification in addition to stratify areas, personnel, and schedules for patients to avoid the rapid transmission.

Declarations

Conflict of interest The authors have declared that no conflict of interest exists.

Ethical approval All procedures performed were in accordance with the ethical standards research committee. Being an action carried out for early detection during the contingency of the COVID-19 pandemic, an approval number was not required. Informed consent was obtained from all individual participants included in the study.

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