Coronavirus Disease 2019 Case Fatality and Parkinson’s Disease

Case fatality rates (CFR) of the novel coronavirus disease 2019 (COVID-19) have been reported ranging from 0% to 40% among patients with Parkinson’s disease (PD). However, because of small sample sizes and the lack of large matched comparison groups, previous studies have not clarified whether PD is an independent risk factor for death. The goal of our study was to determine whether patients with PD had a higher COVID-19 CFR.

We compared COVID-19 CFR in patients with PD with a large, demographically matched population via the TriNetX COVID-19 research network, a health research database with deidentified medical records of >50 million patients mostly from the United States. As of July 15, 2020, this database listed 79,049 adult patients with COVID-19, 694 of whom had PD. On September 9, 2020, we extracted mortality data for this cohort. We included an 8-week delay to allow identified cases to resolve. Among 78,355 patients with COVID-19 without PD, 4290 died compared with 148 of the 694 patients with PD (5.5% non-PD vs. 21.3% PD; \( P < 0.001 \), \( \chi^2 \) test).

The non-PD and PD groups had different age distributions (median age 50 vs. 78), sex balances (female 55.3% vs. 39.8%), and racial compositions (Black 19.7% vs. 9.7%). In addition, CFRs from COVID-19 have been reported as higher in males versus females, Blacks versus Whites, and elderly versus younger patients. We accounted for these differences using logistic regression with age, sex, and race as covariates. This analysis revealed that the risk of dying from COVID-19 was significantly elevated in the PD group (odds ratio, 1.27; 95% confidence interval, 1.04–1.53; \( P = 0.016 \); Fig. 1A,B).

To assess residual confounders, we matched 5 patients with COVID-19 without PD to each patient with PD with the exact age, sex, and race. We then performed a conditional logistic regression and found that patients with PD had a significantly higher risk of dying from COVID-19 compared with patients without PD (odds ratio, 1.30; 95% confidence interval, 1.13–1.49; \( P < 0.001 \)). We further replicated the analysis with 1000 random matchings and found similar results with the effect being statistically significant in all but 2 replications.

In summary, we leveraged the TriNetX database and found that COVID-19-related CFR was increased in patients with PD, independent of age, sex, and race. These results are not without limitations. First, the TriNetX COVID-19 research network includes >40 healthcare organizations primarily in the United States. We were unable to account for confounding regional factors that could increase mortality. In addition, this database lacks information on key comorbidities. Second, CFRs from COVID-19 have decreased with increased access to testing. It will be important to conduct follow-up studies. Third, this study only reports an association between COVID-19-related mortality and a diagnosis of PD; however, in this context, designs better suited to causal inference are challenging. Finally, the TriNetX database does not include information on recovery. Despite these limitations, our results indicate that it will be...
critical to develop effective strategies whereby healthcare providers can prevent the transmission of COVID-19 while provid-
ing neurological care to patients with PD.

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