Session: P-14. COVID-19 Complications, Co-infections, and Clinical Outcomes

Background. Early data suggest that people with substance use disorder (SUD) who develop coronavirus disease 2019 (COVID-19) have increased intubation and mortality rates when compared to those without SUD. Information on other COVID-19-related complications in this population is limited. We evaluated COVID-19 outcomes in patients with and without SUD.

Methods. We created a retrospective cohort of patients with COVID-19 admitted to an urban safety net hospital from 3/16/2020 to 4/28/2020. Inclusion criteria were admission with laboratory-confirmed severe acute respiratory syndrome coronavirus 2 and age greater than 18 years. SUD included alcohol use disorder or heavy alcohol use as defined by the National Institute on Alcohol Abuse and Alcoholism, use of cocaine, non-prescribed opioids or amphetamines. Primary outcome was inpatient mortality. Secondary outcomes were clinical complications (intubation, secondary infections, renal failure, venous thromboembolism, stroke, hepatitis, myocardial infarct, multisystem organ failure) and resource utilization (length of stay, intensive care unit [ICU] admission, ICU days, readmission). We used multivariable regression to assess factors associated with mortality and length of stay, and univariate analyses for other outcomes.

Results. Of 409 included patients, 70 (17.1%) had SUD. Those with SUD were more likely to be male and have pulmonary disease or hepatitis C. There were no differences in other comorbidities, mean age or race/ethnicity. After multivariable analysis, SUD was not associated with mortality (aOR 1.60; 95% CI, 0.60-3.81). Similarly baseline oxygenation defined as the ratio of oxygen saturation to fraction of inspired oxygen (aOR 1.01; 0.13-11.30) and administration of immunomodulatory therapy (tocilizumab, sarilumab or anakinra) (aOR 1.41; 0.65-3.01) did not affect mortality. In contrast, age (aOR 1.06; 1.03-1.09), sex (aOR 2.30; 1.04-5.47) and obstructive sleep apnea (aOR 4.07; 1.64-9.66) were associated with mortality. We did not find any associations with secondary outcomes.

Conclusion. Our findings suggest that substance use alone may not increase COVID-19 adverse outcomes. Future studies should evaluate these results in the current period of improved COVID-19 therapy.

Disclosures. All Authors: No reported disclosures

334. Impact of Overall Dexamethasone Exposure on Development of Invasive Pulmonary Aspergillosis in Hospitalized Patients with COVID-19

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Background. Dexamethasone is one of the etiological agents that can be found during this complication. Spain started to vaccinate its population at the beginning of 2021. We noticed an important increase in Staphylococcus aureus infections and bacteremia during this period of time, leading us to study the relationship with previous vaccination.

Methods. In this case series we present a cohort of twenty patients with Staphylococcus aureus bacteremia (SAB) during the study period (January 1, 2021 through May 31, 2021), attended in our Institution (Hospital Nuestra Señora de Sonsoles, Ávila, Spain). We tried to establish or at least create the debate of a possible relationship between SAB and COVID-19 vaccine.

Results. From January 1, 2021 through May 31, 2021, 20 SAB were identified in our Institution. 13/20 patients were vaccinated (all of them with the mRNA vaccine type). 5/13 (38%) were male and 8/13 (62%) female. 10 of them (77%) received at least one dose of the vaccine before hospital admission, and 3 of them (23%) after admission. From the 10 previously COVID-19-vaccinated patients treated for SAB (CVPSAB), 4 died - 40% (2 deaths directly related to the SAB).

Conclusion. Although SAB may be a rare side effect after intramuscular injections or vaccines, it always implies an outstanding risk due to potential complications. Even if our study is not able to directly establish a link between SAB and previous vaccination, it implies a possible association between the vaccine injection and a threatening disease (SAB). We should be aware of this probable relationship, so that we can maximize preventive measures.

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335. Staphylococcus aureus Bacteremia as a Potential and Severe Complication from Intramuscular COVID-19 Vaccine Injection

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Session: P-14. COVID-19 Complications, Co-infections, and Clinical Outcomes

Background. Abscess formation and bacteremia following intramuscular injections are rare complications from vaccine injections, and they are most commonly seen in immunocompromised individuals. Staphylococcus aureus is one of the etiological agents that can be found during this complication. Spain started to vaccinate its population at the beginning of 2021. We noticed an important increase in Staphylococcus aureus infections and bacteremia during this period of time, leading us to study the relationship with previous vaccination.

Methods. In this case series we present a cohort of twenty patients with Staphylococcus aureus bacteremia (SAB) during the study period (January 1, 2021 through May 31, 2021), attended in our Institution (Hospital Nuestra Señora de Sonsoles, Ávila, Spain). We tried to establish or at least create the debate of a possible relationship between SAB and COVID-19 vaccine.

Results. From January 1, 2021 through May 31, 2021, 20 SAB were identified in our Institution. 13/20 patients were vaccinated (all of them with the mRNA vaccine type). 5/13 (38%) were male and 8/13 (62%) female. 10 of them (77%) received at least one dose of the vaccine before hospital admission, and 3 of them (23%) after admission. From the 10 previously COVID-19-vaccinated patients treated for SAB (CVPSAB), 4 died - 40% (2 deaths directly related to the SAB).

Conclusion. Although SAB may be a rare side effect after intramuscular injections or vaccines, it always implies an outstanding risk due to potential complications. Even if our study is not able to directly establish a link between SAB and previous vaccination, it implies a possible association between the vaccine injection and a threatening disease (SAB). We should be aware of this probable relationship, so that we can maximize preventive measures.

Disclosures. All Authors: No reported disclosures

336. COVID-19 and Pneumocystis jirovecii Pneumonia

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Session: P-14. COVID-19 Complications, Co-infections, and Clinical Outcomes

Background. More accounts of opportunistic infection in COVID-19 patients are emerging. At our institution, we identified 2 COVID-19 patients with Pneumocystis jirovecii pneumonia (PJP) opportunistic infection. This presentation reviews the literature to identify trends in patient characteristics, risk factors, and outcomes in this population.

Methods. A literature review was conducted using PubMed that identified 13 patients with PJP and COVID-19. We performed a review of the literature to identify trends in patient characteristics, risk factors, and outcomes in this population.

Results. Eleven patients were male. The average age was 56 years. All but 2 patients were immunocompromised. At time of PJP diagnosis, seven patients had already diagnosed HIV and one had known, well-controlled HIV. One patient had rheumatoid arthritis receiving leflunomide, 1 patient was a renal transplant recipient immunosuppressed on tacrolimus, and 1 patient had ulcerative colitis receiving budesonide, 1 patient was a renal transplant recipient immunosuppressed on tacrolimus, and 1 patient had ulcerative colitis receiving budesonide, and sulfasalazine, 2 patients had multiple myeloma whereby both were on lenalidomide. 1 patient was a renal transplant recipient immunosuppressed on tacrolimus, mycophenolate, and methylprednisolone, and 1 patient had chronic lymphocytic leukemia getting fludarabine, cyclophosphamide, and rituximab. Nine patients had positive COVID-19 and PJP tests performed within 7 days of one another. One patient tested positive for PJP 54 days into admission for COVID-19. This patient received high dose steroids and tocilizumab for initial COVID-19 infection. Three patients were re-hospitalized with PJP after a recent admission for COVID-19 pneumonia, with a mean time to readmission of 25 days. One of these 3 patients had no treatment for COVID-19, while 2 received steroids. Five of the total 15 patients (33%) died.

Conclusion. COVID-19 treatment with high dose steroids and tocilizumab can make patients vulnerable for opportunistic infection with PJP. Furthermore, COVID-19 is known to cause lymphopenia which may further increase this risk. A diagnosis of concomitant PJP can be especially challenging due to nearly identical radiographical findings. PJP beta-D glucan testing can be especially helpful in this situation, and there should be a low threshold for performing bronchoalveolar lavage.

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