Acute psychosis in COVID-19: Is it due to favipiravir treatment or acute viral illness?

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Case Report

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Abstract

Introduction:

In this report, we present a case of acute psychosis that occurred after the loading dose of favipiravir, one of the most prescribed antiviral drugs in many countries, in a patient with COVID-19.

Case presentation:

A 31-year-old female was readmitted to the emergency department after three days of sleep disturbance, intermittent paranoid delusions, auditory hallucinations, and agitation that started after the favipiravir treatment was initiated. A physical examination revealed disorganized speech, distorted orientation, and agitation. All laboratory tests, including drug screening and cerebrospinal fluid analysis, were normal. After excluding all the other causes of acute psychosis, favipiravir treatment was discontinued, antipsychotic treatment was started, and the patient was admitted to the psychiatric ward. The symptoms resolved on the second day of hospitalization. Although acute viral illness does seldomly cause psychosis in patients with increased inflammatory response, in the presented case, none of the inflammatory markers were positive and acute psychosis was attributed to the loading dose of favipiravir.

Conclusion

In conclusion, emergency department physicians should be aware of this rare adverse effect and prescribe cautiously to patients at a high risk of psychosis.

Introduction

Favipiravir is a selective RNA polymerase inhibitor that prevents viral replication in human cells [1]. It has been shown that favipiravir can be effective in treating infections caused by RNA viruses such as influenza, Ebola, rabies, norovirus, and SARS-Cov2 [2]. Although the use of favipiravir has been approved in many countries for the treatment of COVID-19 during the pandemic, its safety and effectiveness still remain unclear [3]. In an interim report of an observational study, the most common adverse events described were hyperuricemia (15%), abnormal liver function (7%), and diarrhea (1.4%). While no psychotic symptoms that could have been related to favipiravir treatment have been seen, agitation has been reported in one patient (0.05%) [4]. Herein, we present a case of psychosis that progressed gradually over four days in a patient receiving favipiravir.

Case Presentation

A 31-year-old female reporting five days of coughing and muscle aches was admitted to the emergency department (ED). The patient had normal vital signs and no remarkable past medical history. Although
the patient’s physical examination and routine laboratory tests were normal, a computerized tomography of the chest revealed mild peripheral infiltration in the right lung. The rRT-PCR examination of a nasopharyngeal swab yielded a positive result for SARS-CoV-2. Diagnosis of COVID-19 was confirmed and favipiravir (1600 mg twice on day 1 and 600 mg twice per day for another four days) was prescribed. On the fourth day of treatment, the patient was readmitted to the ED after reporting three days of sleep disturbance, intermittent paranoid delusions, auditory hallucinations, and agitation. According to the family, there was no history of trauma, drug or alcohol misuse, or the use of alternative medicine. Upon physical examination, the vital signs were normal, and the Glasgow Coma Scale value was 13 (E4M5V4), but orientation of place, person, and time was distorted. Speech was also disorganized. Further, during the physical examination, the patient became agitated, yelled at the ED staff, and attempted to attack the treating physician. Intramuscular haloperidol (10 mg) and chlorpromazine (25 mg) were administrated to sedate the patient. The laboratory tests were within a normal range. A brain CT and an MRI showed no pathology. No progression in pneumonia was detected by the chest x-ray. A urine drug screen was negative. A lumbar puncture was performed to exclude other central nervous system pathologies. Cerebrospinal fluid (CSF) analysis and cultures were negative. The meningitis/encephalitis panel from the CSF was clear. The patient was admitted to the psychiatric department. Favipiravir was discontinued and risperidone (2 mg once per day) and quetiapine (100 mg once per day) were initiated. The symptoms improved on the second day of admission and the patient was discharged on the fifth day of hospitalization. The acute onset of psychosis was attributed to favipiravir use. The patient did not have any symptoms after one-month follow-up.

Discussion

Psychotic symptoms associated with COVID-19 have been described in several reports [5, 6]. In these reports, it was postulated that increased inflammatory response resulting from COVID-19 could be the trigger for psychotic symptoms. Interestingly, all previously reported psychosis patients had increased inflammatory markers, especially C-reactive protein levels [5, 6]. In our case, inflammatory markers of the patient were negative upon both ED admissions, yet the psychotic symptoms acutely started on the second day of favipiravir administration. Although it was challenging to determine whether the symptoms were due to the new medication or acute viral illness, the mild course of the clinical findings from COVID-19, having normal laboratory and physical examination findings suggested that the psychotic symptoms were triggered by the new antiviral regimen.

Many of the drugs used to treat COVID-19 have been found to be correlated to neuropsychiatric symptoms, including chloroquine, hydroxychloroquine, umifenovir, interferon-alpha, and corticosteroids [7]. In one study, psychiatric reactions were reported only in two patients (1.7%) after favipiravir treatment [8]. However, it is not clear what type of psychiatric symptoms were seen in those patients. To our knowledge, this is the first report that describes a case of psychosis in a patient receiving favipiravir.

Conclusion
Favipiravir may lead to acute psychosis in patients with COVID-19. Emergency physicians should be aware of this rare side effect and prescribe cautiously, especially to those who have underlying psychiatric disorders.

Declarations

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Availability of data and material: Not applicable.

**Code availability:** Not applicable.

**Authors’ contributions:** MD was the treating physicians. MD collected the data. MD and IUO conceptualized the report. IUO drafted manuscript. MD and IUO critically reviewed the manuscript. Both of the authors approved the final version.

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