CASE REPORT

Rare case of intracranial hemorrhage associated with seoul virus infection diagnosed by metagenomic next-generation sequencing

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Abstract
Background: Seoul virus (SEOV) is a Hantavirus and the causative pathogen of Hemorrhagic Fever with Renal Syndrome (HFRS). Diagnosing SEOV infection is difficult because the clinical presentations are often undistinguishable from other viral or bacterial infections. In addition, diagnostic tools including serological and molecular assays are not readily available in the clinical settings.

Case Report: A 57-year-old male presented with fever and a sudden loss of consciousness in November 2019. Computed tomography (CT) scan showed subdural hematoma, subfalcine herniation, and brain infarction. He developed thrombocytopenia and elevated transaminases, but no rashes or obvious kidney damage. He reported having a rat bite. HFRS was suspected. The Hantavirus IgG was positive, and the metagenomic next-generation sequencing (mNGS) detected SEOV sequences directly in the blood.

Conclusion: This report highlights the importance of suspecting SEOV infection in febrile patients with thrombocytopenia and elevated liver enzymes despite the absence of hemorrhagic manifestations of skin and renal syndromes. Next-generation sequencing is a powerful tool for pathogen detection. Intracranial hemorrhage and brain infarction as extrarenal manifestations of HFRS are rare but possible as demonstrated in this case.

KEYWORDS
intracranial hemorrhage, mNGS, seoul virus

1 | BACKGROUND

Seoul virus (SEOV) is a member of Hantavirus family, which has three segments of negative-sense, single-stranded RNA genomes. The small (S) segment encodes a nucleoprotein, the medium (M) segment encodes 2 membrane glycoproteins (Gn and Gc), and the large (L) segment encodes an RNA-dependent RNA polymerase. SEOV infection is a worldwide public health threat in humans. The virus,
harbored by Rattus norvegicus and R. rattus rats, is one of major causative etiologies of Hemorrhagic Fever with Renal Syndrome (HFRS) in China. Clinically, the typical HFRS manifestations include five consecutive phases: febrile, hypotensive shock, oliguric, polyuric, and convalescent. The general clinical features include fever, myalgias, hemorrhage, thrombocytopenia, leukocytosis, renal failure, and shock in most severe cases. Elevation of liver enzymes is characteristic of HFRS caused by SEOV. But these presentations are non-specific for HFRS, which are often undistinguishable from other viral or bacterial infections. The serological assays to detect Hantavirus IgM and IgG offered by the state or regional Centers for Disease Control and Prevention (CDC) can confirm HFRS, but are not specific to SEOV. Molecular assays to detect SEOV viral RNA is also not available in most hospitals. Metagenomic next-generation sequencing (mNGS) is a powerful tool for the detection and identification of pathogens directly from the specimen. In this report, we describe a highly unusual case of SEOV infection diagnosed by mNGS in a patient with intracranial hemorrhage but without typical hemorrhagic manifestations of skin or renal syndromes.

2 | CASE REPORT

A 57-year-old previously healthy male was transferred to our hospital in November 2019 after a sudden loss of consciousness. Four days prior, the patient developed fever (maximum temperature 39°C) of unknown origin. He was admitted to a local hospital and was diagnosed with Dengue fever after the Dengue virus IgM was tested weakly positive. During the hospitalization, he suddenly lost consciousness and started vomiting. He had no significant medical history (particularly no anti-platelet or anticoagulant history), no recent travel history, and no history of cerebral trauma. Physical examination showed blood pressure of 161/82 mmHg, heart rate of 97 beats/minutes, respiratory rate of 25/minutes, and temperature of 37.8°C. He was in a coma and did not respond to voice. The Glasgow Coma Scale (GCS) score was 6 points. The diameter of his left and right pupil was 3.0mm and 4.0mm, respectively. The direct and indirect light reflex was slow. Limb movement was observed, and his chest, cardiac, and abdominal examination was normal. There were no rashes, petechia, ecchymoses, or other skin lesions. Blood tests performed 1 day before admission at the local hospital showed a white blood cell (WBC) count of 8.32 × 10^9/L, hemoglobin (Hb) level of 171 g/L, hematocrit of 52.5%, platelet count of 31 × 10^9/L, aspartate aminotransferase (AST) of 272 IU/L, alanine aminotransferase (ALT) of 225 IU/L, blood urea nitrogen (BUN) of 6.47 mmol/L, and serum creatinine of 83.4 µmol/L. Computed tomography scan of head indicated left subdural hemorrhage, subfalcine hernia, and brain infarction (Figure 1), there was no abnormality in the CT angiography (CTA). CT scan of whole abdomen suggested perirenal inflammation and chest CT showed slight bilateral inflammation of lower lungs (Figure 2). Surgical removal of subdural hematoma and decompressive craniectomy were performed immediately despite the low platelet count (20 × 10^9/L). He was then transferred to the intensive care unit (ICU). The patient received an immediate transfusion of one unit of platelet. He also received fluid resuscitation for hypotension and high lactic acid (8 mmol/L). Meanwhile, noradrenaline (0.3 µg/kg/min) was used to maintain his blood pressure. He had no oliguria. Peripheral blood tests upon ICU admission showed highly elevated WBC of 23.55 × 10^9/L, Hb level of 85 g/L, hematocrit of 24.5%, and platelet count of 42 × 10^9/L. Urinalysis was normal. Biochemical tests showed elevated liver enzymes with peak AST of 3320 IU/L, ALT of 1780 IU/L. The prothrombin time (PT) was 13.7 seconds, the activated partial thromboplastin time (APTT) was 49 (normal range 28-40 at our hospital) seconds and the fibrinogen concentration (Fib) was 2.89 G/L. Hypersensitive C-reactive protein was 8.14 mg/L, and procalcitonin was 1.49 ng/mL. Brain natriuretic peptide (BNP) was 215.54 pg/ml. Hepatitis A, B, C, and E viruses, HIV, Syphilis, Cytomegalovirus, and Epstein-Barr virus were all tested negative by serology. Dengue virus antigen, IgM and IgG, and RNA were also tested negative. Triglyceride was normal

![Figure 1](https://example.com/figure1.png)  Head CT on admission indicated subdural hematoma (red arrow), brain infarction (blue arrow), and subfalcine herniation (dark arrow)
Upon further communication with his family members, his son recalled that the patient had been bitten by a rat one month ago. Peripheral blood sample on ICU day 1 was sent for mNGS test and also sent to Guangzhou CDC for serological testing of leptospirosis and Hantavirus. On ICU day 3, the initial bone marrow biopsy showed scattered hemophagocytic cells. On ICU day 4, the mNGS performed by IngeniGen Biotechnology Inc using Illumina MiniSeq detected SEOV with 28 high-confidence sequence reads and mapped to an SEOV reference genome (S segment: NC_005236.1, M segment: NC_005237.1, L segment: NC_005238.1) (Figure 3). The important laboratory results prior to admission and after ICU admission were summarized in Table 1. Serology of Hantavirus by enzyme-linked immunosorbent assay (ELISA) was also positive for Hantavirus IgG, thus confirming the SEOV infection. Antibody titration test was not performed; therefore, the exact titer of the Hantavirus IgG was unknown. Blood, sputum, and urine cultures were all negative on ICU day 1. The sputum culture performed on ICU day 2 grew multi-drug resistance (MDR) Acinetobacter baumannii. Cerebrospinal fluid culture and blood culture performed on ICU day 11 also grew MDR A. baumannii. He was treated with meropenem and colistin for the bacterial infections. On ICU day 10, head CT indicated new intracranial hemorrhage (Figure 4). On ICU day 15, the patient suddenly developed shortness of breath, hypotension, tachycardia, and respiratory arrest, suggesting foramen magnum hernia. He received the mechanical ventilation. His CSF workup showed markedly increased red blood cells, suggesting an increased intracerebral hemorrhage. The patient died on ICU day 19.
3 | DISCUSSION

The initial differential diagnosis in this case was very challenging, as our patient presented with intracranial hemorrhage but without the three common causes including head trauma, hypertension, and arterial malformations. His presentation of fever, thrombocytopenia, and transaminitis promoted us to investigate an infectious cause. However, the CT scan of his chest did not reveal typical signs of infection, and the patient did not have other typical presentations of infections such as headache, chills, abdominal pain, and diarrhea. Although the abdomen CT scan demonstrated renal damage with perirenal inflammation, he did not have abnormal urination and his urinalysis was normal. Because dengue fever is prevalent in Guangzhou, it was highly suspected initially, especially after the IgM was weakly positive, but the follow-up serological and molecular tests for Dengue were all negative, suggesting the initial IgM was a false-positive result. All other conventional microbiological tests were also negative. mNGS is a genre of technologies that use what is known as shotgun sequencing of clinical samples or pure microbial cultures in which random samples of analyte DNA or RNA are surveyed en masse, which can be used for meaningful clinical diagnostics in microbiology. Fortunately, mNGS was able to detect SEOV directly in the blood which led to a definitive diagnosis confirmed by the positive Hantavirus IgG test.

The typically clinical manifestation of SEOV infection is HFRS, characterized by fever, hemorrhage, and impaired kidney function. The HFRS is more prevalent in Northern and Central Eastern China, but relatively less common in Southern China. HFRS incidence rate in Guangdong was 0.01 to 0.49/100,000. The disease peaks in autumn and winter season, which is consistent with the timing of this case (October – November). Since our patient had a history of rat bite, even though the clinical presentation was atypical: no hemorrhagic manifestations of skin and mucosal membrane and no renal syndromes, the HFRS was suspected.

The extrarenal manifestations of HFRS are rare but have been reported previously. Park KH, et al reported extrarenal manifestations involving the major organs occurred in one-third (23/73) of patients with HFRS during various stages, from the febrile phase to the diuretic phase. The spectrum of the extrarenal manifestations includes pancreato-biliary (8 cases), gastrointestinal (5 cases), cardiovascular (4 cases), and neurological (3 cases) manifestations.
cases), central nervous system (3 cases), retroperitoneal (1 case), intramural (1 case), respiratory (1 case), compartment syndrome (1 case), and hemophagocytic lymphohistiocytosis (HLH) (1 case). Few case reports showed cerebral involvement associated with HFRS including cerebral hemorrhage, cerebral edema and small subarachnoid, epidural and focal brain stem hemorrhages, and brain infarction. Unfortunately, in our case, the hematoma tissue and cerebrospinal fluid were not saved for serological and molecular tests for further investigation, which is the major limitation of this study.

4 CONCLUSION

This report highlights the importance of suspecting SEOV infection in febrile patients with thrombocytopenia and elevated liver enzymes despite the absence of hemorrhagic manifestations of skin and mucosal membrane and renal syndromes. mNGS can be a powerful tool to detect SEOV directly in the blood. Intracranial hemorrhage and brain infarction as extrarenal manifestations of HFRS are rare. Clinicians should be aware of various extrarenal manifestations of SEOV infections.

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CONFLICT OF INTEREST

The authors declare no conflict of interest.

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REFERENCES

1. Vaheri A, Strandin T, Hepojoki J, et al. Uncovering the mysteries of hantavirus infections. Nat Rev Microbiol. 2013;11(8):539-550.
2. Zhang S, Wang S, Yin W, et al. Epidemic characteristics of hemorrhagic fever with renal syndrome in China, 2006–2012. BMC Infect Dis. 2014;14:384.
3. Peters CJ, Simpson GL, Levy H. Spectrum of hantavirus infection: hemorrhagic fever with renal syndrome and hantavirus pulmonary syndrome. Annu Rev Med. 1999;50:531-545.
4. Kim YS, Ahn C, Han JS, et al. Hemorrhagic fever with renal syndrome caused by the Seoul virus. Nephron. 1995;71(4):419-427.
5. Caceres JA, Goldstein JN. Intracranial hemorrhage. Emerg Med Clin North Am. 2012;30(3):771-794.
6. Gu W, Miller S, Chiu CY. Clinical metagenomic next-generation sequencing for pathogen detection. Annu Rev Pathol. 2019;14:319-338.
7. Park KH, Kang YU, Kang SJ, et al. Experience with extrarenal manifestations of hemorrhagic fever with renal syndrome in a tertiary care hospital in South Korea. Am J Trop Med Hyg. 2011;84(2):229-233.
8. Alexeyev OA, Morozov VG. Neurological manifestations of hemorrhagic fever with renal syndrome caused by Puumala virus: review of 811 cases. Clin Infect Dis. 1995;20(2):255-258.
9. Giles RB, Sheedy JA, Ekman CN, et al. The sequelae of epidemic hemorrhagic fever; with a note on causes of death. Am J Med. 1954;16(5):629-638.

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