

# Possible re-infection of SARS-CoV-2 complicated by dengue virus co-infection: report of a rare case from Bangladesh

Wahiduzzaman M<sup>a</sup>, Rahim MA<sup>b</sup>

### ABSTRACT

*Re-infection by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) and co-infection by dengue virus and SARS-CoV-2 are possible. We report a case of dengue haemorrhagic fever, occurring in a young Bangladeshi man, who concurrently tested positive for SARS-CoV-2 infection by reverse transcriptase polymerase chain reaction (RT-PCR). Four months previously, he suffered a mild form of corona virus disease 2019 (COVID-19). This case is reported to make the physicians aware that, co-infections are possible in this COVID-19 pandemic, specially in dengue endemic regions and countries like Bangladesh.*

**Key words:** COVID-19, dengue haemorrhagic fever, re-infection, RT-PCR positivity.

*(BIRDEM Med J 2021; 11(1): 105-106)*

### INTRODUCTION

Starting from Wuhan, China in late 2019, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) has rapidly spread all over the world and claimed huge morbidity and mortality. These are not only due to corona virus disease 2019 (COVID-19) itself but also from other diseases including concurrent infections. In tropics and sub-tropical countries, dengue outbreaks are common during the monsoon. Initial clinical and laboratory features are indistinguishable for dengue and COVID-19.<sup>1</sup> Here, we present a case of dengue haemorrhagic fever, occurring in a young Bangladeshi male patient, who also tested positive for SARS-CoV-2 during the same febrile illness by reverse-transcriptase polymerase chain reaction (RT-PCR), four months after his first clinical event of COVID-19.

### CASE REPORT

A 34-year-old Bangladeshi male patient, with no recent travel history, presented with a 2-day history of high

grade, continued fever, headache, body ache, low back pain and restlessness. He took paracetamol and fexofenadine. Investigation reports revealed a positive natural structural protein 1 (NS1) for dengue virus and RT-PCR for SARS-CoV-2 from nasopharyngeal swabs. His initial platelet count was 1,70,000/cmm of blood. He was started with favipiravir, advised to stay home and monitor oxygen saturation. High resolution computed tomography (HRCT) scan of chest appeared normal. On day 4 of symptom onset, he developed wide spread maculo-papular rash. A repeat blood test revealed low platelet count (42,000/cmm) and he was hospitalized, where subsequently his platelet count further dropped to 28,000/cmm. He remained clinically stable throughout the hospital course; there was no bleeding manifestation, shock or hypoxia. His discharge medications included subcutaneous low molecular weight heparin (started after platelet count had reached over 1,00,000/cmm).

---

### Author information

- Miah Wahiduzzaman, Deputy Chief Medical Officer, Outpatient Department, BIRDEM General Hospital, Dhaka, Bangladesh.
- Muhammad Abdur Rahim, Associate Professor, Department of Nephrology, BIRDEM General Hospital, Dhaka, Bangladesh.

**Address of correspondence:** Miah Wahiduzzaman, Deputy Chief Medical Officer, Outpatient Department, BIRDEM General Hospital, Dhaka, Bangladesh. Email: miahwahiduzzaman@yahoo.com, muradrahim23@yahoo.com

**Received:** November 1, 2020

**Revision received:** November 12, 2020

**Accepted:** December 20, 2020

On June 23, 2020, he first reported with a 3-day history of fever, cough, sore throat and anosmia. That time, he tested positive by RT-PCR for SARS-CoV-2 and was treated at home as mild COVID-19 case, maintaining isolation protocols and took paracetamol, fexofenadine, azithromycin and single dose of ivermectin. He became symptom free in 5 days and tested negative for SARS-CoV-2 after 21 days. He remained in good health throughout the period until October 15, when he again developed febrile illness and subsequently received a diagnosis of dengue haemorrhagic fever and COVID-19.

On follow-up visit, one week after discharge, he was clinically stable and laboratory evaluation revealed positive IgG against dengue virus (15 days after fever onset, IgM and IgG were negative on day 5 after fever onset), improved alanine aminotransferase (151 from 444 U/L) and a normal platelet count and D-dimer level.

## DISCUSSION

Antibodies against SARS-CoV-2 are not long lasting and re-infection by SARS-CoV-2 is possible.<sup>2</sup> Cases of co-infection by dengue virus and SARS-CoV-2 are reported from Reunion Island<sup>1</sup> and Brazil.<sup>3</sup> Care should be taken to make a diagnosis of co-infection by dengue virus and SARS-CoV-2 depending up on serology, as false positive dengue serology in COVID-19 cases are reported.<sup>4</sup>

It is now evident that co-infections are possible by dengue virus and SARS-CoV-2. Some authors warn about the worse combination, while others predict some protective role, at least short-lived ones. Important is to treat the patient by supportive measures, with meticulous monitoring for possible organ involvement/failure, protection of caregivers and hospital staffs from SARS-CoV-2 infection.<sup>5,6</sup>

Repeat infections by SARS-CoV-2 are reported. In our clinical practice, clinical manifestation and second time RT-PCR positivity for SARS-CoV-2 after recovery from the first event are documented but we are unable to go for genome sequencing as routine practice. This limits

the proof for re-infection in resource constrain settings, as in the present case. It is also suggested that, in absence of clinical features, RT-PCR positivity from nasopharyngeal swabs does not necessarily mean persistent infection, re-infection or infectivity.<sup>7</sup>

As COVID-19 cases are increasing with evidences of second waves in some countries and regions, it will not be surprising to face increasing numbers of re-infection by SARS-CoV-2 and co-infection with dengue virus and SARS-CoV-2. A high index of clinical suspicion is warranted to have a better patient outcome and safety for health care workers.

**Authors' contribution:** MW diagnosed the case and planned publication. MAR did literature search and drafted the manuscript. Both authors read and approved the final manuscript for publication.

**Conflicts of interest:** Nothing to declare.

**Consent:** Informed written consent was obtained from the patient for publication of case report and accompanying images.

## REFERENCES

1. Verduyn M, Allou N, Gazaille V, Andre M, Desroche T, Jaffar M-C, et al. Co-infection of dengue and COVID-19: A case report. *PLoS Negl Trop Dis* 2020; 14(8): e0008476.
2. Tillett RL, Sevinsky JR, Hartley PD, Kerwin H, Crawford N, Gorzalski A, et al. Genomic evidence for reinfection with SARS-CoV-2: a case study. *Lancet Infect Dis* 2020 Published Online October 12, 2020 [https://doi.org/10.1016/S1473-3099\(20\)30764-7](https://doi.org/10.1016/S1473-3099(20)30764-7)
3. Bicudo N, Bicudo E, Costab JD, Castroa JALP, Barra GB. Co-infection of SARS-CoV-2 and dengue virus: a clinical challenge. *Braz J Infect Dis* 2020; 24(5): 452-4.
4. Yan G, Lee CK, Lam LTM, Yan B, Chua YX, Lim AYN, et al. Covert COVID-19 and false-positive dengue serology in Singapore. *Lancet Infect Dis* 2020 May; 20(5): 536.
5. Ridwan R. COVID-19 and dengue: a deadly duo. *Tropical Doctor* 2020; 50(3): 270-2.
6. Joob B, Wiwanitkit V. COVID-19 in medical personnel: observation from Thailand. *J Hosp Infect* 2020; 104: 453.
7. Rahim MA, Mostafi M. Repurposive Use of Drugs in COVID-19: A Wake-up Call. *J Bangladesh Coll Phys Surg* 2020; 38 (Suppl. Issue): 3-4.