LETTER TO THE EDITOR:

Esophageal ulcer bleeding due to CMV infection is quite infrequent in clinical practice. We reported two cases presenting with different patterns of esophageal ulcer, which was confirmed as CMV esophagitis. Our report would remind physicians the unusual occurrence of the CMV-related esophageal ulcer bleeding in the intensive care unit patients.

Cytomegalovirus (CMV) usually causes opportunistic infections in immunocompromised patients. Active CMV disease occurred in up to 36% of critically ill patients and may affect their health outcomes [1, 2]. We present upper gastrointestinal (UGI) bleeding with different endoscopic features of CMV esophagitis in two patients hospitalized in intensive care units (ICUs) - one immunocompetent patient who stayed in ICU for a critical illness and a second patient with colon cancer who received tegafur/uracil chemotherapy.

Case 1

A 77-year-old woman with diabetes mellitus and coronary artery disease had short of breath in recent days. Cough with sputum, mild chest tightness, and poor appetite were noted. She was admitted to the ICU and sputum cultures yielded carbapenem-resistant *Acinetobacter baumannii* (CRAB). Antibiotic therapy with doripenem plus sulbactam was given for 10 days. Meanwhile, unstable hemodynamic status and upper gastrointestinal bleeding occurred. Panendoscopic study showed diffuse ulcers of whole esophagus (Fig. 1A). The CMV polymerase chain reaction (PCR) results for blood and stool samples were positive. The CMV antigenemia showed 23 positive cells per 200,000 macrophages. Biopsy of esophageal ulcers revealed some large cells containing eosinophilic inclusion, which was positive in the immunohistochemical staining for CMV. Ganciclovir was added and esophageal bleeding stopped soon. One week later, nevertheless, the patient passed away in the hospital due to worsening CRAB pneumonia.

Case 2

A 75 y/o man of diabetes and colon cancer post hemicolecotmy had received tegafur / uracil chemotherapy. He suffered from nausea, vomiting and hematemesis for one week. The laboratory data showed leukocytosis and panendoscopy revealed pseudomembranous lesions of the esophagus (Fig. 1B). Esophageal biopsy showed CMV infection. The blood and stool CMV-PCR results were positive. A viral load of blood sample was 12,200 IU/mL. After ganciclovir

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therapy, the esophageal bleeding subsided and he was uneventfully discharged.

CMV diseases have been reported in immunocompromised patients especially with acquired immunodeficiency syndrome and solid organ transplant. Beside, in critically ill patients, CMV infection is associated with prolonged ventilator support, nosocomial infections, prolonged hospital stay and increased mortality [1]. Many clinicians neglect CMV colitis in the critically ill patients [3]. But there have been a few reports of serious complications, such as massive intestinal bleeding and intestinal perforation in the critically ill immunocompetent patients [3, 4]. Therefore, a high level of suspicion for CMV disease is required when diagnosing critically ill patients with gastrointestinal symptoms, and endoscopic biopsies should be taken from all gastrointestinal tract lesions to detect CMV in the cytomegalic inclusion bodies.

The CMV esophagitis is rarely highlighted in critically ill patients. The most common lesions were well-circumscribed ulcers, usually located in the middle to distal parts of the esophagus [5]. As we know, the pseudomembranous CMV lesions have been seen in the colitis but not reported in the esophagitis [3]. The intensivists should early diagnose the active CMV disease and to assess those patients who might preferably be given antiviral treatment.

**Conclusion**

We report two cases of esophageal bleeding lesions, which mucosal pathology confirmed CMV esophagitis. Our cases highlight different endoscopic features of CMV esophagitis among non-immunocompromised and immunocompromised patients in the ICUs. Our report would remind physicians the unusual occurrence of esophageal lesions due to CMV infections.
Conflict of interests

We declare no conflict of interest and financial support regarding this letter.

Ethics statement

The above study has been granted exemption from review by the Institutional Review Board of Chi-Mei Medical Center (IRB Serial No. 10410-005).

Financial Disclosure: None

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