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Exanthem and enanthem with fever and dyspnea



Figure 1. Erythematous, maculopapular, blanching rash on the trunk.



Figure 2. Koplik spots (black arrows), whitish, grayish, or bluish elevations of 1 to 3 mm with an erythematous base on the buccal mucosa opposite the molars.

**Koplik spots
are considered
pathognomonic
for measles**

A 49-YEAR-OLD WOMAN PRESENTED with a 5-day history of high fever, productive cough, mild dyspnea, malaise, generalized myalgias, and conjunctivitis. Hours before admission she developed a maculopapular rash. She had been diagnosed 2 years earlier with decompensated liver cirrhosis due to autoimmune hepatitis–primary biliary cholangitis overlap syndrome and was currently being treated with prednisolone, azathioprine, and ursodeoxycholic acid.

doi:10.3949/ccjm.88a.21047

WORKUP AND HOSPITAL COURSE

On examination, her temperature was 38.3°C (100.9°F), heart rate 92 beats per minute, blood pressure 100/68 mm Hg, respiratory rate 24 breaths per minute, and oxygen saturation 92% on ambient air. An erythematous, maculopapular, blanching rash was noted on the face and trunk (**Figure 1**), as well as enanthem on the buccal mucosa opposite the molars (**Figure 2**). Auscultation detected basilar fine crackles in both lungs. A firm liver was palpable 2 to 3 fingers below the right costal margin without tenderness. The rest of the

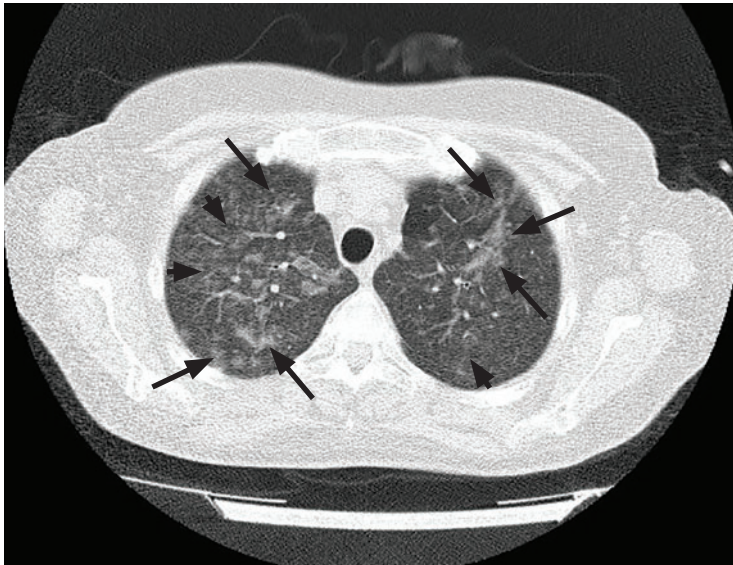


Figure 3. Computed tomography of the lungs revealed bilateral infiltrates (arrows) and ground-glass opacities (arrowheads) mainly in the upper lobes.

The presence of Koplik spots along with symptoms typical for viral infection makes measles the most likely diagnosis, even before the development of exanthem

physical examination was unremarkable.

Laboratory testing revealed leukopenia, thrombocytopenia, and increased serum levels of aspartate aminotransferase, lactate dehydrogenase, creatine kinase, gamma-glutamyl transferase, and total bilirubin.

The patient's dyspnea deteriorated significantly over the next 2 days, accompanied by high fever, with temperatures of 39.6°C to 40.5°C (103.3°F to 104.9°F). The initial findings on auscultation worsened and included diffuse bilateral crackles. Arterial blood gas analysis revealed hypoxemia (P_{aO_2} 59 mm Hg, P_{CO_2} 27 mm Hg), while she was breathing 3 L of oxygen per minute by nasal cannula.

Computed tomography of the lungs revealed bilateral infiltrates and ground-glass opacities in the upper lobes (**Figure 3**). Serologic testing by enzyme-linked immunosorbent assay confirmed the presence of positive immunoglobulin M antibodies against measles virus. She was treated with 40% oxygen through a face mask, intravenous levofloxacin, and oral ribavirin. As the enanthem (Koplik spots) prevented the patient from consuming adequate fluids and food, supportive care was provided including D/W 5% daily by intravenous infusion. The patient was discharged 1 week later in good condition, and at a follow-up visit 1 month after discharge, she was asymptomatic.

DISCUSSION

Measles is a highly contagious viral illness with a 90% transmission rate to susceptible individuals. The majority of deaths are from pneumonitis or encephalitis, while the most common complication is diarrhea. During the 2018 measles outbreak in Europe, tens of thousands of cases were reported, with two-thirds of patients requiring hospitalization.¹

The incubation period for measles is 8 to 12 days² and begins after virus entry via respiratory mucosa or conjunctivae. The virus replicates locally, spreads to regional lymphatic tissues, and is then thought to be disseminated to other reticuloendothelial sites through the blood stream ("primary viremia").³ The prodromal phase is defined by the appearance of symptoms that typically include fever, malaise, and anorexia, followed by conjunctivitis that may be accompanied by lacrimation or photophobia, coryza, and cough.^{2,3} Respiratory symptoms result from mucosal inflammation from viral infection of epithelial cells.

Prodromal symptoms typically intensify a few days before exanthem appears. Around 48 hours before the onset of exanthem, patients may develop enanthem characterized by Koplik spots, whitish, grayish, or bluish elevations of 1 to 3 mm with an erythematous base, typically on the buccal mucosa opposite the molars. Koplik spots are considered pathognomonic for measles.³ They may spread to cover the buccal and labial mucosa and the hard and soft palate.

Although a safe and effective measles vaccine was developed in 1963 and was eventually included in the trivalent measles-mumps-rubella vaccine in 1981 in the Greek National Vaccination Program, this patient had not been vaccinated against measles as a child or as an adult.

CLINICAL DIFFERENTIAL DIAGNOSIS

Once exanthem appears, the differential diagnosis for measles includes varicella, roseola (human herpesvirus 6 and 7), enteroviruses (coxsackievirus A9 and B5), erythema infectiosum (parvovirus B19), rubella, infectious mononucleosis, Epstein-Barr virus (especially during amoxicillin treatment), and group A streptococcal infection.

If dyspnea develops, the differential diagnosis may encompass Rocky Mountain spotted fever, meningococcemia, and *Mycoplasma pneumoniae*-induced rash and mucositis. In immunocompromised individuals like this patient, the differential diagnosis also includes *Candida albicans* mucositis with subsequent fungemia and pneumonia, and herpes simplex virus type 1 infection with viremia and end-organ disease.

The presence of Koplik spots along with the symptoms typical for viral infection makes measles the most likely diagnosis even before the development of exanthem.

SUPPORTIVE TREATMENT IS THE STANDARD

The mainstay of measles treatment is supportive. Treatment of bacterial superinfections such as bacterial pneumonia and otitis, as well as seizures and respiratory failure, may also be necessary. Administration of vitamin A to children with measles is associated with decreased morbidity and mortality.⁴

Early initiation of ribavirin treatment—within the first 5 days of disease onset—seems more effective than later in the disease course.⁵ Given the high risk of measles-associated mortality in immunosuppressed individuals⁶ and the absence of treatment guidelines, some

authors have recommended ribavirin treatment in patients with measles complications (mostly pneumonitis and encephalitis).⁵

In a prospective study of 100 patients with measles, half were assigned to treatment with ribavirin and supportive therapy, and the other 50 patients received only supportive therapy. Those receiving ribavirin had earlier resolution of fever and constitutional symptoms and fewer complications than those receiving only supportive care.⁷

The optimal duration of ribavirin therapy is not known, but 5 to 7 days may be reasonable except for severely immunosuppressed patients, who may require 2 to 3 weeks of treatment.⁸

TEACHING POINT

In patients with short-term high fevers and symptoms of viral infection (eg, malaise, generalized myalgias, conjunctivitis) but no maculopapular rash, a thorough inspection of the buccal cavity for Koplik spots will allow for a timely diagnosis of measles and for prompt administration of ribavirin if dyspnea or other complications develop.⁴

DISCLOSURES

The authors report no relevant financial relationships which, in the context of their contributions, could be perceived as a potential conflict of interest.

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