

Clinical presentation and diagnostic challenges of mycoplasma pneumoniae-Associated Acute Renal Failure.

Houwing Robyn*

Department of Paediatric Haematology, Erasmus University Medical Center, Wytemaweg Rotterdam, Netherlands

Introduction

Mycoplasma pneumoniae is a common respiratory pathogen that can occasionally lead to extrapulmonary complications, including acute renal failure. This article focuses on the clinical presentation and diagnostic challenges associated with Mycoplasma pneumoniae-associated acute renal failure. Understanding the diverse clinical manifestations and addressing the diagnostic complexities is essential for timely recognition and appropriate management of this condition[1].

Mycoplasma pneumoniae is a significant cause of community-acquired respiratory infections. While respiratory symptoms predominate, a range of extrapulmonary complications, such as acute renal failure, have been reported. Recognizing the clinical presentation and navigating the diagnostic challenges associated with Mycoplasma pneumoniae-associated acute renal failure is crucial for healthcare professionals involved in the care of these patients[2].

Clinical presentation

Mycoplasma pneumoniae-associated acute renal failure can present with a variety of clinical manifestations. The onset of renal failure may occur concurrently with or following respiratory symptoms. Common clinical features include decreased urine output, elevated serum creatinine levels, electrolyte imbalances, and signs of renal dysfunction. However, the presentation can vary widely, ranging from asymptomatic renal impairment to severe acute kidney injury requiring renal replacement therapy[3].

Diagnostic challenges

Diagnosing Mycoplasma pneumoniae-associated acute renal failure poses several challenges. The absence of specific clinical markers and the overlap of symptoms with other causes of acute renal failure make accurate diagnosis difficult. Laboratory investigations, including serological tests, may aid in establishing the association between Mycoplasma pneumoniae infection and renal dysfunction, but they often lack sensitivity and specificity. Additionally, distinguishing between primary renal injury caused by Mycoplasma pneumoniae and concurrent renal insults can be challenging[4].

In cases of acute renal failure associated with Mycoplasma pneumoniae infection, it is crucial to consider alternative etiologies. Other causes of acute kidney injury, such as sepsis, glomerulonephritis, drug-induced nephrotoxicity, and obstructive uropathy, should be thoroughly evaluated and excluded. A comprehensive clinical evaluation, including a detailed medical history, physical examination, and appropriate laboratory investigations, is necessary to differentiate Mycoplasma pneumoniae-associated acute renal failure from other potential causes[5].

Establishing a diagnosis of Mycoplasma pneumoniae-associated acute renal failure requires a multidisciplinary approach. Clinical suspicion based on the temporal relationship between respiratory symptoms and renal dysfunction is essential. Laboratory investigations, including serological testing for Mycoplasma pneumoniae-specific antibodies and polymerase chain reaction (PCR) assays, can provide supportive evidence. Serial monitoring of renal function and response to treatment may further strengthen the diagnosis.

Conclusion

Mycoplasma pneumoniae-associated acute renal failure is a rare but important complication of respiratory infections. Recognizing the diverse clinical presentation and addressing the diagnostic challenges is crucial for prompt diagnosis and appropriate management. Healthcare professionals should maintain a high index of suspicion in patients presenting with acute renal failure and a recent or concurrent respiratory infection. Collaboration between clinicians, nephrologists, and infectious disease specialists is essential to navigate the diagnostic complexities and provide optimal care for affected individuals.

References

1. Bajantri B, Venkatram S, Diaz-Fuentes G. Mycoplasma pneumoniae: a potentially severe infection. *J Clin Med Res.* 2018;10(7):535-44.
2. de Groot RC, Sauteur PM, Unger WW et al. Things that could be Mycoplasma pneumoniae. *J Infect.* 2017;74:S95-100.
3. Tong L, Huang S, Zheng C et al. Refractory Mycoplasma pneumoniae pneumonia in children: early recognition and management. *J Clin Med.* 2022 May 17;11(10):2824.

*Correspondence to: Houwing Robyn Department of Paediatric Haematology, Erasmus University Medical Center, Wytemaweg Rotterdam, Netherlands, Japan, E-mail: Brit@

Received: 30-May-2023, Manuscript No. AACNT-23-102789; Editor assigned: 02-June-2023, PreQC No. AACNT-23-102789 (PQ); Reviewed: 16-June-2023, QC No. AACNT-23-102789; Revised: 21-June-2023, Manuscript No. AACNT-23-102789(R); Published: 29-June-2023, DOI: 10.35841/aacnt-7.3.146

4. Wetter DA, Camilleri MJ. Clinical, etiologic, and histopathologic features of Stevens-Johnson syndrome during an 8-year period at Mayo Clinic. In Mayo Clinic Proceedings 2010 (85, 2, 131-138).
5. Kammer J, Ziesing S, Davila LA et al. Neurological manifestations of Mycoplasma pneumoniae infection in hospitalized children and their long-term follow-up. 2016;47(05):308-17.