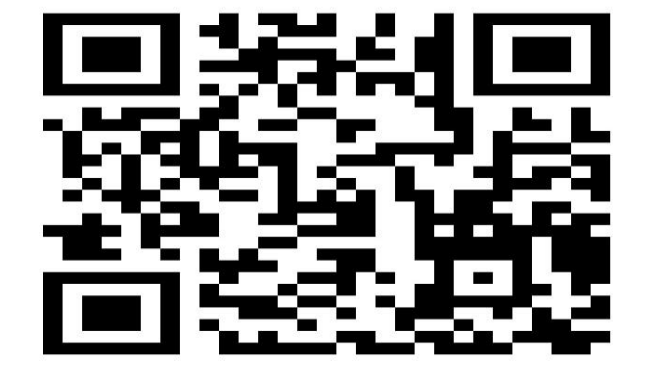


A Ternary Enigma: A Case Report and Literature Review of Myoclonus, Uremia, and Delirium in a Liver Transplant Recipient.

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Introduction

- Background:** Myoclonus is characterized by sudden, involuntary muscle jerks and can be associated with conditions such as uremia and delirium, especially in ICU settings. In this case report, we present a rare instance of myoclonus in a liver transplant recipient. This study aims to aid clinical recognition and management of this complex presentation.
- Objective:** To describe the diagnostic and therapeutic complexities of myoclonus associated with uremia in a liver transplant recipient, emphasizing the role of Consultation-Liaison (CL) psychiatry.

Case Presentation

Patient: 70-year-old female, history of alcohol use disorder, decompensated cirrhosis secondary to hepatitis C, underwent liver transplantation.

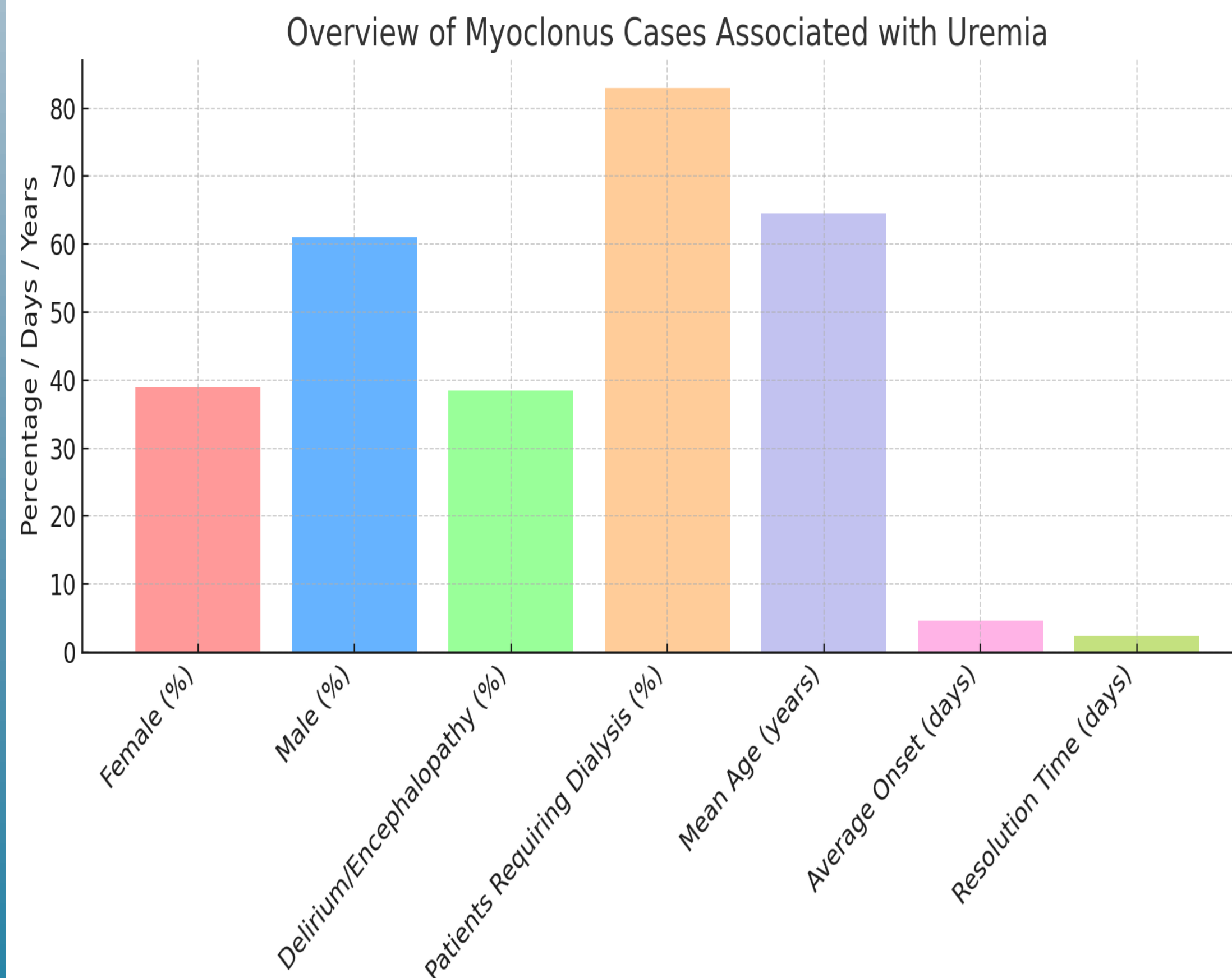
Clinical Course:

- Post-transplant Complications:** ICU admission due to respiratory failure, vancomycin-resistant Enterococcus bacteremia.
- Re-admission:** Blood drainage, hypotension, fever, unstable atrial fibrillation.
- Key Symptoms:** Cognitive impairment, sleep-wake disturbances, agitation, anxiety, myoclonic jerks.
- Diagnostics:** Neurological examination, EEG, laboratory testing indicating uremia.
- Intervention:** Continuous renal replacement therapy (CRRT), stop neurotoxic drugs, symptom management with midazolam.

Methodology

- Systematic Review:** A review of relevant literature was conducted using PubMed and EMBASSE.
- Criteria:** Articles from January 1990 to September 2023 focusing on uremia-associated myoclonus.
- Findings:** 12 manuscripts were included, covering 13 cases of uremia-induced myoclonus, **with no prior reports in solid organ transplant (SOT) recipients**. Most of the articles were case reports, with one case series included.

Systematic Review



Results

- The mean age of the patients was 64.5 years (standard deviation 6 10.92), and 39% were female.
- Myoclonus was diagnosed clinically in all cases, with an average onset of **4.6 days associated with uremia (standard deviation +/- 3.2 days)**.
- Notably, the most common accompanying clinical symptoms were delirium and/or encephalopathy, present in 38.5% of cases.
- Treatment with Hemodialysis resolved myoclonus in approximately 83% of reviewed cases.

Discussion

- Pathophysiology:** Uremia-related myoclonus might involve excitatory effects on NMDA receptors and inhibitory effects on GABA receptors. Metabolic disturbances such as electrolyte imbalances could contribute.
- Role of CL Psychiatry:** Accurate diagnosis and distinction from other movement disorders are crucial for effective treatment. Management requires a multidisciplinary approach.
- Implications:** Early identification and intervention, including dialysis, are critical in reducing morbidity.

Diagnostic & Management

- History/Differential:** Thorough patient history and physical examination. Consider toxic-metabolic, medications, seizures, infections.
- Underlying Cause:** Address primary condition (e.g., dialysis for uremia). Test including Imaging, EEG, electromyography to determine etiology.
- Medication Adjustment:** Modify or discontinue offending medications.
- Symptomatic Treatment:** Benzodiazepines like midazolam for symptom control.
- Multidisciplinary Approach:** Collaborate with neurology and other specialists.

Conclusion

- Key Insights:** This case illustrates the critical role of CL psychiatrists in identifying and managing complex presentations of myoclonus in transplant recipients. A multimodal treatment approach, including CRRT and medication management, is effective.
- Future Directions:** Further research into the neurophysiological mechanisms of uremia-related myoclonus in transplant settings is needed to improve patient outcomes.

References

- Okwuonu, E., & Sher, Y. (2024). Myoclonus, Uremia, and Delirium in a Liver Transplant Recipient: A Case Report and Literature Review. *Journal of the Academy of Consultation-Liaison Psychiatry*. <https://doi.org/10.1016/j.jaclp.2024.07.004>.