According to Rebman and colleagues, writing in the journal *Frontiers in Medicine*, PTLDS is a severe complication of Lyme disease. [1] The authors identified PTLDS using the Infectious Diseases Society of America (IDSA) proposed case definition. [2] “Briefly, this definition relies on prior physician-documented Lyme disease, treatment with standard of care antibiotics, and the development of significant fatigue, widespread musculoskeletal pain, and/or cognitive difficulties that last for a period of at least 6 months, and began within 6 months of a Lyme diagnosis and recommended treatment,” explains Rebman.

Individuals with PTLDS represent a group of patients who have been evaluated for “unexplained fatigue, pain, and neurocognitive symptoms by primary care and sub-specialty physicians,” the authors state. These patients are “highly and clinically significantly symptomatic, with poor health-related quality of life.”

“PTLDS patients,” Rebman states, “exhibited levels of fatigue, musculoskeletal pain, sleep disturbance, and depression which were both clinically relevant and statistically significantly higher than controls.”

Post-treatment Lyme disease syndrome can last for years, study finds.

According to the study findings, PTLDS can last for years. “Our cohort was a median of 3.6 years from onset of PTLDS symptoms to study enrollment, with a range of 8.3 months to 27.7 years,” states Rebman.

And, PTLDS can occur even in Lyme disease patients treated within 30 days of the onset of symptoms. “Time from illness onset to first recommended course of antibiotic treatment was a median of 30 days,” according to Rebman.

Additionally, 59% of the patients with PTLDS reported having a delay in diagnosis or an initial misdiagnosis. Risk factors for PTLDS include a delay in diagnosis and an increased severity of initial illness, including the presence of neurologic symptoms.

The authors did not address other manifestations including chronic neurologic Lyme disease [3] and neuropsychiatric Lyme disease. [4]

The Johns Hopkins study should put to rest the false narrative perpetuated by the 2006 IDSA treatment guidelines which states, “In many patients, post-treatment symptoms appear to be more related to the aches and pains of daily living rather than to either Lyme disease or a tick-borne co-infection.” [2]
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Johns Hopkins’ study supports early identification of Lyme disease patients for re-treatment

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References:
