Chronic Fatigue Syndrome and Fibromyalgia; Now Treatable Diseases

Chronic Fatigue Syndrome (CFS) and Fibromyalgia (FM) are illnesses that often coexist and affect millions of Americans. Symptoms vary amongst individuals and commonly include severe fatigue, sleep disturbances, cognitive problems, commonly called brain fog, muscle pain and multiple infections. Unfortunately, many individuals and physicians continue to deny that these syndromes are legitimate diseases. The medical literature is, however, very clear that these are legitimate diseases and individuals with these syndromes have measurable hypothalamic, pituitary, immune and coagulation dysfunction. These abnormalities then result in a cascade of further abnormalities, in which stress plays a role by suppressing immunity and hypothalamic-pituitary function. The pituitary and hypothalamic dysfunction results in multiple hormonal deficiencies that are often not detected with standard blood tests, and autonomic dysfunction, including neurally mediated hypotension. The immune dysfunction, which includes natural killer cell dysfunction, results in opportunistic infections and yeast overgrowth, resulting in worsening of symptoms. Recent studies have shown that the coagulation dysfunction is usually initiated by a viral infection and has a genetic predisposition. This abnormal coagulation results in increased blood viscosity (slugging) and a deposition of soluble fibrin monomers along the capillary wall. This results in tissue and cellular hypoxia, resulting in fatigue, and decreased cognition (brain fog). Neurotransmitter abnormalities and macro and micro nutrient deficiencies have also been shown to occur with these disorders.

Reactivating viruses combined with suppressed immunity is a common thread amongst CFS and FM patients. Current research suggests that many triggers can initiate a cascade of events, causing the hypothalamic-pituitary-immune and coagulation dysfunction. The most common initiating cause is a viral infection, which is very commonly Epstein Bar Virus, Cytomegalovirus or HHV6. When specialized testing is utilized, these infections are found in 30-80% of CFS and FM patients, and many people with these syndromes can pinpoint the start of their disease to a viral infection that never got better usually during significant life stressors.

Although a concept that is sometimes uncomfortable and foreign to traditional medical styles of thinking, the need for multiple interventions is effective when an illness affects a critical control center.

Effective treatment, with 80 to 90 percent of individuals achieving significant clinical benefits, can be achieved by simultaneously treating the above problems that an individual is found to have. The mix of treatments needed varies from patient to patient based on what abnormalities are present and how progressed the condition, but there but there is consistent abnormal pathophysiology. For instance, a high percentage of individuals with these syndromes have low thyroid. This is, however, usually not picked up on the standard blood tests because the TSH is not elevated in these individuals because of pituitary dysfunction. Many of these individuals will also have high levels of the anti-thyroid reverse T3, which is usually not measured on standard blood tests. In addition, the majority of individuals can also have a thyroid receptor resistance that is not detected on the blood tests. Consequently, thyroid treatment, especially with timed release T3 is effective for many patients. T4 preparations (inactive thyroid) such as Synthroid and Levoxyl do not work well for these conditions. Adrenal insufficiency and growth hormone deficiency are also very common with these disorders, and supplementation with these hormones can often have profound effects. As with thyroid testing, these deficiencies are, unfortunately, usually not detected with the standard screen blood tests and require more specific testing.

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