Lyme Borreliosis: Late Stage Neuropsychiatric and Other Clinical Findings and Clinical Assessment



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Disclosure Statement Robert Bransfield, MD, DLFAPA, PC

- Patients pay me money in return for trying to help them.
- Most of my income is paid directly from patients
- No Lyme, psychoimmunology or infectious disease financial interests.
- No research grants or patents dependent upon a disease definition.
- I have been an expert witness in cases involving Lyme disease.

Differentiating Somatopsychic, Psychosomatic, Multisystem Illness and Medical Uncertainty

Bransfield RC, Friedman KJ. Healthcare (Basel). 2019 Oct 8;7(4). pii: E114. doi: 10.3390/healthcare7040114

Term	DSM-5	ICD-10	ICD-11
	Diagnosis	Diagnosis	Diagnosis
All in your head	No	No	No
Somatic symptom disorder	Yes	Yes	No
Somatoform disorder	No	No	No
Medically unexplained symptoms	No	No	No
Functional neurological symptom disorder	Yes	Yes	No
Conversion disorder	No	Yes	No
Illness anxiety disorder	Yes	No	Yes
Factitious disorder imposed upon another	Yes	Yes	Yes
(Munchausen's by proxy)			
Functional disorders	No	Yes	No
Psychogenic disorders	No	Yes	No
Compensation neurosis	No	No	No
Psychogenic seizures	No	Yes	Yes
Psychogenic pain	No	Yes	No
Psychogenic fatigue	No	No	No
Delusional parasitosis	No	Yes	Yes
Subjective vs. Objective	No	No	No
Non-specific or vague symptoms	No	No	No
Bodily distress disorder	No	No	Yes

Conclusion

• We must recognize there is always some degree of medical uncertainty with any condition. Not everything is well understood or categorized. Complex diseases require complex explanations, and there needs to be recognition of varying degrees of medical uncertainty. Everything is caused by something. Nothing is caused by nothing. When clinical findings are puzzling, the ethical approach is to continue attempting to search for its cause.

Conclusion II

 Medically unexplained symptoms, somatoform disorder, and compensation neurosis are outdated. Terms such as subjective, vague, and non-specific are used inaccurately. *Conversion disorders, functional disorders, psychogenic* illness, Munchausen's by proxy, somatic symptom disorder, psychogenic seizures, psychogenic pain, psychogenic fatigue, and delusional parasitosis are over-diagnosed. *Bodily distress disorder* is *Bodily distress* syndrome is dependent upon "medically unexplained symptoms" and is scientifically unsupported.

Conclusion III

 To properly understand the mind/body connection, a knowledge of general medicine, psychiatry, and the systems which link the soma and the brain are required. No one has a complete knowledge of all fields of medicine. We must have compassion and humility and recognize that not all diseases have been discovered or properly understood, and be aware that much remains to be learned about the brain/body interaction.

Lyme Borreliosis: Late Stage Neuropsychiatric and Other Clinical Findings and Clinical Assessment

 Many late-stage chronic Lyme disease clinical findings are neuropsychiatric. A total clinical assessment is critical in diagnosis, especially since controversy surrounds the reliability of immune-based testing. The clinical findings of one hundred Lyme disease patients with chronic neuropsychiatric symptoms were entered into a database. The prevalence of each clinical finding pre-infection and post-infection were compared and calculated within the 95% confidence interval. Patients had minimal symptoms pre-infection, but a high postinfection prevalence of a broad spectrum of acquired multisystem symptoms.

Clinical Findings Pre-infection, Post-infection, 95% Cl

Sustained attention	7%	(2% - 12%)	84%	(77% - 91%)
Allocation of attention	6%	(1% - 11%)	66%	(57% - 75%)
Distracted by frustration	7%	(2% - 12%)	79%	(71% - 87%)
Hypersensitivity to sound	3%	(0% - 6%)	66%	(57% - 75%)
Hypersensitivity to light	2%	(0% - 5%)	63%	(54% - 72%)
Hypersensitivity to touch	2%	(0% - 5%)	41%	(31% - 51%)
Hypersensitivity to smell	5%	(1% - 9%)	36%	(27% - 45%)
Sensory overload	No data			

N	lemory	Pre-infection		Post-infection	
	Working memory	3%	(0% - 6%)	78%	(70% - 86%)
	Working spatial memory	1%	(0% - 3%)	46%	(36% - 56%)
	Recent memory	5%	(1% - 9%)	77%	(69% - 85%)
	Remote memory	4%	(0% - 8%)	35%	(26% - 44%)
Memory retrieval					
	Words	3%	(0% - 6%)	70%	(61% - 79%)
	Numbers	3%	(0% - 6%)	52%	(42% - 62%)
	Names	6%	(1% - 11%)	68%	(59% - 77%)
	Faces	1%	(0% - 3%)	23%	(15% - 31%)
	Motor memory	1%	(0% - 3%)	10%	(4% - 16%)
	Geographical/spatial	1%	(0% - 3%)	49%	(39% - 59%)

Ρ	rocessing	Pre-infection		Post-infection	
	Letter reversals	2%	(0% - 5%)	45%	(35% - 55%)
	Spelling errors	8%	(3% - 13%)	56%	(46% - 66%)
	Word substitution errors	5%	(1% - 9%)	55%	(45% - 65%)
	Number reversals	1%	(0% - 3%)	39%	(29% - 49%)
	Reading comprehension	6%	(1% - 11%)	59%	(49% - 69%)
A	uditory comprehension	5%	(1% - 9%)	49%	(39% - 59%)
Sound localization		3%	(0% - 6%)	19%	(11% - 27%)
Spatial perceptual distortions		1%	(0% - 3%)	21%	(13% - 29%)
0	ptic ataxia	1%	(0% - 3%)	51%	(41% - 61%)
Tr	ansposition of laterality	2%	(0% - 5%)	22%	(14% - 30%)
Le	eft-Right confusion	6%	(1% - 11%)	30%	(21% - 39%)
Calculation		10%	(4% - 16%)	51%	(41% - 61%)
Fluency of speech		4%	(0% - 8%)	62%	(52% - 72%)
Fl	uency of written language	2%	(0% - 5%)	43%	(33% - 53%)
Η	andwriting	8%	(3% - 13%)	47%	(37% - 57%)

E>	cecutive functioning	Pre-infection		Post- infection	
	Unfocused concentration	4%	(0% - 8%)	81%	(73% - 89%)
	Brain fog	3%	(0% - 6%)	84%	(77% - 91%)
	Prioritizing multiple tasks	6%	(1% - 11%)	76%	(68% - 84%)
	Multitasking	3%	(0% - 6%)	74%	(65% - 83%)
	Racing thoughts	1%	(0% - 3%)	54%	(44% - 64%)
	Obsessive thoughts	4%	(0% - 8%)	56%	(46% - 66%)
	Mental apathy	4%	(0% - 8%)	72%	(63% - 81%)
	Abstract reasoning	3%	(0% - 6%)	51%	(41% - 61%)
Er	Intrusive thoughts notional	no data			
	Decreased frustration tolerance	5%	(1% - 9%)	80%	(72% - 88%)
	Hyperarousal	no data			
	Sudden mood swings	3%	(0% - 6%)	74%	(65% - 83%)
	Hypervigilance	1%	(0% - 3%)	45%	(35% - 55%)
	Paranoia	1%	(0% - 3%)	26%	(17% - 35%)
	Crying spells	0%	(0% - 0%)	50%	(40% - 60%)
	Anhedonia	3%	(0% - 6%)	64%	(55% - 73%)
	Intrusive emotions	No data			

In	nagery				
	Capacity for visual imagery	2%	(0% - 5%)	19%	(11% - 27%)
	Intrusive aggressive images	1%	(0% - 3%)	19%	(11% - 27%)
	Intrusive sexual images	1%	(0% - 3%)	6%	(1% - 11%)
	Intrusive images, other	1%	(0% - 3%)	10%	(4% - 16%)
	Hypnagogic hallucinations	2%	(0% - 5%)	21%	(13% - 29%)
	Vivid nightmares	3%	(0% - 6%)	38%	(28% - 48%)
	Illusions	2%	(0% - 5%)	20%	(12% - 28%)
	Hallucinations (auditory, visual, olfactory, tactile)	2%	(0% - 5%)	18%	(10% - 26%)
E	notional				
	Decreased frustration tolerance	5%	(1% - 9%)	80%	(72% - 88%)
	Hyperarousal	no data			
	Sudden mood swings	3%	(0% - 6%)	74%	(65% - 83%)
	Hypervigilance	1%	(0% - 3%)	45%	(35% - 55%)
	Paranoia	1%	(0% - 3%)	26%	(17% - 35%)
	Crying spells	0%	(0% - 0%)	50%	(40% - 60%)
	Anhedonia	3%	(0% - 6%)	64%	(55% - 73%)
	Intrusive emotions	No data			
D	issociative symptoms				
	Depersonalization	2%	(0% - 5%)	64%	(55% - 73%)
	Derealization	1%	(0% - 3%)	29%	(20% - 38%)
	Dissociative episodes	0%	(0% - 0%)	12%	(6% - 18%)

B	ehavioral				
	Disinhibition	2%	(0% - 5%)	33%	(24% - 42%)
	Exaggerated startle reflex	1%	(0% - 3%)	49%	(39% - 59%)
	Explosive anger	3%	(0% - 6%)	39%	(29% - 49%)
	Suicidal	1%	(0% - 3%)	28%	(19% - 37%)
	Homicidal	0%	(0% - 0%)	1%	(0% - 3%)
	Accident prone	4%	(0% - 8%)	35%	(26% - 44%)
	Decreased social functioning	6%	(1% - 11%)	72%	(63% - 81%)
	Decreased job/school performance	2%	(0% - 5%)	78%	(70% - 86%)
	Marital/family problems	4%	(0% - 8%)	39%	(29% - 49%)
	Substance abuse	1%	(0% - 3%)	12%	(6% - 18%)
	Legal difficulties	1%	(0% - 3%)	8%	(3% - 13%)
	Compensatory compulsions	2%	(0% - 5%)	58%	(48% - 68%)
	Dropping objects	2%	(0% - 5%)	52%	(42% - 62%)

Psychiatric syndromes					
	Depression	9%	(3% - 15%)	79%	(71% - 87%)
	Rapid cycling bipolar	3%	(0% - 6%)	11%	(5% - 17%)
	Panic disorder	2%	(0% - 5%)	49%	(39% - 59%)
	Obsessive compulsive disorder	2%	(0% - 5%)	24%	(16% - 32%)
	Social anxiety disorder	7%	(2% - 12%)	36%	(27% - 45%)
	Generalized anxiety disorder	3%	(0% - 6%)	53%	(43% - 63%)
	Posttraumatic stress disorder	6%	(1% - 11%)	16%	(9% - 23%)

V	egetative				
E	nergy				
	Fatigue	1%	(0% - 3%)	76%	(68% - 84%)
S	eep				
	Non-restorative sleep	4%	(0% - 8%)	76%	(68% - 84%)
In	somnia				
	Insomnia, initial	5%	(1% - 9%)	70%	(61% - 79%)
	Insomnia, mid	1%	(0% - 3%)	72%	(63% - 81%)
	Insomnia, late	1%	(0% - 3%)	58%	(48% - 68%)
	Hypersomnia	2%	(0% - 5%)	73%	(64% - 82%)
	Loss of circadian rhythm	5%	(1% - 9%)	44%	(34% - 54%)
	Delayed sleep phase disorder	no data			
	Sleep apnea, central	no data			
	Sleep apnea, obstructive	no data			
	Sleep paralysis	no data			
	Cataplexy	no data			
	Narcolepsy	no data			

Ea	ating				
	Anorexia	1%	(0% - 3%)	45%	(35% - 55%)
	Weight loss	1%	(0% - 3%)	45%	(35% - 55%)
	Non-appetite over-eating	2%	(0% - 5%)	34%	(25% - 43%)
	Weight gain without increased food intake	1%	(0% - 3%)	27%	(18% - 36%)
	Weight gain with increased food intake	2%	(0% - 5%)	22%	(14% - 30%)
Sexual functioning					
	Decreased libido	4%	(0% - 8%)	60%	(50% - 70%)
	Increased libido	1%	(0% - 3%)	9%	(3% - 15%)
	Decreased arousal	1%	(0% - 3%)	42%	(32% - 52%)
	Decreased orgasm	2%	(0% - 5%)	41%	(31% - 51%)
	Altered sexual imagery	0%	(0% - 0%)	3%	(0% - 6%)

Temperature control					
	Body temperature fluctuations	3%	(0% - 6%)	63%	(54% - 72%)
	Flushing	3%	(0% - 6%)	49%	(39% - 59%)
	Intolerance to heat	2%	(0% - 5%)	58%	(48% - 68%)
	Intolerance to cold	2%	(0% - 5%)	64%	(55% - 73%)
	Decreased body temperature	5%	(1% - 9%)	52%	(42% - 62%)
	Low grade fevers	1%	(0% - 3%)	47%	(37% - 57%)
	Night sweats	2%	(0% - 5%)	60%	(50% - 70%)
	Chills	2%	(0% - 5%)	59%	(49% - 69%)

N	eurological				
	Headache	3%	(0% - 6%)	68%	(59% - 77%)
	Cervical radiculopathy	0%	(0% - 0%)	43%	(33% - 53%)
	Migraine	4%	(0% - 8%)	33%	(24% - 42%)
	Coital cephalgia	0%	(0% - 0%)	4%	(0% - 8%)
	TMJ	2%	(0% - 5%)	41%	(31% - 51%)
	Tension	2%	(0% - 5%)	57%	(47% - 67%)
	Cluster	0%	(0% - 0%)	10%	(4% - 16%)
	Sinus	5%	(1% - 9%)	41%	(31% - 51%)
	Thunderclap	no data			

C	ranial nerves				
	I Olfactory: loss of smell, altered taste	2%	(0% - 5%)	22%	(14% - 30%)
	II Ophthalmologic				
	Blurred vision	2%	(0% - 5%)	50%	(40% - 60%)
	Photophobia to bright light	3%	(0% - 6%)	61%	(51% - 71%)
	Sensitivity to fluorescence and flicker	3%	(0% - 6%)	48%	(38% - 58%)
	Floaters	1%	(0% - 3%)	56%	(46% - 66%)
	Flashes	0%	(0% - 0%)	23%	(15% - 31%)
	Conjunctivitis	0%	(0% - 0%)	19%	(11% - 27%)
	Eye pain	2%	(0% - 5%)	36%	(27% - 45%)
	Dry eyes	0%	(0% - 0%)	32%	(23% - 41%)
	Blind spots	1%	(0% - 3%)	12%	(6% - 18%)
	Night blindness	4%	(0% - 8%)	36%	(27% - 45%)
	Peripheral shadows	2%	(0% - 5%)	18%	(10% - 26%)
	Papilledema	0%	(0% - 0%)	1%	(0% - 3%)
	Iritis	0%	(0% - 0%)	1%	(0% - 3%)
	Uveitis	0%	(0% - 0%)	1%	(0% - 3%)
	Optic neuritis	0%	(0% - 0%)	2%	(0% - 5%)
	Panopsia	no data			

III, IV, VI Double vision or eye drifts when tired, ptosis	2%	(0% - 5%)	36%	(27% - 45%)
V Sensory loss, pain	0%	(0% - 0%)	27%	(18% - 36%)
VII Bell's palsy	2%	(0% - 5%)	16%	(9% - 23%)
VIII Tinnitus	1%	(0% - 3%)	51%	(41% - 61%)
Hearing loss	1%	(0% - 3%)	26%	(17% - 35%)
Dizziness	2%	(0% - 5%)	53%	(43% - 63%)
Vertigo	1%	(0% - 3%)	29%	(20% - 38%)
Motion sickness	9%	(3% - 15%)	40%	(30% - 50%)
Mal de debarquement	no data			
Tullio's	0%	(0% - 0%)	12%	(6% - 18%)
IX, X Episodic loss of speech, choking on food, difficulty swallowing	0%	(0% - 0%)	36%	(27% - 45%)
XI. Sternocleidomastoid and trapezius pain and paresis	0%	(0% - 0%)	44%	(34% - 54%)
XII. Tongue deviates to side	0%	(0% - 0%)	5%	(1% - 9%)

Se	eizures				
	Grand mal	1%	(0% - 3%)	4%	(0% - 8%)
	Partial	2%	(0% - 5%)	8%	(3% - 13%)
0	ther Neurological				
	Numbness	1%	(0% - 3%)	59%	(49% - 69%)
	Tingling	1%	(0% - 3%)	71%	(62% - 80%)
	Sensory loss	1%	(0% - 3%)	40%	(30% - 50%)
	Burning	1%	(0% - 3%)	36%	(27% - 45%)
	Static electric sensation	0%	(0% - 0%)	35%	(26% - 44%)
	Formication, crawling sensation	0%	(0% - 0%)	35%	(26% - 44%)
	Stabbing sensation	0%	(0% - 0%)	28%	(19% - 37%)
	Sensation of wetness	no data			
	Sensation of vibration	no data			
	Paresis	2%	(0% - 5%)	66%	(57% - 75%)
	Tremor	3%	(0% - 6%)	40%	(30% - 50%)
	Twitching	1%	(0% - 3%)	56%	(46% - 66%)
	Muscle tightness	0%	(0% - 0%)	56%	(46% - 66%)
	Restless leg	5%	(1% - 9%)	50%	(40% - 60%)
	Myoclonic jerks	1%	(0% - 3%)	38%	(28% - 48%)
	Torticollis	0%	(0% - 0%)	1%	(0% - 3%)
	Tourette's	0%	(0% - 0%)	2%	(0% - 5%)
	Extrapyramidal symptoms	0%	(0% - 0%)	3%	(0% - 6%)
	Ataxia	1%	(0% - 3%)	6%	(1% - 11%)
	Herniated disc(s)	4%	(0% - 8%)	14%	(7% - 21%)
	Spasticity	1%	(0% - 3%)	1%	(0% - 3%)
	Other neurological	1%	(0% - 3%)	6%	(1% - 11%)
	Romberg	1%	(0% - 3%)	21%	(13% - 29%)

V	lusculoskeletal				
	Joint pain, swelling, tightness, crepitation (specify joints)	2%	(0% - 5%)	81%	(73% - 89%)
	Bone thinning/Fractures	1%	(0% - 3%)	7%	(2% - 12%)
	Periostitis (tibia, ribs, iliac crest, sternum, clavicle, etc.	4%	(0% - 8%)	7%	(2% - 12%)
	Epicondylitis	2%	(0% - 5%)	20%	(12% - 28%)
	Deep bone pain	no data			
	Foot pain	no data			
	Plantar fasciitis	0%	(0% - 0%)	33%	(24% - 42%)
	Fibromyalgia	1%	(0% - 3%)	36%	(27% - 45%)
	Myalgia	1%	(0% - 3%)	54%	(44% - 64%)
	Chondritis (ear, nose, costochondral)	0%	(0% - 0%)	38%	(28% - 48%)
	Tendonitis	3%	(0% - 6%)	17%	(10% - 24%)
	Carpal tunnel	1%	(0% - 3%)	15%	(8% - 22%)

Ca	ardiovascular				
	Chest pain	2%	(0% - 5%)	39%	(29% - 49%)
	Heart block	2%	(0% - 5%)	11%	(5% - 17%)
	Mitral valve prolapse	4%	(0% - 8%)	20%	(12% - 28%)
	Racing pulse	0%	(0% - 0%)	48%	(38% - 58%)
	Episodes rapid and slow heart rate	0%	(0% - 0%)	34%	(25% - 43%)
	Postural orthostatic tachycardia syndrome	no data			
	Postural orthostatic hypotension	0%	(0% - 0%)	12%	(6% - 18%)
	Pericarditis	0%	(0% - 0%)	1%	(0% - 3%)
	Cardiomyopathy	0%	(0% - 0%)	2%	(0% - 5%)
	Murmur	7%	(2% - 12%)	16%	(9% - 23%)
	Hypertension	2%	(0% - 5%)	15%	(8% - 22%)
	Hypertensive crisis	1%	(0% - 3%)	3%	(0% - 6%)

Jpper respiratory and pulmonary				
Swollen glands	0%	(0% - 0%)	41%	(31% - 51%)
Cough	1%	(0% - 3%)	28%	(19% - 37%)
Allergies	7%	(2% - 12%)	35%	(26% - 44%)
Nose bleeds	1%	(0% - 3%)	7%	(2% - 12%)
Tooth pain	0%	(0% - 0%)	32%	(23% - 41%)
Periodontal disease	0%	(0% - 0%)	19%	(11% - 27%)
Shortness of breath	1%	(0% - 3%)	43%	(33% - 53%)
Asthma	4%	(0% - 8%)	14%	(7% - 21%)
Air hunger	no data			

Gastrointestinal				
Upper gastrointestinal distress	6%	(1% - 11%)	25%	(17% - 33%)
Irritable bowel	6%	(1% - 11%)	50%	(40% - 60%)
Abdominal bloating	1%	(0% - 3%)	42%	(32% - 52%)
Gastroparesis	0%	(0% - 0%)	1%	(0% - 3%)
Hepatitis	0%	(0% - 0%)	1%	(0% - 3%)
Pancreatitis	0%	(0% - 0%)	1%	(0% - 3%)
Inflammatory bowel	0%	(0% - 0%)	2%	(0% - 5%)
Cholecystitis	0%	(0% - 0%)	2%	(0% - 5%)
Non-calculous cholecystitis	no data			
Gall stones	0%	(0% - 0%)	1%	(0% - 3%)

G	enitourinary				
	Genital pain	1%	(0% - 3%)	27%	(18% - 36%)
	Breast tenderness, pain	1%	(0% - 3%)	24%	(16% - 32%)
	Lactation	0%	(0% - 0%)	8%	(3% - 13%)
	Menstrual irregularity	3%	(0% - 6%)	30%	(21% - 39%)
	Spastic bladder	1%	(0%-3%)	47%	(37%-57%)
	Urinary incontinence	1%	(0% - 3%)	18%	(10% - 26%)
	Interstitial cystitis	0%	(0% - 0%)	1%	(0% - 3%)
	Recurrent urinary tract infections	1%	(0% - 3%)	11%	(5% - 17%)
	Anesthesia of genitalia	0%	(0% - 0%)	6%	(1% - 11%)
	Atrophy of genitalia	0%	(0% - 0%)	3%	(0% - 6%)

0	ther				
	Chronic pain	0%	(0% - 0%)	41%	(31% - 51%)
	Alcohol intolerance	3%	(0% - 6%)	41%	(31% - 51%)
	Multiple chemical sensitivity	2%	(0% - 5%)	25%	(17% - 33%)
	Mold sensitivity	no data			
	Hair loss	2%	(0% - 5%)	47%	(37% - 57%)
	Thyroid dysfunction	1%	(0% - 3%)	20%	(12% - 28%)
	Wilson syndrome	0%	(0% - 0%)	4%	(0% - 8%)
	Adrenal insufficiency	0%	(0% - 0%)	10%	(4% - 16%)
	Hypoglycemia	2%	(0% - 5%)	20%	(12% - 28%)
	Vasculitis	0%	(0% - 0%)	5%	(1% - 9%)
	Ankle edema	1%	(0% - 3%)	20%	(12% - 28%)
	Splenomegaly	0%	(0% - 0%)	4%	(0% - 8%)
	Ecchymosis	1%	(0% - 3%)	34%	(25% - 43%)
	Acrodermatitis chronicum atrophicans	0%	(0% - 0%)	1%	(0% - 3%)
	Lymphocytoma	3%	(0% - 6%)	3%	(0% - 6%)
	Erythema of palms and soles	0%	(0% - 0%)	0%	(0% - 0%)
	Bartonella tracks	no data			

Symptom patterns				
Herxheimer reaction	0%	(0% - 0%)	73%	(64% - 82%)
Progression of symptoms	0%	(0% - 0%)	86%	(79% - 93%)
Fluctuation of symptoms	0%	(0% - 0%)	82%	(74% - 90%)
Stress increased symptoms	0%	(0% - 0%)	77%	(69% - 85%)
28-day or longer symptom cycle	0%	(0% - 0%)	43%	(33% - 53%)
Antibiotic reduce symptoms	0%	(0% - 0%)	72%	(63% - 81%)

Conclusion

• The results of this study were then used to develop three clinical assessment forms (24 item and full assessment) that can be used when the diagnosis of Lyme disease is suspected. Since controversy surrounds the commonly used immune-based testing for Lyme disease, this use of clinical diagnostic scales can be of value in confirming a diagnosis of Lyme borreliosis. The assessment systems can also be used to track further disease progression, improvement or response to treatment. The use of this assessment, and/or refinement by others can help to further validate these clinical assessment systems for the clinical diagnosis of Lyme borreliosis.

Concentration impairments
Short-term memory problems
Word finding difficulty
Name recall difficulty
Fluency of speech difficulties
Brain fog
Sudden mood swings
Decreased social functioning
Decreased job/school performance
Depression
Fatigue
Insomnia
Night sweats
Low body temperature
Headache
Blurred vision
Floaters
Tinnitus (ringing in the ears)
Sensitive to sound
Dizziness
Numbness
Tingling
Joint pain, swelling
Fluctuation of symptoms
Stress increases symptoms

Proposed Lyme Disease Guidelines and Psychiatric Illnesses. Bransfield RC. Friedman KJ. Healthcare (Basel) 2019. 9;7(3).



Systematic review of Lyme Disease causing psychiatric illness I

- The PubMed electronic search for citations:
- Lyme Disease psychiatric illness: 1054
- Lyme Disease causing psychiatric illness: 384
- Lyme Disease causing mental illness: 413
- Lyme Disease causing developmental disorders children: 134
- Lyme Disease causing behavioral disorders children: 268
- Lyme Disease psychiatric disorders children: 267
- ILADS Lyme/TBD causing psychiatric illness: 377 citations (304 psychiatry, 73 Dementia)
- My archives: 389



Citations in Perspective

- Lyme causing psychiatric symptoms: 387
- Lyme causing dementia: 73
- IDSA: 4 (only 2 failed to show an association)
- Citation total: 467 vs. 2, (0.4% of literature)
- Therefore the guidelines are evidence biased, not evidence based due to the selective reporting of outcomes

Conclusion

- The disclaimer and the manner these guidelines are implemented are insufficient to remove the authors and sponsoring organizations from liability for harm caused by these guidelines.
- The guidelines and supporting citations place improper credibility upon surveillance definition rather than clinical diagnosis criteria.
- The guidelines fail to address the clear causal association between Lyme disease and psychiatric illnesses, suicide, violence, developmental disabilities and substance abuse despite significant supporting evidence.

Conclusion II

- If these guidelines are published without very major revisions, and if the sponsoring medical societies attempt to enforce these guidelines as a standard of care, it will directly contribute to increasing a national and global epidemic of psychiatric illnesses, suicide, violence, substance abuse and developmental disabilities and the associated economic and noneconomic societal burdens.
- The guideline flaws could be improved with a more appropriate disclaimer, an evidence-based rather than an evidence-biased approach, more accurate diagnostic criteria, and recognition of the direct and serious causal association between Lyme Disease and psychiatric illnesses.

Thanks for your attention and your commitment to our patients

