


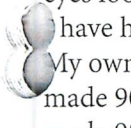
Lymes Disease and Blindness

by Dr. Gregory Bach, D.O., F.A.A.I.M., P.C.

 The diagnosis of Lyme Disease and its coinfections is so easy for the common Patient, but so elusive to the Medical Doctors. The Ideology for the last 37 years is that Lyme Disease is cured in two weeks. There is a Doctor who is asking for a 50 million dollar grant from his friends at CDC/NIH to prove that Lyme Disease is cured in two weeks and after that it's all in your head. It pays well to be part of the so called Holders of the Established Lyme Test and Big Pharma: See Published in Medical Rounds authored by Heather Dewar. "Chronic Lyme in the Cross Hairs" is a recent publication which is free for all PA medical society physicians. This was published to contradict the news of PA Act 83 which now recognizes that there is long term Lyme Disease and a need to be treated longer. For more information, see web site. I was Sub Committee Chairman for Education for the PA Governor's Task Force Act 83. Twenty three (23) other states have similar laws in place including NEW YORK.

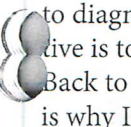
It was once said "It is always better to tell the truth, it will always stand on its own but a lie has to be continually reinforced and the liar has to remember how it was said and has to constantly prop it up and re-support it, but the truth will stand alone." Again through this series follow the Money trail on the 20 Patent holders at the CDC/NIH – who hold over 200 Patents on the Lyme Disease test, along with each of their Universities making Billions of dollars in research monies\$\$\$\$-like the song goes by Abba "Money, Money, Money in a Rich Man's World."

So, we are forced to teach both patient and doctor the clinical method of the signs and symptoms of Lyme Disease and its coinfections. I saw a young 30 year old Ophthalmologist (Eye Doctor) who has a Sub Specialty in Glaucoma yesterday, who was on duty who helped check my eye for an infection. He was most interested, in my research on Lyme Disease because Glaucoma builds up pressure in the eye and eventually can cause you to go blind. Now we know that Lyme Disease causes inflammation, swelling of muscles and different parts of the body including the eyes. As an aside, I told him that I was told myself that 15 years ago I was developing Glaucoma so I used a treatment of antibiotic drops called Ciloxin for an extended period of time, but he looked into my eye yesterday and measured my ocular pressure and he said "you don't have it and your eyes look like you've never had it."

 I have had hundreds of patients diagnosed with Glaucoma and high intraocular eye pressure and over time they improved. My own Mother who was DNA positive for Lyme Disease was diagnosed with it but I treated my mom through her GP. She made 90 years of age without any effects of blindness. She, like my grandfather who made 99 and my grandmother who made 98 had longevity. But both my grandparents cause of death was perforation of the bowels by a nursing home staff using enemas, and my mom who was in stable health died suddenly of pancreatic cancer after being put on a drug called Januvia (Sitagliptin Phosphate), now is being related to rapid pancreatic cancer. My mom was in stable health and could have lived as long as her parents.

Returning to this young Dr. Justin MD, who held me for a long visit and wanted me to explain to him as much as possible about Lyme Disease and blindness. He told me at the ripe old age of 30 and I was around the same age when I learned about Lyme Disease some 30 years ago and said "I always thought it could be linked to something like Lyme Disease but I couldn't get a positive test result."

There you go again, another example of the controlled medical machine (CMM), profiting at the death and suffering of others. This new Doctor and I have agreed to write some future papers together and he has three little children who I promised to send a copy of our children's book which my wife and I co-authored. The Bravest Rescue Puppy can be found at www.mascotbooks.com. Again out of frustration, my wife and I wrote this children's book to help teach children, along with their parents and grandparents who read the book to them to learn about Lyme Disease. Based on a true story about a Puppy we donated for Search and Rescue to a Lady Fire Captain in Ohio (FEMA) Federal Emergency Management Agency. The puppy became one of the top search and rescue dogs in the United States. The illustrator came to me three years ago at age 26, with a misdiagnoses of Multiple Sclerosis (MS). She was in a wheelchair and couldn't use her hands. It turned out to be Lyme Disease and its coinfections. After a couple of months, Amy was starting to walk and I assigned her to draw animal pictures to retrain her hands. She was a Walt Disney trained artist by background. This therapy helped her come back, as did working with my wife and I to put this book together. The second book in the series is complete and book three is started. Future funds will go to support the building of a Lyme Disease Research and Educational hospital on our farm to teach doctors how to diagnose and treat these diseases, and buying more Search and Rescue Puppies to help rescue teams whose mission directive is to search for lost individuals in disaster situations throughout the world.

 Back to the story of eye sight and Lyme Disease cases. 30 years ago, my wife became blind and deaf from Lyme Disease – this is why I'm writing this article on my Saturday afternoon off. God gives us a purpose in life and mine is Helping People with

Lyme Disease. My wife was diagnosed in late 1989 with Macular Degeneration (MD). The macula lutea is an oval-shaped pigmented area near the center of the retina of the human eye. It is estimated that 18 million people will have MD in the future. The saying with MD is you can go and find your way to your movie seat but you'll never be able to watch the picture show because it destroys the central part of your vision. It's a very sad disease.

However, my wife was treated and after getting several doses of antibiotics she went from lying in a bed slowly dying to 48 hours later wanting to paint the bathroom. This is well documented, her sight and hearing loss had returned and she has been quite stable for the last quarter century. Now the eye doctors today say that she has no signs of MD and it looks like she never had it, however her photos and records from the past prove the diagnosis. Her hearing improved until a car accident of 2012 when a construction road sign fell on top of her car and she had a traumatic brain injury and she lost her hearing that we had gotten back for 25 years. So I put the artist and my wife together as a form of physical therapy, to write this series of books with me.

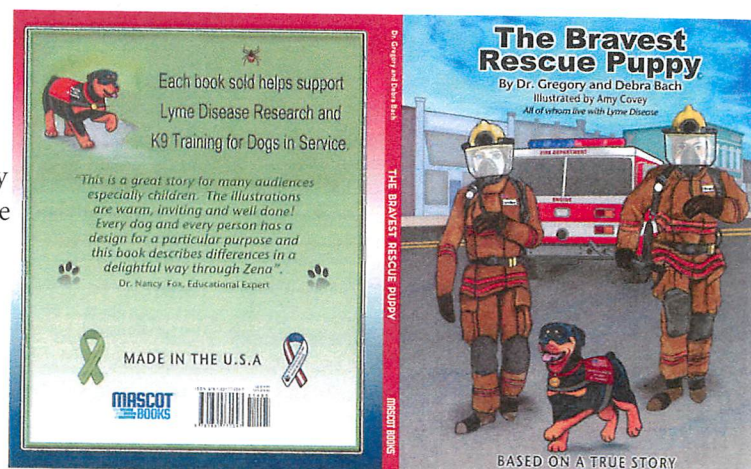
In another case, Brad, a very brilliant steel chemist and a patient for a number of years, was totally blind in his left eye. His retina was detaching which is in the back of the eye (the part of the eye you see with). Retinal detachment and Lyme Disease has a high degree of association. I told him "let's get you treated and have your two eye specialists follow the case." This case is well documented and with treatment for his Lyme Disease and conservative treatment by his eye specialists, which basically they both followed the progress of the eye being restored back to the point that the eye doctors said it regenerated so well it almost looks like he never had it.

Last week, I got a new patient in (I call them respectfully incoming wounded). His diagnosis was missed by under-educated doctors. Under-educated because of information held back by controlled medical machine (CMM) of the 20 patent holders and big Pharma. He had Lyme disease and four years ago he had an operation on his right retina. On his way home from the surgery, he leaned down to pick up a tissue in his car and his eye instantly bled out, they returned to the hospital but the eye could not be saved. The patient reported that he later found out that they never took him off blood thinners used to treat to heart problems from his underlying Lyme Disease.

Strike one – four years later he's reading a symptoms list I published in the Lancaster Farmer (Horse Expo) and comes and sees us to relate his story. Now his left eye, the only good one he has, is starting to lose his vision. After researching his history I asked him, had he been given any steroids recently, and he stated Yes, two months ago he had multiple steroid shots in his back for pain. It took the pain away for a time and then they sent him to a rheumatologist as is common practice to use steroids for inflammation, many of my patients report that they were told they have rheumatoid arthritis (RA) but after testing tens of thousands of Lyme patients for RA factor, I've only found only a hand-full to have RA what they really have is infectious form of arthritis, which used to be called Lyme Arthritis. Over time the name has become Lyme Disease. The worst thing to give a patient who has not been properly, clinically diagnosed for Lyme or a tick borne disease is a Steroid or Immune Suppressant (STEROIDS with Lyme Disease = BLINDNESS). After two weeks of seeing his new rheumatologist who adds more steroids along with his former steroid shots, his vision started to fail within 14 days. Steroids are like putting kerosene on a fire, you don't want suppress the immune system if there is a Possibility of a Bacterial Infection which has not been ruled out properly, including clinically. This is first year medicine. I saw him twice over the past two weeks, he reports his vision is going up and down but better in the AM and the regressing slightly down at 3:00PM. This is a very common cycle in Lyme Disease until the patient stabilizes. The organism wakes up or activates at 3:00AM and goes to sleep or rests at 3:00PM. This has been studied and presented as a scientific fact. It a cyclic organism. This case will be followed for future publication and hopefully it's in God's hands if we are able to restore his sight back fully.

Again, I've seen this cycle in thousands of patients, it is the nature of the disease. I will continue this series next time, but please remember this

Please refuse any steroids if you think you may have Lyme Disease or any tick borne disease infection. They can lead to blindness and make it 10 times worse for the patient. They are only to be used in an emergency crisis when there is no other choice.



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Lymes Disease 103

Children and Lyme Disease



Dr. Gregory Bach

Lyme Disease and children. What a tough topic I have to talk about. We all love our kids and we never want them to be sick, but when the time comes, we are never quite prepared enough. Especially in the very tough and impossible cases that have been linked to Lyme Disease and it's coinfections. (TBD Tick Borne Diseases). *Lyme Disease has been directly linked to a number of childhood diseases over the years. Yet the average Physician and Health Care Worker hasn't a clue, because they're not being trained and taught about Lyme Disease. For example, Autism, deafness, blindness, seizures, paralysis, severe sleeping disorders, nightmares, schizophrenia, asthma, O.C.D. (obsessive compulsive disease), cutting or self-mutilation, drug addiction, aberrant behavior, like children who try to hurt or kill their parents or siblings, hyperactivity, ADD, ADHD, learning disabilities, mental retardation, developmental disabilities, such as, abnormal development, stuttering, Tourette syndrome (where the child cannot help swearing) severe allergies to foods like peanuts, fragrances, soaps, latex, severe anxiety disorders, etc. just to name a few. *Severe anxiety disorders. Let's stop here for a minute. I was watching TV the other day (news flash) Pope blesses child who is going deaf and blind. I got the name off the TV set Fox News story, "Lizzy Myers", the child of Steven and Christine Myers from Ohio is suffering from Ushers syndrome, it causes hearing loss and blindness (R.P. Retinitis Pigmentosa) which causes night blindness and loss

of peripheral vision (side vision). The mother spoke for a second but gave me another clue about this case. "She has always been an ANXIOUS CHILD, so we are afraid to tell her about this disease." *Well let's focus in on this for a second. On my last article on Lyme Disease and blindness, we got to talk about my wife's case from 30 years ago, where she is going deaf and blind. I remember (Usher Syndrome) because they told my wife and Ishe could have this, but if you read the article We discovered her deaf and blindness was coming from her Lyme Disease. When she received the proper treatment she regained her sight and hearing back for over a quarter of a century. Until 2012 when an accident injured her brain and she lost her hearing again. * "What if", "What if" this little five year old girl has a gene for an Usher syndrome, but in the back ground It's Lyme Disease or a tick borne infection causing the gene to turn on and express itself. We know along with other research scientists that genes can turn off and on. I know this for a fact because I had another patient with Huntington's Chorea (HD). This in an inherent condition in which brain cells break down over time. It is very rare. One in 200,000.00 per year. You basically lose control of your mind, body and brain and it can't be cured. Treatment is counseling and physical therapy. It worsens with ANXIETY as per the Mayo Clinic. There's that anxiety word again. It's there for a reason because anxiety makes Lyme Disease patients worse, but it also causes it. The dog chasing its tail. The patient I had with HD was a 35 year old white male, gene positive for HD, was suffering from the above syndrome and

his wife was a nurse and the case was well documented. After two years of treatment his symptoms reversed and never fully went on to be degenerated by his hereditary gene disorder. He simply got better. *Well if anyone can get in touch with this little girls parent, I would like to try and help her, perhaps this could be her story has well! *Let's talk about another little girl, 10 year old Sydney, who is a very special patient in our practice. I met her in late November, she has a biological twin brother who is the smartest 4th grader in his class, but what happened to poor little Sydney? Well we figured it out, at birth she was born first and was given a pint of blood, the other twin did not receive it. I hate to tell our readers that sometimes the blood supply can carry Lyme Disease. The law is, if you're free from Lyme Disease for one year, you can donate blood again. Well 40 to 50 % of my patience who are wonderful, caring people have donated blood before being treated. So it goes on to the next person. But in an emergency if you need blood, you get blood. I always remind my patience if they have a history of Lyme Disease, don't donate. If they need surgery I tell them to Auto donate, meaning set a pint of your own blood a month ahead of the procedure in case they need it. (I have two photos of little Sydney) The first one shows her in a little wheel chair stroller. She was diagnosed in 2014 CDC positive for Lyme Disease and orally treated for a short time and declared cured, but was still sick for you see little Sydney not only couldn't walk, she also really couldn't talk, she also couldn't use her hands and on top of this she had seizures every hour of the day. Her poor mother had been dealing with this for ten years. If you look at the second picture you're going to see her starting to walk only after (8 weeks of the



proper treatment) which consisted of IM (intermuscular antibiotics and oral medication) but besides starting to walk, we had walking braces made for her, which by the way, the insurance company refused to pay for, she also had lost her seizures, and started to string words together like mom - love - you. She started being able to use her hands but not only that, we were able to send Sydney to school. Injust her first week she got a 90% on a vocabulary test, that's and A. *She is very well documented and we will publish her case soon in a medical journal. Ijust ran my pen out of ink, so let's recap - all those above symptoms Ilisted, I had as patience. A quarter of a century later is a long time to learn about these diseases and children, but you must remember which Istressed in all my articles, the local standard Lyme Disease testing is flawed by the 20 Patent Holders and their Universities at the CDC and NIH receiving billions of dollars

in revenue from the old Eliza Test which they kept in place for 37 years, and added an additional 200 patents to that test. Don't you think that 37 years ago when we had cell phones the size of suite cases, and now they fit in your top pocket, we should have a decent test for Lyme Disease by now, but the patent holder are keeping this test in place for their own profit and it's hurting all our children and their families. You have to remember that American Medicine is the world's standard so what happens in American Medicine happens in the whole world. So it's the whole world being affected. *Let's talk about another case of a child. His mother came down from Canada who was a polish immigrant for kidney failure of unknown origin. Her cousin was a board of health nurse who I spoke to on a frequent basis that was very much interested in Lyme Disease. We had made a friendship and one day she asked me if I could help her cousin who was slowly dying from kidney failure. I have been successful with cases of kidney problems in the past because they can be associated with tick borne diseases When I first interviewed her, I said I've been successful with cases like yours before but the case I'm most interested in is that little boy in the waiting room. I said to her, how old is he, he looks about 10 years old? She said to me he is actually 19 and a special needs child. I said that's an unusual scenario. So I took his case on and we found out that like his mother he was suffering from Lyme Disease. When I first met Andrew, he was 40 pounds lighter and one foot shorter and that was 2 Yz years ago. He was supper hyper, couldn't sit still, couldn't look you in the eye and basically was bouncing around the room. He was never educated by the Canadian Government because he

had an IQ in the 50's. They basically taught him how to put his clothes on and tie his shoes. Now I looked at the parents' backgrounds and the father was a Ford Engineer, very smart. His mother was a business person and also very smart. Andrew's sister has her Masters Degree in teaching and his brother is an Air Force pilot. So what's wrong with this story? Andrew had an infection, which didn't allow him to develop to his full potential. So after 2 Yz years of treatment and having him tutored, I was able to petition the Canadian Department of Education for a scholarship to allow him to attend college after he received his high school degree. Please see photo. Andrew was 87 pounds and now is 127 pounds, his height was 4'11Yz" and is now 5'11Yz" tall, in that short span of time we were able to have him catch up with his growth and mental development and he actually went through puberty, his voice changed and has muscles now. *THE DOCTORS MUST TAKE THESE DISEASES SERIOUSLY AND MUST BE EDUCATED PROPERLY, IT IMPACTS ALL OF US AND OUR GENERATIONS TO COME. Until next time as the story continues.

