

Greetings:

I am sorry you or a loved one has this dreadful disease, but now, there is hope for longevity. The Deanna protocol is not a cure but rather a treatment to stop nerve and muscle cells from dying and hence slow down the disease progress.

Please give me some information about yourself such as age, gender and date of disease onset. I would also like to know the following: what is your physical status (limitations and symptoms); are you being treated or followed by a physician; if so by whom; have you had contact with toxic substances.

I am not a treating physician; therefore, I cannot accept you as a patient. However, I can tell you what my daughter is taking. Our treatment consists of natural substances that the body can no longer supply for itself. These substances may be found on line or in health food stores. We recommend you discuss the protocol with your treating physician.

Once I have information for my data base, I will forward the protocol and disclaimer to you. Forgive me for not being able to connect via telephone, but I am inundated with requests and email allows me to handle the volume most efficiently

I would also like feedback from you after 2 to 3 weeks to see how you are responding. It is important to note the frequency and strength of spasms or twitching before and after treatment. Respiratory measurements, muscle strength and any functional improvement before and after treatment.

We have ongoing research being done on the Deanna Protocol using ALS mice at the University of South Florida. We have placed the cart before the horse since we know what works on those affected by ALS, and we are now documenting this in the laboratory to satisfy the requirements of academia. We will also be able to fine tune this treatment protocol by observing the response in ALS mice. At some point, we would like to document the protocol on individuals with ALS in a research project so we can publish our results. However, the academic medical community is not ready to accept our protocol? So your input is extremely valuable in establishing the effectiveness of our treatment.

Remember the treatment is not a cure but has slowed the progression of the disease in individuals. We hope in the near future to have a phase 2, which will repair and regenerate nerve and muscle cells. It would be beneficial to all if you

utilized the blog on www.winningthefight.net so your experiences can be shared by all.

We now have hope,

Dr. T.

Medical Liaison for Winning the Fight

DISCLAIMER

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