Subject: Re: MS and CS
Date: Mon, 8 Sep 2003 22:40:04 -0500
From: "Mike and Nancy" <nancymike@prodigy.net>
To: "Hilary Jacobson" <hjacobson@tiscalinet.ch>

Hi Hilary,

I'm afraid most of the exeeriences have not been compiled anywhere. I need toorganize myself and get someof it down. I just don't know how to go about this, yet.

I do have a short journal I kept which I will add to the end of this email. I will be happy to answer any questions presented to me if I can.

Be sure they understand this is NOT an instant cure. It is slow, subtle and very steady.

Nancy

August 2000

My MS

I have been on Betaseron since it came on the market 6 or 7 years? I would say it did at promised, I have had no exacerbation since I began the injections. However, everyday I hate to get up to see what additional symptom I have to add to my list to get used to.

My right hand is numb, my feet, especially my toes are numb. When I get hot or tired my right leg does not lift well. It drags when I walk. After a day at work, I practically have to crawl to my car. I must hold on to a wall at all times. I really should use a cane. I cannot even go up a curb without holding on to someone or something, No way can I climb a ladder.

When I sit for any length of time, my legs stiffen and get spasms and I have to wait awhile before I can walk. It appears that I have too much to drink. I really should use a cane, but usually I can take my companion's arm to get to my car.

If I sit on the floor for any reason, like play with my grandchildren, I must first get on my knees, then on all four's, then finally I can get up. Just like a cow. I cannot use help getting up from the floor, I need more control. I sit on the floor as little as possible.

When it is hot I must wear a cold pack vest or I cannot walk. My feet are hot all of the time, and I cannot sleep unless my feet are uncovered.

I have night paralysis. I must throw my body in order to turn to another side. My legs are locked I n the fetal position, and it is a real chore to get them unlocked and able to walk. I must use a cane to get to the bathroom during the night. It is about 10 feet from my bed, but it seem like a mile.

Jane Wyman has become my good friend with the Poise pads. I cannot go out without the Ultra Poise pads. If I know I will be away from a bathroom for any length of time, I must use Depends. It goes without saying, I must use the pads at night.

I begin to drink 2 oz. Of CS twice a day - in the morning & at about 3 PM.

Day 4: I begin to drink 8 Oz. Of CS two times per day. I seem to have more energy, and the end of the day seems to come a little later. I do not drag as much to my car.

Day 12: The night paralyzation seems to be easing. I can get out of bed with a little more ease.

Day 14 - day 18: My fingers and toes are tingling more and more. My toes are aching, As the days go by, my fingers seem to be aching, also.

Day 20: I seem to have surreal feelings in my fingers. It's like a far away out of body feeling. They still ache.

Day 21: I am getting out of bed much easier and quicker. I climbed a ladder at work, and I'm not nearly so tired when I leave work. I can actually walk to my car without holding on to the wall. I did some things on the floor at work, and was able to get up without too much trouble.

Week 4: The bottom of my feet are tingling and I could feel whiskers on my husband's face (a surreal feeling).

I could feel the cool bathroom tile on the bottom of my feet.

My legs ached all night. It was very painful, I wanted to scream out. My legs hurt a great deal. The next morning I was able to walk further than I had in years. My husband and I walked about four blocks this morning. I feel stronger and stronger.

Week 5: More and more feeling in both fingers and toes every day. Less surreal and more natural. Both toes and fingers get cold.

Week 10: Seem to have small changes every day. Again my toes ached for several days, then I had more feeling in my toes. It's as though I have a non feeling pad at the bottom of my feet, but feeling all the way around. Like an animal's paw with the padded bottom. It seems I hurt for a few days, then something feels better.

Week 12: I feel like a caterpillar in a cocoon., I wonder if they have pain during the metamorphosis? The bottom of my feet are no longer numb, the fingers on my right hand tingle only at the tips. I don't even think about lifting a heavy container with my right hand. For years, I wouldn't dare lift or I would drop whatever I was holding. I poured coffee from a pot without even thinking about it, until I noticed myself doing it. There is NO WAY I could be working the hours I have this Christmas, if not for the CS. Last year I had to wear my cooling vest all day, every day, and when I when I went home I could barely walk to my car. Some days I literally dragged my right leg to get to my car. I had to hold on to the building to get around the corner and into my car. When I got home I actually crawled on my hands and knees to get up the steps. This year I never once had to wear my cooling vest. I walk normally to my car at the end of the day, and the steps are not too much of a problem. I still go up one leg only, but it is stronger. The fatigue is minimized, also. I've worked many more hours this year than last.

Week 14: I started making my own CS about three weeks ago, and I've had to send samples to San Antonio for testing. It seems the probe they sent me was not working to full potential, and for about a week I was drinking water with a very minimal amounts of silver. After a week, I KNEW IT!!! I was regressing. Things were not working so well again. I was regressing. Thankfully we figured out the problem and within a couple of days I was back on track. Thank God. This set back has convinced me even morel As if that were possible. I have my life back. I will never give up CS.

Week 20: Christmas week. I had 16 people for dinner Christmas eve. I had 7 people for dinner Christmas day, and I worked 11 hours the day after Christmas, and I had 14 people for dinner the next day. I can't remember when I was able to do something like this. I still have night paralysis, but not nearly as bad as it used to be, and I have a lot of stiffness still when I sit a long time, but nothing near as bad as it used to be. My energy level is very high.

Week 24: It seems I was at a standstill for the last several weeks. I didn 't see much improvement, but then I saw NO back sliding - Just status quo. This is OK with me if I don't improve much. I've come such a long way back. However, the stiffness & night paralysis could improve. I drink at least 16 oz. Of CS per day. Some days when I have a lot to do, or am especially busy, I will drink 24 oz.

Week 28: I have been drinking 24 oz. Of CS almost every day. The night paralysis is lessening. It is not taking so long for my legs to get moving in the night or morning. More of the numb parts of my body have feeling.

Week 30: It's hard to believe, but I can still feel better. I can walk almost instantly when I get up from sitting a long time in one place. In the morning my legs are not nearly as stiff as they used to be. I can get moving much quicker then before.

Week 52: It is now a full year and I would say I am about 65% Improved. I have not improved much again, but I have not gone backward either. I can be extremely happy if I stay this way. I feel better than I have in several years.

1 year 4 weeks: Some of my email friends have been discussing drinking CS with Gatorade. They seem to believe the electrolytes in the Gatorade will get the silver into the blood stream quicker. It will not have to go through the entire digestive system to get to work. What have I got to loose?

1 year 8 weeks: They are right. I have been drinking Gatorade two parts to one part CS. It has truly helped. I cannot believe the change. I can't remember ever feeling this good in my entire life. I have some numbness in the very tip of my fingers and the incontinence is still there to some

degree, but then again I drink over 48oz. Of liquid everyday. The morning stiffness is still there to some extent, but it is not as serious. I can get out of bed fairly quick, and it takes only a short time for me to begin walking. If I over do it during the day, my legs do get heavy and tired, but compared to not being able to walk a year ago, it is wonderful.

August 2002

My MS Update

I am sending this to all who have inquired about my original MS journal or helped me along the way.

This is the second anniversary of my long, but wonderful journey with colloidal silver (CS).

I am a 59-year-old female who had relapsing remitting MS for 31 years. About 1995 it changed to secondary progressive MS. Thus began my long road of decline. Everyday I got worse. When I discovered CS I could barely walk. I was beginning to use a cane. I could not even go up on the curb without aid. My prognosis was grim. I had some knowledge of the great properties of silver, so the idea of CS intrigued me. I researched CS. What did I have to loose?

I began drinking 16 oz per day. In about three weeks I began to notice a difference. You already have a log of my first year's progress. I seemed to reach a plateau about this time. I did not improve, BUT I never got worse.

I have since had an MRI and it showed that at this time Aug 2001, I no longer had MS. I have had no new lesions for well over a year. What I was working on at the time is to now repair the damage. Since the damage is to the myelin and not the central nervous system, I was quite confident I could improve.

1 year-6 months: I have researched adding hydrogen peroxide to the CS. One drop of 3% H2o2 per 2 oz of CS. I learned this would cause the tiny silver particles to break up into even more minute particles. After 15 minutes, the peroxide was evaporated out of the CS, so it is not harmful to the body, but the tinier particles of silver got into the blood stream quicker. All this time it was a slow process because by the time the silver got to the myelin where it was needed, it was so diluted, it couldn't penetrate the lesions and kill the mycoplasma (MS virus) Within a week I began to feel old symptoms again. This is what I call a healing crisis: I would get symptoms of the MS as the virus was dying and the dying pathogen aggravated the nerves, so for 2-4 days I would feel like I was having varying degrees of exacerbation. After a short period, it would end and I was improved again.

If I had known about this earlier, I am convinced my recovery time would have increased a great deal.

1 year-9 months: I am sure there is a way to go even quicker... I began to

research IV drips. There are cases of HIV-AIDS infected patients going into complete remission after three infusions. I worked on this project for about six weeks. I finally found someone with a protocol of infusing CS intravenously. I also found a doctor willing to work with me and give this a try.

1 year-11 months: First IV: I had my first IV. By that evening I had my first healing crisis; my legs became extremely heavy (like they were 2 years ago). My fingers tips were still numb, but the numbness was extremely exaggerated. All was better at day four

Second IV a week later: My legs are again aching a great deal, the numbness in my fingers is very intense. It almost feels like they are not attached to me. All better by day three.

Third, fourth, fifth IV: Each time I experienced a reverse of some symptoms I had either forgotten about over the last 40 years, or didn't realize over the years were actually MS symptoms. I've practically no problems at all. I feel better then I have in 15 years. I will have no more IV's, but I will NEVER stop drinking CS.

If I had known about the IV's I probably would have had full recovery even sooner. I am quite sure the old lesions are going away. I am anxious for another MRI to prove this also.

TWO YEAR ANNIVERSARY: very few symptoms.

I would be happy to share what I've learned with anyone. Call me 708-442-6229, snailmail me: Nancy DeLise, 380 Blachawk Road, Riverside, Il. 60546, or email me: nancymike@prodigy.net

I am completely convinced this will help anyone with any autoimmune disease.

	Traditional medicine	vs colloidal silver
Cost:	minimum $1000. per month	$100 per month
Method of Delivery:	injection minimum 1x per week	drink 16 oz. per day
Side effects	page after page	NONE
Prognosis	learn to cope and expect to Continue to deteriorate	kill all MS virus within 12-24 months. Repair damaged

Myelin and regain most functions

12-24 months