Multiple Myeloma

Poly-MVA Cancer Patient Testimonials

Carolyn Huter

No
Picture
Available

Date: December 13, 2011

Diagnosis: Multiple Myeloma Stage 4

4 Updates
9-20-10 – 6-27-11 – 1-12-12 – 10-29-14

(My experience using Poly-MVA the last 15 months)

I was diagnosed with **end stage multiple myeloma in October 2007**. I **began chemo in November and a stem cell transplant in February 2008**. I slowly recovered strength and energy.

In **March 2010 lab test began to show a rise in cancer protein** in blood and urine. My doctors were encouraging another stem cell transplant, **but I felt I would rather take my chances** with the cancer rather than go through recovery of another stem cell transplant when stats showed I could expect even a shorter a period of health than before. I did not feel like it was worth the time it took to recover since I was still feeling good.

I started doing a little research on my own and **Poly-MVA and friends** (other recommended nutritional and detoxification supplements) **seemed like the best alternative method I could find** but the cost was prohibitive.

So, I kept reading and talking to others about methods perhaps less expensive to fight the disease and not tear down healthy cells in my body. **But Poly and friends just kept coming to my attention.**

I **didn't actually start it until September 2010;** at which time my oncologist insisted I must start chemo at least. I did not want to start chemo either because it made me feel so bad.

I wanted to just try the Poly and friends. **He would not consent.** I considered removing myself from his care and proceeding against medical advice. But I consulted with Gary Matson, the consultant assigned to me and **he recommended I take Poly and friends as a complementary to chemo instead of an alternative**. He convinced me that It would **most likely prevent the side effect of chemo** and it has the tendency to **protect my good cells** and **aid the liver** to get rid of all of the cancer cells the chemo was killing more quickly. **It did work that way.**

My oncologist told me when I started chemo again that I would be on **maintenance chemo for the rest of my life.** But **Poly and friends was keeping me from being as sick** as I had felt before so I thought I could follow his advice for a while at least. As months went by, **the cancer protein lab tests started coming closer to normal range**. My oncologist warned me to not get my hopes up. **It would never be 0**.

But by **March of 2011 their tests could not detect the cancer protein**.

The tests had the **same results in June** and the oncologist agreed to let me discontinue chemo.

September's test results were still negative for the cancer protein. I will be tested again this month and I assume every three months thereafter.

I had a wonderful summer(except for a car accident which slowed me down a bit). Even though **I did feel better on chemo this last round** than originally, I really dreaded the thought of continuing for the rest of my life because the steroids they gave along with it made me very nervous with sleep difficulties and fatigue. But the oncologist predicted that chemo(or another stem cell transplant) were the only treatment options when he started it in September 2010. **He had no faith in alternative methods.**

Thank you for the opportunity to share my experience with Poly and friends. I have several loved ones who are battling cancer now but **I have not been able to convince them** to try Poly.

Their doctors, like mine, **think there is no other way to fight it except chemo, surgery or radiation.** I am sorry that the quality of life studies are not taken more seriously in the established medical community.

Carolyn Huter

9-20-2010
Carolyn is struggling to make a decision about what to do **because her numbers are up**. She **doesn't want to do chemo or Velcade** as she knows it only buys a little bit of time. She wanted to prove that Poly could work for her if the Dr. would give her 4 months. **He told her she didn't have 4 months**. She has decided she is **willing to compromise and take his chemo and Velcade to give the Poly more time**. She will go on **12 tsp a day**.

6-27-2011
Carolyn's Doctor says **she has exceeded all of his expectations** so they are **stopping the chemo** and she does not have an appointment until September. She had a car accident and has a broken sternum but **otherwise is great**.

1-12-2012
I realize that MM is considered incurable. **That is why this is all so amazing**. Predictions were **for me to be gone** this coming October. **I don't "feel" like I am going anywhere** just yet and the Lord seems to have placed me in a place of learning and teaching. I am content. But I don't want to take a chance for my oncologist to see a need for chemo. **Dying doesn't scare me as much as slowly killing myself with chemo.**

I did not have the **25 Hydroxy D3 test** with the other testing done on 12/19/11. But I will request it for April. I suppose I should request it each time since it is not in the routine orders.

Carolyn

10-29-2014
Carolyn has been struggling with her Doctor about staying on Poly. **He insisted she stop it for the time being**. She did well for so long but now has a **serious problem with her liver**. He wants her to stay off of Poly for the time being while he tries to get her liver working properly. She told him that the **metals do not stay in her body** because **they are bonded with the other compounds in Poly MVA**….they are not free metals. I told her that she should ask the Doctor about the **negative "long term" side effects caused by Chemo and Velcade** and whether those drugs are what might be causing her liver problems.