

Cystic Fibrosis

Cystic Fibrosis Testimonial # 21:

Hi Wally: Here's something that I received from John Molson that you may not have already. A lady in my Toastmasters group is the President of either the local chapter of the **Cystic Fibrosis Society** or the Canadian President (I got this info second hand so don't know for sure). I had the opportunity to speak with her at the last Toastmasters meeting and asked if she had heard of HMS-90. She gave me the 'what are you trying to sell me' look. Then I asked if she had heard of Dr. Larry Lands because he had done a clinical study using HMS-90 with CF patients. Then she gave me the 'how do you know about him?' look. She told me that she knew 'Larry' quite well. I told her that I would email her some info the next day to read as the meeting wasn't the time or place. The next morning I was thinking that the only info I really had was the 1999 reference to the study that 'was being done' but I had nothing that spoke of the results of this study. I wondered why Dr. Lands hadn't mentioned it to Chris Black (the lady I was speaking to) and if it was because he wasn't happy with the results. So I emailed John, explained my situation and that Chris would certainly be checking any info I sent about Dr. Lands with him as she knew he quite well and asked if he had anything more recent to back up the study. He sent me this! It is just perfect. When I emailed it to Chris, she emailed back a little while later to thank me for the info that said that she will 'love to talk to Larry about this!'. My physiotherapist (who also wants me to do a presentation for her, her staff and some of her patients before too long) knows Chris quite well and said that she will definitely go for something like this if she is satisfied that it is legitimate. A Head Nurse at the hospital here (who is an avid XtraSharp fan and has people waiting to buy it!!) also knows her and had the same opinion. In my email to Chris, I also mentioned that I would be happy to do a presentation to her group if she is interested. So, I'm keeping my fingers crossed that this could turn into something nice. If I read Chris right, if she is impressed with what HMS-90 can do to help CF patients, she will definitely be spreading the word through, not only her group, but all of the groups that she is associated with. Here's hoping. Things are going pretty good here, Wally. I have a presentation lined up for Friday although probably not as many people will show up as I had hoped. The newspaper messed up with my ad and did not put it in this weekend's paper! I have some flyers around town but ... oh well. A few people have phoned and have said they will be there so guess that's better than no one! I signed up a naturopath from Ontario a couple of weeks ago (thanks to Hugh for sending her to me as she asked to speak with someone IN Canada) and a lady from PA last week. My physiotherapist was very interested in the CF study as well and asked if I would forward her the info too (which I have done) as she knows someone who works in the Research dept of the U of BC and who is always looking for new information that can help with asthma. No promises but she was going to be talking (seeing?) her this weekend and wanted the info give her. I have, of course, checked my Immunoweb to see where I stand now and what my commission cheque is for January. Needless to say, I'm a happy camper! A NICE cheque and ... I'm going to get to Montreal for free after all! I thought I was quite a bit away from reaching the 40,000 pts but I guess I don't calculate the points the same way that Immunotec does because, by the end of February, I will be slightly over the 40,000 even if no more new customers or distributors come in! Of course, any new distributors I will put into Paul's downline to try to reach Gold as soon as possible. I am just thrilled, Wally! It feels so good to know that I can continue working this business full time! My confidence has increased by leaps and bounds and I have gone from 'thinking' that I can make this work to 'knowing' I can! What a difference that little change makes! Well, that's my update for now. Talk to you later! **Penny**

Cystic Fibrosis Testimonial # 22:

Eight year-old Zach, a **cystic fibrosis** patient, loved baseball. He was smaller than the rest of the kids, but it was shortness of breath and recurrent respiratory problems, not height, that kept him off the team. He took more care of his nutritional needs and was good about taking his additional vitamins and antioxidant supplementation. His parents learned how to provide him with home aerosol treatments by mask. He has been using both oral and nebulized (by mask) Mucomist (N-acetylcysteine). Although primarily used as a "bench-warmer," Zach is back on the team.

The Celine Dion Cystic Fibrosis Foundation for Kids is using Immunocal.

Irritable Bowel Syndrome (IBS) Testimonial # 23

"Just wanted to let you know that the products really work. I have **Fibromyalgia, osteoarthritis, irritable bowel syndrome (IBS), GERD, headaches and generally exhausted** most of the time. I started initially with 2 packages of Immunocal, vitamins and calcium daily. I was beginning to feel better with this dose but into the 6th week I bumped up to 3 pkg. daily. I now feel great. I had been off of work for 3 mos. after some bladder surgery and just returned back to work as a Registered Nurse this week. Even though it was a busy week putting in approximately 48 hrs, I left work on Friday feeling energetic, a bounce in my step and a smile on my face. My co-workers were amazed at the change---keep in mind they are all nurses or medical professionals and they know how they felt after working less than 5 days in a row. My husband can't believe the change either."

Thanks again, Beverly Pool R.N. December 8, 2001