



It Ain't Easy Being Me: Living With A Rare Disease Known As SCA14

Christopher DeHaven

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Have you ever stopped to think about all the Diseases and Illnesses in the World? Now, take a guess at how many Diseases and Unknown Illnesses there might be in the World... Can't do it, right?

DON'T WORRY, NEITHER CAN I!

If you have ever had a Disease or Illness you know that you need to have at least five things in your favor: GOD in your corner, a sense of humor, thick skin, a great Doctor plus the support of family and friends if you ever plan on fighting your problem.

Now, what kind of person are you? I'm the type of person that likes to stand out in a room filled of People who have Disabilities. I'm not afraid to "make waves" (make a scene) around others just to accommodate someone.

If you ask me I will speak-up and you might not like what I'm going to say but that's your problem, not mine!

Well, by writing this short story I hope that you will learn not to assume anything about anyone because everyone has problems that make people "Jump" to conclusions or assume the "Worst" in someone.

We all have done it! It is just Human Nature and it starts when we are children and as we grow-up by Bullies or even by the people we trust the most, our PARENTS.

This is the story of my life growing up and living with a Very Rare form of Ataxia. I hope that it helps now and even Future Generations by opening the eyes of Parents and even Teachers around the World so they start noticing, talking to and really listening to their kids and children in schools before assuming something is wrong. If I only knew when I was younger that my life would turn out this way I would not have attended School to receive my Degree in Photography and maybe even dropped out of High School to work as much as possible so I could build up my SSDI (Social Security Disability Income).

In this short story I reflect back on many things in my life, some good, bad, crazy and even those "What was I thinking?" moments. I say exactly how I feel about life and I share the "Power of our Mind" as my body starts to break down I begin losing some senses but others are heighten by sharing 2 short stories that I truly believe happened but I have been told otherwise.

Most of the public thinks that we (people with disabilities) just get in the way or disturb their lives but I'm here to tell them all to:

"GET OUT OF OUR WAY!"

I apologize that this book is not longer, it's hard typing a book when you have little to no control of your hands and fingers. I need to "Chicken peck" the keys and my wrist starts hurting after about two hours of typing.

50% from the sale of each book will go to two Charities.

30% will be donated to the National Ataxia Foundation to help in finding treatment, doing research on Ataxia and even finding a cure one day.

The other 20% will go to the MDA Clinic in Oklahoma City for all their help I received from them. I would like to help with the cost of sending kids to camp each year.

90% of all funds made from TV, Movies or other means will go to these Charities, 45%/45%

And the other 10% will be going to the Blanchard High School Alumni Association.

My goal is to present them with a check at the NAF Annual Meeting in Denver, CO each year, during my Yearly Check-up at the MDA Clinic and a check at the Classes Reunion in Blanchard.

To learn more about all the Diseases Associated with Ataxia please go to the National Ataxia Foundation Web Site at www.ataxia.org or you can Email: naf@ataxia.org - You can also reach me on Facebook at www.facebook.com/christopherdehaven1971 to ask questions or talk about the book.

Thank you for your help

This is my book in it's rarest form, right now this book is in Production with Tate Publishing in Mustang, OK, USA and will be at Book Stores soon!

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Eric Alaniz:

In this 21st one hundred year, people become competitive in most way. By being competitive currently, people have do something to make all of them survives, being in the middle of the crowded place and notice by means of surrounding. One thing that at times many people have underestimated the item for a while is reading. That's why, by reading a publication your ability to survive enhance then having chance to stay than other is high. For you who want to start reading any book, we give you this kind of It Ain't Easy Being Me: Living With A Rare Disease Known As SCA14 book as beginning and daily reading book. Why, because this book is usually more than just a book.

Julia Sullivan:

It Ain't Easy Being Me: Living With A Rare Disease Known As SCA14 can be one of your nice books that are good idea. We recommend that straight away because this book has good vocabulary that can increase your knowledge in language, easy to understand, bit entertaining but delivering the information. The article author giving his/her effort to put every word into joy arrangement in writing It Ain't Easy Being Me: Living With A Rare Disease Known As SCA14 nevertheless doesn't forget the main level, giving the reader the hottest along with based confirm resource information that maybe you can be one of it. This great information could drawn you into fresh stage of crucial contemplating.

Leesa Banta:

This It Ain't Easy Being Me: Living With A Rare Disease Known As SCA14 is fresh way for you who has attention to look for some information since it relief your hunger of information. Getting deeper you onto it getting knowledge more you know or else you who still having bit of digest in reading this It Ain't Easy Being Me: Living With A Rare Disease Known As SCA14 can be the light food in your case because the information inside this particular book is easy to get by anyone. These books develop itself in the form that is certainly reachable by anyone, yes I mean in the e-book contact form. People who think that in reserve form make them feel sleepy even dizzy this e-book is the answer. So there isn't any in reading a book especially this one. You can find actually looking for. It should be here for you. So , don't miss that! Just read this e-book sort for your better life and knowledge.

Virgil Santamaria:

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