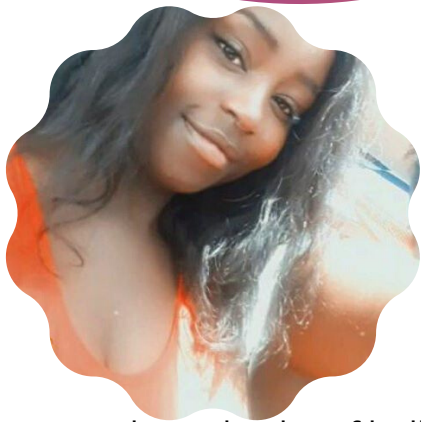


Lupus In Color Leila Muxinga

BUTTERFLIES OF HOPE WARRIOR STORY



My name is Leila Muxinga and I was diagnosed with Lupus at the age of 14 and this is my story.

Back before I got diagnosed, I was misdiagnosed for about 2 years by a doctor who passed on a while ago, he was our family doctor. Later on I got referred to my current physician, who is a Specialist Dr. Goagoseb and he officially diagnosed me with SLE in 2010.

I went through a lot of bullying. I was bullied by the very same people whom I thought were my friends and best friends.

I was bullied for how my fingers were shaped, how they were blue/purple, how I would be in and out of the hospital. They talked about how nasty they thought I was.

But to my high school friends, the true friends, Ralitza Gawises, Bernelee Kahiiko and Trisha Skrywer, I just want to use this opportunity to thank you guys. I love you all still today.

So to make things short, I have been living with Lupus for over 13 years. I am legally disabled, but I have my own company and I am my own BOSS.

I just want to thank the all mighty, my King in Heaven and His one and only begotten Son Jesus Christ for His mercy and His grace. Without Him, I wouldn't have Hope.

Today I have my own Lupus Support group on WhatsApp and also a Christian group whereby I plan on motivating the young people.

THIS IS MY STORY

LUPUS IN COLOR - FLIGHT OF THE BUTTERFLIES - BUTTERFLIES OF HOPE WARRIOR STORIES
SPREADING LUPUS AWARENESS ONE BUTTERFLY AT A TIME
#LUPUSINCOLOR #FLIGHTOFTHEBUTTERFLIES #BUTTERFLIESOFHOPE