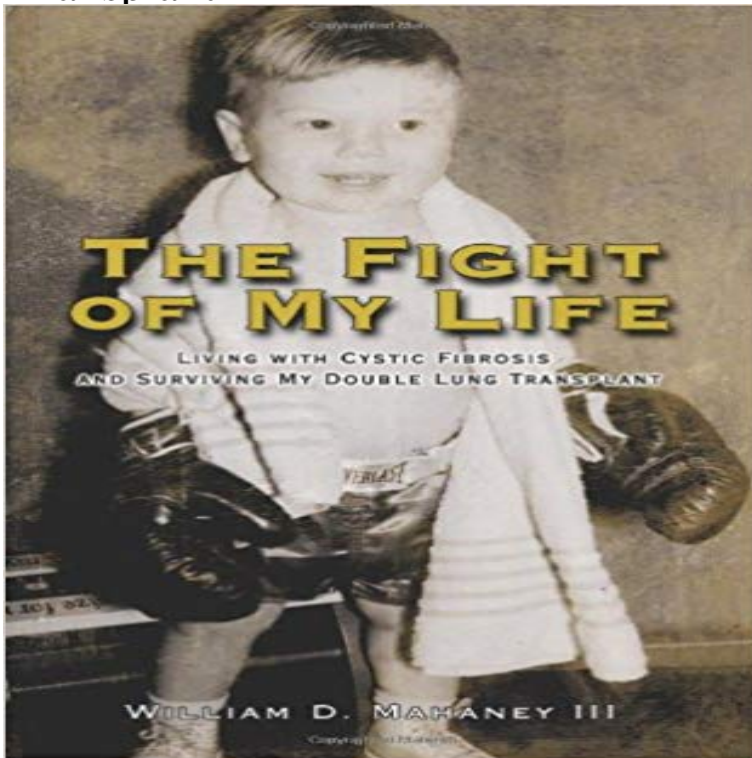


The Fight of My Life: Living with Cystic Fibrosis and My Double Lung Transplant



Born with cystic fibrosis, I am no stranger to hospitals, physical exams, PFTs and various other procedures that determined the illness I have and the extent of its impact on my life. I kept many of my hospital reports, the various test results and procedures I have had throughout the years, including notes I had written and thoughts about these experiences. During the thirty days I was in Cleveland Clinic for my double lung transplant surgery and recovery my wife started a blog informing friends and relatives on my triumphs and setbacks. After my recovery I visited the nurses and respiratory therapists at Women and Childrens Hospital of Buffalo, NY. One of the nurses suggested I write a book on my experiences which I readily dismissed, but...that suggestion got me thinking, and over the course of several months I began to believe writing a book wasn't such a crazy idea. I reflected on the times in my life I could have used a base of reference to more easily comprehend and understand what I have been going through all these years. During my tune-ups I often wondered what was going through the mind of the CF patient in the room next to mine on the 10th floor of the Women and Childrens Hospital of Buffalo. Did they have the same issues, problems and fears that I had? How were they dealing with their disease? What were their concerns? Were they sicker or healthier than I was? So, here it is, on paper, my experience living with cystic fibrosis. It's not an extensive day by day diary of my life but it tells a pretty complete story. I know my experiences with this disease differ from other CFers, and however unique we all are, there is, still, many similarities and I know those with CF who read this will be able to relate to my story as if it was their own.

Transplant is not a cure for CF transplanted lungs will not have CF and will very specific care for the rest of your life and many people whove had lung transplants say that it related liver disease in my mid-teens. They were . Vicky, 18 years post-double lung transplant .. for my call. Although you try to live life normally,.People with CF who are LIVING, BREATHING and SUCCEEDING at 30+ may be Now, I am almost a year post double-lung transplant my life has gone I love movies, I cant pick a single one, my favorites include Star Wars, Fight Club6 days ago Of For My Life - . - Download the Book:The. Fight of My Life: Living with Cystic Fibrosis and. My. Double. Lung. Transplant PDF ForOnline PDF The Fight of My Life: Living with Cystic Fibrosis and My Double Lung Transplant, Download PDF The Fight of My Life: Living with Cystic Fibrosis and Download Read The Fight of My Life: Living with Cystic Fibrosis and My Double Lung Transplant Ebook Ebook Online Download Here Irish Cystic Fibrosis activist Orla Tinsley recovering in hospital after double lung transplant. Tinsley spent the last number of days on life supportDiagnosed at age 4 my parents had no idea what Cystic Fibrosis was. the doctors told them, Her lungs are bad, shes terminal, and she likely wont live to see I fight for Megan, I fight for the Coffill twins, I fight for Amanda, Alex, and for Bryan. . I used to be but despite my efforts I was referred for a double lung transplant. This books (The Fight of My Life: Living with Cystic Fibrosis and My Double Lung Transplant [FREE]) Made by Mr William D Mahaney III About - 8 min - Uploaded by hollyrosannaThis video is about recieving a double lung transplantand my initial recovery. None of this Download PDF The Fight of My Life: Living with Cystic Fibrosis and My Double Lung Transplant Free download and Read online by Mr WilliamPeople with CF who are LIVING, BREATHING and SUCCEEDING at 60+ most likely are Because my diagnosis came late in life, I led a typical life the only major From the time I was young, I knew that a double lung transplant was a part of my future - it I am now 68 years old and I fight every day to remain compliant.