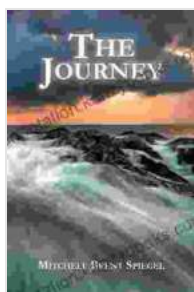


The Journey: A Family's Firsthand Account of Living with ALS

In 2016, my husband, Scott, was diagnosed with ALS, also known as Lou Gehrig's disease. ALS is a progressive, fatal neurodegenerative disease that attacks the motor neurons in the brain and spinal cord. There is no cure for ALS, and the average life expectancy after diagnosis is just two to five years.



The Journey: A Family's Firsthand ALS Account

by Jessica Seraben

★★★★☆ 4.5 out of 5

Language : English
File size : 602 KB
Text-to-Speech : Enabled
Screen Reader : Supported
Enhanced typesetting : Enabled
Word Wise : Enabled
Print length : 71 pages
Lending : Enabled



When Scott was diagnosed, we were devastated. We had two young children, and we couldn't imagine our lives without him. But we were also determined to make the most of the time we had left together.

This book is a memoir of our family's journey through ALS. It is told through the voices of all of us: Scott, me, and our children. We share our experiences with the disease, from the initial diagnosis to the daily

challenges of living with ALS. We also share our hopes, our fears, and our love for each other.

This book is not a medical guide to ALS. It is not a how-to manual for dealing with the disease. It is simply the story of our family's journey. We hope that our story will inspire others who are facing similar challenges. We hope that it will give them hope and strength to keep fighting.

A Father's Perspective

By Scott

When I was diagnosed with ALS, I was scared. I didn't know what the future held for me or my family. But I was also determined to make the most of the time I had left.

I wanted to be there for my wife and children. I wanted to watch them grow up. I wanted to create memories that they would cherish forever.

ALS has taken away my ability to move and speak, but it has not taken away my mind. I am still the same person I was before I was diagnosed. I still love my wife and children. I still have the same hopes and dreams for the future.

This book is my way of sharing my story with the world. I want people to know that even though I have ALS, I am still living. I am still fighting. And I am still loved.

A Mother's Perspective

By Jennifer

When Scott was diagnosed with ALS, my whole world fell apart. I couldn't imagine my life without him. But I quickly realized that I had to be strong for my children.

I have watched Scott fight this disease with courage and grace. He has never given up hope. He has never stopped fighting.

This book is a testament to Scott's strength and determination. It is also a testament to the power of love and family. We have faced this challenge together, and we will continue to face it together.

A Daughter's Perspective

By Sarah

When my dad was diagnosed with ALS, I was 10 years old. I didn't really understand what was happening, but I knew that it was serious.

I have watched my dad go through a lot of changes over the past few years. He can't walk or talk anymore, but he is still the same dad I have always known. He is still kind, funny, and loving.

I am so proud of my dad for fighting this disease with such courage. He is an inspiration to me and to everyone who knows him.

This book is a story of hope and love. It is a story about a family who is facing a difficult challenge together.

A Son's Perspective

By Michael

When my dad was diagnosed with ALS, I was 8 years old. I didn't really know what was going on, but I knew that it was something bad.

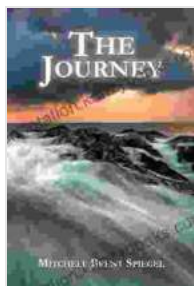
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Free Download Your Copy Today

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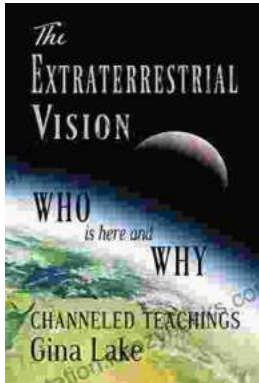
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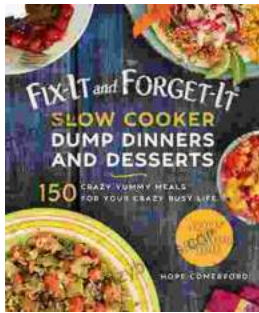
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