

Seizure Mama and Rose: An Epilepsy Memoir

A poignant and inspiring memoir about a mother's journey to understand and embrace her daughter's epilepsy, and the extraordinary bond they share.

When Rose was diagnosed with epilepsy at the age of two, her mother, Jennifer, was devastated. She knew nothing about the condition and was terrified of what the future held for her daughter.



Seizure Mama and Rose: An Epilepsy Memoir

by Flower Roberts

★★★★☆ 4.7 out of 5

Language : English

File size : 426 KB

Text-to-Speech : Enabled

Screen Reader : Supported

Enhanced typesetting : Enabled

Word Wise : Enabled

Print length : 176 pages

Lending : Enabled

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But Jennifer was determined to learn everything she could about epilepsy and to help Rose live a full and happy life. She joined support groups, read countless books, and consulted with numerous doctors.

Over time, Jennifer came to understand that epilepsy is a manageable condition. With the right treatment, Rose could live a normal life. But she

also learned that epilepsy can be a challenging condition, both for those who have it and for those who love them.

In *Seizure Mama and Rose*, Jennifer shares her family's journey with epilepsy. She writes about the challenges they have faced, the triumphs they have celebrated, and the love that has sustained them through it all.

Seizure Mama and Rose is a powerful and inspiring memoir that will resonate with anyone who has been touched by epilepsy. It is a story of love, acceptance, and hope.

Praise for *Seizure Mama and Rose*

"A beautifully written and deeply moving memoir. Jennifer's story is a testament to the power of love and the resilience of the human spirit." —**Dr. Orrin Devinsky**, Director of the Comprehensive Epilepsy Center at New York University Langone Medical Center

"An essential read for anyone who wants to understand epilepsy and its impact on families. Jennifer's honesty and compassion shine through on every page." —**Susan Axelrod**, Executive Director of the Epilepsy Foundation

"*Seizure Mama and Rose* is a gift to the epilepsy community. Jennifer's story will provide hope and inspiration to families who are facing the challenges of epilepsy." —**Kathleen Walsh**, President of the Child Neurology Society

About the Author

Jennifer is a writer and advocate for people with epilepsy. She lives in New York City with her husband and two children.

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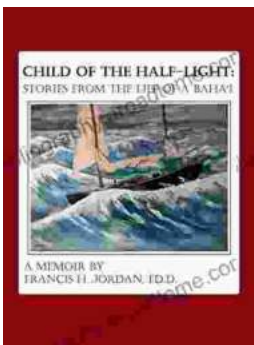


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