



# STRICKEN

VOICES FROM THE HIDDEN EPIDEMIC  
OF CHRONIC FATIGUE SYNDROME

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EDITED BY PEGGY MUNSÖN

# Stricken Voices From The Hidden Epidemic Of Chronic Fatigue Syndrometc

**Joy Selak, Steven Overman**



## **Stricken Voices From The Hidden Epidemic Of Chronic Fatigue Syndrome:**

Stricken Peggy Munson, 2000 This anthology delivers an emotional inside look at people suffering with chronic fatigue immune dysfunction syndrome CFIDS a debilitating disease that involves virtually every organ system and is often wrongly called the yuppie flu Through their distinct voices we feel the grief and hope of those stricken with CFIDS and learn of the complex nature of this misunderstood disorder These are compelling stories about a quiet and baffling epidemic To view an excerpt online find the book in our QuickSearch catalog at [www.HaworthPress.com](http://www.HaworthPress.com) **Stricken** Peggy Munson, 2014-02-04 Develop a better understanding of what CFS CFIDS sufferers are going through In the 1980s a strange emerging epidemic baffled doctors in Incline Village Nevada Dismissed by the media as The Yuppie Flu Chronic Fatigue Immune Dysfunction Syndrome CFIDS turned out to be neither a faddish disease of the wealthy nor a passing trend but rather a growing worldwide epidemic of devastating proportions In the voices of a South African journalist a former marathon runner a teenage girl a public health activist living on the edge of race and gender a cancer patient neglected by doctors because of disdain for her chronic illness and a theologian relearning the art of spiritual empathy the people who share their stories in Stricken Voices from the Hidden Epidemic of Chronic Fatigue Syndrome defy cultural stereotypes and explore the complex social and political dynamics of this hidden epidemic Through their distinct points of view we feel the grief and hope of those stricken with CFIDS and learn of the complex nature of this misunderstood disorder These are compelling stories about a quiet and baffling epidemic The first American anthology to contain stories from a diverse range of people with CFIDS Stricken offers an intimate look at the political and social issues surrounding CFIDS as told by those who are living through this ordeal Stricken addresses several issues such as why some doctors still do not believe CFIDS is real how the disease is mocked in the media myths about this illness the personal fight for medical or public recognition the skepticism and hope that is felt by the ever growing number of CFIDS sufferers Stricken confronts fascinating CFIDS issues such as the Kevorkian suicides accusations of Munchausen Syndrome By Proxy Gulf War Syndrome the role of storytelling in a memory impaired patient movement and the feasibility of mass activism in a disabled population With contributions from Pulitzer prize nominated writer Susan Griffin renowned health writer and radio host Gary Null well known feminist activist Joan Nestle and award winning poet and essayist Floyd Skloot Stricken is an eloquent testament to the heroism defiance and diversity of the CFIDS community **The Fibromyalgia and Chronic Fatigue Resource Book and Life Planner Workbook** Dawn Hughes, 2001-01-01 A must resource for the millions of people affected by Fibromyalgia and Chronic Fatigue Syndromes FMS and CFS unlike well known and understood diseases like cancer spring boards a journey for education understanding and compassion The book powerfully captures this journey and is the first to provide patients family and friends with the imperative resources to find support through their walk In addition The Life Planner is a workbook that maps out a step by step guide for adapting and coping with the lifestyle changes fibromyalgia and chronic fatigue patients must face The book is

divided into four sections The Syndromes Provides an extensive look at the cause diagnosis symptoms and treatments From current research to clinical trials The Resources A comprehensive listing of FMS CFS books medical journals association newsletters periodicals and video s recommended by FMS CFS Associations The Support Worldwide listings of FMS CFS Associations as well as associations for sub categories of these syndromes Including Advocacy pain management alternative therapies and disability issues The Life Planner From diagnoses to acceptance The Life Planner details how to manage life emotionally and physically A compassionate look at What to expect how to work through feelings goal planning worksheets and charts exercise nutritional choices how to educate family and friends and work options Written by a fibromyalgia and chronic fatigue patient the book outlines not only the journey for the patient but gets to the heart of awakening family and friends to the devastating effects these illnesses cause and provides the tools needed for years to come

**Caring for the M.E. Patient** Jodi Bassett,2011 Essential information on the neurological disease M E sourced from the world s leading M E experts Suitable for M E patients their friends and family partners carers or doctors The book includes a foreword by international M E expert Dr Byron Hyde M E is a distinct neurological disease and is not at all the same thing as CFS Learning the facts is not time consuming or complicated Supporting your ill friend or family member or patient more fully by being aware of the basic facts of M E could make all the difference in the world to them This book shows you how Jodi Bassett is the founder of the international M E charity HFME Jodi contracted M E in 1995 when she was just 19 HFME contributors also aim to advocate for those non M E patients who have been given the always meaningless CFS diagnosis and subsequently denied correct diagnosis and treatment

*Tuning the Brain* Jay Goldstein,2013-10-18 In this remarkable volume Dr Jay A Goldstein clearly presents both the theoretical and the practical aspects of this revolutionary approach to treating CFS and other conditions that have often been termed psychosomatic Dr Goldstein will show you how he achieves results for patients with CFS and a variety of other syndromes in days rather than months or years From the most basic questions What is neurosomatic medicine and How can treatments sometimes work so rapidly to specific technical concerns What is receptor profiling and how does it indicate the type of receptor dysregulation in an individual patient Tuning the Brain Principles and Practice of Neurosomatic Medicine provides the answers in a clear and cogent manner You ll learn which abnormalities in brain function produce neurosomatic disorders and how an understanding of these abnormalities can help you provide effective treatment

**An Evocative Autoethnography of Living Alongside Myalgic Encephalomyelitis (ME)** Orlagh Farrell Delaney,2021-07-30 This ground breaking book explores and explains the day to day realities of living long term with Myalgic Encephalomyelitis ME ME is an acquired complex disorder characterised by a variety of symptoms affecting multiple systems of the body Marked fatigue and weakness sickness cognitive dysfunction and symptom flare up can follow any physical or cognitive exertion It is estimated that there are 17 24 million sufferers worldwide The author has lived with moderately severe ME for the last 18 years Utilising autoethnography as a methodology

and drawing on multidisciplinary social science theory the book tells the story of the author's own lived experiences of the illness and how she sought to reimagine a self or a life living alongside the illness that could still be considered a good life This autoethnographic book is beautifully and evocatively written It is a work of scholarship that will be highly accessible to academic and other readers It is also a comprehensive introduction to autoethnography as a methodology but it is much more The images and poetry complement the narrative discussion and are exemplary as part of an approach that integrates creative work with academic argument It illuminates the struggles of living with ME and how there can be sanctuary

**Brilliant Imperfection** Eli Clare, 2017-01-06 In *Brilliant Imperfection* Eli Clare uses memoir history and critical analysis to explore the deeply held belief that body/minds considered broken need to be fixed Cure serves many purposes It saves lives manipulates lives and prioritizes some lives over others It provides comfort makes profits justifies violence and promises resolution to body/mind loss Clare grapples with this knot of contradictions maintaining that neither an anti-cure politics nor a pro-cure worldview can account for the messy complex relationships we have with our body/minds The stories he tells range widely stretching from disability stereotypes to weight loss surgery gender transition to skin lightening creams At each turn Clare weaves race disability sexuality class and gender together insisting on the nonnegotiable value of body/mind difference Into this mix he adds environmental politics thinking about ecosystem loss and restoration as a way of delving more deeply into cure Ultimately *Brilliant Imperfection* reveals cure to be an ideology grounded in the twin notions of normal and natural slippery and powerful necessary and damaging all at the same time

**You Don't LOOK Sick!** Joy Selak, Steven Overman, 2005-03-08 Coming to terms with this reality was a lot like accepting the death of a loved one *You Don't LOOK Sick! Living Well with Invisible Chronic Illness* chronicles a patient's true life accounts and her physician's compassionate commentary as they take a journey through the three stages of chronic illness Getting Sick Being Sick and Living Well This resource helps you focus on building a meaningful life that contains illness as opposed to a life of frustration and fear Designed for patients in at all stages of the chronic illness journey this book will also be illuminating for caregivers and loved ones From the book I've learned that having a chronic illness is not a prison sentence It does not mean I must spend the rest of my life feeling depressed and angry locked away from the world inside my little sick box It does not mean that I am useless and no longer have any gifts to share but it may mean that I must develop some new ones *You Don't LOOK Sick!* addresses practical aspects of chronic illness such as hiring a doctor managing chronic pain coping with grief and the loss of function winning battles with health and disability insurers countering the social bias against the chronically ill recognizing the limitations of chronic illness care and charting a path for change In *You Don't LOOK Sick! Living Well with Invisible Chronic Illness* you will find stories dialogue humor examples and analogy of the three stages to illustrate a challenging but navigable journey You will also find suggested reading materials for learning to live well medical Internet resources illness specific Web sites names and addresses of national associations and a bibliography of medical books by

topic The short chapters and straightforward language of the book will be helpful for readers who are weary and dispirited From the authors I ve learned that having a chronic illness is not a prison sentence It does not mean I must spend the rest of my life feeling depressed and angry locked away from the world inside my little sick box It does not mean that I am useless and no longer have any gifts to share but it may mean that I must develop some new ones Joy H Selak My goal is to work with patients so that like world class athletes they can perform at their peak capacity My job is more than giving answers I must educate counsel and encourage patients to set goals and implement a personal care program as well as take appropriate medications Dr Steven Overman The authors are experienced public speakers If you wish to inquire about their availability to speak to patients or health care professionals please contact Joy Selak by email at JoyWrites austin rr com *Women, Body, Illness* Pamela Moss, Isabel Dyck, 2003-04-14 This provocative and moving work explores concepts of body and space to better understand the daily lives and struggles of women with chronic illness Moss and Dyck show how such women coping with associated notions of illness health and being female restructure their physical and social environments through the strategies they choose to accommodate disabling illnesses such as chronic fatigue syndrome multiple sclerosis or rheumatoid arthritis Strategies might include disclosing or concealing illness from employers and friends seeking or rejecting emotional support through old friends and new contacts and pursuing or resisting specific diagnoses from the biomedical community Featuring a wealth of original research and personal stories *Women Body Illness* tells the tales of chronically ill women forging networks of support redefining themselves and challenging what it is to be ill *The Disability Studies Reader* Lennard J. Davis, 2016-10-19 The fifth edition of *The Disability Studies Reader* addresses the post identity theoretical landscape by emphasizing questions of interdependency and independence the human animal relationship and issues around the construction or materiality of gender the body and sexuality Selections explore the underlying biases of medical and scientific experiments and explode the binary of the sound and the diseased mind The collection addresses physical disabilities but as always investigates issues around pain mental disability and invisible disabilities as well Featuring a new generation of scholars who are dealing with the most current issues the fifth edition continues the Reader s tradition of remaining timely urgent and critical **Race, Gender, Sexuality, and Social Class** Susan J. Ferguson, 2023-06-08 *Race Gender Sexuality and Social Class* Fourth Edition is an anthology of readings that explores the ways these social statuses shape our experiences and impact our life chances in society today Organized around broad topics identity power and privilege social institutions etc rather than categories of difference race gender class sexuality to underscore the idea that social statuses often intersect with one another to produce inequalities and form the bases of our identities in society The text features readings by leading experts in the field and reflects the many approaches scholars and researchers use to understand issues of diversity power and privilege Included with this title LMS Cartridge Import this title s instructor resources into your school s learning management system LMS and save time Don t use an LMS You can still access all of the

same online resources for this title via the password protected Instructor Resource Site

**How to Be Sick (Second Edition)** Toni Bernhard, 2018-09-25 A brand new edition of the best selling classic with added and updated practices In 2001 Toni Bernhard got sick and to her and her partner's bewilderment stayed that way As they faced the confusion frustration and despair of a life with sudden limitations a life that was vastly different from the one they'd thought they'd have together Toni had to learn how to be sick In spite of her many physical and energetic restrictions and sometimes because of them Toni learned how to live a life of equanimity compassion and joy This book reminds us that our own inner freedom is limitless regardless of our external circumstances Updated with new insights and practices hard won from Toni's own ongoing life experience this is a must read for anyone who is or who might one day be sick or in pain

**How to Be Sick** Toni Bernhard, 2010-05-10 This life affirming instructive and thoroughly inspiring book is a must read for anyone who is or who might one day be sick It can also be the perfect gift of guidance encouragement and uplifting inspiration to family friends and loved ones struggling with the many terrifying or disheartening life changes that come so close on the heels of a diagnosis of a chronic condition or life threatening illness Authentic and graceful How to be Sick reminds us of our limitless inner freedom even under high degrees of suffering and pain The author who became ill while a university law professor in the prime of her career tells the reader how she got sick and to her and her partner's bewilderment stayed that way Toni had been a longtime meditator going on long meditation retreats and spending many hours rigorously practicing but soon discovered that she simply could no longer engage in those difficult and taxing forms She had to learn ways to make being sick the heart of her spiritual practice and through truly learning how to be sick she learned how even with many physical and energetic limitations to live a life of equanimity compassion and joy And whether we ourselves are ill or not we can learn these vital arts from Bernhard's generous wisdom in How to Be Sick

**Traveling Blind** Susan Krieger, 2010 TRAVELING BLIND is a deeply reflective description of coming to terms with lack of sight It reveals the invisible work of navigating with a guide dog while learning to perceive the world in new ways The author travels with Teela her lively golden dog through airports city streets and Southwest desert landscapes exploring these surroundings with changed sight

[The Bloomsbury Handbook to the Medical-Environmental Humanities](#) Scott Slovic, Swarnalatha Rangarajan, Vidya Sarveswaran, 2022-07-28 Bringing together two parallel and occasionally intersecting disciplines the environmental and medical humanities this field defining handbook reveals our ecological predicament to be a simultaneous threat to human health The book Represents the first collection to bring the environmental humanities and medical humanities into conversation in a systematic way Features contributions from a wide range of interdisciplinary perspectives including literary studies environmental ethics and philosophy cultural history and sociology Adopts a truly global approach examining contexts including but not limited to North America the UK Africa Latin America South Asia Turkey and East Asia Touches on issues and approaches such as narrative medicine ecoprecarity toxicity mental health and contaminated environments Showcasing and surveying a rich

spectrum of issues and methodologies this book looks not only at where research currently is at the intersection of these two important fields but also at where it is going

**Cure Unknown** Pamela Weintraub, 2013-06-25 This history of Lyme disease is a tale of biological complexities scientific turf battles political intrigue human egos and money lots of it Sacramento Bee Winner American Medical Writers Association Book Award When Pamela Weintraub a science journalist learned that her oldest son tested positive for Lyme disease she thought she had found an answer to the symptoms that had been plaguing her family for years but her nightmare had just begun Almost everything about Lyme disease turned out to be deeply controversial from the microbe causing the infection to the length and type of treatment and the kind of practitioner needed On one side of the fight the scientists who first studied Lyme describe a disease transmitted by a deer tick that is hard to catch but easy to cure no matter how advanced the case On the other side rebel doctors insist that Lyme and a soup of co infections cause a complicated spectrum of illness often dramatically different and far more difficult to treat than the original researchers claim Instead of just swollen knees and a rash patients can experience exhaustion disabling pain and a Lyme fog that leaves them dazed and confused As patients struggle for answers once treatable infections become chronic In this nuanced picture of the intense controversy and crippling uncertainty surrounding Lyme disease Pamela Weintraub sheds light on one of the angriest medical disputes raging today The most comprehensive book ever written about the past present and future of Lyme disease *Cure Unknown* exposes the ticking clock of a raging epidemic and the vulnerability we all share Revised with a new chapter A thoroughly researched and well written account of the disease s controversial history Jane Brody The New York Times A comprehensive and compassionate guide to a dreaded illness named after a bucolic tick infested town on Long Island Sound Hartford Courant

*All in My Head* Paula Kamen, 2005-02-15 A personal cultural and scientific exploration of chronic untreatable pain

*Silicone Injury* Hemitra Elan\*tra Vedenetra, 2013-08-13 In this memoir of courageous survival and endurance author Hermitra Elan tra Vedenetra recounts her harrowing journey through Chronic Fatigue Syndrome Fibromyalgia Connective Tissue Disease Multiple Chemical Sensitivity and Electromagnetic Hypersensitivity as a consequence of chemical poisoning from her silicone breast implants For her breasts implants were supposed to be the solution to a problem Instead a much bigger problem ensued and she was sent careening down a precipice she never could have foreseen Her body was destroyed by silicone and so was her life Her task then became one of survival Her story begins in a small town in Italy unfolds in New Jersey New York City and Dallas Texas and ends in the high mountain desert of Arizona where she had to escape to save her life

**Come, Let Me Guide You** Susan Krieger, 2015 Explores the intimate communication between author Susan Krieger and her guide dog Teela Golden Retriever Yellow Labrador her lively Golden Retriever Yellow Labrador over the 10 year span of their working life together This is a book about being led by a dog to new places in the world and new places in the self a book about facing life s challenges outwardly and within and about reading those clues those deeply felt signals that can help guide the way It is also more broadly about



the importance of intimate connection in human animal relationships academic work and personal life Krieger continues the narrative beginning at the moment she must confront Teela s retirement and then reflecting on the span of their relationship  
Publisher     **Refined in the Furnace of Affliction** R. Paul Gregory,2008

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