

LYME DISEASE



My Search for a Diagnosis

Linda Hanner

*A medical detective story and
an inspirational book in one*

Foreword by John J. Witek, M.D.

Formerly titled *Of Power & Love & Sound Mind*

Lyme Disease My Search For A Diagnosis

Constance A. Bean



Lyme Disease My Search For A Diagnosis:

Lyme Disease Linda Hanner, 1991 Four years into her battle with Lyme disease Linda Hanner's world had become terror filled and seemingly hopeless She had reached the point where she not only longed to die but to wipe out the fact that she had ever existed Linda's story begins in 1981 in the midst of a typically hectic Minnesota summer when there was no time to slow down for illness But the vicious Lyme spirochetes that waged war on her body were unconcerned with her goals and plans One day Linda was energetic in constant motion the next she was knocked off her feet weak helpless confused For six years her illness eluded diagnosis When tests repeatedly failed to reveal any conclusive answers many doctors insisted her problems must be psychosomatic They were the professionals and they seemed so sure Before the illness Linda's faith had been strong She clung to it though it seemed to slip away from her grasp with every step further into the wilderness of chronic undiagnosed illness Finally through the maze of confusion and despair she found a place of comfort rest hope acceptance She learned to believe again in God and in her own self worth Those who have suffered with Lyme disease undiagnosed illness or depression and those who struggle with faith in the midst of pain will identify with Linda's story Book jacket *In the Crucible of Chronic Lyme Disease* Kenneth B. Liegner M.D., 2015-10-08 Following completion of his medical training and a one year stint as attending physician on Howard Champion's Surgical Critical Care Service and MedStar Unit at Washington Hospital Center in the District of Columbia Kenneth Liegner M.D. returned to Westchester County home of his Alma Mater New York Medical College to start a private practice Unwittingly he had plunked himself down in the heart of a burgeoning epidemic of Lyme disease His patients confronted him with puzzling syndromes that defied tidy formulations of the illness and thrust him in to a Maelstrom of medical controversy Lyme disease a new poorly understood disease emerged hand in hand with the rise managed care Physicians caring for persons with Lyme disease loyal to the Hippocratic Oath and serving what they saw as patients best medical interests found themselves on a collision course with a new Corporate Medical Ethic dedicated to maximizing profit One practitioner's work over 25 years is presented here along with correspondence with many principals in the field Documentational in nature and not written as a narrative the materials nonetheless convey the intensity of the struggle to characterize the nature of Lyme disease and the desperate fight for proper diagnosis and treatment upon the outcome of which patients very lives depended The volume includes protocols useful as reference materials for patients and practitioners alike as well as photographic images of many persons important in the history of Lyme disease Foreword by Pam Weintraub Senior Editor of aeon digital magazine and author of award winning book *Cure Unknown Inside the Lyme Epidemic* Preface by Paul W Ewald Professor of Evolutionary Biology at the University of Louisville and author of *Plague Time* Beating Lyme Constance A. Bean, 2008-06-25 More than 30 years after it was first diagnosed Lyme disease remains one of our most misunderstood illnesses This frequently misdiagnosed infection is spreading at an alarming rate and if not treated early can cause debilitating symptoms More than 1.7 million people in the

United States and many others in Europe and Asia currently have Lyme and are unaware or can't find the right treatment. Finally, *Beating Lyme* offers those who struggle with it the guidance to get the help they need. A respected health author and educator, Constance Bean, is an authority on this elusive illness. In 1993 she was diagnosed with Lyme and has spent the past 14 years researching its treatments and diagnoses. In *Beating Lyme*, readers will find comforting, hard-won advice on such topics as what Lyme is and how to recognize the symptoms, what to do after a tick bite, how to protect family and friends, how to get the best treatment, and what to do if insurance won't cover it. Living with long-term Lyme disease is compassionate and thoroughly researched; this is a book that will help both doctors and patients understand and conquer this complex illness.

The Oxford Handbook of the Sociology of Body and Embodiment Natalie Boero, Katherine Mason, 2020-11-13 The Oxford Handbook of the Sociology of Body and Embodiment introduces the sociological research methods and subjects that are key to the growing field of body and embodiment studies. With an emphasis on empirical evidence and diverse lived experiences, this handbook demonstrates how studying the bodily offers unique insights into a range of social norms, institutions, and practices.

When Medicine Goes Awry Juanne Nancarrow Clarke, 2022-01-27 Medical error often results in disability, pain, and suffering, and it is the third leading cause of death in hospitals. Despite its frequency, medical error has been largely invisible to the mainstream public. Within the medical system itself, medical error is often understood as the result of an isolated case of malpractice. *When Medicine Goes Awry* argues that the causes of medical error are not an anomaly but rather the outcome of a number of factors at play, ranging from political to social to economic. *When Medicine Goes Awry* dismisses the common blame perspective associated with medical malpractice, instead asserting that medical error is and will continue to be inevitable given the relentless and expanding processes of medicalization. Shedding light on the ways these forces lead to medicine going awry, the book examines seven well-known cases of medical error. Taking an in-depth look at both patients and medical care providers, Juanne Nancarrow Clarke offers a novel approach to medical error or mishap that applies sociological research and theory to the larger societal forces contributing to a taxing and endemic medical problem.

Departments of Labor, Health and Human Services, and Education, and Related Agencies Appropriations for Fiscal Year 1992: Department of Health and Human Services United States. Congress. Senate. Committee on Appropriations. Subcommittee on Departments of Labor, Health and Human Services, Education, and Related Agencies, 1991

The Silicone Breast Implant Story Marsha L. Vanderford, David H. Smith, 2013-11-05 This volume examines one health issue, breast implants, across a series of contexts often thought to be separate: media coverage, doctor-patient interaction, doctor-doctor professional communication, support group dialogues, public relations campaigns, and more. In so doing, it provides a narrative of how communication shapes the individual perceptions of health, government, and social policy concerning health care. At the core of the silicone breast implant controversy is the need for people to act amid uncertainty about the health risks involved. This need to weigh action in the midst of uncertain risk characterizes a large

number of health issues The attempts of patients physicians drug manufacturers and others to seek and provide both information and influence makes communication central to these issues Consequently the questions explored in this volume will interest a diverse group of readers This audience includes plastic surgeons in particular physicians in general and anyone involved with women s health issues As the medical profession struggles with its identity amid changes in public attitudes government regulations and medical practices this volume s findings concerning media portrayals of doctors and medical devices become even more important Finally this study reveals how interrelated public information and private decisions are and how closely media and interpersonal relationships fit Tracing one medical issue across interpersonal organizational public relations and mediated forums has clearly demonstrated the multiple ways those communication channels overlap and inform one another

In the Kingdom of the Sick Laurie Edwards, 2013-04-09 Thirty years ago Susan Sontag wrote Everyone who is born holds dual citizenship in the kingdom of the well and the kingdom of the sick Sooner or later each of us is obliged at least for a spell to identify ourselves as citizens of that other place Now more than 133 million Americans live with chronic illness accounting for nearly three quarters of all health care dollars and untold pain and disability There has been an alarming rise in illnesses that defy diagnosis through clinical tests or have no known cure Millions of people especially women with illnesses such as irritable bowel syndrome chronic pain and chronic fatigue syndrome face skepticism from physicians and the public alike And people with diseases as varied as cardiovascular disease HIV certain cancers and type 2 diabetes have been accused of causing their preventable illnesses through their lifestyle choices We must balance our faith in medical technology with awareness of the limits of science and confront our throwback beliefs that people who are sick have weaker character than those who are well Through research and patient narratives health writer Laurie Edwards explores patient rights the role of social media in medical advocacy the origins of our attitudes about chronic illness and much more What The Noonday Demon did for people suffering from depression In the Kingdom of the Sick does for those who are chronically ill

Finding Balance Sue Hickey, 2011-03-17 Over 10 million adults in the United States have chronic problems with dizziness and imbalance vestibular problems This compares to 400 000 with multiple sclerosis or 1.5 million with autism for example both chronic illnesses with much greater name recognition Vestibular problems are difficult to diagnose and treat While vestibular problems come with various medical names Meniere s Perilymph Fistula Endolymphatic Hydrops Benign Paroxysmal Positional Vertigo BPPV Mal de Debarquement MdDS and others they present a common challenge for the affected person on how to deal with the dizziness and eventually find balance Finding Balance addresses over a decade of illness from the initial two years searching for a diagnosis through the long years of treatment and finally to the period of rebuilding a life within the confines of what dizziness and vertigo will allow Finding Balance provides a detailed description of symptoms what can trigger them the emotional reactions to symptoms how to recognize their subtle differences and how to track them in order to contribute to your own diagnosis It also emphasizes the

patient's responsibility for healing and eventually how to find the inner balance in life that will be necessary to equitably live with these symptoms over time. And it provides a sense of time of patience. Special Features of Finding Balance Include Call out boxes presenting facts and tips for dealing with the disorder. All of the medical information in the book has been reviewed by vestibular specialists for accuracy. Bulleted lists at the end of chapter with advice to the reader on next steps. *Locked In Locked Out* Shawn Jennings, 2020-01-04. Can there be life after a brainstem stroke? After Dr. Shawn Jennings, a busy family physician, suffered a brainstem stroke on May 13, 1999, he woke from a coma locked inside his body, aware and alert but unable to communicate or move. Once he regained limited movement in his left arm, he began typing his story using one hand and a lot of patience. With unexpected humour and tender honesty, Shawn shares his experiences in his struggle for recovery and acceptance of his life after the stroke. He affirms that even without achieving a full recovery, life is still worth it.

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