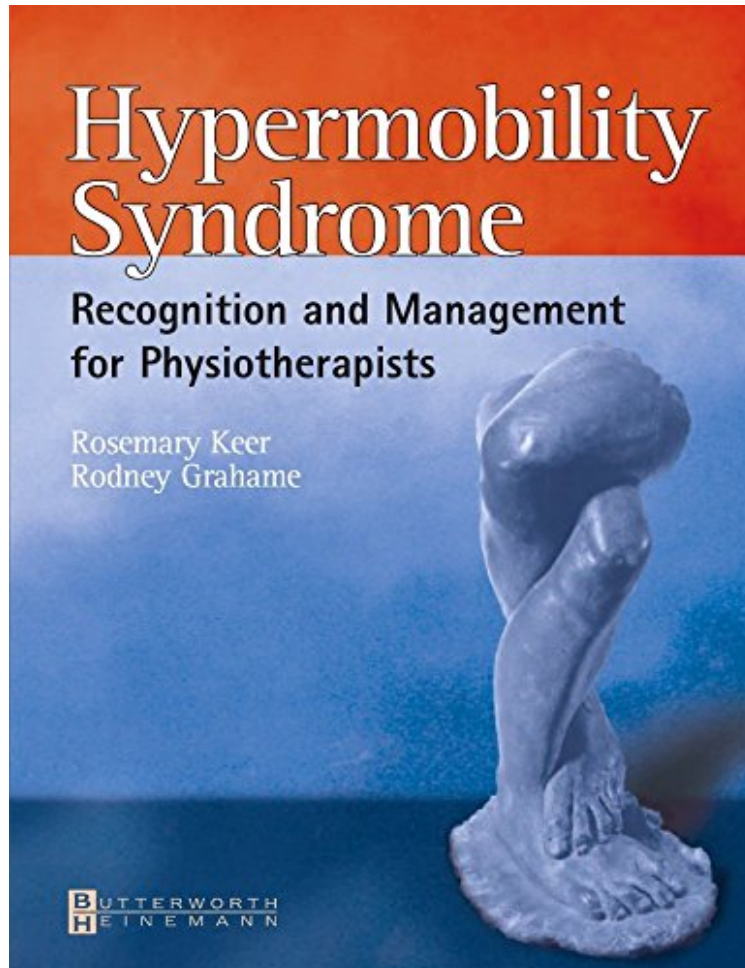


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Hypermobility Syndrome: Diagnosis and Management for Physiotherapists, 1e

*Rosemary J. Keer MSc MCSP MACP, Rodney Grahame CBE MD FRCP FACP
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Rosemary J. Keer MSc MCSP MACP, Rodney Grahame CBE MD FRCP FACP : Hypermobility Syndrome: Diagnosis and Management for Physiotherapists, 1e before purchasing it in order to gauge whether or not it would be worth my time, and all praised Hypermobility Syndrome: Diagnosis and Management for Physiotherapists, 1e:

2 of 2 people found the following review helpful. It's not all in your head...By grandpa jThis is an awesome informative educational book full of helpful management tips for both patient and therapist. It was the first complete medical explanation and validation of my hyper-mobility syndrome. After years of no help from physicians who know little about this syndrome, to orthopedic doctors pretty much telling me there is nothing wrong since symptoms don't match up to common diseases and conditions, this book gave me a sense of relief because even though I knew I wasn't crazy, I could now explain to my doctors exactly what was wrong with me when previously I felt as though they thought it was all in my head.17 of 17 people found the following review helpful. Information for those who need to

knowBy REKSIIf you live in anything less than a large metropolitan area you will need backup to educate your medical and therapy team about the "impossible" symptoms and pain of Hypermobility Syndrome or Ehlers Danlos Syndrome type III. Refer your family practitioner or primary physician, specialists, physical therapists, massage therapist, your family members and your children's teachers [genetic issues]and curious friends to this book. It is the clearest explanation in one place, with an index, of all the myriad issues that folks with our "condition" deal with every minute of every day. Buy your own copy and share it, or buy a separate one for your doctor. Each chapter closes with citations from peer-reviewed medical and therapy journals, an important element to gain respect with folks [including unfortunately physicians and medical specialists] who cannot believe that perfectly fine looking people have major, complicated physical issues. I have referred people to EDS and Hypermobility websites, but this book is more complete and somehow more accessible.2 of 2 people found the following review helpful. it's still good after 10 yearsBy a reviewerI got this to donate to a rehab hospital library. I asked a physical therapist to review it and tell me if it's helpful, and he said yes. Considering it's the only book on the subject that I know of, and very few know anything about it, then it's worth it for basic informatikon

Hypermobility syndrome (HMS) is an inherited connective tissue disorder affecting collagen that is characterized by a collection of features and symptoms. HMS is often encountered in patients, but it is easily overlooked. This practical resource educates readers to its presentation, what to look for, and how best to treat it. Following a brief description of the historical and genetic background of the condition, HMS is described in relation to other connective tissue disorders, such as Ehlers-Danlos syndrome and the Marfan syndrome. Since HMS may occur in childhood, adolescence, adulthood, pregnancy, or old age, each of these stages is covered in the book, with detailed information on the presentation of the condition and its management.Demonstrates how to identify the condition in the clinic.Discusses treatment and management strategies.Provides examples in the form of case studies.Includes details of a comprehensive assessment procedure, which differs depending on the stage of the disorder and the age of the patient.Features contributions by experts in a range of medical fields.

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