

THE 2002 OFFICIAL
PATIENT'S SOURCEBOOK

On

SICKLE CELL
ANEMIA



*A Revised and Updated
Directory for the Internet Age*

JAMES N. FARRAR, M.D.
AND PAUL M. FARRAR, M.D., EDITORS

THE NATIONAL SICKLE CELL ANEMIA FOUNDATION



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The 2002 Official Patients Sourcebook On Sickle Cell Anemia

James N. Parker, Philip M. Parker



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collection or research specific medical options incorporating important data and key concepts about risk factors and symptoms of heart disease *The Official Patient's Sourcebook on Dysphagia* James N. Parker,2002 This book has been created for patients who have decided to make education and research an integral part of the treatment process Although it also gives information useful to doctors caregivers and other health professionals it tells patients where and how to look for information covering virtually all topics related to dysphagia also Hypospadias Dysphagia Syndrome Sideropenic dysphagia from the essentials to the most advanced areas of research The title of this book includes the word official This reflects the fact that the sourcebook draws from public academic government and peer reviewed research Selected readings from various agencies are reproduced to give you some of the latest official information available to date on dysphagia Given patients increasing sophistication in using the Internet abundant references to reliable Internet based resources are provided throughout this sourcebook Where possible guidance is provided on how to obtain free of charge primary research results as well as more detailed information via the Internet E book and electronic versions of this sourcebook are fully interactive with each of the Internet sites mentioned clicking on a hyperlink automatically opens your browser to the site indicated Hard copy users of this sourcebook can type cited Web addresses directly into their browsers to obtain access to the corresponding sites In addition to extensive references accessible via the Internet chapters include glossaries of technical or uncommon terms

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reproduced to give you some of the latest official information available to date on scleroderma Following an introductory chapter the sourcebook is organized into three parts PART I THE ESSENTIALS Chapter 1 The Essentials on Scleroderma Guidelines Chapter 2 Seeking Guidance Chapter 3 Clinical Trials and Scleroderma PART II ADDITIONAL RESOURCES AND ADVANCED MATERIAL Chapter 4 Studies on Scleroderma Chapter 5 Patents on Scleroderma Chapter 6 Books on Scleroderma Chapter 7 Multimedia on Scleroderma Chapter 8 Periodicals and News on Scleroderma Chapter 9 Physician Guidelines and Databases Chapter 10 Dissertations on Scleroderma PART III APPENDICES Appendix A Researching Your Medications Appendix B Researching Alternative Medicine Appendix C Researching Nutrition Appendix D Finding Medical Libraries Appendix E Your Rights and Insurance ONLINE GLOSSARIES SCLERODERMA GLOSSARY INDEX Related topics include Familial Progressive Systemic Sclerosis Morphea Progressive systemic sclerosis Systemic Sclerosis

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In a world driven by information and connectivity, the power of words has been evident than ever. They have the capacity to inspire, provoke, and ignite change. Such is the essence of the book **The 2002 Official Patients Sourcebook On Sickle Cell Anemia**, a literary masterpiece that delves deep into the significance of words and their impact on our lives. Written by a renowned author, this captivating work takes readers on a transformative journey, unraveling the secrets and potential behind every word. In this review, we shall explore the book's key themes, examine its writing style, and analyze its overall effect on readers.

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