

Scleroderma Coping Strategies

Thank you enormously much for downloading **Scleroderma Coping Strategies**. Most likely you have knowledge that, people have seen numerous times for their favorite books subsequent to this Scleroderma Coping Strategies, but stop stirring in harmful downloads.

Rather than enjoying a good book subsequently a cup of coffee in the afternoon, otherwise they juggled gone some harmful virus inside their computer. **Scleroderma Coping Strategies** is handy in our digital library an online permission to it is set as public suitably you can download it instantly. Our digital library saves in multipart countries, allowing you to acquire the most less latency period to download any of our books subsequently this one. Merely said, the Scleroderma Coping Strategies is universally compatible next any devices to read.

Social Support and Psychiatric Disorder T. S. Brugha 1995-08-24 A critical evaluation of research into the impact of social support on mental health with implications for mental health management and intervention.

Multiple Sclerosis Rosalind C. Kalb, MD 2005-09-01 Because most people who are diagnosed with MS are between the ages of 15-55 years, the disease has a significant impact, not only on the individual with the disease, but also on the family members and loved ones whose lives are interwoven with them. Families experiencing multiple sclerosis often find themselves in uncharted territory. Prior patterns of interacting with each other may no longer work; roles often shift with dramatic emotional impact; guilt, anger, sadness, and a sense of burden may create a barrier to intimacy, joy, growth, and family unity. There are ways out of this debilitating situation, and many families have restored their balance, humor, productivity and family solidarity. Multiple Sclerosis: A Guide for Families can lead the way for your family to strengthen its coping skills and to receive targeted information about the disease and its ramifications. This complete yet highly readable guide for families living with multiple sclerosis addresses these issues and more. Families must learn how to strike a reasonable balance between the interests and needs of its members, and to accommodate the limitations imposed by MS without allowing them to impact every aspect of family life. This is easier said than done, and extensive discussions consider how the basic goals of living with this chronic disease can be met.

Multiple Sclerosis: Coping with Complications Barry Farr, MD 2016-06-06 There's no miracle cure for multiple sclerosis. But there are ways to reduce its negative impact. What better source than a physician who battled MS as a patient for 24 years? Unlike the "cure for MS" books published for decades (none yet deemed valid by the scientific community), this one aims to help patients deal with the reality of chronic complications of MS, sharing new strategies. Why "new" ones? Older strategies didn't work so well. For example, using long-recommended conventional, medical wisdom, MS patients still got more urinary tract infections. Dr. Barry Farr tells how he avoided urinary tract infection for 20 years despite being high risk. When conventional approaches didn't relieve severe rib fracture pain, he tried something different. The pain disappeared. After 3 aspiration pneumonias during 7 months, he made a change and had none over 7 years. Being new, such strategies won't be found in other books for MS patients.

Multiple Sclerosis Cynthia L. Sullivan 2000

Supporting People with Scleroderma Katherine Milette 2018 "Reducing the burden of chronic diseases has been identified as a health priority by National and International health organizations around the world, and community-based supportive interventions have been proposed as one possible way of achieving this goal. In general, receiving social support has been found to impact health and benefit many individuals with chronic disease who are engaged in disease management efforts. Scleroderma is a rare and complex rheumatic, autoimmune disease that affects the connective tissues and people who receive a diagnosis can experience a wide variety of challenges in their daily life. Because of the unique challenges that rare disease patients can experience related to the disease impact and navigating health care, people diagnosed with scleroderma may benefit from a variety of supportive interventions to help them better cope. However, no definitive conclusions can be drawn at this time about the unique support needs of people with scleroderma since qualitative investigations directly looking at coping and social support for scleroderma are limited. In the current dissertation research, two distinct focus group studies were conducted using a social constructionist framework. The purpose of Study 1 was to explore differing perspectives regarding the challenges experienced and strategies used while coping with scleroderma, through focus group discussions with people who have a diagnosis of scleroderma (4 focus groups, n=34) and health care professionals working with patients with scleroderma (1 focus group, n=8). The purpose of Study 2 was to perform an exploratory, follow-up investigation to Study 1 to deepen our understanding of patient perspectives about the influence of social support on the ability to cope with scleroderma (4 focus groups, N=19). Findings from Study 1 provided a deeper understanding of the unique challenges experienced by people with scleroderma while actively coping with the disease, including problems accessing information (e.g., from healthcare professionals), dealing with negative emotions (e.g., related to misunderstandings with loved ones), and accessing resources (e.g., effective scleroderma treatments). Study 1 also provided knowledge about helpful strategies to address those coping challenges (e.g., advocating for your needs and learning to problem solve). Next, findings from Study 2 showed that to better cope with their disease, individuals with scleroderma often relied on close social relationships to provide different types of support, such as emotional, informational, and instrumental support. In addition, different relational factors were identified that either enhanced or impeded the ability of people with scleroderma to rely on social support to cope with their disease, including different issues related to communication style, active engagement, and complementarity. Implications of the dissertation findings and future direction for research will be discussed." --

The Information Needs and Information-seeking Patterns of Women Coping with and Adjusting to Multiple Sclerosis Lynda Baker 1994 Multiple sclerosis (MS) is a chronic neurological disease with an unpredictable course and no known cause or cure. Coping strategies of people with MS may therefore yield insights into information-seeking patterns. This study of information preferences was based on Miller's (1980) theory of information-seeking, which states that some people (monitors) cope by actively seeking information, while others (blunters) reject information. In this study, 251 female MS clinic patients were classified through the Miller Behavioral Style Scale (MBSS) as either monitors or blunters and by the length of time since diagnosis. It was hypothesized that monitors (in contrast to blunters) would be more interested in information regardless of phase of the disease, format or specificity of information. In Part One of this study, 160 women completed a mail survey designed to assess their interest in and desire for information on 29 topics relevant to MS, as well as the preferred formats of this information. The results revealed that monitors expressed preferences for more information than did blunters in earlier phases of the disease, while blunters became interested and wanted information in later phases of the disease. Videos, magazines, pamphlets and books were the most preferred formats for information on multiple sclerosis. Physicians and spouses were the most common human resources, though preferences varied with topic and format. In Part Two, an additional 93 women with MS, also classified as monitors or

blunters were asked to assess a general or a specific pamphlet on one of two topics (fatigue or treatment of acute attacks). More monitors than blunters rated the pamphlet they had received as relevant, regardless of the specificity of information. This study contributes to the development of a methodology for studying information-seeking patterns and suggests that information providers (e.g., MS societies, health care professionals and librarians) may need to consider general orientation to information as well as phase of the disease in order to meet the information needs of women with MS.

Families Affected by Multiple Sclerosis Rosalind C. Kalb 1995

Occupational Therapy Practice and Research with Persons with Multiple Sclerosis Marcia Finlayson 2013-04-15 Discover strategies to enhance quality of life and promote social and community participation for people with MS! Occupational Therapy Practice and Research with Persons with Multiple Sclerosis will familiarize you with the complex issues experienced by people who have multiple sclerosis, suggesting ways to enhance your practice or research with this population. This vital resource fills a void in the scarce literature on occupational therapy and multiple sclerosis, providing you with a unique single-source reference on the subject. This book compiles the work and contributions of experts from Europe, Australia, Canada, and the United States—from a variety of fields, including occupational therapy, medicine, physical therapy, and psychology. This thought-provoking book offers new perspectives on potential assessment and intervention ideas and provides information that could be used for broader program planning. This extensive resource will give you a deeper appreciation of the MS disease process and its influence on everyday living for persons with MS and their families. In this collection, you will learn more about: the health-related service needs of older adults with MS the range of fatigue assessment tools that are available for clinical and research applications the effect of wheelchair use on quality of life the implications of tremor on everyday activities the development and use of Lifestyle Management Programs© coping processes used by women with MS as they age the symptom and functional limitation profiles experienced by people with MS that lead to referrals to occupational therapy Occupational Therapy Practice and Research with Persons with Multiple Sclerosis contains charts, figures, graphs, and bibliographies to augment the research and studies found in this book. Also provided are contributions by Dr. Nicholas G. LaRocca—a well-known MS researcher—and Dr. Carol A. Gaetjens—an educator with MS. Occupational therapy students, clinicians, and researchers working with individuals and families who are affected by MS will find this book an important resource in their profession.

Women and Multiple Sclerosis Duane O'Mahony 2010 Multiple sclerosis (MS) is the most common chronic neurological disease of the central nervous system that affects over 2.4 million people world-wide. Disease onset usually occurs in young adults, and it is more common in females. The exact cause of MS is unknown, but an environmental factor is thought to trigger MS in genetically predisposed persons. This book focuses on MS and its effects on women with regards to quality of life, reproduction and bone health. Fatigue, as one of the most common symptoms of multiple sclerosis (MS), is also examined in this book - in particular, the relational coping strategies used by mothers with MS to manage their fatigue while parenting young children. A discussion on the sex related differences concerning MS presentation, signs and symptoms, therapeutics, response to treatment and quality of life among patient populations is included, as well as the relationship between self-efficacy, self-esteem, hope and disability in women with multiple sclerosis.

Miracles on My Watch Bianca Podesta 2016-02-26 A gifted storyteller with long experience in ministry, Bianca Podesta has a heart that recognizes everyday holiness. Here you will read how ordinary events and people have been transformed into light-filled windows that reflect grace and Presence. A just-baptized baby giggles, and an entire congregation is drawn into a new awareness of who they are. A dying man is given several months for catching up with his soul. From time to time you will be surprised by a story that cannot be easily explained. It will take your breath away, more exactly give it back. Not always pretty or easy, these carefully nuanced and honest tales will challenge you to become more aware of the miracles in your own life. Rev. Tilda Norberg, Founder, Gestalt Pastoral Care. Author, *Consenting to Grace* A journalist for twenty years, I am always on the lookout for a good story. Bianca Podestas *Miracles on My Watch*, derived from her work in churches as well as from widely different settings, is a treasure trove. Thought provoking and inspiring. I couldn't put it down. Valerie Zehl, Award-Winning Columnist, Gannett

Navigating Life with Amyotrophic Lateral Sclerosis Mark B. Bromberg 2017 *Navigating Life with Amyotrophic Lateral Sclerosis* provides accessible, comprehensive, and up-to-date information about the challenges patients, family members, and caregivers face when confronted by ALS, a disease that affects approximately 5,600 Americans every year, with as many as 30,000 people managing the disease at any given time. ALS is a difficult disease for the patient and is also challenging for the caregiver and family as there are many questions, issues relating to care, and problems to manage. This guide covers all aspects of managing ALS, from the onset of symptoms, diagnosis, treatments, and coping strategies, to the use of home health care or hospice, and new research in the field. The book also sheds lights on difficult topics, such as end-of-life care and managing legal affairs. *Navigating Life with Amyotrophic Lateral Sclerosis* is unique because it covers two perspectives: one author is a neurologist with 30 years of experience treating ALS patients, and the other author experienced first-hand the issues in providing care for a parent with ALS. Formatted in a question-and-answer style, peppered throughout with patient stories, and with sections devoted to family members and caregivers, this compassionate resource provides guidance to those seeking to understand how to live with this disease.

Benefits of Multiple Sclerosis and Quality of Life. The Mediating Role of Coping Strategies Jose Luis Gonzu00e1lez-Castro 2017 Multiple Sclerosis (MS) is an immune mediated process affecting a person's central nervous system. This illness has a significant impact in social relationships, autonomy, or psychological wellbeing reducing the quality of life of those who suffer the illness. Nevertheless, studies have shown that people with MS may also find positive aspects, or benefits, from the illness. These relate to survival strategies based on the search for meaning aimed towards improving Quality of life (QoL) defined as a subjective and objective feeling of general wellbeing or satisfaction regarding important aspects of one's life. The way people use cognitive and behavioral strategies and mechanisms to try to master, minimize or adapt to stressful situations is termed coping. The aim of this study was to analyze the mediating role of coping strategies in the relationship between perceived benefits of MS and Quality of Life. 250 participants took part in the study with a mean age of 41.74 years (sd = 10.34). Participants answered the following measures: Functional Assessment of Multiple Sclerosis (FAMS) (Cella et al., 1996). The Brief COPE-28 (Carver, 1997: problem based, emotional based and other coping strategies). The Psychosocial impact of multiple sclerosis (Mohr et al., 1999; only the benefits of MS subscale). SPSS v.24 and Process v. 3 were used in the analysis. Results from the mediation analysis showed that benefits had no direct effect on QoL. Benefits did have a positive significant effect on emotional support, planning and self-distraction, and marginal and positive on acceptance. Acceptance and emotional support had a positive effect on QoL while denial, planning and self-distraction had a negative effect on QoL. Indirect effects of the mediation show that denial had no significant mediating role, and that acceptance and emotional support positively mediated between benefits and QoL improving the latter. Planning and self-distraction negatively mediated the relationship between benefits and QoL. This study shows that the idea that positive growth from MS can improve QoL is only found when emotional, and not problem, based, coping mechanisms, are used. These results render support for flexibility theories regarding the adaptive nature of coping strategies or mechanisms.

The Autoimmune Connection: Essential Information for Women on Diagnosis, Treatment, and Getting On With Your Life Rita Baron-

Faust 2016-04-08 The most up-to-date book on the subject gives women all the essential information on diagnosis and treatment of autoimmune medical conditions The Autoimmune Connection explains the links between autoimmune diseases and offers up-to-date information on diagnosis, treatments, and risks for women with one or more autoimmune disease, such as lupus, rheumatoid arthritis, or Crohn's disease. You will learn how to sort out vague and seemingly unrelated early signs and symptoms; which diagnostic tests they may need and what the results can mean; how autoimmune diseases and treatments can affect you at different stages of life; what new treatments and therapies are on the horizon; and where to find the proper specialist and how to navigate the health-care system NEW: Major breakthroughs in the treatment of rheumatoid arthritis, thyroid disease, diabetes, Celiac, multiple sclerosis NEW: Potential environmental triggers for autoimmunity, such as cigarette smoke, crystalline silica, asbestos, mercury, lead, viruses and other microbes, pesticides, pollutants, solvents, and petroleum byproducts NEW: Current drugs on the market and recently discovered side effects and long-term complications connected to existing drug therapies NEW: Healthcare law reform provisions and key changes to Medicare and Medicaid My Story Amelia Davis 2004-03-01 In a series of dramatic essays and photographs by the renowned San Francisco-based photographer Amelia Davis, My Story is an evocative description of what it is like to live with multiple sclerosis (MS), a disease that affects about 350,000 Americans and whose cause is still not entirely known. The essays and accompanying photographs in this highly engaging, beautifully illustrated book poignantly portray the lives of thirty-two men and women from the ages of seventeen to seventy and of various ethnicities, and socio-economic backgrounds who share the challenge of living with MS. Some, like Amelia, use no mobility aids, while others use canes, wheelchairs, or electric scooters. All have had to face the challenges and limitations that MS has imposed upon their lives, and each has devised unique and often creative coping strategies. Accompanying each essay are commentaries by family members and friends that express their own personal feelings and experiences of living with this disease. Here are the inspirational stories of women, men, and children who live with this disease. Many have children; one or two are currently expecting them. Treatments and therapies for slowing the progression of the disease are revealed and shared, from the latest advances in prescription medication to alternative methods of coping, including yoga, exercise and competitive sports, creative activities such as writing and art, and even community activism. A wide range of accompanying stories by spouses, children, and other loved ones depict the ups and downs of living and caring for someone who has MS, from the moment of first diagnosis to dealing with its ongoing challenges. All are strong reminders of the selflessness of the human spirit, and its ability to nurture and remain strong under even adverse circumstances. Highly motivating and deeply inspirational, My Story will be welcomed by anyone who lives with or shares the life of someone who has MS.

Voices of Scleroderma International Scleroderma Network 2004-08 Contains anecdotes and experiences of people with scleroderma or with loved ones who have it.

Understanding the Dynamics of Coping, Psychological Well-being and Health-related Quality of Life in Persons with Relapsing-remitting Multiple Sclerosis Kristine Kelly McGuinn 2006

Stress and Multiple Sclerosis Judith Tschannen Reppel 1986

Interpersonal and Disease-related Coping Strategies Among Patients with Multiple Sclerosis Rodrigo Da Silva Dias 2002

Advances in Integrative Dermatology Katlein França 2019-01-22 An authoritative overview of contemporary approaches to integrative skin health The popularity of integrative medical treatment of skin disorders has increased significantly in recent years—requiring practicing clinicians to keep pace with continuing developments in dermatological research and methods. *Advances in Integrative Dermatology* offers a comprehensive survey of this dynamic field, providing up-to-date information on both preventative and therapeutic approaches to skin health. Combining clinical medicine with complementary treatment plans, integrative dermatology provides an innovative perspective to individual patient care. This essential text explores new research in areas including the effects of stress and pollution on the skin, the importance of high-quality sleep, complementary methods of averting skin conditions, and more. Recognizing the impact skin disorders have on physiological, psychological, and emotional health, editors Katlein França and Torello Lotti illustrate key components of inclusive skin health strategies, such as therapeutic diets and nutritional supplements, topical botanicals, and other complementary therapies. Filling a significant gap in current literature on integrative dermatology, this valuable resource: Answers the common questions asked by patients in real-world dermatology practices Addresses pervasive misconceptions of integrative dermatological methods and principles with evidence-based review of current practices Examines contemporary research in the diagnosis and treatment of dermatological disorders Presents comprehensive treatment options for a wide range of conditions such as rosacea, melanoma, and psoriasis *Advances in Integrative Dermatology* is an indispensable volume for physicians seeking to incorporate holistic techniques into their practices, expand their knowledge of integrative medicine, and provide the best possible care for their patients.

The Relationship Between Coping Strategies and Adjustment to Multiple Sclerosis Dayna M. Davis 1998

Factors Contributing to the Quality of Life Among Individuals with Multiple Sclerosis Margaret Kathryn Howlett 2010

Coping with Multiple Sclerosis: Coping Strategies, Personality, and Cognitive Appraisals as Predictors of Adjustment Among

Multiple Sclerosis Patients Holly McCartney Chalk 2007 Correlational results indicated that coping and cognitive appraisals were systematically related to adjustment, while physical disability was not. This implies that many of the variables which relate to adjustment outcomes among MS patients are controllable, suggesting that practitioners should target these cognitive and behavioral variables to positively affect adjustment to MS.

Practical Management of Systemic Sclerosis in Clinical Practice Marco Matucci-Cerinic 2020-12-18 This book provides a practical guide for managing a variety of problems encountered by the clinician in managing patients with systemic sclerosis. Chapters take a problem-orientated approach to help the reader cut through potential barriers that can arise when working with different medical specialities. Management strategies for a broad range of conditions, including pericardial and pleural effusion, sicca syndrome, calcinosis and watermelon stomach, are presented. *Practical Management of Systemic Sclerosis in Clinical Practice* describes a range of problems and clinical items encountered by a variety of medical professionals who encounter these patients. It is a valuable resource for rheumatologists, immunologists, specialist nurses and primary care professionals.

Multiple Sclerosis T. Jock Murray, MD 2017-10-28 An MS diagnosis is not an endpoint. This is the place to begin your education and manage your future. Now in its fifth edition, *Multiple Sclerosis: A Guide for the Newly Diagnosed* is the best-selling guide to multiple sclerosis (MS) for newly diagnosed patients and their families. Written in plain language by renowned MS expert physician T. Jock Murray, this book guides patients at any stage who want to know more about the disease, its potential impact on your life, and the medical treatments now available for managing it successfully. The fifth edition is updated to reflect recent advances in understanding the underlying mechanisms and disease progression, the diagnostic criteria and classification of MS, and new drugs and therapies. This book opens the window on an MS diagnosis and provides answers to the multitude of questions patients will have so they can take charge and make informed decisions about their health and treatment.

Summary of Gabor Mate's When the Body Says No Milkyway Media 2022-04-28 Please note: This is a companion version & not the original

book. Book Preview: #1 Raynaud's phenomenon is a condition in which the small arteries supplying the fingers are narrowed, depriving the tissues of oxygen. It can lead to gangrene, and in some cases, scleroderma. #2 The idea that people's emotional coping style can be a factor in scleroderma or other chronic conditions is anathema to some physicians. #3 The more specialized doctors become, the less they understand the human being in whom that part or organ resides. The people I interviewed for this book reported that neither their specialists nor their family doctors had ever invited them to explore the personal, subjective content of their lives. #4 Until the advent of modern medical technology and scientific pharmacology, physicians relied on placebo effects to treat their patients. Today, we have lost the ability to treat our patients based on their confidence in their inner ability to heal.

Multiple Sclerosis Dr. Nancy J. Holland, RN, EdD 2007-02-15 The diagnosis of multiple sclerosis (MS) poses potential concerns related to all aspects of life and plans for the future. Family members and other loved ones are similarly concerned, and everyone involved struggles to make sense of life with this permanent intruder. One of the first responses is usually an active search for information about the disease itself and its potential long-term effects. Chapters discuss the nature of MS, its management, and guidelines for dealing with all aspects of the disease and its impact on your life. A chapter on services available from the National Multiple Sclerosis Society, a glossary, a list of resources, and additional reading suggestions make this the place to begin your education about MS. With education and proper care, most people diagnosed with multiple sclerosis will lead full and productive lives. *Multiple Sclerosis: A Guide for the Newly Diagnosed, Third Edition* is an essential resource for everything you need to know about MS, and includes new or updated sections on: The most current medical treatments for the management of MS Complementary and alternative medicine and MS Financial and life planning Children with MS Updated diagnostic criteria

Coping Profiles and Health Outcomes Among Individuals with Systemic Sclerosis 2016 Systemic sclerosis (SSc) is a severe rheumatic disease with extensive implications for quality of life. Physically, the disease causes fatigue, chronic pain, and functional disability. Mentally, the disease is associated with body image distress, anxiety, and depression. Coping represents a mechanism by which individuals living with a chronic illness can exercise control over the situation and adapt more successfully. The primary aims of this study were to (1) identify coping-based profile groups, and (2) compare the coping-based profile groups on physical health and mental health outcomes. Participants included 94 adults with confirmed diagnoses of SSc. Data were drawn from a larger study, and were cross-sectional and archival in nature. A latent profile analysis was conducted to create profiles derived from participants' raw scores on coping subscales including problem-focused (PF), wishful thinking (WT), seeking social support (SS), avoidance (AV), self-blame (SB), blaming others (BO), counting one's blessings (CYB), and religiosity (RG). A three-profile solution was supported statistically and substantively. Low Copers (n = 7) were characterized by a below-average use of coping across all eight subscales. Conversely, High Copers (n = 23) exhibited an above-average use of coping across all eight subscales. Mixed Copers (n = 64) demonstrated infrequent use of BS, SS, AV, BO coping strategies and frequent use of the PF, WT, CYB, RG coping strategies. ANCOVAs were conducted to examine whether the profile groups differed in physical health, and whether the groups differed in psychological health. A significant effect was found for psychological health. Post hoc comparisons revealed significant differences in psychological distress between Low Copers and High Copers, and Mixed Copers and High Copers, with High Copers exhibiting significantly greater levels of distress when compared with the other two groups. This finding raises an interesting question about whether engaging in frequent use of a variety of coping strategies is related positively to one's psychological health, as has been assumed, or whether it may reflect distress. It is recommended that future research on coping with SSc employs an LPA approach, but also collects data at multiple time points to explore the causal relationship of coping and health.

Coping Strategies of Multiple Sclerosis Patients and Their Families ... Donna B. Stauber 1993

The Relationship Between Neuropsychological Test Performance and Coping Strategies in Individuals with Multiple Sclerosis Vickie M. Jean 1996

Occupational and Physical Therapy for Children with Rheumatic Diseases Gay Kuchta 2022-02-27 'This volume comes at an important time in the development of the discipline, and reflects the knowledge gained by the authors through decades of hands-on experience of treating children and adolescents with the entire spectrum of rheumatic diseases. It emphasizes the integration of the expertise of all health care providers in an interactive team' - Ross Petty in the Foreword. This highly practical handbook is easy to read and refer to on a daily basis. The wide-ranging style encourages a multidisciplinary, team-based approach to provide continuity of care, and the handy binding and layout is designed to aid quick access of core information. Featuring over 230 full-colour images and reproducible resources for patients and carers, this guide is a vital resource for Allied Health Professionals, especially those working with children.

Psychological Functioning After a Recent Diagnosis of Multiple Sclerosis Shirley Isobel Anderson 2005

Scleroderma Coping Strategies B. Bianca Podesta 2011-02-15 A kind of "how-to" guide for dealing with scleroderma.

Disorders of Emotion in Neurologic Disease 2021-08-11 Disorders of Emotion in Neurologic Disease, Volume 183 in the Handbook of Clinical Neurology Series, informs clinicians on which neurologic diseases are likely to have a secondary effect on emotion, what to look for in diagnosis, and best practices for treatment. The book begins with an understanding of the neurological basis for emotions in order to better understand what goes awry in neurological disease. It then discusses specific neurologic diseases and disorders affecting emotion. Reviews the neurology of emotions Specifies neurologic diseases that affect emotional expression Informs clinicians on how to diagnose, along with best practices for treatment Includes coverage of stroke, dementia, epilepsy, Huntington's, Parkinson's, TBI, and more

Relationships Between Personality, Coping Strategies, Prosocial Behavior, Empathy, and Quality of Life Among Persons with Multiple Sclerosis and Their Caregivers Donna LeAnn Plemons 2004

Clinical Management in Psychodermatology Wolfgang Harth 2008-11-14 Psychocutaneous Medicine offers an overview of diseases in psychosomatic dermatology and creates a bridge between cutaneous and emotional disorders using extraordinary illustrations and clinical images of psychosomatic dermatology. It covers both common and rare diseases and helps doctors and psychologists recognize and deal with psychosocial features in dermatology and venerology. This superbly illustrated clinical atlas with concise text passages follows the American diagnosis classification DSM-V and current evidence-based guidelines. It allows rapid recognition of masked emotional disorders and thus administration of the most effective and efficient treatment as early as possible. Hone your diagnostic vision for psychosomatic disorders. Treat your patients efficiently and effectively. Psychocutaneous Medicine is a picture atlas and textbook that is indispensable for dermatologists, psychologists, pediatricians and general practitioners.

Scleroderma. An Issue of Rheumatic Disease Clinics Maureen D. Mayes, MD, MPH 2015-08-03 Scleroderma, or systemic sclerosis, is a chronic connective tissue disease generally classified as one of the autoimmune rheumatic diseases. Hardening of the skin is one of the most visible manifestations of the disease. It's estimated that about 300,000 Americans have scleroderma. About one third of those people have the systemic form of scleroderma. Since scleroderma presents with symptoms similar to other autoimmune diseases, diagnosis is difficult. This issue will cover: The Genetic Basis of SSc: Genetics, Epigenetics, Mechanisms of Pathogenesis - linking fibrosis, vasculopathy and immune dysregulation, The Role of Autoantibodies in Diagnosis and Prognosis/survival; Managing Raynaud's phenomenon and ischemic ulcers,

managing SSc Lung Disease, Monitoring for and Managing Pulmonary Arterial Hypertension, and many more!

Rehabilitation of the Hand and Upper Extremity, E-Book Terri M. Skirven 2020-01-14 Long recognized as an essential reference for therapists and surgeons treating the hand and the upper extremity, *Rehabilitation of the Hand and Upper Extremity* helps you return your patients to optimal function of the hand, wrist, elbow, arm, and shoulder. Leading hand surgeons and hand therapists detail the pathophysiology, diagnosis, and management of virtually any disorder you're likely to see, with a focus on evidence-based and efficient patient care.

Extensively referenced and abundantly illustrated, the 7th Edition of this reference is a "must read" for surgeons interested in the upper extremity, hand therapists from physical therapy or occupational therapy backgrounds, anyone preparing for the CHT examination, and all hand therapy clinics. Offers comprehensive coverage of all aspects of hand and upper extremity disorders, forming a complete picture for all members of the hand team—surgeons and therapists alike. Provides multidisciplinary, global guidance from a Who's Who list of hand surgery and hand therapy editors and contributors. Includes many features new to this edition: considerations for pediatric therapy; a surgical management focus on the most commonly used techniques; new timing of therapeutic interventions relative to healing characteristics; and in-print references wherever possible. Features more than a dozen new chapters covering Platelet-Rich Protein Injections, Restoration of Function After Adult Brachial Plexus Injury, Acute Management of Upper Extremity Amputation, Medical Management for Pain, Proprioception in Hand Rehabilitation, Graded Motor Imagery, and more. Provides access to an extensive video library that covers common nerve injuries, hand and upper extremity transplantation, surgical and therapy management, and much more. Helps you keep up with the latest advances in arthroscopy, imaging, vascular disorders, tendon transfers, fingertip injuries, mobilization techniques, traumatic brachial plexus injuries, and pain management—all clearly depicted with full-color illustrations and photographs.

Relationships of Positive and Negative Affect to Coping and Functional Outcomes in Systemic Sclerosis Ingunn Hansdottir 2002 A vast literature has shown that chronic illness has a negative impact on emotional adjustment. To date conceptual models guiding this research have focused on negative emotions and psychopathology. However, most people with chronic illness do not meet criteria for a diagnosable disorder, suggesting that our focus should be on normal emotional processes instead. Based on the two-dimensional model of emotion proposed by Watson and Tellegen (1985), which suggests two distinct factors of emotion, positive affect (PA) and negative affect (NA), the present study extended previous research by examining the role of both PA and NA in adjustment to chronic illness. It was hypothesized that PA and NA would represent distinct aspects of emotional well-being with separate influences on factors related to adjustment to chronic illness. Participants were 96 patients with confirmed diagnoses of Systemic Sclerosis, a severe and chronic rheumatic disease. As part of a larger longitudinal study, participants completed self-report measures assessing PA and NA (Positive and Negative Affective Schedule; PANAS), coping (Revised Ways of Coping Checklist), and functional outcomes (pain and disability measured with the Health Assessment Questionnaire). Confirmatory factor analysis indicated that PA and NA constituted separate, negatively correlated factors. The utility of assessing both PA and NA was further supported by differential relationships of PA and NA to coping and functional outcomes. Hierarchical regression analysis showed that relations among coping, affect and functional outcomes were consistent with a mediational model, in which affect was found to mediate the association between coping and functional outcomes after controlling for disease severity. Adaptive coping was associated with higher PA, and PA was related to better functional outcomes, i.e. lower levels of pain and disability. Maladaptive coping was associated with higher NA, and NA was related to higher levels of pain. Using a latent structural equation modeling approach, the mediational model yielded a marginally acceptable fit to the data. Implications suggest the importance of assessing both PA and NA in order to fully understand the process of adjusting to chronic illness, as well as to identify mechanisms that lead to improved functional outcomes in systemic sclerosis.

Scleroderma John Varga 2016-11-07 Comprised of the authoritative work of international experts, this fully-updated second edition of *Scleroderma* builds upon the well-regarded approach in the first edition to provide integrated, concise, and up-to-date synthesis of current concepts of pathogenesis and modern approaches to management of systemic sclerosis (scleroderma). With a multidisciplinary approach to comprehensive care, this book is easily accessible for health care professionals in many fields. The new edition includes extensive updated material based on major developments in the field, with new chapters on personalized medicine, cancer complications, global perspectives on scleroderma, and more. It presents a succinct and thoughtful synthesis of current pathomechanistic concepts, providing a valuable reference tool for basic and translational investigators working in the field. *Scleroderma: From Pathogenesis to Comprehensive Management* serves as an essential, all-inclusive resource for rheumatologists, pulmonologists, cardiologists, gastroenterologists, nephrologists and all those involved in the care of scleroderma patients.

Psychological Issues in Amyotrophic Lateral Sclerosis Francesco Pagnini 2016-03-02 Amyotrophic lateral sclerosis is a fatal and progressive disease, characterized by progressive muscles weakness, with consequent loss of physical capacities. Patients become relentlessly immobile and, in the late stages of the disease, develop a "locked-in" state in which only residual muscular movement is possible, but the intellect and the personality usually remain unimpaired. At now, there is no cure for ALS. The psychological impact of the disease is huge, on both patients and caregivers. Aim of the present Research Topic is to collect new evidence about quality of life, depression, anxiety, pain, spiritual and existential issues, hope and hopelessness in the ALS field, with attention to both patients and their caregivers. Emphasis will be provided to the investigation of psychological support and the possible role of psychologists in this challenging field. Keywords: Amyotrophic Lateral Sclerosis; Health Psychology; Clinical Psychology, Motor Neuron Disorder; Quality of Life. Subtopics: The subtopics to be covered in the Research Topic include, but not limited to: 1. Assessment of psychological variables in ALS 2. Quality of life during the course of the illness 3. Impact of technological assistance to illness (wheelchairs, NIV...) 4. Interfaces among biological, psychosocial, and social factors 5. Psychological and psychotherapeutic interventions 6. Couple and family relationships 7. Research methodology, measurement and statistics 8. Cultural and social features of ALS 9. Professional issues, including training and supervision 10. Implications of research findings for health-related policy