

THE COSTS OF CARING

Families with disabled children

Sally Baldwin

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**Shane Kavanagh,Louis Opit,Politeia
(Association : London, England)**

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EBOOK: Young People Living With Cancer Anne Grinyer, 2007-02-16 This book is shouting out to every person who is involved with young people's cancer. I hope they listen. Macmillan Cancer Support. This is an absolute must read for all those who care for young people with cancer including the patients themselves, parents, doctors, nurses, psychologists and all of the caring professions. Professor Tim Eden, Christie NHS Trust, Manchester, UK. I particularly liked the layout of the book and the initial chapter is called 'setting the scene'. It does just that and informs the reader of services available and leads the reader into the remaining chapters about diagnosis through to sexuality and finally implication for policy and practice. I would recommend this book for everyone involved with cancer care. As a nurse, I found it excellent. As a mum of two young people, I found it very emotional. Jan Stevens, Nurse Coordinator, Hospice at Home. This book should be compulsory reading for all healthcare professionals who work with young people with cancer and especially for all professionals who may at some stage meet a solitary young person with cancer. Sue Morgan, St James's University Hospital, Leeds, UK. What issues are of most concern to young adults living with cancer? How can a supportive care setting be established? How can we offer the optimum age appropriate care? This book uses original data gathered from in-depth research to present an account of what it is like to be an adolescent or young adult living with cancer. These first-hand accounts contribute to the insight necessary to the provision of age appropriate quality care. The implications of these research findings for policy and practice are also related to NICE Guidance 2005. Though primarily based on a thematic analysis of interview data, the book also provides contextual and statistical information on the current incidence of cancer in young adults. *Young People Living with Cancer* is essential reading for health professionals engaged in the care of young adults with cancer, support workers in the health services, young adults with cancer, their families, academics and students. The text offers a contribution to policy and practice that may enhance compliance and consequently improve outcomes.

EBOOK: Carers Perceived Julia Twigg, Karl Atkin, 1994-01-16 Carers are the bedrock of community care and yet our understanding of how they do and do not fit into the care system is limited. Concern is often expressed about the need to support carers but the best way to do this is not always clear. This book breaks new ground in exploring the reality of how service providers, the doctors, social workers and community nurses respond to carers. It looks at which carers get help and why, analyzing how age, relationship, class and gender structure the responses of service providers and carers. It examines the moral and policy issues posed by trying to incorporate carers' interests into service provision. What would services look like if they took the needs of carers seriously? How far can they afford to do so? Is this only achieved at the expense of disabled people? What is the proper relationship between carers and services? Carers pose in acute form many of the central dilemmas of social welfare and the account presented here has the widest significance for the analysis of community care. Focusing on the views of carers as well as service providers, the book looks at caring across a variety of relationships and conditions including people with mental health problems and learning

disabilities *EBOOK: Cancer In Young Adults* Anne Grinyer, 2002-10-16 The original inspiration for this book was George who died from osteosarcoma at the age of 23 During his illness his parents tried without success to access information on the life stage issues that make life threatening illness during young adulthood particularly difficult to manage They could find no literature relating specifically to this problem and struggled throughout George's 4 years of living with cancer to cope with the additional problems faced by families in this situation After his death they set up a research project to help other families facing these issues This book is the outcome of that research It is heavily based on the use of narrative material written by parents whose young adult children have been diagnosed with cancer The book addresses issues such as sexuality and fertility independence the need for normality the effect on siblings the ownership of medical information financial issues the impact on the parents partnership and the emotional consequences of the illness It is designed to be of practical assistance both to parents and to health professionals involved with the care of young adults with cancer *Mental Health Care, Google eBook* Catherine Hungerford, Donna Hodgson, Richard Clancy, Tony Jones, Anthony Harrison, Chris Hart, 2012-02-02 A text written for non specialist mental health nursing courses and suitable for undergraduate students from nursing and other health disciplines which is supported by integrated online resources This is an adaptation of a Wiley Blackwell title by Harrison Hart editors *Mental Health Care for Nurses* designed to meet the needs of undergraduate students *Mental Health Nursing* is generally a core second year subject and may be taken by nurses paramedics occupational therapists and other health disciplines The focus is not on the general hospital as many nurses and other health professionals in Australia will not necessarily be working in the hospital setting It is important to consider the role of the multidisciplinary team

Supporting the Wellbeing of Children with SEND Kerry Murphy, 2022-03-07 As an early years practitioner you will educate and care for children with a range of developmental needs and differences This essential book introduces you to a play rich approach providing both universal and targeted ideas that will support social and emotional development and ensure that children feel safe secure and nurtured Using the four broad areas of need as a guide each accessible chapter positions wellbeing at the heart of an effective approach to inclusion and offers meaningful and responsive teaching practices that create a sense of belonging and acceptance Founded in the latest research the book presents key knowledge alongside ideas and activities to support wellbeing which can be embedded into the child's everyday experiences and adapted to meet their individual needs This book offers Evidence based strategies and techniques that have a positive impact on the long term social and emotional wellbeing of children with SEND Guidance through the four broad areas of need with a focus on play learning and developing an emotionally healthy early years environment Examples of practice in action Case studies reflective questions and activities that will upskill the reader and empower them in their role Providing up to date transferrable and essential knowledge on SEND in the early years this is an essential resource for any practitioner looking to expand their repertoire and enrich the wellbeing of children with SEND *Taking Care, Making Do* Caring Costs, 1991

Caring Families-- Families Giving Care Robin E. Cooper, National Association of State Directors of Developmental Disabilities Services, 2010

The Impact of Childhood Disability on Family Life Barbara Dobson, Sue Middleton, Alan Beardsworth, 2001 This report follows on from an earlier JRF project on childhood disability which described a minimum budget standard that parents thought necessary to raise a severely disabled child This book describes the actual spending patterns of parents with severely disabled children and presents a detailed examination of the costs involved

The Family Guide to Disability and Personal Finances Ed Arbuckle, 2018-05-02 This book is directed at helping individuals with disabilities and their families find financial solutions to the financial challenges of disability A very common concern of families is their difficulty in getting information that makes sense and lets them move on The search for answers is filled with complexity and while perhaps not a Rubik's Cube it certainly is a jigsaw puzzle with more than a few pieces The book is intended to be a guide for planning Planning will be better if it is based on structures for such things as family wealth transfers good tax and estate planning and the use of proper legal agreements The book will give readers a road map as well as understandable information that can be used to take their own situation to a higher level of resolution with more confidence and less heartache Whatever you may learn from The Family Guide remember that everyone with a disability is a person first and shouldn't be defined by their disability See them as they live their own lives everyday going to a hockey game attending school or being with family

Disability and Caring Among Families with Children Stephen McKay, Adele Atkinson, 2007 This report looks at a cross section of families in relation to disability caring and employment The main aims are to investigate the characteristics of families including their employment propensities where there is one or more disabled adult and or disabled child or child with health problems including families with both disabled adults and children or member with caring responsibilities inside or outside the household P 7

Family Policy and Disability Arie Rimmerman, 2015 This book explores the status and scope of family policies related to households of children with disabilities providing an in depth evidence based review of legal programmatic issues It includes a discussion of the gaps between family needs and contemporary family policies in the United States and European countries as demonstrated in these households surveys In addition the volume offers a comparative analysis of cash benefits tax credits and deductions and in kind provisions between the United States and select European countries UK France and Sweden Most importantly this book identifies and continues the discussion regarding the critical role of family centered policies as expressed in the United Nations Convention on the Rights of Persons with Disabilities UNCRPD as well as the future of family policy toward families of children with disabilities at a time of economic crisis

The Cost of Caring Shane Kavanagh, Louis Opat, Politeia (Association : London, England), 1998

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