

Paediatric Palliative Care Rcpch

Essential Nursing Care for Children and Young People is the definitive guide for all nursing and healthcare students and professionals caring for children and young people. Designed to meet the 2010 NMC competency standards for pre-registration nurses, the textbook supports you through the process of caring for children and young people with varied needs and conditions across all healthcare settings. A practical, patient-centred approach is taken throughout, with the 'voices' of children, their families and their carers used to tie theoretical knowledge to the real experience of providing care. The inclusion of the latest research and health and social care policies ensures that you are fully in line with the latest clinical practice, whilst the book's activities and exercises allow you to regularly check your understanding and develop confidence during your journey from student to nursing professional. Main features: Course-Focused Approach – pedagogy and content designed specifically for all three years of the child nursing degree programme. Contemporary Content – research note features and policy integrated throughout to give you instant access to the latest evidence-based practice. Illustrated Throughout – this highly accessible text regularly uses pictures and diagrams to highlight key issues. Voices – include authentic experiences of children, young people, parents, carers and professionals to help develop a patient-focused attitude to care. Pertinent A&P – includes coverage of all relevant anatomy and physiology for child and young person nursing courses. Practice Guidelines – provide practical guidance on everyday procedures for your quick reference. Activities and Answers – allow you to check your knowledge and build confidence. Specialist Authors – each chapter is written by leading experts in each area. Essential Nursing Care for Children and Young People is the ideal main textbook for all undergraduate child and young person nursing students, as well as professionals wanting to ensure they are using the latest practice. This text is also relevant to any student or professional involved in the health and social care of children and young adults.

This book is designed to provide a comprehensive insight into the key and most prevalent contemporary issues associated with palliation. The reader will find viewpoints that are challenging and sometimes discerning, but at the same time motivating and thought-provoking in the care of persons requiring palliation. This book is divided into three sections. Section 1 examines contemporary practice; Section 2 looks at the challenges in practice; Section 3 discusses models of care. This book is an excellent resource for students, practising clinicians and academics. By reading the book, reflecting on the issues, challenges and opportunities ahead, we hope it will create within the reader a passion to take on, explore and further develop their palliative care practice.

A concise and practical guide to caring for children with life-limiting conditions, Paediatric Palliative Care covers the common symptoms and challenging issues healthcare professionals are likely to encounter, and includes a detailed drug formulary for quick reference.

This comprehensive clinical practice manual emphasises evidence-based best practice, focusing on inter-disciplinary care of the child in hospital and community settings. The additional needs of neonates and adolescents are highlighted.

Children's palliative care has developed rapidly as a discipline, as health care professionals recognise that the principles of adult palliative care may not always be applicable to children at the end of life. The unique needs of dying children are particularly evident across Africa, where the scale of the problem is overwhelming and the figures so enormous that they are barely comprehensible. Written by a group with wide experience of caring for dying children in Africa, this book provides practical, realistic guidance on improving access to, and delivery of, palliative care in this demanding setting. It looks at the themes common to palliative care - including communication, assessment, symptom management, psychosocial issues, ethical dilemmas, end of lifecare, and tips for the professional on compassion and conservation of energy - but always retains the focus on the particular needs of the health care professional in Africa. Whilst containing some theory, the emphasis is on practical action throughout. It will provide health care professionals working in Africa, and other resource-poor settings, with the confidence, knowledge, and capacity to improve care for the terminally ill child in constrained and demanding environments.

This book is a printed edition of the Special Issue "Paediatric Palliative Care" that was published in Children

In this stimulating and provocative book the editors have drawn together a diverse and international range of respected authors, each of whom has taken a critical approach to the contentious question of how you define and achieve quality early childhood services. It is a book designed to provoke and promote critical dialogue and discourse amongst practitioners and students through critical engagement with the position of the authors within the text. I believe anyone who reads this book will be inspired and motivated to challenge and extend their thinking and professional practice, adopting the critical stance which lies at the heart of quality services for children and families. Professor Chris Pascal, Director of Centre for Research in Early Childhood (CREC) Early childhood is a complex and important area of study where it is important to develop your critical thinking and reflect upon key issues. This book will help do both. It explores interrelated topics such as: Child development Play Safeguarding Professionalism Curriculum and Policy Each chapter will not only engage with what you need to know but help you develop your academic skills. The book also comes with lots of online resources which can be found at <https://study.sagepub.com/reedandwalker> and include: Podcasts from the authors of each chapter so you can better understand the key concepts PowerPoints to help you revise the essential information Journal articles related to each chapter provide further reading Michael Reed and Rosie Walker are both Senior Lecturers in Early Childhood at the Institute of Education, University of Worcester.

Providing an in-depth look at the issues surrounding the palliative care of children and their families, this innovative text considers children with both malignant and non-malignant conditions, and the specialist care required. By addressing the complexities of providing palliative care to children, it aims to assist all those who work in this field to enhance their practice. Special features include: ? broad research evidence, from national and international sources, on which to base practice ? absorbing activities to support learning and help build knowledge in a meaningful way ? interprofessional perspectives from the whole palliative care team The contributors to the book are drawn from both education and practice across a range of disciplines. The result is a genuinely engaging, interdisciplinary text that is essential reading for students and professionals aiming to deliver quality palliative care in partnership with children and their families.

"Children's palliative care is an evolving specialty and as such our knowledge base cannot remain static. This book constantly challenges the reader to critically analyze their own practices and beliefs within an evidence-based framework and as such makes a valuable contribution to the growing body of knowledge on this important subject." - Susan Fowler-Kerry, in the Foreword. This book caters for readers from different working environments and levels of experience. It is ideal for paediatric nurses with no specialist palliative care knowledge, and also for palliative care nurses with no specialist paediatric experience. Other healthcare professionals and therapists working with children, young people and their families will also find this book invaluable. It will also be ideal for undergraduate and postgraduate health and social care students, and professionals involved in children's hospices, community services and charity groups. "This book captures not just the rapidly evolving evidence base, but also as many current developments as possible and applies them in a meaningful way to the care of infants, children, and young people living with life-limiting illness, and their families. It offers an overview of contemporary issues and helps to stimulate the type of dialogue that can bring about the actions that will make a real difference for the children, young people and families in our care." - Rita Pfund, in the Preface.

Comprehensive in scope, exhaustive in detail, and definitive in authority, this third edition has been thoroughly updated to cover new practices, current epidemiological data, and the evolving models that support the delivery of palliative medicine to children. This book is an essential resource for anyone who works with children worldwide.

"From its beginning with Eugene Stead, MD at Duke University, the Mission of the Physician Assistant profession has been to create, increase or expand health care access in areas and populations with the greatest need.1 Dr. Henry Silver's Child Health Associate Program

at the University of Colorado is a good example as is Dr. Hu Myer's Alderson Broaddus Program which serves isolated rural communities in West Virginia.² Dr. Richard Smith's MEDEX Program at the University of Washington was also designed specifically to serve rural and medical underserved communities of the Pacific Northwest.³--

Provides a comprehensive overview of good practice in caring for terminally ill children, young people and their families.

The second edition of *Nursing Care of Children and Young People with Long Term Conditions* remains the only nursing-specific text on the care of paediatric patients with chronic illness. Written to meet the needs of nursing students and professionals alike, this comprehensive volume provides authoritative and up-to-date information on the context, theory, and practice of delivering holistic care to children and families in a range of health and social care settings. Contributions from a team of experienced academics, educators, and practitioners offer valuable insight into the impact of chronic illness on children and parents, the practical implications of meeting their physical, psychological, and social needs, empowering them to be 'experts' in their care, and many more vital aspects of long-term paediatric care. This edition features new and revised content reflecting contemporary guidelines and evidence-based practice, including updated clinical case studies and a new chapter examining the impact of having a sibling with a long-term condition. Emphasising a multi-disciplinary approach to managing chronic illness, this important resource: Provides numerous case studies and activities illustrating the application of theoretical principles and current evidence in nursing practice Investigates the genetic basis of chronic illness and the differing onsets of long-term conditions Discusses current political, economic, and social policies that are influencing healthcare for children and bringing challenges to managers and practitioners Examines both classic and contemporary theories of grief, loss, coping, and adaptation Explores ethical, legal, and professional aspects of nursing children and young people with chronic illness Addresses evolving nursing roles, the importance of acute emergency care, and the planning and delivery of effective transition from child to adult services *Nursing Care of Children and Young People with Long Term Conditions* is required reading for student and registered children's nurses, as well as for practitioners in related health and social care disciplines.

The death of a child is a special sorrow. No matter the circumstances, a child's death is a life-altering experience. Except for the child who dies suddenly and without forewarning, physicians, nurses, and other medical personnel usually play a central role in the lives of children who die and their families. At best, these professionals will exemplify "medicine with a heart." At worst, families' encounters with the health care system will leave them with enduring painful memories, anger, and regrets. *When Children Die* examines what we know about the needs of these children and their families, the extent to which such needs are "and are not" being met, and what can be done to provide more competent, compassionate, and consistent care. The book offers recommendations for involving child patients in treatment decisions, communicating with parents, strengthening the organization and delivery of services, developing support programs for bereaved families, improving public and private insurance, training health professionals, and more. It argues that taking these steps will improve the care of children who survive as well as those who do not and will likewise help all families who suffer with their seriously ill or injured child.

Featuring illustrative case histories, the book discusses patterns of childhood death and explores the basic elements of physical, emotional, spiritual, and practical care for children and families experiencing a child's life-threatening illness or injury.

Optimal terminal and palliative care requires consideration of the patient and family unit as well as cultural and religious sensitivities. The patient's well being in terms of mobility, anxiety, stress, social interaction, and pain control needs expert focus and attention. Furthermore, there is an increasing awareness that diet and nutritional support plays an integral part of the patient's holistic well being. The interface between nutritional, emotional, cultural, and medicinal support challenges terminal and palliative care providers to recognize the right thing to do, often in the face of considerable uncertainty. Currently, there is no comprehensive book on nutrition in terminal or palliative care that is suitable for novices and experts alike. *Diet and Nutrition in Palliative Care* addresses this deficiency in the literature. Designed for doctors, nurses, caregivers, and those working within the palliative or end of life domain, each chapter contains sections on applications to other areas of terminal or palliative care, practical methods and techniques, guidelines, key points and ethical issues. The book is divided into six sections: Setting the Scene Cultural Aspects General Aspects Cancer Non-Cancer Conditions Pharmacological Aspects Despite the complexity of the correlation between diet and disease, there is now a sufficient body of evidence to encourage applying nutritional science in everyday clinical practice. Increasingly, a strong interest and up-to-date knowledge and understanding of scientific studies on nutrition enables clinicians to help patients under their care more effectively at every stage of their illness. In response to this need, this handbook presents important information on the holistic use of nutrition and diet therapy in palliative care.

'A Guide to the Development of Children's Palliative Care Services' is a key resource for all organisations and professionals who have an interest in developing or supporting services for life-limited or life-threatening children and their families.

Designed for teachers in cancer and palliative care, this work covers the complex changes in cancer and palliative care delivery. It includes highlighted key points, self help questions for reflection, and references where applicable. It provides guidance for healthcare professionals with palliative care teaching responsibilities.

Handbook of Nutrition and Diet in Palliative Care, Second Edition, is a comprehensive guide, providing exhaustive information on nutrition and diet in terminal and palliative care. It covers physical, cultural and ethical aspects, bridging the intellectual divide in being suitable for novices and experts alike. Following in the tradition of its predecessor, chapters contain practical methods, techniques, and guidelines along with a section on applications to other areas of palliative care. Each chapter features key facts highlighting important areas, summary points, and ethical issues. **FEATURES** • Use of cannabinoids in palliative nutrition care • Pain control in palliative care • Communications in palliative/end-of-life care: aspects of bad news • Anorexia in cancer: appetite, physiology, and beyond • Palliative care in severe and enduring eating disorders • Linking food supplementation and palliative care in HIV • Eating-related distress in terminally ill cancer patients and their family members • Palliative care of gastroparesis • Preoperative nutrition assessment and optimization in the cancer patient • Childhood leukemia, malnutrition, and mortality as components of palliative care • End-of-life decisions in persons with neurodevelopmental disorders • Resources: listing web sites, journals, books and organizations

How does palliative care differ in the Republic of Ireland to other countries? This text provides an overview of palliative care services in Ireland from a multi-professional viewpoint.

Draw upon relevant research and the author's experience to outline and contextualise the key aspects of loss: theoretical concepts, approaches and vocabulary terms. Loss and bereavement is a sensitive area; no two stories will ever be the same. At its core this resource aims to explore the common anxieties your students will encounter in this specific setting and provide strategies for managing the challenges unique to loss counselling. It is a practical guide for anyone taking an award, certificate or diploma in counselling and for experienced practitioners looking to implement best practice.

This book is "innovative and original in assisting the reader to apply the principles of science to paediatric practice .

Professor Neena Modi, President, Royal College of Paediatrics and Child Health. The Science of Paediatrics, MRCPCH Mastercourse, provides essential background preparation for the MRCPCH Theory and Science examination. It contains an up-to-date review of the application of science to everyday paediatric clinical practice, whether it is interpreting clinical signs or investigations, prescribing drugs or identifying best management. Although this understanding is essential in order to make informed clinical decisions, it is difficult to obtain as it is not usually covered in clinical textbooks. Key features • MRCPCH exam-format questions embedded in each chapter to test understanding • Emphasis on embryology to explain many congenital abnormalities • An overview of the relevant anatomy and physiology • Focus on the application and interpretation of investigations • Examples of recent advances in science and clinical research that have benefited the children's care • All clinical specialties covered by paediatric specialists. • Chapters covering evidence-based paediatrics, statistics, ethics and quality improvement.

This textbook is the first to focus on comprehensive interdisciplinary care approaches aimed at enhancing the wellbeing of children with cancer and their families throughout the illness experience. Among the topics addressed are the epidemiology of pediatric cancer distress, including physical, emotional, social, and spiritual dimensions; the role of the interdisciplinary team; communication and advance care planning; symptom prevention and management; care at the end of life; family bereavement care; and approaches to ease clinician distress. The contributing authors are true experts and provide guidance based on the highest available level of evidence in the field. The book has not only an interdisciplinary but also an international perspective; it will appeal globally to all clinicians caring for children with cancer, including physicians, nurses, psychosocial clinicians, and chaplains, among others.

Drawing on a range of approaches developed by paediatric chaplaincy teams worldwide, this edited collection provides best principles, practices and skills of chaplaincy work with neonates, infants, children, young people and their families. By engaging with paediatric chaplaincy from an international, multifaith perspective, contributors from around the world and different faith traditions show what good spiritual, religious and pastoral care for children and their families looks like. The book contains contributions from specialists who work with children with mental health issues or profound disabilities, as well as chapters that focus on how best to provide palliative and bereavement care. Includes resources and activities for use in specialist care situations and tools for assessment, making this a must-have for any paediatric chaplaincy team working in a hospital or hospice.

Issues in Disability, Rehabilitation, Wound Treatment, and Disease Management: 2012 Edition is a ScholarlyEditions™ eBook that delivers timely, authoritative, and comprehensive information about Palliative and Supportive Care. The editors have built Issues in Disability, Rehabilitation, Wound Treatment, and Disease Management: 2012 Edition on the vast information databases of ScholarlyNews.™ You can expect the information about Palliative and Supportive Care in this eBook to be deeper than what you can access anywhere else, as well as consistently reliable, authoritative, informed, and relevant. The content of Issues in Disability, Rehabilitation, Wound Treatment, and Disease Management: 2012 Edition has been produced by the world's leading scientists, engineers, analysts, research institutions, and companies. All of the content is from peer-reviewed sources, and all of it is written, assembled, and edited by the editors at ScholarlyEditions™ and available exclusively from us. You now have a source you can cite with authority, confidence, and credibility. More information is available at <http://www.ScholarlyEditions.com/>.

All children have a right to appropriate prevention, assessment and control of their pain. Managing Pain in Children is an evidence-based, practical guide to care in all areas of children's pain management, providing nurses and other health care practitioners with the skills and expertise necessary to manage children's pain effectively. The text first explores the relevant anatomy and physiology of children, the latest policy guidelines surrounding pain management and ethical issues involved in managing children's pain. It then goes on to look at the various pain assessment tools available for children and non-drug methods of pain relief. The text then goes on to apply these pain management principles to practice in relation to acute pain, chronic pain, palliative care and the management of procedural pain. Each of these chapters covers the evidence base, assessment techniques, pain relieving interventions, and guidance for best practice in both hospital and community settings. Special features: Practical and accessible, with 'best practice' points in each chapter Written in the context of latest policy developments Provides the necessary evidence-base for care throughout With contributions from experts in the field

Palliative Nursing is an evidence-based practical guide for nurses working in areas of practice where general palliative care is provided. This may be in hospitals, nursing homes, dementia units, the community and any other clinical areas which are not classified as specialist palliative care. This book first explores the history and ethos of palliative care, and then looks at palliative nursing across various care settings. It then looks at palliative nursing care for people with specific illnesses, including heart failure, dementia, chronic obstructive pulmonary disease, cancer, and neurological conditions. Palliative care for children and young people is discussed, and then the book finally looks at education and research in palliative nursing. Palliative Nursing will be essential reading for all nurses working with palliative care patients in a non-specialist role, i.e. in hospitals, primary care and nursing homes, as well as nursing students. SPECIAL FEATURES Explores the palliative nursing issues related to specific diseases groups Written in the context of the new national tools, i.e. the end of life initiative, preferred place of care, Liverpool care pathway and Gold standards framework. Each chapter includes practice points and cases to allow the practitioner to undertake guided reflection to improve practice Written by nurses for nurses Provides guidance for nurses working in all four countries of the UK

Pediatric palliative care is a field of significant growth as health care systems recognize the benefits of palliative care in areas such as neonatal intensive care, pediatric ICU, and chronic pediatric illnesses. Pediatric Palliative Care, the fourth volume in the HPNA Palliative Nursing Manuals series, highlights key issues related to the field. Chapters address pediatric hospice, symptom management, pediatric pain, the neonatal intensive care unit, transitioning goals of care between the emergency department and intensive care unit, and grief and bereavement in pediatric palliative care. The content of the concise, clinically focused volumes in the HPNA Palliative Nursing Manuals series is one resource for nurses preparing for specialty certification exams and provides a quick-reference in daily practice. Plentiful tables and patient teaching points make these volumes useful resources for nurses. Taking account of the British government's "End of Life Care Strategy", contributors set out the key issues affecting practice

across a range of health and social care contexts. The book covers topics ranging from dying and death to symptom management and spiritual care, backed up with practical examples. Each entry comprises: a snapshot definition of the topic, key points, a discussion of the main debates, links to practice through thought-provoking case histories, and suggestions for further reading. The book provides comprehensive, practical guidelines on the responsibilities of those who lead, co-ordinate and manage volunteers in small hospices, large specialist palliative care units, and in general hospitals with palliative care teams. Volunteers are key workers, who often perform difficult and important work. In the United Kingdom alone there are thousands of volunteers in hospice work, a small proportion doing work with patients, and the vast majority doing equally valuable work such as driving, sitting with relatives, manning charity shops and telephones. As a result, Europe, Australia, the United States and Canada are very interested in the United Kingdom's use of volunteers. Aimed primarily at Voluntary Services Managers in small hospices, large specialist palliative care units, and in general hospitals with palliative care teams, this updated second edition covers volunteer selection, training, supervision and support, and legal and ethical considerations. Information is presented in an easily accessible way, using key points, summary panels and checklists. This book will also appeal to the volunteers themselves.

This exciting new edition is again structured into four main sections: Organisational facets; Philosophical issues; Dimensions of practice; and Advancing practice, and has been expanded to include detailed guidance on the commissioning and resourcing of services. It provides essential information for implementing the requirements of the children's National Service Framework that will support the expansion of Community Children's Nursing and enable it to move forward and away from fragmented service delivery. Bringing together the work of some of the most distinguished experts in the field, there is comprehensive coverage of the key aspects of Community Children's Nursing, including multi-disciplinary/interagency planning; provision of nursing services to sick children and their families in a range of community setting; and the needs of both the recipients and providers of care within the trajectory of acute, life-limiting and terminal illness. The first - and only - book on this topic, addressed specifically at the Community Children's Nurse (CCN) An authoritative guide to the principles underpinning the development of the specialty, which puts into context the scope of the CCN's work and clearly describes his/her place in the community team) A balance of the theoretical and practical, presented by the key names in this field) Offers up-to-date "evidence" which supports the development of this rapidly expanding specialty) Foreword by Elizabeth Fradd, Independent Health Service Advisor, UK New chapters have been added on the topics of: Developing a national strategy and corporate identity for Community Children's Nursing Delivering and funding care for children with complex needs Strategic planning and commissioning of services Benchmarking Transitional care The following chapters have been rewritten and expanded: Young carers Complementary therapies Partnerships with the voluntary sector Health promotion Information management The role of the Community Children's Nurse Manager is also explored in depth.

Nurses and healthcare professionals are constantly faced with ethical and philosophical dilemmas when working with children in everyday practice. Ethical and Philosophical Aspects of Nursing Children and Young People is a comprehensive text on the ethics and philosophy behind paediatric nursing that reflects the contemporary issues encountered while working with children and young people. The title provides a philosophical and historical analysis of the subject, looking at a review of sociological and political theories concerning the nature of childhood, and providing a critical analysis of contemporary notions about childhood. It then goes on to look at moral theories and their application to paediatric nursing practice, ethical issues when caring for children of all ages, from infancy to adolescence. It considers issues of disabled children, confidentiality, mental health issues, children's rights, and pain management. With case studies and activities throughout, this book will enable students and newly qualified nurses both to understand philosophical concepts and issues but also to articulate their own reflections and observations on these subjects. Written by children's nurses for children's nurses With contributions from internationally recognised experts in the field Reflective scenarios, further reading, extensive referencing, case studies, guided questions, and resources throughout Includes appendices on the RCPCH Guidelines on Withdrawal of Treatment, the ICN Ethical Code, the Summary of the UNCRC, and the RCPCH Guidelines for the Conduct of Ethical Research

This handbook offers a practical, thorough approach to the clinical practice of palliative care. Adding North American authors to its roster of UK contributors, the third edition of this award-winning book addresses important changes in the evidence base of palliative care, as well as an emphasis on end-of-life community-based care. It features new chapters on dementia and advance care planning, a simplified lymphoedema discussion, and an ongoing commitment to providing essential guidance for physicians, nurses, and all primary care providers involved in palliative care in hospital, hospice, and community settings.

The first of its kind, this book describes pediatric palliative care in more than 23 countries. Each region in the world is covered and countries included are both resource poor and rich. Authors are multidisciplinary and regarded nationally and internationally in their field. Clinicians, advocates, policymakers, funders, and researchers will learn how programs were developed and implemented in each country. Authors describe children for whom pediatric palliative care is needed and provided for in their country. When applicable, a brief history of pediatric palliative care is included noting especially policy changes and legislative acts. For example, the chapter on Poland describes how pediatric palliative care grew from the Catholic church into a national movement spearheaded by several health care workers. The Pole national spirit that brought them through a change in political regime has also been a driving force in the pediatric palliative care movement. The chapter on South Africa, for example, illustrates how a resource poor country has been able to leverage philanthropic and government funding to make its dream of having an infrastructure of pediatric palliative care a reality. These are just a few examples of the inspiring stories that are included in this book. Readers from countries who wish to start a pediatric palliative care program, or advance an existing program, will learn valuable lessons from others who have faced similar barriers. Introduction and concluding chapters highlight the strengths and weaknesses of the modern pediatric palliative care movement.

Endorsed by the RCPCH and ESPID, and packed with helpful tips and practical guidance, The Blue Book is an easy to use, easily-accessible, but fully comprehensive and evidence-based reference guide, helping busy paediatricians recognise, investigate and manage both common and rare infectious diseases in children and babies.

A Really Practical Handbook of Children's Palliative Care for Doctors and Nurses Anywhere in the World offers really practical solutions to common problems faced by health professionals caring for dying children and their families, whatever their culture Looking after children with life-limiting conditions can be very difficult for both parents and health care professionals. This second edition of Paediatric Palliative Medicine is full of easily-accessible, detailed information on medical conditions and symptoms and includes specific management plans in order to guide the practicing clinician through treatment of children requiring palliative care. Using the bestselling Oxford Specialist Handbook format to deliver practical and concise information, this handbook facilitates bedside delivery of effective palliative medicine to children by professionals who have not trained or had experience of caring for the dying child, as well as for students and trainees

interested in paediatric palliative care. It includes detailed information on symptom control and the philosophy and models that support delivery of palliative medicine to children, while also covering practical delivery of palliative medicine relating to other professionals and to families, and the learning and coping skills required in palliative care. It also contains a quick-reference drugformulary. Fully updated with an expanded formulary and a new chapter on the intensive care unit, this new edition continues to be the authoritative reference tool in paediatric palliative care.

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