

The purpose of this manual

This manual is intended to be a guide for the medical equipment user to carry out basic maintenance tasks. As the majority of equipment problems are either simple or user-related it is the aim that the better care and regular maintenance enabled by this manual will have a significant positive effect on the delivery of healthcare across India. The tasks are limited to simple ‘first-line’ maintenance, that is:

- tasks that can be done by the user of the equipment
- tasks that take place at the point of equipment use
- tasks that do not require the opening of the main body of the equipment

This manual is not intended as a complete maintenance guide – that is the role of a biomedical technician. Neither is it intended to be a guide to the actual use of equipment – it is assumed that the user is trained in the correct operation of the equipment. Users are asked to note that while every care has been taken to make the contents as clear and accurate as possible, neither the authors, the Ministry of Family Health and Welfare nor Crown Agents can take responsibility for the results of actions taken as a consequence of using this manual.

The format of this manual

The text of the manual is in English and is designed for on-line access as well as hardcopy prints. General topics on maintenance and disposal are covered by individual chapters. Section 7 covers the most commonly found equipment in detail. Each equipment section comprises:

- a brief description of the function and working of the equipment
- a line drawing of the equipment and its parts
- a troubleshooting checklist for common problems and their solution
- a maintenance checklist for daily and weekly tasks

The checklists are on separate pages so they can be copied and laminated for display near the equipment.

The choice of which equipment to include was guided by the 2010 revision of the Indian Public Health Standards. Equipment specified for health institutions up to the size of a 50 bed hospital was included, on the basis that this will cover the vast majority of simple equipment also found elsewhere. More advanced equipment will naturally require more advanced maintenance support.

This manual does not include laboratory equipment, since the recent excellent World Health Organization publication “Maintenance Manual for Laboratory Equipment” covers these in great detail. Similarly, cold chain equipment is covered comprehensively by the Indian MoHFW 2009 publication “Maintenance of Cold Chain Equipment”.

Acknowledgements

This manual draws on work done by many in this field. In particular, the authors acknowledge:
The UK Department for International Development for project funding

Medical Equipments Service Manual

Canada



Medical Equipments Service Manual:

Medical Equipment Service Manual Frank Biloon, 1978 The format is particularly suited to the quick effective repair of equipment malfunctions Wherever possible a description of the equipment is followed by diagrammatic description and numbered check lists for service Also included are photographs and schematics of equipment currently in use **Clinical Engineering Handbook** Joseph F. Dyro, 2004-08-27 As the biomedical engineering field expands throughout the world clinical engineers play an ever more important role as the translator between the worlds of the medical engineering and business professionals They influence procedure and policy at research facilities universities and private and government agencies including the Food and Drug Administration and the World Health Organization Clinical engineers were key players in calming the hysteria over electrical safety in the 1970s and Y2K at the turn of the century and continue to work for medical safety This title brings together all the important aspects of Clinical Engineering It provides the reader with prospects for the future of clinical engineering as well as guidelines and standards for best practice around the world

The Acquisition and Maintenance of Medical Equipment United States. Bureau of Domestic Commerce, 1975
Monthly Catalogue, United States Public Documents, 1993 Center for Devices and Radiological Health Publications Index Center for Devices and Radiological Health (U.S.). Publications Support Branch, 1978 Unit, Direct Support, and General Support Maintenance Manual (including Repair Parts and Special Tools List), 1990 **Current Catalog** National Library of Medicine (U.S.), 1979 First multi year cumulation covers six years 1965 70 *Monthly Catalog of United States Government Publications* United States. Superintendent of Documents, 1993 *Center for Devices and Radiological Health Publications Index* United States. Center for Devices and Radiological Health, 1988 **Investigational Device Exemptions Manual** Center for Devices and Radiological Health (U.S.). Division of Small Manufacturers Assistance, 1996

Investigational Device Exemptions Manual, 1996 **Inspection of Medical Devices** Almir Badnjević, Mario Cifrek, Ratko Magjarević, Zijad Džemić, 2017-10-26 This book offers all countries a guide to implementing verification systems for medical devices to ensure they satisfy their regulations It describes the processes procedures and need for integrating medical devices into the legal metrology framework addresses their independent safety and performance verification and highlights the associated savings for national healthcare systems all with the ultimate goal of increasing the efficacy and reliability of patient diagnoses and treatment The book primarily focuses on diagnostic and therapeutic medical devices and reflects the latest international directives and regulations Above all the book demonstrates that integrating medical devices into the legal metrology system and establishing a fully operational national laboratory for the inspection of medical devices could significantly improve the reliability of medical devices in diagnosis and patient care while also reducing costs for the healthcare system in the respective country Radiology in Global Health Daniel J. Mollura, Matthew P. Lungren, 2014-07-02 The World Health Organization stated that approximately two thirds of the world s population lacks adequate access to

medical imaging The scarcity of imaging services in developing regions contributes to a widening disparity of health care and limits global public health programs that require imaging Radiology is an important component of many global health programs including those that address tuberculosis AIDS related disease trauma occupational and environmental exposures breast cancer screening and maternal infant health care There is a growing need for medical imaging in global health efforts and humanitarian outreach particularly as an increasing number of academic government and non governmental organizations expand delivery of health care to disadvantaged people worldwide To systematically deploy clinical imaging services to low resource settings requires contributions from a variety of disciplines such as clinical radiology epidemiology public health finance radiation physics information technology engineering and others This book will review critical concepts for those interested in managing establishing or participating in a medical imaging program for resource limited environments and diverse cross cultural contexts undergoing imaging technology adaptation **The Canada Gazette**

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Publications Index Center for Devices and Radiological Health (U.S.),1988 **Journal of the Royal Army Medical Corps** Great Britain. Army. Royal Army Medical Corps,1910 **Book Bulletin of the Chicago Public Library** Chicago Public Library,1917 **National Library of Medicine Current Catalog** National Library of Medicine (U.S.),1990

International Labeling Requirements for Medical Devices, Medical Equipment and Diagnostic Products Charles Sidebottom,2003-06-27 Completely revised this second edition provides the practical hands on labeling information needed to secure rapid regulatory approval gain marketplace acceptance and assure user comprehension A complete guide to all aspects of advertising labeling and packaging it explains the relevant laws regulations and requirements in major markets w

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