

factors from the perspective of the patient and family. The flow charts are not designed to be read as text; they are too repetitious. However, with their indexes they are ideal for reference.

While primarily designed for emergency departments, this volume could be used in walk-in clinics or urgent-care centres to help determine which individuals would be better served in another type of facility. It would also be useful as a teaching tool for house staff taking calls in the family medicine setting.

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The Ethics of Human Gene Therapy

LeRoy Walters and Julie Gage Palmer. 209 pp. Illust. Oxford University Press, Oxford, England; Oxford University Press Canada, Don Mills, Ont. 1997. \$42.95. ISBN 0-19-505955-7

Overall rating: Excellent

Strengths:

Easily understood, concise description of science, techniques and possible uses of gene therapy; listing of ethical issues from a clinical

ethics perspective Little analysis of the collec-

Weaknesses:

tive effects of the use of gene therapy on shaping social attitudes and practices; incomplete discussion of social and . ethical issues

Audience:

Health professionals, research ethics committee members, general public

The authors' purpose was to de-**L** scribe the science of gene therapy and to examine the major ethical questions. Toward the end of the volume, they state that their mission has been accomplished if they have stimulated the reader "to think about these possibilities in a calm and rational way and to reach their own judgment." Most of the 153 pages of text and the 43 pages of appendices focus on describing and illustrating the current and potential uses of gene therapy. There are useful summaries of guidelines and opinion surveys.

The authors, a philosopher and an attorney, introduce genes and heredity in the first chapter. The next three chapters discuss somatic and germ-line therapy, and enhancement genetic engineering, and a fifth chapter considers public policy. Each chapter begins with a clear, well-illustrated discussion of historical developments and builds to the current developments.

Technical advantages are clearly described, without ethical assessment. For instance, the advantages of germ-line over somatic therapy are described as more effective when the disease 1) affects many different organs and disparate cell types (e.g., cystic fibrosis); or 2) is expressed in nonremovable and nondividing cells (e.g., Lesch-Nyhan syndrome). Of course, germline therapy is also effective beyond the individual, with the therapeutic effect passed on to subsequent children and a possible reduction of certain inherited diseases in the human gene pool. Technical discussions are followed by an evaluation of ethical arguments. Generally speaking, the authors take a permissive stance with respect to somatic and germ-line therapy and toward enhancement genetic engineering.

Their ethical discussions are probably the weakest point in the volume. The technical discussions receive about twice as much discussion as the ethical, and the approach is implicitly utilitarian. The authors admit that their analysis is shaped by their assumptions about health (speciestypical function or improved function) and human nature (dissatisfaction with the human condition). Their fascination with the details of the technologies seems to connote a bold embrace of genetic technologies without evaluation of alternatives. It is curious that the authors of such a historically and technologically grounded account seem to lack the humility and sense of fallibility that lead other authors to more cautious conclusions. For instance, they conclude: "genetic enhancements are an important part of the overall task of attempting to provide a better life and a better world to our descendants."

Overall, the volume is well written and readable for anyone who has taken a college or high school biology course. It will certainly stimulate reflection by interested physicians, members of research ethics committees, or members of the general public. Ethical reflection would benefit from supplemental reading through the perspective of feminist writers and persons with disabilities.

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Slow Dance: A Story of Stroke, Love and Disability

Bonnie Sherr Klein. 365 pp. Alfred A. Knopf Canada, Toronto. 1997. \$29.95. ISBN 0-394-28161-6

Overall rating:

Strengths:

Weaknesses:

Important insights into the challenges of stroke survivors Fewer "characters" analysed in greater detail might have

had more impact

Audience:

Health care professionals, stroke patients and their fam-

This book is a narrative account lacksquare by its author, an award-winning documentary NFB filmmaker, of survival and partial recovery from a brainstem hemorrhage. It can be



compared to a National Film Board documentary with remembrances by family, friends, physicians and therapists, with voice-over by the patient. The cast is lengthy, with cameo appearances by hospital workers, fellow patients, rabbis and purveyors of alternative therapies. Locations include Montreal, London, Ont., Vancouver, Costa Rica, and the patient's mind. It is "directed" by the author, with her physician husband in a leading role. The reader's imagination supplies sound and images.

Scenes weave through initial symptoms, early diagnostic confusion, clinical deterioration, MRI demonstration of the lesion, surgery and the long road of rehabilitation. Subplots include feminism, family catharsis and the patient as advocate. But the dominant theme is tenacity and the need to maintain dignity while coming to terms with a damaged body trying to navigate through each new day.

There are varying examples of how the system can be manipulated by those who know their way around it. In the words of a friend: "Michael [Klein's husband] made up his own rules for your care." This may be discouraging to readers without such a connection.

The story includes their search for alternative therapies: acupuncture, quartz crystals, folded Japanese paper cranes, meditation tapes, Feldenkrais, Alexander technique ... the list goes on.

She tells the story of a US stroke patient who struggled on her own without insurance as an example of problems with the US health care system. It invites speculation: How different would Klein's story have been if she had to trade places?

Physicians, patients, feminists and health care workers will have varying reactions to this book. But everyone will benefit from a better understanding of the many physical and emotional hurdles faced by the author. Osler was right: "Listen to the patient."

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[See the Jan. 1, 1997, issue for Klein's husband's experience and how he challenged the medical profession to rethink its views toward the physician as family member (page 53). — Ed.]

Books and other media received

Livres et autres documents recus

Endocrinology

Endocrinology: Basic and Clinical Principles. Edited by P. Michael Conn and Shlomo Melmed. 448 pp. Illust. Humana Press, Totowa, NJ. 1997. US\$125. ISBN 0-89603-349-X

Epidemiology

The Global Burden of Disease: A Comprehensive Assessment of Mortality and Disability from Diseases, Injuries, and Risk Factors in 1990 and Projected to 2020. Edited by Christopher J.L. Murray and Alan D. Lopez. Global Burden of Disease and Injury Series, vol. 1. 990 pp. Illust. World Health Organization; Harvard University Press, Cambridge, Mass. 1996. US\$39.95. ISBN 0-674-35448-6

Family medicine

Saunders Review of Family Practice. 2nd ed. Edited by Edward T. Bope, Alvah R. Cass and Michael D. Hagen. 354 pp. Illust. W.B. Saunders Company/Harcourt Brace and Company, Philadelphia; W.B. Saunders Canada, Toronto. 1997. \$71.95. ISBN 0-7216-5817-2

Health care

An Inventory of Quality Initiatives in Canada: Maintaining and Improving Quality in Health Care. 2nd ed. Health Canada. 307 pp. Health Canada, Ottawa; fax 613 941-5366. 1997. Free of charge. ISBN 0-662-24978-X. (Includes inventory of Internet sites relevant to quality in health care issues. Also available online [www.hwc.ca] or on computer diskette or audiocassette, or in large print or braille. Également disponible en français.)

History

Jean I. Gunn: Nursing Leader. Natalie Riegler. Hannah Institute for the History of Medicine Series: Canadian Medical Lives. Series editor, T.P. Morley. 263 pp. Illust. Fitzhenry and Whiteside, Markham, Ont. 1997. \$18.95. ISBN 1-55041-175-6

Neurology

Neurological Investigations. Edited by R.A.C. Hughes. Vol. 3 of *Journal of Neurology, Neurosurgery, and Psychiatry* books. 511 pp. Illust. BMJ Publishing Group, London. 1997. Distributed in Canada by the Canadian Medical Association, Ottawa. \$137.95 (\$114.95 CMA members). ISBN 0-7279-1080-9

Pharmacology

Goodman and Gilman's The Pharmacological Basis of Therapeutics (CD-ROM). 9th ed. Edited by Joel G. Hardman, Lee E. Limbird, Perry B. Molinoff, Raymond W. Ruddon and Alfred Goodman Gilman. Illust. McGraw-Hill Companies, Health Professions Division, NY; McGraw-Hill Ryerson Ltd., Whitby, Ont. 1996. \$181.25. ISBN 0-07-864186-1

Surgery

Practical Minor Surgery (CD-ROM). David Crawford and Christopher Koo. Omni Media Associates and harwood academic publishers, The Netherlands; Overseas Publishers Association, Amsterdam. 1996. Price not stated. ISBN 90-5702-007-6 (single user licence); ISBN 90-5702-006-8 (site/institution licence)