Variations in Hospital Length of Stay: Their Relationship to Health Outcomes

August 1983

NTIS order #PB84-111483

HEALTH TECHNOLOGY CASE STUDY 24: Variations in Hospital Length of Stay: Their Relationship to Health Outcomes

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This case study was performed as a part of OTA's Assessment of

Medical Technology and Costs of the Medicare Program



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Prepared under contract for OTA by: Mark R. Chassin, M. D., M. P. P., M.P.H.

**OTA Case Studies** are documents containing information on a specific medical technology or area of application that supplements formal OTA assessments. The material is not normally of as immediate policy interest as that in an OTA Report, nor does it present options for Congress to consider.

Recommended Citation:

Health Technology Case Study 24: Variations in Hospital Length of Stay: Their Relationship to Health Outcomes (Washington, D. C.: U.S. Congress, Office of Technology Assessment, OTA-HCS-23, August 1983). This case study was performed as a part of OTA's assessment of Medical Technology and Costs of the Medicare Program.

Library of Congress Catalog Card Number 83-600708

For sale by the Superintendent of Documents, U.S. Government Printing Office, Washington, D.C. 20402

#### Preface

Variations in Hospital Length of Stay: Their Relationship to Health Outcomes is Case Study 24 in OTA's Health Technology Case Study Series. It was prepared in response to a request by the Senate Finance Committee, Subcommittee on Health, and is part of OTA's project on Medical Technology and Costs of the Medicare Program, requested by the House Committee on Energy and Commerce and its Subcommittee on Health and the Environment and the Senate Finance Committee, Subcommittee on Health. A listing of other case studies in the series is included at the end of this preface.

OTA case studies are designed to fulfill two functions. The primary purpose is to provide OTA with specific information that can be used in forming general conclusions regarding broader policy issues. The first 19 cases in the Health Technology Case Study Series, for example, were conducted in conjunction with OTA's overall project on The Implications of Cost-Effectiveness Analysis of Medical Technology. By examining the 19 cases as a group and looking for common problems or strengths in the techniques of costeffectiveness or cost-benefit analysis, OTA was able to better analyze the potential contribution that those techniques might make to the management of medical technology and health care costs and quality.

The second function of the case studies is to provide useful information on the specific technologies covered. The design and the funding levels of most of the case studies are such that they should be read primarily in the context of the associated overall OTA projects. Nevertheless, in many instances, the case studies do represent extensive reviews of the literature on the efficacy, safety, and costs of the specific technologies and **as** such can stand on their own as a useful contribution to the field.

Case studies are prepared in some instances because they have been specifically requested by congressional committees and in others because they have been selected through an extensive review process involving OTA staff and consultations with the congressional staffs, advisory panel to the associated overall project, the Health Program Advisory Committee, and other experts in various fields. Selection criteria were developed to ensure that case studies provide the following:

- examples of types of technologies by function (preventive, diagnostic, therapeutic, and rehabilitative);
- examples of types of technologies by physical nature (drugs, devices, and procedures);
- examples of technologies in different stages of development and diffusion (new, emerging, and established);
- examples from different areas of medicine (e.g., general medical practice, pediatrics, radiology, and surgery);
- examples addressing medical problems that are important because of their high frequency or significant impacts (e. g., cost);
- examples of technologies with associated high costs either because of high volume (for lowcost technologies) or high individual costs;
- examples that could provide information material relating to the broader policy and methodological issues being examined in the particular overall project; and
- examples with sufficient scientific literature.

Case studies either are prepared by OTA staff, are commissioned by OTA and performed under contract by experts (generally in academia), or are written by OTA staff on the basis of contractors' papers.

OTA subjects each case study to an extensive review process. Initial drafts of cases are reviewed by OTA staff and by members of the advisory panel to the associated project. For commissioned cases, comments are provided to authors, along with OTA's suggestions for revisions. Subsequent drafts are sent by OTA to numerous experts for review and comment. Each case is seen by at least *30*, and sometimes by 80 or more outside reviewers. These reviewers may be from relevant Government agencies, professional societies, consumer and public interest groups, medical practice, and academic medicine. Academicians such as economists, sociologists, decision analysts, biologists, and so forth, as appropriate, also review the cases.

Although cases are not statements of official OTA position, the review process is designed to satisfy OTA of each case study's scientific quality and objectivity. During the various stages of the review and revision process, therefore, OTA encourages, and to the extent possible requires, authors to present balanced information and recognize divergent points of view.

| Case Series       | J  | Case Study<br>Series<br>number  | Case study title; author(s);<br>OTA publication number <sup>b</sup>  |
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|                   | Formal Analysis, Policy Formulation, and End-<br>Stage Renal Disease;<br>Richard A. Rettig (OTA-BP-H-9 (1)) <sup>c</sup>   | 13 Cardiac Radionuclide Imaging and Cost<br>Effectiveness;<br>William B. Stason and Eric Fortess                    |  |
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capped People. 8Background Paper #2 to Technology and Handicapped People.

mation. bOriginal publication numbers appear in parentheses. 'The first 17 cases in the series were 17 separately issued cases in Background Paper #: Case Studies of Medical Technologies, prepared in conjunction with OTA's August 1980 report The Implications of Cost-Effectiveness Anal-vsis of Medical Technology.

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