VOL II

Gastrointestinal Haemorrhage

EDITED BY

P. S. HUNT MS FRACS

CLINICAL SURGERY INTERNATIONAL

VOL 11

Gastrointestinal Haemorrhage

CHRESCHILL VI (1946 STONE) Wedeel Dicker of Longoun Gr

Liszcigniech in the United States of A-I Liebzeicherhot, 1500 Brodf Aug 1849 by 1891 fled controlnes state bes and throughout the world

6 Longaga Group Limited 1985

EDITED BY

P. S. HUNT MS FRACS

Clinical Associate Professor of Surgery, Monash University, Melbourne, Australia All rights reserved. No pair of this published move by the molecula soft in a critical speed, or transmiss in an former by any means of transmissional pair to young former by any means of transmission of the proceedings of other recording or otherwise, without the price permission of the published (Churchill Livringwons, Kobert Structure, Manne, Barrer's Plage. Leath Wals, Flinburgh CH1 344).

First published 1-85

CHINE O 443 USOTO ?

Bruigh Labraty Caraloguein, in Publication Data, reasonnerunal hasmorthage. — (Clinical current: international, 1888, 026), 4422, v. 117

Gastromiesonal hemorrhis
 Flori, P.S. 11 Series

ibrary of Congress Catalogurg in Publication Data Visin epity nodes other

(Clinical surgery international, ISSN 0263-43

(1) lo

Granourestoral hemorouse—
 Granourestoral state—Surger
 Chaica surgery interneounal v. 11.

 L. Seines: Chaica surgery interneounal v. 11.

WEL CLIPSTE WILLIWE 143 G-25 RENGARIGATE 1986 617.43



EDINBURGH LONDON MELBOURNE AND NEW YORK 1986

CHURCHILL LIVINGSTONE Medical Division of Longman Group Limited

Distributed in the United States of America by Churchill Livingstone Inc., 1560 Broadway, New York, N.Y. 10036, and by associated companies, branches and representatives throughout the world

© Longman Group Limited 1985

All rights reserved. No part of this publication may be reproduced, stored in a retrieval system, or transmitted in any form or by any means, electronic, mechanical, photocopying, recording or otherwise, without the prior permission of the publishers (Churchill Livingstone, Robert Stevenson House, 1-3 Baxter's Place, Leith Walk, Edinburgh EH1 3AF).

First published 1985

ISBN 0 443 03074 X

British Library Cataloguing in Publication Data Gastrointestinal haemorrhage. — (Clinical surgery international, ISSN 0263-4422; v. 11)
1. Gastrointestinal hemorrhage
I. Hunt, P.S. II. Series
616.3'3 RC802

Library of Congress Cataloging in Publication Data Main entry under title:
Gastrointestinal haemorrhage.
(Clinical surgery international, ISSN 0263-4422; vol. 11)
Includes index.
1. Gastrointestinal hemorrhage—Surgery.
2. Gastrointestinal system—Surgery. I. Hunt, P. S. II. Series: Clinical surgery international; v. 11.
[DNLM: 1. Hemorrhage, Gastrointestinal.
W1 CL795U v.11/W1 143 G2564]
RD540.G377 1986 617'.43 85-13259

BDITED BY

C. S. HUNT MS FRACS

Clinical Associate Professor of Surgery, Monach University, Melbomme. Australia

Printed in Great Britain by Butler & Tanner Ltd, Frome and London CHURCHILL IVI EDANBURGH UND

L. B. Angorn

Department of Surgery, Efective of Medicine. University of Matal, Congella, Naral, South Africa

L. W. Baker

Department of Surgery, Faculty of Medican, University Maial, Congella, Natal, South Africa

David C. Carter

Professor of Sorgery, University Departmen of Surger Royal Infirmary, Glasgow, UK

Allstair Cowen MB BS MD FRACE

Consultant Gastrocinerologisi, Royal Brisbane Hospita Brisbane, Australia

V. Buncombe

Gäströinjosna.: Unsty Prince of Waler Hosp of Sydney Adstralia

O. James Garden

Research Fellow, University Department of Surgery, Royal Informaty, Glasgow, UK

mall M. I

Chargointestinal Unit, Prince of Wates Hospital, Sydney Australia

I. Mansky FRACP

Keader in Medicine and Physician to Gasus enterology Unit, Monash University, Prince Henry's Hospital, Melbourne, Victoria, Australia

Owen D. Harris FRACP

Department of Gostroenterology, Frinces, Alexandra Rospital, Woolloongabba, Oncensional, Austra.

H. S. Himal MD FRCS(C) FACE

Associate Professor of Surgery, University of Toronto, Attending, Surgeon, Toronto Western-Hospiral, Toronto, Ontario, Canada

John M. Holman Ir MD

Department of Physiology, Albany, Mudical College At any, New York, USA

P. M. Hunt MS PRACS

Olinical Associate Professor, of Surveyy. Department of burgery, Monagh University; Gastromestical Surgeon. Frince Henry's Flospital, Melbourne, Microria, Australia

In contrast to the postwar period to about 1970, the last decade has seen considerable advances in the management of gastrointestinal haemorrhage. This has arisen mainly from specialized attention to the problem, supported by advances in resuscitation and intensive care. Specialization has produced the attention to detail, timing of established treatment and experience necessary for successful clinical practice. Fibreoptic endoscopy has improved diagnostic accuracy, facilitating the introduction of planned management. As a consequence, morbidity and mortality have been reduced by early control of bleeding in high-risk cases.

The authors have been asked to write chapters on these aspects of management, as well as certain additional topics, including bleeding from the lower gastrointestinal tract. A more rounded view than usual has been sought, the intention being to describe each topic in the context of the author's clinical practice. As a consequence, some overlap will be found, particularly in relation to contentious issues. It has also been planned that chapters be based upon sound pathophysiological principles with speculation about future development. My thanks to the authors for their endeavours.

Melbourne, 1986 P.S.

Snore Hospiell, Sydney, Austrilia

Layron F. Rikkers Mil.

Department of Surgery, Conversity of U. h. 1914. Medicine, Salt Lake (2015) "Utah. USA 1914.

E W. Salzman

Harvand Medical School, Rail Sand-Hosput, Dorton Masker maetrs, USA

laith. C

J. centrel at Medicine, University of Sydnes, Point Son Share Hospitals, Sydney, Australia

Twille II Veomana MD PRACE

st Assistini in Medicine, University of Multioring Epartment of Medicine, Austra Marquial, Hubblewick action, Australia

Contributors

I. B. Angorn

Department of Surgery, Faculty of Medicine, University of Natal, Congella, Natal, South Africa

L. W. Baker

Department of Surgery, Faculty of Medicine, University of Natal, Congella, Natal, South Africa

David C. Carter

Professor of Surgery, University Department of Surgery, Royal Infirmary, Glasgow, UK

Alistair Cowen MB BS MD FRACP

Consultant Gastroenterologist, Royal Brisbane Hospital, Brisbane, Australia

V. Duncombe

Gastrointestinal Unit, Prince of Wales Hospital, Sydney, Australia

O. James Garden

Research Fellow, University Department of Surgery, Royal Infirmary, Glasgow, UK

J. M. Ham

Gastrointestinal Unit, Prince of Wales Hospital, Sydney, Australia

J. Hansky FRACP

Reader in Medicine and Physician to Gastroenterology Unit, Monash University, Prince Henry's Hospital, Melbourne, Victoria, Australia

Owen D. Harris FRACP

Department of Gastroenterology, Princess Alexandra Hospital, Woolloongabba, Queensland, Australia

H. S. Himal MD FRCS(C) FACS

Associate Professor of Surgery, University of Toronto; Attending Surgeon, Toronto Western Hospital, Toronto, Ontario, Canada

John M. Holman Jr MD

Department of Physiology, Albany Medical College, Albany, New York, USA

P. S. Hunt MS FRACS

Clinical Associate Professor of Surgery, Department of Surgery, Monash University; Gastrointestinal Surgeon, Prince Henry's Hospital, Melbourne, Victoria, Australia

Richard H. Hunt FRCP FRCP(C)

Professor, Head Division of Gastroenterology, McMaster University Medical Centre, Hamilton, Ontario, Canada

P. C. Johnson bezilaipeds mort vinian mesirs san

Harvard Medical School, Beth Israel Hospital, Boston, Massachusetts, USA

George W. Johnston MCh FRCS

Royal Victoria Hospital, Belfast, Ireland

Melvyn G. Korman MBBS PhD FRACP

Gastroenterology Unit, Prince Henry's Hospital, Melbourne, Victoria, Australia

Roger J. Leicester FRCS and availation bus

Consultant Surgeon, Head of Surgery, Royal Naval Hospital Haslar, Gosport, UK

Ronald A. Malt MD

Surgical Services, Massachusetts General Hospital; Department of Surgery, Harvard Medical School, Boston, Massachusetts, USA

Luc Michel MD

Surgical Unit, Mont-Godinne Academic Hospital; Department of Surgery, University of Louvain Medical School, Yvoir, Belgium

P. O'Brien

Department of Surgery, Flinders Medical Centre, Bedford Park, South Australia, Australia

D. W. Piper

Professor of Medicine, University of Sydney, Royal North Shore Hospital, Sydney, Australia

Layton F. Rikkers MD

Department of Surgery, University of Utah School of Medicine, Salt Lake City, Utah, USA

E. W. Salzman

Harvard Medical School, Beth Israel Hospital, Boston, Massachusetts, USA

D. Stiel

Lecturer in Medicine, University of Sydney, Royal North Shore Hospital, Sydney, Australia

Neville D. Yeomans MD FRACP

First Assistant in Medicine, University of Melbourne, Department of Medicine, Austin Hospital, Heidelberg, Victoria, Australia

要完整PDF请访问: www.ertongbook.com

Contents

- Natural history and mortality trends of acute upper gastrointestinal haemorrhage 1 D. W. Piper and D. Stiel
- Prospective study and computer evaluation of the management of gastrointestinal haemorrhage 13 H. S. Himal
- Clinical assessment and early endoscopic diagnosis 19
 Hansky
- 4. Anaesthesia and intensive care 27 G. Parkin
- Surgical management of bleeding chronic peptic ulcer 37
 P. S. Hunt
- Acute upper gastrointestinal haemorrhage in elderly patients 47
 J. M Ham and V. Duncombe
- 7. Mallory-Weiss syndrome 55 L. Michel and R. A. Malt
- 8. The pathogenesis, prevention and treatment of stress ulceration 60
 P. O'Brien
- 9. Drugs and alcohol and upper gastrointestinal haemorrhage 68
 N. D. Yeomans
- Modern use of balloon tamponade in reduction of mortality from bleeding oesophageal varices 77 M. G. Korman
- Injection sclerotherapy for oesophageal varices 83
 D. Harris
- 12. Variety of treatment for oesophageal varices and plan of management 92G. W. Johnston
- 13. Selective shunting in esophageal varices 105 J. M. Holman Jr and L. F. Rikkers

- 14. Endoscopic methods in prognosis and treatment of bleeding peptic ulcer 115
 O. J. Garden and D. C. Carter
- 15. Acute colonic haemorrhage 123 R. J. Leicester and R. H. Hunt
- 16. Chronic intestinal blood loss 135 A. Cowen
- 17. Haemobilia 143 L. W. Baker and I. B. Angorn
- Gastrointestinal haemorrhage in generalised haemorrhagic states 153
 P. C. Johnson and E. W. Salzman

Index 167

Natural history and mortality trends of acute upper gastrointestinal haemorrhage

D. W. PIPER and D. STIEL

Relevance of aetiology to mortality

The interpretation of mortality figures in patients with acute upper gastrointestinal haemorrhage is made difficult by the difference in frequency of the causal lesions and the failure of many authors to define the cause and the presence or absence of other risk factors when quoting mortality rates. It is clear that patients may be divided into three groups:

- 1. Those with bleeding due to peptic ulcer where the bleeding commenced prior to admission to hospital, i.e. peptic ulcer patients, where acute stress ulcers have been excluded. This group may be further subdivided into three groups with different mortalities, i.e. chronic gastric ulcer, chronic duodenal ulcer and acute ulcer.
- 2. Those whose bleeding commenced after admission to hospital, i.e. acute stress ulcer. Here a high mortality may exist due to other severe physical disease, i.e. severe trauma, major medical illnesses, burns, etc.
- 3. Bleeding due to oesophageal or gastric varices, carcinoma of the stomach and haematological diseases (i.e. purpura, leukaemia, etc.). In this group, the mortality of the acute episode of bleeding is added to the mortality of the primary lesion, e.g. liver disease, cancer, etc.

Morrallic in recess cories 1778-1982

Mortality rates of haematemesis with time

For the purposes of this discussion, three time periods will be considered:

- 1. Prior to World War II, which covers the period when blood transfusion was not available and when the heterogeneous nature of the causal lesions was poorly defined.
- 2. 1940-1975 which coincides with the development of resuscitative procedures, the introduction of widespread use of modern diagnostic techniques, the selective use of surgery and a general availability of antibiotics and other measures widely used in the treatment of the severely ill. During this period, too, a large series of adequately diagnosed patients were presented which defined the risk factors associated with acute upper gastrointestinal haemorrhage, especially relating to peptic ulcer.

This period saw the developments in the acute medical and surgical management of bleeding varices, notably the use of vasopressin infusions, oesophageal balloon tamponade, emergency portosystemic shunts and transoesophageal varix ligation.

3. 1976-1983. This encompasses the period which includes the best that has been possible in the diagnosis and treatment of acute upper gastro-intestinal haemorrhage. These advances include the widespread use of fibreoptic endoscopy, the better definition of the role of surgery and the treatment of patients in sophisticated intensive care wards.

Mortality due to upper gastrointestinal haemorrhage prior to 1940

well, several series of various beeding were

These data have been summarized by Allan & Dykes (1976) and their, and additional, information is presented in Table 1.1. It is seen that the mortality rate varies between 1% and 22%. Mortality from

Table 1.1 Mortality from acute upper gastrointestinal haemorrhage 1900-1940

rends of dentenan	Region 17.054	Period	Number of patients	Mortality rate (%)
Aitken (1934)	London	1929-1933	255	eant
Bulmer (1932)	Birmingham	1902-1926	467	10
Burger & Hartfall (1934)	London	1921-1930	137	21
Chiesman (1932)	London	1925-1931	1812	10.8
Christiansen (1934)	Copenhagen	1923-1932	289	7.9
Cullinan & Price (1932)	London	1925-1929	109	18
Davies & Nevin (1934)	London	1924-1933	391	21.5
Hellier (1934)	Leeds	1926-1932	202	13
Hurst (1924)	London	1911-1920	600	2.5
Hurst & Ryle (1937)	London	1919-1935	371	1.1
Meulengracht (1935)	Copenhagen	1932-1936	251	1.
Paterson (1924)	London	?-1924	1343	3.8

variceal haemorrhage was, however, considerably greater. Ratnoff & Patek (1942), in a series of 108 patients with bleeding varices treated prior to 1938, reported an immediate mortality of 33%, with 70% of patients dead within 12 months.

Data provided in Table 1.1 is limited by the failure of diagnostic facilities in that era to demarcate variceal from ulcer bleeding. This should not detract from the success of the series with low mortality because better diagnostic procedures would have demarcated the variceal group with its worse prognosis. The most common mortality rate is about 10%.

It is seen in three series in the period that there was a low mortality (3.8%, 1.1% and 1.0%). No reason is obvious for this apparent success.

Mortality due to acute upper gastrointestinal haemorrhage 1940-1975

Data relevant to this period are summarized in Table 1.2. It is seen that the mortality rate in all series was less that 11% and usually 5-9%. In many series, the mortality in the peptic ulcer subgroup and the whole series can be obtained from the reports and if so, both mortalities are presented.

As well, several series of variceal bleeding were reported. Over this period, the mortality rates from variceal haemorrhage in seven series ranged from 45% to 84%: 76% by Higgins (1947), 83% by Atik & Simeone (1954), 59% by Nachlas et al (1955), 74% by Cohn & Blaisdell (1958), 45% by Taylor & Jontz (1959), 76% by Merigan et al (1960), 84% by Orloff (1962). Similar high mortality rates were achieved

with the use of either intravenous or intra-arterial vasopressin: 93% by Merigen et al (1962), 64% by Brant et al (1972), 77% by Conn et al (1972), 50% by Marubbio et al (1972), 61% by Murray-Lyon et al (1973), 61% by Nussbaum et al (1974). Results with the use of balloon tamponade were equally disappointing: 47% by Reynolds et al (1952), 75% by Hamilton (1955), 82% by Conn (1958), 74% by Read et al (1960), 82% by Orloff (1962).

Orloff et al (1974) reported an early mortality rate of 52% in 115 consecutive patients undergoing emergency portacaval shunts, a similar figure to that obtained in an earlier series with transoesophageal varix ligation (46%) and substantially lower than the 83% observed in medically-treated patients (Orloff 1962). Other smaller studies support the finding of lower mortality in shunted patients, with an overall average mortality rate of 44% in 13 series summarized by Conn (1975). Interpretation of the results from these studies is complicated, however, by the considerable variation in mortality rates (23-71%) and by the presence of confounding variables such as heterogeneity regarding the aetiology of portal hypertension, the severity of liver diseasee and the precise timing of intervention.

Mortality in recent series 1976-1983

These series have the advantage that the bleeding site has often been defined and the mortality defined in each diagnostic group. The data for the major series of acute upper gastrointestinal haemorrhage as a group and also where in some series the ulcer

the mortality of the primary lesion, e.g., liver disease

Table 1.2 Mortality from acute gastrointestinal haemorrhage 1940-1975

Author Author	Region	Period	Number patients	of Mortality rate (%)
Allan & Dykes (1976)	Birmingham	1971-1973	300	(avery 9.7% × nsll.
Banning et al (1965)	Essex	1960-1965	223	(58.3%) 8 19 1918
Cates (1959)	Bristol	1953-1956	300	7%
			259	5%*
Cocks et al (1972)	London	1953-1962	1427	8.6%*
Coghill & Wilcox (1960)	London	1947-1958	325	CROTT SIG 13% S. mlora
Cotton (1973)	London	1971-1972	208	3.8%
	124-1426		109	3.7%
Duggan (1956)	Newcastle (Aus.)	1949-1954	363	8.3%
	180		219	8.7%
Fitzherbert & Epps (1950)	Sydney	1947-1949	124	7%
Fraenkel & Truelove (1955)	Oxford ,	1948-1952	377	5.6%*
Hellers & Ihre (1975)	Stockholm	1968-1969	149	20%
	101210		131	11%*
Jones (1947)	London	1940-1947	530	8%*
Jones (1956)	London		1764	7.9%
Jones (1961)	London	. 1	534	4.1%*
Large (1960)	Reading		488	5.9%*
Lewin & Truelove (1949)	Oxford	1938-1947	305	19%
20 20 20 20		_	252	17%*
MacCaig et al (1964)	London	1957-1960	559	7.7%*
Main (1964)	Falkirk	1952-1961	300	8.3%
			220	8.2%*
Needham & McConachie (1950)	Aberdeen	1941-1948	476	14%
Schiller et al (1970)	Oxford	1953-1967	2149	9%
Scott (1940)	Glasgow	1936-1940	110	13%
Smith (1945)	Glasgow	1934-1945	180	10%
Thomas & Rees, (1954)	Swansea	1938-1942	160	11.8%
		1942-1951	218	8.7%
Walls et al (1971)	Leeds	1967-1970	165	8%
Ward-McQuaid et al (1960)	Mansfield	1963-1956	200	4.5%*
and the		1956-1958	200	8%*
Waterson (1956)	Cambridge	1950-1952	154	9.7%

^{*}Series restricted to peptic ulcer.

subgroup has been defined separately, are indicated in Table 1.3. Again, the mortality in most series, where all cases of acute upper gastrointestinal haemorrhage are considered as a single group, lies in the range of 6-12%. The new statistic to appear is a series where, when the bleeding is restricted to ulcer, the mortality is less than 2% (Hoare et al 1979, La Brooy et al 1979, Kang & Piper 1980). Series where the low mortality in bleeding peptic ulcer has been achieved deserve comment, as it could be asked why a low mortality is possible in these series and not in others.

1. La Brooy et al (1979), at Central Middlesex Hospital, London, reported 109 patients with haemorrhage not due to varices, stress ulcer or cancer and only 1 death occurred. The mean age was 58 years with a range of 20 to 87 years; 13 required surgery. This outstanding result was achieved in

patients who started to bleed prior to admission to hospital, without malignancy and without severe liver or renal disease and without severe exsanguination.

- 2. Hoare et al (1979) in Birmingham, England, reported 66 patients with peptic ulcer, all over 45 years, in a trial assessing the effect of cimetidine on bleeding. The latter had no effect. Patients with serious liver and renal disease were excluded, as were those with haemorrhage secondary to other major diseases. Emergency surgery was needed in seven patients.
- 3. Kang & Piper reported 184 patients with acute upper gastrointestinal haemorrhage in the period-1976-77, with 18 deaths (9.8%). If varices and cancer were excluded, the mortality was 5%. Of 148 with bleeding not due to varices, cancer or stress-related ulcer, only 3 deaths were observed (2%): of the 14

Table 1.3 Mortality from upper gastrointestinal haemorrhage due to ulcer since 1975

	edani Z	egion		Period	Num	ber of		Mortality rate (%)
Allan & Dykes (1976)	Bi	irmingham	1071	1970-1973		300	(1975)	9.7%
Barer et al (1983)	N	ottingham		1980	GU		186	11%*
		ort		1982	DU		202	6%*
	1000				Erosi	ons	58	5%
	1			to he had a	Othe		230	6%
Brolin & Stremple (1982)	Pi	ttsburgh		1973-1977			(1) koo	12%*
Brown et al (1981)		ristol		1953-1956		287		5.5%*
. 87.3	964.			1974-1976		272		4.5%
Carstensen et al (1982)	C	openhagen	-04-1	1977-1978		88		10%
Dawson et al (1982)		rmingham		1981		151		6%
Dronfield et al (1977)		ottingham		1976		322		10%
Gilbert et al (1979)		SA		1210		800		9.4%
Himal et al 1978)		ontreal		1963-1971		630		12.5%
Timal et al 1976)	141	Ontreal		1973-1971	2	334		
Hann et al (1079)	D:	rmingham	Luthe 1	1979		66		6.7%
Hoare et al (1978)	LATI	Immgnam		1979		100		1.5%*
H1 (1082)	14	elbourne		1072 1077	T1		720	1%
Hunt et al (1983)	IVI	elbourne		1972–1977	Total		728	8.5%
1 201					DU		225	6.2%*
					GU		76	13.1%*
					Vario		58	34.5%
4.	1.1(4)				Acute		144	7.6%*
					Other		116	6.9%
				Same Language		iagnosi		2%
STAC				1977-1982	Total		588	5.8%
33.				Pietar	DU		219	2.7%*
	0.41				GU	1	113	6.2%*
					Vario	7	.59	16.9%
					Acute		55	1.8%*
		(AP.10)			Other		.126	4.8%
				e e e e e e e e e e e e e e e e e e e	No di	agnosi	s 27	3.7%
Kang & Piper (1980)	Sy	dney		1976-1977		148		2%
Kittang et al (1982)	Os	slo				90		. 0*
La Brooy et al (1979)	Lo	ondon		1979		101		1%*
Morgan et al (1977)	K	eighley		1975-1976		66		11%
Siddiqui et al (1979)	H	artlepool		1979		113	f at pai	8%
Vallon et al (1981)	Lo	ondon		1981		178		11%*
Vellacott et al (1982)	No	ottingham		1975-1977	GU		173	13.9%*
tion tring books a		is odw	trainent	1978-1980	GU		225	12.9%*
and their true servers				1975-1977	DU		224	9.4%*
e to the man Autorities	William VI			1978-1980	DU		260	9.8%*
Wetterfors et al (1982)	Li	nköping	char to	1976-1980	. Lanines	367	izen 15	10%
Wong et al (1980)		ong Kong		To hamer or	A.S0.7	467	n alne	6.4%*

^{*}Series restricted to peptic ulcer.

who started to bleed after admission, i.e. acute stress ulcers, 28% died.

The three units where these results have been obtained have only one feature in common — they are departments that have a long history of interest in the treatment of acute upper gastrointestinal haemorrhage and it is likely the good results reflect this interest and expertise. The role of special units available for the treatment of haematemesis and melaena has been emphasized by Hunt and his colleagues (1983).

The progress in the treatment of bleeding ulcer has not been as obvious when variceal bleeding is reviewed. Several studies since 1975, including some prospective controlled series, have again failed to show reduced mortality with the use of vasopressin infusions: 67% by Johnson & Widrich (1976), 49% by Kaufman et al (1977), 48% by Sherman et al (1979), 46% by Johnson et al (1977), 70% by Chojkier et al (1979). Fogel et al (1982) reported an in-hospital mortality rate of 45%, which was not influenced by randomization to intravenous vasopressin or placebo

therapy. Graham & Smith (1983), in a study of 85 patients with alcoholic cirrhosis and endoscopically proven bleeding varices, reported an early mortality in medically treated patients of 42% and, in a study by Kang & Piper (1980), 63% died in hospital. In one of the few studies examining the course of patients with variceal haemorrhage due to non-alcoholic liver disease, Toouli et al (1983) reported a mortality of 63% in 8 patients undergoing emergency portacaval anstamosis. In an extension of previous studies, Orloff et al (1980) reported a 42% early mortality in 180 unselected cirrhotics undergoing shunts.

The old technique of endoscopic injection sclerotherapy has be reintroduced in recent years, and its efficacy has been reviewed (Allison 1983). In reported series of injection sclerotherapy for acute variceal haemorrhage, mortality rates ranging from 13% to 50% were observed: 13% by Denck (1971), 50% by Raschke & Paquet (1973), 16% by Johnston & Rodgers (1973), 39% by Terblanche et al (1981), 32% by Palani et al (1981), 21% by Barsoum et al (1982), 31% by Kjargaard et al (1982), 14% by Alwmark et al (1982).

In an 8-year (1972-1980) prospective study of 91 patients with bleeding varices, Hunt et al (1982) reported an overall in-hospital mortality rate of only 26%, falling to 18% in the latter 4-year period of the study. Subsequent data from the same unit, shown in Table 1.4, confirm this trend (Hunt et al 1983). The authors attribute these excellent results to adherence to a protocol which comprises early endoscopy, close medical-surgical liaison, management in an intensive therapy unit, extensive use of balloon tamponade and selective employment of decompressive surgery. This study, those of Orloff and his colleagues (1974) and those summarized by Allison (1983), provide

evidence that improved survival can be achieved in patients with variceal haemorrhage.

Has the mortality of acute upper gastrointestinal haemorrhage changed?

Peptic ulcer

A survey of the results of various series showed gross divergence in the mortality rates observed over periods of time. It is seen (Table 1.1-1.3) that a mortality rate for haematemesis overall is usually 8-10% and it could be claimed, therefore, that it has not changed. To state this is to ignore the fact that haematemesis is a disease of the elderly and in all communities the elderly population is increasing; it also ignores the different mortality associated with three common causes of bleeding, i.e. chronic ulcer, acute ulcer and varices. Kang & Piper (1980) have shown that, while there is little change in the overall acute upper gastrointestinal haemorrhage mortality rate in the published series in the English literature from 1939-1977, if the age-related mortality rates are plotted for non-variceal, non-cancer, non-stressrelated ulcer groups, there is a marked reduction in the mortality of those over 60 years in the 12 published series where sufficient data were published to make the comparison possible (Fig. 1.1). Any change in the total series is obscured by the low mortality in the younger age group.

The second series of observations relevant to improved prognosis of acute upper gastrointestinal haemorrhage in the past decade originated from the detailed studies of Hunt and his colleagues (Hunt et al. 1983).

Table 1.4 Mortality rates due to acute upper gastrointestinal haemorrhage at Prince Henry's Hospital, Melbourne over two 5-year periods, 1972-1977 and 1977-1982, in relation to cause of bleeding (NS difference was not significant).

	CARLOTTE BEEF	1972-1977	1977–1982		Statistical	
	Cases	Deaths	Cases	Deaths	comparison	
Total	728	62 (8.5)	588	34 (5.8)	TO NS ATOT FACT	
Duodenal ulcer	225	14 (6.2)	219	6 (2.7)	NS	
Gastric ulcer	76	10 (13.1)	113	7 (6.2	p<0.05*	
Desophageal varices	58	20 (34.5)	59	10 (16.9)	there was a sign fice	
Prosive gastritis	144 1089	3.11 (7.6)	28 55 0000	1 (1.8)	NS NS	
Mallory-Weiss syndrome	54	1 (1.8)	45	3 (6.6)	NS ,	
Other causes	62	7 (11.3)	81	3 (3.6)	NS MS MOU	
No diagnosis	98 2000	2 (2.0)	27	231701 (3.7)	the please grown and	

^{*}For chronic peptic ulcer.

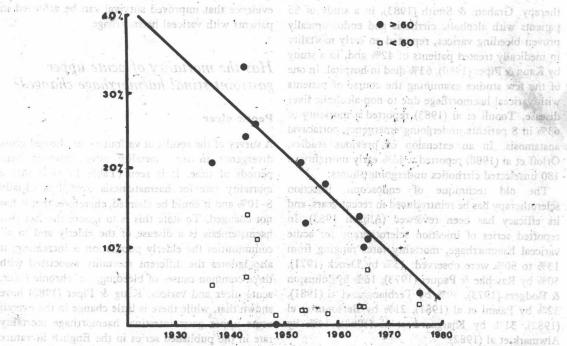


Fig. 1.1 Age-related mortality from selected series of acute upper gastrointestinal haemorrhage (due to ulcer excluding acute stress ulcer). (From Kang & Piper 1980).

728 patients in the period 1972-1977 were compared in each diagnostic group with 588 patients in the period 1977-1982. These results are shown in Table 1.4; it is seen there was an approximate halving of the mortality observed over the whole series and when patients with duodenal ulcer, gastric ulcer or oesophageal varices are considered separately. This drop reached statistical significance only in the gastric ulcer group. The improvement is attributed by the investigators to the establishment of a special unit devoted to the treatment of acute upper gastro-intestinal haemorrhage, adequate resuscitation and a defined plan of manaement.

A similar trend in improvement in mortality with time was observed by Himal et al (1978) in Montreal, comparing two series treated in the periods 1963-1971 and 1973-1976. Using a more aggressive surgical approach in the latter group, it was observed there was a significant drop in overall mortality with time (12.5% versus 6.7% mortality). This drop was not present in the variceal group but only present in the ulcer group and in the total series.

Evidence that the mortality of ulcer has not changed is provided by the studies of Schiller et al (1970) in Oxford. They studied 2149 emergency admissions because of haematemesis or melaena during 3 successive 5-year periods in the interval 1953-1967 inclusive. The fatality rate remained virtually constant throughout the period, in spite of changes in diagnostic measures and management. During this period, the age and sex distribution was similar and the chief diagnostic groups remained identical in size. The overall mortality rate was 8.9%.

The striking feature of the mortality of haematemesis and melaena is the marked variation observed in different series. This is obvious in the three time periods studied and no explanation is available that adequately accounts for this.

Varices

Graham & Lacey, in reporting similar mortality rates in 1981 to those described in series published 27 and 30 years previously, concluded, 'it is apparent that we have learned much and accomplished little'. While the overall mortality from variceal haemorrhage appears to have altered little since the first substantive

report by Ratnoff & Patek in 1942, some investigations suggest improved survival with the use of portacaval anastamosis (Orloff et al 1980), injection sclerotherapy (reviewed by Allison 1983) or a protocol of intensive medical and surgical managements (Hunt et al 1982). We should be guided by studies such as these if any impact is to be made on the depressing mortality statistics relating to variceal haemorrhage.

Factors that could have been expected to produce a decline in mortality of acute upper gastrointestinal haemorrhage

The benefit conferred by new measures has not been demonstrated statistically. Such studies are inhibited by the ethical problems associated with the relevant clinical trials and by the changing natures of the exposed population, the proportion of elderly increasing with their recognized higher mortality.

patient who has not confired blood. In a

Advances in treatment of bleeding peptic

These include a boold toll syllagen startiggs pirities

Blood transfusion social ni the so blooms it inneited

As Grossman et al (1950) have stated, there is no clinical trial data proving that blood transfusion decreases the mortality of haemorrhage from peptic ulcer. However, to deny its use, i.e. failure to transfuse a patient with oligaemic shock, defeats the logic of medicine and physiology.

Definition of risk factors resulting in death in patients with acute upper gastrointestinal haemorrhage due to peptic ulcer

These were defined by Avery Jones (1947, 1956), Truelove and his colleagues (Schiller et al 1970) and Allan & Dykes (1976) and have been summarized by Balint (1977) (Table 1.5). Increasing risk is associated with:

Table 1.5 Factors influencing mortality in patients with bleeding peptic ulcer (based on data reproduced, with thanks, from Balint J A et al 1977).

edying from chrotara been defined (Parsons	o sir dad Mortality org
	Aldridge 1951). These inqui
chronic GII	16, 9.3
chronic DU	1. latic, lons based on
acute ulcers tag only and	mentidet d above and include
10	years of age with recurrent and ways have a chronic ulce
10 37	
vidence indicater-09 iv	2. 2. 18the where the e
reskel with as 08 Cre	bleeding, from a large rec
Age of patient and typ of u	bacmcolepamic changes. In
cinomic oc 400	
	has been defined by S and
chronic DU < 60	44.0
, ,,	
acute ulcer < 60 > 60	a: 177 in blood pressure g
Magnitude of bleed systolic BP	falk in central venous within 15 minutes.
	b. Fresh haemaremesis or i
80-99 mmHg	18
< 80 mmHg	mi pet hour
	c. Continuous bleeding re
single	The best aver 48 hours
recurrent	10, 21.5
continuous	30

1. Chronic ulcer rather than acute, and chronic gastric ulcer has a higher mortality than cronic duodenal ulcer.

Role of endoscopy in charges in mortality of the

- 2. The age of the patient, mortality rising rapidly in those over 60 years.
 - 3. Continued or recurrent bleeding.
 - 4. Those with other serious disease.
- 5. The presence of persistent ulcer pain after admission.

Factors that do not influence the immediate mortality are the length of ulcer history and history of previous bleeding (Lewin & Truelove 1949).

As is seen from the above data, the patient who is over 60 years of age, has a chronic ulcer and who bleeds recurrently or continuously after admission is higher risk. Also in the high risk group is the person with acute ulcer associated with other serious disease, i.e. acute stress ulcer (Hubert et al 1980, Kang & Piper 1980). The definition of risk factors is relevant in that if more than one is present, the treatment of that patient in an intensive care ward of a major hospital is mandatory.

Indications for emergency surgery

Mindful of the factors that place the patient in the group with high risk of dying from bleeding, indications for surgery have been defined (Parsons & Aldridge 1951). These include:

- 1. Indications based on bad prognostic features mentioned above and includes the patients over 60 years of age with recurrent or continuous bleeding and who have a chronic ulcer.
- 2. Those where the evidence indicates free bleeding from a large vessel with associated haemodynamic changes. This group of indications has been defined by Semb & Myren (1974) and includes:
 - a. Fall in blood pressure greater than 50 mmHg or fall in central venous pressure of 5 cm H₂O within 15 minutes.
 - b. Fresh haematemesis or melaena greater than 600 ml per hour.
 - Continuous bleeding requiring more than 4000 ml blood over 48 hours.

Role of endoscopy in changes in mortality of bleeding peptic ulcer

Chromic ulter rather than acute

There are several studies that have shown that endoscopy as a procedure has not reduced the mortality rate of acute upper gastrointestinal haemorrhage (Dronfield et al 1977, Eastwood 1977, Graham 1980). A fall is not expected unless the information so obtained is used with profit in the utilization of the several therapeutic measures applicable to the control of bleeding. It is considered endoscopy serves an essential role in the management of the patient with acute upper gastrointestinal haemorrhage. Firstly, it enables a definition of the bleeding lesion which may influence immediate treatment, i.e. if a bleeding lesion is a varix and not an ulcer. Secondly, the demonstration of a chronic ulcer may provide one of the indications for surgery in the elderly patient who bleeds continuously or recurrently (qv). Thirdly, by the demonstration of an acute lesion, it may shorten the patient's stay in hospital. Fourthly, it may show signs of recent haemorrhage which will provide a guide to the risk of further bleeding. Fifthly, it allows the use of

interventive techniques such as photocoagulation or diathermy which may have a role in the haemostasis of actively bleeding ulcers (Swain et al 1981).

protocol of intensive medical and survicel manage

sindicarsulds as these if any impact is to be made on

Nasogastric suction a SW. (\$801 Is to failt) among

This involves the passage of a nasogastric tube as soon as haemorrhage is suspected. The patient is placed on continuous suction and 100 ml water drunk each hour to facilitate aspiration. This serves the dual purpose of keeping the stomach free of blood clot and, more importantly, of detecting the presence of continuous or recurrent bleeding which, in appropriate circumstances, may constitute an indication for emergency surgery. It enables bleeding to be detected long before it is obvious from the haemodynamic changes as indicated by a rise in the pulse rate, a fall in blood pressure or central venous pressure. Most authorities recommend its use (Alexander Williams 1975, Chandler 1953, Wetterfors et al 1983). Gastric aspiration of blood also establishes the fact that melaena is due to upper gastrointestinal bleeding in a patient who has not vomited blood. In one study of the validity of this method in indicating gastroduodenal bleeding involving over 1000 patients, only 1% of patients with an upper gastrointestinal site of bleeding proximal to the ligament of Trietz had a gastric aspirate negative for blood and these were all duodenal ulcers (Luk et al 1979).

Unless it is causing marked discomfort to the patient, it should be left in place for 24 hours.

Intensive care ward observation

Every patient with proven or suspected acute upper gastrointestinal haemorrhage should be nursed in an intensive care ward for the first 48 hours at least. This environment enables careful observation of the patient's haemodynamic state. It is essential for the bad-risk patient. In this environment, central venous monitoring may be used.

decreases the mostality of haemorribage from per

Central venous monitoring

The nature of this assessment has been established by the work of Andersen (1970). He found this technique was of value in detecting episodes of major arterial haemorrhage and as a guide to the patient's transfusion requirements. As in the case of blood transfusion, its value has not been assessed using double blind clinical trials.

Advances in the treatment of bleeding varices

Blood transfusion when her the transfer and the Landaud

Similar principles apply here as to peptic ulcer (qv). In addition, administration of clotting factors is a logical adjunct in patients with hepatic coagulopathy.

Risk factors and desired and supplied by the hi missing one

Patients with bleeding varices may die from exsanguination, from the consequence of liver failure, such as encephalopathy or from sepsis. Logic, therefore, dictates that patients in Child's group C should fare worse than those in Child's group A or B. This seems to be borne out in most series (Olsson 1972). Other factors influencing survival have been described by Graham & Smith (1981). There is debate as to whether prognosis is better in patients with non-alcoholic liver disease than in those with alcoholic cirrhosis (Woodward & Webster 1972, Toouli et al 1983).

Endoscopy the self-tention of the self-tention

Haematemesis in patients known to have varices emanates from non-variceal sources in over one-third

Himsel H S. Perrault C. Masbi R 1979 Daper gestrointestinal

Surgery 84: 448-452

bedannien de

Heller's G. Thre T 1975 Importance of charges to usery diagno

REFERENCES

Aitken R S 1934 The treatment of profuse bleeding from stomach and duodenum. Lancet i: 839-842

Alexander Williams J 1975 Surgical treatment. In: Topics in gastroenterology 3, Blackwell Scientific Publications, Oxford, p 53-68

Allan R, Dykes P 1976 A study of the factors influencing mortality rates from gastrointestinal haemorrhage. Quarterly Journal of Medicine XLV: 533-550

Allison J G 1983 The role of injection sclerotherapy in the emergency and definitive management of bleeding oesophageal varices. Journal of the American Medical Association 249: 1484-1487

of cases (Novis et al 1976). Hence early endoscopic assessment of the bleeding point is vital if rational therapy aimed at reducing mortality is to be undertaken. Injection sclerotherapy may be performed at the same time, if desired.

Medical and surgical intervention to the state of the sta

The contributions to the mortality rate of vasopressin administration, balloon tamponade, injection sclerotherapy and surgical intervention have been discussed in detail (qv).

Brant B. Rösch I. Krippachne W. W. 1972 Experences with

Intensive care ward observation and will make the remarks

Similar principles apply as outlined for patients with bleeding peptic ulcer.

general memorrhage. A randomised controlled trail

Scandinavian Journal of Gastroenterology 15: 103-105

Conclusion

Where valid data for comparison exist, there is evidence that the mortality of acute upper gastro-intestinal haemorrhage from ulcers and varices has improved with time. This improvement has been predominantly in the ulcer group and here it has been restricted to those over 60 years of age. Moreover, this improvement has only been apparent in series from special units devoted to the care of acute upper gastrointestinal haemorrhage where full use is made of intensive care procedures and selective and skilled surgery.

Alwmark A, Bangmark S, Borjesson B, Gullstrand P, Joelsson B 1982 Emergency and long-term transesophageal sclerotherapy of bleeding esophageal varices. Scandinavian Journal of Gastroenterology 17: 409-412

the ligh massive hemorrhage. Surgery, Gynecology and

with citrhosis of the liver with esophageat varices following

Andersen D 1970 The use of measurement of central venous pressure in the selection of patients with massive gastroduodenal haemorrhage for emergency operation.

Scandinavian Journal of Gastroenterology 5: 25-32

Atik M, Simeone F 1954 Massive gastrointestinal bleeding: a study of 296 patients at City Hospital of Cleveland. Archives of Surgery 69: 355-365

Balint J A, Sarfel I J, Fried M B 1977 Gastrointestinal bleeding.

Diagnosis and management. J Wiley and Sons, New York, p
63

Banning A, Baron A, Kopelman H, Lam K L, Warren P 1965
Bleeding peptic ulcer. British Medical Journal ii: 781-784

Barer D, Oglivie A, Henry D, Dronfield M, Coggan D, French S et al 1983 Cimetidine and tranexamic acid in the treatment of acute upper gastrointestinal tract bleeding. New England Journal of Medicine 308: 1571-1575

Barsoum M'S, Bolous F I, El-Rooby A A, Rizk-Allah M A, Ibrahim A'S 1982 Tamponade and injection sclerotherapy in the management of bleeding oesophageal varices. British

Journal of Surgery 69: 76-78

Brant B, Rösch J, Krippaehne W W 1972 Experiences with angiography in diagnosis and treatment of acute gastrointestinal bleeding of various etiologies. Preliminary report. American Surgeon 176: 419-434

Brolin R E, Stremple J F 1982 Emergency operation for upper gastrointestinal haemorrhage. The American Surgeon 48:

302-308

- Brown S G, Salmon P R, Brown P, Read A E 1981 Upper gastrointestinal haemorrhage. Journal of the Royal College of Physicians of London 15: 265-268
- Bulmer F 1932 Mortality from haematemesis. A supplementary analysis. Lancet ii: 720-722
- Burger G, Hartfall S J 1934 Haumatemusis in peptic ulcer. Guys Hospital Reports 84: 197-209
- Carstensen H E, Bülow S, Hart-Hansen O, Hamilton Jakobsen B, Krarup Pedersen T et al 1980 Cimetidine for severe gastroduodenal haemorrhage. A randomised controlled trial. Scandinavian Journal of Gastroenterology 15: 103-105
- Cates J E 1959 A review of 300 patients with haematemesis or melaena. British Medical Journal i: 206-211
- Chandler G N, Watkinson G 1953 Gastric aspiration in haematemesis. Lancet ii. 1170–1175
- Chiesman W E 1932 Mortality of severe haemorrhage from peptic ulcers. Lancet ii: 722-723
- Chojkier M, Groszmann R J, Atterbury C E 1979 A controlled comparison of continuous intraarterial and intravenous infusions of vasopressin in hemorrhage from esophageal varices. Gastroenterology 77: 540-546
- Christiansen T 1934 cited by Meulengracht E 1935 Treatment of haematemesis and melaena with the mortality. Lancet i: 1220-1222
- Cocks J R, Desmond A M, Swynnerton B F, Tanner N C 1972 Partial gastrectomy for haemorrhage. Gut 13: 331-340
- Coghill N F, Willcox R G 1960 Factors in the prognosis of bleeding chronic gastric and duodenal ulcers. Quarterly Journal of Medicine 29: 575-596
- Cohn R, Blaisdell F W 1958 The natural history of the patient with cirrhosis of the liver with esophageal varices following the first massive hemorrhage. Surgery, Gynecology and Obstetrics 106: 699-701
- Conn H O 1958 Hazards attending the use of esophageal tamponade. New England Journal of Medicine 259: 701-707
- Conn H O 1975 Cirrhosis. In: Schiff L (ed) Diseases of the liver, 4th edn. J B Lippincott, Philadelphia, ch 26, p 876
- Conn H O, Ramsby G R, Storer E H 1972 Selective intraarterial vasopressin in the treatment of gastrointestinal hemorrhage. Gastroenterology 63: 634-645
- Cotton P B, Rosenberg M T, Waldran R P L, Axon A T R
 1973 Early endoscopy of oesophagus, stomach and duodenal
 bulb in patients with haematemesis and melaena. British
 Medical Journal ii: 505-509
- Cullinan E R, Price R K 1932 Haematemesis following peptic ulceration. Prognosis and treatment. St Bartholomew's Hospital Report 65: 185-213

- Davies T A L, Nevin R W 1934 Prognosis of haematemesis. A statistical review. British Medical Journal ii: 858-859
- Dawson J, Cockel R 1982 Ranitidine in acute upper gastrointestinal haemorrhage. British Medical Journal ii: 476-477
- Denck H 1971 Endoesophageal sclerotherapy of bleeding esophageal varices. Journal of Cardiovascular Surgery 12: 146
- Dronfield M W, McIllmurray R, Ferguson R, Atkinson M, Langman M J S 1977 A prospective, randomised study of endoscopy and radiology in acute upper gastrointestinal tract bleeding. Lancet j: 1167-1169

Duggan J M 1956 Haematemesis and melaena: A survey. Medical Journal of Australia 2: 941-949

- Eastwood G.L. 1977 Does early endoscopy benefit patients with active upper gastrointestinal bleeding? Gastroenterology 72: 737-739
- Fitzherbert J C, Epps R G 1950 Management of haematemesis and melaena. Medical Journal of Australia 1: 490-499
- Fogel M R, Knauer C M, Andres L L, Mahal A S, Stein D E T, Kemeny M J et al 1982 Continuous intravenous vasopressin in active upper gastrointestinal bleeding. A placebo controlled study. Annals of Internal Medicine 96: 565-569
- Fraenkel G J, Truelove S C 1955 Haematemesis, with special reference to peptic ulcer, British Medical Journal i: 999-1002
- Gilbert D A, Silverstein F E, Tedesco F J, Buenger N K, 677.
 members of the ASGE 1979 Prognosis of upper
 gastrointestinal bleeding. Preliminary results of ASGE
 National Bleeding Survey. Gastroenterology 76: 1138
 (abstract)
- Graham D Y 1980 Limited value of early endoscopy in the management of acute upper gastrointestinal bleeding.

 Prospective controlled trial. American Journal of Surgery 140: 282-290
- Graham D Y, Smith J L 1981 The course of patients following variceal hemorrhage. Gastroenterology 80: 800-809
- Hamilton J E 1955 Management of bleeding esophageal varices associated with cirrhosis of the liver. Annals of Surgery 141: 637-647
- Hellers G, Ihre T 1975 Importance of changes to early diagnosis and surgery in major upper gastrointestinal haemorrhage.

 Lancet ii: 1250-1251
- Hellier F F 1934 Actiology and mortality rate of haemateinesis

 Lancet ii: 1271-1274
- Higgins H W Jr 1947 The esophageal varix; A report of 115 cases. American Journal of the Medical Sciences 214: 436-441
- Himal H S, Perrault C, Mzabi R 1979 Upper gastrointestinal haemorrhage: Aggressive management decreases mortality. Surgery 84: 448-452
- Hoare A.M, Bradby G U H, Hawkes C T 1979 Cimetidine in bleeding peptic ulcer. Lancet ii: 671-673
- Hubert J P, Kiernan P D, Welch J S, Remine W H, Beahrs O H 1980 The surgical management of bleeding stress ulcers. Annals of Surgery 191: 672-679
- Hunt P S, Francis J K, Hansky J, Hillman H, Korman M G, McLeish J et al 1983 Reduction in mortality from upper gastrointestinal haemorrhage. Medical Journal of Australia 2: 552-555
- Hunt P S, Korman M G, Hansky J 1982 Management of bleeding oesophageal varices. An eight wear prospective study, Medical Journal of Australia & J21-124
- Hurst A 1924 The treatment of severe gastric and duodenal haemorrhage, Lancet i: 1095-1098

- Hurst A F, Ryle J J 1937 The measure of mortality and treatment of haemorrhage in gastric and duodenal ulcer. Lancet i: 1-6
- Ivy A C, Grossman M I, Bachrach W H 1950 Peptic ulcer. The Blakiston Company, Philadelphia, Toronto, p 937
- Johnston G W, Rodgers H W 1973 A review of 15 years experience in the use of sclerotherapy in the control of acute haemorrhage from oesophageal varices. British Journal of Surgery 60: 797-800
- Johnson W C, Widrich W C 1976 Efficacy of selective splanchnic arteriography and vasopressin perfusion in diagnosis and treatment of gastrointestinal hemorrhage. American Journal of Surgery 131: 481-489
- Johnson W C, Widrich W C, Anself J E, Robbins A H, Nabseth D C 1977 Control of bleeding varices by vasopressin: A prospective randomized study. Annals of Surgery 186: 369-376
- Jones F A 1947 Haematemesis and melaena with special reference to bleeding peptic ulcers. British Medical Journal ii: 477-482
- Jones F Avery 1956 Haematemesis and melaena with special reference to causation and to the factors influencing the mortality from bleeding peptic ulcers. Gastroenterology 30: 166-190
- Jones F Avery 1961 In: Mairgot R (ed) Abdominal operations, 4th edn. Lewis, London, p 286
- Kang J Y, Piper D W 1980 Improvement in mortality rates in bleeding peptic ulcer. Medical Journal of Australia 1: 213-215
- Kaufman S L, Harrington D P, Barth K H, Maddrey W C, White R I Jr 1977 Control of variceal bleeding by superior mesenteric artery vasopressin infusion. American Journal of Roentgenology 128: 567-569
- Kittang E, Aadland E, Øyen D, Holm, H, Berstad A, Pytte R et al 1982 Effect of peroral antacid treatment in patients with acute upper gastrointestinal haemorrhage: A randomised controlled trial. Scandinavian Journal of Gastroenterology 17 (supplement 75): 109-112
- Kjaergaard J, Fischer A, Miskowiak J, Lindahl F, Baden H 1982 Sclerotherapy of bleeding esophageal varices. Scandinavian Journal of Gastroenterology 17: 363-367
- La Brooy S J, Misiewicz J J, Edwards J, Smith S J, Haggie S J, Libman L et al 1979 Controlled trial of cimetidine in upper gastrointestinal haemorrhage. Gut 20: 892-895
- Large J M 1960 Gastroduodenal haemorrhage as a surgical emergency. British Medical Journal i: 932-935
- Lewin D C, Truelove S 1949 Haematemesis with special reference to chronic peptic ulcer. British Medical Journal i; 383
- Luk G D, Bynum T E, Hendrix T R 1979 Gastric aspiration in localisation of gastrointestinal haemorrhage. Journal of the American Medical Association 241: 576-578
- MacCraig J N, Strange S L, Norris T St M 1964 Haemorrhage from the upper gastrointestinal tract. Gut 5: 136-141
- Main R G 1964 Haematemesis in a peripheral Scottish hospital. Scottish Medical Journal 9: 152-161
- Marubbio A T Jr, Lombardo R P, Holt P R 1973 Control of variceal bleeding by superior mesenteric artery pitressin perfusions — complications and indications. American Journal of Digestive Diseases 18: 539-543
- Merigan T C Jr, Hollister R M, Gryska P F, Starkey G W B, Davidson C S 1960 Gastrointestinal bleeding with cirrhosis: Study of 172 episodes in 158 patients. New England Journal of Medicine 263: 579-585

- Merigan T C Jr, Plotkin G R, Davidson C S 1962 Effect of intravenously administered posterior pituitary extract on hemorrhage from bleeding esophageal varices. A controlled evaluation. New England Journal of Medicine
- Meulengracht E 1935 Treatment of haematemesis and melaena with food. Lancet ii: 1220-1222
- Morgan A G, McAdam W A F, Walmsley G L, Jessop A, Horrocks J C, de Dombal F T 1977 Clinical findings, early endoscopy and multivariate analysis in patients bleeding from the upper gastrointestinal tract. British Medical Journal ii: 237-240
- Murray-Lyon I M, Pugh R N H, Nunnerley H B, Laws J W, Dawson J L, Williams R 1973 Treatment of bleeding oesophageal varices by infusion of vasopressin into the superior mesenteric artery, Gut 14: 59-63
- Nachlas M M, O'Neil J E, Campbell A J A 1955 The life history of patients with cirrhosis of the liver and bleeding oesophageal varices. Annals of Surgery 141: 10-23
- Needham G D, McConachie J A 1950 Haematemesis and melaena. British Medical Journal ii: 133-138
- Northfield T C, Smith T 1970 Central venous pressure in clinical management of acute gastrointestinal bleeding. Lancet ii: 584-586
- Novis B H, Duys P, Barbezat G O, Clain J, Bank S, Terblanche J 1976 Fibreoptic endoscopy and the use of the Sengstaken tube in acute gastrointestinal haemorrhage in patients with portal hypertension and varices. Gut 17: 258-263
- Nusbeam M, Younis M T, Baum S, Blakemore W S 1974
 Control of portal hypertension. Selective mesenteric arterial
 infusion of vasopressin. Archives of Surgery 108: 342-347
- Olsson R 1972 The natural history of oesophageal varices. A retrospective study of 224 cases with liver cirrhosis.

 Digestion 6: 65-74
- Orloff M J 1962 A comparative study of emergency transesophageal ligation and non-surgical treatment of bleeding esophageal varices in unselected patients with cirrhosis. Surgery 52: 103-116
- Orloff M J, Bell R H, Hyde P V, Skivolocki W P 1980 Long-term results of emergency portacaval shunt for bleeding esophageal varices in unselected patients with alcoholic cirrhosis. Annals of Surgery 192: 326-340
- Orloff M J, Chandler J G, Charters A C, Condon J A, Grambort D E, Modafferi T R et al 1974 Emergency portacaval shunt for bleeding esophageal varices. Prospective study in unselected patients with alcoholic cirrhosis. Archives of Surgery 108: 293-299
- Palini C K, Abuabara S, Kraft A R, Jonasson O 1981 Endoscopic sclerotherapy in acute variceal hemorrhage. American Journal of Surgery 141: 164-168
- Parsons K O, Aldridge L W 1951 Partial gastrectomy in the treatment of bleeding peptic ulcer. British Journal of Surgery 38: 370-378
- Paterson H J 1924 Discussion on the treatment of severe gastric and duodenal haemorrhage. Proceedings of the Royal Society of Medicine 17: 1-22
- Raschke E, Paquet K T 1973 Management of hemorrhage from esophageal varices using the esophagoscopic sclerosing method. Annals of Surgery 177: 99-102
- Ratnoff O D, Patek A J 1942 The natual history of Laennec's cirrhosis of the liver. An analysis of 386 cases. Medicine 21: 207-268
- Read A E, Dawson A M, Kerr D N S, Turner M D, Sherlock S 1960 Bleeding oesophageal varices treated by oesophageal compression tube. British Medical Journal i: 227-231