REVIEW ARTICLE

Consent and anaesthetic risk

K. Jenkins and A B. Baker*

Department of Anaesthetics, University of Sydney, Royal Prince Alfred Hospital, Missenden Road, Camperdown, Sydney, NSW 2050, Australia

Summary

The incidences of mortality and morbidity associated with anaesthesia were reviewed. Most of the published incidences for common complications of anaesthesia vary considerably. Where possible, a realistic estimate of the incidence of each morbidity has been made, based on the best available data. Perception of risk and communication of anaesthetic risk to patients are discussed. The incidences of anaesthetic complications are compared with the relative risks of everyday events, using a community cluster logarithmic scale, in order to place the risks in perspective when compared with other complications and with the inherent risks of surgery. Documentation of these risks and discussion with patients should allow them to be better informed of the relative risks of anaesthetic complications. Depending on specific comorbidities and the severity of operation, these risks associated with anaesthesia may increase for any one individual.

Keywords Anaesthesia; risk. Complication; incidence, mortality, morbidity, perioperative.

Correspondence to: A. B. Baker E-mail: bbaker@usyd.edu.au

*Present address: North Bristol NHS Trust, Southmead Hospital,

Bristol, UK

Accepted: 26 May 2003

'That which is necessary is never a risk' Paul de Gondi. The legal interpretation of the acceptable practice of medicine was changed markedly in Australia by the Rogers v Whitaker [1] decision that overturned the notion of the 'reasonable doctor' as enumerated in traditional British law by the Bolam Principle [2]. This decision was subsequently confirmed by another case, Chappel v Hart [3, 4]. The Australian National Health and Medical Research Council (NHMRC) has also firmly stated that 'Known risks should be disclosed when an adverse outcome is common even though the detriment is slight, or when an adverse outcome is severe even though its occurrence is rare', and 'Complex interventions require more information, as do interventions where the patient has no illness' [5]. Even in Britain, there are winds of change blowing, with recent professional guidelines from the General Medical Council [6] stating that 'existing caselaw gives a guide as to what can be considered minimum requirements of good practice', and that patients 'must be given sufficient information, in

a way that they can understand, in order to enable them to make informed decisions about their care'. These legal concepts have recently undergone further change following Rosenberg v Percival [7], when the High Court of Australia said: 'The more remote the contingency which a doctor is required to bring to the notice of a patient, the more difficult it may be for the patient to convince a court that the existence of the contingency would have caused the patient to decide against surgery'. These decisions have meant that anaesthetists have to decide which anaesthetic risks to declare to patients, and what incidences to quote for those declared risks. The actual risks of anaesthesia are not readily listed anywhere, and those complications that have been recorded often have widely differing variances in different studies. Even with the certain end-point of death, there are varying interpretations as to whether or not anaesthesia was the sole cause or was only contributory [8, 9]. In any event, anaesthetists tend to be ignorant of the overall published mortality figures at one month for patients undergoing

elective surgery of 1:177 (~1:200), or of 1:34 (~1:40) for emergency surgery [10].

In an endeavour to improve this situation, we surveyed the literature and compiled a listing of the risks of anaesthesia for a variety of anaesthetic complications and side effects. The choice and incidence of complications have been influenced by the available literature, and to some extent the inherent risks will be a result of the type of practice that those authors had. As in the literature on medical and surgical risk assessments, anaesthetic literature is more likely to represent best practice than worst practice, and to be the experience of practitioners who are commonly performing the procedures rather than infrequent practitioners of any particular procedure.

Methods

A Medline® search of the literature was conducted via PubMed®, from 1966 to date, for all publications concerned with peri-operative risk and complications associated with anaesthesia. This computerised search identified keywords in the title, abstract and medical subject headings (MeSH). In addition, the 'related articles' feature of PubMed® was used to identify other relevant publications. Key words included anaesthesia/anaesthesia, risk, peri-operative, postoperative, complications, mortality and morbidity. Specific complications were also sought, such as cardiac arrest, respiratory, cardiovascular and neurological complications, awareness, anaphylaxis, ocular complications, deafness, regional anaesthesia morbidity and minor morbidity (pain, postoperative nausea and vomiting, sore throat, headache, drowsiness, dizziness and dental damage). Reference lists from appropriate selected publications were hand searched to identify additional relevant articles.

Results

Traditional measures of adverse outcome in anaesthesia may be divided crudely into peri-operative mortality and morbidity.

Mortality

Anaesthetic-related mortality is rare. Although death is a clearly definable end-point, epidemiological studies of peri-operative mortality over the last 50 years vary in their objectives, study design, populations, definitions and time span, leading to difficulties in comparing studies. In addition, the denominator data are most often only estimated and are often unreliable, whereas the numerator will often be under-reported, particularly in situations with potential medicolegal implications. Table 1 summarises these studies [8, 9, 11–34].

Total peri-operative mortality is not insignificant, with 30-day mortality rates in the UK quoted in 2000 as 1:34 (2.94%) after emergency surgery and 1:177 after elective surgery (0.56%) [10, 35]. However, the contribution of anaesthetic-related mortality is now considered to be < 10% of total operative mortality. Studies of anaestheticrelated mortality in the 1950s reported an incidence of 2.5-6.4:10 000 deaths [11, 21, 36]. Since then, data from South Africa have shown a decrease from 4.3:10 000 anaesthetic-related deaths (1956-71) to 0.7:10 000 (1972-87) [21]; from 1:5500 to 1:20 000 in New South Wales (NSW), Australia, from 1960 to 1990 [25], and from 0.3:10 000 to 0.13:10 000 for the whole of Australia from 1985 to 1999 [9, 25]. These Australian figures exemplify the difficulties inherent in such statistics, as the NSW rate (with compulsory Coroner reporting for all deaths within 24 h of anaesthesia and surgery) is given as 1:66 183 in the Australia and New Zealand College of Anaesthetists (ANZCA) report [9], yet the number of procedures estimated by the NSW Committee for the same period is approximately half those calculated by the ANZCA report (J Warden & R Holland pers. commun.), thus bringing the incidence more into line with other recent NSW reports [25]. The UK Confidential Enquiry into Peri-Operative Death (CEPOD) in 1987 estimated the risk of death within 30 days solely due to anaesthesia as 1:185 000 [8]. Some authors now quote the incidence of death due to anaesthesia in ASA physical status I and II patients as $\sim 1:100\ 000$ (range 1:2500–1:185 000) [8, 20] with risk increased 5–10 times for high-risk patients and emergency surgery [37]. This improvement in mortality may be attributed to several factors: better general medical care in the community; changes in anaesthetic drugs and techniques; better staffing, supervision and training; improvements in monitoring and peri-operative care; changes in surgical techniques; continuing medical education, including the recognition of preventable hazards through mortality surveys.

National studies of mortality that assess the quality of the delivery of care continue to highlight factors that contribute to anaesthetic-related mortality: inadequate pre-operative assessment, preparation and resuscitation; inappropriate anaesthetic technique; inadequate perioperative monitoring; lack of supervision; poor postoperative care [25, 35, 38].

Mortality studies in the outpatient population reflect the safety profile of anaesthesia for this group. Warner et al. [39] studied 38 598 patients having 45 090 procedures and reported four deaths within one month of surgery, comprising two from road traffic accidents and two from myocardial infarction (nonaccidental death rate 1:22 546). Other large studies report no peri-operative deaths in the ambulatory population [40–43].

Table 1 Studies of anaesthetic and peri-operative mortality.

Author	Location	Study Period	Number of Procedures	Deaths	Deaths:10 000
Beecher & Todd [11]	10 university hospitals, USA	Hospital	599 548	7977	133.1 total
	1948–1952	stay			6.4 anaesthesia-related
Vacanti <i>et al.</i> [12]		48 h	68 388		ASA1 8 total
					ASA5 940 total
Marx et al. [13]	Bronx Municipal hospital New York, US	5 days	34 145	645	189 total
Hovi-Viander [14]	100 Finland	3 days	338 934	626	18 total 2 anaesthesia-related
Turnbull et al. [15]	General Hospital, Vancouver,	48 h	195 232	423	22 total
ramban et an [15]	Canada	10 11	155 252	123	2 preventable
	Carrada				anaesthesia-related
Lunn & Mushin [16]	5 regions in UK	6 days	1 147 362	3736	62.5 total
	g	,-			5.9 anaesthesia-related
					1 attributed to anaesthesia
Gibbs [17]	1979–1984	Hospital stay	1 100 000*		0.5 anaesthesia-related
Buck et al. [8]	3 NHS regions	30 days	555 168	1:185 056 solely	70 total
Duck et an [e]	5 Time regions	oo aayo	333 .00	attributed to anaesthesia	7.7 anaesthesia-related
Tiret <i>et al.</i> [18]	460 French public and private	24 h	198 103	67 (16 coma)	4.2 total
11101 01 01. [10]	hospitals 1978–1982		150 105	or (10 coma)	1.26 anaesthesia-related
Olsson & Hallen [19]	1967–1984	7 days	250 543		2.4 total
0.550.1 @ 1.00.1 [1.5]	.50, .50.	, aays	250 5 .5		0.4 anaesthesia-related
Pederson & Johansen [20]	Prospective 1 year study		7306		4 attributed to anaesthesia
Harrison [21]	Groote Schur Hospital, Cape Town,	24 h	782 182		18.3 total
	South Africa, 1956–1987				1.9 anaesthesia-related
NH & MRC [22]	Australia 1985-1987	2 days	5 470 000*		0.3 anaesthesia-related
NH & MRC [23]	Australia 1988-1990	2 days	7 800 000*		0.2 anaesthesia-related
Wang & Hagerdal [24]	1979–1989	24 h	262 850		0.3 anaesthesia-related
Warden et al. [25]	Australia 1984-1990	24 h	3 500 000*	1503	4.4 total
					0.5 anaesthesia-related
Tikkanen &	Finland 1986	Hospital	325 585		17.5 total
Hovi-Viander [26]		stay			0.2 anaesthesia-related
McKenzie [27]	Zimbabwe teaching hospital 1992	24 h	34 553	89	25.8 anaesthesia-related 3.3 avoidable anaesthetic
Eagle & Davis [28]	Western Australia 1990-1995	48 h	166 000		6 total
			per year*		0.25 anaesthesia-related
ANZCA [29]	Australia 1991-1993	2 days	7 800 000*	116	0.15 anaesthesia-related
ANZCA [30]	Australia 1994–1996	2 days	8 500 000*	135	0.16 anaesthesia-related 0.067 attributable to anaesthesia
NHSE [31]	NHS performance indicators	30 days	2 300 000	32 956	140 emergency total
	1998:1999	Ju days	_ 555 555		50 elective total
Arbous et al. [32]	Prospective study 1995–1997	24 h	869 483	811	8.8 total
	, ,			-	1.4 anaesthesia-related
Kawashima et al. [33]	Training hospitals, Japan 1999	7 days	793 840		7.19 total
•	· · ·	•			0.13 attributable to anaesthesia
Kawashima et al. [34]	Training hospitals, Japan 2000	7 days	941 217		7 total
• •	· · ·	•			0.1 attributable to an anaesthesia
ANZCA [9]	Australia 1997–1999	2 days	10 336 000	130	1:79 509 anaesthesia-related 0.13 attributable to anaesthesia

^{*}Estimated.

NH & MRC = National Health & Medical Research Council; NHSE = National Health Service Executive; ANZCA = Australian and New Zealand College of Anaesthetists.

The UK Confidential Enquiry into Maternal Deaths (CEMD) has recorded mortality in pregnant women since 1952. The direct death rate associated with Caesarean section has decreased from 4:1000 (1952–54) to ~ 0.1:1000 in the last triennium (1997–99) [44].

Improvements in maternal safety in the UK have taken place in association with the development of specialist obstetric anaesthesia services and a move towards the increased use of regional anaesthesia. The contribution of anaesthesia to peri-operative maternal mortality in the UK

has been recorded since 1970. Of the 104 anaesthetic deaths reported since then, 75% were associated with emergency procedures and 25% with elective procedures; 96% were associated with general anaesthesia [44]. Maternal mortality rates in the developing world are much higher, with sub-Saharan Africa recording average mortality rates of 100 times that in the UK (980:100 000 live births) [45, 46]. In South Africa, a confidential enquiry into maternal deaths noted that anaesthetic accidents contributed to 5% of maternal deaths, with complications resulting from general anaesthesia, particularly difficult or failed tracheal intubation, being the commonest cause [47].

The risk of peri-operative death increases with age. Jin and Chung quoted an overall mortality rate of 1.2% within 30 days of surgery for the general population. This increased to 2.2% in 60-69 years olds, 2.9% in 70-79 years olds, 5.8-6.2% in those aged > 80 years and 8.4% in those aged > 90 years. Major surgery further increases this risk, leading to a 19.8% mortality rate in the latter group [37,48-51]. The 1999 CEPOD report found that > 90% of peri-operative deaths were in the over 60s, with 38% in those aged > 80 years. The majority of these procedures were urgent or emergencies (65%) in a highrisk population (84% ASA physical status III or higher). Most elderly patients underwent general (42%), orthopaedic (22%) or vascular (14%) procedures [52]. As the elderly constitute a growing proportion of the elective and emergency surgical workload, anaesthetists need to be aware of their higher risk of morbidity and mortality in order to ensure optimal peri-operative care.

Mortality rates associated with anaesthesia in children have decreased steadily from 1.8–3.3:10 000 in the 1960s to 0.18–0.25:10 000 by 1990 [53].

Table 2 Studies of intra-operative and immediate postoperative morbidity.

Author	Study period	No. of Procedures	Intra-operative events (%)	Recovery events (%)
Cohen <i>et al</i> . [56]	1975–1978	52 197	7.6	3.1
	1979–1983	60 524	10.6	5.9
Cooper et al. [57]	1985-1986	12 088	13.8	7.1
Zelcer & Wells [58]	Nov 1985	443		30.0
Pederson et al. [48]	1986-1987	7306	4.5	7.4
Hines et al. [59]	1986-1989	18 473	5.1	23.7
Moller et al. [60]	1989-1990	20 802	14.9	13.5
Rose <i>et al.</i> [61]	1991-1993	24 157		1.3 respiratory
Ouchterlony et al. [62]	1985-1988	1361	18.7	47.4
Schwilk et al. [63]		18 350	23.2 (1.2 serious)	
Hunter & Molinaro [64]	1990-1994	1126	2.3	
Schwilk et al. [65]		26 907	27.9 (0.9 serious)	
Chung et al. [66]	3 years	17 638	4.0	9.6
Bothner et al. [55]	1992–1997	96 107	22 overall peri-operatively (1.0 serious)	
Fasting & Gisvold [67]	1996–2000	83 844	15.7	

Morbidity

Anaesthetic morbidity ranges from major permanent disability to minor adverse events causing distress to the patient but no long-term sequelae. As with studies of mortality, there is a lack of uniformity in reporting perioperative adverse events between institutions and countries. Criteria for reporting vary enormously, from limited details due to medicolegal necessity [54] to comprehensive computerised data acquisition for national benchmarking processes [55]. Table 2 summarises some of the larger studies of intra-operative and recovery adverse events in both ambulatory patients and inpatients [48, 55–67]. Both the methods of data acquisition and definitions of criteria for adverse events differ between studies, making comparisons difficult. Other methods of investigation have been established to examine closed legal claims and critical incidents or sentinel events in anaesthesia. These national studies highlight problems or system failures and recommend improvements in patient care [38, 44, 54, 68, 69].

Cardiac arrest

The incidence of peri-operative cardiac arrest has decreased significantly over the last 25 years (Table 3) [14, 19, 24, 33, 34, 56, 70–77]. Keenan *et al.* noted that the cardiac arrest rate halved over two decades at their institution (21:10 000 in 1969–1978 vs 1.0:10 000 in 1979–1984), predominantly because of a decrease in respiratory complications [73]. Most studies in the last 10 years quote incidences of anaesthesia–related cardiac arrest of 0.12–1.4:10 000, with associated mortality rates of 0.06–0.6:10 000 [24, 33, 34, 76, 77]. Morray's study of children quotes a similar incidence (1.4:10 000), with 55% of events occurring in infants less <1 year old [75]. The commonest causes of cardiac arrest included

Table 3 Studies of peri-operative cardiac arrest.

Author	Study period	Procedures	Cardiac arrest	Remarks
Hovi-Viander [14]	1975 Finland	338 934	3:10 000	
Pottecher et al. [70]	1978–1982 France		6:10 000	3.5:10 000 anaesthesia-related mortality rate
Cohen et al. [56]	1975-1983 Canada		7.1:10 000 recovery	·
Olsson & Hallen [19]	1967–1984	250 543	6.8:10 000	0.3:10 000 anaesthesia-related mortality 2.4:10 000 overall mortality
Pederson et al. [71]	Prospective	7306	10.9:10 000 intra-operative	,
Aubas <i>et al</i> . [72]	1983–1987 France	102 468	2.8:10 000 anaesthesia-related	1.1/10 000 anaesthesia-related 1.2 mortality rate
Keenan & Boyan [73]	1969–1978 1979–1988	241 934	2.1:10 000 1.0:10 000	0.8:10 000 1:10 000 due to respiratory cause
Wang & Hegerdal [24]	11 years	262 850	0.04:10 000 intra-operative 0.08:10 000 recovery	
Auroy <i>et al</i> . [74]	5 months, France	103 730 (regional anaesthesia)	3.1:10 000 total 6.4 (1.2):10 000 in spinals	23% fatal in spinals
Morray et al. [75]	Multicentre POCA <18 years old	•	1.4 (0.45):10 000 related to anaesthesia	26% fatal 55% <1 year old
Biboulet et al. [76]	6 years, France	101 769	1.1:10 000	0.6:10 000 anaesthesia-related mortality rate
Kawashima et al. [33]	1999, Japan training hospitals	793 840	6.53:10 000 total 0.78:10 000 attributable to anaesthesia	0.1:10 000 anaesthesia-attributed mortality rate
Kawashima et al. [34]	2000, Japan training hospitals	941 217	6.52:10 000 total 0.53:10 000 attributable to anaesthesia	0.06:10 000 anaesthesia-attributed mortality rate
Newland et al. [77]	1989–1999	72 959	1.37:10 000 contributory anaesthesia 0.69:10 000 attributable to anaesthesia	0.55:10 000 anaesthesia-attributed mortality rate

POCA: Paediatric peri-operative cardiac arrest register.

medication-related events, cardiovascular causes including hypovolaemia, and poor airway management. In patients undergoing regional anaesthesia, including spinal, epidural, peripheral nerve blocks and intravenous regional anaesthesia, the overall cardiac arrest rate has been quoted at 3.1:10 000. Spinal anaesthesia alone accounted for 6.4:10 000 events, of which 23% were fatal [74].

Respiratory complications

Postoperative respiratory complications, such as pneumonia, remain a major cause of surgical morbidity and mortality [78]. The contribution, if any, of anaesthesia to these events is not often recorded. Respiratory complications due to anaesthesia are more commonly those acute events closely associated in time to the operation. Engelhardt & Webster [79] recently reviewed pulmonary aspiration of gastric contents associated with anaesthesia. The incidence of this complication in the general surgical population has been reported in three large studies. Olsson *et al.* [80] found an aspiration incidence of 1:2131

during anaesthesia in 185 385 patients, with a mortality rate of 1:45 454. Forty-seven per cent of the patients who aspirated developed pneumonitis and 17% required lung ventilation. A retrospective review of > 200 000 patients in the Mayo Clinic revealed an aspiration rate of 1:3216, with a mortality rate of 1:71 829 [81]. In a study of 85 594 adult surgical patients by Mellin-Olsson *et al.* [82] the incidence of pulmonary aspiration was 2.9:10 000, all in patients undergoing general anaesthesia. The incidence was four times greater in emergency cases. In 30 199 patients undergoing regional anaesthesia, none aspirated.

In children, the risk of regurgitation and pulmonary aspiration may be greater but it is rarely associated with pneumonitis [79, 83]. Large studies have reported an incidence between 1:10 000 (n = 40 240) and 10:10 000 cases (n = 50 880), with no associated deaths or serious morbidity [84, 85]. Aspiration occurred more frequently in patients with poorer ASA physical status (III or IV) and in emergency cases.

With regard to obstetric anaesthesia, patients undergoing Caesarean section under general anaesthesia have at least twice the risk of pulmonary aspiration when compared with the general population. Two Italian studies quoted an incidence of aspiration between 1:1431 and 1:1547, whereas a more recent study reported aspiration in 1:900 patients undergoing Caesarean section [86–88]. There were no fatalities in this group.

The reported incidence of a Cormack and Lehane grade 3 or 4 view at laryngoscopy is 2–8%. Difficulty with tracheal intubation (defined as three or more attempts) in general surgical patients is reported as occurring in 1.15–3.8% of patients, with failure to intubate the trachea seen in 0.13–0.3%. Inability to intubate a patient's trachea or ventilate a patient's lungs happens much less commonly, and is estimated to occur in 1–3:10 000 cases [89]. In the obstetric population, the incidence of difficult or failed tracheal intubation is more common, reported at between 1:250 and 1:300 patients, presumably due to anatomical and physiological airway changes and, more recently, the relative lack of training opportunities in obstetric general anaesthesia [90–92].

Other cardiovascular complications

Peri-operative myocardial infarction (MI) has been recognised as a major problem since the 1950s [93]. Recent MI and congestive cardiac failure were two early risk factors identified as being associated with peri-operative MI. Shah *et al.* [94] reported that the peri-operative MI rates were 5% if the time from MI to operation was > 6 months, 15% if between 3 and 6 months, and 37% if < 3 months. In a study by Rao *et al.* [95] in which patients with a previous MI were aggressively monitored and managed peri-operatively, lower peri-operative morbidity and mortality rates were produced. Re-infarction occurred in 5.7% of patients who were 0–3 months after an MI, and in 2.3% of patients who were 4–6 months after an MI.

Pre-operative predictors of cardiovascular risk in non-cardiac surgery have been identified, notably in Goldman et al.'s cardiac risk index [96], Detsky et al.'s [97] modified cardiac risk index (incorporating unstable angina, history of pulmonary oedema and Canadian Cardiovascular Society angina classes III and IV), and more recently Lee et al.'s [98] revised cardiac risk index derived from 4315 patients undergoing major noncardiac procedures in a teaching hospital. Goldman et al.'s criteria and risk stratification for elective noncardiac surgery are summarised in Table 4. Lee et al.'s study [98] found that major cardiac complications occurred in 2% of patients in the derivation cohort (56:2893). The six independent predictors of complications were: high-risk surgery, history of ischaemic heart disease, history of congestive

Table 4 Goldman's Cardiac Risk index [96].

Criteria				
Age >70 years o	old		5	
Myocardial infra	action within :	x months	10	
S3 gallop or rais	sed jugular ve	ous pressure	11	
Important aorti	c stenosis		3	
Rhythm other tl	nan sinus or p	emature atrial coi	tractions 7	
>5 premature vo surgery	entricular con	ractions per minu	e before 7	
Poor general m	edical status			
P_aO_2 <8.0 kPa o				
		arbonate <20 <mark>m</mark> m	ol.l ^{–1}	
Urea >18 mmol	.I ^{–1} or <mark>creatini</mark>	ie >240 mmol.l ⁻¹		
Abnormal aspar	tate transami	ase		
Signs of chronic	liver disease			
Bedridden from	non-cardiac o	auses	3	
Intraperitoneal, intrathoracic or aortic operation				
Emergency open	ration		4	
TOTAL			53	
	Life-thr	atening		
Points Grou	p complic	tions (%) Mo	tality (%)	
0–5 I	0.7	0.:		
6–12 II	5	1.5	;	
13–25 III	11	2.:		
26–53 IV	22	56		

cardiac failure, history of cerebrovascular disease, preoperative treatment with insulin, pre-operative serum creatinine >166 μ mol.1⁻¹. Combining the derivation and validation cohorts, rates of major cardiac complication with 0, 1, 2 or \geq 3 of these factors were approximately 0.5, 1, 5 and 10% respectively (Table 5) [99].

The American College of Cardiologists and the American Heart Association have published guidelines for perioperative cardiac evaluation of patients undergoing noncardiac surgery [100]. They classified minor, intermediate and major clinical predictors of risk for MI, cardiac failure and peri-operative death that are summarised in Table 6, but gave no indication of the relative risks.

Risks for invasive monitoring depend on the type of monitoring and the site of access. Scheer et al. [101] reviewed complications associated with peripheral arterial catheters in anaesthesia and intensive care and found that major complications, such as permanent ischaemic damage, sepsis and pseudo-aneurysm formation, occurred in < 1% of cases. Temporary arterial occlusion occurred in 1.5–35% (mean 19.7%) of radial arteries in which catheters had been placed, and in 1.45% of femoral arteries. They suggested that risk increases with an increase in catheter diameter and if duration of cannulation is > 48 h. Local site infection occurred in 0.74% of radial arteries and 0.78% of femoral arteries. Duration of cannulation > 96 h was associated with an increased risk of infection. Cannulation of the ulnar artery had a similar complication rate to radial artery cannulation.

Major medical risk factors for noncardiac surgery

Actual rates of cardiac complications % [range]

No. of simple	Revised cardiac	Ref. 99			Approximate rates of cardiac complications	
	risk index		No blocker	Blocker	No blocker	
0	Class I	0.4 [0.05–1.5]	1.0 [0.6–1.9]	0.4 [0.1–0.9]	\sim 0.5%	
1	Class II	0.9 [0.3-2.1]	2.2 [1.4–3.3]	0.8 [0.3–1.7]	\sim 1.0%	
2	Class III	6.6 [3.9–10.3]	4.5 [3.2-6.3]	1.6 [0.8–3.3]	$\sim 5.0\%$	
3	Class IV	11.0 [5.8–18.4]	9.2 6.5-13]	3.4 1.7-6.7]	\sim 10.0%	
4	Class V		18 [12–20]	7.0 [3.4–14]	\sim 15.0%	
=5	Class VI		32 [19–47]	14 [6.5–27]	\sim 30.0%	

Simple risk factors: High-risk surgery; ischaemic heart disease; congestive cardiac failure; cerebrovascular disease; insulin-dependent diabetes; creatinine >170 mmol. l^{-1} .

Major Intermediate Minor Unstable coronary syndromes Mild angina pectoris **Elderly** patients Decompensated congestive Compensated or prior Low functional capacity heart failure congested heart failure (< 4 metabolic equilvalents) Significant symptomatic Prior myocardial infarction Non-sinus rhythm arrhythmias Severe valve disease Diabetes mellitus Stroke history Renal insufficiency Uncontrolled hypertension Abnormal electrocardiogram

Table 5 Lee's revised cardiac index [98, 99].

Table 6 Clinical predictors of increased peri-operative cardiovascular risk [100].

Deaths have been reported after central venous cannulation with an incidence of up to 1:252 using infraclavicular subclavian lines [102]. A subsequent series reported no deaths in 13 800 patients after central venous cannulation [103]. The incidence of pulmonary artery perforation has been quoted as 0.06% in a series of 6245 patients [104]. Looking at nonfatal complications, Ruesch et al. [105] analysed 17 prospective comparative trials of internal jugular vein (n = 2085) and subclavian vein (n = 2428) cannulation. Arterial puncture was more common with internal jugular catheters (3 vs. 0.5%), as was bloodstream infection (8.6 vs. 4%). Malposition of the venous catheter was seen more commonly via the subclavian route (9.3 vs. 5.3%), as were haemopneumothorax (1.5 vs. 1.3%) and vessel occlusion (1.2 vs. 0%). Comparing the subclavian route with the femoral venous route in critically ill patients, femoral lines caused a higher incidence of infections (19.8 vs. 4.5%) and thrombotic complications (21.5 vs. 1.9%) [106].

A meta-analysis of trials looking at the effect of ultrasound guidance suggested that real-time ultrasound guidance improved success rates and decreased complications associated with internal jugular and subclavian vein catheter placement when compared with anatomical landmark techniques for placement [107].

The incidence of peripheral venous thrombophlebitis was studied extensively in the 1970s and 1980s when

intravenous preparations of drugs solubilised in propylene glycol and 'Cremophor EL' were available. In a study of 519 patients, venous sequelae were reported in 12% of patients receiving diazepam alone or fentanyl and methohexital [108]. Other authors have reported thrombophlebitis in 24–43% patients given etomidate and in 4–23% of those given thiopental [109, 110]. A review by Clarke [111] suggested that venous sequelae depended on the solubilizing agent, with the frequency of complications in water-soluble and Cremophor EL-based anaesthetics varying between 5 and 10%. Diazepam or etomidate dissolved in propylene glycol could produce venous reactions in > 25% of patients.

Postoperative neurological dysfunction

In the elderly surgical patient, postoperative cognitive dysfunction (POCD), most commonly manifested as a lack of concentration and problems with memory, may persist for a long period. The ISPOCD1 multicentre study looked at 1218 patients aged > 60 years undergoing major abdominal, thoracic or orthopaedic surgery [112]. There was a 26% incidence of POCD at one week after surgery and a 10% incidence at three months after surgery, compared with 3.4 and 2.8% in controls. Risk factors identified for early POCD included increasing age, duration of anaesthesia, limited education, second operation, postoperative infections and respiratory

complications. Only age was found to be a risk factor for late POCD. There was no difference in long-term cognitive function between patients having general or regional anaesthesia [113]. Follow-up at 1–2 years showed that POCD is a reversible condition in the majority of patients, but may persist in ~ 1% [114].

The reported incidence of peri-operative delirium varies widely due to differences in diagnostic criteria. Up to 14% of general surgical patients develop postoperative delirium, with the risk increasing to 40% if intensive care is required [115, 116]. Another study showed a 44% incidence of delirium in the elderly after fixation of a fractured neck of femur [117, 118]. Significant risk factors for postoperative delirium include increasing age and poor medical condition. After 75 years of age, there is a threefold increased risk of developing postoperative delirium [115].

The incidence of peri-operative cerebrovascular accident (CVA) varies between 0.08 and 2.9% in general surgical patients, and is as much as 4.8% in patients undergoing head and neck surgery, with a reported mortality rate of 46%. As a reference point, the annual incidence of stroke in the UK is 0.1–0.2% [116, 119]. Most peri-operative CVAs occur between the second and tenth postoperative day (mean = seventh day). Risk factors include advancing age, previous cerebrovascular disease (10-fold increased risk), hypertension (fourfold increased risk), peripheral vascular disease, chronic obstructive airways disease, atrial fibrillation, carotid artery stenosis and obstructive sleep apnoea. The incidence of peri-operative CVA in a patient with a previous stroke is 2.1%, with an associated increased risk of mortality of 60% [116].

The risks associated with carotid endarterectomy have been recently summarised in a review by Barnett et al. [120]. Several randomised controlled trials were published in the 1990s comparing best medical treatment with best medical and surgical treatment [121–123]. The combined risk of stroke and death within 30 days of surgery was 6.2% in symptomatic patients and 4.4% in asymptomatic patients [122, 123]. The risk of a disabling peri-operative CVA or death was 2.1%, the majority occurring within 24 h. Cranial nerve injuries (involving the facial nerve, superior laryngeal branch of vagus nerve, spinal accessory nerve or hypoglossal nerve) were seen in up to 6.8% of patients [120]. When cerebral angiography was used, 0.6% of the 2885 patients in the North American Symptomatic Carotid Surgery Trial had nondisabling strokes within 24 h, and an estimated 0.1% suffered disabling strokes [120, 124, 125].

Awareness

The fear of awareness is common in patients, with up to 54% concerned about waking up during their surgery

[126]. In the 1970s, anaesthetic studies using 60–70% nitrous oxide alone as a maintenance anaesthetic showed incidences of awareness in up to 7% of patients [127]. More recently, the incidence of conscious awareness with explicit recall and severe pain has been estimated at < 1:3000 general anaesthetics [128]. Conscious awareness with explicit recall but without pain is more common, with a reported incidence of 0.1–0.7% [128–130]. Most large clinical studies nowadays estimate the risk of explicit awareness as < 0.3% [127]. The incidence may be increased in general anaesthesia using neuromuscular blockade compared with general anaesthesia without paralysis (0.18 vs. 0.1%) [130]. A study using total intravenous anaesthesia with neuromuscular blockade in 1000 patients reported an incidence of awareness of 0.2% [131]. Cases of awareness come not only from cardiac surgery and obstetrics, but also from all surgical specialities. After surgery, patients who experience intraoperative awareness may develop a post-traumatic stress disorder, particularly those patients who have experienced severe pain. The Australian Incident Monitoring Study recently reviewed 81 cases of awareness and found the commonest causes to be drug error resulting in inadvertent paralysis of an awake patient, and failure of delivery of a volatile anaesthetic. The authors concluded that an objective central nervous system depth of anaesthesia monitor and an improved drug administration system might have prevented the majority of these cases [132].

Anaphylaxis

Life-threatening allergic reactions associated with anaesthesia occur in < 1:10 000 patients. Studies of anaphylaxis associated with anaesthesia show an incidence between 1:10 000 and 1:20 000 in Australia (1993) [133] and 1:13 000 in France (1996) [134]. A French survey of 467 patients with a history of anaphylaxis in 1997–98 revealed the main causal agents to be neuromuscular blocking drugs (69.2%), latex (12.1%) and antibiotics (8%). Clinical features included cardiovascular collapse (53.7%), cutaneous symptoms (69.6%), bronchospasm (44.2%), angiooedema (11.7%) and cardiac arrest (4%) [135].

Ocular complications

Peri-operative visual changes may vary in severity from transient diplopia or blurring of vision to irreversible blindness. A recent prospective study of 671 patients undergoing general anaesthesia or central neuraxial blockade reported the new onset of blurred vision lasting at least three days in 4.2% of patients, most of which resolved in one to two months. However, seven patients (1%) required eye care intervention for permanent blurred vision [136]. The incidence of ocular injury in

large prospective studies of nonocular surgery varies from 0.056% (n = 60~965) [137] to 0.17% (n = 4652) [138]. The commonest injury sustained is corneal abrasion, in keeping with the findings of the ASA Closed Claims Analysis of ocular injury. They found 35% of all claims against anaesthesiologists for eye injuries were a result of corneal abrasions occurring during general anaesthesia [139].

Vision loss and blindness after surgery occur rarely, with a retrospective study of noncardiac surgery reporting an incidence of 1:125 234 (n = 410 189) [140]. In cardiac surgical patients, the incidence increases, with 0.1–2% reporting loss of vision [141, 142].

Deafness

Hearing loss may occur after general anaesthesia or spinal anaesthesia. Sudden sensorineural hearing loss (SNHL) has been extensively reported following cardiopulmonary bypass (incidence 1:1000 cases) presumably due to microemboli. In nonbypass cases, there are only 18 case reports in the literature of sudden SNHL following general anaesthesia in nonotological surgery. These are presumed to be due to cochlear or middle ear membrane breaks. Nitrous oxide has been implicated in this process because of pressure effects in the middle ear. The incidence of idiopathic sudden SNHL is 5–20:100 000 per year. However, there is a high rate of spontaneous recovery (47–78%) [143].

There have been reports in the literature of up to 16% of patients suffering transient hearing loss after spinal anaesthesia. It is hypothesised that the hearing loss is due to middle ear changes as a result of cerebrospinal fluid (CSF) leakage. Studies have shown a threefold increase in mild hearing loss in young patients (< 30 years) compared with their elders (> 60 years), suggesting that CSF leak after dural puncture occurs more frequently in younger patients [144, 145].

Minor morbidity

Incidences of relatively minor morbidity, such as pain and postoperative nausea and vomiting, have not changed significantly over the last 30 years despite improvements in anaesthesia drugs and techniques [146, 147]. Minor sequelae following surgery often have a significant impact on patient recovery, leading to decreased function and slower resumption of daily activities after discharge [148, 149]. Much of the available data on minor sequelae are from studies in outpatients that attempt to identify areas for quality improvement [150].

Postoperative pain

Inadequate management of acute pain in recent years has led to the development of multidisciplinary acute pain

services in an attempt to improve quality of care [151– 154]. The UK Audit Commission proposed in 1997 that < 20% of patients should experience severe pain following surgery after 1997, and that this should be decreased to < 5% by 2002 [155]. However, Dolin et al. [156] recently reviewed published studies on the incidence of moderate to severe pain after major surgery, and concluded that the overall incidence of moderate to severe pain was 30%, and that of severe pain was 11%. When looking at commonly used analgesic techniques, the incidence of moderate to severe and severe pain using intramuscular analgesia was 67 and 29%, respectively. For patientcontrolled analgesia (PCA), the incidence of moderate to severe pain was 36% and severe pain 10%. In patients receiving epidural analgesia, the incidence of moderate to severe pain was 21% and severe pain 8%.

In ambulatory surgery, postoperative pain has been shown to be a major cause of delayed discharge, unplanned hospital admission and readmission [157]. A systematic review of postdischarge symptoms in outpatients reported an overall incidence of pain of 45% (range 6–95%) [150]. Severity of postoperative pain is dependent on both the type and length of surgical procedure. Of note, up to half the patients undergoing orthopaedic, laparoscopic and general surgical procedures still have significant pain at 24 h [158].

Patients' understanding of postoperative pain is poor. More than 50% of patients assume that pain is a normal part of the postoperative course and the healing process. Most are prepared to suffer pain rather than complain, and despite this > 80% are satisfied with their pain management [159]. The Australian National Health & Medical Research Council suggests that 'changes are called for in training, knowledge, attitudes and practice of medical, nursing and allied professionals along with greater public awareness and expectations in the treatment of pain' [154]

Postoperative nausea and vomiting

Postoperative nausea and vomiting (PONV) is one of the most common and distressing complications following anaesthesia with an average reported incidence between 20 and 30% [160], but it may occur in up to 80% of patients [161]. In outpatients, the incidences of nausea and vomiting after discharge have been reported as 17 and 8%, respectively [150]. PONV has a multifactorial aetiology including type and duration of anaesthesia, drug therapy, type of surgery and patient characteristics. It is seen particularly in the young, females (three times the risk for males [162]), overweight, nonsmokers and in those with a history of motion sickness and previous PONV [40, 146, 160]. Surgical procedures such as laparoscopy, strabismus surgery, ear, nose and throat

(ENT), dental, orthopaedic and plastics operations have been associated with the highest incidence, suggesting operations that are associated with significant postoperative pain may lead to PONV [163–165]. Sinclair *et al.* [162] and Apfel et al. [166] have attempted to quantify the risks of PONV in individual patients by statistical analyses taking these factors into account.

The debate over optimal PONV management continues, in particular anti-emetic prophylaxis vs. symptomatic treatment. Looking at studies of efficacy and cost-effectiveness, Tramer *et al.* [166, 167] suggested that treatment of PONV with the 5HT₃ antagonist ondansetron (1 or 4 mg) might be more cost-effective and safer than prophylaxis (4 or 8 mg). Other authors [169] now recommend routine anti-emetic prophylaxis in all patients at >10% risk of PONV. Combination therapy, using a 5HT₃ antagonist, dexamethasone and/or low-dose droperidol (0.625–1.25 mg), has been recommended if PONV risk exceeds 30% [169]. Rather than looking at a 'surrogate' outcome, such as percentage of patients vomiting, it has been suggested that true outcomes, such as patient satisfaction and delayed discharge, are more relevant [170].

Sore throat

The incidence of sore throat following general anaesthesia has been studied prospectively in 5264 ambulatory patients, with a reported incidence of 12.1% overall at 24 h after surgery. The type of airway used affected the incidence, with 45% of patients complaining of sore throats after tracheal intubation, 18% after laryngeal mask (LMA) insertion and 3% following the use of a facemask [171]. Other studies quote a 14–64% incidence of sore throat in association with tracheal intubation, 9–29% with the LMA and 48% using a Combitube [172–177].

Headache

The incidence of nonspecific headache after anaesthesia in outpatients is quoted as 17% (range 2–30%) [150]. Nikolajsen *et al.* [178] identified risk factors associated with postoperative headache, which included a daily caffeine consumption of > 400 mg in 24 h, pre-operative headache, those who normally experience two or more headaches per month, and a longer duration of fasting. They suggested caffeine withdrawal might cause symptoms within 12–16 h.

Drowsiness and dizziness

Drowsiness has been reported as occurring after discharge in 42% (range 11–62%), and dizziness in 18% (range 7–41%) of outpatients [150]. A review by Holte *et al.* [179] concluded that intra-operative fluid administration of 1 leads to a decrease in incidence of postoperative drowsiness and dizziness.

Dental and oral damage

Oral tissue and dental damage are common complications of general anaesthesia and account for a significant proportion of all medicolegal claims against anaesthetists. A 10-year study of 598 904 procedures reported 132 cases (1:4500) of dental injury that required intervention. Nearly half of these injuries occurred during laryngoscopy and tracheal intubation [180]. A New Zealand survey of anaesthetists' practice estimated the incidence of dental damage to be 10.4:1000 [181]. Looking at all types of oral trauma, a prospective study of 404 patients having general anaesthetics with tracheal intubation showed an incidence of 6.9%, ranging from soft tissue laceration to tooth fracture or avulsion [182].

Peripheral nerve injuries

Peripheral nerve injuries associated with anaesthesia are not uncommon. Dhuner's [184] retrospective review of > 30~000 cases over a 6-year period found nerve injuries in $\sim 1:1000$ cases, most commonly involving the ulnar nerve (83%). More recently, prospective studies of postoperative ulnar neuropathy (n=1502;~n=6538) revealed incidences of 1:200–1:350. Fifteen per cent of claims in the ASA Closed Claims Study followed perioperative nerve injury, in particular the ulnar nerve (> 30%), brachial plexus (23%) and lumbosacral nerves (16%) [183–187].

Regional anaesthesia morbidity

Morbidity due to regional anaesthesia has been comprehensively reviewed [188]. Risks include neurological injury, death, cardiac arrest, local anaesthetic toxicity and infection. With central neuraxial blockade, which accounts for >70% of regional anaesthesia practice, the risks of headache, backache and urinary dysfunction must also be considered.

In the 1950s, the incidence of permanent nerve injury following subarachnoid block was reported as 1:10 098 [189], and 2:10 000 after epidural anaesthesia (1969) [190]. More recently, Auroy et al.'s [74] prospective study of 103 730 regional anaesthetics found that long-term neurological injury was three times more common after subarachnoid block (1:10 000) than epidural block (0.3:10 000). In contrast, Dahlgren & Tornebrandt. [191] reported more long-term sequelae in their epidural group (10:10 000) than following spinals (3:10 000), with an overall incidence of 7:10 000 in 17 733 patients. The incidence of paraplegia in Auroy et al.'s study was 0.1:10 000 [74]. Transient neurological complications are more common. After subarachnoid blocks, the incidence varies between 4:10 000 (in 40 640 cases) [74] and 80:10 000 (10 098 cases) [189]. With epidurals, this varies between 1:10 000 and 10:10 000 [74, 190].

The incidence of spinal or epidural haematoma associated with central neuraxial blocks has been quoted as 1:150 000 for epidurals and 1:220 000 for spinals, from a study of 1.5 million patients [192]. The main risk factors are the presence of a coagulopathy, difficult insertion, presence of an epidural catheter and timing in relation to anticoagulant administration [193]. Regional differences have been noted in association with the dose of low molecular weight heparin (LMWH) used. In the USA, where a larger dose of LMWH tends to be used, the incidence of reported haematoma is 1:14 000 compared with 1:2 250 000 in Europe [194]. Interestingly, the risk of spontaneous spinal or epidural haematoma has been estimated as 1:1 000 000 [195].

In the obstetric population, demand for anaesthesia is increasing, with 22% of pregnant women undergoing Caesarean section in the UK [196]. Of these, > 90% of elective and 77% of emergency caesarean sections are performed under regional anaesthesia. Neurological deficits occurring after labour and delivery may be due to the obstetric process itself, secondary to regional analgesia or anaesthesia, or may occur spontaneously. Neurological complications after obstetric epidural analgesia or anaesthesia occur after 0.8-36.2:10 000 epidural blocks [197-202]. After spinal anaesthesia, these complications occur in 5.4-35.4:10 000 blocks [197, 202, 203]. There are no data from large-scale studies on neurological complications after combined spinal-epidural techniques. Cranial nerve palsies follow regional anaesthesia in 1–3.7:100 000 obstetric patients. The abducens nerve is most commonly affected, leading to diplopia [200, 204]. Looking at maternal obstetric palsies, the nerves most commonly injured include the lumbosacral plexus, femoral nerve, obturator nerve and common peroneal nerve. These occur due to nerve compression by the foetal head, during forceps delivery or due to patient positioning during labour. They occur more often after longer labours and high-risk deliveries and are mostly unilateral [197]. Ong et al. [201] reported maternal palsies in 18.9:10 000 deliveries (n = 23 827), manifested as paraesthesia and/or motor dysfunction, all of which resolved within 72 h with supportive care only. A common condition in pregnancy that may be wrongly attributed to regional analgesia or anaesthesia is meralgia paresthetica, resulting from compression of the lateral femoral cutaneous nerve (spinal roots L2-3). Anterolateral thigh numbness or paraesthesia may occur from 30 weeks gestation but is self-limiting and usually resolves within three months of childbirth [197].

Transient radicular irritation has been described in association with spinal anaesthesia in the general population. It is characterised by mild to severe radiating back and buttock pain typically starting within 24 h, and lasting

< 48 h. Risk factors include day surgery, lithotomy position and knee arthroscopy, and the incidence varies with the type of local anaesthetic used [205–207]; with 5% hyperbaric lignocaine, the incidence varies between 10 and 37%, with bupivacaine 0–3%, with mepivacaine 30–37% and with tetracaine 6.8% [188].

Auroy et al. [74] reported the mean (SD) incidence of cardiac arrest associated with spinal anaesthesia as 6.4 (1.2):10 000. In comparison, the incidence of cardiac arrest following epidural anaesthesia was much less at 1 (0.4):10 000. The risk of systemic local anaesthetic toxicity with epidural techniques is higher, with an incidence of 1:10 000, but there are no reported cases in subarachnoid blocks. Epidural abscesses and infection associated with central neuraxial blockade occur in between 1:1930 [208] and 1:7500 cases [209]. However, spontaneous epidural abscesses account for 0.2-2:10 000 hospital admissions per year [210]. The risk of infection is increased in the immunocompromised patient, patients on steroids and when epidural catheters are introduced. In the obstetric population, the incidence of epidural infection is reported between 0.2 and 3.7:100 000 following epidural anaesthesia [200, 204].

Post-dural puncture headache (PDPH) is one of the most common complications associated with spinal and epidural anaesthesia in inpatients, and the incidence is now quoted as $\sim 1\%$ for both, although individual rates may vary widely (range: 0.6-4.2%) [188, 211]. The incidence of spinal headache can be decreased by the use of smaller gauge noncutting needle types [212]. Central neural blockade in outpatients has a reported 9% incidence of headache (range 1-37%) [150]. In obstetric practice, following the use of small gauge noncutting needles for spinal anaesthesia, the incidences of mild, moderate and severe headaches have been reported as 5.9, 4.7 and 0.75%, respectively [213]. Most authors quote an incidence of PDPH of 70-90% after inadvertent dural puncture with epidural needles, and an epidural blood patch success rate of 70-100%. Headache may recur in 30-50% [87, 214, 215]. Any persisting or recurring headaches must raise the suspicion of cranial subdural haematoma, which has an estimated incidence of 2:1 000 000 [200]. The incidence of accidental dural puncture in UK obstetric practice have been reported overall as 0.85% in 294 268 epidural insertions [216]. The fluid used for the loss of resistance technique may affect this rate, with dural puncture rates of 0.69% using saline and 1.11% using air. Long-term morbidity, such as headache and backache, after accidental dural puncture has also been reported [217, 218].

Prospective studies of postpartum back pain have found no association with epidural analgesia or anaesthesia, and the incidence of backache two months after delivery has been reported as being the same regardless of anaesthetic technique used [219–221]. In fact, one of the most important factors associated with back pain following any surgery is the duration of the operation. Brown & Elman [222] reported an 18% incidence of back pain after surgery lasting <1 h, increasing to 50% with surgery lasting 4–5 h.

Urinary dysfunction is a common complication of neuraxial blocks with both opioids and local anaesthetics. Asantila et al. [223] compared different methods of postoperative analgesia after thoracotomy and reported a 90% incidence of urinary retention with epidural morphine 6 mg, and a 60% incidence with epidural bupivacaine 0.25%. Other authors have reported a 1-3% incidence of urinary dysfunction in patients receiving epidural infusions of bupivacaine and fentanyl [224]. In obstetric anaesthesia, studies of the effects of epidural analgesia on postpartum urinary retention have shown that although epidurals may be associated with an increase in residual urine in the bladder, postpartum urinary retention seems to be related more to prolonged or difficult labour than to the effects of epidural analgesia itself, with an incidence of up to 18% [225].

Systemic toxicity is another side effect of local anaesthetic use. This results most usually from inadvertent intravascular injection of the local anaesthetic. Auroy et al. [74] prospectively studied 21 278 regional anaesthetics that included 11 229 intravenous regional anaesthetics (IVRA). The risk of systemic toxicity to local anaesthetic overall was 1:10 000, with peripheral nerve blocks having the highest incidence of toxicity at 7.5:10 000. Seizures occurred in 2.7:10 000 intravenous regional anaesthetics [74]. The mean (SD) incidence of cardiac arrest was 1 (0.4):10 000 patients. Brown et al. [226] reported seizures during or after brachial plexus blocks in 20:10 000 patients. The incidence was lower in patients having an axillary block (12:10 000) compared with a supraclavicular block (79:10 000).

Auroy et al. [74] reported an incidence of permanent nerve injury after peripheral nerve block as 1.9:10 000, with all affected patients experiencing pain or paraesthesia during block insertion. Other authors report a 2% incidence of neurapraxia lasting up to three months following brachial plexus block [227]. Pneumothorax may also occur in association with supraclavicular brachial plexus blocks in up to 6.1% of patients [228, 229].

The use of local anaesthetic blocks for eye surgery is now common practice. The risk of retrobulbar haemorrhage after retrobulbar block has been reported as being between 0.5 and 44:10 000 [230, 231]. Brainstem anaesthesia occurs in \sim 7–29:10 000 cases [232]. Globe perforation occurs in 1:12 000 cases [233], particularly in

eyeballs with an axial length > 26 mm. Transient complications, such as ptosis and diplopia, are also seen. The incidence of postoperative ptosis at 24 h after surgery is reported to be up to 50%, with residual problems in 20% of patients at one month [234]. Between 8 and 70% of patients suffer from diplopia at 24 h after surgery, with wide variations due to the different local anaesthetics used [235, 236].

Discussion

Anaesthesia is generally perceived to be safe by the public, by surgeons and physicians, and by anaesthetists. This perception is, as we have shown, somewhat optimistic. Nevertheless, such a perception will exacerbate the anger of patients who suffer ill effects related to anaesthesia, particularly if the surgery for which anaesthesia is indicated is itself of marginal benefit or is only an optional or cosmetic procedure. Thus, if the surgery is judged by the anaesthetist to be of 'marginal' benefit or to be only of cosmetic benefit, the anaesthetist would be wise to be most diligent in explaining the attendant risks of anaesthesia, the reasons for particular techniques, and to allow the patient to become involved in the choices of anaesthetic management whenever possible.

Discussion with the patient of the risks of anaesthesia could potentially encourage anxiety about anaesthesia. Alfidi [237] showed that about a third of patients had increased anxiety after being given detailed information about angiography. However, a more recent Australian study on patient responses to detailed information about anaesthetic complications indicated that there was no increase in anxiety levels when detailed information was made available [238]. Thus, in the current medicolegal climate, patients should rarely have detailed information about the risks of anaesthesia or surgery withheld on the grounds that they are likely to suffer adversely from such information.

When detailing the specific risks of anaesthesia and surgery, patients should also be reminded of the risks they are subjected to in daily life in order to place these medical risks in perspective. Calman [239] has highlighted that the perceptions of any given risks are subjective, personality dependent, often subconscious and without any logical or rational basis. Both the patients' and the anaesthetists' perceptions will contribute to the discussion of risks. Anaesthetists should recognise that their own bias may influence the presentation of anaesthetic risks, and that 'informed consent' may suffer as a consequence.

The perception of risk is modified by a number of factors [240]:

Probability of occurrence. The true incidence requires a large population sample, and may be susceptible to regional bias in techniques, exposure bias (catastrophic or dramatic overpublicity) and compression/expansion bias (underestimates of large risks or overestimates of small risks). The specific objective of this review is to develop a list of the 'best guess' estimates of incidences for the important and common complications of anaesthesia based on the published literature. We have done this in Tables 7 and 8 with full realisation that there are many inadequacies in these illustrations, in which the numerators are dependent upon publication bias, medicolegal constraints and the reporting reticence inherent with medical complications. There are also often difficulties inherent in estimating the denominators. We hope that deficiencies and inaccuracies will be noted and improvements made that will more accurately reflect the true incidences of anaesthetic complications. In any event, these are global 'best guess' estimates, and each anaesthetist should substitute his or her own data for these global estimates wherever possible. Individual anaesthetists may have better or worse results for various anaesthetic complications depending on different levels of experience, frequency of use of various techniques, type of surgical and anaesthetic subspecialisation, patient population, etc.

Severity. High severity risks such as death, paraplegia and permanent organ failure, even though of very low probability, are perceived as higher overall risks than more common complications with a much greater incidence, such as PONV, sore throat or thrombophlebitis.

Vulnerability. Often denial or optimism and a feeling of immunity or invincibility allow us to go through life dismissive of the risks we take daily. Patients may feel more vulnerable because they are not in control, or, as is the situation with general anaesthesia, they are unconscious and thus have totally lost control of their circumstances. These concerns often magnify the importance of particular risks such as awareness.

Controllability. Loss of conscious choice, with a feeling of loss of control over events, increases the feeling of vulnerability and this is very pertinent to anaesthesia, particularly where general anaesthesia or heavy sedation is used. The issue of consent, together with a choice of clinical alternatives, is important, as patients who perceive that they have had adequate and realistic information, with the choice of different anaesthetic options, will be less resentful of any subsequent complications, just as smokers or motorcyclists accept the increased risks of their activities. Familiarity. Patients who have had many major anaesthetic procedures before may be less worried about any inherent risks during future anaesthetics, even though those risks may increase with progression of disease processes and with ageing. Conversely, patients having their first anaesthetic experience may be more worried.

Acceptability and dread. Fear of paraplegia may feature more prominently with anaesthetists than stroke, major myocardial infarction or a patient's death. Cultural or regional expectations may alter these perceptions for both patients and anaesthetists. This particularly applies to the use of regional anaesthesia vs. general anaesthesia, where one or the other is dominant geographically, and hence the expected norm.

Framing or presentation. Particularly when relative risks are discussed with patients, positive framing [241] is better than negative framing. One may quote 90% survival rather than 10% mortality, or that outcomes are twice as good with one management regimen than with another, although the actual differences may only be between 0.005% and 0.01% mortality! However, such 'bias' should not impede discussion of the true incidence or real clinical significance with patients. Some anaesthetists may feel immune to complications that could occur in their practice either because of bravado or due to a misguided sense of the true incidence of a particular complication. Such anaesthetists may present too optimistic an opinion of the real risk to their patients. Unless anaesthetists have their own quality control figures to prove that their practice is better than the published data, we consider that they should use published data for risks of anaesthesia rather than anecdotal evidence. Where individual results are worse than published data, there may be good reasons, but it behoves the particular anaesthetist to be aware of this fact and to be honest with their patients, or be subject to the criticisms levelled at the Bristol Royal Infirmary paediatric cardiac surgeons [242].

Patients, particularly at times of anxiety related to surgery, will have great difficulty assimilating and retaining details such as incidences of complications. It is unlikely that detail will be assimilated or retained from a short visit the night before operation or just before the day of admission for surgery. Even a more leisurely preadmission consultation, remote in time from surgery, may be too detailed for full comprehension. Overload of information is sometimes used as an excuse for a very limited discussion of anaesthetic complications. Such an excuse is unlikely to satisfy an aggrieved patient or be used as defence against the legal challenge that follows. There would appear to be a place for a succinct leaflet given in advance to patients, listing the complications and placing them generally in perspective to everyday risks [243]. Such a leaflet may then be studied at leisure by the patient, who could then seek further clarification as

Presentation of relative everyday risks to patients may help to place the risks of potential anaesthetic complications in perspective. The logarithmic community cluster classification (Table 9) [244], which is based on Calman's

Table 7 Predicted incidence of complications of anaesthesia.

Mortality and morbidity	Incidence	Rate per 10 000 population	Remarks	
Total peri-operative deaths (within 30 days)	\sim 1:200 (elective surgery)	50		
, , , , , , , , , , , , , , , , , , ,	~ 1:40 (emergency surgery)	250		
	\sim × 2 (60–79 years)			
	\sim ×5 (80–89 years)			
	\sim ×7 (>90 years)			
Death related to anaesthesia	\sim 1:50 000 (anaesthesia-related)	0.2		
	\sim 1:100 000 (ASA physical status I and II)	0.1		
Cardiac arrest	1:10 000 - 1:20 000 (general anaesthesia)	0.5-1.0	Mortality ~ 1:15 000-1:150 000	
	~ 1:3 000 (local anaesthesia)	3	•	
	~ 1:1 500 (Spinal: 25% fatal)	7		
Myocardial re-infarction	~ 1:20 (0-3 months after myocardial	500		
,	infarction			
	\sim 1:40 (4-6 months after myocardial	250		
	infarction)	230		
Respiratory complications	marctiony			
Aspiration during general anaesthesia	~ 1:3 000	3	×4 in emergencies	
rispiration during general anaestriesia	1.5 000	3	×3 in obstetrics	
	\sim 1:60 000 (Death)	0.16	×3 iii obstetiits	
Difficult intubation	~ 1:50	200		
Failure to intubate	~ 1:500 ~ 1:500		Obstetrics ∼ 1:250	
		20	Obstetrics ~ 1:250	
Failure to intubate & ventilate	~ 1:5000	2		
Postoperative cognitive dysfunction	\sim 1:4 at 1 week	2500	Regional anaesthesia	
(> 60 years)	\sim 1:10 at 3 months	1000	pprox General anaethesia	
	\sim 1:100 permanent	100		
Postoperative delirium	\sim 1:7 (general surgery)	1400	× 3 >75 years	
	up to \sim 1:2 for elderly fractures neck	5000	×3 if requiring intensive care	
	of femur			
Drowsiness	∼ 1:2	5000	Day surgery	
Dizziness	∼ 1:5	2000	Day surgery	
Headache	~ 1:5	2000	, , ,	
Cerebrovascular accident (CVA)	\sim 1:50 if previous stroke	200	46% mortality	
, ,	\sim 1:100 general surgery	100	60% mortality if previous CVA (~ 1:700 in the non-surgical population)	
	\sim 1:20 head and neck surgery	500	population	
	(\sim 1:700 in the non-surgical population)	300		
C		700		
Carotid endarterectomy (CVA + death)	~ 1:15 if symptomatic	700		
(5)	\sim 1:25 if asymptomatic	400		
(Disabling CVA + death)	∼ 1:50	200		
Awareness				
with pain	~ 1:3000	3	2/3 with neuromuscular blockade	
without pain	~ 1:300	30	1/3 without neuromuscular	
			blockade	
Total intravenous anaesthesia	~ 1:500	20		
Anaphylaxis	~ 1:10 000	1		
Deafness				
'idiopathic' (general anaesthesia)	~ 1:10 000	1	\sim 1:1 000 cardiac surgery	
transient after spinal anaesthesia	∼ 1:7	1500		
oss of vision	~ 1:125 000	0.08		
	\sim 1:100 (cardiac surgery)	100		
Pain	~ 1:3 (moderate)	3000		
(after major surgery)	~ 1:10 (severe)	1000		
(day surgery)	~ 1:2	5000		
Postoperative nausea and vomiting (PONV)	~ 1:4	2500	2/3 nausea and 1/3 vomiting	
ostoperative nausea and voilliting (FONV)	· · · 1.47	2300	Female:male 3:1	
Sore throat	\sim 1:2 (if tracheal tube)	5000	i cinale.iliale 3.1	
ooie unoat	· ·	5000		
	~ 1:5 (if laryngeal mask)	2000		
Dental damage	\sim 1:10 (if facemask only)	1000		
Dental damage	4.5000			
requiring intervention	~ 1:5000	2		
all dental damage \sim 1:100		100		
all oral trauma after tracheal intubation \sim 1:20		500		

Table 7 (Continued).

Mortality and morbidity	Incidence	Rate per 10 000 population	Remarks
Peripheral nerve injury	\sim 1:300 ulnar neuropathy	30	
(general anaesthesia)	\sim 1:1 000 (other nerves)	10	
Thrombophlebitis	\sim 1–2:20 water-soluble drugs	500-1000	
•	\sim 1:4 propylene glycol-based	2500	
Arterial cannulation complications	< 1:100 (permanent)	<100	
Pulmonary artery perforation	\sim 1:2000 $^{\circ}$	5	
Arterial Puncture (during central venous cannulation)			
internal jugular vein cannulation	∼ 1:35	350	
subclavian vein cannulation	~ 1:200	50	

 Table 8 Predicted incidence of complications of regional anaesthesia.

	Incidence	Rate per 10 000 population	Remarks
Paraplegia	~ 1:100 000	0.1	
Permanent Nerve Injury			
spinal	1–3:10 000	1–3	
epidural	0.3–10:10 000	0.3–10	
peripheral nerve block	~ 1:5000	2	2% brachial plexus neuropraxia lasting >3 months
Epidural haematoma	\sim 1:150 000 (epidural)	0.07	\sim 1:1 000 000 (spontaneous)
	~ 1:200 000 (spinal)	0.05	~ 1:14 000 (USA) ~ 1:2 250 000 (Europe) ~ 1:10 000 (spontaneous)
Epidural abscess	1:2000-1:7500	0.7–5.0	,
Transient neural complications	1:1000-1:10 000 (epidural)	1-10 (epidural)	
•	1:125–1:2 500 (spinal)	4–80 (spinal)	
Transient radicular irritation (spinal)	Up to 1:3 (heavy lidocaine and mepivacaine)	3000	
Cardiac arrest	\sim 1:1500 (spinal)	5 (spinal)	
	\sim 1:3000 (local anaesthesia)	3 (local anaesthesia)	
	\sim 1:10 000 (epidural)	1 (epidural)	
	\sim 1:10 000 (regional blocks)	1 (regional)	
Post-dural puncture headache	~ 1:100	100	80% following inadvertent
	\sim 1:10 (day surgery)	1000	dural tap
			Blood patch 70–100% Immediate success but headaches recur in 30–50%
Backache	< 1 h surgery \sim 20%	2000	GA pprox LA
	$>$ 4 h surgery \sim 50%	5000	
Urinary dysfunction	∼ 1:50	200	
Pneumothorax	\sim 1:20 (supraclavicular blocks)	500	
Systemic LA toxicity	\sim 1:10 000 (epidural)	1	
	\sim 1:1500 (regional blocks)	7	
Cerebral seizures	\sim 1:4000 (intravenous regional anaesthesia)	2.5	Axillary ∼1:1000
	\sim 1:500 (brachial plexus)	20	Supraclavicular ∼ 1:125
Eye blocks			·
Retrobulbar haemorrhage	1: 250-1:20 000	0.5–40	
Brainstem anaesthesia	~ 1:700	15	
Globe perforation	\sim 1:10 000	1	
Ptosis – transient after eye block	\sim 1:2 at 24 h	5000	
•	\sim 1:5 at 1 month	2000	
Diplopia – transient after eye block	8–70%	800–7000	

verbal scale [239], is one way to describe the relative risks that may also be presented graphically [245].

Another comparison is with the lottery probability scale [246], but this seems unnecessarily complex and has the

wrong emphasis, as gamblers are invariably too optimistic and patients are usually too pessimistic.

Whilst we have focused on the risks of anaesthesia, there are other aspects as indicated by Paul de Gondi's

Table 9 Community cluster logarithmic scale of risk classification (modified from Calman & Roystone [244] and Admas & Smith [247]).

Risk level	Calman's Verbal Scale	Community	Community examples	Anaesthetic or medical examples
1:1–9	Very High	Sibling	Genetic dominant	Postoperative nausea and vomiting 1:4 Dizziness 1:5 Headache 1:5
1:10–99	High	Family	Genetic recessive	All oral trauma following intubation 1:20 Emergency surgery death 1:40 Difficult intubation 1:50
1: 100–999	Moderate	Street	Deaths per year 1: 100	Peri-operative death 1:200 Awareness without pain 1:300 Failure to intubate 1:500
1: 1000–9999	Low	Village	Traffic deaths per year 1:8000	Awareness with pain 1:3000 Aspiration 1: 3000 Cardiac arrest 1:3000 (local anaesthesia) Epidural abscess \sim 1:5000 (local anaesthesia) Failure to intubate and ventilate 1:5000
1:10 000–99 999	Very Low	Small Town	Accidental deaths at home per year 1:11 000	Anaphylaxis 1:10 000 Spontaneous epidural abscess 1:10 000 Cardiac arrest ~ 1:15 000 (general anaesthesia) Death (related to anaesthesia) 1:50 000
1:100 000–999 999	Minimal	Large Town	Rail accidents per year 1:140 000	Loss of vision (general anaesthesia) 1:125 000 Paraplegia (local anaesthesia) 1:100 000 Epidural haematoma 1:150 000–1:200 000 Death due solely to anaesthesia 1:180 000
1:1 000 000–9 999 999	Negligible	City	Six balls in UK National Lottery 1:2 796 763	Spontaneous epidural haematoma 1:1 000 000
1:10 000 000–99 999 999	Minute	Country	Lightning deaths per year 1:10 000 000	

quotation at the beginning of this article. The benefits of different management options (medical, surgical and anaesthetic) also have varying probabilities and 'spin' which we have not attempted to address here. The mnemonic *BRAN* offers a useful approach when assessing the risks of a course of action, and includes the **Benefits**, **Risks**, **Alternatives** and what would happen if **Nothing** were done [240]. We believe we have made a start by establishing a listing of the incidences of various anaesthetic complications. Much work remains to be done to establish incidences of the inherent benefits in different clinical situations, and also for alternatives including the option of doing nothing. We have no doubt that better and more complete listings of incidences of anaesthetic complications will be developed. This is but a start.

References

- 1 Rogers v. Whitaker (1990) 4 Medical LR79 (High Court of Australia); 1992 109 ALR265; 67 ALJR 47; Medical Law R 331; 175 LLR 479.
- 2 Sidaway v Board of Governors of Bethlem Royal Hospital and the Maudsley Hospital (1985). AC 871.
- 3 Chappel v Hart (1998). 72 ALJR 1344.

- 4 Scott RJ. Chappel v Hart: the High Court considers causation of damage from a surgeon's negligent failure to warn. *Medical Journal of Australia* 2000; **172**: 134–6.
- 5 National Health and Medical Research Council. General Guidelines for Medical Practitioners on Providing Information to Patients. Canberra: Australian Government Publishing Service, 1993.
- 6 General Medical Council. Seeking Patients' Consent: The Ethical Considerations. London: General Medical Council, 1999.
- 7 Rosenberg v Percival (2001) HCA 18.
- 8 Buck N, Devlin HB, Lunn JN. The Report of a Confidential Enquiry into Peri-Operative Deaths. London: Nuffield Provincial Hospitals Trust and the King's Fund, 1987.
- 9 Australian and New Zealand College of Anaesthetists. A Review of Anaesthesia Related Mortality 1997–99 (Appendix 3). Melbourne, Australia: ANZCA, 2002.
- 10 Department of Health. NHS performance indicators. Available at: http://www.doh.gov.uk/nhsperformance indicators
- 11 Beecher HK, Todd DP. A study of the deaths associated with anaesthesia and surgery. Annals of Surgery 1954; 140: 2–34
- 12 Vacanti CJ, Van Houten RJ, Hill RC. A statistical analysis of the relationship of physical status to postoperative

- mortality in 68,388 cases. *Anesthesia and Analgesia* 1970; **49**: 564–6.
- 13 Marx GF, Mateo CV, Orkin LR. Computer analysis of postanesthetic deaths. *Anesthesiology* 1973; 39: 54–8.
- 14 Hovi-Viander M. Death associated with anaesthesia in Finland. *British Journal of Anaesthesia* 1980; **52**: 483–9.
- 15 Turnbull KW, Fancourt-Smith PF, Banting GC. Death within 48 hours of anaesthesia at the Vancouver General Hospital. Canadian Anaesthetic Society Journal 1980; 27: 159–63.
- 16 Lunn JN, Mushin WW. Mortality associated with anesthesia. Anaesthesia 1982; 37: 856.
- 17 Gibbs JM. The anaesthetic mortality assessment committee 1979–1984. New Zealand Medical Journal 1986; 99: 55–9.
- 18 Tiret L, Desmonts JM, Hatton F, Vourc'h G. Complications associated with anaesthesia: a prospective survey in France. Canadian Anaesthetic Society Journal 1986; 33: 336–44.
- 19 Olsson GL, Hallen B. Cardiac arrest during anaesthesia. A computer aided study in 250 543 anaesthetics. Acta Anaesthesiologica Scandinavica 1988; 32: 653–4.
- 20 Pederson T, Johansen SH. Serious morbidity attributable to anaesthesia. Considerations for prevention. *Anaesthesia* 1989; 44: 504–8.
- 21 Harrison GG. Death due to anaesthesia at Groote Schuur Hospital, Cape Town 1956–87. Part 1. Incidence. South African Medical Journal 1990; 77: 412–15.
- 22 National Health and Medical Research Council. Report on Deaths Associated with Anaesthesia in Australia 1985–87. Canberra: Australian Government Publishing Service, 1990.
- 23 National Health and Medical Research Council. Report on Deaths Associated with Anaesthesia in Australia 1988–90. Canberra: Australian Government Publishing Service, 1992.
- 24 Wang L, Hagerdal M. Reported anaesthetic complications during an 11-year period. A retrospective study. *Acta Anaesthesiologica Scandinavica* 1992; 36: 234–40.
- 25 Warden JC, Borton CL, Horan BF. Mortality associated with anaesthesia in NSW 1984–1990. Medical Journal of Australia 1994; 161: 585–93.
- 26 Tikkanen J, Hovi-Viander M. Death associated with anaesthesia and surgery in Finland in 1986 compared to 1975. Acta Anaesthesiologica Scandinavica 1995; 39: 262–7.
- 27 McKenzie AG. Mortality associated with anaesthesia at Zimbabwean teaching hospitals. South African Medical Journal 1996; 86: 338–42.
- 28 Eagle CC, Davis NJ. Report of the Anaesthetic Mortality Committee of Western Australia 1990–1995. Anaesthesia and Intensive Care 1997; 25: 51–9.
- 29 Horan BF (Committee on Anaesthetic Mortality). Australian and New Zealand College of Anaesthetists: Anaesthesia Related Mortality in Australia 1991–1993. Melbourne, Australia: ANZCA, 1998.
- 30 Davis NJ (Committee on Anaesthetic Mortality) Australian and New Zealand College of Anaesthetists: Anaesthesia Related Mortality in Australia 1994–1996. Melbourne, Australia: ANZCA, 1999.

- 31 NHS Executive. Quality and Performance in the NHS. NHS Performance Indicators. London: NHSE, July 2000.
- 32 Arbous MS, Grobbee DE, Van Kleef JW *et al.* Mortality associated with anaesthesia: a qualitative analysis to identify risk factors. *Anaesthesia* 2001; **56**: 1141–53.
- 33 Kawashima Y, Seo N, Morita K et al. Annual study of perioperative mortality and morbidity for the year of 1999 in Japan: the outlines – report of the Japan Society of Anesthesiologists Committee on Operating Room Safety. Masui 2001; 50: 1260–74.
- 34 Kawashima Y, Seo N, Morita K et al. Annual study of anesthesia-related mortality and morbidity in the year 2000 in Japan: the outlines – Report of Japanese Society of Anesthesiologists Committee on Operating Room Safety. Masui 2002; 51: 1032–47.
- 35 Paramesh K, Dunkley C. Lessons from the National Confidential Enquiry into Perioperative Deaths. In: McConachie I, ed. *Anaesthesia for the High Risk Patient*. London: Greenwich Medical Media, 2002: 41–9.
- 36 Clifton BS, Hotten WT. Deaths associated with anaesthesia. *British Journal of Anaesthesia* 1963; **35**: 250–9.
- 37 Jin F, Chung F. Minimizing perioperative adverse events in the elderly. *British Journal of Anaesthesia* 2001; 87: 608–24.
- 38 NCEPOD. Then and Now. The 2000 Report of the National Confidential Enquiry into Perioperative Deaths. London: NCEPOD, 2000.
- 39 Warner MA, Shields SE, Chute CG. Major morbidity and mortality within 1 month of ambulatory surgery and anesthesia. *Journal of the American Medical Association* 1993; 270: 1437–41.
- 40 Duncan PG, Cohen MM, Tweed WA et al. The Canadian four-centre study of anaesthetic outcomes. III. Are anaesthetic complications predictable in day surgical practice? Canadian Journal of Anesthesia 1992; 39: 440–8.
- 41 Osborne GA, Rudkin GE. Outcome after day-care surgery in a major teaching hospital. *Anaesthesia and Intensive Care* 1993: 21: 822–7.
- 42 Chung F, Mezei G, Tong D. Adverse events in ambulatory surgery. A comparison between elderly and younger patients. *Canadian Journal of Anesthesia* 1999; 46: 309–32.
- 43 Natof HE. Complications associated with ambulatory surgery. *Journal of the American Medical Association* 1980; 244: 1116–18.
- 44 HMSO, TSO. Reports on Confidential Enquiries in to Maternal Deaths in England and Wales 1952–1984, United Kingdom 1985–1999. London: HMSO and TSO, 2002.
- 45 Rout C. Maternal mortality and anaesthesia in Africa: a South African perspective. *International Journal of Obstetric Anaesthesia* 2002; **11**: 77–80.
- 46 UNICEF. The progress of nations: league tables of maternal death Available at: http://www.unicef.org/pon96/leag/wom.htm.
- 47 Pattinson B. Saving mothers. Report on Confidential Enquiries into maternal deaths in South Africa 1998. Available at: http://www.doh.gov.za/docs/reports/mothers/contents.html

- 48 Pederson T, Eliasen K, Henriksen E. A prospective study of mortality associated with anaesthesia and surgery: risk indicators of mortality in hospital. *Acta Anaesthesiologica Scandinavica* 1990; 34: 176–82.
- 49 Ackerman RJ, Vogel RL, Johnson LA, Ashley DW, Solis MM. Surgery in nonagenarians: morbidity, mortality and functional outcome. *Journal of Family Practice* 1995; 40: 129–35.
- 50 Djokovic JL, Hedley-White J. Prediction of outcome of surgery and anaesthesia in patients over 80. Journal of the American Medical Association 1979; 242: 2301–6.
- 51 Hosking MP, Lobdell CM, Warner MA, Offord KP, Melton LJ. Anaesthesia for patients over 90 years of age. Outcomes after regional and general anaesthetic techniques for two common surgical procedures. *Anaesthesia* 1989; 44: 142–7
- 52 NCEPOD. Extremes of Age. The 1999 Report of the National Confidential Enquiry into Perioperative Deaths. London: NCEPOD, 1999.
- 53 Van Der Walt J. Searching for the Holy Grail: measuring risk in paediatric anaesthesia. *Paediatric Anaesthesia* 2001; 11: 637–41.
- 54 Cheney FW, Posner K, Caplan RA, Ward RJ. Standard of care and anaesthesia liability. *Journal of the American Medical Association* 1989; 261: 1599–603.
- 55 Bothner U, Georgieff M, Schwilk B. Building a large scale perioperative anaesthesia outcome tracking database: methodology, implementation and experiences from one provider within the German quality project. *British Journal* of Anaesthesia 2000; 85: 271–80.
- 56 Cohen MM, Duncan PG, Pope WD, Wolkenstein C. A survey of 112 000 anaesthetics at one teaching hospital 1975–1983. Canadian Anaesthetic Society Journal 1986; 33: 22–31.
- 57 Cooper JB, Cullen DJ, Nemeskal R et al. Effects of information feedback and pulse oximetry on the incidence of anesthesia complications. Anesthesiology 1987; 67: 686–94.
- 58 Zelcer J, Wells DG. Anaesthetic-related recovery room complications. *Anaesthesia and Intensive Care* 1987; 15: 168–74.
- 59 Hines R, Barash PG, Watrous G, O'Connor T. Complications occurring in the postanaesthesia care unit: a survey. *Anesthesia and Analgesia* 1992; **74**: 503–9.
- 60 Moller JT, Pederson T, Rasmussen LS et al. Randomized evaluation of pulse oximetry in 20 802 patients: 1. Design, demography, pulse oximetry failure rate and overall complication rate. Anesthesiology 1993; 70: 135–40.
- 61 Rose DK, Cohen MM, Wigglesworth DF, De Boer DP. Critical respiratory events in the post anaesthesia care unit: patient, surgical and anaesthetic factors. *Anesthesiology* 1994; 81: 410–81.
- 62 Ouchterlony J, Arvidsson L, Sjostedt L, Svardsudd K. Peroperative and immediate postoperative adverse events in patients undergoing elective general and orthopaedic surgery. Acta Anaesthesiologica Scandinavica 1995; 39: 643–52.

- 63 Schwilk B, Muche R, Bothner U, Goertz A, Friesdorf W, Georgieff M. Quality control in anesthesiology. Results of a prospective study following the recommendations of the German Society of Anesthesiology and Intensive Care. *Anaesthesist* 1995; **44**: 242–9.
- 64 Hunter MJ, Molinaro AM. Morbidity and mortality with outpatient anesthesia: the experience of a residency training program. *Journal of Oral Maxillofacial Surgery* 1997; 55: 684–7; discussion 687–8.
- 65 Schwilk B, Muche R, Treiber H, Brinkmann A, Georgieff M, Bothner U. A cross-validated multifactorial index of perioperative risks in adults undergoing anaesthesia for non-cardiac surgery. Analysis of perioperative events in 26907 anaesthetic procedures. *Journal of Clinical Monitoring and Computing* 1998; 14: 283–94.
- 66 Chung F, Mezei G, Tong D. Adverse events in ambulatory surgery. A comparison between elderly and younger patients. *Canadian Journal of Anesthesia* 1999; 46: 309–32.
- 67 Fasting S, Gisvold SE. Serious intraoperative problems a five year review of 83 844 anesthetics. *Canadian Journal of Anesthesia* 2002; **49**: 545–53.
- 68 Gaba DM. Anaesthesiology as a model for patient safety in health care. British Medical Journal 2000; 320: 785–8.
- 69 Webb RC, Currie M, Morgan CA et al. The Australian Incident Monitoring Study: an analysis of 2000 incident reports. *Anaesthesia and Intensive Care* 2002; **21**: 520–8.
- 70 Pottecher T, Tiret L, Desmonts JM, Hatton F, Bilaine J, Otteni JC. Cardiac arrest related to anaesthesia: a prospective survey in France 1978–1982. European Journal of Anaesthesiology 1984; 1: 305–18.
- 71 Pederson T, Eliasen K, Henriksen E. A prospective study of risk factors and cardiopulmonary complications associated with anaesthesia and surgery: risk indicators of cardiopulmonary morbidity. *Acta Anaesthesiologica Scandinavica* 1990; 34: 144–55.
- 72 Aubas S, Biboulet P, Daures JP, Cailar J. Incidence and etiology of cardiac arrest occurring during the preoperative period and in the recovery room. Apropos of 102 468 anaesthesia cases. *Annales Francaises d'Anesthesie et de Reanimation* 1991; **10**: 436–42.
- 73 Keenan RL, Boyan CP. Decreasing frequency of anesthetic cardiac arrests. *Journal of Clinical Anesthesia* 1991; 3: 354–7.
- 74 Auroy Y, Narchi P, Messiah A, Litt L, Rouvier B, Samii K. Serious complications related to regional anaesthesia.
 Results of a prospective survey in France. Anesthesiology 1997; 87: 479–86.
- 75 Morray JP, Geiduschek JM, Ramamoorthy C et al. Anesthesia-related cardiac arrest in children: initial findings of the Pediatric Perioperative Cardiac Arrest (POCA) Registry. Anesthesiology 2000; 93: 6–14.
- 76 Biboulet P, Aubas P, Dubourdieu J, Rubenovitch J, Capdevila X, d'Athis F. Fatal and non-fatal cardiac arrests related to anesthesia. *Canadian Journal of Anesthesia* 2001; 48: 326–32.
- 77 Newland MC, Ellis SJ, Lydiatt CA et al. Anesthetic-related cardiac arrest and its mortality: a report covering 72,959

- anesthetics over 10 years from a US teaching hospital. *Anesthesiology* 2002; **97**: 108–15.
- 78 Brooks-Brunn JA. Postoperative atelectasis and pneumonia. Heart and Lung 1995; 24 (2): 94–115.
- 79 Englehardt T, Webster NR. Pulmonary aspiration of gastric contents. *British Journal of Anaesthesia* 1999; **83**: 453–60.
- 80 Olsson GL, Hallen B, Hambraeus-Jonzon K. Aspiration during anaesthesia. A computer-aided study of 185 358 anaesthetics. Acta Anaesthesiologica Scandinavica 1986; 30: 84–92.
- 81 Warner MA, Warner ME, Weber JG. Clinical significance of pulmonary aspiration during the perioperative period. *Anesthesiology* 1993; **78**: 56–62.
- 82 Mellin-Olsen J, Fasting S, Gisvold SE. Routine preoperative gastric emptying is seldom indicated. A study of 85 594 anaesthetics with special focus on aspiration pneumonia. Acta Anaesthesiologica Scandinavica 1996; 40: 1184–8.
- 83 Phillips S, Daborn AK, Hatch DJ. Preoperative fasting for paediatric anaesthesia. *British Journal of Anaesthesia* 1994; 73: 529–36.
- 84 Borland LM, Sereika SM, Woelfel SK, Saitz EW, Carrilo PA, Motoyama EK. Pulmonary aspiration in pediatric patients during general anesthesia: incidence and outcome. *Journal of Clinical Anesthesia* 1998; **19**: 95–102.
- 85 Tiret L, Nivoche Y, Hatton F, Desmonts JM, Vourc'h G. Complications related to anaesthesia in infants and children. A prospective survey of 40 240 anaesthetics. *British Journal of Anaesthesia* 1988; 61: 263–9.
- 86 Dindelli M, La Rosa M, Rossi R. Incidence and complications of the aspiration of gastric contents syndrome during cesarean section in general anaesthesia. *Annali Di Ostetricia, Ginecologica, Medicina Perinatale (Milano)* 1992; 112: 376–84.
- 87 La Rosa M, Piva L, Dindelli M, Pagnoni B. Aspiration syndrome in Cesarean section. Our experience from 1980 to 1990. *Minerva Anestesiologie* 1992; **58**: 1213–20.
- 88 Soreide E, Bjornestad E, Steen PA. An audit of perioperative aspiration pneumonitis in gynecological and obstetric patients. *Acta Anaesthesiologica Scandinavica* 1996; **40**: 14–19.
- 89 Crosby ET, Cooper RM, Douglas MJ. The unanticipated difficult airway with recommendations for management. *Canadian Journal of Anesthesia* 1998; **45**: 757–76.
- 90 Yentis SM, Brighouse D, May A, Bogod D, Elton C. Section 2 – pregnancy. IV. Anaesthetic complications. In: WB Saunders, ed. Analgesia, Anaesthesia and Pregnancy – A Practical Guide. London: Harcourt, 2002.
- 91 Barnardo PD, Jenkins JG. Failed tracheal intubation in obstetrics: a 6-year review in a UK region. *Anaesthesia* 2000; **55**: 690–4.
- 92 Harmer M. Difficult and failed intubation in obstetrics. *International Journal of Obstetric Anesthesia* 1997; **6**: 25–31.
- 93 Wroblewski F, La Due JS. Myocardial infarction adds a postoperative complication of major surgery. *Journal of the American Medical Association* 1952; **150**: 1212–16.
- 94 Shah KB, Kleinman BS, Rao TLK et al. Angina and other risk factors in patients with cardiac diseases undergoing

- non-cardiac operations. *Anesthesia and Analgesia* 1990; **70**: 240–7.
- 95 Rao TL, Jacobs KH, El-Etr AA. Reinfarction following anesthesia in patients with myocardial infarction. *Anesthesiology* 1983; **59**: 499–505.
- 96 Goldman L, Caldera DL, Nussbaum SR. Multifactorial index of cardiac risk in non-cardiac surgical procedures. New England Journal of Medicine 1977; 297: 845–50.
- 97 Detsky AS, Abrams HB, Forbath N, Scott JG, Hillard JR. Cardiac assessment for patients undergoing non-cardiac surgery. A multifactorial clinical risk index. *Archives of International Medicine* 1986; **146**: 2131–4.
- 98 Lee TH, Marcantonio ER, Mangione CM et al. Derivation and prospective validation of a simple index for prediction of cardiac risk of major noncardiac surgery. *Circulation* 1999; 100: 1043–9.
- 99 Boersma E, Poldermans D, Bax J et al. for the DECREASE Study Group. Predictors of cardiac events after major vascular surgery: role of the clinical characteristics, dobutamine echocardiography, and beta-blocker therapy. *Journal* of the American Medical Association 2001; 285: 1865–73.
- 100 Eagle KA, Berger PB, Calkins H et al. ACC:AHA guideline update for perioperative cardiovascular evaluation for non-cardiac surgery: a report of the American College of Cardiology: American Heart Association Task Force on Practice Guidelines (Committee to Update the 1996 Guidelines on Perioperative Cardiovascular Evaluation for Non-Cardiac Surgery). Available at: http://www.acc.org/ clinical/guidelines/perio/dirIndex.htm.
- 101 Scheer BV, Perel A, Pfeiffer UJ. Clinical review. Complications and risk factors of peripheral arterial catheters used for haemodynamic monitoring in anaesthesia and intensive care medicine. Critical Care 2002; 6: 199–204.
- 102 Matthews NT, Worthley UG. Immediate problems associated with infraclavicular subclavian catheterisation; a comparison between left and right sides. *Anaesthesia and Intensive Care* 1982; 10: 113–15.
- 103 Eerola R, Kaukinen L, Kaukinen L. Analysis of 13 800 subclavian vein catheterizations. *Acta Anaesthesiologica Scandinavica* 1985; **29**: 193–7.
- 104 Kamlesh BS, Tadikonda LKR, Laughlin S, Adel AEE. A review of pulmonary artery catheterisations in 6245 patients. Anesthesiology 1984; 61: 271–5.
- 105 Ruesch S, Walder B, Tramer MR. Complications of central venous catheters: internal jugular versus subclavian access – a systematic review. *Critical Care Medicine* 2002; 30: 454–60.
- 106 Merrer J, De Jonghe B, Golliot F et al. and the French Catheter Study Group in Intensive Care. Complications of femoral and subclavian venous catheterization in critically ill patients: a randomized controlled trial. *Journal of the American Medical Association* 2001; 286: 700–7.
- 107 Randolph AG, Cook DJ, Gonzales CA, Pribble CG. Ultrasound guidance for placement of central venous catheters: a meta-analysis of the literature. *Critical Care Medicine* 1996; 24: 2053–8.
- 108 Driscoll EJ, Gelfman SS, Sweet JB, Butler DP, Wirdzck PR, Medlin T. Thrombophlebitis after intravenous use of

- anaesthesia and sedation: its incidence and natural history. *Journal of Oral Surgery* 1979; **37**: 809–15.
- 109 Schou Olesen A, Huttal MS, Hole P. Venous sequelae following the injection of etomidate or thiopentone IV. British Journal of Anaesthesia 1984; 56: 171–3.
- 110 Kortilla K, Aromaa U. Venous complications after intravenous injection of diazepam, flunitrazepam, thiopentone and etomidate. Acta Anaesthesiologica Scandinavica 1980; 24: 227–30.
- 111 Clarke RS. Adverse effects of intravenously administered drugs used in anaesthetic practice. *Drugs* 1981; 22: 26–41.
- 112 Moller JT, Cluitmans P, Rasmussen LS et al. Long term postoperative cognitive dysfunction in the elderly: ISPOCD 1 study. Lancet 1998; 351: 857–61.
- 113 Williams-Russo P, Sharrock NE, Mattis S, Szatrowski TP, Charlson ME. Cognitive effects after epidural vs. general anesthesia in older adults. A randomized trial. *Journal of the American Medical Association* 1995; 274: 44–50.
- 114 Abildstrom H, Rentowl P, Hanning CD, Rasmussen H, Kristensen PA, Moller JT. Cognitive dysfunction 1–2 years after non-cardiac surgery in the elderly. ISPOCD group. International Study of Post-Operative Cognitive Dysfunction. Acta Anaesthesiologica Scandinavica 2000; 44: 1246–51.
- 115 O'Keefe ST, Chonchubhair AN. Post-operative delirium in the elderly. *British Journal of Anaesthesia* 1994; **73**: 673–87.
- 116 Sharpe P, Hanning C. Neurological sequelae. In: Van Aken H, ed. Best Practice and Research Clinical Anaesthesiology, London: Bailliere Tindall, 1999: 451–63.
- 117 Dodds C, Allison J. Postoperative cognitive deficit in the elderly surgical patient. *British Journal of Anaesthesia* 1998; 81: 462.
- 118 Berggren D, Gustafson Y, Eriksson B et al. Postoperative confusion after anaesthesia in elderly patients with femoral neck fractures. *Anesthesia and Analgesia* 1987; **66**: 497–504.
- 119 Hinton R.C. Thrombosis and cerebrovascular disease. Medical Clinics of North America 1998; 82: 523–44.
- 120 Barnett HJM, Meldrum HE, Eliasziw M. The appropriate use of carotid endarterectomy. *Canadian Medical Association Journal* 2002; **166**: 1169–79.
- 121 European Carotid Surgery Trialists Group. Randomised trial of endarterectomy for recently symptomatic carotid stenosis: final results of the MRC European Carotid Surgery Trial (ECST). *Lancet* 1998; 351: 1379–87.
- 122 Ferguson GG, Eliasziw M, Barr HWK et al. for the North American Symptomatic Carotid Endarterectomy Trial Collaborators. The North American Symptomatic Carotid Endarterectomy Trial. Surgical results in 1415 patients. Stroke 1999; 30: 1751–8.
- 123 Taylor DW, Barnett HJM, Haynes RB et al. for the ASA and Carotid Endarterectomy (ACE) Trial Collaborators. Low-dose and high-dose acetylsalicylic acid for patients undergoing carotid endarterectomy: a randomized controlled trial. Lancet 1999; 353: 2179–84.
- 124 Hankey GJ, Warlow CP, Molyneux AJ. Complications of cerebral angiography for patients with mild carotid territory ischaemia being considered for carotid endarterectomy.

- Journal of Neurology, Neurosurgery and Psychiatry 1990; 53: 542–8.
- 125 Eliasziw M, Rankin RN, Fox AJ et al. for the NASCET Group. Accuracy and prognostic consequences of ultrasonography in identifying severe carotid artery stenosis. *Stroke* 1995; 26: 1747–52.
- 126 Klafta JM, Roizen MF. Current understanding of patients' attitudes toward and preparation for anaesthesia: a review. *Anesthesia and Analgesia* 1996; **83**: 1314–21.
- 127 Heier T, Steen PA. Awareness in anaesthesia: incidence, consequences and prevention. *Acta Anaesthesiologica Scandinavica* 1996; **40**: 1073–86.
- 128 Schwender D, Klasing S, Daunderer M, Madler C, Poppel E, Peter K. Awareness during general anesthesia. Definition, incidence, clinical relevance, causes, avoidance and medicolegal aspects. *Anaesthesist* 1995; 44: 743–54.
- 129 Aitkenhead A. Awareness during anaesthesia: what should the patient be told? *Anaesthesia* 1990; **45**: 351–2.
- 130 Sandin RH, Enlund G, Samuelsson P, Lennmarken C. Awareness during anaesthesia: a prospective case study. *Lancet* 2000; 355: 707–11.
- 131 Nordstrom O, Engstrom AM, Persson S, Sandin R. Incidence of awareness in total i.v. anaesthesia based on propofol, alfentanil and neuromuscular blockade. *Acta Anaesthesiologica Scandinavica* 1997; 41: 978–84.
- 132 Bergman IJ, Kluger MT, Short TG. Awareness during general anaesthesia. A review of 81 cases from the Anaesthetic Incident Monitoring Study. *Anaesthesia* 2002; 57: 549–56.
- 133 Fisher MM, Baldo BA. The incidence and clinical features of anaphylactic reactions during anaesthesia in Australia. Annales Francaises d'Anesthesie et de Reanimation 1993; 12: 97–104.
- 134 Laxenaire MC. Epidemiologie des reactions anaphylactoides peranesthetiques. Quatrieme enquete multicentrique (Juillet 1994–Decembre 1996). *Annales Francaises d'Anesthesie et de Reanimation* 1999; **18**: 796–809.
- 135 Laxenaire MC, Mertes PM. Anaphylaxis during anaesthesia. Results of a 2 year survey in France. *British Journal of Anaesthesia* 2001; 87: 549–58.
- 136 Warner ME, Fronapfel PJ, Hebl JR et al. Perioperative visual changes. Anesthesiology 2002; 96: 855–9.
- 137 Roth S, Thisted RA, Erickson JP, Black S, Schreider BD. Eye injuries after nonocular surgery. A study of 60,965 anesthetics from 1988 to 1992. *Anesthesiology* 1996; 85: 1020–7.
- 138 Cucchiara RF, Black S. Corneal abrasion during anaesthesia and surgery. Anesthesiology 1988; 69: 978–9.
- 139 Gild WM, Posner KL, Caplan RA, Cheney FW. Eye injuries associated with anaesthesia. A Closed Claims analysis. *Anesthesiology* 1992; 76: 204–8.
- 140 Warner ME, Warner ME, Garrity JA, Mackenzie RA, Warner DO. The frequency of perioperative vision loss. *Anesthesia and Analgesia* 2001; **93**: 1417–21.
- 141 Sweeney PJ, Breuer AC, Selhorst JB *et al.* Ischemic optic neuropathy: a complication of cardiopulmonary bypass surgery. *Neurology* 1982; **32**: 560–2.

- 142 Shaw PJ, Bates D, Cartlidge NEF *et al.* Neurologic and neuropsychological morbidity following major surgery: comparison of coronary artery bypass and peripheral vascular surgery. *Stroke* 1987; **18**: 700–7.
- 143 Evan KE, Tavill MA, Goldberg AN, Silverstein H. Sudden sensorineural hearing loss after general anaesthesia for nonotologic surgery. *Laryngoscope* 1997; 107: 747–52.
- 144 Gultekin S, Ozcan S. Does hearing loss after spinal anaesthesia differ between young and elderly patients? *Anesthesia and Analgesia* 2002; 94: 1318–20.
- 145 Lamberg T, Pitkanen MT, Marttila T, Rosenberg PH. Hearing loss after continuous or single shot spinal anaesthesia. *Regional Anaesthesia* 1997; 22: 539–42.
- 146 Chung F, Mezei G. Adverse outcomes in ambulatory anesthesia. Canadian Journal of Anesthesia 1999; 46: R18–R26.
- 147 Ogg TW. An assessment of postoperative outpatient cases. *British Medical Journal* 1972; **4**: 573–5.
- 148 Tong D, Chung F, Mezei G. Which specific postoperative symptoms predict postoperative functional level in ambulatory patients? *Anesthesiology* 1997; 87: A37.
- 149 Swan BA, Maislin G, Traber KB. Symptom distress and functional status changes during the first seven days after ambulatory surgery. *Anesthesia and Analgesia* 1998; 86: 739–45.
- 150 Wu CL, Berenholtz SM, Pronovost PJ, Fleisher LA. Systematic review and analysis of postdischarge symptoms after outpatient surgery. *Anesthesiology* 2002; 96: 994–1003.
- 151 Commission on the Provision of Surgical Services. *Pain After Surgery*. London: Royal Colleges of Surgeons and Anaesthetists, 1990.
- 152 American Society of Anesthesiologists. Practice Guidelines for the Management of Acute Pain in the Perioperative Setting. Park Ridge, Illinois, USA: ASA, 1995.
- 153 Ready B, Oden R, Chadwick H et al. Development of an anesthesiology-based postoperative pain management service. Anesthesiology 1988; 68: 100–6.
- 154 National Health, Medical Research Council. Acute Pain Management: Scientific Evidence. Canberra: Australian Government Publishing Service, 1999.
- 155 Audit Commission. Anaesthesia Under Examination. London: Audit Commission, 1997.
- 156 Dolin SJ, Cashman JN, Bland JM. Effectiveness of acute postoperative pain management: 1. evidence from published data. *British Journal of Anaesthesia* 2002; 89: 409–23.
- 157 Chung F, Ritchie E, Su J. Postoperative pain in ambulatory surgery. Anesthesia and Analgesia 1997; 85: 808–16.
- 158 Chung F, Un V, Su J. Postoperative symptoms 24 hours after ambulatory anaesthesia. *Canadian Journal of Anesthesia* 1996; 43: 1121–7.
- 159 Scott NB, Hodson M. Public perceptions of postoperative pain and its relief. *Anaesthesia* 1997; **52**: 438–42.
- 160 Watcha MF, White PF. Postoperative nausea and vomiting. Its etiology, treatment and prevention. *Anesthesiology* 1992; 77: 162–84.
- 161 Fortney JT, Gan TJ, Gracyk S *et al.* A comparison of the efficacy, safety, and patient satisfaction of ondansetron versus

- droperidol as antiemetics for elective outpatient surgical procedures. *Anesthesia and Analgesia* 1998; **86**: 731–8.
- 162 Sinclair D, Chung F, Mezei G. Can postoperative nausea and vomiting be predicted? *Anesthesiology* 1999; 91: 109–18.
- 163 Sinclair D, Chung F, Mezei G. Relation of postoperative nausea and vomiting to the surgical procedure. *Canadian Journal of Anesthesia* 1998; 45: A25A.
- 164 Scuderi PE, James RL, Harris L, Mims GR. Antiemetic prophylaxis does not improve outcomes after outpatient surgery when compared to symptomatic treatment. *Anesthesiology* 1999; 90: 360–71.
- 165 Chia YY, Kuo MC, Liu K, Sun GC, Hseih SW, Chow LH. Does postoperative pain induce emesis? *Clinical Journal* of Pain 2002; 18: 317–23.
- 166 Apfel CC, Greim CA, Haubitz I et al. The discriminating power of a risk score for postoperative vomiting in adults undergoing various types of surgery. Acta Anaesthesiologica Scandinavica 1998; 42: 502–9.
- 167 Tramer MR, Reynolds JM, Moore RA, McQuay HJ. Efficacy, dose–response, and safety of ondansetron in prevention of postoperative nausea and vomiting. A quantitative systematic review of randomized placebo-controlled trials. *Anesthesiology* 1997; 87: 1277–89.
- 168 Tramer MR, Phillips C, Reynolds DJ, McQuay HJ, Moore RA. Cost-effectiveness of ondansetron for postoperative nausea and vomiting. *Anaesthesia* 1999; 54: 226–34.
- 169 White PF, Watcha MF. Postoperative nausea and vomiting: prophylaxis versus treatment. Anesthesia and Analgesia 1999; 89: 1337–9.
- 170 Fisher DM. Surrogate outcomes: meaningful not! Anesthesiology 1999; 90: 355–6.
- 171 Higgins PP, Chung F, Mezei G. Postoperative sore throat after ambulatory surgery. *British Journal of Anaesthesia* 2002; 88: 582–4.
- 172 Joshi GP, Inagaki Y, White PF. Use of the laryngeal mask as an alternative to the tracheal tube during ambulatory surgery. *Anesthesia and Analgesia* 1997; **85**: 573–7.
- 173 Kloub R. Sore throat following tracheal intubation. *Middle East Journal of Anesthesiology* 2001; **16**: 29–40.
- 174 Splinter WM, Smallman B, Rhine EJ, Komocar L. Postoperative sore throat in children and the laryngeal mask airway. *Canadian Journal of Anesthesia* 1994; 41: 1081–3.
- 175 Dingley J, Whitehead MJ, Wareham J. A comparative study of the incidence of sore throat with the laryngeal mask airway. *Anaesthesia* 1994; 49: 251–4.
- 176 Oczenski W, Krenn H, Dahaba AA *et al.* Complications following the use of the Combitube, tracheal tube and laryngeal mask airway. *Anaesthesia* 1999; **54**: 1161–5.
- 177 Christensen AM, Willemoes-Larsen H, Lundby L, Jakobsen KB. Postoperative throat complaints after tracheal intubation. *British Journal of Anaesthesia* 1994; 73: 786–7.
- 178 Nikolajsen L, Larsen KM, Kierkegaard O. Effect of previous frequency of headache, duration of fasting and caffeine abstinence on perioperative headache. *British Journal of Anaesthesia* 1994; 72: 295–7.

- 179 Holte K, Kehlet H. Compensatory fluid administration for preoperative dehydration – does it improve outcome? Acta Anaesthesiologica Scandinavica 2002; 46: 1089–93.
- 180 Warner ME, Benenfeld SM, Warner MA, Schroeder DR, Maxson PM. Perianesthetic dental injuries: frequency, outcomes, and risk factors. *Anesthesiology* 1999; 90: 1302–5.
- 181 Burton JF, Baker AB. Dental damage during anaesthesia and surgery. *Anaesthesia and Intensive Care* 1987; **15**: 262–8.
- 182 Fung BK, Chan MY. Incidence of oral tissue trauma after the administration of general anaesthesia. *Acta Anaesthesiologica Sinica* 2001; **39**: 163–7.
- 183 Sawyer RJ, Richmond MN, Hickey JD, Jarratt JA. Peripheral nerve injuries associated with anaesthesia. *Anaes*thesia 2000; 55: 980–91.
- 184 Dhuner KG. Nerve injuries following operations: survey of cases during a 6 year period. Anesthesiology 1950; 11: 289–93.
- 185 Alvine FG, Schurrer ME. Postoperative ulnar nerve palsy. Are there predisposing factors? *Journal of Bone and Joint Surgery* 1987; 69: 255–9.
- 186 Kroll DA, Caplan RA, Posner K, Ward RJ, Cheney FW. Nerve injury associated with anaesthesia. *Anesthesiology* 1990; 73: 202–7.
- 187 Warner MA, Warner DO, Matsumoto JY, Harper CM, Schroeder DR, Maxson PM. Ulnar neuropathy in surgical patients. Anesthesiology 1999; 90: 54–9.
- 188 Faccenda KA, Finucane BT. Complications of regional anaesthesia. Incidence and prevention. *Drug Safety* 2001; 24: 413–42.
- 189 Dripps RD, Vandam LD. Long term follow up of patients who received 10 098 spinal anaesthetics. *Journal of the American Medical Association* 1954; 16: 1486–91.
- 190 Massey Dawkins CJ. An analysis of the complications of extradural and caudal block. *Anaesthesia* 1969; 24: 554–63.
- 191 Dahlgren N, Tornebrandt K. Neurological complications after anaesthesia. A follow up of 18 000 spinal and epidural anaesthetics performed over three years. Acta Anaesthesiologica Scandinavica 1995; 39: 872–80.
- 192 Tryba M. Epidural regional anaesthesia and low molecular heparin: pro (German). Anaesthesia Intensivmed Notfallmed Schmerzther 1993; 28: 179–81.
- 193 Vandermeulen EP, Van Aken H, Vermylen J. Anticoagulants and spinal–epidural anesthesia. *Anesthesia and Analgesia* 1994; 79: 1165–77.
- 194 Tryba M, Wedel DJ. Central neuraxial block and low molecular weight heparin (enoxaparin): lessons learned from different dosage regimes in two continents. *Acta Anaesthesiologica Scandinavica* 1997; **41**: 100–3.
- 195 Holtas S, Heiling M, Lonntoft M. Spontaneous spinal epidural haematoma findings at MR imaging and clinical correlation. *Radiology* 1996; 199: 179–81.
- 196 Thomas J, Paranjothy S, Royal College of Obstetricians and Gynaecologists Clinical Effectiveness Support Unit. *The National Sentinel Caesarian Section Audit Report.*London: RCOG Press, 2001. Available at: http://www.rcog.org.uk/effectiveness/nscs_audit.html

- 197 Loo CC, Dahlgren G, Irestedt L. Neurological complications in obstetric regional anaesthesia. *International Journal of Obstetric Anaesthesia* 2000; 9: 99–124.
- 198 Holdcroft A, Gibberd FB, Hargrove RL, Dawkins DF, Dellaportas CI. Neurological complications associated with pregnancy. *British Journal of Anaesthesia* 1995; 75: 522–6.
- 199 Paech MJ, Godkin R, Webster S. Complications of obstetric epidural analgesia and anaesthesia. a prospective analysis of 10 995 cases. *International Journal of Obstetric* Anaesthesia 1998; 7: 5–11.
- 200 Scott DB, Hibbard BM. Serious non-fatal complications associated with extradural block in obstetric practice. *British Journal of Anaesthesia* 1990; 64: 537–41.
- 201 Ong BY, Cohen MM, Esmail A et al. Paresthesia and motor dysfunction after labor and delivery. Anesthesia and Analgesia 1987; 66: 18–22.
- 202 Scott DB, Tunstall ME. Serious complications associated with epidural: spinal blockade in obstetrics: a two year prospective study. *International Journal of Obstetric Anesthesia* 1995; 4: 133–9.
- 203 Phillips OC, Ebner H, Nelson AT, Black MH. Neurologic complications following spinal anesthesia with lidocaine. a prospective review of 10 440 cases. *Anesthesiology* 1969; 30: 284–9.
- 204 Crawford JS. Some maternal complications of epidural analgesia for labour. *Anaesthesia* 1985; 40: 1219–25.
- 205 Tarkkila P, Huhtala J, Tuominen M. Transient radicular irritation after spinal anaesthesia with hyperbaric 5% lignocaine. *British Journal of Anaesthesia* 1995; 74: 328–9.
- 206 Freedman K, Li D-K, Drasner K et al. and the Spinal Anaesthesia Study Group. Transient neurological symptoms after spinal anaesthesia. An epidemiological study in 1863 patients. *Anesthesiology* 1998; 89: 633–41.
- 207 Pollock JE, Neal JM, Stephenson CA, Wiley CE. Prospective study of the incidence of transient radicular irritation in patients undergoing spinal anesthesia. *Anesthesiology* 1996; 84: 1361–7.
- 208 Wang LP, Hauerberg J, Schmidt JF. Incidence of spinal epidural abscess after epidural analgesia. *Anesthesiology* 1999; 91: 1928–36.
- 209 Kindler C, Seeberger M, Schneider M. Extradural abscess complicating lumbar extradural anaesthesia and analgesia in an obstetric patient. *Acta Anaesthesiologica Scandinavica* 1996; 40: 858–61.
- 210 Baker AS, Ojemann RG, Swantz MN et al. Spinal epidural abscess. New England Journal of Medicine 1975; 293: 463–8.
- 211 Toyama TM, Ranasinghe JS, Siddiqui MN, Steadman JL, Lai M. Incidence of post dural puncture headache and epidural blood patch following dural puncture with epidural needle in 15,411 obstetric patients in a large tertiary care teaching hospital. *Anesthesiology* 2002; 96: 100.
- 212 Halpern S, Preston R. Postdural puncture headache and spinal needle design. Metaanalyses. *Anesthesiology* 1994; 81: 1376–83.
- 213 Hopkinson JM, Samaan AK, Russell IF, Birks RJS, Patrick MR. A comparative multicentre trial of spinal needles for Caesarean section. *Anaesthesia* 1997; 52: 998–1014.

- 214 Stride PC, Cooper GM. Dural taps revisited: a 20-year survey from Birmingham Maternity Hospital. *Anaesthesia* 1993: 48: 247–55.
- 215 Taivanen T, Pitkanen M, Tuominen M, Rosenberg PH. Efficacy of epidural blood patch for post-dural puncture headache. Acta Anaesthesiologica Scandinavica 1993; 37: 702–5.
- 216 Gleeson CM, Reynolds F. Accidental dural puncture rates in UK obstetric practice. *International Journal of Obstetric* Anesthesia 1998; 7: 242–6.
- 217 Macarthur C, Lewis M, Knox EG. Accidental dural puncture in obstetric patients and long term symptoms. *British Medical Journal* 1993; 306: 883–5.
- 218 Jeskins GD, Moore PAS, Cooper GM, Lewis M. Long-term morbidity following dural puncture in an obstetric population. *International Journal of Obstetric Anesthesia* 2001; 10: 17–24.
- 219 Patel M, Fernando R, Gill P et al. A prospective study of long term backache after child birth in primigravidae – the effect of ambulatory epidural analgesia during labour. International Journal of Obstetric Anesthesia 1995; 4: 187.
- 220 Russell R, Dundas R, Reynolds F. Long term backache after childbirth: prospective search for causative factors. *British Medical Journal* 1996; 312: 1384–8.
- 221 Breen TW, Ransil BJ, Groves PA et al. Factors associated with back pain after childbirth. Anesthesiology 1994; 81: 29–34.
- 222 Brown EM, Elman DS. Postoperative backache. *Anesthesia* and *Analgesia* 1961; **40**: 683–5.
- 223 Asantila R, Rosenber PH, Scheinn B. Comparison of different methods of postoperative analgesia after thoracotomy. Acta Anaesthesiologica Scandinavica 1986; 30: 421–5.
- 224 Breivik H, Hogstrom H, Niemi G et al. Safe and effective postoperative pain relief: introduction and continuous quality improvement of comprehensive post-operative pain management programmes. Bailliere's Clinical Anaesthesiology 1995; 9: 423–60.
- 225 Liang C-C, Wong S-Y, Tsay PT et al. The effect of epidural analgesia on postpartum urinary retention in women who deliver vaginally. *International Journal of Obstetric Anesthesia* 2002; 11: 164–9.
- 226 Brown DL, Ransom DM, Hall JA et al. Regional anaesthesia and local anaesthesia induced systemic toxicity: seizure frequency and accompanying cardiovascular changes. Anesthesia and Analgesia 1995; 81: 321–8.
- 227 Selander D, Edshage S, Wolf T. Paresthesiae or no paresthesiae? Acta Anaesthesiologica Scandinavica 1979; 23: 27–33.
- 228 Brand L, Papper EM. A comparison of supraclavicular and axillary techniques for brachial plexus blocks. *Anesthesiology* 1961; 22: 226–9.
- 229 Hickey R, Garland TA, Ramamurthy S. Subclavian perivascular block: influence of location of parasthesiae. *Anesthesia and Analgesia* 1989; 68: 767–71.
- 230 Hamilton RC. Retrobulbar haemorrhage after retrobulbar blocks. *Anesthesia and Analgesia* 1994; **78**: 608.

- 231 Edge KR, Nicoll JMV. Retrobulbar haemorrhage after 12 500 retrobulbar blocks. Anesthesia and Analgesia 1993; 76: 1019–22
- 232 Hamilton RC. Brainstem anaesthesia as a complication of regional anaesthesia for ophthalmic surgery. *Canadian Journal of Ophthalmology* 1992; 27: 323–5.
- 233 Hamilton RC, Gimbel HV, Strunin L. Regional anaesthesia for 12 000 cataract extraction and intraocular lens implantation procedures. *Canadian Journal of Anesthesia* 1988; 35: 615–23.
- 234 Gillart T, Bazin JE, Montetagaud M, Bevillard F, Amara S, Schoeffler P. The effects of volume and speed of injection in peribulbar anaesthesia. *Anaesthesia* 1998; 53: 486–91
- 235 McLure HA, Rubin AP, Westcott M, Henderson H. A comparison of 1% ropivacaine with a mixture of 0.75% bupivacaine and 2% lignocaine for peribulbar anaesthesia. *Anaesthesia* 1999; 54: 1178–82.
- 236 Sarvela PJ, Paloheimo MPJ, Nikki PH. Comparison of pH-adjusted bupivacaine 0.75% and a mixture of bupivacaine 0.75% and lidocaine 2%, both with hyaluronidase, in day case cataract surgery under regional anesthesia. *Anesthesia and Analyesia* 1994; 79: 35–9.
- 237 Alfidi R. Informed consent: a study of patient reaction. Journal of the American Medical Association 1971; 216: 1325–9.
- 238 Inglis S, Farnill D. The effects of providing preoperative statistical anaesthetic risk information. *Anaesthesia and Intensive Care* 1993; 21: 799–805.
- 239 Calman KC. Concern. Science and society and the communication of risk. *British Medical Journal* 1996; 313: 799–802.
- 240 Adams A. The meaning of risk. In: McConachie I, ed. Anaesthesia for the High Risk Patient. London: Greenwich Medical Media, 2002: 239–47.
- 241 Malenka DJ, Baron JA, Johansen S et al. The framing effect of relative and absolute risk. Journal of General International Medicine 1993; 8: 543–8.
- 242 Bristol Royal Infirmary Inquiry, July 2001. Learning from Bristol: the report of the public inquiry into children's heart surgery at the Bristol Royal Infirmary 1984–1995. Available at: http://www.bristolinquiry.org.uk/final report.
- 243 Edwards A, Elwyn G, Mulley A. Explaining risks: turning numerical data into meaningful pictures. *British Medical Journal* 2002; 324: 827–30.
- 244 Calman KC, Royston HD. Risk language and dialects. British Medical Journal 1997; **315**: 939–42.
- 245 BMA. Guide to Living with Risks. Harmondsworth, UK: Penguin, 1990.
- 246 Barclay P, Costigan S, Davies M. Lottery can be used to show risk (letter). British Medical Journal 1998; 316: 124.
- 247 Adams AM, Smith AF. Risk perception and communication: recent developments and implications for anaesthesia. Anaesthesia 2001; 56: 745–55.