time of the operation but also because inpatient investigations and treatment are necessary.

Revision operations take longer to perform than the equivalent primary procedure and are certainly more expensive in terms of the consumable items used. Twenty five out of 36 of the exchange arthroplasties used cement impregnated with antibiotic, and 27 required non-standard implants.

The cost of antibiotics was over 10 times that incurred with primary surgery because combined, often parenteral, treatment with expensive drugs was given for long periods.

To reduce the costs to the patient and the NHS all efforts must ensure that a primary joint replacement is

as successful as possible. We caution against short term economies, which may well produce considerable clinical and financial burdens in the long term.

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Retention of skills by advanced trained ambulance staff: implications for monitoring and retraining

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In the United Kingdom there are no formal recommendations for monitoring ambulance staff who have training in advanced cardiac life support skills. Although the NHS Training Authority's guidelines for extended training of ambulance staff describe the content of courses in detail, recommendations for monitoring standards are limited.¹ To assess the need for monitoring we evaluated the performance of ambulance staff who had been trained in advanced cardiac life support skills but had not been formally monitored or retrained over two years.

Subjects, methods, and results

In 1985, 17 ambulance staff in London were trained in advanced cardiac life support skills. The table shows the training programme and protocols of treatment. After the training they submitted reports and electrocardiograms/audiocassette tapes (dual channel cassette tapes providing an electrocardiogram and audible record of the incident) on all patients requiring advanced skills. In 1987 all reports and tapes available for these patients were assessed for recognition of rhythms; correct treatment of arrhythmias according to the protocols; compliance with the sequence of treatment stated in the protocols; quality of the electrocardiogram; time to direct current shock (for patients found in ventricular fibrillation); and continuity of basic life support.

From 99 cases of cardiac arrest 57 tapes were available for assessment. In 100 out of 111 episodes the rhythm was accurately recognised. Drugs were given correctly in 49 out of 57 cases. Defibrillation was attempted on 54 occasions: 34 were appropriate and nine inappropriate, and in 11 cases assessing whether

the shock was appropriate was impossible because of inadequate or missing electrocardiograms. In nine patients some of the treatment was given on the basis of inadequate electrocardiograms. On two occasions the electrocardiograph's "gain" seemed to have been turned down. On four occasions "quick look" paddles were used but the resulting trace was inadequate, and on three occasions the trace was distorted by artefact. In 17 cases the patient was intubated or moved to the ambulance before electrocardiograph leads were applied. The mean time from arrival of the ambulance crew to delivery of the first shock was 4.3 (SD 2.7) minutes (range 1-9 minutes; 27 cases assessed). The mean time from the first appearance of ventricular fibrillation on the oscilloscope to delivery of the first shock was 66 (46) seconds (range 22-221 seconds). Interruptions to basic life support of longer than 30 seconds occurred in 23 cases.

Comment

The ambulance staff showed a good knowledge of the protocols of treatment and the electrocardiographic rhythms encountered. Errors in applying skills were more common and resulted in delays in defibrillation (often secondary to the patient being intubated or moved before electrocardiograph leads were applied), interruptions to basic life support, and the risk of inappropriate treatment as a result of an inadequate electrocardiogram. Improvement in these would have optimised the lifesaving capabilities of the staff, but as the staff did not receive any feedback they were probably unaware of how they might improve their performance.

The results suggest that the initial training programme was sufficient in theoretical content but required more emphasis on practical aspects. The protocol was not clear about what equipment and procedures should be used first when attending a possible cardiac arrest. Monitoring of staff is therefore necessary to assess the adequacy of the initial training programme, to evaluate how training translates into practice, and to identify aspects in which retraining is required.

Review of electrocardiograms/audiocassette tapes in

Details of advanced training programme (full programme lasts six weeks) and protocols of treatment for ambulance staff

Staff eligible for programme	Content of course	Assessment	Training given by	Practical skills acquired
Selected staff	Two weeks' theoretical work	Written examination	Hospital consultants and ambulance instructors	Cardiac monitoring
Staff trained in intubation and infusion	One week's practical work in classroom	Oral examination		Use of defibrillator
Staff able to pass written test before course	Three weeks' practical work in hospital	Practical assessment with manikin		Administration of selected drugs by endotracheal tube and intravenous routes according to set protocols*

*Ventricular fibrillation: precordial thump → shock 200 J → shock 200 J → shock 400 J → intubation → lignocaine 200 mg → shock 400 J → adrenaline (20 ml) 1/10 000 → shock 400 J. Asystole: intubation → atropine 2 mg → adrenaline (20 ml) 1/10 000 → isoprenaline 100 µg.

addition to written reports is required to gain a comprehensive view of performance. In the United States this type of monitoring by a physician is recommended for all cases of cardiac arrest treated by emergency medical technicians using advanced skills.² No such recommendations exist in the United Kingdom. Monitoring of electrocardiograms and cassette tapes is time consuming and requires additional equipment.³ Lack of resources or medical support, in addition to the lack of official directives, may result in ambulance services failing to monitor standards, and the full potential of advanced trained ambulance staff will not be achieved.

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Screening for colonic cancer in patients with Barrett's oesophagus

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In 1985 Sontag *et al* reported an association between Barrett's oesophagus and colonic neoplasia, but their study did not include a control population.¹ In another uncontrolled series no colorectal cancers were seen in 36 patients with Barrett's oesophagus, although 33% of these patients had colorectal adenomas.² We performed colonoscopy prospectively in our patients with Barrett's oesophagus, none of whom had symptoms of colonic neoplasia, and compared the results with those obtained in a control group of patients thought to have the irritable bowel syndrome.³

Subjects, methods, and results

Barrett's oesophagus was defined as columnar epithelium occurring in a tubular oesophagus at least 3 cm above the manometrically defined lower oesophageal sphincter.⁴ We studied 32 patients (21 men and 11 women), whose mean age was 61 (range 29-81). None had a history of colorectal neoplasia. Two control patients thought on clinical grounds to have the irritable bowel syndrome, and matched for age and sex, were chosen for each patient with Barrett's oesophagus; their mean age was 55 (range 28-81). No additional screening tests, such as testing for faecal occult blood, were done to exclude colonic neoplasia.

Eight patients with Barrett's oesophagus (25%) had 12 colorectal tumours, three of which were malignant and nine benign (table). All were successfully treated, two by local resection, one by subtotal colectomy, and

five by endoscopic polypectomy. Three of the 64 controls (5%) had single adenomatous polyps, which were removed endoscopically. The relative risk of colonic tumour was 5.3 in the patients with Barrett's oesophagus (95% confidence interval 1.94 to 8.69).

Comment

We found a high prevalence of colorectal tumours in our patients with Barrett's oesophagus, in keeping with the findings of Sontag *et al*.¹ Our study was entirely prospective, and the association between colonic neoplasia and Barrett's oesophagus seemed to be genuine and not due to some hidden bias; this may not have been the case in the uncontrolled study by Sontag *et al*, in which seven of 10 carcinomas were identified retrospectively.

Selecting an appropriate control group was difficult, and we considered that colonoscopic examination of asymptomatic people was unjustified because of the risks and discomfort entailed. We thus selected patients who had been referred for investigation and in whom the irritable bowel syndrome had been diagnosed on clinical grounds. The controls were on average six years younger than the study group, and this could have affected the prevalence of colonic tumours: the prevalence of colonic neoplasia related to age in asymptomatic subjects is unknown. The difference in age between the two groups, however, is unlikely to have accounted for all of the observed differences in prevalence.

The aetiology of both Barrett's oesophagus and carcinoma of the colon remains unknown. The symptoms of dyspepsia or chest pain are unlikely to have been caused by the colonic carcinoma or adenoma, and in each case the finding of a colonic lesion was incidental. An inherited or acquired common aetiological agent, such as a dietary factor or a tendency to neoplastic or metaplastic change throughout the gastrointestinal tract, may have been the cause. There was no evidence of dysplasia in biopsy specimens or resected colonic epithelium not affected by adenomatous or carcinomatous change.

It has been recommended that frequent oesophagoscopy should be undertaken in patients with Barrett's oesophagus to detect early oesophageal carcinoma or dysplasia. None of our patients had high grade dysplasia or oesophageal carcinoma but eight had colonic neoplasia. Colonoscopy may therefore be a more worthwhile screening procedure for cancer than oesophagoscopy in Barrett's oesophagus.

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Clinical details of patients and controls with colorectal tumours

Case No	Age (years) and sex	Site	Histological findings	Treatment
Patients				
1	79 M	{ Sigmoid colon	Tubulovillous adenoma with severe	Endoscopic polypectomy
2	66 M	{ Descending colon	dysplasia at both sites	
		{ Descending colon	Tubulovillous adenoma with severe	Endoscopic polypectomy
3	53 M	Sigmoid colon	dysplasia	
4	72 F	Sigmoid colon	Tubular adenoma and mild	Endoscopic polypectomy
5	74 M	{ Hepatic flexure	dysplasia at two sites	
		{ Sigmoid colon	Adenomatous polyp	Endoscopic polypectomy
		{ Transverse colon	Adenomatous polyp	Endoscopic polypectomy
6	52 M	{ Sigmoid colon	Tubular adenoma with mild	Right hemicolectomy
		{ Hepatic flexure	dysplasia	
7	44M	{ Hepatic flexure	Dukes's A carcinoma	Subtotal colectomy
		{ Splenic flexure	Dukes's B carcinoma	
8	71M	{ Sigmoid colon	Tubular adenoma	Sigmoid colectomy
		{ Sigmoid colon	Dukes's A carcinoma	
Controls				
9	52 M	Sigmoid colon	Adenomatous polyp	Endoscopic polypectomy
10	64 F	Descending colon	Adenomatous polyp	Endoscopic polypectomy
11	66 M	Rectum	Adenomatous polyp	Endoscopic polypectomy