Commentaries

Life, death, and varices

In patients with cirrhosis and portal hypertension, gastrointestinal haemorrhage is a major complication and cause of death. Oesophageal varices are present in approximately 50% of patients with cirrhosis; the prevalence is higher in Child-Pugh C patients than in those with Child-Pugh A-B. Among patients with varices, the risk of gastrointestinal haemorrhage ranges from 30% to 50% and half of these patients die within six weeks after bleeding. Thus 10–15% of unselected patients with cirrhosis die from gastrointestinal bleeding. The number of deaths has decreased in the last few decades as a result of modern techniques (see below). The exact mechanisms of variceal rupture have not yet been determined and thus no ideal treatment has been found. However, haemorrhage has been shown to cause different complications such as sepsis or renal failure which may be responsible for death. In patients admitted for variceal haemorrhage, certain prognostic factors for death have been determined such as age, Child-Pugh C, hepatocellular carcinoma, early rebleeding, hepatic encephalopathy, and renal failure.1

Ideally, patients admitted for acute variceal haemorrhage must be taken to a medical centre with an emergency facility and an intensive care unit with experience in variceal bleeders. In the last 20 years, various medical, surgical, radiological, pharmacological, and endoscopic treatments have been used and most are effective in controlling variceal haemorrhage compared with no treatment or placebo. Most of these treatments, except endoscopic therapy, act by decreasing portal pressure. There is a negative relationship between treatment efficacy and invasiveness of the method. For example, surgical shunts or transjugular intrahepatic portosystemic stent shunts are more effective than pharmacological and endoscopic treatments but are more invasive and thus associated with more side effects. Although all of these treatments stop variceal haemorrhage, a significant difference in mortality has only been found with endoscopic sclerotherapy or terlipressin administration compared with placebo or no treatment. The discrepancy between the efficacy of treatment on variceal bleeding and survival rate suggests that other factors play a role in survival. Emergency endoscopic sclerotherapy is now the gold standard in the management of acute variceal haemorrhage as it increases hospital survival.1 A recent meta-analysis showed however that endoscopic sclerotherapy and variceal ligation are equally effective, suggesting that variceal ligation may also improve survival rate with fewer complications.1 Two controlled studies have shown that terlipressin administration is effective both in controlling bleeding and improving survival rate. In one trial, terlipressin stopped bleeding in 90% of patients with a significant difference in survival rate before discharge.1 In a second study, terlipressin was administered with gyceryl trinitrate by an intensive care team, within one hour after an emergency call before admission.1 In this trial, mortality due to bleeding episodes was significantly lower in the terlipressin group than in the placebo group at day 15 and day 42. These findings suggest that this drug should be administered as soon as possible before endoscopic investigation. Moreover, it is interesting to note that these two treatments (sclerotherapy and terlipressin administration) are equally effective in the initial control of variceal bleeding and in the prevention of early rebleeding.5

The second type of therapy is for the prevention of the complications of haemorrhage. Patients with gastrointestinal bleeding are at a high risk of bacterial infection and thus antibiotics must be prescribed. A meta-analysis confirmed that the use of prophylactic antibiotics prevents bacterial infection but also the risk of early rebleeding and more important, significantly improves survival rate.3 Prevention of liver failure, including encephalopathy and renal failure, may also play a major role in the improvement in survival rate. Administration of blood products must be limited as it has been suggested that excess blood products may induce early rebleeding. Similarly, limiting certain procedures that are more or less invasive investigations may reduce severe complications. Accordingly, improvements in survival rate may depend in part on the type of treatment. This is relatively clear for terlipressin administration1 but no explanation exists for endoscopic treatment.

Finally, mortality due to acute variceal haemorrhage has significantly decreased in the last few decades. Preliminary results of one study showed that inhospital mortality has decreased by 50% over the past 15 years.7 Moreover, the authors showed that although there were no deaths in Child-Pugh A patients following gastrointestinal bleeding, 25% of Child-Pugh C patients still died from haemorrhage. A second recent study followed 5000 patients in two cohorts separated by an interval of 11 years, who were followed for more than six years.10 The authors demonstrated that mortality in the late cohort significantly declined at 30 days (approximately 30%) and at six years (approximately 6%). These results were despite the fact that patients in the late cohort had more severe liver disease. Thus improvement in survival rate is even more significant with multivariate analysis. This study also showed that patients treated with sclerotherapy during the first hospitalisation had a lower mortality rate than the rest of the late cohort. In this issue of Gut, McCormick and O’Keefe have calculated the survival rate in patients with cirrhosis admitted for a first episode of variceal haemorrhage (see page 682).11 They compared the survival rate from “control” groups in approximately 1500 patients from 1960 to 2000. The results show a significant reduction (approximately 40%) in bleeding related mortality.

Although this improved survival rate after variceal haemorrhage might be expected, variceal haemorrhage was controlled for more than 20 years without any significant effects on survival rate. The only treatments known to improve survival rate are pharmacological or endoscopic treatments and antibiotic administration (see above). Nevertheless, a certain number of patients still die of variceal haemorrhage, suggesting that other factors must be studied. Thus new ideas, hypotheses, and approaches are needed to further our understanding of the treatment and mechanisms of variceal haemorrhage.
Can endoscopic papillary balloon dilation really preserve sphincter of Oddi function?

The technique of endoscopic balloon dilation of the major duodenal papilla with extraction of bile duct stones was first reported in 1982 but an unacceptably high rate of acute pancreatitis following the procedure impeded its acceptance until further reports in 1994 and 1995 reported its safety. A number of randomised controlled trials have demonstrated its effectiveness in clearing the bile ducts of stones compared with endoscopic sphincterotomy, with variable short term complication rates similar to sphincterotomy. Currently, this technique has not become popular worldwide, but for reasons that are not clear has been adopted much more readily in Japan and Korea than elsewhere.

Those who advocate this technique over sphincterotomy do so on the premise that: (1) it is less traumatic, (2) it is simpler to perform, (3) it is as effective, (4) it avoids the immediate complications, such as bleeding and perforation, (5) it is cheaper or at least equivalent cost, (6) it preserves sphincter of Oddi function, and (7) it avoids the long term complications.

Many of these apparent advantages have been questioned in terms of technical performance and short term outcome and now there is new evidence from the report by Yasuda et al in this issue of Gut that the long term advantages may also be in doubt (see page 686).

The above selection criteria in favour of balloon dilation of the papilla have been criticised and are discussed below under the same categories.

(1) Forcible balloon dilation of the papilla well beyond its normal resting diameter may be just as traumatic or more traumatic than its incision by sphincterotomy and indeed, the description of the degree of trauma is "in the eye of the beholder". (2) The technique is not necessarily simpler to perform than sphincterotomy, especially as current over-the-guidewire hydrostatic balloons are not designed for this application and may be difficult to keep in position during the dilation process. (3) The overall effectiveness of clearing the bile ducts is statistically equivalent to sphincterotomy although this is at the expense of a far greater need for mechanical lithotripsy and a small proportion of patients requiring additional procedures to achieve clearance. Recent use of intraductal ultrasound has cast doubt on the completeness of ductal clearance after balloon dilation with a retained stone rate of up to 33%. (4)
stones (4–24 mm), and diameter of the bile duct (6–30 mm) but the use of mechanical lithotripsy was significantly higher in the balloon dilation group (17% compared with 9% in sphincterotomy patients).

With this new additional long term evidence it seems likely that endoscopic balloon dilation of the papilla for the treatment of choledocholithiasis would be considered as an alternative to sphincterotomy in certain circumstances but not as a replacement for it on a routine basis. Many centres have already adopted the practice of using balloon dilation in patients with previous surgically altered anatomy where access to the papilla is from the distal side and also in patients where there may be an uncorrectable coagulopathy which might carry increased risks of post-sphincterotomy bleeding. The converse of this discussion is the question of which might carry increased risks of post-sphincterotomy bleeding. The possible unsubstantiated differences in patient populations with choledocholithiasis in different countries may account for some of the differences in early morbidity with balloon dilation between the Far East and the West. I would currently advocate caution in the use of endoscopic dilation of the papilla for routine treatment of choledocholithiasis, except for patients in special circumstances, as detailed above, and then only for those with stone diameters up to the size of the balloon being employed to avoid excessive use of mechanical lithotripsy which in itself may have some influence on the complication rate. Unquestionably, further long term follow up studies are required to substantiate the findings by Yasuda and colleagues but for the present time, sphincterotomy, a tried and tested technique now well into its third decade, would still seem to be the standard endoscopic modality to be offered to most patients with choledocholithiasis.

D L CARR-LOCKE
Division of Gastroenterology, Brigham and Women’s Hospital, Boston, Massachusetts, USA
dcarrlocke@partners.org

6 DiSario JA, Freeman ML, Bjorkman DJ, et al. Endoscopic balloon dilation compared to sphincterotomy (EDES) for extraction of bile duct stones. Gas troenter Endosc 1997;45:AB129.