Small for size liver remnant following resection: prevention and management

Rony Eshkenazi¹, Yael Dreznik², Eylon Lahat², Barak Bar Zakai¹, Alex Zendel³, Arie Ariche¹

¹Department of HPB Surgery, ²Department of Surgery B, ³Department of Surgery C, Chaim Sheba Medical Center, Tel-Hashomer, Israel

Correspondence to: Dr. Arie Ariche. Department of HPB Surgery, Chaim Sheba Medical Center, Tel Hashomer, Ramat Gan 52621, Israel.
Email: Arie.Ariche@sheba.health.gov.il.

Abstract: In the latest decades an important change was registered in liver surgery, however the management of liver cirrhosis or small size hepatic remnant still remains a challenge. Currently post-hepatectomy liver failure (PLF) is the major cause of death after liver resection often associated with sepsis and ischemia-reperfusion injury (IRI). “Small-for-size” syndrome (SFSS) and PFL have similar mechanism presenting reduction of liver mass and portal hyper flow beyond a certain threshold. Few methods are described to prevent both syndromes, in the preoperative, perioperative and postoperative stages. Additionally to portal vein embolization (PVE), radiological examinations (mainly CT and/or MRI), and more recently 3D computed tomography are fundamental to quantify the liver volume (LV) at a preoperative stage. During surgery, in order to limit parenchymal damage and optimize regenerative capacity, some hepatoprotective measures may be employed, among them: intermittent portal clamping and hypothermic liver preservation. Regarding the treatment, since PLF is a quite complex disease, it is required a multi-disciplinary approach, where it management must be undertaken in conjunction with critical care, hepatology, microbiology and radiology services. The size of the liver cannot be considered the main variable in the development of liver dysfunction after extended hepatectomies. Additional characteristics should be taken into account, such as: the future liver remnant; the portal blood flow and pressure and the exploration of the potential effects of regeneration preconditioning are all promising strategies that could help to expand the indications and increase the safety of liver surgery.

Keywords: Liver surgery; small for size liver remnant; post-hepatectomy liver failure (PLF); liver resection

Submitted Sep 02, 2014. Accepted for publication Sep 09, 2014.
doi: 10.3978/j.issn.2304-3881.2014.09.08
View this article at: http://dx.doi.org/10.3978/j.issn.2304-3881.2014.09.08

Introduction

In the latest decades an important change was registered in liver surgery, related to the progress of surgical techniques, anesthesiology and postoperative treatment, allowing a sharp decrease in mortality and morbidity. However, management of liver cirrhosis or small size hepatic remnant still remains a challenge (1).

The liver presents regenerative capacity, allowing performance of repeated resections. In certain cases, when this capacity is impaired, or where extensive resections were performed with small remnant liver, these patients may develop small for size syndrome (SFSS) with the presence of reduced liver mass insufficient to maintain normal liver function.

The term SFSS was first employed in liver transplantation to describe the development of acute liver failure (ALF) (hyperbilirubinemia, coagulopathy, encephalopathy and refractory ascites) resulted from the transplantation of a donor liver that was too small for a given recipient (2). A similar syndrome, called “post-hepatectomy liver failure (PLF)” was also described in hepatic surgery involving extended resections of liver mass. The last one is characterized by postoperative liver dysfunction, with clinical signs of prolonged cholestasis, coagulopathy, portal hypertension and ascites. PLF is the major cause of death after liver resection often associated with sepsis and ischemia-
reperfusion injury (IRI) (3).

The patho-physiological mechanisms of the SFSS and PLF are very similar, both presenting reduction of liver mass and portal hyper flow beyond a certain threshold (4).

The aim of this review is to discuss applicable peri-operative methods to prevent the SFSS or PLF and highlight the main treatment types.

**Pathophysiology**

The liver should contain minimum amount of parenchymal hepatic cells to assure its functions and the maintenance of its regeneration capacity. The hepatic parenchyma should be able to accommodate the hemodynamic changes that occur after liver resection, avoiding venous congestion. Factors such as decrease of hepatic parenchyma cells, infection and different causes that might jeopardize regeneration should be absent (5).

Decrease in parenchymal volume results in a hyper perfusion of the liver, causing dilation of sinusoids, hemorrhagic infiltration, shear stress, centro lobular necrosis, prolonged cholestasis impaired synthetic function and inhibition of cell proliferation (6).

Hepatic resections have higher risks of infection (above 50%). The number of Kupffer cells after hepatic resection decreased and thus the liver's ability to fight against infection as well. The sepsis possesses the ability to complicate or precipitate PLF. A relative increase in the production of endotoxins in the remnant liver is beneficial, once it activates the Kupffer cells, trigging the liver regeneration. This prolonged state may cause Kupffer's cellular dysfunction, resulting in difficulty of regeneration and even liver necrosis (7).

The parenchymal damage occurs following vascular occlusion or after hemorrhagic shock, causing IRI. After a period of ischemia, the complement cascade is triggered, leading to the activation of Kupffer cells, reactive oxygen appearance of species (ROS) and endothelial cell lesion. During reperfusion a release of cytokines, cell adhesion, activation and recruitment of T cell and polymorphonuclear cell occurs, resulting in microvascular lesion, inflammation and cell death (8).

**Preoperative period—prevention**

**Liver function tests and scores (9)**

The liver function tests can be divided into three types:

- Conventional tests, i.e., serum bilirubin, albumin, alkaline phosphatase, gamma glutamyl transpeptidase, prothrombin time (PT) and platelet count;
- Quantitative tests, i.e., aminopyrine breath test, antipyrine clearance, caffeine clearance, lidocaine clearance, methacetin breath test, galactose elimination capacity, low-dose galactose clearance, clearance sorbitol, indocyanine green disappearance, albumin synthesis, urea synthesis and 99mTc-GSA;
- Scores, i.e., Child-Turcotte-Pugh and MELD.

One of the best tests today to check liver function before surgery is liver retention of indocyanine green. Widely used since the decade of the 70 in Asian countries, and not yet widespread in the west.

Based on the decisional tree [established by Seyama et al. (Figure 1)] identify before the operation which hepatic volume can be resected in cirrhotic patients depending on their liver function (9).

**Liver volume (LV) manipulation and liver parenchymal protection**

The ideal volume of the hepatic remnant was exhaustively discussed in the literature and some formulas to calculate it were described (10) (Table 1).

The radiological examinations (mainly CT and/or MRI) before surgery are fundamental to quantify the LV. More recently 3D computed tomography reconstructions could define more accurately the hepatic volume allowing preoperative studies. Through this exam, the surgeon can simulate a resection, making possible the planning and the choice of the best way to do the procedure (15,18).

Measurement of volume ratios correlated with the etiology and severity of chronic liver disease (CLD) constitute a reliable predictors of patient survival (19). Although, the reliability of this ratio might be compromised by the presence of dilated bile ducts, multiple tumors, undetected lesions. Additionally, due to cholestasis or previous chemotherapy, cholangitis, vascular obstruction, steatosis or cirrhosis, or segmental atrophy and/or hypertrophy from tumor growth, negatively impacts the liver function (16).

Values calculated from graft weight-to-recipient body weight ratio (GRBWR), or standardized liver volume (SLV) based on recipient body surface area (BSA) are used to predict minimum adequate graft volume (15). But in presence of steatosis, particularly >30%, graft weight alone is not a suitable guide (10).

Extended resection of 80% of functional parenchyma can be performed in the absence of CLD for hepatobiliary
malignancies (20). Recommended minimal functional remnant LV following extended hepatectomy is 25% in a normal liver, and 40% in a “sick” liver, with moderate to severe steatosis, cholestasis, fibrosis, cirrhosis, or following chemotherapy (15).

There are some strategies that allow volume manipulation, such as portal vein embolization (PVE) and two-stage hepatectomy (16,21). PVE is usually performed percutaneously by transhepatic PVE, but may also be achieved by surgical ligation and injection of alcohol or other products to prevent the recanalization of the portal vein. PVE increases the functional capacity of the liver remnant and can increase contralateral lobe volume by up to 20 per cent, with the peak in growth occurring within 2–4 weeks (22).

Patients, in which the liver does not have a good result after PVE are selected as no good candidates for large resections due to the difficulty of regeneration (22). Patients with bilateral tumors when proceeding PVE may stimulate the growth of neoplastic cells in the non-embolized lobes, in this cases surgical treatment or ablation [radiofrequency (23,24), microwave (25) and NanoKnife® (personal experience)] of such lesions prior to the embolization are required (26). Neoadjuvant chemotherapy (27) and intra-arterial chemotherapy (28) also can be used in combination with PVE to control tumor load before resection (20,29).

Patients with bilateral liver lesions, where resection is not feasible under one procedure, the two stage hepatectomy is applied, allowing the remaining liver to be resected to achieve the suitable LV at the second stage.
**Intraoperative period—prevention**

In order to limit parenchymal damage and optimize regenerative capacity, two hepatoprotective measures may be employed: intermittent portal clamping and hypothermic liver preservation.

Intermittent portal clamping with intervals allowed for reperfusion is preferred to continuous clamping, usually applying a 15-min clamp-5-min release regimen (30-32).

Total vascular exclusion of the liver should be used when we have no choice to do the resection without it. When chosen, we can utilized hepatic vascular exclusion with preservation of the caval flow (33).

Hypothermic liver preservation in conjunction with total vascular exclusion attenuates IRI. The future remnant is infused with a preservative fluid and surrounded by crushed ice to maintain the liver at 4°C. This approach is a useful adjunct to complex resections when total vascular exclusion and vascular reconstructions are programmed (34). During surgery it is still possible to apply techniques to prevent the SFSS, if other procedures were not considered on the pre-operative period.

**Association liver partition and portal vein ligation (ALPPS)**

ALPPS, a newer strategy to increase resectability of hepatic malignancies, has been described for the first time in 2010 (35). This method relies on the fact (proved in clinical trials) that any closure of portal branch will be followed by a reactive perfusion through intrahepatic branches and collaterals present between two lobes. Hence, partition of the liver along the falciform ligament line, for example, will enhance regeneration compared to traditional methods. ALPPS has shown high hypertrophy rates compared to PVE/PVL (40% to 80% within a week compared to 8% to 27% within 2 to 60 days by PVL/PVE), however it is associated with high morbidity rates (16-64% of patients) and mortality rates (12-23% of patients), therefore a careful selection of surgical candidates should be done prior to surgery. Further investigation if ALPPS approach accelerates tumor growth is still required (35-37).

Recently, a number of comparisons between ALLPS and standard methods (PVE followed by liver resections) have been published (38-40). One of the proposed benefits of ALLPS, for example, is rapid removal of tumor(s), thus preventing patient dropout due to disease progression of existing liver tumors. This assumption, however, failed to achieve clinical relevance in a recent publication that compared right PVE + segment 4 to ALPPS, demonstrating mainly extra-hepatic location of metastasis in the patient’s drop-out group. In addition, using PVE in this study yielded sufficient growth in 96.5% of the patients, with median hypertrophy of 62%, comparable to the FLR hypertrophy rates associated with the ALLPS approach (38).

Although none of the studies published with this technique provide measurements of portal pressure or portal blood flow, the clinical data suggest that the acceleration of the hypertrophy of the residual parenchyma occurs due to the reduction of intra-hepatic communicants, once the *in situ* split procedure leads to complete portal devascularization of segment 4, preventing formation of collaterals between the left and the right liver that could otherwise undermine the completeness of right portal vein occlusion alone (41).

A second and not mutually exclusive explanation would be the “regenerating liver” hypothesis proposed by Nagano *et al.* (42).

**Modulation of portal pressure**

Intraoperative Doppler ultrasonography has been used in combination with hepatic portal inflow modulation to detect and offset hyperperfusion in a small-for-size graft. Importantly, numerous interventions that modulate the portal blood flow have been shown to prevent the development of the SFSS in experimental models, such as: the performance of a portocaval anastomosis (43,44), the ligation of the splenic artery (45), banding of the portal vein (46) or the infusion of adenosine (47), somatostatin (48), pentoxifylline (49) or endothelin-1 (50). It is important to highlight that the role for inflow modulation at the time of major liver resection or as a salvage therapy in humans remains undefined.

After all these studies cited above we can conclude that the development of SFSS or PLF are not strictly determined by the “size” of the liver graft or remnant. It is determined by the hemodynamic parameters of the hepatic circulation and, specifically, by a portal blood flow that, when excessive for the volume of the liver parenchyma leads to over-pressure, sinusoidal endothelial denudation and hemorrhage. Perisinusoidal and periportal hemorrhage occurs in the first minutes after an extended hepatic resection as well as after the reperfusion of a small graft, while arterial vasoconstriction and ischemic cholangitis are observed at later stages (6).

Also, experimental and clinical studies consistently show that an increased portal blood flow relative to the weight of the liver results in an inverse relationship between portal and arterial blood flows that is known as the arterio-portal...
buffer (51). The arterio-portal buffer occurs when the portal blood flow increases, the adenosine concentration in the space of Disse decreases leading to arterial vasoconstriction and decrease of arterial blood flow, which is responsible for the late damage (52).

Studies performed in patients undergoing liver transplantation in which the portal and hepatic arterial blood flows were measured intra-operatively have provided further insights into the pathophysiology of the SFSS (6,53,54). A portal blood flow of 300 mL/min/100 g was established by Jiang et al. as the threshold above that the incidence of the SFSS increases significantly (54).

In living donor liver transplantations involving grafts with GWRW below 0.8, Troisi et al. showed that the construction of a portal-systemic shunt whenever the portal blood flow exceeded 250 mL/min/100 g was able to prevent the histological alterations characteristic of the SFSS and to improve the overall patient and graft survival (43,54).

Several studies indicate that additionally to blood flow, portal pressure can also be considered a good parameter for predicting the failure of the graft. For example, patients with a portal pressure higher than 20 mmHg show a decrease from 85% to 38% in their 6-month survival (55). Yagi et al. also described that a portal pressure above 20 mmHg was associated with the development of ascites, coagulopathy and hyperbilirubinemia as well as with an early hypertrophy of the graft, higher values of hepatocyte growth factor (HGF) and diminished levels of vascular epithelial growth factor (VEGF), suggesting that an increased portal pressure also influences liver regeneration (56). Kaido et al. reported their experience with small grafts (GWRW of 0.6) in combination with portal pressure control (targeting final portal pressures below 15 mmHg), showing that the survival of recipients of small grafts and standard-size grafts was similar and that the portal pressure control strategy resulted in a decreased rate of complications in the donors (57).

As in liver transplantation, studies involving extended hepatic resections also indicate that the increased portal blood flow with diminished residual parenchyma are a critical factor determining the development of PLF (47,58,59). The performance of a portocaval anastomosis in a patient with liver cirrhosis undergoing a major hepatectomy effectively prevented the syndrome, probably by reducing shear stress and damage to the sinusoids (60).

**Post-operative period-treatment (61)**

PLF is a quite complex disease, that requires a multi-disciplinary approach, where it management must be undertaken in conjunction with critical care, hepatology, microbiology and radiology services (1).

After liver resection, clinical and laboratory assessment should be proceeded. Normally, the level of serum bilirubin and the INR rises in the first 48-72 h after resection. It is possible to identify liver dysfunction, whenever bilirubin concentration is above 50 µmol/L (3 mg/dL) or INR greater than 1.7 beyond 5 days of surgery (3). The most sensitive variable is serum bilirubin as predictor of outcome in PLF (62). PT and INR are also relevant, but the interpretation may be compromised if patients have received clotting factors.

Serum albumin, although an indicator of hepatic synthetic function, will vary in response to inflammation and administration of intravenous fluids (63,64). Increased levels of liver enzymes are common after liver resection and do not predict outcome (3).

Ascites and hepatic encephalopathy are important markers for liver failure, although it may be difficult to assess in the immediate postoperative period. The first occurs as a result of surgery (portal hypertension, dissection, gross fluid overload), while the second is a result of mental state as collateral effect of drugs such as opiates (62).

Several studies assessed the role of postoperative functional of the liver. This task still consist a challenge, once the ICGR15 is capable to predicts PLF (65), but its value diminishes once liver failure is established, since the changes in hepatic blood flow impacts ICGR15. In the absence of controlled trials for PLF, management relies on data from experience with ALF, secondary to paracetamol toxicity (66-68).

The pattern of organ dysfunction that occurs as a result of PLF is similar to that in sepsis (1). Once the following symptoms occur: cardiovascular failure, characterized by reduced systemic vascular resistance and capillary leak; acute lung injury, due to pulmonary edema and acute respiratory distress syndrome may ensue and acute kidney injury can progress rapidly in PLF. In those cases, fluid balance should be managed judiciously with avoidance of salt and water overload (64). Identifying and treating underlying sepsis is key in managing patients with PLF. Sepsis may exacerbate PLF, and bacterial infection is present in 80 per cent of patients with PLF (69) and in 90 per cent of those with ALF (70).

Therefore, any acute deterioration should be attributed to sepsis until proven otherwise. Management of sepsis should be in accordance with the surviving sepsis guidelines (71). A trial of prophylactic antibiotics after liver
resection failed to show a reduction in liver dysfunction or infective complications (72). A study of ALF have shown that prophylactic antibiotics reduce infections, but the impact on a long-term outcome is inconclusive (70). In critically ill patients with PLF, chest radiography and cultures of blood, urine, sputum and drain site/ascitic fluid should be performed (68). Current guidelines for ALF propose that broad-spectrum antibiotics should be administered empirically to patients with progression to grade 3 or 4 of hepatic encephalopathy, renal failure and/or worsening SIRS parameters (68).

Additionally coagulopathy may occur transiently after major resection and is found in all patients with PLF. As in ALF, coagulation parameters can be used to chart the progress of PLF, provided blood products have not been given. In the absence of bleeding it is not necessary to correct clotting abnormalities, except for invasive procedures or when coagulopathy is severe. The level at which a coagulopathy should be corrected before an interventional procedure in ALF has yet to be defined (66,68,73). Vitamin K may be given, but this is not supported by clinical trials (66). Thrombocytopenia may complicate liver failure (74). Indications for platelet transfusion in ALF include bleeding, severe thrombocytopenia (less than 20×10⁶/L), or when an invasive procedure is planned. A platelet count above 70×10⁶/L is deemed safe for interventional procedures (75). Recombinant factor VIIa (rFVIIa) has been used to treat coagulopathy in patients with ALF (76). In a large controlled trial of rFVIIa following major liver resection, no reduction in bleeding events was observed (77). Its role in PLF is yet to be defined.

Gastrointestinal hemorrhage is a recognized complication of liver failure. In ALF, H2-receptor blockers and proton pump inhibitors (PPIs) reduce gastrointestinal ill patients ensuring euglycemia improves survival and reduces morbidity (78).

The role of imaging in PLF is to assess hepatic blood flow, identify reversible causes of liver failure and locate sites of infection. Hepatic blood flow can be evaluated using non-invasive imaging. Doppler ultrasonography may identify portal vein, hepatic artery and hepatic vein thrombosis. Contrast CT or MRI can be used to establish hepatic blood flow, provide more details of vascular abnormalities and identify sites of infection. If patency of hepatic vessels is still in doubt on cross-sectional imaging, angiography is the “gold standard” (79).

Portal vein thrombosis has also been implicated in the development of PLF. In these rare cases of inflow and outflow thrombosis with PLF, a decision must be taken regarding the benefit of surgical or radiological thrombectomy or dissolution versus anticoagulation (80,81). The use of terlipressin also can reduce the portal venous pressure helping to hepatic regeneration (82). Cerebral edema and intracranial hypertension may occur as a result of PLF. It is unlikely in patients with grade 1 or 2 of liver encephalopathy. When achieving grade 3 encephalopathy, a head CT should be performed to exclude intracranial hemorrhage or other causes of declining mental status.

In patients with established ALF and encephalopathy, enteral lactulose might prevent or treat cerebral edema, although the benefits remain unproven. Progression to grade 3/4 encephalopathy warrants ventilation and may require intracranial pressure monitoring (68).

The concept of hepatocyte transplantation has been investigated as a strategy to boost residual liver function. Intrahepatic hepatocyte transplantation (83) has been used successfully to treat patients with metabolic disorders of the liver. However, results in liver failure (including patients with PLF) have been poor due to insufficient delivery of functional cells. The potential for stem cell therapies has yet to be established (84).

The use of salvage hepatectomy and orthotopic liver transplantation for PLF has been reported in seven patients who underwent liver resection for cancer (85). Although the indications for transplantation in this study were questionable, overall 1-year (88 per cent) and 5-year (40 per cent) survival rates were promising.

Extracorporeal liver support (ELS) devices fall into two categories: artificial and bioartificial systems. Artificial devices use combinations of haemodialysis and adsorption over charcoal or albumin to detoxify plasma. Bioartificial devices use human or xenogenic hepatocytes maintained within a bioreactor to detoxify and provide synthetic function. These systems have not been evaluated extensively in patients with PLF. A recent meta-analysis and systematic review showed that ELS may improve survival in patients with ALF, but not acute-on-chronic liver failure, in comparison with standard medical therapy (86).

**Conclusions**

The increased use of small liver grafts and the expansion of indications of curative liver surgery in patients with hepatic tumors allows a step change in the knowledge of the mechanisms responsible for the development of the SFSS
and PLF.

It became evident that the size of the liver cannot be considered the main variable in the development of liver dysfunction after extended hepatectomies. Additional characteristics should be taken into account, such as: the future liver remnant; the portal blood flow and pressure and the exploration of the potential effects of regeneration preconditioning are all promising strategies that could help to expand the indications and increase the safety of liver surgery.

Acknowledgements

Disclosure: The authors declare no conflict of interest.

References


51. Eipel C. Regulation of hepatic blood flow: The hepatic...


57. Kaido T, Mori A, Ogura Y, et al. Lower limit of the graft-to-recipient weight ratio can be safely reduced to 0.6% in adult-to-adult living donor liver transplantation in combination with portal pressure control. Transplant Proc 2011;43:2391-3.


63. Lobo DN, Stanga Z, Aloysius MM, et al. Effect of volume loading with 1 liter intravenous infusions of 0.9% saline, 4% succinylated gelatine (Gelofusine) and 6% hydroxyethyl starch (Voluven) on blood volume and endocrine responses: a randomized, three-way crossover study in healthy volunteers. Crit Care Med 2010;38:464-70.