Posthepatectomy Liver Failure

Teresa Schreckenbach  Juliane Liese  Wolf O. Bechstein  Christian Moench
Department of General and Visceral Surgery, Johann Wolfgang Goethe University, Frankfurt am Main, Germany

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Posthepatectomy liver failure • Risk analysis • Prevention • Future liver remnant • Cirrhosis

Abstract
Background: Posthepatectomy liver failure (PHLF) is one of the most serious complications after liver resection and is still reported in up to 8% after liver resection. Aims: To provide an overview about the current status of risk analysis and definition of PHLF. Prevention and treatment is also discussed. Methods: A literature review was carried out on PubMed using the terms 'liver failure', 'posthepatectomy' and 'liver surgery' to search relevant papers. Discussion: PHLF remains a serious problem in patients undergoing major liver resection. Adequate preoperative risk assessment and an optimal postoperative treatment are essential for PHLF prevention.

Introduction
Posthepatectomy liver failure (PHLF) is one of the most serious complications after liver resection. Although morbidity and mortality after liver surgery improved over the past 10 years, PHLF is still reported in up to 8% depending on the patient's condition and on functional reserve of the liver before resection [1–3].

Knowledge of the incidence as well as risk factors is essential to prevent PHLF. Scoring systems can be useful to stratify between functional respectability and non-respectability. After surgery, these systems may be useful for the early diagnosis of PHLF resulting in adequate treatment at an early time point.

This article gives a short overview about a current status of risk analysis, prevention and treatment of liver failure after liver surgery.

PHLF – The Problem of Definition
So far, there is no standardized definition or terminology for the PHLF [4].

In 2010, Rahbari and other members of the International Study Group of Liver Surgery (ISGLS) defined PHLF as 'a postoperative acquired deterioration in the ability of the liver to maintain its synthetic, excretory and detoxifying functions, which are characterized by an increased international normalized ratio and concomitant hyperbilirubinemia on or after postoperative (POD) 5 (table 1) [4].

The ISGLS differentiated also the severity of PHLF in 3 grades from A to C. The classification is shown in table 2.

There are still numerous other definitions of PHLF in the literature. These definitions usually depend on countries and hospitals. They contain blood chemistry results and clinical examination in different variations. For example, in the study of Eguschi et al. [5] in 2000, PHLF was diagnosed when three findings were present in the patient: (1) hepatic encephalopathy, (2) progressive hyperbilirubinemia and (3) reduced heparplastin test.

Since the ISGLS definition is easily comparable, it should be used widely.
The incidence of PHLF varies in the literature between 1.2 and 32%; in the most recent literature, the incidence is up to 8% \[3, 4\]. The wide range reflects the differences in patient populations and performed procedures. Decrease in PHLF in the last years can be explained by the improvement in surgical techniques and in intensive care medicine. But also risk factors of the patient population seem to have a big influence on the occurrence of PHLF.

**Incidence of PHLF – Still a Serious Complication**

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**Risk Factors and Risk Assessment for PHLF**

Identification of risk factors is important to decrease the incidence of PHLF. Three groups of risk factors can be differentiated (table 3):

- **Patient related** – right selection of the patients can prevent PHLF
- **Surgery related** – improvement in technique can prevent PHLF
- **Related to postoperative management** – avoiding of machine ventilation, vasopressors and hepatotoxic drugs may decrease the incidence of PHLF.

**Patient-Related Risk Factors**

Various patient-related risk factors have been described such as age or comorbidity.

Important risk factors for PHLF are pre-existing liver diseases, e.g. steatosis, cirrhosis, cholangitis and chemotherapy-associated hepatotoxicity \[6\]. But also active viral hepatitis can increase the risk of PHLF.

Steatosis of the liver seems to be associated with a higher perioperative rate of complications and increased incidence of PHLF. Patients with a biopsy-proven moderate steatosis showed a higher incidence of PHLF compared to patients with normal liver parenchyma \[7\].

Although morbidity increased in patients with preoperative chemotherapy resulting in CASH (chemotherapy-associated steatohepatitis) or sinusoidal obstruction is a defined risk factor for PHLF as well.

In their study, Tanaka et al. \[11\] analyzed the influence of chemotherapy on liver regeneration. They found no difference in liver regeneration between patients with colorectal liver metastases after chemotherapy who underwent portal vein embolization or first hepatectomy. There was only a significant lower liver hypertrophy between patients with steatosis and patients without steatosis of the liver \(p = 0.04\) \[11\].

**Table 1. Consensus definition and severity grading of posthepatectomy liver failure by the ISGLS** \[4\]

<table>
<thead>
<tr>
<th>Grade</th>
<th>Description</th>
</tr>
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<tbody>
<tr>
<td>A</td>
<td>PHLF resulting in abnormal laboratory parameters but requiring no change in the clinical management of the patient</td>
</tr>
<tr>
<td>B</td>
<td>PHLF resulting in a deviation from the regular clinical management but manageable without invasive treatment</td>
</tr>
<tr>
<td>C</td>
<td>PHLF resulting in a deviation from the regular clinical management and requiring invasive treatment</td>
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</table>

INR = International normalized ratio.
Surgery-Related Risk Factors

Risk factors for PHLF which are related to surgical procedure and perioperative management are the extent of the resection with influence on the size of the future liver remnant (FLR), the time of the Pringle maneuver and operation, and the need for blood transfusion during operation.

The minimal functional liver mass needed for adequate liver function is estimated to be 20–25% if patients have a normal liver parenchyma. Patients with abnormal parenchyma (steatosis, fibrosis or cirrhosis) need an FLR up to 40% as minimal functional liver mass [12].

But also intraoperative blood loss >1,000 ml increases the risk of PHLF [6, 13]. This effect is caused by a fluid shift following excessive blood loss and causes a systemic inflammation because of bacterial transmission. But also coagulopathy following blood loss seems to increase the risk of intra-abdominal hematomas and bacterial infections [14].

A third surgical risk factor which was identified for increased PHLF is prolonged operation time.

Risk Factors Related to Postoperative Management

To prevent the occurrence of PHLF, adequate risk assessment at the right time is necessary. It is important to differentiate between pre- and postoperative risk assessment.

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**Table 2. Grading criteria of PHLF defined by ISGLS**

<table>
<thead>
<tr>
<th>Grade A</th>
<th>Grade B</th>
<th>Grade C</th>
</tr>
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<tbody>
<tr>
<td>Specific treatment</td>
<td>Fresh frozen plasma, Albumin, Daily diuretics, Noninvasive ventilation, Transfer to IMC/ICU</td>
<td>Transfer to ICU, Circulatory support (vasoactive drugs), Hemodialysis, Intubation and mechanical ventilation, Extracorporeal liver support, Rescue hepatectomy/liver transplantation</td>
</tr>
</tbody>
</table>

| Hepatic function | Adequate coagulation (INR <1.5), No neurological symptoms | Inadequate coagulation (INR ≥1.5 <2.0), Beginning of neurologic symptoms (i.e. somnolence, confusion) | Inadequate coagulation (INR ≥2.0), Severe neurologic symptoms/hepatic encephalopathy |

| Renal function | Adequate urine output (≥0.5 ml/kg/h), BUN <150 mg/dl, no symptoms of uremia | Inadequate urine output (≤0.5 ml/kg/h), BUN <150 mg/dl, no symptoms of uremia | Renal dysfunction not manageable with diuretics, BUN ≥150 mg/dl, symptoms of uremia |

| Pulmonary function | Arterial oxygen saturation >90%, May have oxygen supply via nasal cannula or oxygen mask | Arterial oxygen saturation <90% despite oxygen supply via nasal cannula or oxygen mask | Severe refractory hypoxemia (arterial oxygen saturation ≤85% with high fraction of inspired oxygen) |

| Additional evaluation | Not required | Abdominal ultrasonography/CT, chest radiography, sputum, blood, urine culture, brain CT | Abdominal ultrasonography/CT, chest radiography, sputum, blood, urine culture, brain CT, ICP monitoring device |

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**Table 3. Risk factors for PHLF [6]**

<table>
<thead>
<tr>
<th>Patient related</th>
<th>Pre-existent liver diseases (cirrhosis, steatosis, fibrosis, cholangitis)</th>
<th>Neoadjuvant treatment because of chemotherapy-associated hepatotoxicity</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Male gender</td>
<td>Comorbidity</td>
</tr>
<tr>
<td></td>
<td>Advanced age (&gt;65 years)</td>
<td>Malnutrition</td>
</tr>
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| Surgery related | Small remnant liver volume, Excessive intraoperative blood loss (>1,250 ml), Prolonged operation time |

| Miscellaneous | Hepatic parenchymal congestion, Ischemia-reperfusion injury, Infection |

**Preoperative Risk Assessment**

Preoperative risk assessment should include physical examination to clarify the following questions (table 4):

- Comorbid conditions such as diabetes mellitus, pulmonary, renal or cardiovascular diseases
- Pre-existing liver diseases like steatosis, cirrhosis, fibrosis, hepatitis
intensive care course, in addition to the subjective assessment, objective markers are necessary to initiate appropriate treatments as early as possible.

Easy for daily use is the determination of C-reactive protein (CRP). Because CRP is produced in the liver, Rahman et al. [19] postulated a dampened CRP could be sign of liver failure. They were able to show that patients developing a PHLF had a significantly lower CRP level on POD 1 than patients without PHLF [19].

Another useful tool for determining PHLF is the so called ‘50–50’ criteria. They are composed of a prothrombin time less than 50% and a serum bilirubin level >50 μmol/l on POD 3 and 5 as a predictive marker for increased mortality [20]. It was shown in recent studies that these criteria allow an early diagnosis of PHLF. But it was also shown that the 50–50 criteria are not a significant marker for increased morbidity after liver surgery [18].

The MELD (model for end-stage liver disease) system can be used for assessment of PHLF as well. MELD is used for patients with HCC undergoing liver resection to assess their perioperative morbidity and mortality [21]. Today, it is especially used for allocation of liver transplantation candidates. Rahbari et al. [18] could show in a comparison of MELD, 50–50 criteria and PHLF definition of ISGLS that MELD score >8 on POD 5 helps to identify patients with increased morbidity. They showed a sensitivity of 55% for prediction of morbidity and 71% for prediction of mortality. The same study also showed the sensitivity for morbidity and mortality for the ISGLS criteria of PHLF of 23 and 65% [18], respectively. In conclusion, the MELD score gave the best early prediction of increased morbidity and mortality after liver surgery, and the PHLF definition was able to assess the risk for increased mortality [18].

During treatment in the ICU, the Acute Physiology and Chronic Health Evaluation (APACHE) score is used to predict the outcome of patients in the ICU. The APACHE system was introduced in 1981 by Knaus et al. [22] for prediction of mortality in the ICU. Because of advances in intensive care medicine, the system had to be updated, and currently APACHE III score is used. In 1998, Hamahata et al. [23] demonstrated that APACHE III score gives a sufficient prediction of mortality after liver resection. Unfortunately, this study was only performed on patients with cholangiocellular carcinoma, and up to now there have been no further analyses for other groups of patients after liver resections.
Prevention of PHLF – What Can Be Done to Avoid PHLF?

First of all, there are risk factors which cannot be influenced like age, gender, existence of cirrhosis or fibrosis and diagnosis of the patient. This has led to a careful indication for liver resections. Patients in poor general condition have a higher risk for perioperative complications and PHLF. But there are possibilities to improve patient general condition and to decrease risk factors for PHLF like small liver remnant volume.

Improving Patient’s Clinical Condition

Liver surgery candidates should fulfill the criteria for general operational capability. Comorbid conditions should be optimized before surgery as much as possible. The Classification of the American Society of Anesthesiology gives a good overview of general capability of the patient.

No improvement in postoperative course and outcome may be prevented with preoperative weight reduction in patients with overweight or obesity. In a study of Cuchetti et al. [24], it was shown that liver surgery can be performed safely in overweight and obese patients with HCC. There was only a significantly increase in mild respiratory complications (p = 0.044) in these patients.

On the other hand, many patients undergoing liver surgery suffer from malignant diseases or from advanced liver diseases and have a bad nutritional status [25]. To decrease the risk of general complications, optimization of nutritional status especially in patients with cirrhosis may be helpful [26], but no relationship between malnutrition and PHLF has been demonstrated [6].

Removal of cholestasis is not a guarantee that PHLF will be avoided either. Different studies tried to show an advantage in patients with jaundice getting percutaneous transhepatic drainage (PTC) before liver surgery, but there was never a significant improvement in outcome. Normally, PTC-related complications led to an increase in overall complications [27].

Improving Size and Function of FLR

As mentioned above, patients with normal liver parenchyma need about 25% FLR in order to have an adequate liver function after surgery. Preoperative strategies to increase the size of the FLR are a useful tool in PHLF prevention.

First of all, it is possible to be cautious by the extent of resection. Further possibilities are two-stage hepatectomy, downsizing with neoadjuvant chemotherapy and hypertrophy induction.

In patients with normal liver function, portal vein embolization of the diseased side of the liver is one way to avoid small liver remnant. With portal vein embolization, a volume increase from 28 up to 46% is obtainable depending on pre-existing liver disease [28, 29].

Improving Intraoperative Course

As excessive intraoperative blood loss is a risk factor for PHLF, improvement in dissection technologies in the last years has led to a decrease in the volume of blood loss during liver resections and improved postoperative outcome. An easy method to lower blood loss is the use of portal vein clamping. An older study reported a lower blood loss due lowering central venous pressure during dissection to ≤5 mm Hg [30]. A possible way to avoid ischemic damage in the liver remnant seems to be the so-called ischemic precondition. It means a temporal clamping of portal triad before a prolonged clamping. It was shown that this helps to reduce hepatocyte damage and is associated with less morbidity, but there was no influence on PHLF occurrence [31].

Improving Postoperative Management

Adequate postoperative monitoring is essential to predict postoperative complications early enough. Postoperative liver enzymes, albumin, creatinine and blood coagulation should be monitored, and patients should be clinically reevaluated on a regular basis.

Patients that develop complications like encephalopathy, hypocoagulation or jaundice should be placed in intermediate care (IMC) or in the ICU for better monitoring, and should be checked for PHLF development.

Treatment of PHLF Remains Difficult

Although surgical techniques and postoperative treatment has improved in the last years, treatment of PHLF still remains difficult.

If PHLF is detected in a patient, it should be scored by the ISGLS system. PHLF grade A should be monitored well, but will normally not be in need of specific treatment. If PHLF grade B occurs, it has to be evaluated if the patient should be placed in IMC or in the ICU. Patients with PHLF grade C need invasive treatment and have to be placed in the ICU [4].

Pulmonary, renal and circulatory disturbances should be treated with a goal-directed therapy regime. Diuretics
and renal replacement therapy have to be used as indicated. In cases of pulmonary insufficiency, noninvasive or invasive ventilation has to be used [4].

For specific treatment of hepatic insufficiency, there are still little options. Goal-directed therapy means application of human albumin, fresh frozen plasma PPSB or AT III to cover the liver function [6].

Liver support systems have been available for some years now:
- Molecular absorbent recirculating system (MARS®)
- Modified fractionated plasma separation and adsorption (Prometheus®)
- Bioartificial liver and extracorporeal liver assist device.

**Molecular Absorbent Recirculating System**
MARS is an extracorporeal albumin dialysis where blood is dialyzed against an albumin-enriched solution. There are only few studies dealing with the use of MARS in cases of PHLF. Van de Kerkhove et al. [32] reported on 5 patients in the Netherlands who were treated with MARS because of PHLF. Three patients showed significant improvement in clinical course, and all patients showed improvement in biochemical parameters. But only one patient survived. Also a study from Hong Kong Hospitals showed a poor outcome in patients with PHLF and MARS therapy despite of improvement in biochemical parameters [33].

It seems that MARS therapy improved clinical and biochemical parameters, although patients died of other complications before their liver function could recover. Since there is a persistent lack of randomized data, all questions concerning the modality of MARS treatment, like time point or duration, are still unanswered.

**Modified Fractionated Plasma Separation and Adsorption (Prometheus)**
The Prometheus system also uses albumin dialysis as principle. Albumin-bound toxins are eliminated through an albumin-permeable membrane, and cleaned albumin is returned to a patient. Prometheus seems to have a better detoxifying capacity than MARS, but there is also a lack of data regarding its use in PHLF, so future studies have to show if it will become a standardized therapy for PHLF.

**Liver Transplantation**
Patients that have no contraindications for liver transplantation can be considered as liver transplantation candidates if severe PHLF occurs. In the daily praxis, however, most indications for liver resections are contraindications for liver transplantation, and therefore this option usually is not an alternative.

**Conclusion**
PHLF remains a serious and life-threatening problem in patients with major liver resections or limited functional reserve due to preexisting liver disease. Adequate preoperative risk assessment of liver function and general condition, gentle and parenchyma-sparing surgery and optimal postoperative treatment are essential in PHLF prevention. Early diagnosis of PHLF can help initiate treatment in the ICU to optimize and recover both hepatic and extrahepatic organ function. Extracorporeal liver devices are still experimental medicine, and their use can only be recommended in prospective randomized trials.

**References**
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