

## **Changes of immune function and its correlation with biliary tract infection before and after biliary stent in malignant obstructive jaundice.**

**Chengli Liu<sup>1\*</sup>, Cheng Wang<sup>1</sup>, Hui Zhang<sup>1</sup>, Xiaoxia Zang<sup>2</sup>, Gang Zhao<sup>1</sup>, Yalin Kong<sup>1</sup>, Yingjie Wang<sup>3</sup>, Hongqi Li<sup>3</sup>**

<sup>1</sup>Department of Hepatobiliary Surgery, Air Force General Hospital of Chinese People's Liberation Army, Haidian, Beijing, PR China

<sup>2</sup>Department of Stomatology, Air Force General Hospital of Chinese People's Liberation Army, Haidian, Beijing, PR China

<sup>3</sup>Department of Oncological Radiotherapy, Air Force General Hospital of Chinese People's Liberation Army, Haidian, Beijing, PR China

### **Abstract**

**Objective:** To investigate the changes of immune function and its correlation with biliary tract infection before and after biliary stent in malignant obstructive jaundice.

**Methods:** A total of 148 patients from September 2014 to 2016 were collected in our hospital, the level of glycosylated haemoglobin, white blood cells, CD4+ T cell count, CD8+ T cell count and CD4+/CD8+ ratio, neutrophil count and percentage of neutrophil etc., total bilirubin, alanine aminotransferase, plasma hs-CRP, TNF- $\alpha$ , IL-1 $\beta$ , IL-6, IL-8 were recorded and compared before and after surgery.

**Results:** After a follow-up of 6 weeks, the CD4+ T lymphocytes count was significantly increased than that of before surgery (F=10.23, P=0.02). After follow-up of 6 weeks, the proportion of CD4+/CD8+ increased to  $2.37 \pm 1.09\%$ , compared with that of before biliary stent implantation  $0.71 \pm 0.21\%$ , the difference was statistically significant (F=21.74, P=0.02). 3 weeks after operation, hs-CRP, TNF- $\alpha$ , IL-1 $\beta$ , IL-6, IL-8, the levels of total bilirubin, alanine aminotransferase were significantly lower than that before operation (F=12.83, 18.45, 29.4, 12.8, 10.6, 9.29, 10.08; P=0.03, 0.01, 0.01, 0.02, 0.02, 0.00, 0.01, respectively). High risk factors for increasing CD4+/CD8+ ratio were investigated with patients of malignant obstructive jaundice, which includes blood glucose, glycosylated hemoglobin, ALT, AST, LDL-C, HDL-C and triglycerides, hs-CRP, neutrophil count, neutrophil percentage, T lymphocyte count, ratio of CD4+/CD8+ and biliary tumor size.

**Conclusion:** For patients with malignant obstructive jaundice, biliary stent could improve cholestasis by increasing the proportion of CD4+/CD8+ T lymphocytes, which can improve immune system, function for patients with malignant obstructive jaundice.

**Keywords:** Malignant obstructive jaundice (MOJ), Biliary tract infection, Immune function, Biliary stent.

*Accepted on August 03, 2017*

### **Introduction**

Malignant Obstructive Jaundice (MOJ), is a series of clinical signs caused by bile duct obstruction due to hepatic and external bile duct obstruction, which will result in growth, invasion, or metastasis of tumors [1,2]. At present, percutaneous biliary stent implantation is a mainly surgery characterized by minimally invasion, higher efficiency and success rate of reducing jaundice [3]. However, postoperative complications such as infection, bleeding, bile leakage, pancreatitis and so on are more common, especially the postoperative infection [4,5]. According to statistics, the incidence of complications can be as high as 30-50%, while the endotoxin was as high as 50-80% [6]. After injury of the intestinal mucosal barrier, jaundice will probably cause the

imbalance of intestinal flora. If infection occurs at the same time, it can easily cause liver failure and multiple organ dysfunction syndrome [7,8]. In this condition, the efficiency of anti-infection treatment is very low. If the MOJ induced enterogenous endotoxin and got into the circulation, it will further cause sepsis, MODS and even death during the peri-operative period [9,10].

In this study, 148 patients with malignant obstructive jaundice were enrolled. Logistic regression analysis was performed on the patients with CD4+/CD8+ ratio and biliary tract infection, and the related factors were also investigated. In addition, we performed the biliary stent procedures for the enrolled patients and observed its effect on immune function changes after

surgery. Finally, we evaluated the clinical efficacy of biliary stent implantation in the treatment of jaundice.

## Materials and Methods

### Clinical data

148 cases of patients with malignant obstructive jaundice were collected from September 2014 to 2016, including 78 males and 70 females with an average age of  $61.8 \pm 4.6$  y old. The course of the disease varied from 3 months to 5 y, and the average course was  $2.7 \pm 1.3$  y. Average operation time was  $65.2 \pm 12.6$  min. Average blood loss was  $56.4 \pm 4.7$  ml. Average hospital stay was  $6.8 \pm 1.3$  d. All the patients in this research group were informed and signed the informed consent. This study has been approved by the Ethics Committee of our hospital.

### Inclusion and exclusion criteria

**Diagnostic criteria:** Biochemical indicators: total bilirubin exceeds the upper limit of the normal value; direct bilirubin/total bilirubin > 50%; Radiographic parameters: Intrahepatic tumor, with or without extra hepatic bile duct dilatation, clinical manifestation of invasion or compression in extra hepatic bile duct caused by malignant tumor. Other indicators: Positive urine bilirubin and clay colored stools.

**Inclusion criteria:** (A) The clinical or pathological examination identified as malignant obstructive jaundice; (B) No surgical indications; (C) Patients agreed to perform percutaneous biliary stent implantation and had signed informed consent prior to surgery.

**Exclusion criteria:** (A) Accompanied with duodenal obstruction, patients are unable to take food orally preoperatively; (B) Unsuccessful biliary stent implantation; (C) Hemorrhage and other non-infectious causes leads to exacerbations; (D) Before biliary obstruction, liver dysfunction caused by primary liver diseases such as viral hepatitis; (E) Comorbidities such as serious diseases of heart, lung and kidney; (F) After surgery, patients are unwilling to cooperate to continue the examination and treatment.

## Methods

### Observational parameters

Venous sterile whole blood 4-6 ml was extracted, peripheral blood leukocyte and neutrophil were detected. Quantification of T lymphocyte subsets by monoclonal antibody assay. Detection method: HbA1c detection, with fasting, peripheral blood 4-6 ml of patients was detected by Variant II hemoglobin detector (Bio-Rad Company, United States). Fasting Blood Glucose (FBG): Medtronic (Medtronic, USA) portable blood glucose meter was applied to detect FBG. Other biochemical indicators: Blood lipid, aminotransferase and other biochemical indicators were detected by Roche Modular automatic biochemical analyzer. CRP was determined to use

the immune assay (Zhongshan golden bridge, China). The determination of TNF- $\alpha$ , IL-1 beta, IL-6, IL-8, etc., were detected by enzyme-linked immunosorbent assay (Cenzyme Kit).

### Biliary stent implantation

(1) Take the eighth intercostal space of right midaxillary line as puncture point, inject the 37% iopromide injection to reveal intrahepatic bile duct dilatation through percutaneous transhepatic. (2) Exit the stylet then intubate the guide wire, adjust the guide wire to get it through the stricture bile duct. (3) Implant the puncture trocar through the fine guide wire. (4) Implant the stiff type guide wire to get it through the stricture bile duct then into the duodenum. (5) Use the stiff type guide wire to reach biliary stenting, and then reveal that the contrast agent passed successfully and got into the duodenum. (6) Indwell the external drainage catheter through the guide wire, implant the end of the guide wire in common bile duct, exit the guide wire, then fix the external drainage catheter to skin (Figures 1a-d).

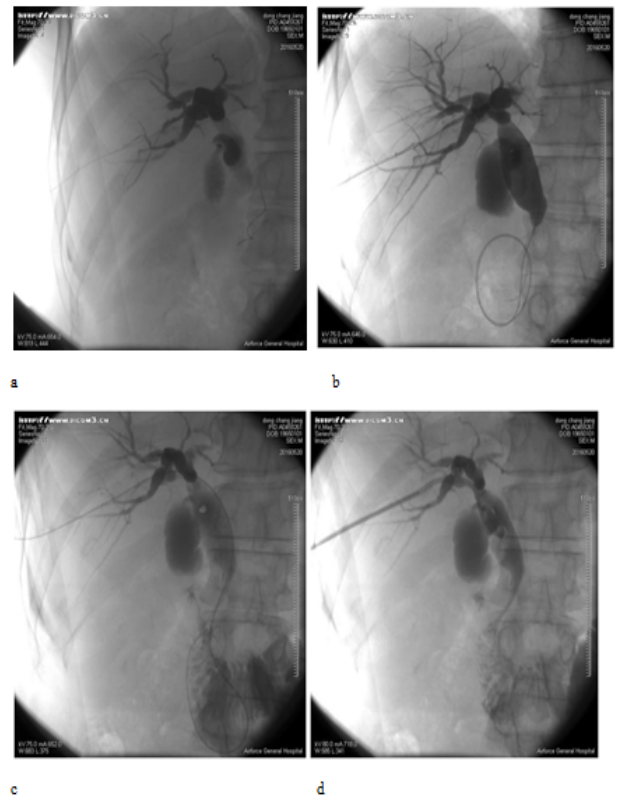


Figure 1. Biliary stent implantation.

### Statistics

SPSS19.0 software was used for statistical analysis in this study, quantitative data were compared by analysis of variance (ANOVA), fisher exact method. Correlation between postoperative lymphocyte changes and biliary tract infection were evaluated by Pearson's correlation, the difference was statistically significant when  $P < 0.05$ .

## Results

### Comparison of results of T cell subsets

After operation, the total number of leukocytes, neutrophils and lymphocytes, CD4+ T cell count, percentage of CD4+ T

cells, CD8+ T cell counts, the percentage of CD8+ T cells were all recorded and statistically analyzed. 6 weeks after operation, the CD4+ T cell count and percentage, ratio of CD4+/CD8+ showed significantly difference ( $P<0.05$ ) compared with the preoperative records (Table 1).

**Table 1.** Comparison of results of T cell subsets ( $\bar{x} \pm s$ ).

Items	Cases (n)	Before treatment	3 weeks after operation	6 weeks after operation	F value	P value
CD4+ T cell count (cells/ $\mu$ L)	148	421.7 $\pm$ 102.5	512.6 $\pm$ 79.3*	669.4 $\pm$ 102.4*#	10.23	0.02
Percentage of CD4+ T cells (%)	148	37.4 $\pm$ 14.3	55.6 $\pm$ 14.7*	58.9 $\pm$ 12.6*#	2.24	0.02
CD8+ T cell count (cells / $\mu$ L)	148	337.3 $\pm$ 64.5	435.7 $\pm$ 35.8*	473.2 $\pm$ 56.4*#	0.87	0.39
Percentage of CD8+ T cells (%)	148	430.4 $\pm$ 102.4	521.7 $\pm$ 111.2*	1489.4 $\pm$ 148.4*#	0.97	0.33
CD4+/CD8+ (%)	148	0.71 $\pm$ 0.21	0.89 $\pm$ 0.24*	2.37 $\pm$ 1.09*#	21.74	0.02

Note: \*Compared with before treatment,  $P<0.05$ ; #Compared with 3 weeks after operation,  $P<0.05$ .

### Comparison of percentage of white blood cell count, neutrophil count and percentage of neutrophil

White blood cell count, neutrophil count and percentage of neutrophil after biliary stent implantation were analyzed

statistically, and there was no significant difference ( $P>0.05$ ) (Table 2).

**Table 2.** Comparison of percentage of white blood cell count, neutrophil count and percentage of neutrophil ( $\bar{x} \pm s$ ).

Items	Cases (n)	Before treatment	3 weeks after operation	6 weeks after operation	F value	P value
White blood cell count ( $\times 10^9$ /L)	148	17.65 $\pm$ 11.16	8.87 $\pm$ 2.32*	7.85 $\pm$ 1.13*	28.23	0
Neutrophil count ( $\times 10^9$ /L)	148	12.23 $\pm$ 1.39	7.12 $\pm$ 1.21*	6.87 $\pm$ 0.86*	18.38	0
Percentage of neutrophil (%)	148	82.43 $\pm$ 7.59	73.23 $\pm$ 6.32*	72.40 $\pm$ 6.64*	9.14	0

Note: \*Compared with before treatment,  $P<0.05$ .

### Comparison of inflammatory cytokines and biochemical indexes ( $\bar{x} \pm s$ )

We recorded and statistically analyzed the inflammatory factors and the level of serum biochemical indexes in patients after operation. 6 weeks after treatment, the hs-CRP, TNF- $\alpha$ , IL-1 $\beta$ , IL-6,IL-8 and other indicators were significantly lower

than that before treatment, the difference was statistically significant ( $P<0.05$ ), which is specifically shown in Table 3. Serum creatinine, blood urea nitrogen, serum total bilirubin, alanine aminotransferase, serum amylase, etc., were significantly lower than that before treatment ( $P<0.05$ ) (Table 4).

**Table 3.** Comparison of inflammatory cytokines ( $\bar{x} \pm s$ ).

Items	Cases (n)	Before treatment	3 weeks after treatment	6 weeks after treatment	F value	P value
Hs-CRP (mg/L)	148	217.6 $\pm$ 23.4	168.7 $\pm$ 10.8*	91.2 $\pm$ 3.1*#	12.83	0.03
TNF- $\alpha$ (pg/mL)	148	647.9 $\pm$ 201.3	1004.5 $\pm$ 121.7*	921.7 $\pm$ 22.8*#	18.45	0.01
IL-1 $\beta$ (pg/mL)	148	728.4 $\pm$ 21.8	403.4 $\pm$ 22.8*	271.6 $\pm$ 12.7*#	29.4	0.01
IL-6 (pg/mL)	148	602.4 $\pm$ 31.7	315.4 $\pm$ 12.9*	104.7 $\pm$ 21.7*#	12.81	0.02
IL-8 (pg/mL)	148	602.3 $\pm$ 22.3	366.4 $\pm$ 12.7*	217.3 $\pm$ 8.2*#	10.6	0.02

Note: \*Compared with before treatment,  $P<0.05$ ; #Compared with 3 weeks after operation,  $P<0.05$ .

**Table 4.** Comparison of biochemical indexes ( $\bar{x} \pm s$ ).

Items	Cases (n)	Before treatment	3 weeks after treatment	6 weeks after treatment	F value	P value
Scr ( $\mu\text{mol/L}$ )	32	439.4 $\pm$ 38.6	377.3 $\pm$ 32.4*	236.5 $\pm$ 50.7*#	3.89	0.02
BUN (mmol/L)	32	27.4 $\pm$ 13.4	18.3 $\pm$ 3.4*	14.4 $\pm$ 23.5*#	17.65	0.01
TBIL ( $\mu\text{mol/L}$ )	32	56.4 $\pm$ 21.8	77.8 $\pm$ 23.5*	68.4 $\pm$ 12.7*#	9.29	0
ALT (U/L)	32	91.4 $\pm$ 21.5	69.4 $\pm$ 13.3*	47.3 $\pm$ 12.8*#	10.08	0.01
AMS (U/L)	32	817.8 $\pm$ 212.5	611.3 $\pm$ 1.4*	510.3 $\pm$ 1.3*#	10.87	0.02

Note: \*Compared with before treatment, P<0.05; #Compared with 3 weeks after operation, P<0.05.

### Correlation analysis of risk factors of postoperative infection

Logistic regression analysis of relationship between multiple risk factors and the CD4+/CD8+ ratio were analyzed after biliary stent implantation. The risk factors included gender, age (y), SBP (mmHg), DBP (mmHg), AST (U/L), ALT (U/L), logGGT (U/L), FPG (mmol/L), BMI (kg/m<sup>2</sup>), WBC ( $\times 10^9/\text{L}$ ),

Scr ( $\mu\text{mol/L}$ ), BUN (mmol/L), TBIL ( $\mu\text{mol/L}$ ), AMS (U/L), Hs-CRP (mg/L), TNF- $\alpha$  (pg/mL), IL-1 $\beta$  (pg/mL), IL-8 (pg/mL), IL-6 (pg/mL), Neutrophil count ( $\times 10^9/\text{L}$ ), neutrophil percentage (%), CD4+ T cell count (cells/ $\mu\text{L}$ ), percentage of CD4+ T cells (%), CD8+ T cell count (cells/ $\mu\text{L}$ ) and percentage of CD8+ T cells (%) (Tables 5-7).

**Table 5.** Correlation between CD4+/CD8+ and Age, SBP, DBP, etc.

Items	Age	SBP (mmHg)	DBP (mmHg)	AST (U/L)	ALT (U/L)	Log GGT (U/L)	FPG (mmol/L)	BMI (kg/m <sup>2</sup> )
CD4+/CD8+	r	0.351	0.116	0.328	0.124	0.415	0.106	0.238
	P	>0.05	>0.05	>0.05	>0.05	>0.05	>0.05	>0.05

**Table 6.** Correlation between CD4+/CD8+ and WBC, Scr, BUN, TBIL, etc.

Items	WBC ( $10^9/\text{L}$ )	( $\times$ Scr ( $\mu\text{mol/L}$ ))	BUN (mmol/L)	TBIL ( $\mu\text{mol/L}$ )	AMS (U/L)	Hs-CRP (mg/L)	TNF- $\alpha$ (pg/mL)	IL-1 $\beta$ (pg/mL)	IL-8 (pg/mL)	IL-6 (pg/mL)
CD4+/CD8+	r	-0.021	-0.351	-0.161	-0.482	-0.283	-1.027	1.063	-0.381	-0.191
	P	>0.05	>0.05	>0.05	>0.05	>0.05	<0.05	<0.05	>0.05	>0.05

**Table 7.** Correlation between CD4+/CD8+ and neutrophil count, neutrophil percentage, etc.

Items	Neutrophil count ( $10^9/\text{L}$ )	The neutrophil percentage (%)	CD4+ T cell count (cells/ $\mu\text{L}$ )	Percentage of CD4+ T cells (%)	CD8+ T cell count (cells/ $\mu\text{L}$ )	The percentage of CD8+ T cells (%)
CD4+/CD8+	r	0.02	0.35	0.16	0.28	0.24
	P	>0.05	>0.05	>0.05	>0.05	>0.05

## Discussion

Previous studies have indicated that MOJ may increase the risk of infection after biliary stent implantation. The reasons can be categorized as follows [11,12]: Retrograde infection of biliary tract, low immunity, disruption of intestinal mucosal barrier, etc. In this study, we found various indicators of postoperative infection 3 weeks after surgery, the absolute value of CD4+ T lymphocytes in patients significantly increased (P<0.05). After 6 weeks of follow-up, the proportion of CD4+/CD8+ increased

compared with that before biliary stent implantation, the difference was statistically significant (P<0.05), while the white blood cells, neutrophils were not significantly improved (P>0.05). As a good medium of bile bacteria, when the intestinal juice reflux, it becomes extremely easy to break out retrograde infection from biliary tract [13]. The characteristics of tumor, malnutrition, hyperbilirubinemia, endotoxemia and preoperative radiotherapy and chemotherapy had complicated effects on the immune function of patients. We speculated that the bile deficiency in the gut when bile duct obstruction occurs,

## *Changes of immune function and its correlation with biliary tract infection before and after biliary stent in malignant obstructive jaundice*

the CD4<sup>+</sup>, CD8<sup>+</sup> T lymphocytes and IgA were significantly decreased in the intestinal mucosa [14,15]. It decreased concentration of intestinal secretory IgA and reduced the intestinal immune function, leading to bacterial translocation [16,17]. It has been reported that cytokines also play an important role in this process. Endotoxin stimulates the production of Tumor Necrosis Factor (TNF), which inhibits the immune function of the host cells [18].

In addition, we found in our study, 6 weeks later, hs-CRP, TNF- $\alpha$ , IL-1 $\beta$ , IL-6, IL-8 and other indicators were significantly lower than that before treatment, the difference was statistically significant ( $P < 0.05$ ). Serum creatinine, blood urea nitrogen, serum total bilirubin, alanine aminotransferase, serum amylase and so on were significantly lower than that of before treatment ( $P < 0.05$ ). Many studies have indicated that the inflammatory mediators play an important role in malignant obstructive jaundice [19]. Prostaglandin E (PGE) [20] inhibited T lymphocyte to produce Interferon (INF) and Interleukin-2 (IL-2), which inhibits the activation of T lymphocyte proliferation and the T lymphocyte subsets, so that the activity of NK cells was decreased consequently. In addition, endotoxin could damage the function of mononuclear phagocytic system, and the activity of Kupfer cells in liver was inhibited. Phagocytosis and killing activity of liver cells decreased, which inhibits the immune function [21]. It has been indicated that both inflammatory response and immune dysfunction are the key steps in the development of traumatic (including surgery) sepsis and MODS.

In our study, we found that the risk factors of postoperative infection included gender, age (y), SBP (mmHg), DBP (mmHg), AST (U/L), ALT (U/L), logGGT (U/L), FPG (mmol/L), BMI (kg/m<sup>2</sup>), WBC ( $\times 10^9/L$ ), Scr ( $\mu\text{mol/L}$ ), BUN (mmol/L), TBIL ( $\mu\text{mol/L}$ ), AMS (U/L), Hs-CRP (mg/L), TNF- $\alpha$  (pg/mL), IL-1 $\beta$  (pg/mL), IL-8 (pg/mL), IL-6 (pg/mL), neutrophil count ( $\times 10^9/L$ ), neutrophil percentage (%), CD4<sup>+</sup> T cell count (cells/ $\mu\text{L}$ ), percentage of CD4<sup>+</sup> T cells (%), CD8<sup>+</sup> T cell count (cells/ $\mu\text{L}$ ) and percentage of CD8<sup>+</sup> T cells (%) [22-25]. In MOJ, due to the lack of bile, intestinal mucosal chemical barrier is easily impaired [26]. Bile salt deficiency is a reason of nutrient absorption, intestinal mucosal atrophy, changes of intestinal mucosal morphology and decreased electrical activity of smooth muscle [27]. In MOJ, under the light microscope, the thin ileum mucosa, blunt and sparse villi, edematous epithelium, and the separated epithelial layer from laminae propria of the mucosa can be seen. Under the electron microscope, it characterized as the shed partial epithelial tissue, vacuole cytoplasm, formation of phagocytosis of dissolved body, broken tight junctions between cells, swollen mitochondria and ridge fracture, which caused damage to the mechanical barrier [28].

In summary, we believe that biliary stent implantation in patients with malignant obstructive jaundice will significantly improve the cholestasis. The proportion of CD4<sup>+</sup>/CD8<sup>+</sup> T lymphocytes was increased, by which induced and improved the immune system function, thus reducing the incidence of

infection of patients with malignant obstructive jaundice, and will significantly improve their quality of life.

### **Acknowledgement**

This study was supported by Capital Citizen Health Project Nurture of Beijing (clinical research of preoperative intra-intestinal immune nutrition intervention for biliary stent implantation in malignant obstructive jaundice, Z131100006813010).

### **References**

1. Yamamoto R, Takahashi M, Osafune Y. Comparison of endoscopic stenting for malignant biliary obstruction: A single-center study. *World J Gastrointest Endosc* 2015; 7: 889-894.
2. Andruszkow H, Fischer J, Sasse M. Interleukin-6 as inflammatory marker referring to multiple organ dysfunction syndromes in severely injured children. *Scand J Trauma Resusc Emerg Med* 2014; 22: 16.
3. Phillip LC, Emmanuel SK, Mabula M. Etiological spectrum and treatment outcome of Obstructive jaundice at a University teaching Hospital in North-western Tanzania: A diagnostic and therapeutic challenges. *BMC Res Notes* 2011; 4: 1-7.
4. Kallis Y, Phillips N, Steel A. Analysis of endoscopic radiofrequency ablation of biliary malignant strictures in pancreatic cancer suggests potential survival benefit. *Dig Dis Sci* 2015; 60: 3449-3455.
5. Wang SB, Wu HB, Wang QS. 18F-FDG PET/CT in differentiating malignant from benign origins of obstructive jaundice. *Hepatobiliary Pancreatic Dis Int* 2015; 14: 5166-5222.
6. Yu H, Wu S, Yu X. Single-incision laparoscopic biliary bypass for malignant obstructive jaundice. *J Gastrointest Surg* 2015; 19: 1132-1138.
7. Fan WD, Nie HF, Chen YB. Influence of malignant severe obstructive jaundice on immune function and therapeutic effects of immunopotentiator. *J Hepatopancreatobiliary Surg* 2013; 25: 35-38.
8. Patel P, Rangarajan B, Mangat K. Improved accuracy of percutaneous biopsy using "Cross and Push" technique for patients suspected with malignant biliary strictures. *Cardiovasc Interventional Radiol* 2015; 38: 1005-1010.
9. Aswad MG, Dennison AR, Neal CP. Biliary stenting for benign and malignant obstructive jaundice: safe use of extended stent-change intervals. *Surg Laparosc Endosc Percutan Tech* 2014; 24: 385-390.
10. Brunson BA, Hawes R, Hoffman B. Poor 'real-life' negative predictive value of cross-sectional imaging in obstructive jaundice. *Can J Gastroenterol Hepatol* 2014; 28: 385-390.
11. Semi P, Jeong YP, Moon JC. The efficacy of endoscopic palliation of obstructive jaundice in hepatocellular carcinoma. *Yonsei Med J* 2014; 55: 1267-1272.

12. Artifon EL, Otoch JP, Tchekmedyian AJ. A review on endoscopic palliation of obstructive jaundice before surgery. *Rev Gastroenterol Perú* 2014; 34: 225-228.
13. Pan H, Liang Z, Yin TS. Hepato-biliary-enteric stent drainage as palliative treatment for proximal malignant obstructive jaundice. *Med Oncol* 2014; 31: 853.
14. Li TF, Ren KW, Han XW. Percutaneous transhepatic cholangiobiopsy to determine the pathological cause of anastomotic stenosis after cholangiojejunostomy for malignant obstructive jaundice. *Clin Radiol* 2014; 69: 13-17.
15. Xu C, Lv PH, Huang XE. Internal-external percutaneous transhepatic biliary drainage for patients with malignant obstructive jaundice. *Asian Pac J Cancer Prev* 2014; 15: 9391-9394.
16. Javia SB, Patel R, Singhal S. Endoscopic closure with an over-the-scope clip of a duodenocolonic fistula caused by a migrated biliary stent. *Gastrointest Endosc* 2016; 83: 845-846.
17. Moon JH, Rerknimitr R, Kogure H. Topic controversies in the endoscopic management of malignant hilar strictures using metal stent: side-by-side versus stent-in-stent techniques. *J Hepatobiliary Pancreat Sci* 2015; 22: 650-656.
18. Matsumoto K, Kato H, Tsutsumi K. Successful biliary drainage using a metal stent through the gastric stoma. *World J Gastroenterol* 2015; 21: 7594-7597.
19. Sejjal DV, Vamadevan AS, Trindade AJ. Removal of an embedded, migrated plastic biliary stent with the use of cholangioscopy. *Gastrointest Endosc* 2015; 81: 1482-1483.
20. Sanaka MR, Wadhwa V, Patel M. Retrieval of proximally migrated biliary stent with direct peroral cholangioscopy with an ultraslim endoscope. *Gastrointest Endosc* 2015; 81: 1483-1484.
21. Assimakopoulos SF, Tsamandas AC, Louvros E. Intestinal epithelial cell proliferation, apoptosis and expression of tight junction proteins in patients with obstructive jaundice. *Eur J Clin Invest* 2011; 41: 117-125.
22. Miyazaki M, Shibuya K, Tokue H. Percutaneous transhepatic biliary drainage assisted by real-time virtual sonography: a retrospective study. *BMC Gastroenterol* 2013; 13: 127.
23. Bun Teoh AY, Ning Chong CC, Wong Lau JY. Video of the month: eus-guided choledochoduodenostomy with a cautery-equipped lumen-apposing stent allows future biliary access in patients with type 2 duodenal obstruction. *Am J Gastroenterol* 2015; 110: 800.
24. Zi-Kai W, Jian-Guo X, Xue-Fei H. Effect of biliary drainage on inducible nitric oxide synthase, CD14 and TGR5 expression in obstructive jaundice rats. *World J Gastroenterol* 2013; 19: 2319-2330.
25. Lyon M, Menon S, Jain A. Use of biliary stent in laparoscopic common bile duct exploration. *Surg Endosc* 2015; 29: 1094-1098.
26. Joseph PM, Joseph C. The changing pattern and implications of Multiple Organ Failure (MOF) after blunt injury with hemorrhagic shock. *Crit Care Med* 2012; 40: 1129-1135.
27. Griffith M, Peter JV, Karthik G. Profile of organ dysfunction and predictors of mortality in severe scrub typhus infection requiring intensive care admission. *Indian J Crit Care Med* 2014; 18: 497-502.
28. Naitoh I, Nakazawa T, Ban T. 8-mm versus 10-mm diameter self-expandable metallic stent in bilateral endoscopic stent-in-stent deployment for malignant hilar biliary obstruction. *J Hepatobiliary Pancreat Sci* 2015; 22: 396-401.

**\*Correspondence to**

Chengli Liu

Department of Hepatobiliary Surgery

Air Force General Hospital of Chinese People's Liberation Army

Haidian

Beijing

PR China