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

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Abstract

Objective: To investigate the changes of immune function and its correlation with biliary tract infection before and after biliary stent implantation 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-α, IL-1β, 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 ± 1.09%, compared with that of before biliary stent implantation 0.71 ± 0.21%, the difference was statistically significant (F=21.74, P=0.02). 3 weeks after operation, hs-CRP, TNF-α, IL-1β, IL-6, IL-8 were recorded and compared before and after surgery.

Conclusions: 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.

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 perioperative 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 ± 4.6 y old. The course of the disease varied from 3 months to 5 y, and the average course was 2.7 ± 1.3 y. Average operation time was 65.2 ± 12.6 min. Average blood loss was 56.4 ± 4.7 ml. Average hospital stay was 6.8 ± 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-α, 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).

**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.
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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>
<thead>
<tr>
<th>Items</th>
<th>Cases (n)</th>
<th>Before treatment</th>
<th>3 weeks after operation</th>
<th>6 weeks after operation</th>
<th>F value</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>CD4+ T cell count (cells/μL)</td>
<td>148</td>
<td>421.7 ± 102.5</td>
<td>512.6 ± 79.3</td>
<td>669.4 ± 102.4*</td>
<td>10.23</td>
<td>0.02</td>
</tr>
<tr>
<td>Percentage of CD4+ T cells (%)</td>
<td>148</td>
<td>37.4 ± 14.3</td>
<td>55.8 ± 14.7</td>
<td>58.9 ± 12.6*</td>
<td>2.24</td>
<td>0.02</td>
</tr>
<tr>
<td>CD8+ T cell count (cells /μL)</td>
<td>148</td>
<td>337.3 ± 64.5</td>
<td>435.7 ± 35.8</td>
<td>473.2 ± 56.4*</td>
<td>0.87</td>
<td>0.39</td>
</tr>
<tr>
<td>Percentage of CD8+ T cells (%)</td>
<td>148</td>
<td>430.4 ± 102.4</td>
<td>521.7 ± 111.2*</td>
<td>1489.4 ± 148.4*</td>
<td>0.97</td>
<td>0.33</td>
</tr>
<tr>
<td>CD4+/CD8+ (%)</td>
<td>148</td>
<td>0.71 ± 0.21</td>
<td>0.89 ± 0.24*</td>
<td>2.37 ± 1.09*</td>
<td>21.74</td>
<td>0.02</td>
</tr>
</tbody>
</table>

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>
<thead>
<tr>
<th>Items</th>
<th>Cases (n)</th>
<th>Before treatment</th>
<th>3 weeks after operation</th>
<th>6 weeks after operation</th>
<th>F value</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>White blood cell count (× 10^9/L)</td>
<td>148</td>
<td>17.65 ± 11.16</td>
<td>8.87 ± 2.32*</td>
<td>7.85 ± 1.13*</td>
<td>28.23</td>
<td>0</td>
</tr>
<tr>
<td>Neutrophil count (× 10^9/L)</td>
<td>148</td>
<td>12.23 ± 1.39</td>
<td>7.12 ± 1.21*</td>
<td>6.87 ± 0.86*</td>
<td>18.38</td>
<td>0</td>
</tr>
<tr>
<td>Percentage of neutrophil (%)</td>
<td>148</td>
<td>82.43 ± 7.59</td>
<td>73.23 ± 6.32*</td>
<td>72.40 ± 6.64*</td>
<td>9.14</td>
<td>0</td>
</tr>
</tbody>
</table>

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

Comparison of inflammatory cytokines and biochemical indexes (x̄ ± 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-α, IL-1β, 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>
<thead>
<tr>
<th>Items</th>
<th>Cases (n)</th>
<th>Before treatment</th>
<th>3 weeks after treatment</th>
<th>6 weeks after treatment</th>
<th>F value</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hs-CRP (mg/L)</td>
<td>148</td>
<td>217.6 ± 23.4</td>
<td>168.7 ± 10.6*</td>
<td>91.2 ± 3.1*</td>
<td>12.83</td>
<td>0.03</td>
</tr>
<tr>
<td>TNF-α (pg/mL)</td>
<td>148</td>
<td>647.9 ± 201.3</td>
<td>1004.5 ± 121.7*</td>
<td>921.7 ± 22.8*</td>
<td>18.45</td>
<td>0.01</td>
</tr>
<tr>
<td>IL-1β (pg/mL)</td>
<td>148</td>
<td>728.4 ± 21.8</td>
<td>403.4 ± 22.8*</td>
<td>271.6 ± 12.7*</td>
<td>29.4</td>
<td>0.01</td>
</tr>
<tr>
<td>IL-6 (pg/mL)</td>
<td>148</td>
<td>602.4 ± 31.7</td>
<td>315.4 ± 12.9*</td>
<td>104.7 ± 21.7*</td>
<td>12.81</td>
<td>0.02</td>
</tr>
<tr>
<td>IL-8 (pg/mL)</td>
<td>148</td>
<td>602.3 ± 22.3</td>
<td>366.4 ± 12.7*</td>
<td>217.3 ± 8.2*</td>
<td>10.6</td>
<td>0.02</td>
</tr>
</tbody>
</table>

Note: *Compared with before treatment, P<0.05; #Compared with 3 weeks after operation, P<0.05.
Table 4. Comparison of biochemical indexes (x ± s).

<table>
<thead>
<tr>
<th>Items</th>
<th>Cases (n)</th>
<th>Before treatment</th>
<th>3 weeks after treatment</th>
<th>6 weeks after treatment</th>
<th>F value</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Scr (μmol/L)</td>
<td>32</td>
<td>439.4 ± 38.6</td>
<td>377.3 ± 32.4*</td>
<td>236.5 ± 50.7*</td>
<td>3.89</td>
<td>0.02</td>
</tr>
<tr>
<td>BUN (mmol/L)</td>
<td>32</td>
<td>27.4 ± 13.4</td>
<td>18.3 ± 3.4*</td>
<td>14.4 ± 23.5*</td>
<td>17.65</td>
<td>0.01</td>
</tr>
<tr>
<td>TBIL (μmol/L)</td>
<td>32</td>
<td>56.4 ± 21.8</td>
<td>77.8 ± 23.5*</td>
<td>68.4 ± 12.7*</td>
<td>9.29</td>
<td>0</td>
</tr>
<tr>
<td>ALT (U/L)</td>
<td>32</td>
<td>91.4 ± 21.5</td>
<td>69.4 ± 13.3*</td>
<td>47.3 ± 12.8*</td>
<td>10.08</td>
<td>0.01</td>
</tr>
<tr>
<td>AMS (U/L)</td>
<td>32</td>
<td>817.8 ± 212.5</td>
<td>611.3 ± 1.4*</td>
<td>510.3 ± 1.3*</td>
<td>10.87</td>
<td>0.02</td>
</tr>
</tbody>
</table>

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/m²), WBC (× 10⁹/L), Scr (μmol/L), BUN (mmol/L), TBIL (μmol/L), AMS (U/L), Hs-CRP (mg/L), TNF-α (pg/mL), IL-1β (pg/mL), IL-8 (pg/mL), IL-6 (pg/mL), Neutrophil count (× 10⁹/L), neutrophil percentage (%), CD4+ T cell count (cells/μL), percentage of CD4+ T cells (%), CD8+ T cell count (cells/μL) and percentage of CD8+ T cells (%) (Tables 5-7).

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

<table>
<thead>
<tr>
<th>Items</th>
<th>Age</th>
<th>SBP (mmHg)</th>
<th>DBP (mmHg)</th>
<th>AST (U/L)</th>
<th>ALT (U/L)</th>
<th>Log GGT (U/L)</th>
<th>FPG (mmol/L)</th>
<th>BMI (kg/m²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CD4+/CD8+</td>
<td>r</td>
<td>0.351</td>
<td>0.116</td>
<td>0.328</td>
<td>0.124</td>
<td>0.415</td>
<td>0.106</td>
<td>0.238</td>
</tr>
<tr>
<td>P</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
</tr>
</tbody>
</table>

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

<table>
<thead>
<tr>
<th>Items</th>
<th>WBC (× 10⁹/L)</th>
<th>Scr (μmol/L)</th>
<th>BUN (mmol/L)</th>
<th>TBIL (μmol/L)</th>
<th>AMS (U/L)</th>
<th>Hs-CRP (mg/L)</th>
<th>TNF-α (pg/mL)</th>
<th>IL-1β (pg/mL)</th>
<th>IL-8 (pg/mL)</th>
<th>IL-6 (pg/mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CD4+/CD8+</td>
<td>r</td>
<td>-0.021</td>
<td>-0.351</td>
<td>-0.161</td>
<td>-0.482</td>
<td>-0.283</td>
<td>-1.027</td>
<td>1.063</td>
<td>-0.381</td>
<td>-0.191</td>
</tr>
<tr>
<td>P</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&lt;0.05</td>
<td>&lt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
</tr>
</tbody>
</table>

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

<table>
<thead>
<tr>
<th>Items</th>
<th>Neutrophil count (10⁹/L)</th>
<th>The neutrophil percentage (%)</th>
<th>CD4+ T cell count (cells/μL)</th>
<th>Percentage of CD4+ T cells (%)</th>
<th>CD8+ T cell count (cells/μL)</th>
<th>The percentage of CD8+ T cells (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CD4+/CD8+</td>
<td>r</td>
<td>0.02</td>
<td>0.35</td>
<td>0.16</td>
<td>0.28</td>
<td>0.24</td>
</tr>
<tr>
<td>P</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
<td>&gt;0.05</td>
</tr>
</tbody>
</table>

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+, CD8+ 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-α, IL-1β, 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), WBC (× 10⁹/L), Scr (μmol/L), BUN (mmol/L), TBIL (μmol/L), AMS (U/L), Hs-CRP (mg/L), TNF-α (pg/mL), IL-1β (pg/mL), IL-8 (pg/mL), IL-6 (pg/mL), neutrophil count (× 10⁹/L), neutrophil percentage (%), CD4+ T cell count (cells/μL), percentage of CD4+ T cells (%), CD8+ T cell count (cells/μL) and percentage of CD8+ T cells (%) [22-25]. In MOI, 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 MOI, 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+/CD8+ 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

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References


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