Research about NG-PDT Influence to Thymus-Dependent Lymphocyte Subpopulation from Peripheral Blood of Bronchogenic Carcinoma

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Abstract

Introduction: To research about PDT influence to thymus-dependent lymphocyte subpopulation from peripheral blood of bronchogenic carcinoma.

Method: Taking forty patients as study objects who suffering the bronchogenic carcinoma, the researchers use semi-conductive laser therapy apparatus (made in China) to irradiate the diseased region, and then observe the changed information of thymus-dependent lymphocyte subpopulation from peripheral blood during pre-therapy and post-treatment by immunohistochemical techniques of streptavidin-peroxidase (abbreviation: S-P method).

Result: The level of some medical data in pre-therapy is higher than the data in post-therapy (like CD3+, CD4+, CD8+, CD4+/CD8+, CD45RA+, the absolute value of CD45RA+, the absolute value of CD45RA+ and CD45RA+/RO+); And the level of CD45RO+ (P<0.05) is lower than the data in pre-therapy.

Conclusion: Photodynamic therapy provides the technical support with clinic treatment for bronchogenic carcinoma through immune adjustment to enhance the immunologic function.

Keywords: Photodynamic Therapy; Bronchogenic Carcinoma; T Lymphocyte Subpopulation (Pb)

Introduction

The theory of Photodynamic Therapy (PDT) depends on biological-photodynamic sensitization [1], which has high tissue specificity, low virulence and no immunosuppression. The chemical reaction of luminescence is widely applied to clinical therapy for the reason that it would not take interference into surgical operation, radiotherapy, and chemotherapy, and besides this therapy could be taken repeatedly in limited time of therapy. Because the progression of tumor, especially malignancy, is closely related to immunologic function, it is necessary to discuss the medical effect of immunologic function that photodynamic therapy caused. The key point in this paper is aim to research the medical change of thymus-dependent lymphocyte subpopulation (PB) which the photodynamic therapy caused.

Information and method

General Information

Take forty patients who were suffering bronchogenic carcinoma as study objects, among them there are 26 males and 14 females; The ages are between 47s-75s, and the average age is 60.11±8.09. Pathological classification: 29 cases for lung cancer of gland cell, 11 cases for squamous carcinoma; TNM staging: 7 cases for II stage, 22 cases for III stage, 11 cases for IV stages.

Inclusion criteria

i. Ages75s.
ii. No taking any inhibitor medicine at pre-therapy.
iii. No infectious diseases within one month at pre-therapy.
iv. No fever.
v. No taking blood transfusion within three months at pre-therapy or taking operation (including post operation) under 400ml (or without blood transfusion).
vi. No symptomatic infection at post-therapy.
vii. The one who sign his name on the informed consent.
Exclusion criteria:

viii. Age>75s; with history of immunological diseases; getting complication like infection.

Therapeutic Method

Before taking therapy, all the patients take blood sampling through cubital vein to check T lymphocyte subpopulation on morning with empty stomach. And then 2mg/kg photosensitizer was injected into vein firstly. After 48 hours, 630nm (wave length) laser was irradiated into nidus under endoscope and photodynamic doses was 200J/cm. The range of irradiation should be 1-2cm over the superior and inferior margin of nidus. According to the condition of the disease, the same drug should be injected after 72 hours and 96 hours and reuse the same photodynamic dose [2] several times to take laser radiation.

Detection Method of Thymus-Dependent Lymphocyte Subpopulation

Take 3ml peripheral venous blood at pre-therapy and post-treatment from the patients who suffer bronchogenic carcinoma in the morning; take heparin sodium for anticoagulation as standby application; based on the operating instructions, use immunohistochemical techniques of streptavidin-peroxidase (abbreviation: S-P method) strictly, and the reagent was produced by ShenZhen Kebio Company.

Statistical Method

Analyze the data statistically by software SPSS18.0 ; Measurement data is showed by ( ͞x ±s ), verified by ‘t’ , standard: P<0.05.

Results

Situation Analysis for Variation of Thymus-dependent Lymphocyte Subpopulation

The level of CD3+, CD4+, CD8+ and CD4+/CD8+ at post-therapy is higher than the level at pre-therapy (P<0.05) (Table 1).

Table 1: Comparison for TC Subpopulation at Pre-therapy and Post-therapy ( X ±s )

<table>
<thead>
<tr>
<th>Time</th>
<th>CD3+ (%</th>
<th>CD4+ (%</th>
<th>CD8+ (%</th>
<th>CD4+/ CD8+</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pre</td>
<td>55.29±</td>
<td>45.92±</td>
<td>32.17±</td>
<td>1.09±</td>
</tr>
<tr>
<td></td>
<td>11.20</td>
<td>8.73</td>
<td>6.53</td>
<td>0.29</td>
</tr>
<tr>
<td>Post</td>
<td>63.71±</td>
<td>55.68±</td>
<td>38.20±</td>
<td>1.66±</td>
</tr>
<tr>
<td></td>
<td>14.05</td>
<td>9.20</td>
<td>7.09</td>
<td>0.34</td>
</tr>
</tbody>
</table>

Situation Analysis for Variation of CD45RA+/RO+ at Pre-therapy and Post-therapy

The level of CD45RA+, the absolute value of CD45RA+, the absolute value of CD45RA+/CD45RA+ at post-therapy is higher than the level at pre-therapy, the level of CD45RO+ is lower than pre-therapy (P<0.05) Reference (Table 2).

Table 2: Variation of CD45RA+/RO+ at Pre-therapy and Post-therapy ( X ±s )

<table>
<thead>
<tr>
<th>Time</th>
<th>CD45 RA+ (%</th>
<th>CD45 RA+ (%</th>
<th>CD45 R+ (%</th>
<th>CD45RO+ (%</th>
<th>CD45RA+/ RO+</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pre</td>
<td>49.09± 0.78±</td>
<td>46.37± 0.56±</td>
<td>1.24±</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>12.03</td>
<td>10.22</td>
<td>0.27</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Post</td>
<td>58.78± 1.25±</td>
<td>42.15± 0.98±</td>
<td>1.58±</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>14.55</td>
<td>8.09</td>
<td>0.38</td>
<td></td>
<td></td>
</tr>
<tr>
<td>P</td>
<td>0.028</td>
<td>0.022</td>
<td>0.015</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Discussion

The diagnostic rate of patient with lung cancer on early stage is very low. Most of the patients who go to hospital for treatment are in the middle and advanced stage, so most of the patients miss the optimal cure time. The cancer cell would produce non-tumor-specific immune suppressor so that the immune cell could be suppressed extensively, and then immunologic function of the patients would get decay. CD3+ is representative of total amount of T-lymphocyte; CD4+, which could secrete IL-2,FN-γ and could get synergism with NK cell, is one kind of helper T-lymphocytes. According to practical function, CD8+ could divide into Suppressor T Cell (Ts) and Cytotoxic T Cell (Tc) and it would lead to suppression of immune response [3]. CD4+/CD8+ could be the important indicator for measuring cellular immune function and judging the information of prognosis. NK cell could secrete the cytotoxin straightly so that the tumor cell could be killed. If the activity gets lower, one reason is that the tumor gets progress and another reason is that the tumor gets metastasis. If the specific value gets lower, it means that the immunologic balance gets destroy so that it causes the cellular immune function gets lower.

Photodynamic Therapy is one kind of cold optics reaction, which is consist of photosensitizer, visible light (common laser) and oxygen. The basic action of PDT is biological-photodynamic sensitization which uses the tumor affinity [4] toward Photofrin, to inject into the patients, and then the ground state of tumor cell, where gathered photosensitizer densely, after irradiated by the laser radiation (wave length: 630nm), would translate into sensitized singlet state which would cause chemical reaction with the sounding oxygen subsequently. The research result show that after taking Photodynamic Therapy, some numerical value, like CD3+, CD4+, CD8+, CD4/CD8+, CD45RA+, the absolute value of CD45RA+, the absolute value of CD45RA+ and CD45RA+/ RO+, are all getting distinct variation (P<0.05). It means that PDT could influence thymus-dependent lymphocyte subpopulation from peripheral blood of bronchogenic carcinoma.

References

