

STUDY ON THE EFFECT OF POLYSACCHARIDES FROM *SOLANUM NIGRUM LINNE* ON CELLULAR IMMUNE FUNCTION IN TUMOUR-BEARING MICE

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*E-mail: chenhai10@126.com**Abstract**

We investigated the anti-tumour effect of polysaccharides from *Solanum nigrum Linne*, and its relationship with the immune function of tumour-bearing organisms. MTT assay was used to observe the effect of different doses of polysaccharides from *Solanum nigrum Linne* on proliferation of lymphocytes in tumour-bearing mice. ELISA assay was also used to detect the levels of IL-2 in mice, and a laser scanning confocal microscope was used to detect the effect of polysaccharides from *Solanum nigrum Linne* on intralymphocytic free calcium ion concentration in tumour-bearing mice. Different doses of polysaccharides from *Solanum nigrum Linne* significantly inhibited the growth of mouse H22 solid tumours, improved the survival time of tumour-bearing mice, increased the proliferation of lymphocytes, elevated the levels of IL-2, and increased the concentration of calcium ions in the lymphocytes. Polysaccharides from *Solanum nigrum Linne* have certain anti-tumour effect, which is related with the cellular immune function that regulates the body.

Keywords: *Solanum nigrum* polysaccharides; tumour-bearing mice; cellular immunity

Introduction

Chinese herbal medicine, Long Kui, is the whole plant of *Solanum nigrum Linne*, which is in the genus *Solanum*, family of Solanaceae. It is widely distributed worldwide. The whole plant of *Solanum nigrum Linne* can be used as medicine. It is cold in nature, bitter and slightly sweet in taste, slightly toxic, and has the effects of clearing heat, detoxification, invigorating blood circulation, inducing diuresis, and dispersing swelling. *Solanum nigrum Linne* contains polysaccharides, minerals, vitamins, pigments, and amino acids (Liu et al, 2003; Mi et al, 2002). *Solanum nigrum Linne* fruit also contains esters, sterols, carboxyl compounds, and saponins (Eldridge, 1983). As an ancient medicine, *Solanum nigrum Linne* has been playing an important role in the treatment of cancer in China since ancient times. It is reported in the literatures that *Solanum nigrum Linne* can inhibit tumour cell growth and promote apoptosis of tumour cells, and has immunomodulatory and cytotoxic effects (Son et al, 2003; Yen et al, 2001). Its pharmacological importance also include anti-inflammatory, anti-shock (Jiang, 1977), anti-bacterial, anti-viral (Li et al, 1998), antipyretic, analgesic, cardiovascular protective (Compilation, 1978) as well as expectorant, and antitussive effects (Wang et al, 2007).

Materials and Methods**Medicinal materials**

Solanum nigrum Linne, purchased from Beijing PLA, which was identified by a Professor Sun with voucher No. 2012-5A-52

Experimental animals and cell lines

Kunming mice, weighing 18-22 g, half male and half female, were provided by the Laboratory Animal Center of Beijing PLA; tumour lines: liver cancer (H22) was provided by the Affiliated Hospital of Beijing PLA. All experimental procedures were approved by the Animal Research Ethics Committee of PLA.

Reagents and instruments

ConA: Shanghai Huashun Bioengineering Co., Ltd.; MTT: Sigma, USA; DMSO: Tianjin Damao Chemical Instrument Supply Station; RPMI1640 culture medium: GIBCO, USA; IL-2ELISA kit: Nanjing Jiancheng Reagent Co., Ltd. Other reagents were all of analytical grade: Olympus inverted microscope (OLYMPUS, Japan), low-temperature refrigerated centrifuge (Eppendorf, Germany), clean bench (Suzhou Purification Equipment Co., Ltd.), CO₂ incubator (SANYO, Japan), continuous wavelength microplate reader (Bio-RAD).

Experimental Methods

Preparation of polysaccharides from *Solanum nigrum* Linne

Referring to the method in Su et al (2011), *Solanum nigrum* Linne was decocted twice with an 8-fold volume of water with 2 h each time, and filtered. The filtrates were then combined and dried and *Solanum nigrum* polysaccharides were obtained.

Effect of polysaccharides from *Solanum nigrum* Linne on life prolongation rate of tumour-bearing mice

Kunming mice were randomly divided into 5 groups, with each group containing 10 mice. Under sterile conditions, well-grown ascites were extracted from H22 tumour-bearing mice 7 d after tumour inoculation, and cell morphology and cell count were observed. When the number of tumour cells exceeded 97%, the ascites were eligible for use and were diluted 1:2 with sterile saline, and inoculated intraperitoneally with 0.3 ml per mouse. 24 h later, the mice were weighed, and subcutaneously administered with the drug with 0.4 ml per mouse once daily for 7 consecutive days. The doses in *Solanum nigrum* polysaccharides high-, medium-, and low-dose groups were 120 mg/kg, 60 mg/kg and 30 mg/kg respectively. Negative control group was given equivalent volume of normal saline, and positive control group was given 100 mg/kg Astragalus polysaccharides. Life prolongation rate was observed and recorded.

Effect of polysaccharides from *Solanum nigrum* Linne on lymphocyte proliferation in H22 tumour-bearing mice (Ma et al, 2004; Zhang et al, 2005)

Mice were divided into *Solanum nigrum* polysaccharides high-, medium-, and low-dose groups, positive control group, negative control group and normal group, with each group containing 10 mice. Under sterile conditions, well-grown ascites were extracted from H22 tumour-bearing mice 7 d after tumour inoculation, and cell morphology and cell count were observed. When the number of tumour cells exceeded 97%, the ascites were eligible for use and were diluted 1:2 with sterile saline, and inoculated intraperitoneally with 0.3 ml per mouse. 24 h later, the mice were weighed, and the method of administration was the same as in 2.2. On the second day after administration, the mice were sacrificed, spleens were harvested, and spleen tissues were gently crushed in a plate filled with sterile Hank's solution, and sieved with 200 mesh nylon screen to collect spleen cell suspension. Cell concentration was adjusted to 1×10^6 cells/mL with RPMI1640 culture medium. The experimental groups included those containing ConA and those without ConA, each well of 96-well plate was added with 100 μ L of cell suspension plus 100 μ L of ConA group that did not contain ConA serving as a control. Plates were

incubated in a CO₂ incubator set at 37°C for 48 h. 20 µL of MTT solution was added to each well. After incubating for another 4 h, the supernatant was discarded and 150 µL of DMSO was added to each well, then the plate was shaken in a micromixer for 10 min. After the crystalline was fully dissolved, absorbance of each well was measured at 570 nm.

Effect of polysaccharides from *Solanum nigrum* Linne on IL-2 production in H22 tumour-bearing mice

100 µL of spleen cell suspension were placed in the culture plate, added with 100 µL of ConA, and incubated in a CO₂ incubator set at 37°C for 48 h. Then, the supernatant was collected and cryopreserved at -20°C. OD value at 450 nm was measured with a microplate reader by referring to the mouse IL-2ELISA kit instructions (Shi et al, 2002).

Effect of polysaccharides from *Solanum nigrum* Linne on intralymphocytic Ca²⁺ concentration in H22 tumour-bearing mice

Referring to the method in Zhang et al (2000), after the mice were sacrificed, spleens were harvested under sterile conditions to obtain the spleen cell suspension, and cell concentration was adjusted to 1×10⁹ cells/L with RPMI16640 culture medium. 1 mL of the cell suspension was placed in the eppendorf tube, added with Fluo-3 to make the final concentration 4 µmol·L⁻¹. Then, the suspension was incubated at 37°C for 40 min. After washing three times in PBS, the suspension was resuspended in 1 mL. One drop of the suspension was added on the glass slide, allowed to stand for 10 min, and then covered with coverslip. The sample was placed on a laser scanning confocal microscope with 40X eyepiece, with the fluorescence intensity representing the Ca²⁺, and experimental data were directly input into the computer for processing and image analysis.

Data Processing

Data were all presented as $\bar{x} \pm s$; one-way ANOVA and comparison among groups were performed using the SPSS 10.0 statistical software.

Results

Effect of polysaccharides from *Solanum nigrum* Linne on life prolongation rate of H22 tumour-bearing mice

The experimental results are shown in Table 1. Each *Solanum nigrum* polysaccharides dose group can all significantly prolong the survival time of H22 tumour-bearing mice, which were all higher than that of the negative control group, indicating that the *Solanum nigrum* polysaccharides can extend the survival time of H22 tumour-bearing mice, of which the medium- and high-dose groups had significant survival time prolongation effects. However, the survival times of each dose group were all lower than that of the positive control group.

Effect of polysaccharides from *Solanum nigrum* Linne on lymphocyte proliferation in H22 tumour-bearing mice

The results are shown in Table 2. The number of lymphocytes in tumour-bearing mice was significantly lower than that in the normal group. Different doses of *Solanum nigrum* polysaccharides all improved the lymphocyte proliferation, of which the medium dose group had better effect. The *Solanum nigrum* polysaccharides could also enhance the ConA-stimulated lymphocyte proliferation.

Table 1: Effect of polysaccharides from *Solanum nigrum* Linne on survival time of H22 tumour-bearing mice ($x \pm s$, $n = 10$)

Group	Dose (mg/kg)	Route of administration	Survival time (d)
Negative control group		Ih	9.87±2.56
Positive control group	100	Ih	18.62±2.31**
<i>Solanum nigrum</i> polysaccharides low-dose group	30	Ih	12.98±2.42
<i>Solanum nigrum</i> polysaccharides medium-dose group	60	Ih	16.31±2.01**
<i>Solanum nigrum</i> polysaccharides high-dose group	120	Ih	17.80±2.78**

Note: Comparison with the negative control group: ** $p < 0.01$

Table 2: Effect of polysaccharides from *Solanum nigrum* Linne on lymphocyte proliferation in tumour-bearing mice ($x \pm s$, $n = 10$)

Group	Route of administration	Dose (mg/kg)	Not added with ConA	Added with ConA
Normal group	Ih		0.5483±0.1552	0.6732±0.1783
Negative control group	Ih		0.3422±0.1124	0.3791±0.0912
Positive control group	Ih	100	0.5242±0.1887**	0.6662±0.1052**
<i>Solanum nigrum</i> polysaccharides low-dose group	Ih	30	0.5027±0.1046**	0.5827±0.1083**
<i>Solanum nigrum</i> polysaccharides medium-dose group	Ih	60	0.5184±0.2011**	0.6098±0.1059**
<i>Solanum nigrum</i> polysaccharides high-dose group	Ih	120	0.5019±0.1322**	0.5710±0.1036**

Note: Comparison with the negative control group: ** $p < 0.01$

Effect of polysaccharides from *Solanum nigrum* Linne on IL-2 production in H22 tumour-bearing mice

The results are shown in Table 3. The IL-2 production in tumour-bearing mice was significantly reduced than that in the normal group. After treatment with *Solanum nigrum* polysaccharides, IL-2 production was significantly improved, of which the effect was better in the medium dose group.

Effect of polysaccharides from *Solanum nigrum* Linne on intralymphocytic Ca^{2+} concentration in H22 tumour-bearing mice

The results are shown in Table 4. Intralymphocytic Ca^{2+} concentration in tumour-bearing mice was significantly reduced than that in the normal group, while the Ca^{2+} concentrations in *Solanum nigrum* polysaccharides groups were significantly increased, and the effect was better in the medium dose group.

Table 3: Effect of polysaccharides from *Solanum nigrum* Linne on IL-2 production in tumour-bearing mice ($x \pm s$, n = 10)

Group	Route of administration	Dose (mg/kg)	IL-2
Normal group			0.4274±0.1984
Negative control group	Ih		0.2051±0.0862
Positive control group	Ih	100	0.4025±0.0823**
<i>Solanum nigrum</i> polysaccharides low-dose group	Ih	30	0.3234±0.1045*
<i>Solanum nigrum</i> polysaccharides medium-dose group	Ih	60	0.3921±0.0842**
<i>Solanum nigrum</i> polysaccharides high-dose group	Ih	120	0.3074±0.0833*

Note: Comparison with the negative control group: * p<0.05, ** p<0.01

Table 4: Effect of polysaccharides from *Solanum nigrum* Linne on intralymphocytic Ca²⁺ concentration in tumour-bearing mice ($x \pm s$, n = 10)

Group	Route of administration	Dose (mg/kg)	[Ca ²⁺]
Normal group			2745.52±1621.68
Negative control group	Ih		722.23±1263.21
Positive control group	Ih	100	1732.75±1375.37**
<i>Solanum nigrum</i> polysaccharides low-dose group	Ih	30	1694.47±1679.58**
<i>Solanum nigrum</i> polysaccharides medium-dose group	ih	60	2949.94±1906.74**
<i>Solanum nigrum</i> polysaccharides high-dose group	Ih	120	1285.86±904.43*

Note: Comparison with the negative control group: * p<0.05, ** p<0.01

Discussion

The above experiments showed that the *Solanum nigrum* polysaccharides can prolong the survival time of H22 tumour-bearing mice, promote lymphocyte proliferation, enhance anti-tumour immune function, and thereby achieve the anti-tumour effect. *Solanum nigrum* polysaccharides can also significantly elevate the level of IL-2, which may be accomplished through promoting lymphocytes to produce lymphokines, thus promoting lymphocyte proliferation and IL-2 production. Chinese medicine polysaccharides mainly improve the host's immune function, and enhance the anti-tumour activity. In the treatment of cancer, immune function enhancement plays an important role in the process of tumour cell elimination (HANSK et al, 2006), while *Solanum nigrum* polysaccharides have significant survival time prolongation and immunomodulatory effects in tumour-bearing mice, which thus have a good prospect in cancer treatment.

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