

Ganoderma lucidum polysaccharides eradicates the blocking effect of fibrinogen on NK cytotoxicity against melanoma cells

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Received August 25, 2011; Accepted November 28, 2011

DOI: 10.3892/ol.2011.515

Abstract. Natural killer (NK) cell cytotoxicity is an effective defense against metastatic tumor cells or viruses in the blood. However, NK cytotoxicity against tumor cells may be inhibited by a fibrinogen coat adhered to the surface of tumor cells. *Ganoderma lucidum* (*G. lucidum*) polysaccharides have been reported for their inhibitory ability on the adhesion of type I collagen, hyaluronan, fibronectin and laminin to integrins that were highly expressed on melanoma cells, and were therefore capable of enhancing NK cytotoxicity to tumor cells. In this study, we investigated the effect of *G. lucidum* polysaccharides on fibrinogen's adhesion to melanoma cells and NK cytotoxicity to tumor cells. Melanoma cells B16 and A375 were cultured and analyzed using flow cytometry. Human NK cells were isolated and analyzed using an NK cytotoxic assay. The results showed that polysaccharides extracted from *G. lucidum* inhibit the adhesion of fibrinogen to melanoma cells, and reverse the blocking effect of the fibrin coat on NK cytotoxicity against melanoma cells.

Introduction

It is well known that metastasis of tumor cells is the main cause of mortality in patients with cancer, and this is a hot topic for investigators. Metastasis is a multi-step and highly regulated cascade process, in which tumor cells separate from a primary foci, cross the wall of vessels, circulate through the whole organism within the blood, and eventually form new colonies at remote sites following adhesion to the endothelium (1). The majority of tumor cells are killed by the immune system during circulation in the vessels (2,3), and natural killer cells (NK cells) are one of the main cellular immune factors. The anti-metastasis function of NK cells has been reported in various models of experimental and spontaneous metastasis (4,5). In these reports, it is well accepted that

direct contact between NK cells and tumor cells is crucial (6). Therefore, the mechanism by which the surviving tumor cells avoid lethal contact with NK cells is a key topic for anti-cancer research. Various studies have demonstrated that a number of blood coagulation factors facilitate the hematogenous metastasis of tumor cells (7,8). These factors contribute to tumor angiogenesis, stroma formation, growth or metastasis (9-11). Of all the known effects of cancer progression, the protective effect of fibrinogen to tumor cells has been well probed. A mass of fibrinogen may adhere to tumor cells tightly, and coat the tumor cells to form a protective shield, which inhibits the effector-target interaction and subsequently blocks NK cytotoxicity against tumor cells (12,13).

Ganoderma lucidum (*G. lucidum*) has been widely used as a traditional herb for disease treatment and health promotion, particularly by cancer patients. *G. lucidum* polysaccharides have been reported that may significantly promote the immune parameters of patients with advanced cancer (14). In their study, Sliva *et al* demonstrated that *G. lucidum* inhibits the migration of various cancer cells (15,16). The antitumor activities of the polysaccharides appear to be due to different mechanisms, such as inhibition of adhesion of tumor cells to type I collagen, hyaluronan, fibronectin and laminin, promotion of the expression of cytokines, promotion of tumor cells to induce lymphocyte proliferation and suppression of tumor-induced angiogenesis (17-20). This study was designed to observe the effect of *G. lucidum* polysaccharides on the adhesion of fibrinogen to melanoma cells and NK cytotoxicity to tumor cells. We found that fibrinogen is capable of adhering to the melanoma cells B16 and A375, and that $\alpha_5\beta_1$ and $\alpha_v\beta_3$ integrin are essential for adhesion. Coated fibrin may protect melanoma cells from NK cytotoxicity, and *G. lucidum* polysaccharides are capable of eliminating the adhesion of fibrinogen to tumor cells, and then eradicating the blocking effect of fibrinogen on NK cytotoxicity against melanoma cells and decreasing the lung metastasis of melanoma cells in mice.

Materials and methods

Reagents. Fibrinogen conjugated with Alexa Fluor 488 and IMDM culture medium was purchased from Invitrogen (OR, USA), and fibrinogen was purchased from Sigma-Aldrich (St. Louis, MO, USA). Blocking antibodies of mouse β_1 (Biolegend, CA, USA, HMB1-1) and mouse β_3 integrins (Santa Cruz Biotechnology, Santa Cruz, CA, USA, 2C9.G2), human

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Key words: *Ganoderma lucidum* polysaccharides, NK cytotoxicity, fibrinogen, integrin

$\alpha_5\beta_1$ (Chemicon, Millipore, Billerica, MA, USA, HA5) and human $\alpha_v\beta_3$ integrins (Santa Cruz Biotechnology, HM2034) were prepared according to the manufacturer's instructions. *G. lucidum* polysaccharides were purchased from Johnsun Mushroom (Hangzhou, China), and dissolved in phosphate-buffered saline (PBS).

Cells and animals. Melanoma cells B16 and A375 were obtained from the Shanghai Cell Bank (Chinese Academy of Science, Shanghai, China). Cells were cultured in IMDM medium and 10% fetal bovine serum (FBS) was added in a humidified cell incubator (37°C, 5% CO₂). These cells were harvested with trypsin and EDTA. Subsequently, PBS-washed cells were suspended in culture medium for further use. Six- to eight-week-old C57Bl/6J healthy male mice (specific pathogen free) were obtained from the Jilin University Animal Center (Changchun, China). All of the protocols applied in the animal experiments were approved by the Animal Care and Use Committee of Northeast Dianli University (Jilin, China).

Flow cytometric assay. Melanoma cells were harvested and resuspended in IMDM. For the adhesion assay, Alexa Fluor 488 conjugated fibrinogen (green fluorescence) was used. Each aliquot (0.1 ml) of tumor cells was mixed with fibrinogen (1 mg/ml) and incubated at 37°C for 30 min. After washing twice, cells were suspended in 0.5 ml of PBS for further analysis by a flow cytometer (FACScan, Beckman-Coulter, Miami, FL, USA).

Preparation of NK cells. Human NK cells were isolated from human peripheral blood mononuclear cells (PBMCs) with a negative magnetic bead NK cell isolation kit (Miltenyi Biotec, Bergisch Gladbach, Germany). In brief, B cells, T cells, stem cells, monocytes, dendritic cells, granulocytes and erythroid cells were labeled by a cocktail of magnetically biotin-conjugated antibodies. Highly pure NK cells were separated by depletion of magnetically labeled cells in the MicroBead. Mouse NK cells were separated from spleen cells with a similar kit (Miltenyi Biotec).

NK cytotoxic assay. An NK cytotoxic assay was performed in the presence or absence of fibrinogen. In brief, Na⁵¹CrO₄ pre-labeled tumor cells in 0.05 ml (1x10⁴) culture medium were pre-incubated with 0.05 ml of various concentrations of fibrinogen in the presence (0.1 mg/ml) or absence of *G. lucidum* polysaccharides for 30 min at 37°C. Subsequently, 100:1 of NK cells (0.05 ml) were added. Following 4 h of co-incubation at 37°C, cell culture plates were centrifuged at 1500 rpm for 10 min, and supernatants (0.05 ml) were collected for the radioactivity measurement. The cytotoxicity was measured using the formula % = (A - B)/(C - B) x 100, where A is radioactivity in the test well, B is spontaneous radioactivity from the well target cells without effect cells, and C is the total target released radioactivity.

Statistical analysis. Data were presented as the mean ± SD. The significance of differences between the means was calculated by the analysis of variance. When the difference of the means was shown to be significant, multiple comparisons by pairs were calculated by the T-test. Probability values of P<0.001,

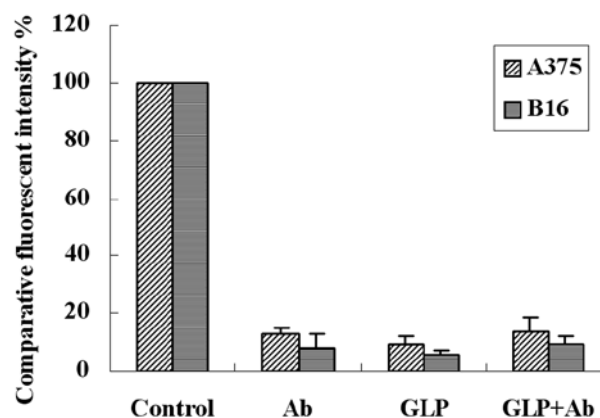


Figure 1. Fibrinogen binds to melanoma cells in an integrin-dependent manner. A total of 0.1 ml of 2x10⁵ B16 or A375 cells co-incubated with the same volume of 2 mg/ml fibrinogen conjugated with Alexa Fluor488. Function blocking antibodies of $\alpha_v\beta_3$ and $\alpha_5\beta_1$ integrins were added separately ($\alpha_v\beta_3$ and $\alpha_5\beta_1$) or together (Ab). The average fluorescent intensity was compared with the assays to which PBS had been added (control).

P<0.01 or P<0.05 were considered to indicate a statistically significant difference.

Results

Fibrinogen binds to A375 and B16 cells in a $\alpha_v\beta_3$ and $\alpha_5\beta_1$ integrin-dependent manner. Fibrinogen is known to adhere to $\alpha_v\beta_3$ and $\alpha_5\beta_1$ integrins on the surface of tumor cells. To confirm the adhesion of fibrinogen to the tumor cells we used, fibrinogen conjugated with Alexa Fluor 488 was incubated with B16 or A375 cells at 37°C for 30 min with or without sufficient blocking antibodies. The results showed that fibrinogen strongly bound to the A375 (~94%) and B16 (~98%) cell surface. Both $\alpha_v\beta_3$ and $\alpha_5\beta_1$ integrins bind to fibrinogen. In A375 and B16 cells, total eradication of the adhesion appeared when the functions of the two integrins were blocked (Fig. 1). Our results revealed that these two types of integrins mediated the adhesion of melanoma cells to fibrinogen.

Fibrinogen protects tumor cells from NK cytotoxicity. Fibrinogen may be coated on tumor cells and may protect these cells from NK cytotoxicity. Moreover, the adhesion of fibrinogen to tumor cells is essential. Melanoma cells strongly adhere to fibrinogen (Fig. 1). Therefore, we examined the ability of fibrinogen to protect melanoma cells from NK cytotoxicity. The results revealed that fibrinogen was capable of protecting tumor cells from NK cytotoxicity in a concentration-dependent manner (Fig. 2).

***G. lucidum* polysaccharides block the adhesion of fibrinogen to melanoma cells.** To observe the effects of *G. lucidum* polysaccharides, we added them into the flow cytometric adhesion system. The results revealed that *G. lucidum* polysaccharides inhibited adhesion to the level of the blocking antibodies, and the combination of *G. lucidum* polysaccharides and antibodies (GLP+Ab) had no further blocking effect (Fig. 3). The results suggest that *G. lucidum* polysaccharides eradicate melanoma cell-fibrinogen adhesion by blocking $\alpha_v\beta_3$ and $\alpha_5\beta_1$ integrins.

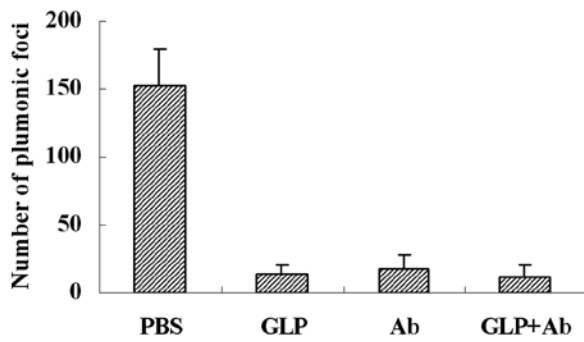


Figure 2. Fibrinogen blocks the NK cytotoxicity against melanoma cells. A total of 0.05 ml of 1×10^4 ^{51}Cr -labeled B16 or A375 cells were co-cultured with various densities of fibrinogen for 30 min. Cells were then co-incubated with 0.05 ml of 1×10^6 NK cells for 4 h to check the effect of fibrinogen on NK cytotoxicity.

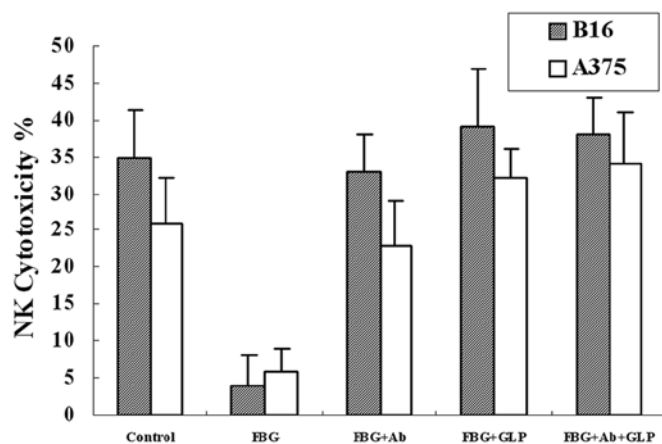


Figure 3. *G. lucidum* polysaccharides block the adhesion of fibrinogen to melanoma cells. *G. lucidum* polysaccharides (GLP) or antibodies (Ab, a combination of antibodies to $\alpha_v\beta_3$ and $\alpha_5\beta_1$ integrins) of integrins were added into the flow cytometric system prior to the addition of fluorescent fibrinogen, separately or together (GLP+Ab).

G. lucidum polysaccharides eradicates the blocking effect of fibrinogen on NK cytotoxicity against melanoma cells. Fibrinogen coats are capable of blocking NK cytotoxicity against melanoma cells, and fibrinogen adhesion mediated by $\alpha_v\beta_3$ and $\alpha_5\beta_1$ integrins are eliminated by *G. lucidum* polysaccharides. Therefore, we examined the effect of *G. lucidum* polysaccharides on NK cytotoxicity affected by fibrinogen. The results showed that *G. lucidum* polysaccharides (FBG+GLP) reduced the blocking effect to almost the level of the control (Fig. 4). The results were also similar to the samples with antibodies added (FBG+Ab), and a combination of *G. lucidum* polysaccharides and antibodies (FBG+Ab+GLP) revealed no further improvement. Our results suggested that *G. lucidum* polysaccharides are able to eliminate the blocking effect of fibrinogen on NK cytotoxicity to melanoma cells.

G. lucidum polysaccharides decrease the lung metastasis of melanoma cells in mice. NK cytotoxicity is an important immune defense for tumor metastasis, which may be blocked by fibrinogen, and *G. lucidum* polysaccharides may eliminate

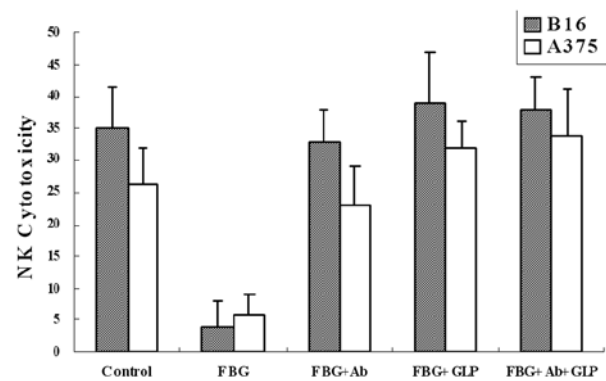


Figure 4. *G. lucidum* polysaccharides block the adhesion of fibrinogen to melanoma cells. *G. lucidum* polysaccharides (FBG+GLP) or antibodies (FBG+Ab) of integrins were added into the wells containing radio-labeled melanoma cells, separately or together (FBG+GLP+Ab). All of the wells, with the exception of the control (control) had fibrinogen added 30 min prior to injection of NK cells.

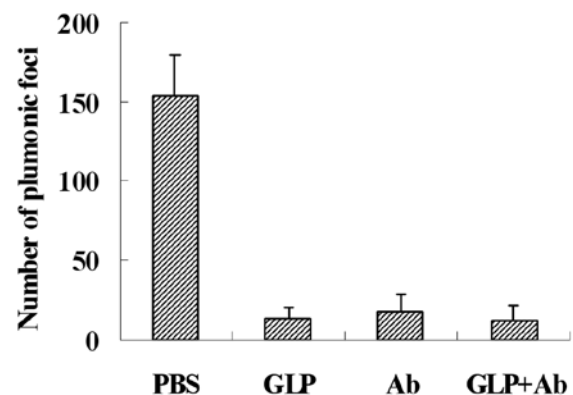


Figure 5. *G. lucidum* polysaccharides decrease the lung metastasis of B16 cells. B16 cells were injected into mice through the lateral tail vein following the injection of *G. lucidum* polysaccharides (GLP), antibodies (Ab) or both (GLP+Ab). Visible melanoma foci on the surface of murine lungs were counted 20 days following injection.

the blocking effect of fibrinogen on NK cytotoxicity against tumor cells. To detect the effect of *G. lucidum* polysaccharides on the metastasis of melanoma cells in a mouse model, we injected *G. lucidum* polysaccharides or function-blocking antibodies 5 min prior to injection of B16 tumor cells through the lateral tail vein. Our results, as shown in Fig. 5, shown that *G. lucidum* polysaccharides (GLP) and antibodies (Ab) decreased metastasis efficiently. The inhibitive rate of *G. lucidum* polysaccharides and antibodies was almost identical, and the combination of *G. lucidum* polysaccharides and antibodies (GLP+Ab) provided no further inhibition.

Discussion

It has been reported that *G. lucidum* polysaccharides is capable of eliminating the adhesion of tumor cells to various matrix proteins, including collagen, hyaluronan, fibronectin and laminin, and subsequently affecting the physiological phenomenon invoked by these proteins. Blood is rich in fibrinogen, and fibrinogen is capable of strongly blocking NK cytotoxicity.

Although investigators have reported that *G. lucidum* polysaccharides improve the NK cell profile or cytokine secretion, the effects of *G. lucidum* polysaccharides on NK cytotoxicity blocked by fibrinogen have not previously been reported. In this study, we found that *G. lucidum* polysaccharides eliminate melanoma cell-fibrinogen adhesion mediated by $\alpha_5\beta_1$ and $\alpha_3\beta_1$ integrins, and eradicate the blocking effect of fibrinogen on NK cytotoxicity against melanoma cells. We suggest that it is a new area of study of anti-cancer activity of *G. lucidum* polysaccharide.

Acknowledgements

This study was supported by grants from the National Natural Science Foundation of China (31101009), Doctoral Initiating Project of Northeast Dianli University of China (BSJXM-200905), and 'Twelfth Five' scientific and technological research projects from Jilin Provincial Department of Education (2011.74).

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