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Introduction

Gastric cancer is one of the most common malignant tumors worldwide, with 989,600 new diagnosed cases (8% of the total malignancy) and 738,000 deaths annually (10% of total cancer deaths) (Jemal et al., 2011). Over 70% of new cases and deaths occur in developing countries. The highest incidence rates are in Eastern Asia (Kamangar et al., 2006), Eastern Europe, and South America (Jemal et al., 2011). Half of all gastric cancer patients are from Eastern Asia (463,000 gastric cancer patients in China alone), and approximately two thirds of all cases occur in developing countries (Choi et al., 2014). Therefore, the development of novel approaches and effective anticancer strategies is critically needed for prolonged survival of stomach cancer.

*Rabdosia rubescens* are used in Chinese folk medicine for treatment of esophageal cancer in Taihang Mountains area of China for a long time. In 1970s, Research from Chinese researcher showed *Rabdosia rubescens* had better effect in the treatment of Gush door cancer, liver cancer and esophagus cancer, and oridonin was one of the most important antitumor active ingredient of *Rabdosia rubescens*.

Oridonin, molecular formula C_{20}H_{28}O_{6} (Figure 1), is a diterpenoid compound. Previous studies have shown that oridonin has anti-tumor activities *in vivo* and *in vitro*, but little is known about cell cycle effects of oridonin in gastric cancer.

Materials and Methods

Material

Oridonin (>98%) was purchased from National Institutes for Food and Drug Control in China. Hydroxycamptothecin (HCPT) was provided by Shanghai Longxiang Biological Medicine Development Co. Ltd. in China, Ltd. MTT (3-(4, 5-Dimethylthiazol-2-yl)-2, 5-diphenyltetrazolium bromide) was purchased from the...
United States Molecular Probe company. RPMI-1640 medium was purchased from GIBCO Company. Dimethyl sulfoxide (DMSO), trypsin, and propidium bromide (PI) were purchased from Sigma. Rabbit polyclonal anti-Cyclin B1 antibody, anti Cdk1/Cdc2 rabbit polyclonal antibody and alkaline phosphatase-labeled goat anti-rabbit IgG were provided by Beyotime institute of Biotechnology in China.

**Cell culture**

Human gastric cancer SGC-7901 cell line was obtained from Institute of Tumor Research of Harbin Medicine University (Harbin, China). Cells were cultured in RPMI 1640 medium (Gibco, 31800-022) supplemented with 10% (v/v) fetal bovine serum (Gibco, 10099-141), 100 U/mL penicillin, 100µg/mL streptomycin and 1 mM L-glutamine at 37℃ in an atmosphere of 5% CO₂. The medium was renewed two or three times/week. Cells in logarithmic growth phase were used for further experiments.

**Cell Viability Assay**

Cell viability was measured by the MTT assay, which was based on the conversion of MTT to formazan crystals by mitochondrial dehydrogenases (Mosmann, 1983). Briefly, SGC-7901 cells were plated at a density of 1×10⁴ cells/well in 96-well plate, which was in 100µL RPMI 1640 medium containing 10% (v/v) fetal bovine serum (Gibco, 10099-141), 100 U/mL penicillin, 100µg/mL streptomycin and 1 mM L-glutamine at 37℃ in an atmosphere of 5% CO₂. The medium was renewed two or three times/week. Cells in logarithmic growth phase were used for further experiments.

**Flow cytometry assay**

1 ml/well (1×10⁴ cells/well) SGC-7901 cell suspension was added into 6-well plate and cultured for 24h at 37℃ in an atmosphere of 5% CO₂. After 24h, oridonin in different concentrations were added into the wells and cultured for 48h. Then, 70% cold ethanol fixed the collected cells in

**Western blot assay**

5 ml/well (2×10⁴ cells/ml) SGC-7901 suspensions were added into 150 ml flask, cultured for 24h. Then, oridonin in different concentrations were added into the flasks and cultured for 48h. After 48h, the cells were collected, and proteins were extracted with lysis buffer [50mM Tris-Cl, pH 8.0, 120mM NaCl, 50mM NaF, 200µM sodium vanadate, 0.5% NP-40, 10 mM phenylmethylsulfonyl fluoride (PMSF), 2µg/mL aprotinin 0.2 µL, 10µg/mL Leupeptin 10µL], and then the cells were centrifuged at 12000×g for 10min at 4℃, the supernatant was saved in -80℃. The protein concentration of the supernatant was detected by Bradford assay. The protein samples were separated by SDS-PAGE and then transferred to the NC membrane with TBST containing 5% skim milk at room temperature for 2h, and the membranes were incubated with the primary antibodies overnight at 4℃. Next day, TBST solution sufficiently washed the membrane for 10
Effect of oridonin on Cdk1 and CyclinB1 protein expression

To determine whether the Cdk1 protein and CyclinB1 protein are involved in the cell cycle arrest effects of oridonin on SGC-7901 cells, Western blot assay was applied. As shown in Figure 3 and Figure 4, after 48h of the SGC-7901 cells being exposed to oridonin for different concentrations (0.375, 0.75, 1.5μmol/L), the expression levels of Cdk1 protein and Cyclin B1 protein in SGC-7901 decreased (p<0.01), compared with negative group (show in Figure 3 and Figure 4).

Discussion

Gastric cancer is one of the high incidence of tumor. Establishment of effective therapies for stomach cancer is very important for cancer research (Hu et al., 2012). Although there are a lot of therapeutic options available for patients with tumor, the efficacy is not curative. Cytotoxicity is believed as one of the major goals of cancer chemotherapy. It is thought that arresting the cell cycle of cancer is important factor in the development of cancer cell (Alabsi et al., 2012; Yang et al., 2013). Therefore, it appears that exploiting the cell cycle blocker is good idea about anti-cancer drug discovery.

A lot of antitumor drugs were derived from plants or traditional medicine, which were used to treat the disease as different folk medicine for a long time, so new drug discovery from herbs is more likely to succeed (Shin et al., 2012; Huang et al., 2013). Rabdosia rubescens has been used as an herbal remedy for various ailments including cancer in China, and oridonin is one of the anticancer components of Rabdosia rubescens. The paper was initiated with the purpose of evaluating the action of oridonin on human stomach cancer.

The experimental results show that oridonin had good cytotoxicity against the SGC-7901 cells based on the high IC_{50} value as similar as hydroxycamptothecine (Table 1). Mang documents show that oridonin could block MCF-10A, HepG2, laryngeal carcinoma and other tumor cell cycles (Hsieh et al., 2005; Kang et al., 2010; Wang et al., 2010). However, there is little report about the SGC-7901. So, this paper explored SGC-7901 from the perspective of cell cycle. The result showed that oridonin in different concentrations could block the SGC-7901 cells in G_{2}/M phase.

Cell cycle involves four sequential phases that go from quiescence (G_{0} phase) to proliferation (G_{1}, S, G_{2}, and M phases), by which all living things reproduce. In all eukaryotic cells, including human cells, cell cycling is driven by sequential activation of cyclin-dependent kinases (CDKs) and its cofactor cyclins (Díaz-Moralli et al., 2013). Among the countless elements taking part in this process, the sequential activation of heterodimeric
CDK-cyclin complexes (cyclins and their counterpart cyclin-dependent kinases (CDKs)) has been described as the key regulatory events. Cyclin-dependent kinases (CDKs) are a family of mammalian heterodimeric serine/threonine protein kinases composed of two subunits, the catalytic one known as cyclin and the regulatory one known as cyclin (Shapiro, 2006; Malumbres and Barbacid, 2007). The kinase activity of CDKs is tightly regulated by the binding to cyclins, the activating subunits which are expressed in an oscillatory way, the binding to negative regulators (CDK inhibitors, CKI) and phosphorylation/depiphosphorylation events (Manchado et al., 2012). Progression through each cell cycle phase and transition from one phase to the next are monitored by sensor mechanisms, called checkpoints, which maintain the accurate sequence of events (Hartwell and Weinert, 1989).

CDK1-cyclin B complexes are essential for initiating mitosis and can phosphorylate a broad spectrum of proteins involved in regulatory and structural processes required for mitosis such as nuclear envelope breakdown, chromosomal condensation, fragmentation of the Golgi apparatus, formation of the spindle and attachment of chromosomes to it (Malumbres and Barbacid, 2005; Mahadevan et al., 2011). Exitin mitosis requires the inactivation of CDK1-cyclin B, which is carried out by the ubiquitin-dependent proteolysis of B-type cyclins by the APC/C-CDC20 complex (Harper et al., 2002; Peters, 2002). So, cell cycle proteins (cyclinsB1) and cyclin-dependent protein kinase (Cdk1) were closely related with G2/M phase. CyclinB1, as the key factor to switch on mitosis, could compose compound with Cdk1 to adjust the G2/M phase.

Our study showed that oridonin arrested the SGC-7901 cell cycle in G2/M phase. To explore the relation of G2/M phase arrestion to CDK1 and cyclin B1, we tested the effect of oridonin on Cdk1 and Cyclin B1 proteins. The results showed that oridonin could significantly decreased the expression of Cdk1 and Cyclin B1 proteins in SGC-7901 cells, which would led to a significant reduction in the formation of CDK1-cyclin B complexes, and then block the cells in G2/M phase.

The above analysis showed that oridonin blocked SGC-7901 cells in G2/M phase by down-regulating the protein expressions of CDK1, CyclinB1.

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References


