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Cucurbitacin B inhibits growth, arrests the cell cycle, and potentiates antiproliferative efficacy of cisplatin in cutaneous squamous cell carcinoma cell lines

WEIKAI CHEN^{1,2}, AMANDA LEITER², DONG YIN², MURIEL MEIRING¹, VERNON J. LOUW⁴, and H. PHILLIP KOEFFLER^{2,3}

¹Department of Hematology and Cell Biology, Faculty of Health Sciences, University of the Free State, Bloemfontein 9301, Free State, South Africa

²Division of Hematology and Oncology, Cedars-Sinai Medical Center, University of California-Los Angeles School of Medicine, Los Angeles, CA 90048, USA

³National Cancer Institute of Singapore and CSI, Singapore

⁴Division of Clinical Haematology, Department of Internal Medicine, Faculty of Health Sciences, University of the Free State, Bloemfontein 9301, Free State, South Africa

Abstract

Cutaneous squamous cell carcinoma (CSCC) is the second most common skin cancer with a substantial risk of metastasis which causes clinical treatment failure. This study investigated the anti-CSCC effects of a triterpenoid compound, Cucurbitacin B (CuB). Dose-response studies showed that CuB inhibited 50% growth (ED₅₀) of the CSCC cell lines (SRB1, SRB12, SCC13, COLO16) in liquid culture at 4×10^{-7} – 10^{-5} M. Soft-agar assays demonstrated that nearly all of the CSCC clonogenic cells were inhibited at 10^{-7} M CuB. FACS analysis found that the compound (10^{-7} M, 48 h) caused G2/M arrest. The CSCC cells underwent profound morphologic changes within 60 min after exposure to CuB (10^{-7} M), rounding up and losing their pseudopodia. CuB (10^{-7} M) caused prominent multinucleation of the cells after they were pulse-exposed (24 h) to the drug, washed and cultured in normal medium for an additional 24 h. The drug (10^{-8} – 10^{-6} M, 3–24 h) decreased levels of CDC2 and cyclin B1 in SRB1 and SRB12 cell lines as seen by Western blot analysis. Migration of SRB1 and SRB12 cells was inhibited by 10^{-7} M CuB. Interestingly, CuB synergistically potentiated the anti-proliferative effect of cisplatin in CSCC. In summary, CuB has a prominent anti-proliferative activity on CSCC cells. *In vivo* studies and clinical trials of this drug should be pursued in CSCC.

Keywords

cucurbitacin B; cell cycle; cisplatin; cutaneous squamous cell carcinoma

Introduction

CSCC is the second most common skin cancer representing 20% of all cutaneous malignancies (1-3). In addition, the incidence of CSCC appears to be rising (3). CSCC tends to locate on the skin of the head and neck (4) and has a substantial risk of metastasis to local lymph nodes and to distant organs (3,5). For those patients with metastatic disease as well as those with a poor prognosis (5), systemic chemotherapy is the standard treatment (5,6). The platinum compound cisplatin is regarded as a standard chemotherapeutic agent used either alone or in combination with other agents against CSCC (7). However, cisplatin is associated with a variety of side-effects, including the relapse of most cancers after therapy and eventual resistance to cisplatin. Therefore, new effective anti-CSCC therapeutic regimens are needed.

Natural herbal medicines have been used as anticancer agents (8). Cucurbitacins are a group of highly structurally diverse tetracyclic triterpenes isolated from numerous plant families such as the Cucurbitaceae and Cruciferae, which possess a wide spectrum of pharmacological effects such as cytotoxicity, chemoprevention, antitumor properties, hepatoprotective, analgesic, anti-inflammatory, antimicrobial, anti-helminthic, as well as procardiovascular activities (8,9). This family of compounds has been used as folk medicines for centuries in Asia, especially China and India (9,10). Chemically, cucurbitacins are highly diverse compounds and are arbitrarily divided into twelve categories (cucurbitacins A-T) (11). Of these compounds, CuB is one of the most abundant and has been most widely used (10,12). Recent studies have suggested that CuB inhibits the growth of numerous human cancer cell lines and tumor xenografts, including leukemias, lymphomas, multiple myelomas, melanomas and cancers of oral cavity, nasopharynx, breast, prostate, lung, uterine cervix, liver, and brain (10,11,13). The anti-tumor mechanism of CuB involves a variety of effects including induction of apoptosis, and/or differentiation, arrest of the cell cycle, disruption of the cytoskeleton as well as interruption of several signal transduction pathways (14-17). However, the effect of CuB and its underlying mechanism of action against CSCC are unexplored.

In the present study, we evaluated the action of CuB on CSCC cell lines. CuB significantly inhibited the proliferation as well as metastatic potential of CSCC cell lines. CuB caused a marked G2-M phase arrest and increased apoptosis. It also produced a rapid alteration of cell morphology including rounding of the outer cell membrane and prominent multinucleation. Moreover, the compound significantly potentiated the anti-proliferative effects of cisplatin against CSCC cells.

Materials and methods

CSCC cell lines, cell proliferation and migration

Human CSCC cell lines SRB1, SRB12, SCC13, COLO16 (kindly provided by Dr Reuben Lotan, MD Anderson Cancer Center) were maintained in Dulbecco's modified Eagle's medium (Gibco-BRL) with 10% fetal calf serum (Gemini Bio-Products, Calabasas, CA), 10 U/ml penicillin-G, and 10 mg/ml streptomycin (Gemini Bio-Products). The cells were incubated at 37°C in 5% CO₂. For pulse-exposure experiments, cells were exposed to CuB

(10^{-7} M) for either 2, 9 or 24 h, washed extensively with culture medium and grown in CuB-free medium for 1, 2, 3 days. For clonogenic assays, cells were plated into 24-well flat-bottomed plates using a two-layer soft agar system with a total of 3×10^3 cells/well in a volume of 400 μ l/well, as described previously (15). After 14 days of incubation, the colonies were counted. For proliferation measurements, the cells were placed into 96-well plates, and cell growth was measured at various times by 3-(4,5-Dimethyl-thiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay according to the protocol provided by Roche Molecular Biochemicals (Basel, Switzerland). For cell migration assays, SRB1 and SRB12 cell lines were seeded into 12-well flat-bottomed plates until 80% confluence. Distance of migration was measured microscopically, one half of the dish was physically cleared of cells, the leading edge of cells was marked microscopically, fresh medium was added and incubated either with or without CuB for 24 h, then the distance of migration was measured on the bottom of the plate.

Chemicals

CuB was provided by CK Life Sciences Ltd. (Hong Kong) and was dissolved in DMSO at a stock concentration of 10^{-2} M and stored at -20°C . Fresh dilutions in medium were made for each experiment.

Assessment of apoptosis

To study induction of apoptosis, Annexin V assays (Annexin V-FITC Apoptosis Detection kit; Pharmingen, San Diego, CA) were performed according to the manufacturer's instructions. Briefly, cells were harvested after exposure to reagents, washed twice with PBS, incubated with FITC-conjugated annexin V and Propidium Iodide (PI) for 15 min; staining was measured by FACScan (Becton-Dickinson).

Cell cycle analysis

CSCC cells were placed in 6-well dishes and cultured for 24 h with CuB. After trypsinization and washing twice with ice-cold phosphate-buffered saline (PBS), cells were fixed in 70% ice-cold ethanol overnight. The samples were treated by RNAase A and exposed to propidium iodide (PI), then analyzed by flow cytometry (FACScan, Becton-Dickinson).

Western blot analysis

Cells were harvested for total cell lysates with RIPA buffer (1% Nonidet P-40, 0.5% sodium deoxycholate, 0.1% SDS, 50 mM Tris-HCl, pH 7.5) containing a mixture of protease inhibitors (Roche Diagnostics GmbH, Mannheim, Germany), as well as 1 mM NaF and 1 mM NaVO_4 . Cell lysates were centrifuged at 13,000 rpm for 10 min at 4°C . Supernatants were collected, and the concentration of proteins was measured. The proteins in the lysates (30 μ g) were denatured in the sample buffer (10% glycerol, 5% β -mercaptoethanol, 2.3% SDS, 62.5 mM Tris-HCl, pH 6.8) by boiling and then subjected to 4-15% SDS-PAGE followed by electro-transfer to polyvinylidene difluoride membrane. The immune complexes were visualized with Supersignal west dura extended duration substrate (Pierce, Rockford,

IL) and normalized by an internal control (GAPDH). The antibodies were purchased from Santa Cruz Biotechnology, CA.

Statistical analysis

Differences between the results of experimental treatments were evaluated by the t-test. Differences were considered significant at values of $p < 0.05$.

Results

CuB inhibits proliferation in liquid and soft gel culture of CSCC cells as well as decreases their migration

We tested the ability of CuB to inhibit the proliferation of CSCC cell lines (SRB1 and SRB12) in liquid culture using the MTT assay (Fig. 1A). The SRB12 cells were the most sensitive to CuB with an ED_{50} of about 4×10^{-7} M at 72 h of CuB exposure. Two other CSCC cell lines (SCC13 and COLO16) were less sensitive to this compound with an ED_{50} about 10-5 M CuB at 72 h exposure (data not shown).

Clonogenic assay depends on individual colony-forming cells proliferating actively in soft gel culture to form a colony. Clonal growth of CSCC was inhibited significantly by the drug in a dose-dependent manner compared with the control cells. At 10^{-7} M CuB, nearly all of the clonogenic SRB1 cells were inhibited (Fig. 1B, data not shown). Furthermore, migration of SRB12 CSCC cells cultured with CuB (2.5×10^{-7} M, 24 h) was significantly decreased compared with diluent-treated control cells (Fig. 1C).

CuB rapidly alters cell morphology, produces cell cycle arrest, and increases apoptosis of CSCC cells

The CSCC cell lines (SRB1 and SRB12) underwent rapid morphologic changes within 15-30 min after exposure to CuB (5×10^{-7} M), rounding up and losing their pseudopodia; by 120 min, the cells were very irregular and refractile (Fig. 2A).

The ability of CuB to induce apoptosis in a time- and dose-dependent manner was measured by Annexin V-FITC and propidium iodide double labeling assay. A total of 9% (SRB12), 16% (SCC13), and 26% (COLO16) of the CSCC cells were apoptotic after a 48 h of drug exposure (10^{-6} M), whereas less than 2.5% of the diluent treated cells had undergone apoptosis (Fig. 2B).

The effect of CuB on the cell cycle was studied by flow cytometry (Fig. 2C). The compound caused a time- and dose-dependent G2-M phase arrest of the cells (SRB12, SCC13 and COLO16). After 48 h of drug exposure (10^{-6} M), the percentage of cells in the G2-M phase increased by 5-10 fold compared with the controls. We further examined the expression of several proteins involved in cell cycle regulation. The protein levels of CDC2 and cyclin B1 decreased remarkably in a time- and dose-dependent fashion (Fig. 2D).

Influence of pulse-exposure to CuB on cell proliferation and cell morphology of CSCC cells

We examined the proliferative activity of CSCC cell lines after they were pulse-exposed to CuB (10^{-7} M) for either 2, 9, or 24 h, washed extensively, and grown in liquid culture in the absence of CuB for an additional 24, 48 or 72 h (Fig. 3A). A significant decrease in proliferation of the CSCC cells (SRB12) occurred after exposed for 24 h to CuB compared with diluent-exposed control cells (Fig. 3A; $p < 0.01$). Only a slight decrease in proliferation of these cells was noted when exposed for either 2 or 9 h.

In additional experiments, SRB12 cells were exposed to either CuB (2.5×10^{-7} M) or diluents for 24 h, washed and recultured in drug-free media for another 24 h (Fig. 3B). CuB caused prominent multinucleation of the cells. After 24 h pulse-exposure to CuB [SRB1 (10^{-6} M), SRB12 (10^{-7} M)] followed by extensive washing and 24 h of culture in CuB-free medium, a mean $23 \pm 3\%$ SRB1 and $11 \pm 5\%$ SRB12 cells developed marked polyploidy (predominantly tetraploidy) compared with $2 \pm 2\%$ SRB1 and $4 \pm 3\%$ SRB12 untreated control cells (Fig. 3C).

CuB synergistically potentiates the anti-proliferative effects of cisplatin

The growth of SRB1 and SRB12 cells treated with CuB (10^{-7} M and 2.5×10^{-7} M), cisplatin (5, 10 and 20 μ M), or a combination of both was determined by MTT assay (Fig. 4A and B). A significant reduction in growth was observed in the cell lines (SRB1 and SRB12) treated with a combination of CuB (2.5×10^{-7} M) and cisplatin (10 and/or 20 μ M) ($p < 0.05$).

Furthermore, the morphology of SRB12 cells was examined under optical microscopy after 36 h exposure to CuB (2.5×10^{-7} M), cisplatin (10 μ M), or a combination of both (Fig. 4C). No prominent alterations in cell morphology were observed in cells exposed to either CuB or cisplatin alone compared with diluent control cells. However, after treatment with both CuB and cisplatin at these concentrations, the cells lost their pseudopodia and became refractile (apoptotic appearing) (Fig. 4C).

Discussion

CuB is extracted from *Trichosanthes kirilowii* Maximowicz (Cucurbitaceae family), the fruits and roots of which have long been utilized in traditional Asian medicine for their anti-inflammatory, anti-diabetic and abortifacient effects (15). In modern medicine, its anti-proliferative potential is of particular interest, which have been studied in leukemia and a variety of solid tumors (10,12,14-16,18). In this study, we examined for the first time the anti-proliferative effect of CuB against CSCC cell lines. Sensitivity to the anti-proliferative activity of CuB differed largely among the cell lines (ED₅₀ at 72 h: averaged between 4×10^{-7} M and 10^{-5} M). The clonogenic assay also confirmed the growth inhibitory effect of CuB using a soft agar assay.

To elucidate the mechanism by which CuB inhibited cell growth of CSCC, Annexin V assay and cell cycle analyses were conducted. A moderate increase in apoptosis (up to 24%) occurred after exposure to 10^{-6} M CuB for 48 h. Flow cytometry analysis revealed that CuB caused significant accumulation of CSCC cells at the G₂-M phase with a decrease of cells in S-phase. CuB also caused a G₂-M cell cycle arrest in laryngeal squamous cell carcinoma

cells, glioblastoma multiforme cells, and pancreatic cancer cells (12,14,15). In contrast, CuB induced a S phase arrest of leukemia cells and hepatocellular carcinoma cells (10,16). The explanation for these differences is unclear. Western blot analysis showed that cell cycle regulators, CDC2 and cyclin B1, were down-regulated in a dose- and time-dependent manner in CSCC cells. These proteins are responsible for the G2-M phase transition. Our findings paralleled those of Thoennesses *et al* (14) showing that CuB decreased expression of Cyclin B1 and arrested the cell cycle in the G2-M phase in pancreatic cancer cells.

Previous studies from our group demonstrated CuB can disrupt F-actins and microtubules in glioma, leukemia, as well as breast cancer cells (15,16,18). Given that the F-actin and the microtubule cytoskeleton is a key component involved in cell migration, we tested the ability of CuB to affect cell migration and found that CuB inhibited migration of CSCC cells. In addition, a short exposure (<60 min) to CuB made the CSCC cells become round and refractile (Fig. 2A). These CSCC cells could recover to their normal morphology, if after 60 min of CuB exposure the cells were washed free of the drug and cultured for an additional 24 h. This is similar to morphologic changes noted in glioma, pancreatic and breast cancer cells (14,15,18).

Our pulse-exposure experiments showed that a 24-h exposure to CuB (10^{-7} M) largely inhibited growth of CSCC cells, while a shorter (2 or 9 h) exposure did not produce much growth inhibition. Additionally, CSCC cells which were pulse-exposed to CuB developed prominent multinucleation, a sign of failed mitosis. This multinucleation phenomenon has also been observed in glioma and pancreatic cancer cells after pulse-exposure to CuB (14,15).

Metastatic CSCC is poorly treated by conventional therapies including cisplatin. Our *in vitro* studies showed that CuB (2.5×10^{-7} M) could potentiate the cell killing mediated by cisplatin (10-20 μ M). Further *in vivo* studies are warranted to extend these findings.

In summary, the present study showed for the first time that CuB has profound anti-CSCC activity *in vitro*. The agent inhibits cellular growth and migration as well as arrests the cell cycle at the G2-M phase in concert with decreasing the expression of CDC2 and cyclin B1. This compound also produces multinucleation and potentiates the anti-proliferative activity of cisplatin. Our studies provide a rationale for the development of CuB as a therapeutic agent against human CSCC.

Acknowledgments

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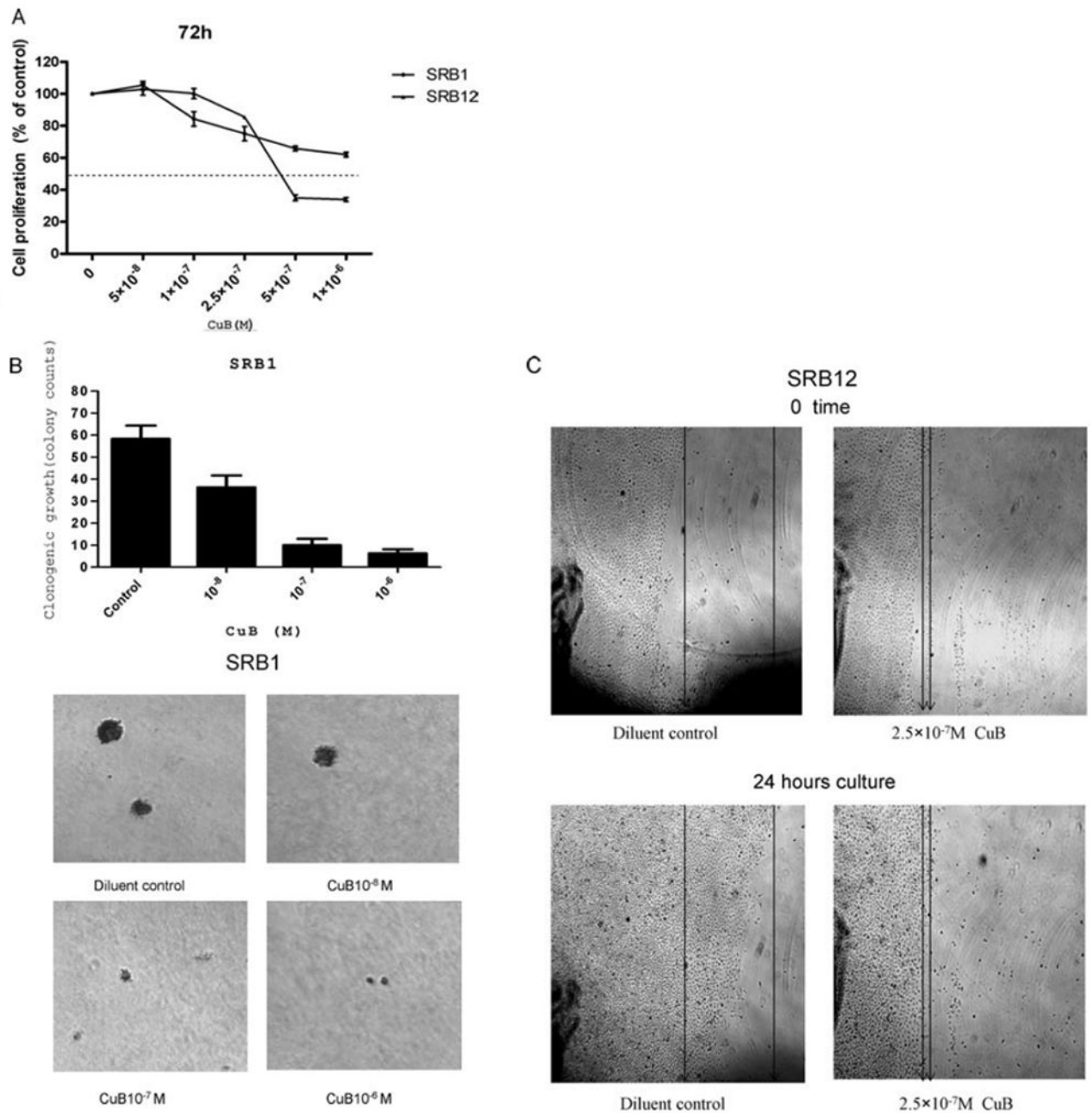


Figure 1.

Effect of CuB on proliferation, clonal growth and migration of CSCC cells. (A) SRB1 and SRB12 CSCC cell lines were cultured in the presence of various concentrations (5×10^{-8} - 1×10^{-6} M) of CuB for 72 h. Cell proliferation was measured by MTT assay. Results are expressed as a mean percentage of MTT activity present in the diluent only control wells. Each point represents a mean \pm standard deviation (SD) of 3 independent experiments done in triplicate. (B) CSCC cell line SRB1 was cultured in 2 layer soft agar either without or with CuB (10^{-8} - 10^{-6} M), and colonies were counted after 14 days. Photographs show representative results of the colony assay. (C) Migration assay: SRB12 CSCC cells were

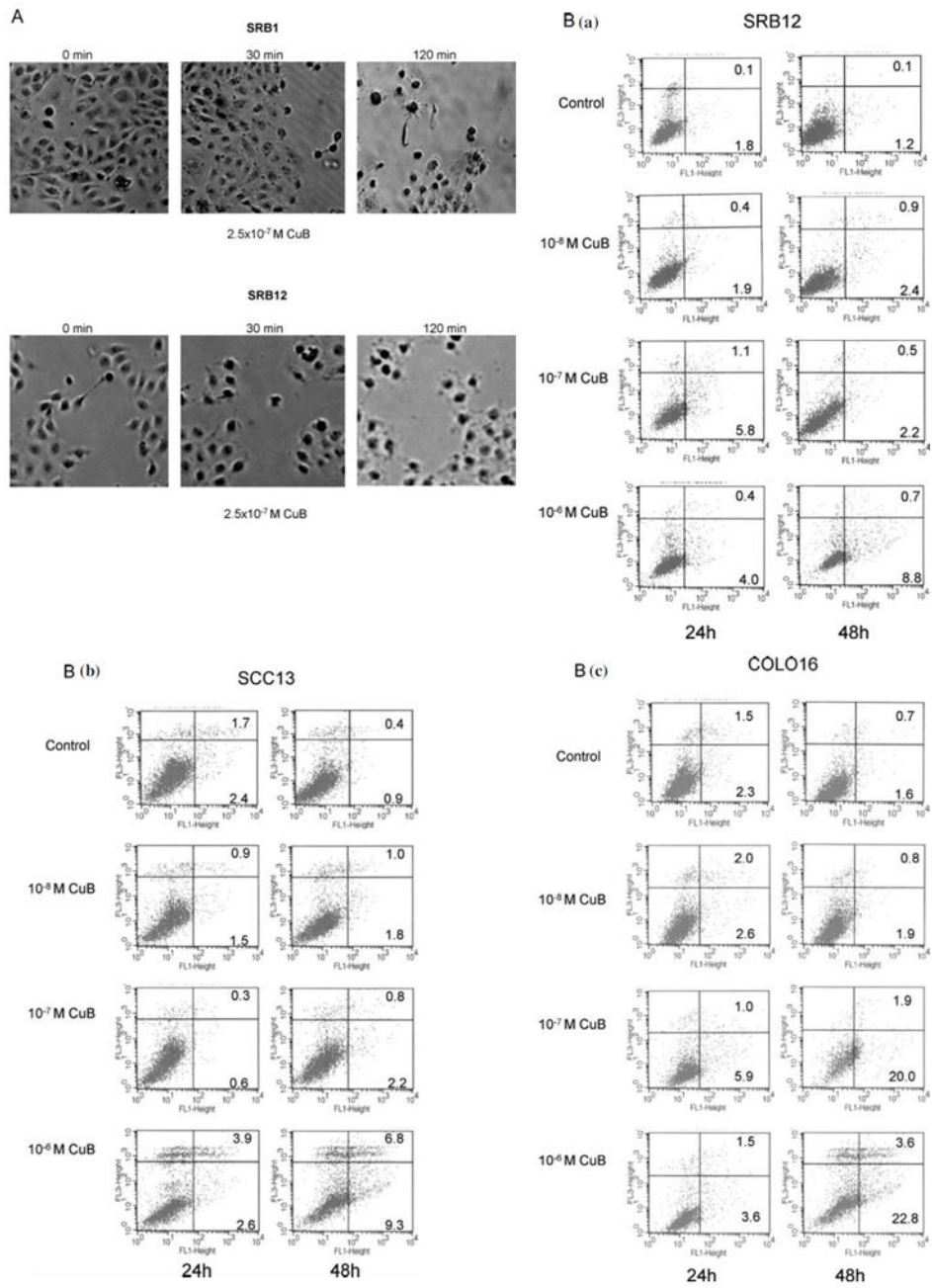
cultured either without or with CuB (2.5×10^{-7} M) for 24 h, and cellular migration was measured. Photographs were taken at the same position and magnification with marks at 0 and 24 h (representative results shown for each cell line).

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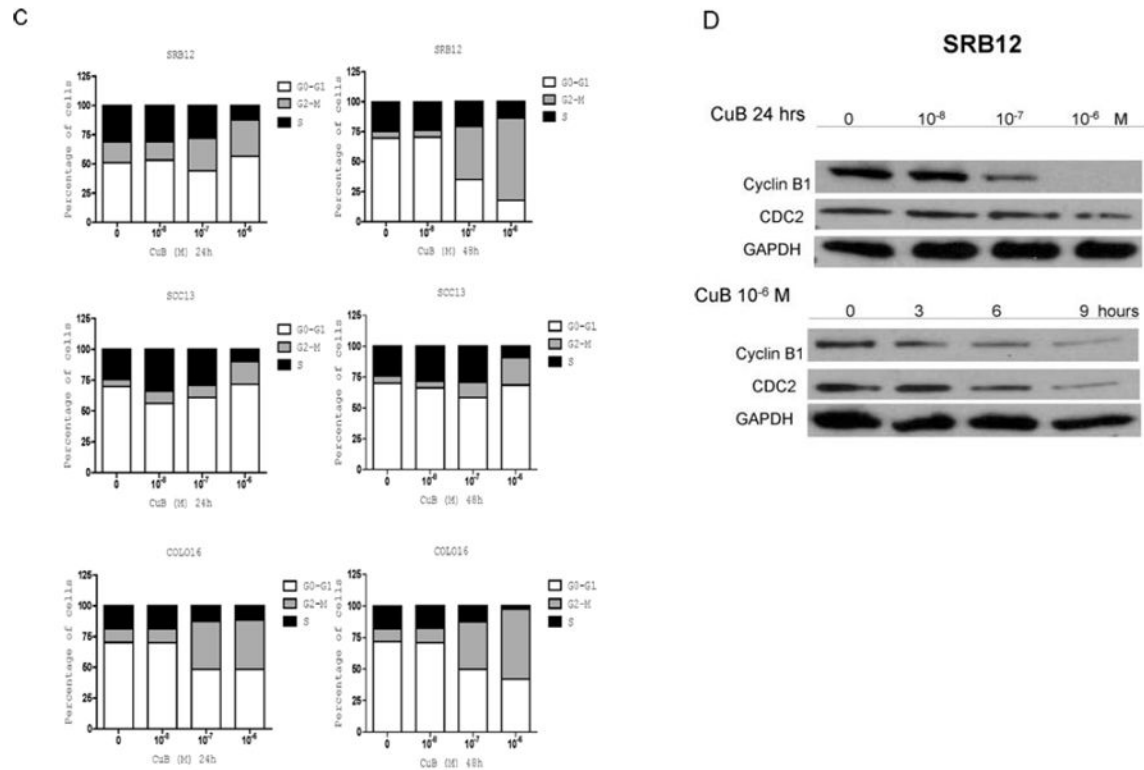
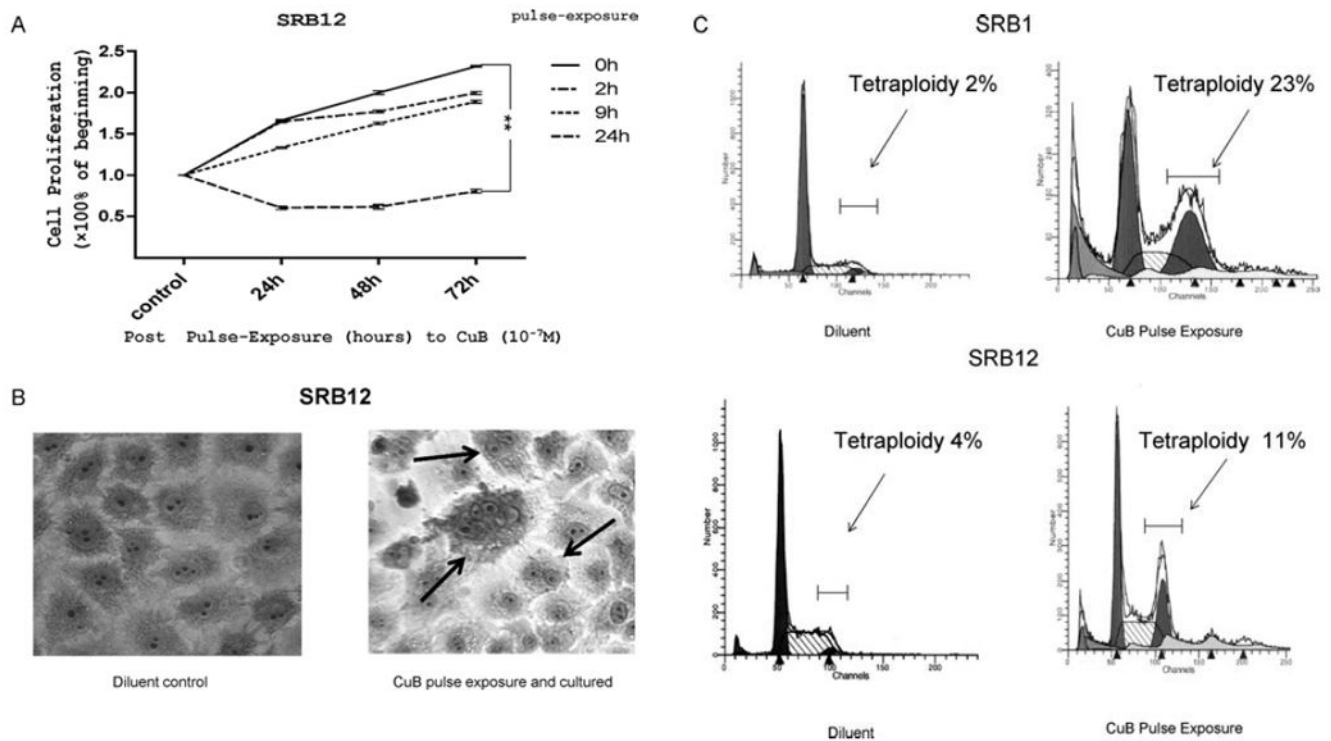


Figure 2.

CuB alters cell morphology, arrests cells in G2-M cell cycle, and induces apoptosis. (A) After exposed to CuB (5×10^{-7} M for either 0, 30 or 120 min), morphology of SRB1 and SRB12 CSCC cells were examined. (B) CSCC cells (SRB12, SCC13 and COLO16) were cultured either without or with CuB (10^{-8} - 10^{-6} M). After 24 and 48 h, apoptosis was measured by Annexin V/PI assay. Representative results are shown (three experiments were performed, each with similar results). (C) Cell cycle distribution of CSCC cells (SRB12, SCC13 and COLO16) was analyzed after culturing for 24 and 48 h either without or with CuB (10^{-8} - 10^{-6} M). Cell cycle was assessed by propidium iodide staining and FACS analysis. For each sample, the percentage of cells in G0/G1, S or G2/M phase of the cell cycle is indicated. Representative result is shown (two additional experiments gave similar results). (D) Western blot: SRB12 cells were exposed to either CuB (10^{-8} - 10^{-6} M) for 24 h (upper panel) or to 10^{-6} M CuB for 3, 6, and 9 h (lower panel), lysates were made and Western blot analysis was performed using antibodies against cyclin B1, cdc2 and GAPDH (loading control).

**Figure 3.**

Influence of pulse-exposure to CuB on growth and cell morphology of CSCC cells. (A) SRB12 cells were exposed to CuB (10^{-7} M) for 2, 9 or 24 h, washed extensively, cultured in CuB-free culture medium for 24, 48 and 72 h, and cell growth was measured by MTT assay. Each point represents a mean \pm SD of triplicate wells. (B) CuB causes multinucleation: SRB12 cells were treated with CuB (2.5×10^{-7} M for 2 days), washed and cultured an additional 2 days with no CuB, multinucleation developed, see arrows. (C) SRB1 or SRB12 cells were cultured with CuB (10^{-6} M or 10^{-7} M for 24 h), washed extensively and cultured in regular media for an additional 24 h and examined for DNA content per cell by FACS after staining with propidium iodide. Cells were gated using FL2-W versus FL2-A bivariate graphs.

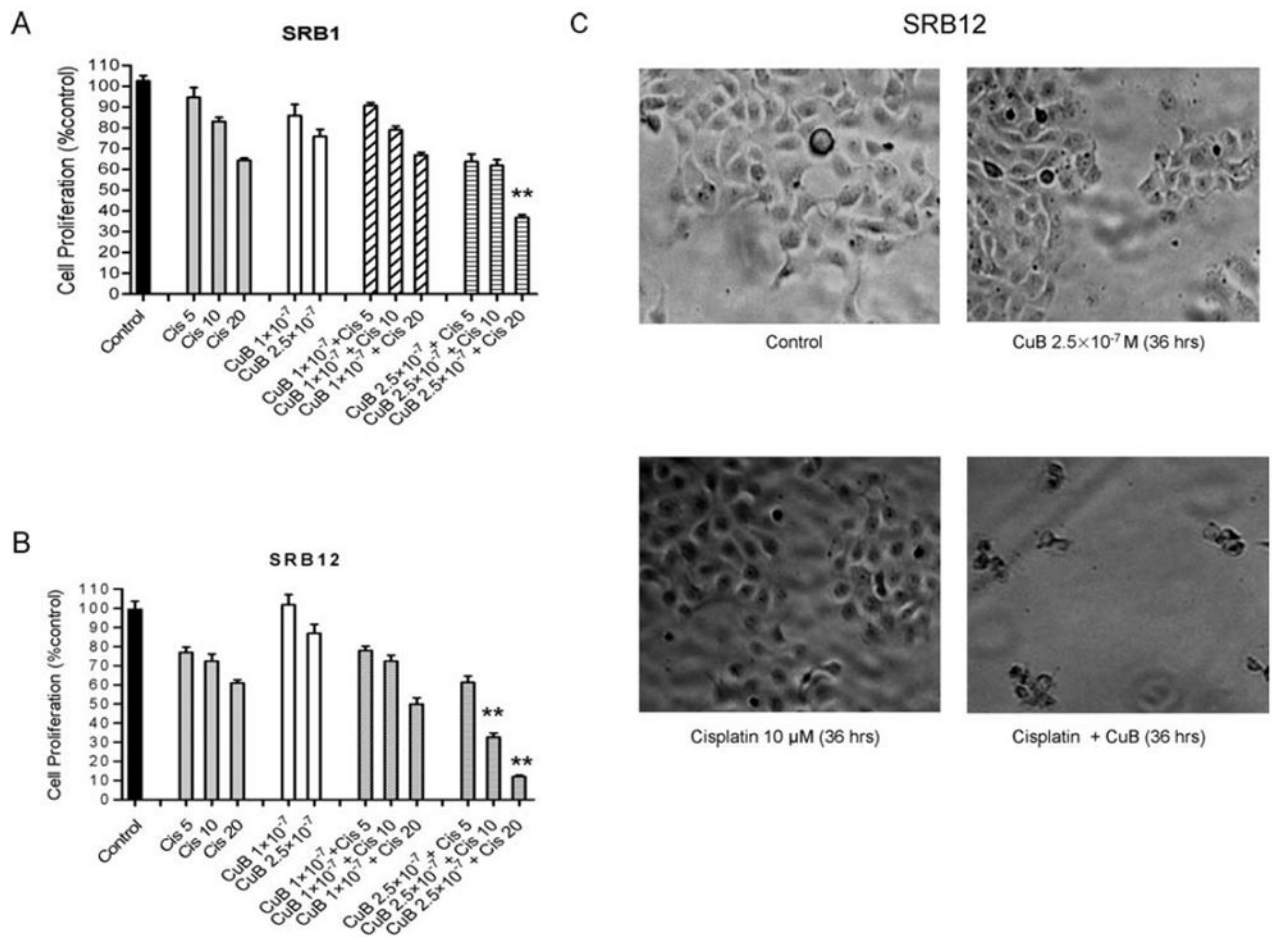


Figure 4.

CuB synergistically potentiates the anti-proliferative effects of cisplatin. (A and B) Proliferation of SRB1 and SRB12 cells cultured with either CuB (10^{-7} M or 2.5×10^{-7} M), cisplatin [(cis), 5, 10 and 20 μ M], or a combination of both was determined by MTT assay. ** $p < 0.01$ (C) SRB12 cells were cultured either without or with CuB (2.5×10^{-7} M) and/or cisplatin (10 μ M) for 36 h and examined by light microscopy ($\times 40$).