Survival advantages of multicellular spheroids vs. monolayers of HepG2 cells in vitro

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Abstract. Mammalian cells grow in three-dimensions (3-D) in vivo. Commonly used two-dimensional (2-D) cell cultures are inadequate to recreate the biological microenvironment of tumor cells. The potentially different outcomes from 2-D and 3-D culture systems may have a significant impact on the relevance of experimental findings. The purpose of this study was to characterize the human hepatoma cell line HepG2 in 2-D and 3-D cultures. HepG2 cells in 2-D and 3-D cultures were treated with cisplatin, 5-fluorouracil, and adriamycin and were analyzed by scanning electron microscopy and transmission electron microscopy. Cell cycle progression and apoptosis were detected by flow cytometry. Inhibition of cell proliferation was quantified by MTT assay. The expression of E-cadherin, CD44v6, VEGF, KDR, endostatin, Bax, and cytochrome-c were analyzed by immunohistochemical (IHC) staining. As compared to the 2-D monolayer culture, the 3-D multicellular spheroids (MCS) of HepG2 cells featured a greater fraction of cells in G1 phase and were organized with more abundant cell-cell adhesion. In addition, cells in MCS were significantly less apoptotic in maintenance culture media and were more resistant to drug-induced apoptosis. E-cadherin, CD44v6, VEGF, KDR, endostatin, and cytochrome-c levels were increased in MCS as compared to 2-D cell cultures. In conclusion, MCS conferred differentiated phenotypes including increased cell-cell adhesion and G1 phase cell cycle arrest, enhanced cellular resistance to apoptosis, and upregulated angiogenic potential. Based on our data, a multicellular morphological hierarchy may sustain the growth/survival advantages of cancer cells in vivo. Therefore, a 3-D culture system should be the preferred technique for cancer biology investigation.

Introduction

Hepatocellular carcinoma (HCC) is the most common malignancy of the liver. It is the third most common cause of cancer mortality globally, and thus a major health concern worldwide. Therapeutic options for HCC include surgical resection, local ablative therapies, systemic treatment, and liver transplantation (1). Surgery, being the most effective treatment, is unfortunately not applicable to the majority of HCC patients since the diagnosis is most often made at unresectable stages. Even for patients eligible for hepatic resection, the postoperative recurrence rate is as high as 50% two years after surgery. As a result, the median life span of HCC patients after diagnosis is <6 months (2). Chemotherapy is an area that has yet to be explored as an alternative to improve the prognosis of HCC. Unfortunately, current chemotherapies for HCC generally have low response rates (3-6). It is important to note that cytotoxicity in established cancer cell lines cultured in a two-dimensional (2-D) monolayer is a standard first line drug screening criterion. However, as suggested by clinical observations, HCC, in their primary and disseminated states, tend to be multicellular. More importantly, tumor cells always grow in a three-dimensional (3-D) space in vivo. Thus, the commonly used 2-D cell culture system is probably inadequate to recreate the biological microenvironment of tumor cells (7). The potentially different outcomes from 2-D and 3-D culture systems may have a significant impact on the relevance of experimental findings.

In light of this, 3-D culture has gained significant approval as a more advantageous in vitro system than 2-D culture, for studying various molecular processes and screening therapeutic agents. In the 3-D culture model, many established human cancer cell lines form multicellular spheroids (MCS) which mimic solid tumors more closely than the monolayer cells in 2-D cultures (8). The 3-D system seems to be particularly valuable in the assessment of tumor cell drug...
sensitivity since MCS manifest elevated resistance to chemotherapeutic agents compared with monolayer cell cultures. The increased drug resistance of MCS may be a result of increased cell-cell interactions and polarized cell-matrix interactions, which may not be supported by the monolayer cell culture system. It is well documented that molecules that regulate cell-cell and cell-matrix interactions, including E-cadherin, catenins, CD44 family members and integrin, play critical roles in cell proliferation, differentiation and apoptosis (9). In addition to the histological organization and the intrinsic characteristics of tumor cells, tumor angiogenesis also regulates drug resistance. To this end, tumor cells are considered to be the major contributor of the angiogenic growth factor, vascular endotheliosis growth factor (VEGF). It is intriguing to speculate that MCS may provide a convenient in vitro model to screen and characterize anti-cancer drugs and to study how tumor cells may contribute to angiogenic responses in their natural environment. These possibilities are addressed in the current study.

Materials and methods

2-D and 3-D cell culture. The human hepatoma cell line HepG2 (ATCC, USA) was maintained in a 2-D monolayer culture in RPMI-1640 supplemented with 10% fetal bovine serum (Gibco, USA). MCS of HepG2 cells were obtained by a liquid overlay technique as described previously (10). Briefly, for 2-D cultures, a HepG2 single cell suspension in complete media was seeded at 2x10^5 cells/ml in each culture flask. The conditions for 3-D cell culture were the same as for monolayer cultures, except that the culture flasks were coated with 2% agarose before cell plating. Fifty percent of the culture medium was renewed every two days.

Scanning electron microscopy (SEM) and transmission electron microscopy (TEM) studies of 2-D and 3-D cell organization. The monolayer cells and MCS were washed with PBS and fixed in 2.5% glutaraldehyde for 2 h. The cells were then post-fixed on the plate with 1% OsO4, and dehydrated by graded ethanol. The cells were then covered with gold palladium and examined by SEM (Hitachi, H-600, Japan), or embedded in Epon812 expoy resin and examined by TEM (Hitachi, H-600, Japan).

MTT cell viability assay. The cytotoxicity of drugs was measured by 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide (MTT) assay using a previously described method (11). Briefly, 1500 cells in either a monolayer or MCS in 150 μl of maintenance medium were seeded into a 96-well plate. Next, the cells were treated with 50 μl of cisplatin (CDPP) (Qilu Pharmaceutical Company, Shandong, P.R. China), or 5-fluorouracil (5-FU) (Xudonghaipu Pharmaceutical Company, Shanghai, P.R. China), or adriamycin (ADM) (Haizheng Pharmaceutical Company, Zhejiang Province, P.R. China) for 48 h at the indicated concentrations. The cells were then incubated with 20 μl of MTT reagent (5 mg/ml, Sigma, USA) at 37°C for 4 h. The media was collected and measured at the spectrometric absorbance of 492 nm, yielding A_{492_{nm}} values, using a Multi-function microplate reader (POLARstarOPTIMA, BMG, Germany). The inhibition of cell viability was calculated by the formula (1-A_{492_{nm}}/A_{492_{nm}}) x100%, where A_{492_{nm}} is from the treated cells and A_{492_{nm}} is from the untreated cells. All sample measurements were replicated 5 times.

Cell sorting by flow cytometry. Monolayer cells and MCS were harvested, suspended, and fixed in 70% alcohol at 4°C overnight. The cells were then stained by propidium iodide (PI, 50 μg/ml) at 37°C in the dark for 30 min and analyzed by flow cytometer (FACScan, BD, USA). In parallel, cells harvested from monolayer or MCS cultures were stained with PI and assayed for Annexin V. Briefly, cells were resuspended in a 200 μl solution containing FITC-conjugated Annexin V antibody (Beijing Zhongshan Golden Bridge Company, P.R. China) and PI (50 μg/ml) for 15 min and analyzed by flow cytometry. The percentage of Annexin V-positive/PL-negative apoptotic cell populations was calculated using the Cell Quest software.

IHC. HepG2 cells were plated on glass coverslips in a monolayer and fixed with 4% polyoxymethylene. The MCS were fixed with 4% polyoxymethylene and embedded in paraffin. Blocks were sectioned at a thickness of 4 μm, deparaffinized and rehydrated in graded alcohol. After quenching endogenous peroxidase with 3% H2O2 in methanol for 30 min, the sections were incubated overnight at 4°C with mouse monoclonal antibodies against human E-cadherin (1:200 dilution), CD44v6 (1:200 dilution) (Maxin Biotechnology Co.), VEGF (1:200 dilution), VEGFR-2 (1:200 dilution), endostatin (1:200 dilution), Bax (1:200 dilution), or cytochrome-c (1:200 dilution). The antibodies against E-cadherin and CD44v6 were from P.R. China. All other primary antibodies were from Santa Cruz Biotechnology Inc, (Santa Cruz, CA, USA). After the slides were blocked, the sections were incubated with the anti-mouse secondary antibody for 40 min at 37°C. The bound secondary antibody was visualized by the activity of the horse radish peroxidase conjugate using a 3,3-diaminobenzidine tetrahydrochloride substrate (Sigma, UK). The sections were counter-stained with hematoxylin and fixed. For semi-quantitative evaluation of the IHC, 20 random visual fields were examined using the LeicaQ550cw imaging analysis system (Germany).

Statistical analysis. Data were reported as means ± SD. The paired t-test was used for statistical analysis, and p<0.05 was considered statistically significant.

Results

Morphological characteristics. After 3-4 days of incubation, monolayer cells were adhered to the bottom of the flask and showed significant cell body extension, while MCS were floating in the media (Fig. 1A). SEM showed that the MCS were in different sizes, ranging from 160-200 μm in diameter and their shapes were either oval spheroid, polyhedron or tightly packed (Fig. 1B). TEM showed abundant desmosome and intermediate junctions only in MCS. In parallel, the monolayer cells were typically polygonal-shaped with more microvilli on their surfaces, and had wider intercellular spaces
compared to those in the MCS. Interestingly, cells in MCS had significantly higher numbers of mitochondria than monolayer cells (Fig. 1C).

The relationship of multicellular organization to proliferation and apoptosis. As shown in Fig. 2, more cells in the MCS accumulated in the G1 phase of the cell cycle (59.23±0.41%) as compared to the monolayer cells (39.45±0.27%) (p<0.01). Consequently, the MCS had a smaller cell population in the G2 phase (12.47±0.65%) than the monolayer cells (36.15±0.69%) (p<0.01). The percentage of cells in S phase was similar between monolayer cultures (22.37±0.76%) and MCS (26.82±0.18%) (p>0.05). MCS also have altered kinetics of drug-induced cell death. Specifically, the early stage cell death (ED) for MCS and

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Figure 1. Morphological characterization of 2-D and 3-D HepG2 cell cultures. (A) Phase contract microscopy, x100. (B) Scanning electron microscopy, x30000 and (C) transmission electron microscopy, x60000.

Figure 2. The effects of cellular organization on HepG2 cell cycle progression. Summarizes the quantitative differences of monolayer cultures and MCS in different phases of the cell cycle. Each experiment was repeated 5 times (*p<0.05).
monolayer cells were 1.62±0.50% and 6.29±1.61%, respectively (p<0.05). Late stage cell death (LD) was not significantly different between the two types of cell cultures (Fig. 3).

The drug sensitivity of MCS. The sensitivity of the cells cultured in monolayers and MCS to CDDP, 5-FU, and ADM was investigated by MTT assay. Cell inhibition ratios in monolayer cultures exposed to various concentrations of drugs were higher than those in MCS (p<0.05) (Fig. 4A), indicating the HepG2 cells in MCS became resistant to CDDP, 5-FU, and ADM. When drug concentrations were increased, cell inhibition ratios also increased.

On the basis of MTT assay results, an Annexin V/PI staining method was used to evaluate the apoptosis rate at 3 different concentrations of each drug. As expected, apoptosis rates were much lower in MCS than those in monolayer cells after exposure to 5-FU and ADM for 48 h (p<0.05), but were not significantly different after exposure to CDDP for the same amount of time (p>0.05) (Fig. 4B).

The protein expression profile of MCS. IHC analysis demonstrated that the cell adhesion molecules E-cadherin and CD44v6 were upregulated in MCS compared to monolayer cells (Table I). E-cadherin protein was detected predominantly in the cellular membrane and in the cytoplasm (Fig. 5A). The positive signals of CD44v6 expression were observed mainly in the cellular membrane (Fig. 5B). Further analyses in this study showed that Cyt-C levels were higher in cells in MCS than in monolayer cells. The cytoplasm of MCS was rich in mitochondria as revealed by transmission electron microscopy. Thus, cells in MCS seemed to have preserved a high level of integrity. In parallel, Bax expression

**Table I. Immunohistochemical pictorial analysis of protein expression.**

<table>
<thead>
<tr>
<th>Protein</th>
<th>Mean grey value</th>
<th>Monolayer cells</th>
<th>MCS</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>E-cadherin</td>
<td>232.28±7.80</td>
<td>199.93±16.58</td>
<td>&lt;0.01</td>
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<tr>
<td>CD44v6</td>
<td>256.82±13.87</td>
<td>207.48±8.79</td>
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<tr>
<td>VEGF</td>
<td>209.81±7.11</td>
<td>194.71±13.23</td>
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<tr>
<td>KDR</td>
<td>174.40±19.12</td>
<td>140.98±19.92</td>
<td>&lt;0.01</td>
<td></td>
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<tr>
<td>Endostatin</td>
<td>162.42±6.88</td>
<td>148.48±11.06</td>
<td>&lt;0.01</td>
<td></td>
</tr>
<tr>
<td>Cyt-C</td>
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<td>211.12±21.02</td>
<td>&lt;0.05</td>
<td></td>
</tr>
<tr>
<td>Bax</td>
<td>172.56±17.80</td>
<td>171.76±14.69</td>
<td>&gt;0.05</td>
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</tbody>
</table>

Figure 3. Cell death analyses by flow cytometry. Cell death of monolayer cultures and MCS detected at early (ED) and late (LD) time points were quantified and summarized. Data are the average of five experiments (*p<0.05).

Figure 4. Drug sensitivities of HepG2 cells in a monolayer culture and MCS. (A) Cell viability measured by MTT assay. The dose-dependent effects of the indicated drugs on cell proliferation are presented as percent inhibition of cell viability. (B) Quantification of the apoptotic cell population is based Annexin V-positive/PI-negative flow cytometry. Data represent the average of five experiments (*p<0.05).
Figure 5. IHC profiling of protein expression. Representative staining results are shown for E-cadherin (A), CD44v6 (B), VEGF (C), KDR (D), endostatin (E), cytochrome-c (F), and Bax (G). IHC performed with PBS in the presence of the primary antibody was used as the negative control (H). The magnifications for monolayer cultures and MCS are x200 and x400, respectively.
Compared with monolayer culture, MCS was more differentiated phenotypes. We showed that, in MCS of HepG2 cells cultured as 3-D cultures not found in 2-D cultures. Cells in 3-D cultures i) express more extracellular matrix proteins (22); ii) are linked by membrane protein anchorage with more cell adhesion molecules such as E-cadherin and CD44v6 (10,23-25); and iii) express more proteins involved in heat shock response or hypoxia (26-29). The overexpression of cell adhesion molecules seems to be particularly advantageous for survival, since inhibition of these molecules by specific antibodies or siRNA-mediated gene silencing disrupted the spheroid formation (10) and rendered cells more sensitive to drug-induced apoptosis (30).

MCS may favor angiogenesis. The growth and metastasis of tumors depends on angiogenesis. Pathological angiogenesis plays a pivotal role during hepatic carcinogenesis. Due to their 3-D architecture, multicellular tumor spheroids may also manifest the biochemical constraints found in a fast growing tumor mass that eventually develops the dependence on angiogenesis. Indeed, angiogenesis-related molecules VEGF, VEGFR-2, and endostatin were increased in the cells with 3-D cultures not found in 2-D cultures. Cells in 3-D cultures i) express more extracellular matrix proteins (22); ii) are linked by membrane protein anchorage with more cell adhesion molecules such as E-cadherin and CD44v6 (10,23-25); and iii) express more proteins involved in heat shock response or hypoxia (26-29). The overexpression of cell adhesion molecules seems to be particularly advantageous for survival, since inhibition of these molecules by specific antibodies or siRNA-mediated gene silencing disrupted the spheroid formation (10) and rendered cells more sensitive to drug-induced apoptosis (30).

In summary, we provide in vitro evidence that MCS of hepatoma cells mimic the heterogeneous multicellular structures and the more differentiated/quiescent cellular behavior that are common characteristics of solid tumors in vivo. More importantly, this MCS structure restored the resistance of tumor cells to drug-induced apoptosis. These findings call for a greater adoption of 3-D culture systems for cancer drug screening and for better mechanistic studies of MDR.
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13. Ponce de León V and Barrera-Rodríguez R: Changes in P-glycoprotein activity are mediated by the growth of a tumour cell line as multicellular spheroids. Cancer Cell Int 5: 20, 2005.