Size-dependent resistance of human tumour spheroids to photodynamic treatment

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Summary Spheroids derived from the human colon adenocarcinoma cell line, WiDr, were exposed to 10 μg ml⁻¹ Photofrin II and irradiated with light (700 nm, 50 mW cm⁻²). Compared with exponentially growing monolayer cultures, cells in spheroids of 100, 250 and 500 μm diameter were respectively 1.8, 2.5 and 22-fold less sensitive. The small resistance of plateau-phase cultures (1.3-fold) was insufficient to account for this marked spheroid size-dependent resistance. For monolayer cultures and for spheroids of 100 and 250 μm diameter, the results were the same whether irradiations were carried out pre- or post-trypsinisation. However, there was a difference for the largest spheroid size: when irradiations were carried out pre-trypsinisation, spheroids were more resistant than when irradiations were given post-trypsinisation. Drug extraction studies showed there was no difference in the average drug uptake between cultures of exponentially growing or plateau-phase cells, and 100 μm diameter spheroids while 250 and 500 μm diameter spheroids took up proportionally 0.5 and 0.4 as much drug. Cell contact effects, drug heterogeneity between cells, hypoxia and problems in drug penetration are suggested as possible reasons for the resistance of large spheroids to photodynamic treatment.

During the past decade there has been increasing interest in the potential of photodynamic therapy (PDT) as a new modality for the treatment of cancer. Although many of the factors determining response have been investigated, there are still many areas of the biology of PDT yet to be studied. The use of multicellular spheroids as a tool for the investigation of tumour response to other therapies is well established. They are useful, for example, for assessing the contribution of hypoxia (Sutherland et al., 1970; West et al., 1984; West & Sutherland, 1987), diffusion limitations (Sutherland et al., 1979; Durand, 1981), cell contact effects (Durand & Sutherland, 1972; West & Stratford, 1987), and repair processes (Durand & Sutherland, 1972; West et al., 1984) in the response of cells to a particular treatment.

The resistance of spheroids to photodynamic treatment has been documented previously (Christensen et al., 1984). However, to date, there has been no systematic study of the effect of spheroid size on the development of this resistance. In addition, there have been conflicting reports of the effect of cell growth phase on sensitivity to photodynamic treatment. Christensen et al. (1984) reported that log- and plateau-phase NHIK 3025 cells were equally sensitive to haematoporphyrin derivative (HPD) plus light, while Ben-Hur et al. (1987) showed that V79 cells in the plateau-phase of growth were more sensitive than exponentially growing cells to photosensitisation with chloroaluminium phthalocyanine.

The following work describes a study of the effect of spheroid size on photosensitisation by Photofrin II. In addition, the role of the growth phase of cells during drug exposure has been investigated. The end-point used was clonogenic cell survival.

Materials and methods

Monolayer cultures

The human colon adenocarcinoma cell line, WiDr, was obtained from the American Type Culture Collection (Rockville, MD, USA) at passage 24 from the original tumour. The cells were maintained in exponential growth in Eagle's basal medium (BME, Gibco, Paisley, Scotland) plus 10% fetal calf serum (FCS, Gibco), supplemented with penicillin (Gibco, 100 IU ml⁻¹) and streptomycin (Gibco, 0.1 mg ml⁻¹). Cultures were incubated in a humidified 5% CO₂ atmosphere and subcultured weekly using 0.01% trypsin (180 IU ml⁻¹). Worthington Diagnostics Ltd, Freehold, NJ, USA) in 0.02% ethylenediamine tetra-acetic acid (EDTA, BDH Chemicals Ltd, Poole, England) made up in phosphate buffered saline (PBS, pH 7.3). Under these conditions the cells grew with a doubling time of 0.8 days and a plating efficiency of around 90%. Stocks of cells were stored frozen and experiments were carried out on cell passages 28-44 from the original tumour. Cultures were tested routinely for Mycoplasma.

Spheroid cultures

Multicellular spheroids were initiated by inoculating 5 × 10⁵ cells in 10 ml growth medium on to 100 mm Petri dishes which had been underlayered previously with 1% agar (Noble; Difco, Detroit, MC, USA) in BME without serum. After 5 days' incubation, spheroids were filtered through 95 and 100 μm nylon screens to obtain a homogeneous population and approximately 1,000 were placed in 95 mm diameter spinner flasks (Techne) containing 250 ml growth medium. Flasks were gassed with an air plus 5% CO₂ gas mixture and placed on magnetic stirrers set at 60 r.p.m. in a 37°C warm room. The medium was replaced thrice-weekly and when spheroids had reached approximately 300 μm in diameter the number of spheroids per flask was reduced to 500 and the stir rate increased to 80 r.p.m. Spheroids were dissociated to single cells using a 15 min exposure to 0.05% trypsin in EDTA.

Drug exposure

Photofrin II (batch number PC238C), 2.5 mg ml saline⁻¹, was a gift from Photomedica Inc. (New York, USA). On arrival the drug was thawed and aliquoted into small quantities which were stored frozen at −20°C and thawed immediately before use. Exponentially growing (days 4–7 cultures) and unfed plateau-phase (days 12–13) monolayer cultures were exposed to 10 μg ml⁻¹ drug in BME plus 10% FCS. After 24 h the drug containing medium was removed, the cells washed and fresh BME plus 10% FCS added for a further 1 h. The cells were trypsinised, resuspended at 3–5 × 10⁵ cells ml⁻¹ in Hepes-buffered BME plus 10% FCS at room temperature, and 1 ml aliquots were placed into 35 mm Petri dishes. Spheroids were selected for the appropriate size and the size distribution estimated by measuring 30. Then size-selected
spheroid cultures were exposed to drug, 10 μg ml⁻¹ in 70 ml growth medium, in small Techni flasks (65 mm diameter). After 24 h, the spheroids were harvested by sedimentation, rinsed in BME plus 10% FCS, placed in fresh growth medium in a Petri dish and incubated at 37°C. After 1 h, one-half of the spheroids were trypsinised and then either intact or disaggregated spheroids were placed in 1 ml Hepes buffered BME plus 10% FCS in 35 mm Petri dishes.

Light irradiation

The light source used has been described in detail elsewhere (Moore et al., 1986). Briefly, light irradiations were carried out using 12 V, 100 W quartz tungsten halogen lamps (Xenophot HLX; Woton, London) with a KG1 infra-red filter (Schott, Mainz, FR Germany). This set-up produced light within the wavelength range 300–1,100 nm, with peak spectral irradiations at 700 nm. A circular light beam of 50 mm diameter uniform to 10% across the beam was produced with a power density at the working surface of 50 mW cm⁻². Single cell suspensions, 1 ml at 3–5 × 10³ cells ml⁻¹. Hepes buffered BME plus 10% FCS, were irradiated at room temperature in 35 mm Petri dishes with the lids removed. For some experiments exponentially growing monolayers were irradiated before trypsinisation. Cells were seeded on to 35 mm dishes and exposed to drug and washed as described above. At 1.5 h post-wash, the cells were irradiated and immediately trypsinised. Likewise, spheroids irradiated pre-trypsin were irradiated in 35 mm dishes in 1 ml Hepes buffered growth medium 1.5 h after removal of the drug, and then trypsinised. The longest irradiation time was 11 min 40 s and during this time the maximum recorded temperature rise was from 20 to 24°C.

Cell survival

Following illumination, the cells were counted, serially diluted and appropriate numbers plated in triplicate in BME plus 10% FCS on to 60 mm Petri dishes. The dishes were incubated at 37°C in an air plus 5% CO₂ atmosphere. After 21 days colonies were stained with methylene blue and counted. Curves were fitted to all data using a single-hit, multi-target type equation (e.g. Gilbert, 1969). Use of such equations, derived initially to describe survival after exposure to sparsely ionising radiation, gave adequate empirical fits to the data. However, the interpretation of the curve shapes may well differ for the two modalities. From the fitted curves, values were obtained for the parameter D₅₀ (the reciprocal of the slope of the final exponential portion of the curve). In order to test for differences between curves, analysis of variance was carried out. χ² values were obtained for sets of curves fitted independently or pooled. An F test was then used to examine for significant differences between the χ².

Drug uptake

The method used for measuring drug uptake by cells has been described elsewhere (Pantelides et al., 1989). Monolayer cultures and spheroids were exposed to drug, washed and trypsinised as described above. After washing in PBS, 10³ cells were resuspended in 1 ml 0.1 M Hepes and 10 mm cetyltrimethylammonium bromide at pH 7 and vortexed for 30 s. Samples were protected from light and stored at −20°C. On the day of analysis, samples were thawed and 5 ml of a 1:4 solution of ethyl acetate:glacial acetic acid was added, followed by 4 ml 1 M HCl. Two layers resulted and the volume of the lower, containing the extracted porphyrins in HCl, was recorded. Fluorescence was measured at room temperature, using a spectrofluorophotometer (Shimadzu) before and after the sample had been incubated in a water bath at 90°C for 10 min (to monomerise aggregated drug). Excitation was at 404 nm with fluorescence emission measured at 594 nm. A standard curve was produced by adding known amounts of drug to 1 ml HCl. This was linear up to at least 0.4 μg ml⁻¹. There was no fluorescence quenching by the cells up to at least a cell density of 2 × 10⁷ cells ml⁻¹. Fluorescence measurements were made on cells that had not been exposed to Photofrin II, and the values were found to be negligible. All measurements were carried out in triplicate on four separate occasions and the mean (of the means) ± standard error of the mean (s.e.m.) was calculated.

Results

Figure 1 illustrates the sensitivity of exponentially growing WiDr cells that had been exposed to 10 μg ml⁻¹ Photofrin II for 24 h, washed for 1 h in BME plus 10% FCS and irradiated with light. Light alone, up to 10⁴ J cm⁻² has been shown previously to be non-toxic to the cells (West & Moore, 1988) and so light-only controls were not carried out routinely. Drug-only exposed cells formed the control for all experiments and plating efficiencies did not fall below the range of values expected for cells without the drug. It has been shown already that, under the experimental conditions employed, only Photofrin II levels greater than 150 μg ml⁻¹ were toxic to WiDr cells (West & Moore, 1988). The sensitivity of the cells to Photofrin II plus light was the same whether irradiations were carried out before or after trypsinisation (Figure 1). A single experiment is shown in which cells were irradiated pre- and post-trypsinisation on the same day. There was no significant difference in the survival curves shown (P=0.735). This suggests that (1) treatment of WiDr cells with trypsin does not make them more fragile and so more sensitive to subsequent light treatment and (2) the washing procedure was sufficient to remove all loosely bound drug from the cells.

![Figure 1](image-url)

Figure 1 The sensitivity of exponentially growing WiDr cells to a 24 h exposure to 10 μg ml⁻¹ Photofrin II and a 1 h wash (both in BME + 10% FCS), trypsinisation and irradiation with light. A single experiment is shown in which cells were irradiated both before (open circles) and following (filled circles) trypsinisation to a single cell suspension. Individual data points are shown which are the average of three replicate plates. Different symbols represent independent experiments.
Unfed plateau-phase cultures were more resistant to Photofrin II photosensitisation (by a factor of 1.3; taken from the ratio of the $D_0$s) than exponentially growing cells (Figure 2). Two experiments are shown in which log- and plateau-phase cells were compared on the same day. Analysis of the curves fitted to both sets of data showed them to be significantly different ($P<0.01$).

Cells in small spheroids, approximately 100 and 250 $\mu$m in diameter, were 1.8- and 2.5-fold more resistant than exponentially growing monolayers (Figure 3). For both spheroid sizes, the sensitivity to photodynamic treatment was the same whether irradiations were carried out pre- or post-trypsinisation. Large spheroids, 500 $\mu$m in diameter, showed markedly greater resistance to photodynamic treatment (22-fold compared with exponentially growing cells). For these spheroids there was a difference between the curves for irradiating pre- and post-trypsinisation (Figure 3c). Back-extrapolation of the higher-dose portion of the curve for 500 $\mu$m spheroids irradiated intact, suggested a resistant sub-population of about 40%.

Drug uptake studies were carried out (Table I). There was no statistical difference in the average amount of drug uptake by cells from exponentially growing and plateau-phase cultures and 100 $\mu$m diameter spheroids. Cells from larger spheroids, 250 and 500 $\mu$m in diameter, took up proportionally 0.5 and 0.4 as much drug respectively.

**Discussion**

The relative resistance of plateau-phase cultures to photodynamic treatment is not in accordance with observations published previously by others. It has been reported that log- and plateau-phase NHK cells are equally sensitive to HPD plus light (Christensen et al., 1984) and that plateau-phase V79 cells are more sensitive than those in log-phase to chloroaluminium phthalocyanine (Ben-Hur et al., 1987). Although these studies involved different photosensitisers, the results may parallel the situation seen for sensitivity to sparsely ionising radiation where some cell lines are equally sensitive across the growth phases, others are more sensitive in plateau-phase and others more sensitive in log-phase (Hahn & Little, 1972). Interestingly, cultures of WiDr cells are more sensitive to sparsely ionising radiation when irradiated in the plateau-phase of growth (West et al., 1988). The resistance of plateau-phase WiDr cells to photodynamic

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**Figure 2** The sensitivity of plateau-phase cells (filled symbols) to Photofrin II plus light (conditions as in Figure 1). Individual data points from four separate experiments. For two experiments, exponentially growing (open symbols) and plateau-phase cells were investigated on the same day.

**Figure 3** The sensitivity of 100 (a), 250 (b) and 500 (c) $\mu$m diameter spheroids to Photofrin II plus light. Spheroids were irradiated both pre- (filled symbols) and post- (open symbols) trypsinisation. Broken lines in a and b indicate response of exponentially growing monolayer. The curves are compared in d.

**Table I** A comparison, for different WiDr cultures, of the number of cells per spheroid, the median cell volume, the drug uptake following exposure to 10 $\mu$g ml$^{-1}$ Photofrin II and the slope of the survival curve ($D_0$) after subsequent irradiation with light.

| Culture          | Cells per spheroid | Median cell volume ($\mu$m$^3$) | Drug uptake (ng 10$^6$ cells$^{-1}$) | $D_0$ (J cm$^{-2}$) |
|------------------|--------------------|---------------------------------|-------------------------------------|---------------------|
| Log-phase cells  | -                  | 2,140 ± 96                     | 297 ± 34                            | 0.17 ± 0.02         |
| Plateau-phase cells | -            | 1,807 ± 93                     | 310 ± 26                            | 0.22 ± 0.03         |
| 100 $\mu$m spheroids | 8.3 ± 4.2 x 10$^3$ | 1,967 ± 237                    | 261 ± 8                             | 0.30 ± 0.04         |
| 250 $\mu$m spheroids | 5.8 ± 1.3 x 10$^3$ | 1,943 ± 92                     | 115 ± 12                            | 0.42 ± 0.03         |
| 500 $\mu$m spheroids | 3.5 ± 0.56 x 10$^3$ | 1,818 ± 88                     | 115 ± 18                            | 3.80 ± 0.31$^*$     |

Mean ± s.e.m. for four independent experiments. *Mean of two independent measurements; *Obtained by computer-fitting of survival curves; *Spheroids irradiated post-trypsinisation.
treatment may be related to a cell cycle effect. There is an increase in the proportion of G₂-like cells with increasing WiDrc culture age from 68 (day 6 cultures) to 78% (day 12) (15% increase (West et al., 1980)). It has been shown that, for some cell lines, G₂ cells are the most resistant of cells in the cycle to photodynamic treatment (Christensen et al., 1981). It is possible that WiDr cells show a variable cell cycle response with G₂ resistance and that the increase in proportion of G₂ cells in plateau-phase cultures accounts for resistance to PDT. This is supported by the similarity in drug uptake between exponentially growing and plateau-phase cells (Table 1). In contrast to the results of Christensen et al. (1981), Gomer & Smith (1980) showed no cell cycle response, further supporting the notion of cell line dependent, growth phase related differences.

Although the presence of oxygen is essential for photosensitiser-induced light cytotoxicity (Moan & Sommer, 1985), the size-dependent spheroid resistance seen here may not be related to the presence of hypoxia. For small spheroids there was no difference in photodynamic sensitivity when light irradiations were carried out both before and after trypsinisation to single cell suspensions. This indicates that there was no induced hypoxia as a result of static culture conditions (using static culture conditions for irradiation with sparsely ionising radiation can lead to an increase in the level of hypoxia in spheroids (Durand, 1980)). Secondly, irradiations were carried out in fresh medium and at room temperature, both of which are known to be important in spheroid reoxygenation (e.g. Franko & Koch, 1983). Thirdly, WiDr spheroids, up to 500 μm in diameter, do not contain central necrosis or radiobiological hypoxia, i.e. there is no hypoxic fraction on the radiation survival curve for 500 μm spheroids (West, unpublished data). The induction of a hypoxic region in the larger spheroids during light exposure cannot be totally ruled out because of the difference in sensitivity for spheroids irradiated pre- and post-trypsinisation. However, this may be due also to inefficient light penetration through spheroids. The depth of penetration of light in tissue has been described as being 200 μm at 400 nm and 1 mm at 700 nm (Eichler et al., 1977).

A broad spectrum light source was used (West & Moore, 1988) and absorption by Photofrin II of shorter, less penetrating wavelengths (350-550 nm) dominates its cytotoxic effect (West & Moore, 1989).

The question of interest is what causes the marked spheroid size dependent resistance? The small resistance of 100 μm diameter spheroids may be related to a cell contact effect and there is some evidence that this can occur in monolayer cultures (Christensen & Moan, 1980). The resistance of plateau-phase cells and reduced uptake may be involved also. However, although these may account for the resistance of the smaller spheroids (100, 250 μm), because of the magnitude of the effect, other factors must be involved in the resistance of large spheroids.

There is some indication that heterogeneity of drug uptake may be important. The majority of in vitro photodynamic survival curves have broad and clearly defined shoulders similar to that illustrated in Figure 1 (see also Gomer & Smith, 1980). Spheroids, 100 and 250 μm in diameter, gave similar shaped curves while the curve for 500 μm diameter spheroids was a straight line (Figure 3d). It is interesting to compare this spheroid survival curve with that for exponentially growing cells exposed to 1 μg ml⁻¹ drug (Figure 4). Comparison of drug uptake showed that cells exposed to 10-fold less drug accumulated 10-fold less drug (West, unpublished data) and were 10-fold less sensitive (taken from a ratio of D₅₀ (West & Moore, 1988)) Spheroids of 500 μm diameter took up 3-fold less drug but were 22-fold less sensitive than exponentially growing monolayers. The straight line of the 500 μm spheroid 'curve' and the shallower slope than would be expected on the basis of reduced average drug uptake suggest an increase in heterogeneity. It is possible that there is a wide spectrum of intracellular drug levels following the exposure of large spheroids to Photofrin II. A compilation of a number of PDT survival curves for different drug-exposed cell populations would produce a survival curve with a shallower slope than each of the composite curves (see Dutreix et al., 1988) and a curve without a shoulder.

A drug penetration problem is the obvious explanation and our initial expectation was of a gradient in drug concentration from the peripheral to the central region. This has been demonstrated in spheroids exposed to the chemotherapeutic agent adriamycin (Sutherland et al., 1979). However, preliminary studies of fluorescence micrographs of centrally cut frozen sections of Photofrin II-exposed spheroids suggest good overall penetration but a patchy distribution between cells with some fluorescing brightly and others dimly regardless of position across the spheroid (West, unpublished data). In addition, there is no tail to the survival curve for spheroids irradiated post-trypsinisation which would be seen if there was a drug-free sub-population of cells.

If drug penetration is not thought to be the cause of heterogeneity of drug uptake within spheroids, what is? It could be related to a cell contact effect. The degree of cell coupling has been shown to decrease as a function of spheroid age and size (Dertinger et al., 1982). Alternatively, or in addition, decreased drug uptake in some cells of spheroids may be related to membrane changes that might occur with increasing spheroid size. Serum lipoproteins have a high affinity for haematoporphyrins and drug uptake may be via receptor-mediated endocytosis of complexes of porphyrin with low density lipoprotein (LDL; see Barel et al. (1986) for discussion). There may be a decrease in the number of LDL receptors in some cells when grown in close contact, leading to an increase in the heterogeneity of drug intake due to the different drug uptakes.

In conclusion a marked spheroid size dependent resistance to photodynamic treatment has been demonstrated. This

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**Figure 4** A comparison of the survival curves for exponentially growing WiDr cells exposed to 10 (from Figure 1) and 1 (from West & Moore, 1988) μg ml⁻¹ and 500 μm diameter spheroids (from Figure 3e) exposed to 10 μg ml⁻¹ Photofrin II followed by irradiation with light.
resistance cannot be explained simply in terms of hypoxia, reduced drug uptake or the limited penetration of Photofrin II into spheroids. There is evidence to suggest that heterogeneity of intracellular drug levels between cells is important. This may be mediated via changes in the degree of cell contact between cells or in the membrane composition of cells when grown as three-dimensional multicellular spheroids.

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