Characterization of radiation resistant hypoxic cell subpopulations in KHT sarcomas. (ii) Cell sorting

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Summary  Hypoxic cells in KHT sarcomas were characterized using fluorescence activated cell sorting based on the diffusion properties of the fluorochrome Hoechst 33342. Tumour-bearing female C3H/HeJ mice were injected i.v. with 10 µg g⁻¹ Hoechst 33342 and the cells derived from the tumours sorted on the basis of their staining intensities. For each sorted fraction the DNA histogram was evaluated using FCM analysis. The results indicated that the bright and dim cells were not equally distributed about the cell cycle. For example, a greater proportion of S phase cells were in the bright subpopulations whereas the dim subpopulations contained an increased proportion of cells in G₁. When the tumours were irradiated with a single dose of radiation prior to cell sorting, the dim cells survived preferentially. Dose response curves for the 20% most dim and 20% most bright cells, sorted on the basis of fluorescence intensity, then were determined. The survival curves of the dim and bright cells were found to have slopes similar to those of KHT cells irradiated in situ in dead animals or in vitro under fully oxic conditions, respectively. In addition, when KHT sarcoma-bearing mice were given a 2.5 mmol kg⁻¹ dose of mitomycin (MSO) prior to irradiation and cell sorting, the dim subpopulation was sensitized whereas the bright subpopulation was not. These findings suggest that (i) compared to well-oxygenated areas, hypoxic regions of KHT tumours contain a smaller percentage of cells actively proliferating and (ii) Hoechst 33342 sorting may allow the detailed in situ evaluation of agents acting directly against hypoxic cells in solid tumours.

Since the milestone observations of Thomlinson and Gray in 1955, which suggested that hypoxia might possibly severely hamper the success of clinical radiotherapy, hypoxic tumour cells and approaches to eliminating them have dominated much of experimental radiobiology (Adams, 1977; Guichard et al., 1980; Fowler, 1985). Similarly, with the mounting evidence that oxygen-deficient tumour cells may be refractory to certain clinically active anti-cancer agents (Teicher et al., 1981; Siemann, 1984; Sartorelli, 1986; Adams & Stratford, 1986), the role of hypoxia also has received considerable attention in both the assessment of the efficacies of conventional chemotherapeutic agents and in new agent development (Moore, 1977; Kennedy et al., 1980; Teicher & Sartorelli, 1980; Teicher et al., 1981). It is clear that most rodent tumours contain hypoxic cells (for review, see Moulder & Rockwell, 1984) and the evidence is fairly convincing that such cells may influence the clinical outcome in at least some human neoplasias treated by radiotherapy (Bush et al., 1978; Guichard et al., 1980; Dische et al., 1983; Overgaard et al., 1986). Consequently, there has been considerable interest in the in situ characteristics of this, with respect to therapeutic outcome, potentially critical tumour cell subpopulation. Yet this has been difficult primarily as a consequence of an inability to isolate or characterize directly from solid tumours this relatively small subpopulation (typically 10–20%) of radiobiologically hypoxic cells in the presence of an overwhelming majority of well oxygenated tumour cells. Conventional cell proliferation studies, utilizing the technique of thymidine autoradiography have demonstrated that both labeling and mitotic index decreased marked with increasing distance from blood vessels (Tannock, 1968; Tannock & Steel, 1970). Consistent with these classic observations we previously used centrifugal elutriation in conjunction with FCM and cell survival curve analysis to show that hypoxic cells in the KHT sarcoma were located primarily in the G₁ phase of the cell cycle (Siemann & Keng, 1987). Taken together, these observations imply that hypoxic cells in tumours demonstrating diffusion limited hypoxia may be resistant to therapy not only due to their lack of oxygen but also a consequence of their cell cycle or proliferation state.

A novel approach, which utilizes fluorescence activated cell sorting to study hypoxic cells in solid tumours, was recently developed by researchers at the British Columbia Cancer Research Centre (Chaplin et al., 1985; Olive et al., 1985; Chaplin et al., 1986). This technique allows cells to be sorted on the basis of their proximity to blood vessels by making use of the diffusion properties of the fluorochrome stain Hoechst 33342. This stain has a very short distribution half-life after injection and remains bound within cells (Olive et al., 1985). Olive et al. (1985) and Chaplin et al. (1985, 1986) have utilized this method extensively to study hypoxic cells in SCCVII tumours. In the present investigations, this technique was applied to the KHT sarcoma. In particular, the aim of these studies was to characterize more definitively and gain a better understanding of the hypoxic cell subpopulation in this solid tumour model.

Materials and methods

Animals and tumour model

The KHT sarcoma (Kallman et al., 1987) was used in all experiments. KHT cells are passaged in vivo every two weeks by preparing single cell suspensions from solid tumours using a mechanical dissociation procedure (Thomson & Rauth, 1974). For experiments, 2 x 10⁵ tumour cells were injected into the hind limbs of 8–14 week-old female C3H/HeJ mice obtained from Jackson Laboratories (Bar Harbor, Maine). Tumours were used when they reached a size of 0.5–0.7 g.

Irradiation

Unanaesthetized mice were irradiated whole body using a ¹³⁷Cs source operating at a dose rate of 4.75 Gy min⁻¹. The animals breathed air during the irradiation procedure. Immediately after treatment, the tumours were dissociated to single cell suspensions using a combination of mechanical and enzymatic (0.2% trypsin plus DNase) dissociation procedure (Thomson & Rauth, 1974).

Sorting cells on the basis of fluorescence intensity

KHT sarcoma-bearing mice were injected i.v. with a 10 µg g⁻¹ dose of Hoechst 33342 twenty minutes prior to irradiation. Immediately following treatment, the mice were...
killed and the tumours dissociated to single cell suspensions as described above. Cells were prepared in 4°C phosphate buffered saline at a concentration of 5 × 10^6 cells per ml and sorted according to fluorescence intensity using an EPICS V flow cytometer (Loeffler et al., 1987). The cells were stirred and maintained at 4°C during sorting. Laser excitation at 100–150 mW, generated from an Argon laser, was used throughout the experiments. A minimum of 1 × 10^5 cells was collected for each sorted fraction.

**Sensitizer treatment**

MISO was received from Dr Ven Narayanan, Drug Synthesis and Chemistry Branch of the National Cancer Institute. The sensitizer was dissolved in sterile saline at a concentration of 20 mg ml^-1 and was injected i.p. according to body weight 30–45 min prior to irradiation.

**Flow cytometric analysis**

For each sorted fraction, FCM analysis was used to determine the percentage of cells in the G_i, S and G_i+M phases of the cell cycle (Keng et al., 1981; Siemann & Keng, 1986). Briefly, FCM measurements were made with an EPICS V dual laser (argon and krypton) flow cytometer (Coulter Electronics Inc.), using a TERAK 8600 minicomputer for data storage and analysis. Cells were fixed in 70% ethanol and stained with mithramycin (1.0 mg ml^-1) according to the methods of Crisman & Tobey (1974). DNA histograms were analyzed using the model of Fried & Mandel (1979) implemented as the 'CCycle' program on the TERAK 8600 system. Previously obtained results using the KHT sarcoma have demonstrated a close agreement between the estimates of cells in the various phases of the cell cycle based on FCM analysis and tritiated thymidine uptake (Keng et al., 1981).

**Clonogenic cell survival assay**

After sorting, the cells in fractions of differing fluorescence intensities were counted using a haemocytometer. Various dilutions were mixed with lethally irradiated cells in 0.2% agar containing alpha-minimal essential medium supplemented with 10% foetal calf serum plated into 24-well multiwell plates. The plates were maintained in a 5% CO_2 in air atmosphere at 37°C for two weeks. By this time, the surviving tumour cells formed colonies which were counted with the aid of a dissecting microscope. For each sort fraction, survival values were calculated on the basis of the number of non-neoplastic cells actually plated as determined from different counts performed on cytocentrifuge slides (Siemann et al., 1981).

**Results**

In the initial experiments tumour-bearing mice were irradiated with a single dose of either 10 or 15.5 Gy 20 min after Hoechst dye injection. After tumour dissociation, the cells were sorted on the basis of fluorescence intensity into 5 fractions. Cells from each sort fraction were prepared for FCM and cytocentrifuge slide analysis and also plated for clonogenic cell survival.

DNA histograms of cells dispersed from solid KHT sarcomas and sorted on the basis of fluorescence intensity are illustrated in Figure 1a–e. Cell suspensions prepared from KHT sarcomas contain a mixture of both non-neoplastic infiltrating host cells and neoplastic cells (Siemann et al., 1981). In untreated tumours, the former typically represent ~40–60% of the total cell population. These host cells readily can be identified by morphological analysis or by their diploid DNA content. The latter contrasts with the near-tetraploid DNA content of neoplastic KHT cells. The results in Figure 1 and Table I illustrate that the sort fractions contained non-neoplastic cells, but not necessarily in equal proportions. Further, the data in Table I demonstrate that the percentage of normal host cells determined on the basis of differential counts was in good agreement with the percentage of diploid cells determined using FCM analysis.

A comparison of the histograms of the sorted cell populations (Figure 1) further suggested that the cells in the various fractions were not equally distributed about the cell cycle. Consequently, FCM analysis was performed on all the sort fractions. The data show (Figure 2) that the proportion of

![Figure 1](https://via.placeholder.com/150)

**Table 1** Host cell distributions in the 20% most dim and 20% most bright sort fractions sorted on the basis of Hoechst 33342 fluorescence intensity

| Sort fraction | FCM analysis | Differential counts |
|---------------|--------------|---------------------|
| 20% most bright | 15.4 ± 8.1 | 18.9 ± 4.2 |
| 20% most dim  | 53.3 ± 12.4 | 45.0 ± 8.0 |

*Data are the mean ± s.d. of 5–8 experiments.
cells in the S phase of the cell cycle (○) increased from 18.9 ± 3.2% in the most dim fraction to 34.2 ± 4.9% in the most bright. For the same sort fractions, the proportion of G1 cells (●) decreased from 75.2 ± 7.2 to 51.3 ± 6.8%. G2 + M cells (▲) increased from 5.9 ± 2.1% in the dimmest sort fractions to 14.5 ± 4.3% in the most bright.

Survival of cells in the isolated fractions obtained from tumours irradiated with 10 or 15.5 Gy prior to sorting is shown in Figure 3. Results presented have been corrected for the presence of non-neoplastic host cells as determined from differential counts made of cytospin slides (Siemann et al., 1981). The data indicate that for both radiation doses investigated, radiosensitivity increased with increasing fluorescence intensity. Survival of cells from unirradiated tumours was constant in all the sort fractions (data not shown).

On the basis of these initial results, investigations were undertaken in which tumour-bearing mice were irradiated with a range of doses prior to tumour dissociation and cell sorting. In these experiments, only the 20% most dim and 20% most bright cells obtained by sorting were analyzed in detail. Host to tumour cell ratio, cell cycle distribution and clonogenic cell survival were measured in each experiment.

Cell survival in the isolated cell fractions following tumour irradiation with doses ranging between 10 and 18 Gy are shown in Figure 4a. All data were again corrected for the presence of host cells. The results demonstrate that the brightly fluorescent tumour cells (○) were 2- to 3-fold more radiosensitive than those in the dimmest cell fraction (●). The Do values of the most dim and most bright cell subpopulations were calculated to be 1.25 ± 0.2 Gy and 3.84 ± 0.2 Gy, respectively. These values are comparable to the Do values of KHT sarcoma cells irradiated under fully aerobic or anoxic conditions (Hill et al., 1979; Siemann & Kochanski, 1981; Siemann & Keng, 1984; Siemann & Mulcahy, 1984). Despite the differences in cell cycle distributions (Figures 1 and 2), the data in Figure 4 implied that the difference in radiosensitivity between the cells in the brightest and dimmest sort fractions was predominantly a consequence of a difference in the cellular oxygenation state at the
evaluate independently the host and neoplastic cells derived from KHT tumours. Cells were stained either in vivo prior to tumour dissociation and cell separation or in vitro following tumour dissociation and elutriation. The G2 tumour cells were found to fluoresce ~2-fold more brightly than tumour cells in the G1 phase; likely due to their being twice as large. Consequently it is possible that the proportion of G2+M cells calculated to be in, for example, the 0–20% sort fraction (Figure 2) may be somewhat of an underestimate since some of these cells that should have been in this fraction may actually have appeared in the next (20–40%) sort fraction. However, it is likely that this factor had only a relatively minor effect on the observed cell cycle distributions in the various sort fractions (Figure 2), because the difference in the fluorescence of the 0–20% fraction and the 20–40% fraction was >6-fold. The present results therefore indicate that the bright and dim staining cells were not equally distributed about the cell cycle. Rather, the proportion of S cells decreased with distance from the vessels while the proportion of G2 cells increased. These data are entirely consistent with our previous results obtained using centrifugal elutriation to isolate hypoxic cells (Siemann & Keng, 1987) and imply that chronic hypoxia is the dominant form of hypoxia in this tumour model. This conclusion also is consistent with the data shown in Figure 3 which illustrate increasing tumour cell killing with increasing fluorescence intensity (i.e. the proportion of cells staining brightly) over time. Similar results have been observed previously by Chaplin et al. (1986) in small SCCVII tumours containing chronically hypoxic cells.

In addition to the cell cycle distribution differences, the results illustrated in Figure 1 and Table I demonstrate that the proportion of diploid non-neoplastic host cells was not constant in the various sort fractions. The DNA profiles and cytocentrifuge analyses indicate the existence of a larger percentage of host cells in the dimmer sort fractions than in those staining more brightly; suggesting a larger proportion of host cells in the hypoxic regions of the tumour. The interpretation assumes that the two cell types (host and neoplastic) have similar Hoechst 33342 staining patterns. However, similar to observations made in the EMT-6 tumour model (Loeffler et al., 1987), when the host and neoplastic cell subpopulations were stained in vitro with Hoechst 33342 after separation by centrifugal elutriation, the average fluorescence intensity of the host cells was 3–4-fold lower than that of the isolated neoplastic KHT cells. These staining pattern differences could be due to differences in the cells' size, surface area or a number of other factors. Irrespective of the mechanism, the data imply that host cells sorted into a given fraction according to their DNA content intensity may not be derived from the same physical location within a tumour as the neoplastic cells sorted into the same fraction. Consequently, it is not possible to draw firm conclusions from the present findings about the distribution of host cells relative to aerobic or hypoxic KHT tumour cells. Although significant cell cycle distribution differences between the bright and dim staining cell populations were observed (Figure 2), particularly with respect to the S and G2 cell cycle phases, it is unlikely that these differences had a major impact on the cell survival illustrated in Figures 3 and 4. This is because in the KHT sarcoma, there is little difference between the radiation sensitivity of the most resistant S and G1 cell subpopulations (Siemann & Keng, 1984; Keng et al., 1984) i.e., the cell subpopulations which dominate the dimmest and brightest sort fractions. However, when Hoechst 33342 sorting is used to study hypoxic cell subpopulations in other tumour models, differences in the cell cycle distributions in the sorted fractions may need to be considered when interpreting the results.

When the nitroimidazole MIS0 was administered to tumour-bearing mice prior to irradiation, the dimmest tumour cell sort fraction was brightest, whereas the brightest cell fraction was not (Figure 4b). These data support previous results (Chaplin et al., 1985, 1986; Olive et
al., 1985) which indicated that Hoechst 33342 could be used to sort radiobiologically hypoxic cells directly from solid tumours or multi-cell spheroids. In addition, these findings offer direct in situ evidence that MISO sensitizes hypoxic tumour cells preferentially.

In summary, the present results suggest that the cell isolation technique based on Hoechst 33342 stain diffusion may be useful in both the characterization of hypoxic cell subpopulations in vivo and the evaluation of the efficacy of therapeutics against oxic and hypoxic tumour cell subpopulations. The latter may be of particular importance in (i) mechanistic studies of drug actions and (ii) the development of new agents with modes of action directed specifically (either as radiosensitizers or as cytotoxic agents) against hypoxic tumour cells.

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