Human hepatoma cells rich in P-glycoprotein are sensitive to aclacuribin and resistant to three other anthracyclines

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Summary Drug resistance is a major obstacle to successful chemotherapy of primary liver cancer, which is associated with high expression of the multidrug resistance (MDR) gene product P-glycoprotein (Pgp), a multidrug efflux transporter. The most effective single agents in treatment of primary liver carcinoma belong to the anthracycline family, yet several anthracyclines are known to be substrates for Pgp. In the present study, we compared four anthracyclines with respect to cell growth inhibition, intracellular accumulation and cellular efflux using the HB8065/R human hepatoma cell line which is rich in Pgp, and the Pgp-poor parental line HB8065/S. The anthracyclines were also administered in conjunction with the Pgp-modifying agents verapamil and SDZ PSC 833 to assess modulation of resistance. The HB8065/R cells were sensitive to aclacuribin (ACL) and highly resistant to etoposide (EPI), doxorubicin (DOX) and daunorubicin (DNR). SDZ PSC 833 enhanced cytotoxicity of DOX, DNR and ACL to the HB8065/R cells, but none of these effects was seen with ACL. In conclusion, ACL is apparently not transported by Pgp and retains its activity in a multidrug-resistant human hepatoma cell line; such properties can be exploited for clinical purposes.

Keywords: P-glycoprotein; multidrug resistance; flow cytometry; anthracycline; hepatoma; human cell line

Cancer chemotherapy has major limitations because multidrug resistance (MDR) frequently prevents successful treatment outcome in various human cancers. One important mechanism of MDR is expulsion of certain anti-cancer drugs (MDR drugs) from the interior of the malignant cells by means of the multidrug efflux transporter P-glycoprotein (Pgp), which enables the malignant cells to maintain tolerable intracellular levels of these drugs (Juliano and Ling, 1976; Endicott and Ling, 1989). Pgp expression has been shown to correlate negatively with chemosensitivity and survival in leukemias (Campos et al., 1992; Marie et al., 1991), lymphomas (Yuen and Sikic, 1994), childhood sarcomas (Chan et al., 1990), neuroblastomas (Chan et al., 1991) and ovarian carcinomas (Baeckelandt et al., 1994).

Primary liver cancer is one of the most frequently fatal human malignancies, and the response rate to chemotherapy is less than 20% for a series of drug regimens (Falkson et al., 1984). Overexpression of Pgp has been reported in 33–75% of patients at diagnosis (Teetier et al., 1993; Itsuno et al., 1994). Therefore, circumvention of Pgp-mediated MDR might improve the prognosis of unresistable hepatocellular carcinoma. Unfortunately, most of the clinically important anthracyclines appear to be substrates for Pgp (reviewed by Scambia et al., 1994). Among these are the type I anthracyclines epirubicin (EPI), doxorubicin (DOX) and daunorubicin (DNR), which preferentially inhibit the synthesis of DNA. The type II anthracyclines aclacuribin (ACL), which consists of a 9-alkyl-aglycone (aklavinone) and a trisaccharide (rhodamine, 2-deoxyfructose and L-cinerulose A), differs from EPI, DOX and DNR with respect to mechanism of action, as ACL preferentially inhibits RNA synthesis (Muggia and Green, 1991). The 9-alkyl substitution of the anthrancene A ring and certain sugar modifications have also been associated with reduced affinity for Pgp and retention of cytotoxic activity in certain MDR tumour cell lines (Coley et al., 1990). Thus, ACL has some structural characteristics in favour of MDR circumvention as opposed to EPI, DOX and DNR (Figure 1).

ACL has a wide range of activity in tumours and human xenografts (Hori et al., 1977; Oki et al., 1981), and shares the toxic effects of the other anthracyclines, although it appears to be less cardiotoxic (Dantchev et al., 1979; Mortensen, 1987). The cytotoxic activity of ACL is reported to be retained in MDR variants of a mouse mammary tumour line and a human small-cell lung cancer line (Coley et al., 1989, 1993). In clinical trials, several cases of complete remissions have been reported with ACL in DNR-resistant acute myeloid leukaemia (Machover et al., 1984; Pedersen-Bjergaard et al., 1984). In unresactable hepatocellular carcinoma, tumour-targeting chemotherapy with ACL has been reported as highly effective, and patients with unknown Pgp status have achieved response rates of 43–50% (Beppu et al., 1991; Ichikara et al., 1989). The possible relationship between a favourable response to ACL and circumvention of Pgp-mediated MDR warrants study, which could be done in vivo using a hepatoma cell line expressing high levels of Pgp.

In our laboratory, resistance to the second-generation anthracycline epirubicin has been developed in a human hepatoma cell line (HB8065) that has retained differentiated liver cell functions (Knowles et al., 1980; Hall et al., 1991). This resistant subline expresses high levels of Pgp compared with the parental line (Lehne et al., 1994). In the present study, the cytotoxicity and the cellular pharmacokinetics of ACL, EPI, DOX and DNR were compared in the absence or presence of the calcium channel blocker verapamil and the novel cyclosporin D analogue SDZ PSC 833. Apparently, certain calcium channel blockers and cyclosporins bind to Pgp and counteract the active outward transport of MDR drugs (Cornwell et al., 1987; Foxwell et al., 1989; Boesch et al., 1991; Friche et al., 1992). The objective of the present study was to investigate the selectivity of Pgp-mediated resistance to different anthracyclines in human hepatoma cells by assessments of cell growth inhibition, intracellular drug accumulation, cellular drug efflux and response to resistance modifiers.

Materials and methods

Chemicals

The cells were propagated in Eagle’s modified minimum essential medium (EMEM; Bio Whittaker, Walkersville, MA,
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USA, supplemented with 10% fetal calf serum, t-glutamine (0.05 mM mmol ml⁻¹), streptomycin (100 μg ml⁻¹), penicillin (100 U ml⁻¹), and nystatin (40 U ml⁻¹). Trypsin-EDTA (Bio Whittaker) was used to make single-cell suspensions from monolayer cultures. The primary antibody MRK16, which recognises a surface domain of Pgp (Hamada and Tsuruo, 1986), was a gift from Professor Takashi Tsuruo, Institute of Molecular and Cellular Biosciences, The University of Tokyo. The IgG2a isotypic control antibody mouse IgG2a was purchased from Monosan, Uden, The Netherlands. Verapamil hydrochloride was purchased from Knoll (Ludwigshafen, Germany). SDZ PSC 833 was a gift from Sandoz Pharma (Basle, Switzerland). Epirubicin (EPI) and doxorubicin (DOX) were provided by Pharmacia (Milan, Italy), daunorubicin (DNR) from Rhône Poulenc Rorer (Birkerød, Denmark) and aclacinon (ACL) from Medac (Hamburg, Germany). A 500 μg ml⁻¹ stock solution of each anthracycline was prepared in sterile physiological saline solution and kept frozen at -20°C for experimental use within 6 months.

**Scanning fluorimetry of anthracyclines**

The emission and excitation spectra of native anthracycline fluorescence were demonstrated by scanning fluorimetry, using a Hitachi F-4500 fluorescence spectrophotometer (Nissel Sangyo, Tokyo, Japan).

**Cells and culture conditions**

Human hepatoma cells HB8065 (American Type Culture Collection), originating from biopsies of a primary hepatocellular carcinoma (Aden et al., 1979), were made resistant to EPI by stepwise increase of the drug concentration in the culture medium (Hall et al., 1991). The resistant cells (HB8065/R) and the sensitive parental cells (HB8065/S) were maintained and propagated as previously described (Lehne et al., 1994).

**Flow cytometry: immunofluorescence assay of Pgp expression**

Specific immunofluorescence was obtained by a three-layer staining technique. Cell suspensions were washed with phosphate-buffered saline (PBS)/bovine serum albumin (BSA) and incubated on ice for 60 min with MRK16 (25 μg ml⁻¹) or mouse IgG2a (25 μg ml⁻¹) in PBS/BSA. The second and third layer-staining protocols were carried out with 100 μl of biotinylated horse anti-mouse IgG (1:35 dilution in PBS/BSA) and 100 μl of fluorescein isothiocyanate (FITC)-c conjugated streptavidin (1:35 dilution in PBS/BSA) for 20–30 min each, with one PBS/BSA wash between them. Immunofluorescence distributions were generated using a FACScan flow cytometer (Becton Dickinson, San José, CA, USA) with a 15 mW argon ion laser tuned to 488 nm. FITC fluorescence of gated populations was collected through a bandpass filter (FL1; bandwidth 515–545 nm). Calculations
of logarithmically amplified fluorescence values were performed in arithmetic mode using the LYSIS (Becton Dickinson) computer program.

Cell growth inhibition assay
Approximately 50 x 10^3 cells were plated in 16 mm-diameter wells (Costar, Cambridge, MA, USA) and grown in 1 ml of drug-free medium (EMEM) for the first 24 h. The wells were then supplemented with the appropriate anthracycline at certain dose levels either alone or together with SDZ PSC 833 (1.5 µg ml^-1). Six replicate cultures were made from each of three dose levels and from untreated controls from both cell lines (HB8065/R and HB8065/S). After 72 h the cells were harvested by trypsinisation and counted in a Coulter Counter ZM (Coulter Electronics, Luton, UK). The dose level required for 50% inhibition of cell growth (GI50) was calculated from linear plots of dose vs cell number. The resistance factor (RF) was defined as the ratio between the GI50 values obtained in HB8065/R and HB8065/S cells. Modulation of growth inhibition was assessed by co-incubation with SDZ PSC 833. The modulating factor (MF) was defined as the ratio between the GI50 values of non-modified and modified HB8065/R cells.

Flow cytometry: intracellular accumulation of anthracyclines
The cells were grown in drug-free medium for 24 h before flow cytometric analysis. FACScan (Becton Dickinson) and Ortho Cytofluorograf 50-H (Ortho Diagnostic Systems, MA, USA) laser flow cytometers, both tuned to 488 nm laser excitation wavelength and running at 15 mW and 250 mW respectively, were used to generate anthracycline fluorescence. In the FACScan, fluorescence was transmitted through a bandpass filter of 564–606 nm (FL2) and logarithmically amplified. In the Cytofluorograf, fluorescence emitted above 510 nm was collected on log scale and transformed to linear values (Lehne et al., 1995). Correlated forward angle (a relative measure of cell size) and right angle (a measure of cell granularity) light scatter measurements were generated to exclude dead cells and debris from analyses. Acquired data from 3000–10 000 events were analysed using LYSIS (Becton Dickinson) and MULTI2D (Phoenix Flow Systems, San Diego, CA, USA) computer programs. Changes in the fluorescence intensity of EPI, DOX, DNR and ACL were recorded by repeated measurements during incubation with 1–10 µg ml^-1 of the drugs at 37°C for up to 180 min. Co-incubation with verapamil (5 µg ml^-1) and PSC 833 (1.5 µg ml^-1) was carried out to study any potential modification of anthracycline membrane transport.

Cellular efflux of anthracyclines
Approximately 200 x 10^3 cells were plated in 16 mm-diameter wells (Costar) and grown in 1 ml of medium (EMEM) for 6–7 days. The selection pressure of EPI (125 ng ml^-1) was maintained in the HB8065/R culture until 24 before the experiment, at which point they were grown in drug-free medium. The cells were then incubated in media containing 5 µg ml^-1 of either EPI, DOX, DNR or ACL for 90 min at 4°C. The temperature was kept low in order to load the cells with the drug at conditions which minimise active drug transport by Pgp. Afterwards, the cells were washed once and incubated at 37°C for 120 min in Hank's balanced salt solution 1X (HBSS) without phenol red (Bio Whittaker, Walkersville, MA, USA). After incubation, samples of HBSS were withdrawn from each well and placed in a Hitachi F-4500 fluorescence spectrophotometer (Nissei Sangyo, Tokyo, Japan). In series of 4–6 parallels, the drug fluorescence from each anthracycline was measured using 488 nm excitation and 594 nm emission wavelengths. Half of the cell samples were co-incubated with SDZ PSC 833 (1.5 µg ml^-1) at 37°C to assess possible modification of drug efflux.

Confocal laser scanning microscopy
Suspensions of HB8065/R and HB8065/S cells were incubated with anthracyclines (5–10 µg ml^-1) for 60 min at 37°C. Viable cell suspensions were examined in a Nikon Labophot microscope (Nikon, Tokyo, Japan) with an epifluorescence attachment and equipped with a Bio-Rad MRC 600 confocal laser scan unit with a krypton/argon laser using the 488 nm line, a K1 double dichroic excitation filter block and a K2 dichroic emission filter block (Biorad Microscience, Hertfordshire, UK). A Polaroid freeze frame unit (Polaroid, Cambridge, MA, USA) was attached for photographic documentation of acquired images. The confocal microscopic images were taken at ×72 objective magnification.

Results
Scanning fluorimetry
The native fluorescence of the anthracyclines allowed photodetection of these drugs when excited with light of certain wavelengths. Using scanning fluorimetry, the excitation and emission maxima were 490 nm and 590 nm for EPI and DOX respectively, 490 nm and 594 nm for DNR, and 440 nm and 520 nm for ACL. The argon ion lasers in the flow cytometers and in the confocal laser scanning microscope delivered coherent excitation light of 488 nm wavelength, which yielded almost 100% of maximal fluorescence for EPI, DOX and DNR and approximately 50–75% of maximal fluorescence for ACL at the emission wavelengths collected.

Pgp expression
The distributions of Pgp expression in HB8065/R and HB8065/S cells were determined by flow cytometric immuno-fluorescence detection using the anti-Pgp monoclonal antibody MRK16. A low expression of Pgp was seen in the HB8096/S cells compared with the 12-fold higher expression in the HB8065/R cells in terms of Pgp-specific immunofluorescence (Figure 2).
Cell growth inhibition

The HB8065/S and HB8065/R cell lines also differed with respect to susceptibility to anthracyclines, using growth inhibition as a measure of cytotoxicity. After continuous treatment of HB8065/R cells with EPI, DOX, DNR and ACL for 3 days, the GI_{50} values were 2118 ng ml^{-1}, 1784 ng ml^{-1}, 362 ng ml^{-1} and 38 ng ml^{-1} respectively. The corresponding GI_{50} values for HB8065/S cells were 46 ng ml^{-1}, 39 ng ml^{-1}, 40 ng ml^{-1} and 29 ng ml^{-1}, which resulted in RF values of 46, 46, 9 and 1 respectively (Figure 3). Hence, the HB8065/R cells were sensitive to ACL but resistant to EPI, DOX and DNR.

To test if the variation in anthracycline resistance was correlated with Pgp activity, growth inhibition studies were carried out in the presence of SDZ PSC 833, which is a potent modifier of Pgp-mediated MDR (Boesch et al., 1991). After treatment with SDZ PSC 833, the GI_{50} of HB8065/R increased by a factor of 46 (modifying factor, MF) for EPI (GI_{50} untreated, 1878 ng ml^{-1}; GI_{50} treated, 41 ng ml^{-1}), 45 for DOX (GI_{50} untreated, 1967 ng ml^{-1}; GI_{50} treated, 44 ng ml^{-1}) and 11 for DNR (GI_{50} untreated, 375 ng ml^{-1}; GI_{50} treated, 34 ng ml^{-1}) (Figure 4a–c). There was practically no change in GI_{50} for ACL (GI_{50} untreated, 43 ng ml^{-1}; GI_{50} treated, 36 ng ml^{-1}) (Figure 4d). Thus, the resistance to EPI, DOX and DNR was eliminated by SDZ PSC 833, while there was no change in the cytotoxicity of ACL. Even when the treatment duration was reduced from 3 days to 3 h, the sensitivity of HB8065/R cells to ACL persisted, the RF values being 1 for ACL (GI_{50} HB8065/S, 356 ng ml^{-1}; GI_{50} HB8065/R, 467 ng ml^{-1}), 20 for EPI (GI_{50} HB8065/S, 250 ng ml^{-1}; GI_{50} HB8065/R, 5000 ng ml^{-1}), 17 for DOX (GI_{50} HB8065/S, 286 ng ml^{-1}; GI_{50} HB8065/R, 4900 ng ml^{-1}) and 23 for DNR (GI_{50} HB8065/S, 184 ng ml^{-1}; GI_{50} HB8065/R, 4295 ng ml^{-1}). SDZ PSC had absolutely no effect on the cytotoxicity of EPI or ACL in the sensitive HB8065/S cells and the growth curves of modified and non-modified cells were practically identical (data not shown).

**Figure 3** The growth inhibition curves for 72 h continuous treatment with EPI (a), DOX (b), DNR (c) and ACL (d) show the correlations between the anthracycline dose levels and the relative cell numbers of HB8065/S cells (□) and HB8065/R cells (△). The error bars show the 95% confidence intervals. The calculated resistance factors (RF) are given in the upper right corner of each diagram.

**Figure 4** The growth inhibition curves for 24 h continuous treatment with Aclarubicin (Aj) and Aclarubicin + SDZ PSC 833 (A(a)). The lower sensitivity of HB8065/R to Aclarubicin in the presence of SDZ PSC 833 (A(a)) is shown. The red line (RF = 9) represents the growth inhibition of HB8065/R treated with Aclarubicin alone, while the blue line (RF = 1) represents the growth inhibition of HB8065/R treated with Aclarubicin + SDZ PSC 833. The error bars show the 95% confidence intervals. The calculated resistance factors (RF) are given in the upper right corner of each diagram.

**Figure 5** The growth inhibition curves for 24 h treatment with EPI (A), DOX (B), DNR (C) and ACL (D) show the correlations between the anthracycline dose levels and the relative cell numbers of HB8065/S cells (□) and HB8065/R cells (△). The error bars show the 95% confidence intervals. The calculated resistance factors (RF) are given in the upper right corner of each diagram.
appropriate drug. Cells incubated with EPI or DOX (10 μg ml⁻¹) demonstrated weakly detectable fluorescence in the nuclear membrane and the chromoplasm of HB8065/R cells, whereas a strikingly brighter fluorescence with similar distribution was seen in the majority of HB8065/S cells. Fluorescent vesicles were seen in the cytoplasm of both cell types (micrographs not shown). Cells incubated with DNR (5 μg ml⁻¹) showed predominantly cytoplasmic fluorescence, both diffuse and vesicular, and the fluorescence was clearly brighter in the HB8065/S cells (Figure 5a) than in the HB8065/R cells (Figure 5b). On the other hand, indiscriminately bright cytoplasmic fluorescence of ACL (5 μg ml⁻¹), with a mixed diffuse and vesicular distribution, was seen in both cell types (Figure 5c and d). There was no detectable nuclear fluorescence of ACL.

**Cellular accumulation of anthracyclines**

The intracellular drug fluorescence was quantified by flow cytometry after 60 min of incubation and correlated by linear regression with the drug amount added to the medium. The drug fluorescence increased linearly with increasing concentration of all anthracyclines in the medium (range 0.999–1.000) within a dose range of 1–10 μg ml⁻¹ in both HB8065/R and HB8065/S cells (Figure 6a–d). Assuming that the slope of the regression line represents a measure of intracellular drug retention, the HB8065/S cells accumulate 2.6 times more EPI, 2.3 times more DNR and 1.6 times more DOX than HB8065/R cells. In contrast, incubation with ACL for 1 h led to 1.7 times higher intracellular levels in the HB8065/R cells, despite the relatively high expression of Pgp in these cells compared with the parental HB8065/S cells. The total fluorescence from each anthracycline differed in intensity, and at a drug concentration of 5 μg ml⁻¹ the fluorescence intensity of DNR was 4.3-fold that of ACL, 3.6-fold that of DOX and 3.5-fold that of EPI in HB8065/R cells.

Individual profiles of drug fluorescence in resistant (R) and sensitive (S) hepatoma cells during 3 h incubation with each anthracycline are presented in Figure 7. Flow cytometric measurements revealed a rapid influx of all four anthracyclines in both HB8065/R and HB8065/S cells. The rate of anthracycline accumulation appeared to be similar for the two cell types in the first few minutes of incubation. After 15–30 min, the accumulation rate in the HB8065/R
cells subsided, as the fluorescence of the anthracyclines reached a plateau at this stage. In HB8065/S cells, the fluorescence of EPI and DOX continued to increase for 45 and 150 min, respectively, whereas the increase in fluorescence of DNR and ACL had already subsided within 15 min of incubation. Thereafter, only a slight increase in drug accumulation continued throughout the incubation period. After 3 h, the sensitive cells had accumulated EPI,
DOX and DNR at intracellular levels of 1.9, 1.7 and 2.1 times those in the resistant cells. On the other hand, the resistant cells accumulated 1.4 times more ACL than the sensitive cells.

Modification of membrane transport

The resistant HB8065/R cells were incubated for 90 min with 5 μg ml⁻¹ of each anthracycline alone or in combination with verapamil (5 μg ml⁻¹) or SDZ PSC 833 (1.5 μg ml⁻¹). The two modifying agents were added to the cells after 30 min and pronounced changes in drug fluorescence appeared immediately (Figure 8). The addition of VPL and SDZ PSC 833 resulted in 2- to 3-fold enhancement of the cellular fluorescence of DNR, DOX and EPI. In contrast, essentially no change was seen in the fluorescence of ACL after treatment with either VPL or SDZ PSC 833.

If the changes in intracellular drug fluorescence were induced by interaction between the modifying agent and Pgp, a corresponding change in drug efflux should occur as well. The results of the drug efflux studies are summarised in Table I. We measured significant reductions of effuxed EPI, DOX and DNR from HB8065/R cells after treatment with SDZ PSC 833 (1.5 μg ml⁻¹), but no change was seen in the efflux of ACL. On the other hand, identical treatment of the sensitive HB8065/S cells was not accompanied by significant changes in the efflux of any of the anthracyclines.

Discussion

Several mechanisms of MDR have been described in murine and human cancer cell lines. Apparently, the most common form of MDR is Pgp-mediated increased drug efflux resulting in decreased intracellular drug concentrations. Pgp has a broad specificity to multiple hydrophobic xenobiotics (reviewed by Licht et al., 1994), and many anthracyclines have been shown to be substrates of Pgp (Mulder et al., 1995). In tumour cell lines, non-Pgp-mediated resistance to anthracyclines has been linked to reduced activity of topoisomerase II (Cole et al., 1991; Eijdems et al., 1995), increased activity of glutathione S-transferase (Batist et al., 1986) and reduced activity of cytochrome P450 reductase (Mimaugh et al., 1989). Recently, a new multidrug efflux transporter, multidrug resistance protein (MRP), has been identified (Cole et al., 1992), and overexpression of Pgp and MRP may coexist (Brock et al., 1995).

Our results show that the HB8065/R cells remain sensitive to ACL despite a pronounced overexpression of Pgp, and
that ACL accumulates unabated in these cells. The Pgp-modifying agents used in this study did not influence the cytotoxicity, accumulation or efflux of ACL. On the other hand, treatment of HB8065/R cells with SDZ PSC 833 increased the accumulation and decreased the efflux of EPI, DOX and DNR, resulting in multiplied sensitivity, which became equal to that of the parental HB8065/S cells. Thus, Pgp seemed to be the key determinant to explain the difference in cytotoxicity and cellular pharmacokinetics seen in the four anthracyclines. Our findings strongly indicate that ACL is a poor substrate for active outward transport by Pgp, which is in agreement with previous reports of MDR circumvention with 9-alkyl or morpholinyl substituted anthracyclines in cell lines (Scott et al., 1986; Streeter et al., 1986; Coley et al., 1990). Two recent reports suggest that ACL may also circumvent drug resistance due to altered expression of topoisomerase II and glutathione S-transferase (Jensen et al., 1993; Okuyama et al., 1994).

The assessment of cytotoxicity was performed using growth inhibition assays that measure cell growth by electronic cell counting. Previous attempts to perform colony-forming assays in HB8065/R and HB8065/S cells have been unsuccessful because both lines failed to clone reproducibly on soft agar (Hall et al., 1991). Although electronic assessment of growth inhibition does not discriminate between cells of poor or of good proliferating capacity, the results from growth inhibition assays of anthracycline toxicity have been shown to correlate with those of clonogenic assays in certain pairs of parental and MDR cell lines (Bhalla et al., 1985; Hall et al., 1991). Following a 3 h treatment course the growth inhibition assay revealed a 20-fold resistance to EPI, DOX and DNR in the resistant human hepatoma cell line relative to the parental line, which corresponded with approximately 2-fold lower intracellular drug accumulation in the resistant cells than in the parental cells. By extending the treatment duration to 3 days, the resistance to EPI and DOX was doubled and the resistance to DNR was halved. Thus, the dose–response relationship was schedule dependent. Correspondingly, it has previously been demonstrated that cell killing by doxorubicin is an exponential function of drug exposure time in Chinese hamster ovary cells (Bates et al., 1985), and that lengthening the drug exposure time reduces the relative resistance to vincristine in two human colon carcinoma cell lines (Bates et al., 1994).

Using flow cytometry, we demonstrated rapid accumulation of all the anthracyclines in both cell types within minutes of incubation, which is in agreement with previous in vitro and in vivo findings of rapid influxes of anthracyclines across cellular membranes (Meriwether & Bachur, 1972; Egorin et al., 1974; Bachur, 1976). In both hepatoma cell lines, the fluorescence of DNR and ACL reached a plateau within 15 min. It was previously shown that ACL accumulates more rapidly than DNR or DOX in L1210 ascitic cells (Zenebergh et al., 1982), and that the rate of cellular uptake increases by increasing lipophilicity of the anthracyclines (Skovsgaard, 1987; Wheeler and Kessel, 1980). EPI and DOX reached a plateau within 15 min in the resistant cells but continued to accumulate for 45–150 min in the parental cells. Thus, the drug accumulation was not only governed by physicochemical characteristics of the drugs, but also by cellular characteristics.

The confocal laser microscopic images confirmed the flow cytometric findings by showing less fluorescent brightness of EPI, DOX and DNR in the resistant cells than in the sensitive ones, and by showing no difference in fluorescent brightness of ACL in the two cell types. It has been demonstrated by others that the fluorescence intensity of intracellular DNR parallels measurements of uptake of the corresponding radioactive drug (Bhalla et al., 1985). However, the intracellular anthracycline fluorescence does not accurately reflect the drug content because binding to DNA may cause quenching of the fluorescence, which is particularly pronounced for ACL (Skovsgaard, 1987;
cytoplasmic in cells, drug (Millot et al., 1989). The fluorescence intensity of free ACL has been measured to be 200 times that of nucleus-bound drug (Millot et al., 1989). In the cytoplasm of the HB8065/S cells, and to a much lesser extent in the resistant HB8065/R cells, the anthracycline fluorescence appeared in numerous cytoplasmic vesicles. Similar vesicles have been demonstrated in both MDR cell lines and corresponding parental lines (Weaver et al., 1991), and may represent drug accumulation in lysosomes. It is possible that higher drug concentrations may activate non-specific adsorptive endocytosis, which has been demonstrated in anthracycline-resistant Ehrlich ascites tumour cell lines (Sehested et al., 1987).

Our results showed that low accumulation of EPI and DOX paralleled high efflux of these cytotoxics in the pair of cell lines studied. Because SDZ PSC 833 significantly reduced the efflux of EPI, DOX and DNR from the resistant human hepatoma cells but not from the sensitive ones, it is likely that Pgp was mainly responsible for the variations in the efflux of these anthracyclines. The efflux of ACL was not altered by SDZ PSC 833 in any of the cell types, and the greater efflux of ACL from the resistant human hepatoma cells was obviously unrelated to Pgp, but correlated with an increased accumulation of ACL in the resistant cells. Interestingly, it has been reported that there is no difference in the efflux of ACL between parental and resistant lines of human small-cell lung cancer and mouse mammary tumour (Coley et al., 1993). Thus, there may exist alternative transport mechanisms for ACL in certain cell types that may also express high levels of Pgp.

The applied anthracycline doses in this study were in the upper range or slightly above achievable plasma concentrations in patients after standard bolus i.v. injections (Martini et al., 1984; Paul et al., 1989). An i.v. bolus injection of 60–120 mg m⁻² ACL yields initially 2–3 nmol l⁻¹ (Egorin et al.,

Table 1 Modification of drug efflux by SDZ PSC 833 in HB8065/R and HB8065/S cells. The modified drug efflux was compared with the natural efflux by means of independent samples t-test

| Drug efflux (ng ml⁻¹) (No modifier) | Drug efflux (ng ml⁻¹) SDZ PSC 833 | P    |
|------------------------------------|-----------------------------------|------|
| HB8065/R                          |                                   |      |
| EPI                               | 350 ± 2                           | 235 ± 18 | 0.001 |
| DOX                               | 154 ± 11                          | 80 ± 3  | 0.003 |
| DNR                               | 712 ± 36                          | 515 ± 37 | 0.009 |
| ACL                               | 2554 ± 80                         | 2472 ± 87 | NS   |
| HB8065/S                          |                                   |      |
| EPI                               | 183 ± 12                          | 173 ± 8 | NS    |
| DOX                               | 66 ± 4                            | 46 ± 11 | NS    |
| DNR                               | 719 ± 35                          | 682 ± 20 | NS   |
| ACL                               | 1065 ± 72                         | 1023 ± 62 | NS  |

NS, not significant.

Tarasiuk et al., 1989). The fluorescence intensity of free ACL has been measured to be 200 times that of nucleus-bound drug (Millot et al., 1989). In the cytoplasm of the HB8065/S cells, and to a much lesser extent in the resistant HB8065/R cells, the anthracycline fluorescence appeared in numerous cytoplasmic vesicles. Similar vesicles have been demonstrated in both MDR cell lines and corresponding parental lines (Weaver et al., 1991), and may represent drug accumulation in lysosomes. It is possible that higher drug concentrations may activate non-specific adsorptive endocytosis, which has been demonstrated in anthracycline-resistant Ehrlich ascites tumour cell lines (Sehested et al., 1987).

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Figure 8 Changes in the intracellular drug fluorescence HB8065/R cells during 180 min of incubation with EPI (a), DOX (b), DNR (c) and ACL (d). The solid line represents cells treated with anthracycline alone (☐), and the dashed line represents anthracycline-treated cells modified with either SDZ PSC 833 (●) or verapamil (▲). The time points for addition of anthracyclines and modifiers are indicated by open and closed arrows respectively, (FACScan analysis).
demonstrated between ACL and the type I anthracyclines EPI, DOX and DNR. Therefore, we may conclude that the 9-alkyl trisaccharide ACL appears to be a poor substrate of Pgp in cultured human hepatoma cells. Translation of this unique feature of ACL into clinical practice would mean that treatment of unresectable hepatocellular carcinoma may benefit from tailoring the chemotherapeutic regimens according to the Pgp status of the tumour.

Acknowledgements

The authors are grateful to Dr Henrik S Huitfeldt for valuable help with the laser confocal microscopy and to Karen Johanne Beckstrom, Reidun Hauge and May Ellen Lauritsen for excellent technical assistance. This work was financially supported by Medinnova, the Norwegian Cancer Society and the Research Council of Norway.

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