Cisplatin [cis-diaminedichloroplatinum (II); DDP] is an electrophilic platinum coordinate compound which causes DNA damage by forming platinum–DNA coordination complexes (Zwelling & Kohn, 1980). It is one of the most effective antineoplastic agents clinically used against human small-cell lung cancer (SCLC) (Aisner, 1988). However, its clinical efficacy as an antineoplastic agent is curtailed in many cancers owing to inherent or acquired resistance to its cytotoxicity (Loecher & Einhorn, 1984). Attempts are currently under way to improve the efficacy of DDP by using non-cytotoxic drugs to defeat cellular defence mechanisms that participate in mediating resistance to DDP (Timmer-Bosscha et al., 1992). Cellular defence mechanisms that protect cells from DDP-induced DNA damage (Kelley & Rozenweig, 1989; Perez et al., 1990; Timmer-Bosscha et al., 1992) can be conceptualised as those that repair DDP-induced DNA damage (Teicher et al., 1987; Masuda et al., 1990), those that decrease the accumulation of DDP into the cell (Teicher et al., 1987; Andrews et al., 1988; Bungo et al., 1990) or those that increase the capacity of cytoplasmic defence mechanisms which interfere with the ability of electrophilic toxins, such as DDP, to interact with DNA (Meijer et al., 1990; Kasahara et al., 1991; Muller et al., 1991). Glutathione (GSH), the chief intracellular nucleophilic, function as a scavenger of electrophilic toxins, and its cellular concentrations have been shown to be increased in some DDP-resistant cell lines (Batist et al., 1986; Meijer et al., 1990; Mistry et al., 1991). Glutathione S-transferases (GSTs) are multifunctional cellular GSH enzymes which can detoxify electrophiles by conjugating them with GSH (Jakoby, 1978). Increased GST activity has been linked with DDP resistance (Teicher et al., 1987; Saburi et al., 1989; Miyazaki et al., 1990; Puchalski & Fahl, 1990; Sharma et al., 1993). Because GSTs are susceptible to inhibition by a number of non-cytotoxic drugs (van Bladeren & van Ommen, 1991), they are an attractive target for attempts to enhance DDP efficacy using non-cytotoxic inhibitors, such as ethacrynic acid (a non-cytotoxic diuretic drug). Ethacrynic acid has been shown to enhance the cytotoxic effects of certain alkylating agents (Tew et al., 1988). However, the GST-π isoenzyme, which has been linked with malignant transformation and with resistance to alkylating agents (Tsuchida & Sato, 1992), has been suggested to decrease the efficacy of ethacrynic acid in enhancing alkylating agent cytotoxicity (Kuzmich et al., 1992). Furthermore, ethacrynic acid has been found not to enhance the cytotoxicity of DDP towards resistant malignant cell lines (Plumb et al., 1990).

During studies aimed at finding alternative non-cytotoxic GST inhibitors, we have found that sulphasalazine (SS), a drug commonly used for the treatment of inflammatory bowel disorders, is an effective inhibitor of GSTs including GST-π (Ahmad et al., 1992). We have also studied electrophile defence mechanisms, including GSH levels, and activities of enzymes which participate in detoxification of electrophilic toxins, including GSTs, glutathione peroxidase (GPx), glutathione reductase (GR) and glucose-6-phosphate dehydrogenase (G6PD), and have found that among these defence mechanisms GST activity correlated best with degree of DDP resistance in these cell lines (Sharma et al., 1993). The present studies were designed to test the role of GSTs in mediating DDP resistance in two human SCLC cell lines, NCI H-69 and H-2496, in which GST activities parallel DDP resistance, by studying the effect of SS on their GSTs and sensitivity to DDP. The inhibitory effects of SS of GST activity were studied using GSTs purified from these cell lines. A spectrophotometric cytotoxicity assay (Carmichael et al., 1987; Twentyman & Luscombe, 1987) which utilises 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) was used in conjunction with isobologram analysis (Steel & Peckham, 1979) to determine whether SS is able to enhance the cytotoxicity of DDP in a synergistic manner. The major GST isoenzyme in both cell lines was found to be GST-π. SS was found to be a competitive inhibitor of GST-π at concentrations which can be achieved in human serum with its conventional doses (Das & Dubin, 1976). SS was found to enhance the cytotoxicity of DDP in a synergistic manner in both cell lines, but the degree of synergy was considerably greater towards the H-69 cell line, which was relatively more resistant to DDP and which had a higher GST-π content than the H-2496 cell line. These results support the results of our previous studies (Sharma et al., 1993) which have implicated GST-π as a significant determinant of DDP resistance and suggest that SS could be clinically used to enhance the anti-tumour efficacy of DDP in malignant cells overexpressing GST-π.
SULPHASALAZINE MODULATES CISPLATIN CYTOTOXICITY

Materials and methods

Reagents and chemicals

Reagents including SS and 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT), 1-chloro-2,4-dinitrobenzene (CDNB), 5,5-dithiobis-2-nitrobenzoic acid (DTNB), dimethylsulphoxide (DMSO), diethylpyrocarbonate, formamide, guanidinium thiocyanate and other reagents were purchased from Sigma (St Louis, MO, USA). A random primed labelling kit was purchased from Boehringer-Mannheim (Indianapolis, IN, USA). Radiolabelled [γ-32P]CTP (specific activity 6 × 10⁶ c.p.m. μg⁻¹) was purchased from DuPont NEN Products (Billerica, MA, USA). Zetabind nylon membrane was purchased from Cuno Laboratory (Meriden, CT, USA). DDP was obtained from Bristol Laboratories (Evansville, IN, USA). Fetal bovine serum (FBS) was purchased from Intergen Company (Purchase, NY, USA). Cell culture supplies including RPMI-1640 medium, penicillin-streptomycin (P/S) solution and Dulbecco's phosphate-buffered saline (PBS) were purchased from Gibco Laboratories (Grand Island, NY, USA).

Culture conditions

Two human SCLC cell lines, NCI H-69 and NCI H-2496, were generous gifts from H. Oie at the National Cancer Institute, Baltimore, MD, USA. These SCLC cell lines, which had never been exposed to chemotherapy drugs in vitro, grew in suspension cultures in RPMI-1640 medium containing 10% FBS and 1% P/S at 37°C in a 5% carbon dioxide atmosphere. Cells were maintained in the log phase of growth by diluting them 1:3 with medium every 2–3 days.

Non-protein sulphydryl content of cell lines

Cells growing in log phase were washed with PBS and homogenate prepared by sonication in 10 mM potassium phosphate buffer, pH 7.0. Non-protein sulphydryl (NPSH) content in the acid-soluble fraction of the homogenate was determined spectrophotometrically using DTNB (Beutler et al., 1963).

GST activity, purification, Western blot analysis and kinetics of inhibition by sulphasalazine

GST activity towards CDNB was determined (Habig et al., 1974) on 28,000 g supernatants of the homogenate prepared as described above. One unit of GST activity was defined as 1 μmol of dinitrophenyl-S-glutathione (Dnp-SG, ε₂₅⁰ = 9.6 mM⁻¹ cm⁻¹) formed per minute at 25°C. GSTs were purified from cell homogenate of both cell lines using GSH affinity chromatography as used by us previously (Sharma et al., 1993). Western blot analysis on the purified GSTs was performed (Towbin et al., 1979) using polyclonal antibodies specific for the α, μ and π classes of GST. To determine a dose–response curve for inhibition of GST, purified enzyme was incubated with varying concentrations of SS followed by measurement of initial rate of Dnp-SG formation using spectrophotometrically. Two concentrations of SS (one above and one below the concentration which caused 50% inhibition of GST activity) were used with varying concentrations of CDNB to determine initial velocity for generating double reciprocal plots. Vₘₐₓ and Kₘ for CDNB and nature of inhibition of GST by SS were determined from these plots. Replots of the slopes from the double reciprocal plots were used to determine Kᵢ.

Northern blot analysis of GST RNA expression in cell lines

cDNA probes for the α, μ and π class GSTs were prepared by restriction endonuclease digestion of plasmids containing cDNA clones of the α-class GST Ha-1 (a gift from C.P.D. Tu, Department of Molecular and Cell Biology, The Pennsylvania State University, University Park, USA), the μ-class GST a gift from M. Muramatsu, Department of Biochemistry, University of Tokyo, Japan) and the π-class GST (a gift from I. Listowsky, Department of Biochemistry, Albert Einstein Medical College, New York, USA). The cDNA fragments were purified by a method utilising low melting temperature agarose gel electrophoresis, followed by ethanol precipitation (see Luscombe & Peckham, 1979). The probes were labelled by the random primed labelling method according to the manufacturer's (Boehringer-Mannheim) recommendations using [γ-32P]CTP, and the unincorporated nucleotides were removed by filtration through Millipore Ultra Free MC filter units [10,000 nominal molecular weight limit (NMWL)]. RNA was isolated from 4 × 10⁶ cells by acid guanidinium thiocyanate–phenol–chloroform extraction (Chomczynski & Sacchi, 1987) and the concentration and purity were determined by absorbance at 260 and 280 nm. A 25 μg aliquot of the isolated RNA from both cell lines was subjected to formaldehyde–agarose gel electrophoresis in 1.2% agarose gel at 35 V overnight and transblotted to Zetabind nylon membrane. Nylon filters were prehybridised overnight at 50°C in hybridisation buffer (0.2 M sodium phosphate pH 7.2, 1% bovine serum albumin, 7% SDS, 1 mM EDTA and 15% formamide) and hybridised in the same buffer with the corresponding [32P]-labelled DNA probes at 50°C overnight. Filters were washed once at 50°C for 25 min, and twice at room temperature for 5 min each in washing buffer (40 mM sodium phosphate pH 7.2, 1 mM EDTA and 1% SDS). Filters were exposed to Kodak XAR-5 X-ray film at −80°C using intensifying screens.

Determination of IC₅₀ for cisplatin and SS

A modified MTT cytotoxicity assay similar to that previously described (Carmichael et al., 1987; Twentyman & Luscombe, 1987) was standardised to determine the numbers of viable cells after cisplatin and/or SS treatments. Briefly, aliquots containing approximately 5,000 cells were taken from flasks containing cells in log phase growth at a density of 3–5 × 10⁶ cells ml⁻¹ and inoculated into 96-well plates. The cells were diluted with medium containing various concentrations of DDP and/or SS. After 72 h, 20 μl of MTT (2 mg ml⁻¹) was added to each well followed by incubation for 1 h at 37°C. The cells were centrifuged in Eppendorf tubes at 28,000 g for 10 min. The supernatant was discarded and the pellet thoroughly solubilised in 50 μl of DMSO, followed by addition of 1 ml of isopropanol. The absorption of this extract was recorded at 560 nm using a Gilford Response spectrophotometer. Surviving cell number was identified using a standard curve of cell number versus absorbance at 560 nm. The IC₅₀ values for DDP and SS were determined from a plot of percentage surviving cells (compared with control cells) versus drug concentrations. All cytotoxicity experiments were conducted four times, each with triplicate determinations, and the average values and standard deviations were used to construct log survival curves and isobolograms.

Isobologram analysis

Isobolograms and envelopes of additivity were constructed according to the previously described method (Steel & Peckham, 1979). Briefly, the IC₅₀ values obtained for DDP and SS alone were plotted on a Cartesian plane at equal distances from the origin on the x- and y-axes respectively. The line connecting these points was designated the theoretical isotoxic dose line, or the combination of DDP and SS concentrations at which 50% of cells would survive provided the log survival curves of both drugs were linear. However, this theoretical isotoxic dose line, which represents simple additivity of cytotoxicity of both drugs at any combination of concentrations, may not apply to non-linear log survival curves of one or both drugs. Since, the survival curve for SS was not linear, an envelope of additivity of DDP and SS was calculated from the cell survival data as described previously (Steel & Peckham, 1979). The IC₅₀ values of DDP determined
cytotoxic effects of additivity (SS).

at several fixed concentrations of SS (below the IC₅₀ value of SS) were plotted against the corresponding concentration of SS. A curve thus generated and lying below the envelope of additivity was taken as the evidence of synergy between the cytotoxic effects of two drugs.

Results and discussion

The average NPSH contents of the H-69 and H-2496 cell lines from three separate determinations were found to be 60.4 ± 5.8 and 44 ± 7.2 nmol mg⁻¹ protein respectively. One hour after exposure to 1 mM SS, average NPSH contents of the H-69 and H-2496 cell lines from three separate determinations (62.2 ± 6.4 and 41.0 ± 5.8 nmol mg⁻¹ protein respectively) were not significantly altered acutely by exposure to SS. These results are consistent with a previous report that SS is not conjugated with GSH (Das & Dubin, 1976) and suggest that any effect of SS on DDP sensitivity is not caused by SS-induced changes in NPSH. Western blot analysis of purified GSTs revealed the presence of only the GST-α isoenzyme in these cell lines (Figure 1). The results of Northern blot analysis revealed that only GST-π was present in both cell lines, and GST-α or -μ RNA was not detected (figures not presented). These results were consistent with previous studies on the composition of GST isoenzymes in human SCLC cell lines (Miyazaki et al., 1990). The average GST activity, determined in homogenates of the H-69 and H-2496 cell lines in three separate experiments (9 ± 1 μU mg⁻¹ protein respectively), indicated that the H-69 cell line contained 34-fold higher GST activity than the H-2496 cell line (Sharma et al., 1993). SS inhibited GSTs of both cell lines with 50% inhibition of activity towards CDN B as seen at 10 and 12 μM for the H-69 and H-2496 cell lines respectively (Figure 2). SS was found to be a competitive inhibitor of the GSTs isolated from both cell lines, with Kᵢ values of 6.5 and 7.9 μM for the H-69 and H-2496 cell lines respectively.

An MTT cytotoxicity assay was used to determine the sensitivity of the SCLC cell lines to DDP and sulphasalazine alone and in combination. This assay is based on the optical measurement of a formazan dye which is cleaved from MTT by mitochondrial dehydrogenase of actively respiring mitochondria and has been shown to correlate with results of colony-forming cytotoxicity assays in SCLC cell lines (Carmichael et al., 1987). The calibration curve of the MTT cytotoxicity assay revealed that absorbsances at 560 nm were linear in the range 0.05–1.0 with respect to cell number determined by counting disaggregated cells in a haemocytometer (data not presented). Using this assay, the H-69 cell line was found to be approximately 6-fold more resistant to DDP than the H-2496 cell line (average IC₅₀ values from four separate experiments of 25 ± 3.9 and 4.5 ± 1.0 μM for the H-69 and H-2496 lines respectively). The log survival curves for the H-69 and H-2496 cell lines for DDP (Figure 3a) and SS alone (Figure 3b) are presented. These survival curves were linear with respect to DDP concentrations, but non-linear with respect to SS concentrations, for both cell lines. On the basis of these results, the use of envelopes of additivity became necessary for the analysis of synergy in isobolograms rather than the theoretical isotoxic dose line, which represented simple additivity of cytotoxicity between the two drugs. Isobolograms showing the theoretical envelopes of additivity and the actual isotoxic dose curves for combinations of SS and DDP are shown for the H-69 (Figure 4a) and H-2496 (Figure 4b) cell lines respectively. The isotoxic dose curves for both cell lines were found to lie below the theoretical isotoxic dose line, indicating that SS was able to enhance the cytotoxicity of DDP in a synergistic manner. The degree of synergy, reflected in the degree of deviation from the envelope of additivity, was significantly greater for the more resistant H-69 cell line having higher GST activity than the H-2496 cell line. Greater synergy between SS and DDP in the H-69 cell line, which contains significantly higher GST activity than the DDP-sensitive H-2496 cell line, suggested that GST may play a prominent role in mediating DDP resistance. Since GSTs have not been
shown to catalyse the conjugation of GSH with DDP. Further studies are required to elucidate the mechanisms through which GSTs may participate in defence of cells towards DDP and define the mechanism of enhancement of DDP cytotoxicity by SS. In addition to GST inhibition, SS has been shown to affect the synthesis of leukotrienes and prostaglandins from arachidonic acid contained in plasma membranes (Tornhamre et al., 1989), but its effects on protein kinase C have not been reported. Since inhibitors of protein kinase C can potentiate DDP cytotoxicity (Timmer-Bosscha et al., 1992), future studies on the effect of sulphasalazine on protein kinase C activity may be helpful in delineating an additional mechanism for the observed enhancement of DDP cytotoxicity by SS.

Our studies suggest that SS should be added to the long list of agents which can enhance DDP cytotoxicity (Timmer-Bosscha et al., 1992). The marked degree of enhancement of DDP toxicity in the H-69 cell line at the lower concentrations of SS is interesting. Based on SS pharmacokinetics in humans, concentrations of SS up to 25 μM can be achieved in human plasma at relatively non-toxic doses of SS (Das & Dubin, 1976). Furthermore, animal pharmacokinetic studies of SS show that its concentrations in lung and connective tissues are significantly higher (in the range of 0.5 mM) than in serum, and they can persist for prolonged periods after bolus doses of SS (Hanngren et al., 1963). These findings can be helpful in designing chemotherapeutic regimens to test the efficacy of SS in modulating the anti-cancer efficacy of DDP in human malignancy, particularly lung cancer.

This work was supported in part by Grant GM-32304 awarded by the National Institute of General Medical Sciences (to Y.C.A.). S.A. wishes to express thanks to Don W. Powell, MD, Chair, Department of Internal Medicine, University of Texas Medical Branch, for providing funds for these studies. The secretarial assistance of Mrs Alicia Woods is acknowledged.
MISTRY, P., KELLAND, L.R., ABEL, G., SIDHAR, S. & HARRAP, K.R. (1991). The relationship between glutathione, glutathione S-transferase and cytotoxicity of platinum drugs and melfalan in eight human ovarian carcinoma cell lines. Br. J. Cancer, 64, 215–220.

MIYAZAKI, M., KOHNO, K., SABURI, Y., MATSUO, K., ONO, M., KUWANO, M., TSUCHIDA, S., SATO, K., SAKAI, M. & MURAMATSU, M. (1990). Drug resistance to cis-diaminedichloroplatinum (II) in Chinese hamster ovary cell lines transfected with glutathione S-transferase cDNA. Biochem. Biophys. Res. Commun., 166, 1358–1364.

MULLER, M.R., WRIGHT, K.A. & TWENTYMAN, P.R. (1991). Differential properties of cisplatin and tetraplatin with respect to cytotoxicity and perturbation of glutathione levels. Cancer Chemother. Pharm., 28, 273–276.

PEREZ, R.P., HAMILTON, T.C. & OZOLS, R.F. (1990). Resistance to alkylating agents and cisplatin: insights from ovarian carcinoma model systems. Pharmacol. Ther., 48, 19–27.

PLUMB, J.A., MILROY, R., BICKNELL, S.R. & KAY, S.B. (1990). Glutathione S-transferase, P-glycoprotein and drug resistance in small cell lung cancer cell lines. Proc. Am. Assoc. Cancer Res., 31, 369.

PUCHALSKI, R.B. & FAHIL, W.E. (1990). Expression of recombinant glutathione S-transferase x, Ys or Yh, confers resistance to alkylating agents. Proc. Natl Acad. Sci. USA, 87, 2443–2447.

SABURI, Y., NAKAGAWA, M., ONO, M., SAKAI, M., MURAMATSU, M., KOHNO, K. & KUWANO, M. (1989). Increased expression of glutathione S-transferase gene in cis-diammine-dichloroplatinum(II)-resistant variants of Chinese hamster ovary cell lines. Cancer Res., 49, 7020–7025.

SHARMA, R., SINGHAL, S.S., SRIVASTAVA, S.K., BAJPAI, K.K., FRENKEL, E.P. & AWASTHI, S. (1993). Glutathione and glutathione linked enzymes in human small cell lung cancer cell lines. Cancer Lett., 75, 171–177.

STEEL, G.G. & PECKHAM, M.J. (1979). Exploitable mechanisms in combined radiotherapy–chemotherapy: the concept of additivity. Int. J. Radiat. Oncol. Biol. Phys., 5, 85–91.

TEGER, B.A., HOLDEN, S.A., KI, O.M., SHEA, T.C., CUCIL, C.A., ROSOWSKY, A., HEINNER, W.D. & FREI, III, E. (1987). Characterization of a human squamous carcinoma cell line resistant to cis-diaminedichloroplatinum (II). Cancer Res., 47, 388–393.

TEW, K.D., BOMBER, A.M. & HOFFMAN, S.J. (1988). Ethylenic acid and parprost as enhancers of cytotoxicity in drug resistant and sensitive cell lines. Cancer Res., 48, 3622–3625.

TIMMER-BOSSCHA, H., MULDER, N.H. & DE VRIES, E.G.E. (1992). Modulation of cis-diaminedichloroplatinum (II) resistance: a review. Br. J. Cancer, 66, 227–238.

TORNHAMRE, S., EDENIUS, C., SMEDGARD, G., BIRGITTA, S. & LINDGREN, J.A. (1989). Effects of sulfasalazine and a sulfa-salazine analogue on the formation of lipooxygenase and cyclooxygenase products. Eur. J. Pharmacol., 169, 225–234.

TOWBIN, H., STAHELIN, T. & GORDON, J. (1979). Electrophoretic transfer of proteins from polyacrylamide gels to nitrocellulose sheets: procedure and some applications. Proc. Natl Acad. Sci. USA, 76, 4350–4354.

TSUCHIDA, S. & SATO, K. (1992). Glutathione transferases and cancer. Crit. Rev. Biochem. Mol. Biol., 27, 337–384.

TWENTYMAN, P.R. & LUSCOMBE, M. (1987). A study of some variables in a tetrazolium dye (MTT) based assay for cell growth and chemosensitivity. Br. J. Cancer, 56, 279–285.

VAN BLADEREN, P.J. & VAN OMMEN, B. (1991). The inhibition of glutathione S-transferases: mechanisms, toxic consequences and therapeutic benefits. Pharmacol. Ther., 51, 35–46.

ZWELLING, L.A. & KOHN, K.W. (1980). Effects of cisplatin on DNA and the possible relationships to cytotoxicity and mutagenicity in mammalian cells. In: Cisplatin: Current Status and New Developments. Prestayko, A., Crooke, S. & Carter, S. (eds) pp. 21–37. Academic Press: New York.