Inhibition of proliferation and induction of apoptosis in soft tissue sarcoma cells by interferon-α and retinoids

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Summary Uncontrolled proliferation and a defect of apoptosis constitute crucial elements in the development and progression of tumours. Among many other biological response modifiers known to influence these mechanisms, the efficacy of retinoids and interferons in the treatment of various malignant entities is currently matter of discussion. In the present study, we have investigated the effects of 9-cis-retinoic acid (9cRA), 13-cis-retinoic acid (13cRA), all-trans-retinoic acid (tRA) and interferon-α on proliferation and apoptosis of human soft tissue sarcoma (STS) cell lines HTB-82 (rhabdomyosarcoma), HTB-91 (fibrosarcoma), HTB-92 (liposarcoma), HTB-93 (synovial sarcoma) and HTB-94 (chondrosarcoma) in relation to p53 genotype as well as p53 expression. HTB-91, HTB-92 and HTB-94 STS cells exhibited mutant p53, whereas wild-type p53 was found in HTB-93 STS cells, and a normal p53 status in HTB-82 STS cells, carrying a silent point mutation only. Interferon-α, irrespective of p53 status, inhibited the proliferation of all five cell lines dose- and time-dependently. Similarly, 9cRA, 13cRA and IRA decreased the proliferation of HTB-82 and HTB-93 STS cells, whereas the proliferation of p53-mutated HTB-91, HTB-92 and HTB-94 STS cells remained unchanged. Furthermore, only 9cRA and IRA were capable of inducing apoptosis in HTB-82 and HTB-93 STS cells, whereas HTB-91, HTB-92 and HTB-94 STS cells did not undergo apoptosis under the influence of 9cRA or IRA. Retinoic acid receptor (RAR)-α and RAR-β mRNA were not detectable by Northern blot analysis in the five STS cell lines, whereas mRNA for the universal retinoic acid receptor, RAR-γ, was expressed in all STS cell lines indicating that retinoid resistance was not associated with a lack of RAR expression. Apoptosis was not induced by interferon-α or 13cRA in any of the five STS cell lines tested. Our results indicate that within the panel of tested STS cell lines, inhibition of proliferation and induction of apoptosis result from different mechanisms which differ in their dependence upon the presence of intact p53.

Keywords: apoptosis; soft tissue sarcoma; interferon-α; retinoids; p53

Soft tissue sarcomas (STS) represent a rare entity of malignant disorders of various histologies with mostly aggressive characteristics both locally and in the formation of distant metastases. In contrast to a variety of other malignancies and with exceptions reported in only a few trials and/or analyses (Gherlinzoni et al, 1986; Ravau et al, 1990; Antman, 1997; Sarcoma Meta-Analysis Collaboration, 1997), adjuvant chemotherapy has not shown to significantly improve relapse-free and/or overall survival in patients with STS. Similarly, chemotherapeutic interventions in metastatic STS were generally also disappointing (Borden et al, 1990; Antman et al, 1993). Thus, cure of early disease obviously depends upon appropriate, well-designed interdisciplinary therapy of the primary tumour, including surgery, radiotherapy and – in some cases (Rosenberg et al, 1982; Brodowicz et al, 1997; Sarcoma Meta-Analysis Collaboration, 1997) – cytotoxic chemotherapy.

Based upon the above considerations of an obviously limited efficacy of chemotherapeutic drugs upon the control of proliferation and/or induction of apoptosis in STS, alternate approaches were sought. Although the underlying reasons have not been clarified, certain STS cell lines have been demonstrated to be inhibited in their proliferation by retinoids (Gabbert et al, 1988; Crouch and Helman, 1991) and/or interferon-α (IFN-α; Rosso et al, 1992) in vitro and in vivo. In order to follow and further expand these early experiments, and put them into perspective with recent insights into the regulation of proliferation inhibition and generation of apoptosis on the molecular level, we have investigated the induction of these phenomena by a series of retinoids (9-cis-retinoic acid (9cRA), 13-cis-retinoic acid (13cRA), all-trans-retinoic acid (tRA)) and IFN-α in various human STS cell lines including HTB-82 (rhabdomyosarcoma), HTB-91 (fibrosarcoma), HTB-92 (liposarcoma), HTB-93 (synovial sarcoma) and HTB-94 (chondrosarcoma) in vitro and put them into context with p53 genotype as well as p53 protein expression. This was of particular interest, as growth arrest and induction of apoptosis resulting from appropriate chemo- and/or radiotherapeutic measures have been demonstrated to be dependent upon the intact function of the p53 gene in various models (Bergh et al, 1995; Elledge et al, 1995; Sarkis et al, 1995). Thus, patients with testicular tumours with regular p53 function have been shown to respond particularly well to chemotherapy (Lutzker and Levine, 1996), whereas patients with malignancies exhibiting high frequency of p53 mutations are known to present often with resistance to chemotherapeutic agents (Aas et al, 1996). In the present paper, we report that IFN-α induced inhibition of proliferation of all cell lines irrespective of...
their p53 status, whereas retinoid-induced apoptosis was dose- and time-related, depended upon the presence of regular p53 and differed between retinoid preparations and, finally, STS cell lines.

MATERIALS AND METHODS

Cell cultures

Cell lines were obtained from American Type Culture Collection (ATCC). HTB-82 (A-204, human rhabdomyosarcoma) were cultured in McCoy’s 5A medium supplemented with 10% heat-inactivated fetal calf serum (FCS; all from Gibco, Life Technologies Ltd, Paisley, UK), 50 U ml⁻¹ penicillin, 50 μg ml⁻¹ streptomycin and 2 mM L-glutamine (all from HyClone, Europe Ltd, Cramlington, UK). The human STS cell lines HTB-91 (fibrosarcoma, SW 684), HTB-92 (liposarcoma, SW 872), HTB-93 (synovial sarcoma, SW 982) and HTB-94 (chondrosarcoma, SW 1353) were cultured in Leibovitz’s L-15 medium with L-glutamine (PAA Laboratories Gmbh, Linz, Austria) supplemented with 10% heat-inactivated FCS (Gibco), 50 U ml⁻¹ penicillin and 50 μg ml⁻¹ streptomycin (all from HyClone) per ml. HTB-91, HTB-92, HTB-93 and HTB-94 were grown as monolayers (standard conditions) in T75 flasks (Falcon, Becton Dickinson, NJ, USA) at 37°C in a humidified atmosphere with free gas exchange without carbon dioxide (CO₂) by seeding 5 · 10⁶ cells in 25 ml of appropriate medium. HTB-82 cells were cultured in a humidified atmosphere containing 5%.

Chemicals

9cRA, 13cRA and tRA (Biomol Research Laboratories, Inc., Plymouth Meeting, PA, USA) were dissolved in 1 ml dimethyl sulphoxide (Sigma, St Louis, MO, USA) at a concentration of 8.3 · 10⁻³ M and then diluted with the appropriate culture medium (final concentrations: 10⁻³ M, 10⁻⁴ M and 10⁻⁵ M). Retinoid stock solution was kept at –20°C. Experiments with retinoids were performed in subdued yellow light. Then, 5 · 10⁶ IU IFN-α-2b (Schering-Plough NJ, USA) was dissolved in 1 ml of distilled water and subsequently diluted in appropriate culture medium (final concentrations: 25 U ml⁻¹, 50 U ml⁻¹, 100 U ml⁻¹ and 1000 U ml⁻¹).

DNA probes

Hybridization probes were isolated from plasmids containing: 0.6-kb PstI cDNA fragment from human retinoic acid receptor (RAR)-α (clone p63) in pTZ19R (ATCC); 0.6-kb EcoRI cDNA fragment from human RAR-β in pSG5 (kind gift from Dr P Chambon, INSERM, Illkirch, France); 1.3-kb EcoRI/Awel cDNA fragment from human RAR-γ in pSG5 (kind gift from Dr P Chambon); 1.3-kb EcoRI/HindIII cDNA fragment from rat glyceraldehyde 3-phosphate dehydrogenase (GAPDH) in pSP65. Isolation of inserts were performed as described previously (Harant et al, 1995).

Cell proliferation assay [(³H) Thymidine incorporation assay]

Tumour cells were plated in 96-well microtitre plates (Costar, Cambridge, MA, USA) at a density of 5 · 10⁵ cells per well. Subsequently, retinoid and/or IFN-α-2b preparations in varying concentrations were added to tumour cell lines (see above), which had adhered for 1 h, and subsequently cultured for 48, 72, 96 and 120 h at 37°C under the appropriate conditions (HTB-82 cells were cultured in a humidified atmosphere containing 5% CO₂, whereas all other cell lines were cultured in a humidified

| Cell line       | p53 Mutation* | Proliferation inhibition | Apoptosis induction |
|-----------------|---------------|--------------------------|---------------------|
|                 | Interferon-α  | Retinoids                | 9cRA    | tRA   | 13cRA  |
| HTB-82 (rhabdomyosarcoma) | No            | Yes                      | Yes     | No    | No     |
| HTB-91 (fibrosarcoma)       | Yes           | Yes                      | No      | No    | No     |
| HTB-92 (liposarcoma)        | Yes           | Yes                      | No      | No    | No     |
| HTB-93 (synovial sarcoma)    | No            | Yes                      | Yes     | Yes   | No     |
| HTB-94 (chondrosarcoma)     | Yes           | Yes                      | No      | No    | No     |

*As assessed by p53 sequence analysis.
atmosphere without CO₂). Medium containing retinoids and/or IFN-α-2b was replaced every 24 h. (³H)Thymidine (Amersham International Life Science, plc, Buckinghamshire, UK), at a concentration of 0.5 µCi per well, was included for the final 16 h. The incorporation of (³H)Thymidine into DNA was measured by a Direct Beta Counter-Matrix 96 (Packard, Groningen, The Netherlands) after the cells were harvested with the Harvester Micromate 196 (Packard, Groningen, The Netherlands) onto Glass Fiber Filters (Packard, Groningen, the Netherlands). Experiments were always done in triplicate. Data are presented as a percentage of proliferation of untreated cells for each respective time point.

DNA fragmentation assay analysed by flow cytometry

The 3´OH termini in DNA breaks were measured by attaching fluorescent tagged deoxyuridine triphosphate nucleotides FITC-dUTP, in a reaction catalysed by terminal deoxynucleotidyl transferase (TdT) using the Apo-Direct™ Kit (Phoenix Flow Systems, San Diego, CA, USA) purchased from Pharmingen (San Diego, CA, USA). The amount of incorporated fluorescein was detected by flow cytometry.

HTB-82, HTB-91, HTB-92, HTB-93 and HTB-94 soft tissue sarcoma cell lines (1·10⁶ in T-25 flasks) were incubated with 9cRA, 13cRA or tRA (final concentrations: 10⁻⁷ M, 10⁻⁶ M, 10⁻⁵ M) for 48, 72, 96 and 120 h respectively. In addition, the same experiments were performed with IFN-α-2b (final concentration: 1000 U ml⁻¹) – 9cRA, 13cRA or tRA (final concentrations: 10⁻⁷ M, 10⁻⁶ M, 10⁻⁵ M).

Untreated and treated cells were harvested, washed twice in phosphate-buffered saline (PBS), fixed in 1% (w/v) paraformaldehyde in PBS (pH 7.2), for 15 min on ice. After sedimentation and two more washing steps, cells were resuspended in ice-cold 70% (v/v) ethanol and stored at −20°C until further use (maximum: 3 weeks). According to the manufacturer’s instructions, cells were washed twice in wash buffer, resuspended in 50 µl staining solution (10 µl reaction buffer, 0.75 µl TdT, 8 µl FITC-dUTP and 32 µl distilled water) and incubated for 1 h at 37°C. Afterwards, 1 ml rinsing buffer was added. Cells were centrifuged (1000 g) and rinsed again. Subsequently, cells were resuspended in 1 ml propidium iodine (PI)/RNAase solution and incubated in the dark for 30 min at room temperature. Subsequently, cell samples were analysed by flow cytometry on a FACScan (Becton Dickinson, CA, USA).

Morphological evaluation of apoptosis

Cell morphology was performed by staining with May–Gruenwald–Giemsa. For this reason, cells were cultured in chamber slides (‘Lab-Tek Chamber Slide w/cover Glass Slide 2 Well’) (Nalge Nunc International, Naperville, USA). Treated and untreated cells were stained with May–Gruenwald (Merck, Darmstadt, Germany) for 5 min. Subsequently, cells were washed with aqua dest (Leopold Pharma, Graz, Austria) for 5 min and stained with Giemsa (Merck, Darmstadt, Germany) (diluted 1:10) for 20 min. After a final wash with aqua dest, samples were viewed under light microscopy. Cells were considered to be apoptotic according to the criteria introduced by Kerr et al (1972) which included: compaction of the nuclear chromatin, fragmentation of nuclei, condensation of the cytoplasm and separation of the cell into apoptotic bodies.

p53 immunohistochemistry

Immunohistochemical staining for the p53 protein was carried out using a mouse monoclonal IgG2a antibody (Clone: DO-1; Immunotech, Marseille, France; diluted 1:20) directed against wild-type and mutant p53 protein. For p53 staining, cytocentrifugates were fixed with Merckofix fixation spray (Merck, Darmstadt, Germany). After blocking with horse serum, samples were incubated with the primary antibody for 1 h. Further immunohistochemical staining was performed according to the ABC-method, using products from Vector Laboratories (Burlingame, CA, USA). Briefly, after incubation with the primary antibody and incubation with a biotinylated antiamouse antibody, incubation with the ABC complex for 45 min followed. The reaction product was developed with diaminobenzidine tetrahydrochloride. Finally, slides were counterstained with Gill’s haematoxylin. All steps of incubation were performed at room temperature.

p53 sequence analysis

Total genomic tumour DNA was extracted from 2 × 10⁷ cells using

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**Figure 1** Immunohistochemistry of soft tissue sarcoma cell lines for p53 protein. No overexpression of p53 was found in cell line HTB-91 (fibrosarcoma: A); whereas cell line HTB-92 (liposarcoma) showed intense nuclear staining (B).
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Figure 2  Proliferation of soft tissue sarcoma cell lines during incubation with varying concentrations of the following substances (³H-thymidine incorporation assay): lane 1: interferon-α; lane 2: 9-cis-retinoic acid; lane 3: all-trans-retinoic acid; lane 4: 13-cis-retinoic acid. Data are presented as relative counts per minute of untreated cells (100%) for each respective time point.
standard phenol–chloroform extraction methods. Exons 2–11 of the p53 gene were amplified separately using oligonucleotide primers placed in the adjacent intron regions as described previously (Lehman et al., 1991). Polymerase chain reaction (PCR) products were controlled for purity, quantity and quality by subjecting 5 μl of PCR products to pre-cast 6% acrylamide/bis-acrylamide gels (Novex, San Diego, CA, USA) using the pBR322 DNA-MspI digest as reference standard (Clontech Labs Inc., Palo Alto, CA, USA). To remove residual single-stranded primers, 5 μl of PCR products were enzymatically treated with combination of exonuclease I and shrimp alkaline phosphatase (United States Biochemical, Cleveland, OH, USA). Pretreated PCR products were then directly sequenced using the Cycle Sequencing Kit (Roche Molecular Systems, Inc. Branchburg, NJ, USA), utilizing Ampli Taq® DNA Polymerase and ε35S labelled dATP (DuPont NEN, Brussels, Belgium). For each reaction (for the four nucleotides), 0.5 μl of the thermostable DNA polymerase provided in the kit were added at last to the reaction mix containing 2 μl reaction buffer, 2–4 μl of enzymatically pretreated PCR product, 0.5 pmol unique primer (separate reaction for each exon) and water to adjust total volume to 20 μl. After running, the 8% acrylamide/bis-acrylamide gels are dried for 90 min at 70°C and directly subjected to autoradiography against the Bio Max-MR film (Kodak, New Haven, CT, USA). Mutations found were confirmed by at least one complete reanalysis.

Detection of bcl-2 by immunofluorescence

Cells (1 × 10⁶ in T-25 flasks) were incubated with 9cRA, 13cRA or tRA (final concentrations: 10⁻⁷ M, 10⁻⁶ M, 10⁻⁵ M) for 48, 72, 96 and 120 h respectively. In addition, the same experiments were performed with IFN-α-2b (final concentration: 1000 U ml⁻¹) ± 9cRA, 13cRA or tRA (final concentrations: 10⁻⁵ M, 10⁻⁶ M, 10⁻⁷ M). After harvesting and three washes with PBS cells were fixed and permeabilized with the fix and perm cell permeabilization kit (An der Grub, Bio Research Gmbh, Kaumberg, Austria). According to the manufacturer’s instructions, 10⁶ cells per sample were resuspended in 100 μl of the fix and perm cell permeabilization kit (An der Grub, Bio Research Gmbh, Kaumberg, Austria). After fixing and washing the cells, they were treated with anti-human bcl-2 antibody (clone 124; DAKO, Glostrup, Denmark), washed with PBS and incubated with FITC-conjugated monoclonal mouse IgG anti-human bcl-2 antibody (clone 124; DAKO, Glostrup, Denmark), vortexed at low speed for 2 s and incubated for 15 min at room temperature. After one more washing step, cells were analysed by flow cytometry on a FACSScan (Becton Dickinson, CA, USA).

Northern blotting for RAR mRNA expression

STS cells were plated in T-75 tissue culture flask (Costar) and were grown to sub-confluence (see above). The total RNA was extracted by a one-step acid guanidinium isothiocyanate–phenol procedure using RNAzol (Biotecx, Houston, TX, USA) and 20 mg of the RNA were loaded into each lane and were electrophoresed for 4 h at 50 V in a 1% formaldehyde-containing agarose gel. The separated RNA was then diffusion-blotted overnight onto nylon membranes (Schleicher & Schuell, Dassel, Germany) and immobilized by UV-cross-linking. RAR- and GAPDH-specific cDNA probes were labelled with ³²P-dCTP (New England Nuclear, Vienna, Austria) using the random oligonucleotide primer extension kit (Stratagene, La Jolla, CA, USA). The membranes were hybridized overnight at 65°C, washed twice with 2 × saline-sodium citrate (SSC) and 0.1% sodium dodecyl sulphate (SDS) at 37°C and twice with 0.1 × SSC and 0.1% SDS at 65°C and were exposed at −80°C to Kodak X-OMAT AR film (Eastman Kodak, Rochester, NY, USA) between two intensifying screens. The blots were then stripped and reprobed (overnight, 65°C) with the GAPDH cDNA as an internal control to monitor the quantity and integrity of the loaded RNA samples.

RESULTS

p53 genotype and protein expression

Native cell lines HTB-82, HTB-91, HTB-92, HTB-93 and HTB-94 were screened for their p53 genotype as well as their p53 protein expression. p53 gene mutations are characterized in Table 1: HTB-91, HTB-92 and HTB-94 cell lines had mutations in the p53 gene, whereas wild-type p53 was found in HTB-82 and HTB-93 cell lines. Immunohistochemistry for p53 revealed intense nuclear staining in HTB-92 (Figure 1B) and HTB-94 cells, whereas HTB-82, HTB-91 (Figure 1A) and HTB-93 cells showed no nuclear staining. The p53 mutation of HTB-91 cells was probably associated with loss of protein staining.

Proliferation inhibition

To test the influence of retinoids and IFN-α on proliferation, STS cell lines were incubated with IFN-α and/or retinoids for 48, 72, 96 and 120 h respectively. Figure 2 shows that IFN-α dose- and time-dependently inhibited the proliferation of all five STS cell lines at concentrations ranging from 25 U ml⁻¹ to 1000 U ml⁻¹ and incubation periods from 48 to 120 h. It has to be noted, however, that the inhibitory potential of IFN-α exerted upon various tested cell lines varied with ranges between 49 and 77% (Figure 2).

Similarly, 9cRA, 13cRA or tRA showed a dose- (10⁻⁷ M, 10⁻⁶ M and 10⁻⁵ M) and time-dependent proliferation inhibitory effect on cell lines HTB-82 and HTB-93 (Figure 2). In contrast, HTB-91, HTB-92 and HTB-94 could not be inhibited in their proliferation by either 9cRA or 13cRA or tRA (Figure 2). Coincubation of HTB-82, HTB-91, HTB-92, HTB-93 and HTB-94 STS cells with retinoids and IFN-α did not further increase the antiproliferative effect of IFN-α (data not shown). In control experiments, DMSO only had no inhibitory influence upon either of the five cell lines after any duration of incubation.

Induction of apoptosis: DNA fragmentation, analysis of cell cycle position and DNA content

Induction of apoptosis by retinoids or IFN-α in STS cell lines was investigated by DNA strand-break analysis. In order to analyse cell cycle position of STS cells, global DNA content was measured with PI counterstaining. HTB-82, HTB-91, HTB-92, HTB-93 or HTB-94 STS cells were incubated with 9cRA, 13cRA or tRA (final concentrations: 10⁻⁷ M, 10⁻⁶ M and 10⁻⁵ M) and/or IFN-α (1000 U ml⁻¹) for 48, 72, 96 or 120 h respectively. DNA histograms obtained from STS cell lines and the respective percentages of apoptotic and non-apoptotic cells after 96 h incubation with 10⁻⁵ M of the appropriate retinoid are shown in Figure 3. In summary, apoptosis could be induced by retinoids 9cRA or tRA in cell lines HTB-82 and HTB-93 dose-dependently with an optimal duration of incubation with 10⁻⁵ M of the appropriate retinoid for 96 h (Figure 3), whereas no apoptosis was found in cell lines HTB-91, HTB-92 and HTB-94 during an
incubation period of up to 120 h. In addition, treatment of HTB-82 and HTB-93 cells with 9cRA or tRA was associated with some G1–S arrest. Finally, neither 13cRA nor IFN-α induced apoptosis in either cell line. In detail, the results of induction of apoptosis in correlation with cell cycle positions were as follows.

**HTB-93 (synovial sarcoma)**

After 96 h of incubation with 9cRA (10⁻⁵ M), the percentage of apoptotic cells was 41% (G1: 35%; G2 and S: 6%) versus 59% non-apoptotic cells (G1: 43%; G2 and S: 16%). At lower doses of 9cRA (10⁻⁶ M and 10⁻⁷ M) we observed 29 and 20% apoptotic cells respectively. After incubation with tRA (10⁻⁵ M) 50% of the cells (G1: 45%; G2 and S: 5%) underwent apoptosis versus 50% non-apoptotic cells (G1: 38%; G2 and S: 12%). At a concentration of 10⁻⁶ M of tRA, 36% of cells showed apoptosis and 25% at 10⁻⁷ M. No apoptosis was induced by either retinoid by incubation for 48 h.

**HTB-82 (rhabdomyosarcoma)**

Only tRA was capable of inducing apoptosis. After 96 h of incubation 89% (G1: 67%; G2 and S: 22%) of the cells were apoptotic.

*Figure 3* FACS histograms of specifically for apoptosis stained soft tissue sarcoma cell lines after incubation without (control) or with 10⁻⁵ M 9-cis-retinoic acid (9cRA) or all-trans-retinoic acid (tRA) for 96 h. FACS histograms show on the x-axis the log fluorescence intensity of non-apoptotic and apoptotic cells within the tested cell population. The y-axis represents the relative number of cells.
Eleven per cent of the cells did not undergo apoptosis (G1: 9%; G2 and S: 2%). At lower concentrations of tRA (10^{-6} M and 10^{-7} M) we observed 50 and 41% apoptotic cells respectively.

HTB-91 (fibrosarcoma), HTB-92 (liposarcoma) and HTB-94 (chondrosarcoma)

No apoptosis was measured after treatment with 9cRA, 13cRA, tRA and/or IFN-\(\alpha\). Neither 13cRA nor IFN-\(\alpha\) in any concentration (10^{-7}–10^{-5} M; 25–1000 U ml^{-1}) and after any length of incubation (48–120 h) were capable of inducing apoptosis in either cell line. Moreover, the addition of IFN-\(\alpha\) to retinoids which were able to induce apoptosis in either cell line did not increase their apoptotic potency (data not shown). Finally, incubation of all five cell lines with DMSO only did not induce apoptosis after any time interval (data not shown).

Morphologic evaluation of apoptosis

Morphology of cells shown to undergo apoptosis by flow cytometry exhibited typical apoptotic features such as nuclear-chromatin condensation, cytoplasmic condensation around the nucleus, cell shrinkage and apoptotic ‘bodies’ (Figure 4).

bcl-2 protein expression

The effects of 9cRA, 13cRA and tRA on bcl-2 protein expression were also studied. All five native cell lines (HTB-82, HTB-91, HTB-92, HTB-93 and HTB-94) weakly expressed bcl-2 protein (10–20%). Treatment of these cell lines with 9cRA, 13cRA and tRA (final concentrations: 10^{-7} M, 10^{-6} M, 10^{-5} M) for 48, 72, 96 and 120 h respectively, did not modify the expression of bcl-2 protein.

Expression of RAR isoforms

None of the five STS cell lines expressed RAR-\(\alpha\) and RAR-\(\beta\) mRNA at levels detectable by Northern blot analysis. However, RAR-\(\gamma\)-specific transcripts with molecular sizes of approximately 3.3 kb were found in variable amounts in all cell lines (Figure 5). Thus, the differential response of STS cell lines to retinoids was not associated with a lack of RARs in general, or with a defect in the expression of the universal retinoid receptor RAR-\(\gamma\) in particular, or with a specific pattern of RAR isoform expression.

Correlation of p53 status with proliferation inhibition and apoptosis

Normal p53 was found in cell lines in which retinoids 9cRA or tRA were capable of inducing apoptosis and in which retinoids 9cRA, 13cRA and tRA led to proliferation inhibition. In contrast, mutant p53 was present in cell lines in which retinoids were not able to induce either apoptosis or proliferation inhibition (Table 2). However, proliferation inhibition induced by IFN-\(\alpha\) occurred in all cell lines irrespective of their p53 status (Table 2). In control experiments, p53 genotype as well as p53 protein expression did not change after incubation of STS cell lines HTB-82,-91, -92, -93 and -94 with 9cRA, 13cRA or tRA (final concentrations: 10^{-7} M, 10^{-6} M and 10^{-5} M) and/or IFN-\(\alpha\) (final concentrations: 25, 50, 100 and 1000 U ml^{-1}) for 48, 72, 96 or 120 h (data not shown).

DISCUSSION

In response to DNA damage, wild-type p53 has been shown to induce either cell cycle arrest in the G1 phase (Kastan et al, 1991; Di Leonardo et al, 1994), allowing for DNA repair, or apoptosis (Yonish-Rouach et al, 1991). In addition, wild-type p53 is also responsible for the regulation of the cell cycle (Harvey et al, 1993) during which p53-mediated arrests have been reported to occur not only in the G1, but also in the G2 phases (Ryan et al, 1993; Agarwal et al, 1995; Stewart et al, 1995). The final decision whether induction of p53 triggers G1 arrest or apoptosis...
(Slichenmyer et al, 1993) is thought to be dependent on the presence of a growth factor which provides a survival signal (Cannan et al, 1995). However, additional p53-independent apoptotic pathways exist in normal as well as malignant cells which differ in their sensitivity towards apoptosis-inducing agents (Delia et al, 1993; Bracey et al, 1995; Shao et al, 1995; Thompson, 1995).

Retinoids exert a broad spectrum of biologic effects, including inhibition of cell growth and induction of apoptosis in a variety of cancer cell lines (Lotan, 1995). These biological effects are thought to result from modulation of gene expression mediated by inhibition of cell growth and induction of apoptosis in a variety of cancer cell lines (Lotan, 1995; Bracey et al, 1995; Shao et al, 1995; Thompson, 1995).

However, additional p53-independent apoptotic pathways exist in normal as well as malignant cells which differ in their sensitivity towards apoptosis-inducing agents (Delia et al, 1993; Bracey et al, 1995; Shao et al, 1995; Thompson, 1995).

In the present investigation, we have shown that the mechanisms of STS lines varying in p53 mutations can act independently from the used agents. Thus, IFN-α was found to induce inhibition of proliferation independently from the cells’ p53 status, whereas the inhibition of proliferation achieved by retinoids was associated with intact p53 and was not found in cells with p53 mutations. Considering its important regulatory potential upon blocking apoptosis (Hockenbery et al, 1990), no influence of bcl-2 within the context of the reported findings was detectable in the present model. Nevertheless, the degree of inhibition of proliferation varied between cell lines depending upon the chosen retinoid, its concentration and time. In order to illustrate the actual inhibition of cellular proliferation rather than the induction of apoptosis, a thymidine incorporation assay was used which proved DNA synthesis occurred thus providing a direct parameter for cellular proliferation and its actual inhibition. These findings suggest that p53 status and its influence upon proliferation inhibition in STS cell lines acted independently from and changed with the used preparation. It is conceivable, therefore, that proliferation inhibition induced by IFN-α used a different pathway independent from p53 (Dmitrovsky, 1997; Giandomenico et al, 1997; Oridate et al, 1997) the functionality of which is important, however, for the activity exerted by retinoids in the same context.

In contrast to its antiproliferative effect, IFN-α may not be the ability to induce apoptosis in the tested STS cell lines. In contrast, tRA was capable of inducing apoptosis in cell lines HTB-82 and HTB-93, and 9cRA in cell line HTB-93, whereas 13cRA did not induce apoptosis, at all. However, no induction of apoptosis by otherwise effective retinoids 9cRA or tRA occurred in cell lines HTB-91, HTB-92 and HTB-94 in which mutations of p53 were found. Data obtained from Northern blot analysis demonstrated that resistance to induction of apoptosis induced by retinoids in the three STS cell lines harbouring mutations of the p53 gene did not result from a defect in the expression of RARs indicating that p53 did not regulate RAR-α, -β or -γ expression in these instances. As RAR-γ constitutes a universal receptor for the used retinoids (Repa et al, 1997; Fisher et al, 1998), a defect in the presence of a relevant receptor can be excluded as an explanation for our findings. Thus, no simple correlation of retinoid sensitivity with the expression pattern of RARs could be established for STS cell lines which is in accordance with previous reports (Van der Leede et al, 1993; Engers et al, 1996). Although p53-mutated HTB-92 and HTB-94 cell lines showed intensive nuclear staining by immunohistochemistry, the p53 mutated HTB-91 cell line showed no nuclear staining. The p53 mutation in HTB-91 is probably related to a loss of protein expression in these instances. As a consequence, the p53 mutated HTB-91 cell line showed no nuclear staining (Lotan, 1995). It is, therefore, interesting to note that the inhibition of proliferation by retinoids was closely linked to their ability to induce apoptosis, suggesting the importance of p53 in both contexts. It is noteworthy, however, that 13cRA – while being able to inhibit cellular proliferation – did not induce apoptosis, thus further suggesting a possible divergence of mechanisms regulating the two phenomena. This assumption is supported by recent reports that have demonstrated that 13cRA did not bind to RARs on its own (Wu et al, 1994; Berggren et al, 1995). In summary, an overview of our data suggests that only mutations of p53 correlated with an abrogation of proliferation inhibition and of induction of apoptosis by retinoids 9cRA and tRA.

Regarding the possible clinical impact of our data, we report an antiproliferative influence of IFN-α and certain retinoids as well as the induction of apoptosis by the latter group of drugs, which could constitute the basis for clinical studies in patients with STS, especially when cytotoxic chemotherapy has failed. In similarity to this assumption, recent reports have already reported about the successful inclusion of retinoids in the treatment of certain malignancies (Tallman et al, 1997). Concerning STS, however, our results have shown that the histological subtype and p53 status might predict the efficacy of this therapy. The latter aspect could also be valid for the use of cytotoxic therapy which often has failed to produce convincing results in STS in the adjuvant or palliative setting.

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