Lack of Bcl10 mutations in testicular germ cell tumours and derived cell lines

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Summary Aberrations within Bcl10, a gene involved in execution of apoptosis, has most recently been found in a variety of cancers, including cell lines of testicular germ cell tumours of adolescents and adults (TGCTs). To study this in more detail, we screened exons 2 and 3 of this gene for mutations in a larger series of cell lines as well as primary TGCTs by single-strand conformation polymorphism and endonuclease restriction analysis. Because no aberrations were detected, we conclude that inactivation of Bcl10 by mutation is at least far less important in the development of TGCTs than proposed.

Keywords: testicular cancer; Bcl10; mutation-analysis

One possible event in the development of cancer is a disturbed induction of apoptosis, i.e. programmed cell death. This will lead to an enlarged population of target cells for the subsequent events, finally leading to the fully malignant phenotype. The classical example of this mechanism, eventually resulting in follicular B-cell non-Hodgkin’s lymphomas, is the outcome of a translocation of chromosome 18 band q21 and chromosome 14 band q32. This translocation places Bcl2 under control of the IgH promoter, leading to overexpression, and subsequent inhibition of apoptosis (Tsujimoto et al, 1984). More recently, an alternative and intriguing mechanism has been reported in B-cell lymphomas of mucosa-associated lymphoid tissue (MALT lymphomas). Bcl10, a gene mapped to chromosome 1 band P22, encodes a protein with caspase activity, which is involved in the execution of the apoptotic machinery. Translocation of this gene to chromosome 14 band q32, as has been found in MALT lymphomas, results in a disrupted caspase activity, thereby preventing apoptosis (Willis et al, 1999). Interestingly, aberrations within this gene, like point-mutations and small deletions, were also found in a variety of cancers, including colon cancers, mesotheliomas and testicular germ cell tumours of adolescents and adults (TGCTs). In total, 87 cell lines were screened for aberrations, 81 on genomic DNA level, and six using cDNA clones. The high incidence of aberrations in Bcl10 found, for example, in three out of three TGCT-derived cell lines, suggests that inactivation of this gene is a general and crucial mechanism in the pathogenesis of different types of malignancies.

TGCTs are the most common malignancy in young Caucasian males aged between 15 and 45 years (Swerdlow, 1998), and the incidence is still increasing. It has generally been accepted that these cancers originate from an early germ cell, of which the malignant counterpart is referred to as carcinoma in situ (Skakkebæk, 1972; Jørgensen et al, 1997). Histologically and clinically, TGCTs are divided into non-seminomatous-TGCTs (NS-TGCTs) and seminomas (Ulbright, 1993). While seminomas mimic carcinoma in situ, the NS-TGCTs reflect embryonal development to a certain extent. They can be composed of embryonal carcinoma, being the undifferentiated cells, or teratoma, mimicking somatic differentiation. In addition, the extra-embryonal elements yolk sac tumour and choriocarcinoma can be found. Total DNA content analysis showed that all TGCTs, including carcinoma in situ, are aneuploid (Oosterhuis et al, 1989; De Graaff et al, 1992). Karyotyping, allelic imbalances analysis and, more recently, comparative genomic hybridization revealed a relatively consistent pattern of gains and losses (Mathew et al, 1994; Van Echten-Arends et al, 1995; Sandberg et al, 1996; Looijenga and Oosterhuis, 1999, and references cited therein). For example, loss of (part of) the short arm of chromosome 1 has been found both in seminomas and NS-TGCTs, suggesting the presence of a tumour suppressor gene.

To investigate the suggested role of Bcl10 in the development of TGCTs more extensively, we studied a series of derived and related cell lines, as well as primary tumours for genomic aberrations within exon 2 and 3 of this gene. Therefore we applied the technique of single-strand conformation polymorphism (SSCP) analysis, as well as a more straightforward analysis for a number of reported aberrations by restriction endonuclease digestion analysis.

MATERIALS AND METHODS

The NS-TGCT-derived cell lines Tera-1, Tera-2, GCT44, 2102 EP, NTera-2, GCT27 and NCCIT, were included. In addition, the cell lines Haz and Bos, derived from two extragonadal germ cell tumours, were studied (Sinke et al, 1994; Van Echten-Arends et al, 1995), as well as the cell line S2, of which the origin is still a matter of debate (von Keitz et al, 1994; Andrews et al, 1996). The mesothelioma-derived cell line M25 was also investigated (Versnel et al, 1989). The cell lines were cultured as described.
before (Andrews et al, 1996). Four primary testicular seminomas and four NS-TGCTs, with the histology of pure embryonal carcinoma, pure teratoma, pure yolk sac tumour and mixed containing embryonal carcinoma, teratoma and yolk sac tumour, were investigated. In addition, one gonadoblastoma (reviewed in Hussong et al, 1997), one Leydig cell tumour and one spermatocytic seminoma (reviewed in Rosenberg et al, 1998) were studied. The tumours were snap-frozen upon orchidectomy. Peripheral blood DNA of three unrelated individuals were included in every experiment as control, and all experiments were done at least twice. DNA was isolated from the in vitro cultured cell lines, peripheral blood and tumours using standard procedures as described before (Sambrook et al, 1989).

Exons 2 and 3 were screened for the presence of aberrations on genomic DNA level by SSCP and restriction endonuclease digestion analysis on polymerase chain reaction (PCR)-amplified fragments. Positions within the genomic sequence of the primers used for amplification are schematically represented in Figure 1 and are identical to those described (Willis et al, 1999). Amplification was done in a total volume of 25 μl of buffer (10 mM Tris–HCl pH 8.8, 50 mM potassium chloride, 0.1% Triton X-100, 2 mM magnesium chloride; 5 pmol of each primer; 0.2 mM dNTP and 1 U Taq (Promega, Madison, WI, USA)), containing 25 ng high molecular weight DNA. The PCR programme consisted of an initial denaturation of 4 min at 94°C, 35 cycles of 1 min at 94°C denaturation, 1 min at 60°C annealing and 1 min at 72°C elongation, respectively, followed by 6 min at 72°C.

SSCP was carried out at 4°C on a 6% non-denaturing polyacrylamide gel using radioactive labelled PCR fragments. Briefly, 1 μl of α-32P-dATP (ICN, Zoetermeer, The Netherlands) was added to the PCR-mix and, after amplification, 25 μl sequencing loading buffer was added. After denaturation, 4 μl was loaded on the gel and run for 5 h at 60 Watt, whereafter the gel was dried. BioMax film (Kodak, Rochester, NY, USA) was exposed overnight at –80°C. To exclude misinterpretation of the SSCP patterns due to the presence of double-stranded DNA fragments, non-denatured PCR fragments were included.

To confirm the absence of specific aberrations, endonuclease restriction digestion analysis was done on amplification products of (part of) exon 2 by the primer sets as given in Figure 1 (either exon 2 or exon 2.1, which is the same primer set as used for SSCP analysis; see above). Therefore 5 μl PCR product was digested with 1 unit of StuI or TaqI according to the manufacturer’s instructions (Pharmacia Biotech, Uppsala, Sweden) in a total volume of 10 μl. After digestion, the samples were run on a 2% agarose gel stained with ethidium bromide and analysed using the IS-500 digital imaging system and AlphaImager 2000 (Alpha Innotech, San Leandro, CA, USA). Aberrations within the recognition sites of these specific restriction endonucleases will result in the presence of undigested fragments.

Direct sequencing of amplified fragments of exon 2 and 2.1 using end-labelled primers 2.1r and 2.2r (Figure 1) was performed using the cycle-sequencing kit (Perkin-Elmer Cetus, Norwalk, CT, USA).

**RESULTS**

To screen DNA isolated from NS-TGCT-derived and related cell lines, as well as primary testicular tumours, including both seminomas and NS-TGCTs, for aberrations within exon 2 and 3 of the Bcl10 gene, initially SSCP was performed. For both exons, two
sets of primers were used, resulting in two overlapping fragments, designated 2.1, 2.2, 3.1 and 3.2 respectively (see Figure 1). The expected sizes are indicated in the Figure. In contrast to the reported length of 200 bp (Willis et al, 1999), the 2.1 fragment is found to be 236 bp, both in controls, cell lines and tumours. Direct sequencing of the amplification products demonstrated that a 54 bp (and not 18 bp) intronic sequence separates the 2.1 forward primer and exon 2 (see Figure 2 for the additional sequence). All cell lines and tumours showed a similar SSCP pattern, identical to the pattern detected in the control samples (not shown). The presence of double-stranded fragments, possibly leading to misinterpretation, was excluded within the protocol. Within the series of NS-TGCT-derived cell lines screened in this study, Tera-1 and GCT44 were also investigated (see Materials and Methods), and were not found to contain aberrations on the genomic DNA level, although these cell lines were shown to contain mutations in the original study by Willis et al (1999; see Fig. 1).

To exclude that the presence of specific aberrations within exon 2 remained undetected by SSCP, as can be the case (Landegren et al, 1998), we additionally performed a more direct analysis. Therefore restriction endonuclease digestions were performed on the amplification products of exon 2 (both 2 and 2.1) using Stul and TaqI (see Figure 1). These restriction endonucleases digest the wild-type sequence, resulting in fragments of respectively 79 + 157 and 79 + 338 bp for Stul, depending on the primer sets applied, and 192 + 44 and 192 + 225 bp for TaqI respectively. In case of a mutation within either the Stul or TaqI recognition site, an undigested fragment of 236 or 417 bp will be found. A disrupted Stul recognition site can be expected for Tera-2, although this mutation was only reported in a cDNA clone. None of the cell lines and tumours tested in this study showed the presence of undigested fragments after Stul and TaqI digestion, indicating the absence of mutations within these sequences, of which representative examples are illustrated in Figure 3. These results were confirmed by direct sequencing of the amplification products (data not shown). Moreover, we also studied the mesothelioma cell line M25, found to contain a deletion of 16 bp within exon 3. In our hands, PCR showed the wild-type length of this fragment in this particular cell line (not shown).

**DISCUSSION**

Apoptosis is found to be crucial for normal development and maintenance of an organism. In this process, caspase (‘cysteinyl aspartate-specific proteinases’) activity is important for proper execution. Several caspases are identified so far (Salvesen and Dixit, 1997). In addition to its importance during normal development and maintenance, a disturbed apoptosis has also been identified as a possible event in the development of cancer. This can be the result of activation of apoptosis-inhibiting genes, like Bcl2, or, more recently, also due to inactivation of apoptosis-inducing genes. Disruption of the caspase activity of Bcl10 has most
recently been found in MALT lymphomas (Willis et al. 1999). In addition, this study also identified mutations within exons 2 and 3 of this gene in a variety of cancers of different origin, including three out of three NS-TGCT-derived cell lines. It is of note, that we suggested inhibition of apoptosis being one of the mechanisms involved in the pathogenesis of TGCTs, i.e. lack of induction of apoptosis of the precursor cell of this cancer (Looijenga and Oosterhuis, 1999). Moreover, we showed that RAS mutation or amplification leads to an enhanced in vitro survival of seminoma cells, related to a reduced sensitivity for apoptosis upon disruption of the micro-environment (Oosterhuis et al., 1998; Looijenga and Oosterhuis, 1999).

To study the role of Bcl10 in the pathogenesis of TGCTs in more detail, we performed in principle a similar approach as Willis et al. (1999). DNA of a series of cell lines were investigated by SSCP using the same primer sets as reported before. However, no indications for the presence of aberrations within exons 2 and 3 were obtained in our hands. The presence of independent and multiple controls without denaturation before SSCP prevented misinterpretation of the data due to double-stranded fragments in the samples under investigation. Besides the cell lines, a series of primary TGCTs also showed no aberrant bands by SSCP. In fact, even no aberrant migration patterns of the fragments derived from the cell lines Tera-1 and GCT 44 were observed, in spite of the results reported before (Willis et al., 1999). In addition, the restriction endonuclease digestion analysis for specific sequence abnormalities, also showed no undigested fragment, i.e. only wild-type alleles were amplified, which was confirmed by direct sequencing. There is no reason to assume the inability of amplification of mutant alleles, because of the use of flanking sets of primers (see Figure 1). Because in the initial study the majority of aberrations were detected on the genomic DNA level, we limited our survey to genomic DNA analysis. The formerly reported mutations in the cDNA clones of the Tera-2 cell line could not be confirmed by us on genomic DNA.

Theoretically, the discrepancies between the study of Willis et al. (1999) and our results (this study) could be due to the extended in vitro culturing of the different cell lines, both from NS-TGCT- and Tera-2 cell line could not be confirmed by us on genomic DNA. Therefore, we limited our survey to genomic DNA analysis. The formerly reported mutations in the cDNA clones of the Tera-2 cell line could not be confirmed by us on genomic DNA.

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