High expression of Mcl-1 in ALK positive and negative anaplastic large cell lymphoma

R Rust, G Harms, T Blokzijl, M Boot, A Diepstra, J Kluiver, L Visser, S-C Peh, M Lim, W A Kamps, S Poppema, A van den Berg

Aim: To gain more insight into the genes involved in the aetiology and pathogenesis of anaplastic large cell lymphoma (ALCL).

Methods: Serial analysis of gene expression (SAGE) was undertaken on the CD4+ALK+ (anaplastic lymphoma kinase positive) ALCL derived cell line Karpas299 and as comparison on CD4+ T cells. Quantitative reverse transcription polymerase chain reaction (RT-PCR) and immunohistochemistry were performed on five ALCL derived cell lines and 32 tissue samples to confirm the SAGE data.

Results: High expression of Mcl-1 was seen in the Karpas299 cell line, whereas the two other antiapoptotic Bcl-2 family members, Bcl-2 and Bcl-X\textsubscript{L}, were not detected in the SAGE library. Quantitative RT-PCR confirmed the high expression of Mcl-1 mRNA and low expression of Bcl-2 and Bcl-X\textsubscript{L} in Karpas299 and in four other ALCL cell lines. To expand on these initial observations, primary tissue samples were analysed for Mcl-1, Bcl-X\textsubscript{L}, and Bcl-2 by immunohistochemistry. All 23 ALK+ and nine ALK- ALCL cases were positive for Mcl-1. Bcl-2 and Bcl-X\textsubscript{L} were expressed infrequently in ALK+ ALCL cases, but were present in a higher proportion of ALK- ALCL cases.

Conclusion: The consistent high expression of Mcl-1 in ALK+ and ALK- ALCL suggests that Mcl-1 is the main antiapoptotic protein in this disease. The high frequency of Mcl-1, Bcl-2, and Bcl-X\textsubscript{L} positive ALCL cases in the ALK- group compared with the ALK+ group indicates that ALK induced STAT3 activation is not the main regulatory pathway in ALCL.

Abbreviations: ALCL, anaplastic large cell lymphoma; ALK, anaplastic lymphoma kinase; $\beta_2$m, \beta_2 microglobulin; Ct, threshold cycle; RT-PCR, reverse transcription polymerase chain reaction; SAGE, serial analysis of gene expression

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MATERIALS AND METHODS

Cell lines and tissues

CD4+ T cells were isolated from the buffy coats of healthy donors using a fluorescence activated cell sorter (MoFlo Cytomation, Fort Collins, Colorado, USA) and were used directly for RNA isolation and SAGE analysis. The CD4+ALK+ ALCL derived cell line Karpas299 was obtained from ATCC (Rockville, Maryland, USA) and the NPM/ALK positive ALCL derived cell lines SU-DHL-1, SR786, SUP-M2, and DEL were obtained from DSMZ (Braunschweig, Germany). Based on the presence of the (2;5) translocation, expression of CD4 and ALK, and lack of expression of B cell markers, we selected the Karpas299 cell line for the SAGE analysis. Frozen and paraffin wax embedded ALCL tissue specimens from 23 ALK+ and nine ALK- ALCL cases were obtained from the departments of pathology of the Groningen University Medical Centre, the Netherlands and the University of Malaysia. All protocols for obtaining and studying human...
tissues and cells were approved by the institution’s review board for human subject research.

**SAGE**

A detailed protocol for the SAGE procedure and a computer program (SAGE2000 version 4.12) for the analysis of gene specific tags were kindly provided by Dr KW Kinzler (John Hopkins Oncology Center, Baltimore, Maryland, USA). The SAGE procedure was described previously.14 The SAGE libraries were compared and linked to the Unigene library to identify the corresponding genes.

**Gene specific real time RT-PCR**

Total RNA was isolated with Trizol (Life Technologies Inc, Gaithersburg, Maryland, USA) and first strand cDNA synthesis was performed using the protocol provided by the manufacturer (Life Technologies Inc). Assays-on-Demand™ gene expression products (Applied Biosystems, Foster City, California, USA) were used for Bcl-2 (Hs00236808_s1), Bcl-XL (Hs00236329_m1), and Mcl-1 (Hs00172031_m1). Real time PCR was performed in 1 x Taqman® Universal PCR master mix (Applied Biosystems). β2 Microglobulin (β2m) was used as a positive control and for normalisation; β2m forward (5’-gaaatagcctcagcatctgtg-3’), β2m reverse (5’-agtgctcagcatctgctg-3’), and probe (5’-agcacagttcagcagcgacc-3’) were dual labelled with 6-carboxyfluorescein (FAM) and 6-carboxytetramethylrhodamine (TAMRA). PCR was performed in triplicate in 1 x qPCR master mix (Eurogentec, Liege, Belgium), using 900nM primers and 200nM probe. Reactions were performed on an ABI7900HT Sequence Detection System device (PE Applied Biosystems) using the standard program. Fluorescence was measured by means of sequence detection system software (SDS; version 2.0; Applied Biosystems). Mean cycle threshold values (Ct) and SDs were calculated for all genes. The amounts of the Bcl-2, Bcl-XL, and Mcl-1 targets were normalised relative to the amount of β2m target (ΔCt = ΔCt(gene) - ΔCt(β2m)) and the SD of ΔCt (SD(ΔCt)) was calculated (SD(ΔCt) = [(SD(gene))² + (SD(β2m))²]). The relative amount of target gene was measured by determining ΔΔCt (ΔΔCt = ΔCt(calibrator) − ΔCt(example)) and the factor difference was calculated (2^−ΔΔCt). The range is given as 2^−ΔΔCt±SD(ΔΔCt) and 2^−ΔΔCt±SD(ΔΔCt).

**Immunostaining**

ALK, Bcl-2, Bcl-XL, and Mcl-1 positivity was assessed by immunohistochemistry using standard protocols and appropriate dilutions of monoclonal mouse antihuman ALK antibody, monoclonal mouse antihuman Bcl-2 antibody, polyclonal rabbit antihuman Mcl-1 antibody (Dako, Copenhagen, Denmark), and monoclonal rabbit antihuman Bcl-XL antibody (Zymed, San Francisco, California, USA). Peroxidase activity was visualised with diaminobenzidine and H2O2. For all analyses, negative controls (first incubation step without primary antibody) and positive control tissue sections were included. Differences in the number of positive cases between the ALK+ and ALK− group were tested using Fisher’s exact test.

**RESULTS**

We constructed gene expression profiles of Karpas299 and CD4+ T cells using the SAGE technique. For Karpas299 we sequenced 10 678 tags representing 5090 different genes and for the CD4+ T cells we obtained 8425 tags representing 4467 different genes. These expression profiles were compared and

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**Figure 1** Real time reverse transcription polymerase chain reaction results for Bcl-2, Bcl-XL, and Mcl-1 in tonsil, CD4+ T cells, and the Karpas299, SUP-M2, SR786, SU-DHL-1, and DEL cell lines. The bars on a logarithmic scale indicate the relative amount of mRNA. It is clear that Mcl-1 is the main antiapoptotic protein expressed in anaplastic large cell lymphoma derived cell lines.

**Figure 2** Immunohistochemistry for Mcl-1 in anaplastic large cell lymphoma (ALCL). (A) Mcl-1 staining in the Karpas299 cell line. (B) Mcl-1 staining in an anaplastic lymphoma kinase negative (ALK−) ALCL (case 31). (C) ALK− ALCL (case 25) with tumour cells positive for Bcl-2. (D) ALK− ALCL (case 23) with tumour cells negative for Bcl-2. (E) ALK− ALCL (case 28) with tumour cells positive for Bcl-XL. (F) ALK− ALCL (case 8) with tumour cells negative for Bcl-XL. Original magnification, ×800.
linked to the Unigene library to identify the corresponding genes. The SAGE tag belonging to the Mcl-1 gene was detected at a frequency of 0.04% in Karpas299, and 0.01% in the CD4+ T cells. The tag for the shorter splice variant of Mcl-1 was not detected in the SAGE libraries. No SAGE tags were seen for the other antiapoptotic members of the Bcl-2 family (Bcl-2 and Bcl-XL) in Karpas299 or in CD4+ T cells. To confirm this differential expression, ALK+ ALCI cell lines Karpas299, SUP-M2, SR786, SU-DHL-1, and DEL were analysed using β2m, expressed at similar levels based on the SAGE libraries (0.29% in Karpas299 and 0.24% in CD4+ T cells), as a housekeeping gene. All five cell lines highly expressed Mcl-1 compared with tonsil and CD4+ T cells. Bcl-XL expression was moderate in the SUP-M2, SR786, SU-DHL-1, and DEL cell lines but was low in Karpas299 compared with CD4+ T cells (fig 1).

Immunocytochemical staining for Mcl-1 revealed strong cytoplasmic staining in most of the tumour cells in all cases, but only in three of 18 ALK+ cases (table 1; fig 2). The ALK+ group included significantly more Bcl-2 (p = 0.004) and Bcl-XL (p = 0.000) positive cases than the ALK− group.

Table 1

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- No staining in the tumour cells; +, moderate staining in most of the tumour cells; ++, strong staining in most tumour cells; NR, no result; few cells, staining in <10% of the tumour cells; part of cells, staining in 10–30% of the tumour cells.

ALCL, anaplastic large cell lymphoma; ALK, anaplastic lymphoma kinase.

DISCUSSION

Using SAGE analysis, the antiapoptotic member of the Bcl-2 family Mcl-1 was shown to be highly expressed in the ALCI derived cell line Karpas299 compared with CD4+ T cells. In contrast, tags corresponding to the other two members of the antiapoptotic Bcl-2 family, Bcl-XL and Bcl-2, were not identified in the SAGE libraries. Screening of the SAGE library for tags corresponding to proapoptotic genes revealed the presence only of a low frequency of tags corresponding to the Bak gene. The SAGE results were confirmed with quantitative RT-PCR, which indicated increased expression of the Mcl-1 gene in Karpas299 and in the four other ALCI derived cell lines. Moreover, quantitative RT-PCR for Bcl-XL and Bcl-2 clearly showed that expression was much lower than that of Mcl-1, consistent with the SAGE results.

Immunohistochemical analysis of 32 ALCI cases (23 ALK+ and nine ALK−) demonstrated the presence of the Mcl-1 protein in most tumour cells in all cases. No difference in Mcl-1 staining intensity or percentage of positive tumour cells was seen between ALK+ and ALK− cases. Mcl-1 protein expression was reported previously in 10 of 10 and 10 of 11 ALCIs.30 31 In a more recent study, Rasidakis et al detected Mcl-1 positivity in 16 of 26 ALK− ALCIs using a 10% cutoff.31 These data confirm the consistent high expression of Mcl-1 and strongly suggest that Mcl-1, rather than Bcl-2 or Bcl-XL, is the main antiapoptotic protein of the Bcl-2 family expressed in ALCI, and in particular in ALK+ ALCI. Analysis of the apoptotic rate in ALCI revealed low levels of apoptotic cells, ranging from 1.2% to 3.2%, in ALK+ and ALK− cases.32 33

A possible role for Mcl-1 in lymphomagenesis is supported by the finding of a variety of lymphomas in Mcl-1 transgenic mice.34 Moreover, high Mcl-1 expression has been reported in various other lymphoma subtypes, including angioimmunoblastic T cell lymphoma, mantle cell lymphoma, myeloma cell lines, cutaneous T cell lymphoma, diffuse large B cell lymphoma, and mantle cell lymphoma.35 36 37 38 Treatment with the cyclin dependent kinase inhibitor flavopiridol, which results in a strong downregulation of Mcl-1 expression, has been shown to be effective in multiple myeloma, leukaemia cell lines, and in chronic lymphocytic leukaemia samples,39 40 indicating the importance of Mcl-1 as an antiapoptotic protein.

“Our findings argue against a role for ALK induced activation of STAT3 and suggest involvement of other pathways leading to the induction of Bcl-XL and Bcl-2 expression”

The inhibition of JAK3 in two ALK+ ALCI derived cell lines resulted in downregulation of activated STAT3, decreased levels of Bcl-2 and Bcl-XL, and no changes in Mcl-1 values.31 This suggests that other mechanisms are involved in the induction of Mcl-1 in ALCI cases. However, in a more recent study Amin et al demonstrated the downregulation of Bcl-2, Bcl-XL, and Mcl-1 upon transfection of a dominant negative STAT3 in two ALCI cell lines, supporting the STAT3 mediated induction of Mcl-1 in these cells.32 Similar results were obtained in macrophages and large granular lymphocyte leukaemia, which showed reduced Mcl-1 expression upon treatment with JAK inhibitors.33 34 Our data indicate the presence of the Mcl-1 protein in all ALCI cases, independent of expression of the ALK protein, which suggests that besides the ALK induced activation of STAT3, other pathways might also contribute to the induction of Mcl-1 expression in ALCI cases that lack ALK expression. This is supported by two studies that show the involvement of the phosphatidylinositol 3-kinase pathway in the upregulation of Mcl-1 expression.35 36

In addition to positive regulation via survival signals such as activated STAT3 and the phosphatidylinositol 3-kinase pathway, Mcl-1 can also be downregulated via E2F1.36 Treatment with flavopiridol results in stabilisation of E2F1,
Mcl-1 expression in ALCL

Take home messages

- Mcl-1 was consistently highly expressed in ALK+ and ALK− anaplastic large cell lymphomas (ALCL), suggesting that Mcl-1 is the main antiapoptotic protein in this disease.
- The high frequency of Mcl-1, Bcl-2, and Bcl-XL positive ALCL cases in the ALK− group compared with the ALK+ group indicates that ALK induced STAT3 activation is not the main regulatory pathway in ALCL.
- Treatment with cyclin dependent kinase inhibitors such as flavopiridol may provide a novel tool for the treatment of both ALK+ and ALK− patients with ALCL.

which acts as a transcriptional repressor of Mcl-1, and induces an effective downregulation of Mcl-1. Based on the balance between survival pathways and the effectiveness of Mcl-1 downregulation upon flavopiridol treatment, beneficial effects might be achieved by treatment of ALCL with flavopiridol.

As mentioned above, several studies suggest a role for ALK induced activation of STAT3 in the induction of Bcl-2 and Bcl-XL expression in ALCL cases.31 17–20 Indeed, the presence of activated STAT3 was demonstrated in most ALK+ cases, but also in approximately half of the ALK− cases.31 These data indicate that STAT3 activation correlates with, but is not strictly dependent on, ALK expression in ALCL. The analysis of Bcl-2 and Bcl-XL in ALCL demonstrated the complete absence of Bcl-2 in ALK+ cases and the expression of Bcl-2 in four of nine ALK− cases; whereas Bcl-XL was expressed in only three of 18 ALK+ versus eight of nine ALK− cases. These data indicate that Bcl-2 and Bcl-XL are only infrequently expressed in ALK+ ALCL, but are present in a higher proportion of ALK− ALCLs. An inverse relation between ALK and Bcl-2/Bcl-XL expression was previously noted in ALCL.20 40–42 These findings argue against a role for ALK induced activation of STAT3 and suggest involvement of other pathways leading to the induction of Bcl-2 and Bcl-2 expression.

In summary, similar amounts of Mcl-1 protein are expressed in both ALK− and ALK+ ALCLs, whereas the expression of Bcl-2 and Bcl-XL is limited to ALK− cases. Because activated STAT3 is not detectable in a small proportion of ALK+ cases and in half of the ALK− cases, it is very likely that other regulatory pathways are involved in the expression of these antiapoptotic Bcl-2 family members. The consistent expression of Mcl-1 as demonstrated in our study indicates that treatment with agents such as flavopiridol may provide a novel tool for the treatment of both ALK+ and ALK− patients with ALCL.

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