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K Ha-Kawa, J Hara, Y Keiko, A Muraguchi, N Kawamura, S Ishihara, S Doi, H Yabuuchi

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Research Article

We describe a 10-yr-old boy with T-lineage non-Hodgkin's lymphoma. He had a mediastinal mass, swollen supraclavicular lymph nodes, and pleural effusion. A supraclavicular lymph node biopsy under light microscopy showed a malignant lymphoma of diffuse lymphoblastic type. Most of the cells taken from the malignant pleural effusion expressed T cell-associated antigens such as Leu-1 and OKT 8. To confirm these antigens as T-lineage lymphoma, we examined genomic DNA from malignant cells obtained from the pleural effusion. As was expected, T cell receptor beta-chain gene rearrangements were demonstrated. However, when the immunoglobulin gene organization was analyzed, we detected rearrangements in both the heavy- and kappa-chain genes. To our knowledge, this is the first case in which kappa-chain gene rearrangement was detected in apparent T-lineage cells. These findings provide important information relating to determination of the cellular lineage of lymphoid malignancy.

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Kappa-chain Gene Rearrangement in an Apparent T-lineage Lymphoma

Kyungsae Ha-Kawa,* Junichi Hara,* Yumura Keiko,* Atsushi Muraguchi,†
Nobuaki Kawamura,‡ Shigehiko Ishihara,* Satoru Doi,* and Hyakuji Yabuuchi*

*Department of Pediatrics, Osaka University Hospital; and †Institute for Molecular and Cellular Biology, Osaka University, Osaka, Japan

Abstract

We describe a 10-yr-old boy with T-lineage non-Hodgkin's lymphoma. He had a mediastinal mass, swollen supraclavicular lymph nodes, and pleural effusion. A supraclavicular lymph node biopsy under light microscopy showed a malignant lymphoma of diffuse lymphoblastic type. Most of the cells taken from the malignant pleural effusion expressed T cell-associated antigens such as Leu-1 and OKT 8. To confirm these antigens as T-lineage lymphoma, we examined genomic DNA from malignant cells obtained from the pleural effusion. As was expected, T cell receptor β -chain gene rearrangements were demonstrated. However, when the immunoglobulin gene organization was analyzed, we detected rearrangements in both the heavy- and kappa-chain genes. To our knowledge, this is the first case in which kappa-chain gene rearrangement was detected in apparent T-lineage cells. These findings provide important information relating to determination of the cellular lineage of lymphoid malignancy.

Introduction

Histologic and immunological evaluation of tissue biopsies and/or cell suspension have proven valuable in the diagnosis and classification of lymphoid neoplasms. More recently, detection of Ig and T cell receptor β -chain (T_β)¹ gene rearrangements has been shown to be an effective procedure for identifying monoclonality and the cellular lineage of lymphoid cells even when conventional studies give an ambiguous diagnosis (1-7).

After we reported the first case of T cell acute lymphoblastic leukemia (ALL) with C_μ gene rearrangement in newly diagnosed patients (8), analogous cases were reported. The incidence of this event among T-lineage leukemias can be estimated at ~20% (2, 9). Similarly, C_μ gene rearrangement was also observed in acute nonlymphocytic leukemia (10, 11). However, such Ig gene rearrangements in non-B-lineage cells reported so far are restricted to heavy-chain genes. For the most part, a unique pattern

of single allelic rearrangement has been observed (12). Based on these findings, it is clear that the mere demonstration of rearranged heavy-chain genes is not sufficient to establish a B-cell lineage.

In contrast, light-chain gene rearrangements have been thought to represent a more definitive assessment of B-lineage commitment because of the developmental hierarchy of Ig genes (13). Here, we present the first case of apparent T-lineage lymphoma, whose malignant cells showed heavy- and light-chain gene rearrangements in addition to T cell receptor β -chain gene rearrangements.

Methods

A 10-yr-old boy was presented with a large mediastinal mass, swollen supraclavicular lymph nodes, and pleural effusion. Biopsy specimens of a supraclavicular lymph node revealed non-Hodgkin's lymphoma of diffuse lymphoblastic type. Cells obtained through thoracentesis were also positive for the Papanicolaou's staining. No malignant cells were detected in the peripheral blood or bone marrow.

Immunologic phenotype. Immunologic studies were performed on the cells obtained from the pleural effusion. Mononuclear cells were prepared after Ficoll-Hypaque gradient centrifugation, and reactivity with a panel of monoclonal antibodies was carried out as previously described (8).

DNA analysis. Genomic DNA was extracted from the mononuclear cells and was digested with Eco RI or Bam HI restriction endonuclease, which are known to permit demonstration of both rearranged and germ-line configuration of Ig and T_β genes (1, 2, 5, 7). Digested DNA was size-fractionated by agarose gel electrophoresis and transferred to nitrocellulose filter (14). Such filter-bound DNA fragments were then hybridized to nick-translated ³²P-DNA probes of Ig genes and T_β genes (15). The human Ig gene probes used are the heavy-chain-joining gene (J_H) probe (3-kb embryonic Eco RI-Hind III J_H -containing fragment) and the κ probe (2.5-kb embryonic Eco RI kappa-chain gene (C_κ)-containing fragment). The human T_β gene probe was the Bgl II-Eco RV fragment of the cDNA clone YT-35 that contained the constant region of the T cell receptor gene (0.8-kb Bgl II-Eco RV fragment). J_H and C_κ germ-line clones were kindly provided by Dr. P. Leder (16, 17) (Department of Genetics, Harvard Medical School, Boston, MA). The cDNA clone YT-35 was kindly provided by Dr. T. Mak (18) (Department of Medical Biophysics, Ontario Cancer Institute, Toronto, Canada).

RNA analysis. Total cellular RNA was extracted from cells in the presence of guanidine thiocyanate (19). 10 μ g of each RNA sample was separated to size by electrophoresis in 1% agarose gel. The size-separated fragments were transferred to nitrocellulose filter by Northern techniques and hybridized to radiolabeled ³²P-DNA probes of the T_β and the C_κ described above.

Results

As shown in Table I, most of the patient's lymphoma cells expressed Leu-1 (64.8%) and OKT 8 (70.3%). A small fraction of

Address correspondence to Kyungsae Ha-Kawa, Department of Pediatrics, Osaka University Hospital, 1-1-50, Fukushima, Osaka, Japan 553.

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1. Abbreviations used in this paper: ALL, acute lymphoblastic leukemia; CALLA, common acute lymphoblastic leukemia antigen; C_κ , kappa-chain gene; Ia, I-region-associated antigen; J_H , heavy-chain-joining gene; T_β , T cell-receptor β -chain; T_β mRNA, T cell-receptor β -chain gene mRNA.

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Table I. Results of Immunologic Studies

	Surface marker	% Fluorescent cells
T-lineage	OKT 3	6.0
	4	33.6
	8	70.3
	11	35.6
	Leu-1	64.8
B-lineage	B 1	9.6
	B 4	15.7
	sIg	7.7
Myeloid	My 7	6.5
	Mo 2	12.3
Others	Ia	6.5
	CALLA	5.0

sIg, surface immunoglobulin; CALLA, common acute lymphoblastic leukemia antigen.

the cells reacted with OKT 4 and OKT 11. No other antigens, such as B-lineage and myeloid-lineage, were detected in most of the patient's cells.

The J_H probe recognized a rearrangement band with a faint band that corresponds to the germ-line configuration when genomic DNA was digested with Eco RI (Fig. 1). The C_μ probe (8) also detected two rearranged bands (data not shown). This J_H probe identified an 18-kb Eco RI fragment in the germ-line DNA. In parallel studies, seven cases of T-lineage leukemia showed the germ-line configuration of Ig genes, and 19 of 20 cases with common ALL (I-region-associated antigen [Ia⁺], common acute lymphoblastic leukemia [CALLA⁺]) demonstrated Ig gene rearrangements (20). When the C_μ probe was used after digestion with Bam HI, a single rearranged band was also found with deletion of the other allele (Fig. 1). This C_μ probe consistently detected a 12.5-kb Bam HI fragment for the germ-line k genes (2).

In the human germ-line T cell-receptor β -chain genome, there are two constant-region genes ($C_{\beta 1}$ and $C_{\beta 2}$) (18). The Bgl II-Eco RV fragment of the cDNA clone YT-35 (T_β probe) detected 11-kb ($C_{\beta 1}$) and 4-kb ($C_{\beta 2}$) Eco RI germ-line fragments, and one 24-kb Bam HI germ-line fragment containing both $C_{\beta 1}$ and $C_{\beta 2}$ (5, 21). After Eco RI digestion and hybridization with the T_β probe, the patient's DNA showed one rearrangement band with a germ-line configuration of the 4-kb fragment. This indicates rearrangement of one allele and deletion of the other $C_{\beta 1}$ allele, because there is an Eco RI restriction site just before $C_{\beta 2}$. The 4-kb band does not move, even in $C_{\beta 2}$ rearrangement (Fig. 1) (21). In addition, we detected two rearranged bands in the experiment using Bam HI digestion (Fig. 1). These findings indicate that both alleles of the T_β gene were used in the present case.

Fig. 2 shows the results of Northern blot analysis. Similar to RNA from a T cell line (Fig. 2, lane *HSB*), the RNA sample from the present case (lane *P*) contained a 1.3-kb T cell-receptor β -chain gene mRNA (T_β mRNA), which has been shown to be a functional mRNA transcript (22, 23). However, unlike T_β mRNA, no C_μ mRNA could be detected in this particular case. RNA from a B-cell line (lane *Jijoye*; sIg μ^+ , k⁺) demonstrated 1.0-kb C_κ mRNA. RNA from a non-T-lineage cell line (lane *U-937*) and a newly diagnosed common ALL case (lane *ALL*) were also examined as controls. This common ALL had rearrangements of the Ig heavy-chain and T_β genes, and the germ-line configuration of the Ig light-chain gene. As shown in Fig. 2, no transcripts of the T_β and C_μ genes were observed in either instance.

Discussion

Our finding of kappa gene rearrangement in an apparent T-lineage lymphoma provides new information relating to determination of the cellular lineage of lymphoid malignancies. Ig gene rearrangement is an essential property of cells of B-lineage, and detection of Ig gene rearrangements has been used to define the origin of immature cells and find monoclonality in mixed populations (1-4). Ig gene rearrangement is also observed in

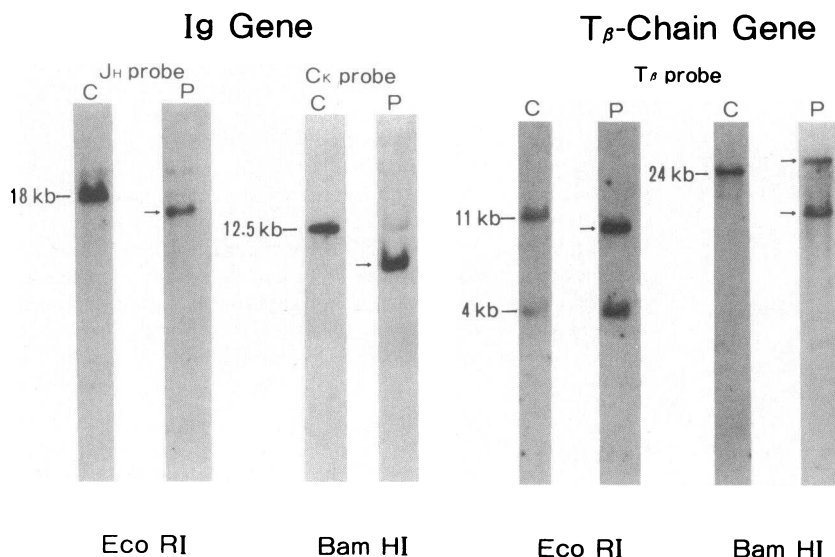


Figure 1. Southern blot analysis to detect germ-line and rearranged Ig genes and T_β -chain genes in control (C) and lymphoma cell DNA (P). The J_H probe, containing a 3-kb germ-line Eco RI-Hind III fragment, detected an 18-kb Eco RI fragment as germ-line (solid line) and rearranged allele (arrow). The C_κ probe, containing a 2.5-kb embryonic Eco RI C_κ -containing fragment, recognized a 12.5-kb Bam HI fragment for the germ-line genes (control, C, solid line). An arrow indicates rearranged allele. Similarly, the T_β probe, containing a 0.8-kb Bgl II-Eco RV fragment of the DNA clone YT-35, detected 11-kb and 4-kb Eco RI germ-line fragments and one 24-kb Bam HI germ-line fragment (solid) and rearranged alleles (arrow).

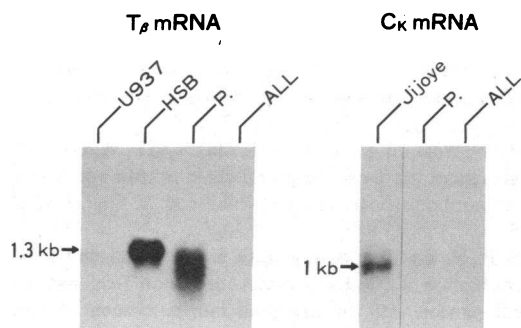


Figure 2. Northern blot analysis to detect transcript of T cell receptor β -chain genes and kappa-chain genes. RNA from a T cell line (lane HSB) and the present case (lane P) expressed a functional 1.3-kb T_{β} gene transcript. No T_{β} mRNA was detected in a non-T-lineage cell line (lane U937) and a common ALL case in which IgH and T_{β} genes were rearranged (lane ALL). Jijoye (B cell line, $slg\mu^{+}$, k^{+}) expressed a 1-kb mRNA of IgK genes, while no transcript was detected in lane P and lane ALL.

non-B-lineage cells, although such rearrangement is fairly restricted to the heavy-chain gene (12). Therefore, Ig gene rearrangement alone is not sufficient to establish B cell lineage. For assignment of a B-lineage commitment, the light-chain gene must undergo rearrangements on the basis of the developmental hierarchy of Ig genes (1, 13).

The β -chain of the T cell-receptor gene shares certain homologies with Ig genes, including rearrangement during T cell development (5–7, 18, 21). With the cloning of T_{β} genes, it has become possible to study the organization of T_{β} genes in individual cells and determine the cellular lineage of given cells. Rearrangements of T_{β} genes are consistently observed in T-lineage leukemia and lymphoma (5, 7, 21).

Tawa et al. reported that T_{β} gene rearrangements were also detected in 10 of 39 patients with common ALL (Ia^{+} , $CALLA^{+}$). The patterns of rearrangement, however, were somewhat unique compared with those of T-lineage leukemia. None of the 10 cases had rearranged bands of $C_{\beta}1$ genes, which seems to be more characteristic of T-lineage leukemia (21).

Using morphological and immunological studies, we diagnosed the present case as having T-lineage non-Hodgkin's lymphoma. Phenotypically, only T-lineage-associated markers such as Leu-1 and OKT 8 were detected on the cell surface. In addition, we found a characteristic rearrangement pattern of T_{β} genes in the Eco RI experiment. More definitively, we were able to show a functional 1.3-kb T_{β} gene mRNA in Northern blot analysis (Fig. 2).

Contrary to these findings, when the patient's DNA was further analyzed using the J_H and C_{κ} probes, both the heavy- and light-chain genes were found to be rearranged. The retention of a faint germ-line band in each experiment (Fig. 1) could have been due to contamination by normal cells.

To our knowledge, this represents the first example of detection of k-gene rearrangement in cells of apparent T-lineage. This finding is indeed different from the previous studies in non-B-lineage cells, where the rearrangement of Ig genes was restricted to the heavy-chain gene (8–12). In this particular case, we could not detect a real message of the rearranged k-genes.

The results presented here suggest that one should always

exercise caution when the cellular lineage is determined at the DNA level and is used to predict the prognosis and determine the therapy.

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